Contents

1 Introduction
   1.1 System overview ................................................................. 7
   1.2 Support for use in regulated environments ......................... 8
   1.3 Associated documentation ................................................. 8
   1.4 Biacore terminology ......................................................... 8

2 Control Software – general features
   2.1 Operational modes ............................................................. 13
   2.2 User interface ................................................................. 14
       2.2.1 Software help ............................................................. 15
   2.3 Basic operation ............................................................... 15
       2.3.1 Selecting cycles and sensorgrams ............................. 15
       2.3.2 File menu ................................................................. 15
       2.3.3 Edit menu ................................................................. 17
       2.3.4 View menu ............................................................... 17
       2.3.5 Run menu ................................................................. 19
       2.3.6 Tools menu ............................................................... 19
       2.3.7 Right-click menus ................................................... 20
   2.4 File storage ............................................................... 21
       2.4.1 Wizard templates and methods ................................ 21
       2.4.2 Result files .............................................................. 21

3 Manual run
   3.1 Preparing for a manual run ............................................... 23
       3.1.1 Instrument preparations ........................................... 23
   3.2 Starting a manual run ........................................................ 24
   3.3 Controlling a manual run .................................................. 25
   3.4 Ending a manual run ......................................................... 27

4 Application wizards
   4.1 Wizard templates ............................................................. 29
       4.1.1 Creating and editing wizard templates ...................... 29
       4.1.2 Running wizards ......................................................... 30
   4.2 Common wizard components ............................................. 30
       4.2.1 Injection sequence ......................................................... 30
       4.2.2 Assay setup ................................................................. 32
       4.2.3 Injection parameters ................................................... 33
       4.2.4 Sample and control sample tables ......................... 34
       4.2.5 System preparations .................................................... 35
       4.2.6 Rack positions ............................................................. 37
       4.2.7 Prepare Run protocol ................................................. 41
   4.3 Wizard groups ................................................................. 42
   4.4 Immobilization pH scouting ............................................. 42
4.5 Immobilization .................................................................45
4.6 Regeneration scouting .......................................................50
4.7 Buffer scouting .................................................................54
4.8 Surface performance ..........................................................56
4.9 Binding analysis .................................................................58
4.10 Concentration analysis .......................................................61
4.11 Kinetics/Affinity ...............................................................65
4.12 Thermodynamics ..............................................................70
4.13 Control experiments .........................................................73
  4.13.1 Mass transfer control ....................................................73
  4.13.2 Linked reactions control ................................................73
  4.13.3 Evaluation of control experiments ..............................74
4.14 Immunogenicity ..............................................................76

5 Methods
5.1 Opening methods .............................................................77
5.2 Method structure ..............................................................78
5.3 Method overview .............................................................79
5.4 General settings ...............................................................80
5.5 Assay steps ......................................................................81
  5.5.1 Base settings ...............................................................82
  5.5.2 Assay step preparations .................................................84
  5.5.3 Recurrence .................................................................84
  5.5.4 Number of replicates ....................................................85
5.6 Cycle types .....................................................................86
  5.6.1 Commands .................................................................87
  5.6.2 Variables .................................................................93
  5.6.3 Report points .............................................................95
5.7 Variable settings .............................................................97
5.8 Verification .................................................................98
5.9 Setup Run .................................................................98
  5.9.1 Detection ..............................................................98
  5.9.2 Variables ..............................................................98
  5.9.3 Cycle run list ..........................................................100
  5.9.4 System preparations .................................................100
  5.9.5 Rack positions ........................................................101
  5.9.6 Prepare Run Protocol ...............................................101
  5.9.7 Starting the run .......................................................101
5.10 Requirements for assay-specific evaluation .................101
  5.10.1 Concentration analysis .............................................101
  5.10.2 Kinetics/Affinity .......................................................102
  5.10.3 Thermodynamics ....................................................102
  5.10.4 Affinity in solution ..................................................102
  5.10.5 Immunogenicity .....................................................103
  5.10.6 Other requirements ..............................................103
6 Evaluation software – general features

6.1 User interface ........................................................................................................ 107
  6.1.1 Organization .................................................................................................. 107
  6.1.2 The Evaluation Explorer ........................................................................... 108

6.2 Basic operations .................................................................................................... 108
  6.2.1 Opening files ................................................................................................. 108
  6.2.2 Printing evaluation results ........................................................................... 109

6.3 Common display functions ................................................................................... 109
  6.3.1 Zooming the display .................................................................................... 109
  6.3.2 Right-click menus ....................................................................................... 109

6.4 Predefined evaluation items ................................................................................. 112
  6.4.1 Sensorgram .................................................................................................. 112
  6.4.2 Plots ............................................................................................................... 112

6.5 Custom report points .......................................................................................... 113
  6.5.1 Adding report points .................................................................................... 114
  6.5.2 Editing and deleting report points ................................................................. 115

6.6 Keywords .............................................................................................................. 115

6.7 Solvent correction ................................................................................................ 117
  6.7.1 Background ................................................................................................... 117
  6.7.2 When solvent correction should be used .................................................... 118
  6.7.3 How solvent correction works ...................................................................... 119
  6.7.4 Applying solvent correction ......................................................................... 119

6.8 Evaluation methods ............................................................................................... 121
  6.8.1 Creating evaluation methods ...................................................................... 121
  6.8.2 Applying evaluation methods ...................................................................... 122

7 Data presentation tools

7.1 Sensorgram items ................................................................................................. 123
  7.1.1 Selecting sensorgrams for display ............................................................... 124
  7.1.2 Removing data ............................................................................................. 124
  7.1.3 Sensorgram adjustment ................................................................................ 125
  7.1.4 Markers .......................................................................................................... 126

7.2 Plot items .............................................................................................................. 127
  7.2.1 Selector functions ......................................................................................... 128
  7.2.2 Table functions ............................................................................................. 129
  7.2.3 Sorting the plot ............................................................................................ 130
  7.2.4 Fitting curves to points ............................................................................... 130
  7.2.5 Adjusting plots for controls ......................................................................... 132
  7.2.6 Ranking ......................................................................................................... 134

7.3 Bar chart items ..................................................................................................... 135
  7.3.1 Selector functions ......................................................................................... 135
  7.3.2 Display options ............................................................................................. 136

7.4 Report point table ............................................................................................... 136
  7.4.1 Displaying the report point table .................................................................. 136
8 Concentration analysis

8.1 Requirements for concentration evaluation .......................... 141
  8.1.1 Calibrated measurements ............................................. 141
  8.1.2 Calibration-free measurements ....................................... 141

8.2 Evaluating calibrated concentration analyses .................... 142
  8.2.1 Calibration curves ..................................................... 142
  8.2.2 Calibration trends ..................................................... 144
  8.2.3 Control samples ....................................................... 146
  8.2.4 Samples ................................................................. 147
  8.2.5 Custom models for calibration curves ............................ 148
  8.2.6 Evaluating combined result sets .................................... 149

8.3 Evaluating calibration-free measurements ......................... 150
  8.3.1 Selecting samples ..................................................... 150
  8.3.2 Performing the evaluation .......................................... 155
  8.3.3 Interpreting the results .............................................. 156
  8.3.4 Fitting model ......................................................... 157

9 Kinetics and affinity analysis

9.1 Requirements for kinetics and affinity evaluation .................. 160

9.2 Evaluating kinetics and affinity in single mode ................. 161
  9.2.1 Basic procedure ..................................................... 161
  9.2.2 Multiple ligand densities ............................................. 169

9.3 Batch mode evaluation .................................................. 170

9.4 Quality assessment for kinetics evaluation ........................ 172
  9.4.1 The Quality Control tab .............................................. 172
  9.4.2 Statistical parameters .............................................. 175
  9.4.3 Components of the fit .............................................. 177
  9.4.4 Check kinetic data ................................................... 177

9.5 Quality assessment for affinity evaluation ........................ 179

9.6 Summarizing kinetics and affinity results ......................... 180
  9.6.1 Creating kinetic summaries ....................................... 180
  9.6.2 Basic summary presentation ....................................... 180
  9.6.3 On-off rate maps ..................................................... 183

9.7 Curve fitting principles .................................................. 184
  9.7.1 Fitting procedure ..................................................... 184
  9.7.2 Local and global parameters ....................................... 185

9.8 Predefined models ....................................................... 186
  9.8.1 Kinetics – 1:1 binding .............................................. 187
  9.8.2 Kinetics – Bivalent Analyte ....................................... 188
  9.8.3 Kinetics – Heterogeneous Analyte ............................... 189
  9.8.4 Kinetics – Heterogeneous Ligand ................................ 191
  9.8.5 Kinetics – Two State Reaction ................................... 193
  9.8.6 Affinity – Steady State 1:1 ....................................... 195

9.9 Creating and editing models ........................................... 195
  9.9.1 Interaction models for kinetics ................................... 196
  9.9.2 Equation models for kinetics .................................... 200
  9.9.3 Models for steady state affinity .................................. 202
10 Thermodynamic analysis
10.1 Background ................................................................. 203
10.1.1 Equilibrium thermodynamics ................................. 203
10.1.2 Transition state thermodynamics ............................. 204
10.2 Performing thermodynamic analysis .............................. 205

11 Affinity in solution
11.1 Conventions and background ....................................... 209
11.1.1 Experimental setup .................................................. 209
11.1.2 Evaluation principles ................................................ 209
11.2 Requirements for affinity in solution .............................. 210
11.3 Evaluation of affinity in solution .................................... 211

12 Immunogenicity

Appendix A Data import and export
A.1 Exporting data ............................................................... 217
  A.1.1 Export functions ..................................................... 217
A.2 Importing data ............................................................... 218
  A.2.1 Control Software ................................................... 218
  A.2.2 Evaluation Software ................................................ 221

Appendix B Method examples and recommendations
B.1 Affinity in solution .......................................................... 223
B.2 Calibration-free concentration analysis ............................ 224
  B.2.1 Assay steps and general settings ............................... 224
  B.2.2 Cycle types ........................................................... 224
  B.2.3 Variable settings ..................................................... 225
  B.2.4 Setup Run .............................................................. 226
B.3 CAP single-cycle kinetics .............................................. 227
  B.3.1 Assay steps ............................................................ 227
  B.3.2 Sample analysis cycle for Sensor Chip CAP ............... 228
B.4 GST kinetics ................................................................. 230
B.5 Inject and Recover ........................................................ 232
B.6 Kinetics heterogeneous analyte ...................................... 234
B.7 L1 liposome capture ..................................................... 234
B.8 LMW kinetics and LMW Screen ................................... 235
B.9 LMW single-cycle kinetics ........................................... 237
B.10 Single-cycle kinetics .................................................. 237

Index .................................................................................. 239
1 Introduction

Biacore™ T200 is a high performance system for analysis of biomolecular interactions, based on GE Healthcare’s surface plasmon resonance (SPR) technology. The Control Software supplied with the system offers easy-to-use wizards for assay development and common applications together with flexible facilities for designing custom analysis methods using a graphical interface called Method Builder. Results are evaluated in separate Evaluation Software designed for efficient and flexible evaluation, with dedicated functions for common applications.

This Handbook describes in detail how to use the Control and Evaluation Software.

1.1 System overview

Instrumentation in the Biacore T200 system is described in full in the Biacore T200 Instrument Handbook. Important features relevant to software operation include:

- Biacore T200 supports simultaneous analysis in up to four flow cells connected in series. The flow cells are arranged in pairs (Fc1-2 and Fc3-4) with minimum dead volume between the flow cells in a pair to provide accurate reference subtraction.

- The sample compartment accommodates one microplate (96- or 384-well, regular or deep-well capacity) and one reagent rack for reagent vials. A combined sample and reagent rack can be used in place of the separate microplate and reagent rack.

- Material that binds to the sensor surface during sample injection can be recovered in a small volume of liquid for further analysis by e.g. mass spectrometry.

- The temperature in the sample compartment is controlled separately from the analysis temperature, allowing samples to be kept at one temperature while analysis is performed at another. Samples equilibrate to the analysis temperature during injection into the flow cell. The analysis temperature can be varied during a run, and the sample compartment temperature can be set to follow the analysis temperature if desired.

- The system includes a buffer selector valve, allowing analysis to be performed in up to four different buffers in the same unattended run.
1 Introduction

1.2 Support for use in regulated environments

Support for use in regulated (GxP) environments is provided in an optional package that adds appropriate functionality to the Biacore T200 software. Functions for GxP support are described in a separate Biacore T200 GxP Handbook. Descriptions of software in the current Handbook apply to installations both with and without the GxP package unless otherwise stated.

1.3 Associated documentation

This Handbook describes Biacore T200 Control Software and Evaluation Software, version 1.0. Any functionality that is added in optional add-on modules is described in separate documentation.

Biacore T200 Instrument Handbook describes the instrumentation in the Biacore T200 system, with instructions for operation, maintenance and troubleshooting.

Biacore T200 GxP Handbook describes functionality added with the optional GxP package, together with some recommendations for using the system in a regulated environment.

Biacore T200 Immunogenicity Handbook describes the use of specialized functions in the software for immunogenicity studies.

Other general handbooks and documentation describing the technology are available from GE Healthcare. Information may also be found on the Internet at www.gelifesciences.com/biacore.

1.4 Biacore terminology

Biacore monitors the interaction between two molecules, of which one is attached to the sensor surface and the other is free in solution. The following terms are used in the context of work with Biacore systems (see Figure 1-1):

- The partner attached to the surface is called the ligand. Attachment may be covalent or through high affinity binding to another molecule which is in turn covalently attached to the surface. In the latter case the molecule attached to the surface is referred to as the capturing molecule.

  Note: The term “ligand” is applied here in analogy with terminology used in affinity chromatography contexts, and does not imply that the surface-attached molecule is a ligand for a cellular receptor.

- The analyte is the interacting partner in solution for which the concentration is to be measured. In direct binding assays, the analyte

---

1 GxP is used as a generic abbreviation for GLP (Good Laboratory Practice), GMP (Good Manufacturing Practice) and GCP (Good Clinical Practice).
binds directly to the ligand. In inhibition assays, the concentration of analyte is measured indirectly through binding of an additional molecule.

![Figure 1-1. Ligand, analyte and capturing molecule in relation to the sensor surface.](image)

- **Regeneration** is the process of removing bound analyte from the surface after an analysis cycle without damaging the ligand, in preparation for a new cycle.
- Response is measured in *resonance units* (RU). The response is directly proportional to the concentration of biomolecules on the surface.
- A *sensorgram* is a plot of response against time (see Figure 1-2), showing the progress of the interaction. This curve is displayed directly on the computer screen during the course of an analysis. Sensorgrams may be analyzed to provide information on the rates of the interaction.
- In many assay situations, sample passes over two or more flow cells in series, where one flow cell (usually the first) serves as a reference while ligand is attached in the other flow cell(s). Surfaces with ligand are referred to as **active**: blank surfaces used for reference purposes are **reference**.
- A particular sensorgram is referred to as a *curve* in several contexts in the software. This terminology is used to distinguish between different classes of sensorgram that recur within a run: for example, measurements on one active and one reference surface can generate separate curves for each of the two flow cells and a third *reference-subtracted* curve (active minus reference).
- A *report point* records the response on a sensorgram at a specific time averaged over a short time window, as well as the slope of the sensorgram over the window. The response may be absolute (above a fixed zero level determined by the detector) or relative to the response at another specified report point.
1 Introduction
1.4 Biacore terminology

Figure 1-2. Schematic illustration of a sensorgram. The bars below the sensorgram curve indicate the solutions that pass over the sensor surface.
Control Software
2 Control Software – general features

2.1 Operational modes

Biacore T200 Control Software offers three modes of operation:

- **Manual run** provides interactive control of the instrument operation, executing commands singly as they are issued. This mode is most useful for *ad hoc* experiments involving one or a few injections, such as testing the response obtained from injection of a single sample.

- **Application wizards** provide guidance in setting up experiments for assay development and execution. Separate wizards are offered for different purposes such as ligand immobilization, concentration determination or measurement of kinetic constants. Each wizard consists of an ordered series of dialog boxes, ensuring that the essential features of the application setup are correctly defined.

- **Methods** provide greater flexibility (and conversely less guidance) in setting up applications, allowing customized applications that are not covered by wizards. Methods are defined in a graphical interface called Method Builder, which is designed to provide full flexibility in method definition while retaining a simple interface for running assays based on established methods. Application wizard templates may be opened in Method Builder to provide a starting point for further refinement of application setup. Predefined methods are also provided as help in defining methods for selected purposes (see Appendix B).

Each of these modes of operation is described in more detail in the following chapters.
2 Control Software – general features
2.2 User interface

The main screen in the control software is divided into the following areas:

- **Menu and toolbar** provide access to control commands.
- **Event log** records settings at the start of the run and instrument control events during the run. The event log is displayed in a separate window, opened by clicking on the **Event Log** button at the right of the toolbar.
- **Sensorgram window** displays the sensorgrams for the current run or the currently open file.
- **Report point table** lists report points for the currently displayed cycle. Report points record the response at a set time and are defined automatically; custom report points can also be added in methods, or after the run in either the Control Software or the Evaluation Software.
- **Keyword table** lists keywords for the currently displayed cycle. Keywords are defined automatically in wizard runs, or in the method for method runs.
- **Status bar** displays the instrument status, including the temperature of the detector and the sample compartment. The content of the status bar varies between different situations: for wizard- and method-based runs, the elapsed run time and the estimated total run time are included.
2.2.1  Software help

Software help is available at any time from the Help menu. Context-specific help for dialog boxes is provided through Help buttons in the boxes.

2.3  Basic operation

2.3.1  Selecting cycles and sensorgrams

During a run, the current cycle is displayed by default. You can choose which cycle to display in the Cycle selector, but the display will revert to the current cycle when a new cycle is started. For a completed run, choose which cycle to display with the Cycle selector in the toolbar:

The Curve selector determines which curve in the cycle is current in the display. Options in the View menu (Section 2.3.4) control which curves are displayed in the sensorgram window.

2.3.2  File menu

The Open/New options for wizard templates and methods create new wizard templates and methods, and open existing templates and methods for editing or for starting a run.

Open opens result files. Most result files just display the sensorgrams and tables. Files from immobilization and regeneration scouting wizards also display a summary window showing the results of the run (see Sections 4.5 and 4.6).

Save and Save As save the results as a Biacore results file (extension .blr).

Export exports the current results to a file in Microsoft Excel or XML format, or exports the contents of the report point table to a tab-separated file. See Appendix A for details of the export format.
Print prints a hard-copy of the results. Select the printer to use and check the items you wish to print.

Sensorograms may be printed as follows:

- **None**
  - No sensorgrams will be printed.

- **Current cycle**
  - The current cycle will be printed with the View:Show... setting and scale as shown on the screen.

- **Range**
  - Multiple cycles will be printed. For Range, enter a range or cycle numbers separated by commas (e.g. 4-16,19,22). All curves will be included in each cycle regardless of the View:Show... setting. Sensorograms will be printed at full scale unless the Lock Scale box is checked in the sensorgram window, in which case the current scaling will be applied to all cycles (with this setting, some sensorgrams may appear to be empty).

- **All cycles**
  - All curves will be included in each cycle regardless of the View:Show... setting. Sensorgrams will be printed at full scale unless the Lock Scale box is checked in the sensorgram window, in which case the current scaling will be applied to all cycles (with this setting, some sensorgrams may appear to be empty).

**Note:** In order to maintain report layout, the print orientation is fixed regardless of the printer settings in Windows.

**Properties** shows detailed properties of the currently opened run, including the properties of the sensor chip used in the run.

When you close the software with Exit while the instrument is still switched on, you may choose to shut down the instrument for a shorter or longer period if required. See the Biacore T200 Instrument Handbook or the on-line help for more details.
2.3.3 Edit menu

Options in the Edit menu allow you to add, edit and delete report points. Report points are created automatically and are used in various evaluation contexts. You should in general avoid editing or deleting report points that are created automatically.

Editing operations for report points in the Control Software may be applied to single report point instances or to all instances of the report point in the current cycle. Note that editing operations are not applied to multiple cycles.

Report points created in the Control Software cannot be edited in the Evaluation Software. The Evaluation Software offers functions for creating and editing custom report points that can be applied to all cycles in the run in a single operation. This is usually preferable to adding report points in the Control Software.

2.3.4 View menu

Chip Properties opens a dialog box that displays the properties of the currently docked sensor chip. The Ligand column is empty for flow cells that have not been used, and shows [Blank] for flow cells that have been prepared as a blank reference surface by activation and deactivation. The text [Incomplete results] indicates that the immobilization run was interrupted (by for instance user intervention or power failure) before it could be completed.
Properties for the sensor chip used in a currently open run may be found under **File:Properties** (Section 2.3.2).

**Title** sets a title in the sensorgram window. The default title is the assay step name.

**Scale** sets the scale of the sensorgram window:

If you set **Auto scale**, the scale will be adjusted if necessary to accommodate the full data range of the currently displayed cycle. During a run, the scale is adjusted at intervals as more data is collected. Check the **Lock scale** box in the top right corner of the sensorgram window to lock the scale to the current settings.

**Adjust Scale** sets the scale to the full data range. This will not affect the **Auto scale** setting in the **Scale** dialog. **Adjust Scale** overrides but does not turn off the **Lock scale** setting.

To scale the sensorgram display interactively, drag with the cursor over the area to be scaled. Double-clicking in the display or choosing **View:Unzoom** restores the previous zoom setting.

**Reference line** toggles display of a movable vertical line in the sensorgram window, together with a separate small window that shows the response and time coordinates at the reference line for the current curve. Use the **Curve** selector in the toolbar (see Section 2.3.1) to set the current curve. Drag the reference line to move it. When the reference line is displayed, choosing **Baseline** sets a baseline at the current reference line position, and the coordinates window shows the response relative to that baseline.
The options **Show Only Current Curve**, **Show Curves of Same Type** and **Show All Curves** control which curves are displayed in the sensorgram window. Curve types distinguish between unsubtracted and reference-subtracted curves.

Choose the **Event Log** option or click on the **Event Log** button at the right of the toolbar to display the event log window.

Choose the **Wizard Template** or **Method** options to display the wizard or method definition for the run. You can edit the definition and save it as a new wizard template or method. You cannot however change the original definition that is saved together with the result file.

**Notebook** opens a notebook window where details of the run may be recorded. The notebook is only available during a run or for a completed result file. The run notebook is saved with the result file and can be viewed in the Evaluation Software.

For some wizard runs and for test tools, the **Wizard Results** option opens a window showing the results of the run. All other runs are evaluated in the Evaluation Software.

**Sensorgram Markers** controls display of report point and event markers and labels in the sensorgram window.

### 2.3.5 Run menu

The options in the **Run** menu are used to start the different types of runs (see Chapters 3–5).

### 2.3.6 Tools menu

Options in the **Tools** menu control instrument operations outside the context of runs.

**Prime** flushes the flow system with fresh buffer. There is an option to include **Prime** at the beginning of each wizard- or method-based run. Use the menu option when you want to flush the system at other times (e.g. before a manual run).

**Shutdown** starts the procedure for shutting down the instrument for long periods of time (more than 7 days). The procedure displays necessary instructions on the screen. Details of the shutdown procedure are given in the Biacore T200 Instrument Handbook.

**Standby** puts the instrument in standby mode, which maintains a low buffer or water flow through the flow system for up to 7 days. Leaving the instrument in standby mode when not in use is generally recommended. The instrument is
automatically put in standby mode at the end of a run. Use the menu option if standby has been stopped and you want to restart it.

**Stop Standby** stops standby mode.

**Eject Rack** ejects the rack tray from the sample compartment. The rack may be ejected during setup for wizard- and method-based runs, and at any time during a manual run. Use the menu option or the toolbar button when you want to eject the rack at any other time.

---

**CAUTION**

The rack tray automatically moves into the instrument a preset time after it has been ejected. The time to auto-close is set in **Tools:Preferences**. A timer in the dialog indicates when the rack tray will be automatically moved into the instrument.

---

**Rack Illumination** switches the illumination in the sample compartment on or off. The illumination helps you to see in the sample compartment but does not otherwise affect instrument function.

**Insert Chip** and **Eject Chip** are used for docking and undocking the sensor chip respectively. More details are given in Chapter 4 of the Biacore T200 Instrument Handbook.

**Set Temperature** sets the sample compartment and analysis temperature. More details are given in Chapter 4 of the Biacore T200 Instrument Handbook.

**Preferences** controls aspects of file storage and data import (see Section 2.4), and the time to automatic retraction of the rack tray.

**More Tools** provides access to maintenance, test and service tools. Details are given in Appendix B of the Biacore T200 Instrument Handbook.

---

2.3.7 **Right-click menus**

Right-clicking with the mouse in many windows opens context menus specific for the window.

**Sensorgram window**

**Scale** opens the same dialog as the **View:Scale** option (Section 2.3.4).

**Copy Graph** copies the sensorgram window exactly as displayed to the Windows clipboard. Use this option to insert a copy of the sensorgram window into other programs such as presentation software.

**Export Curves** exports data for the currently displayed curves to a text file. Entire curves are exported regardless of the scale of the display. The exported data includes report points and event marker times if these are displayed in the sensorgram window. See Appendix A for more details of the export format.
Gridlines controls display of gridlines in the sensorgram window.

**Report point table**
The right-click menu options for the report point table correspond to the **Edit:Report Points** menu options.

**Notebook**
Right-click menu options in the notebook represent standard Windows editing functions.

## 2.4 File storage

### 2.4.1 Wizard templates and methods
Wizard templates are saved in files with a file name extension `.bw**`, where `**` represents an abbreviation that identifies the wizard (e.g., a wizard template for concentration analysis has the extension `.bwConc`).

Methods are saved in files with the file name extension `.Method`.

**Note:** The extension will not be displayed if the setting **Hide file extensions for known file types** is selected in the Windows Explorer folder options. Turning this setting off can help you to identify file types in dialog boxes.

Templates and methods may be saved in any location when the optional Biacore T200 GxP Package is not installed. A folder structure under the default location as specified on the **Folders** tab in **Tools:Preferences** is however recommended, since files in this location are handled preferentially in the **Open/New** dialog boxes for wizards and templates (see Section 4.1.1).

### 2.4.2 Result files
Results are saved in files with the file name extension `.blr`. Result files from wizard- or method-based runs contain a copy of the wizard template or method as well as the results of the run.
2 Control Software – general features
2.4 File storage
3 Manual run

Manual run allows you to control a run interactively. All settings except temperature and choice of microplate and/or reagent rack can be changed during the run. Commands are placed in a queue if the instrument is busy when a command is issued: queued commands that have not yet been started can be edited or deleted from the queue.

The results of a manual run are saved in a normal result file, and can be evaluated in the Evaluation Software. There are however no predefined keywords associated with the run, and the results cannot be evaluated with the Evaluation Software tools for concentration, kinetics/affinity, thermodynamics or immunogenicity.

3.1 Preparing for a manual run

3.1.1 Instrument preparations

The integrated instrument preparation steps that are included with wizard- and method-based runs are not supported for manual run. The instrument should therefore be prepared using options from the Tools menu.

1 Dock the chip that you want to use, and immobilize ligand on the surface (see Section 4.5) if this has not already been done.

2 Choose Tools:Prime to flush the flow system with fresh buffer.

3 Choose Normalize from the Maintenance Tools section of Tools:More Tools if the detector has not been normalized since the chip was docked. (In many cases, the detector will have been normalized in connection with ligand immobilization. However, you may need to perform the operation again if the chip has been undocked and re-docked after immobilization.)

4 Choose Tools:Set Temperature and set the analysis and sample compartment temperatures. Wait until the analysis temperature is stable (as shown in the status bar) before starting the run.

5 Prepare your samples and reagents in the microplate and/or reagent rack. Note the rack positions and volumes of samples that you prepare: there is no software support in manual run for identifying samples or monitoring the volume of liquid in the autosampler positions. You insert the samples as part of the starting procedure for the run. You can also add samples during the run.
3.2 Starting a manual run

Choose Run: Manual Run to start a manual run.

Choose the initial settings for flow rate, flow path and reference subtraction. You can change the flow rate at any time during the run. You can change the flow path at any time: during a cycle, the available options are restricted by the choice made when the cycle is started.

Choose the rack and microplate settings. These will apply throughout the run and cannot be changed.

Click Eject Rack to eject the rack tray so that you can load your samples.

Click Start to start the run. You will be asked to specify a result file name before the run actually starts.
3.3 Controlling a manual run

Control the manual run from the command buttons in the main window or the options in the Command menu:

Commands are executed immediately if the instrument is idle. With a few exceptions (noted in the detailed descriptions below), commands issued when the instrument is busy are placed at the end of a queue. The queue is listed in the left-hand panel, with commands that have been executed in gray text and those that are pending in black text. The command currently being executed is marked with a “working” icon.

Right-click on a pending command for a menu with options for:

- editing the command
- inserting a new command before the selected command (you choose the command to insert from a dialog box)
- deleting the command

You can also use the right-click menu to copy selected command or commands and paste them elsewhere in the queue. The Copy function works with both completed and pending commands.
3 Manual run
3.3 Controlling a manual run

**Flow rate**
Sets the flow rate to a new value.

**Flow path**
Changes the flow path. During a cycle, you can only select a flow path within a range allowed by the setting chosen when the cycle was started (for example, if the current setting is *Flow path 1-2*, you cannot extend it to *Flow path 1-2-3-4*).

**Sample injection**
Injects sample. Choose the position from which the sample will be taken and specify a contact time. Positions that can be chosen are determined by the rack settings in the manual run start-up dialog. Make sure that the chosen position contains enough sample for the injection. The required volume for the specified contact time is indicated in the dialog box.

**Regeneration injection**
Injects regeneration solution. Choose the position from which the solution will be taken and specify a contact time. Positions that can be chosen are determined by the rack settings in the manual run start-up dialog. Make sure that the chosen position contains enough solution for the injection. The required volume for the specified contact time is indicated in the dialog box.

Check *High viscosity solution* if your regeneration solution has a relative viscosity higher than about 3 (corresponding to about 35% glycerol or 40% ethylene glycol at 20°C). This will adjust the injection procedure to ensure correct handling of viscous solutions, and will limit the maximum contact time that can be specified.

**Wait**
Inserts a *Wait* command in the queue, causing the instrument operation to pause for the specified time period. Buffer continues to flow over the sensor surface during the *Wait* period and data collection continues.

**Eject Rack Tray**
Ejects the rack tray so that you can load more samples. Do not change the type of microplate or reagent rack on the tray.

This command is inserted immediately after the command currently under execution, rather than at the end of the queue, so that the rack tray will be ejected as soon as the current command is completed. If you want to place the command later in the queue, use the right-click menu in the queue panel to insert the command at the appropriate place.
New Cycle
Starts a new cycle. You can choose a new flow path and reference subtraction setting for the new cycle, independently of the setting in the current cycle.

Stop <command>
Stops the command currently being executed. The icon changes to show the command that will be stopped, or is gray if the current command cannot be stopped (e.g. it is not possible to stop an Eject Rack Tray command).

Stop Run
Finishes the run.

Pause Run
Pauses the run until a Resume Run command is issued. Buffer continues to flow over the sensor surface while the run is paused. Data collection continues during the pause.

Resume Run
Resumes a run that is paused.

Add report point
Adds a report point to the sensorgram.

Help
Displays help for the manual run.

3.4 Ending a manual run
To end a manual run:

1. Issue a Stop Run command. The command will normally be placed at the end of the queue. If you want to stop the run before the queue is completed, use the right-click menu in the queue panel to delete commands from the queue or to insert the Stop Run command in the appropriate position.

2. Choose Tools:Eject Rack to eject the rack tray and remove your samples and reagents.

3. Choose Tools:Eject Chip to undock the chip if desired.
3 Manual run
3.4 Ending a manual run
4 Application wizards

Application wizards guide you through the procedure of setting up common applications, with recommendations and settings based on GE Healthcare’s expertise in the field of SPR-based interaction studies. Wizards are an ideal starting point for inexperienced or infrequent users, since they offer a structured sequence of settings that covers all essential aspects of the assay in question. Wizard settings can be saved in templates for later use. Advanced users can open wizard templates in Method Builder for more flexible assay design (see Chapter 5).

4.1 Wizard templates

An application wizard consists of a series of dialog boxes that takes you through the steps in setting up the application. Settings in the dialog boxes may be saved in wizard templates, so that opening a template will present the saved settings in each dialog box.

Normally, a wizard template is saved when all steps have been defined, so that the template represents a complete assay definition including sample details if desired. If a wizard sequence is closed before reaching the last step, however, you are given an opportunity to save the template, which will then contain settings as far as they have been defined.

4.1.1 Creating and editing wizard templates

To create a new wizard template or edit an existing template, choose File: Open/New Wizard Template and select the type of wizard in the dialog box. Click New to create a new template, or navigate to the folder where your template is stored, select the template and click Open to edit an existing template.

The top-level folder for wizard templates is defined under Tools: Preferences (see Section 2.4). You can navigate between subfolders under the top level in the dialog box, but you cannot access templates outside the top-level folder directly from within the dialog box. Click Browse to navigate freely in the computer file structure and open wizard templates stored in other locations.

Note: The Open/New Wizard Template dialog box only lists templates of the selected type, but the Browse dialog may list all types. Template types are identified by the file extension, which may or may not be displayed according to your Windows Explorer settings (see Section 2.4.1).
4 Application wizards
4.2 Common wizard components

4.1.2 Running wizards
When you start a run based on a wizard template, the template settings are displayed as you step through the wizard. You can change settings for the particular run if desired: the changes are not saved in the wizard template unless you explicitly request this with the Save wizard template option. Settings used for the run are stored in the result file and can be examined and saved as a new wizard template from the completed run.

4.2 Common wizard components
Several dialogs are common to a number of wizards, with equivalent functions and only minor differences. These dialogs are described in the current section. Any wizard-specific variations in these common components are described in the sections on the respective wizards below.

4.2.1 Injection sequence
This dialog determines the sequence of injections in the wizard analysis cycle. Some injections are not supported in certain wizards (e.g. the kinetics wizard does not support enhancement injections).

Detection settings
Select the flow path for the analysis. The setting will apply throughout the whole wizard run. The available flow paths vary between the different wizards according to wizard purpose.
The detection is automatically set to the same settings as the flow path, so that sensorgrams are recorded only from the flow cells used.

When reference subtraction is used together with ligand capture (Section 4.2.1), the captured ligand passes over the active surface but not the reference surface (for example, with Flow path set to 2-1, the ligand is injected in flow cell 2 but not flow cell 1). If the Flow path setting does not use reference subtraction, ligand is injected in all flow cells included in the flow path.

**Chip type**

Select the sensor chip type for the analysis. This choice will determine certain assay settings in accordance with the requirements of the sensor chip (for example, selecting Sensor Chip NTA will automatically check the Ligand capture option and include an injection of nickel before the ligand capture step, and will suggest 0.35 M EDTA as a conditioning solution).

Choose chip type **Custom** if you are using a chip type that is not listed.

**Injections**

Check the injections that you want to include. The illustration panel shows the sequence of included injections. Injections have the purposes listed below. There may be additional injections for some sensor chip types (for example injection of nickel for Sensor Chip NTA).

**Ligand capture** Intended for ligand solution in applications that use a capturing approach to attach the ligand to the surface. The same solution will be used for the capture injection in all cycles: you cannot vary the captured ligand within the context of one wizard run.

The flow path for capture solution depends on the settings for detection (see above).

**Sample** This is the sample to be analyzed. The solution used for the sample injection is normally different in different cycles, and is specified in the sample table at a later stage in the wizard. The sample injection is required in all wizards.

**Enhancement** Intended for injection of a secondary reagent that binds to analyte on the surface, typically used either to amplify the response obtained from the analyte or to enhance the specificity of analyte detection. The same solution will be used for the enhancement injection in all cycles in the run.

**Regeneration** One or two regeneration injections may be included, which may use the same or different solutions. The same regeneration procedure is used throughout the run.
4 Application wizards
4.2 Common wizard components

4.2.2 Assay setup

Common features of the assay setup dialog are choice of conditioning and start-up cycles at the beginning of the run.

Conditioning cycle
A conditioning cycle prepares the sensor chip for the assay by washing with one or more injections of the specified solution. The surface is not regenerated after the conditioning injections. The conditioning cycle is run once at the beginning of the assay.

Conditioning cycles are recommended for certain chip types, to prepare the surface before starting the assay. Examples are Sensor Chip NTA which should be conditioned with 0.35 M EDTA to remove any bivalent metal ions, Sensor Chip L1 and HPA which may be conditioned with for example octylglucoside to remove any lipids on the surface, and Sensor Chip CAP which should be conditioned with regeneration solution. Conditioning cycles are generally not appropriate for sensor chips where the ligand or capturing molecule is attached to the surface before the assay wizard is started.

If you run several assays after each other without undocking the chip between assays, conditioning is generally only required for the first assay.

Start-up cycles
Start-up cycles are identical to analysis cycles except that the sample is replaced by a dummy sample (often buffer). The response from a newly prepared or newly docked sensor chip often shows some instability during the first few cycles, and start-up cycles allow the response to stabilize before the first analysis cycle is performed. Three start-up cycles are generally recommended for most assay purposes, to ensure a stable response in the analysis. Start-up cycles are treated separately from analysis cycles in the evaluation software.
Start-up cycles are run at the beginning of the experiment (after the conditioning cycle), and also directly after buffer change (in the *Buffer Scouting* wizard) and temperature change (in the *Thermodynamics* wizard).

### 4.2.3 Injection parameters

The *Injection parameters* dialog specifies details of injections selected in the *Injection sequence*. Injections for which the conditions are fixed in the software are not listed.

![Injection parameters dialog](image)

Details of this dialog box may vary according to the injections selected and the particular wizard. Some features may be generalized:

**Parameter limits**

Flow rates can be set between 1 and 100 µl/min in increments of 1 µl/min.

Permitted ranges for injection contact times are determined by the flow rate together with the limits for injected volumes, which are 2–350 µl for normal solutions and 5-100 µl for viscous regeneration solutions (see below).

**Note:** *The injected volume of solution is determined by the combination of flow rate and contact time, rounded to the nearest whole number. At low flow rates, this can result in actual contact times that differ from the requested times: for example, at 1 µl/min a requested contact time of 200 s (requiring 3.3 µl solution) will result in an actual contact time of 180 s (solution volume rounded to 3 µl).*
4 Application wizards
4.2 Common wizard components

Stabilization time after injection
This function is available after a capture injection and after the last injection in the sequence. For capture injections, a stabilization time can be useful if a fraction of the ligand dissociates rapidly. Including a stabilization time to allow for such dissociation can help to improve reproducibility.

A stabilization time may be used after the last injection instead of regeneration for systems where analyte dissociates completely from the surface.

Exposure of the surface to regeneration solution can often lead to transient changes in the baseline. Inclusion of a stabilization time after regeneration helps to ensure a stable baseline for the next cycle.

Sample injection
Normally, the injected sample solution is specified in a separate sample table. Some wizards (e.g. Surface Performance) use only a single sample solution that is specified together with the other injection parameters in this dialog box.

Regeneration
The parameters for regeneration include a check-box for High viscosity solution. Check this box if the regeneration solution has a relative viscosity higher than about 3 (corresponding to about 35% glycerol or 40% ethylene glycol at 20°C). This will modify the injection procedure for better handling of viscous solutions. The maximum injected volume is limited to 100 µl for viscous solutions.

4.2.4 Sample and control sample tables
Details of samples and control samples (where applicable) are entered in the Sample and Control Samples steps respectively. The details of these steps differ according to the wizard purpose, but the following general features may be noted. Further details are given in the respective wizard descriptions later in this chapter.
• The number of completed rows in the sample and control sample tables determine the number of cycles that will be run in the assay. The rack position requirements and the required volumes of common solutions such as regeneration (Section 4.2.6) are calculated on the basis of the number of samples that will be run.

• For some wizards, control samples are defined in a step in the dialog box sequence. For others, the **Control samples** dialog is accessed through a button in the **Samples** step.

• Sample details can be imported from an external file if this option is enabled under **Tools**:**Preferences**. See Appendix A for details of import formats and procedures.

### 4.2.5 System preparations

This dialog box specifies how the system will be prepared before the first cycle.
4 Application wizards
4.2 Common wizard components

Prime before run
This option flushes the flow system with running buffer to make sure that all buffer is fresh. You should generally prime the system before each run to ensure fresh buffer throughout the flow system.

Normalize
This option adjusts the detector response to compensate for small variations in reflectance characteristics between individual sensor chips. For best results, you should normalize the detector whenever the chip is changed. You do not need to run normalization if the same chip remains docked between runs. Normalization injects BIAnormalizing solution (70% glyceroll over the surface: if your ligand does not withstand exposure to this solution, normalize the detector before you immobilize the ligand.

Temperature settings
The Analysis temperature is the temperature at the flow cell. If the specified value differs from the current temperature, the system will wait at the beginning of the run until the analysis temperature is stable at the new value. You can choose to ignore temperature instability, but the response will drift as the temperature stabilizes. The absolute response decreases by about 150 RU for a 1°C increase in temperature.

The Sample compartment temperature is the temperature in the sample compartment. Equilibration of the sample compartment to a new temperature will start when the run is started. The system will not wait for a stable sample compartment temperature at the beginning of the run: samples equilibrate to the analysis temperature during passage through the IFC, so that the sample compartment temperature is not critical for the measured SPR response.

Note: Both analysis temperature and sample compartment temperature can be set in advance with the Set Temperature option from the main Tools menu (Section 2.3.6), to allow the temperature to equilibrate before setting up the assay wizard.

Cycle run list
Click Cycle Run List to display a preview of the cycles that will be run in the assay.
The **Assay step name** is generated automatically as a type identifier for each cycle. Other keywords may also be generated according to the wizard purpose. Keyword information is used in the Evaluation Software.

### 4.2.6 Rack positions

This dialog box shows where samples and reagents are to be placed in the microplate and/or rack. Positions are color-coded by region according to sample and reagent categories; you can change the color-coding in the **Automatic positioning** dialog, accessed through the *Menu* button.
Use the pull-down lists above the respective illustrations to change the reagent rack and microplate types. If change a rack or microplate type, all positions in the affected rack or plate will be cleared and must be reassigned either manually or automatically.

Positions are described by tool tips (hold the cursor on the position to display the tool tip). Empty positions show the position capacity and dead volume. Used positions show in addition the content name and the volume that will be used.

**Note:** The volumes listed in the table are minimum volumes. Use slightly larger volumes if material is available to allow for slight variations in the dead volume in microplates and vials.

You can change sample and reagent positions manually in two ways:

- Click on a sample or reagent in the sample plate and rack illustration and drag it to a new (empty) position. You cannot drag to a position that does not have sufficient capacity for the required volume of sample or reagent.
- Enter an unused position directly in the **Position** column in the table.

Positions can also be reorganized using the **Automatic positioning** dialog (see below).

**Menu functions**

Use the **Menu** button to access additional functions for rack positioning.

**Clear Positions**

This option clears the entries in the **Positions** column for the selected rack or plate.

Positions that are cleared must be reassigned before the run can be started. To reassign positions one by one, select a row in the positioning table and click on the required (empty) position in the illustration. To reassign all positions in one operation, choose **Default Positions** or **Automatic positioning** from the menu.

**Default Positions**

This option restores all entries to default positioning. The default positioning is determined from the type and volume of solution in combination with the currently selected rack type. Choosing **Default Positions** overrides any changes that have been made in the rack positions, even if the changed positions have been saved in the wizard template.
**Automatic Positioning**
This option controls the way samples and reagents are positioned automatically. Samples and reagents are placed by region, and samples within regions are kept together as far as possible.

<table>
<thead>
<tr>
<th>Region</th>
<th>This column lists the sample and reagent regions.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Color</td>
<td>This option controls the display color for the region.</td>
</tr>
<tr>
<td>Orientation</td>
<td>This column determines whether samples are arranged by column (vertically in the rack and plate diagram) or row (horizontally in the diagram).</td>
</tr>
<tr>
<td>Anchor</td>
<td>This column determines the position for the first sample in the region.</td>
</tr>
<tr>
<td>Rack</td>
<td>This option controls whether the samples and reagents will be placed in the reagent rack or the sample microplate. If Auto is chosen, placement is decided on the basis of number and volume of solutions in the region.</td>
</tr>
<tr>
<td>Vial size</td>
<td>Use this option to determine the vial size for reagents. If Auto is chosen, placement is decided on the basis of the volume of solution.</td>
</tr>
<tr>
<td>Pooling</td>
<td>This option allows you to combine solutions with the same name into one position or to split combined solutions into separate positions for each cycle. Choose Yes to pool solutions if suitable vial positions are available, or No if you always want separate positions for each cycle. Choose Auto to set the pooling according to the default settings for the type of region.</td>
</tr>
<tr>
<td>Sort by</td>
<td>Solutions within a region may be sorted by one or two parameters.</td>
</tr>
</tbody>
</table>

Use the **Move up** and **Move down** buttons to change the order in which regions are listed. Regions are placed in the specified rack or plate in the order listed, so that changing the order of the table can change the automatic positioning of samples and reagents.
4 Application wizards
4.2 Common wizard components

**Save Wizard Template/Save Wizard Template As**
Saves the wizard template, with either the same or a different file name. The corresponding function is also available for methods.

**Custom Position Import**
This option imports positioning information from an external source. The option is only available if *Enable custom position import* is checked in **Tools:Preferences**. Choosing the option first exports the rack positions table to a temporary tab-separated text file which is processed by the import program specified in the **Tools:Preferences** dialog. The output of the import program is then imported to the **Rack Positions** table, replacing the existing positioning information. See Appendix A for more details.

**Simple Position Import**
Imports positioning details from an external file. Details of the import settings and file format are described in Appendix A.

**Export Positions**
Exports the data in the positioning table to a tab-separated text file. See Appendix A for details of the exported file format.

**Print Rack Positions**
Prints a copy of the rack positions diagram and table.

**Print Wizard Template/Print Method**
Prints a copy of the currently open wizard template or method.
**4.2.7  Prepare Run protocol**

This dialog box allows you to enter a run protocol to provide instructions to the user when the run is started. The text in the *Prepare Run Protocol* is saved with the wizard template. A suggested general protocol is provided.

Select text and use the controls at the top of the dialog box to control the appearance (typeface, size and style) of the text.

The estimated run time and buffer requirement are shown at the bottom of the dialog box. For all wizards except buffer scouting, only buffer A is used. For the buffer scouting wizard (Section 4.7) and Method Builder-based runs (Section 5.4), buffer names are shown in the *Prepare Run Protocol* dialog.

**Notes:** The estimated run time and buffer consumption are minimum values, that do not include any time that cannot be predicted when the wizard is set up. This includes time for temperature equilibration at the beginning of the run or between cycles, for ligand injection in immobilization runs using *Aim for immobilized level* (Section 4.5) and for conditional statements (Section 5.6.1) in Method Builder-based runs.

The estimated buffer requirement includes a dead volume of at least 50 ml in the buffer bottle and is rounded up to the nearest 100 ml. The actual consumption will often be considerably less than the estimate. Make sure there is sufficient buffer in the bottle to allow for standby time after the run.

The *Menu* button provides options for saving and printing the wizard template (Section 4.2.6).
4.3 Wizard groups

The application wizards are organized into 5 groups:

- **Surface preparation**, covering immobilization pH scouting and immobilization
- **Assay development**, covering regeneration scouting, buffer scouting and surface performance tests
- **Control experiments** for kinetic measurements, covering tests for linked reaction mechanisms and mass transfer limitation
- **Assay wizards**, covering kinetics/affinity, binding analysis, concentration analysis and thermodynamics
- **Immunogenicity**, covering screening, confirmation and isotyping for immunogenicity investigations.

4.4 Immobilization pH scouting

The **Immobilization pH scouting** wizard helps you to find the optimal pH for immobilizing your ligand, by testing ligand pre-concentration at a range of pH values. See the Biacore Sensor Surface Handbook for further details. The injection sequence for immobilization pH scouting is fixed.

**Step 1. Setup**

Choose the flow path for the pH scouting. Immobilization pH scouting is restricted to a single flow cell within a run. The sensor surface in the flow cell should be unmodified.
Enter the buffers and pH values to be used for scouting. The default list covers sodium acetate buffers in the pH range 4 to 5.5, available as ready-to-use solutions from GE Healthcare. Buffers will be tested in the order listed.

**Note:** The buffers listed here are buffers in which the ligand should be prepared. They are not used as running buffers: you should use the same running buffer for pH scouting as you intend to use during immobilization.

**Step 2. Injection parameters**

Enter the name of the ligand to be tested and the contact time and flow rate. Recommended settings are a contact time of 120 seconds at 5 or 10 µl/min; you may need to use a longer contact time if preconcentration of ligand on the sensor surface proves to be slow.

The surface is washed with a “regeneration” injection at the end of each cycle to remove any ligand that might remain on the surface. The recommended solution for this procedure is 50 mM NaOH.

**Step 3. System preparations**

See Section 4.2.5. Run immobilization pH scouting at the same temperature as you intend to run the immobilization (electrostatic preconcentration is however usually fairly insensitive to temperature).

**Step 4. Rack positions**

See Section 4.2.6. Immobilization pH scouting requires one position for ligand solution at each pH tested and one for the surface wash solution. Accept or change the rack positions for the various solutions required (see Section 4.2.6).

**Step 5. Prepare run protocol**

Edit the **Prepare Run Protocol** text if desired (Section 4.2.7). This text will be displayed at the start of a run when the wizard template is used. Save the wizard template if required and start the run.
4 Application wizards
4.4 Immobilization pH scouting

**pH scouting results**

When the wizard run is completed, the results are opened automatically in the Evaluation Software, with an overlay plot of the sensorgrams adjusted to the start of the sample injection. The same overlay plot is created when a saved run from Immobilization pH Scouting is opened in the Evaluation Software.

**Note:** Opening a saved run in the Control Software does not generate the overlay plot automatically.

Choose the optimum buffer pH on the basis of the binding behavior: at pH suitable for immobilization, the ligand binds rapidly to the surface during the injection and dissociates completely after the end of the injection. The optimum is generally the highest pH value (i.e. the mildest condition) that gives sufficient ligand binding, not necessarily the value that gives the highest ligand binding. Beware of conditions that give irregular sensorgrams with incomplete dissociation: this behavior often indicates aggregation or denaturation of the ligand.
4.5 Immobilization

The Immobilization wizard supports immobilization of ligand in any combination of the four flow cells in one run. Immobilization in each flow cell is performed independently in a separate cycle, so that different ligands and/or immobilization conditions can be used in the different flow cells. See the Biacore Sensor Surface Handbook for more information about ligand immobilization.

**Step 1. Immobilization setup**

The choice of Chip type determines the predefined methods that are available for immobilization. The type chosen when the chip was docked is chosen by default: if you change the chip type you will be able to create and save an immobilization wizard template, but you must dock a corresponding chip type before the immobilization can be performed.

Check the flow cells where you want to perform immobilization. For each flow cell, set the parameters as follows:

Choose the immobilization method. Predefined methods are provided for standard immobilization chemistries. Customized methods can be defined by clicking on the **Custom Methods** button (see below). Predefined methods are marked with a T200 icon (t²) in the selection lists.
Choose the way in which immobilization will be controlled:

- If you choose **Aim for immobilized level**, you specify a target level. The immobilization procedure will attempt to reach this level as described below.

- If you choose **Specify contact time and flow rate**, enter the settings in the respective fields.

- If you choose **Blank immobilization**, the surface will be activated and deactivated in accordance with the immobilization method but no ligand will be injected.

Enter the ligand name. To dilute the ligand solution immediately before injection, check **Dilute ligand** and enter a percentage value and a solution name. This option can be used for ligands that have limited stability in immobilization buffer, and that are diluted from a stock solution just before immobilization. A setting of 90% will mix one part of ligand solution with 9 parts of the specified diluent.

**Aiming for immobilized level**

The option **Aim for immobilized level** injects a pulse of ligand over the unactivated surface in order to estimate the rate of preconcentration. The surface is washed to remove traces of ligand and then activated. The procedure then uses ligand contact times based on the preconcentration estimate to attempt to reach the specified target level. If preconcentration is either too fast or too slow to permit the target level to be reached, this will be reported and immobilization will not be performed.

The preconcentration injection injects 10 µl ligand solution at a flow rate of 5 µl/min, giving a contact time of 2 minutes. This injection is included in predefined methods for CM-series sensor chips but is optional in customized methods (see below). If **Aim for immobilized level** is chosen together with a custom method that does not include a preconcentration injection, the immobilization procedure will activate the surface and then inject short pulses of ligand until either the target level or the maximum total ligand volume of 150 µl is reached. This can be used to conserve valuable ligand without losing the benefits of aiming for a target immobilization level, and can be useful for sensor chips where preconcentration cannot be performed, e.g. Sensor Chip SA.
**Custom methods**

Click **Custom Methods** to define customized immobilization methods.

Click **New** to create a new blank method. Select an existing method and click **Copy** to make a copy of the method or **Delete** to delete the method. You cannot delete the predefined methods (marked with a ![icon](image) icon).

For a new method, enter a name in the **Method name** field. Construct the sequence of injections for the immobilization method using the buttons to the right of the main panel. The ligand injection is created automatically and cannot be deleted: solution and contact time for the ligand injection are specified in the main wizard dialog. A method may only contain one ligand injection. Other injections have the following functions:
### Step 2. System preparations

Check the **System preparations** options as required (see Section 4.2.5).

### Step 3. Rack positions

Accept or change the rack positions for the various solutions required (see Section 4.2.6).

---

**Pre-conc** injects 10 µl of ligand solution at 5 µl/min to estimate the rate of preconcentration. This step is only performed if the option **Aim for immobilized level** is chosen when the immobilization method is used. A method may only contain one **Pre-conc** injection. The **Pre-conc** injection should always be placed before surface activation: it will usually be first in the method, although it may be preceded by a surface conditioning injection if required. If you place the **Pre-conc** injection after the surface activation, it will be executed there and the ligand will be immobilized on the activated surface.

After the **Pre-conc** injection, the surface is washed with a solution specified in the immobilization setup dialog, to remove any ligand that may remain on the surface.

Do not use a **Pre-conc** injection with Sensor Chip SA, since biotinylated ligand will bind to the surface and cannot be removed.

**Inject** performs an injection of a specified solution with a specified contact time and flow rate. Values are entered in the dialog box that appears when you click **Inject**.

**Mix & Inject** mixes two specified solutions and performs an injection of the mixture. Details are entered in the dialog box that appears when you click **Mix & Inject**.

**Wash** washes the flow system (but not the sensor surface). The wash solution is specified in the dialog box that appears when you click **Wash**.

Select an injection and use the **Edit**, **Delete**, **Move up** and **Move down** buttons to edit the injection details, remove the injection from the method and change the order of injections in the method.

Custom methods are stored in the immobilization wizard template: if you need the same or slightly modified method in a different template, save a copy of the template and then edit the method.
**Step 4. Prepare run protocol**

Edit the **Prepare Run Protocol** text if desired (Section 4.2.7). This text will be displayed at the start of a run when the wizard template is used. Save the wizard template if required and start the run.

**Immobilization results**

The results of an immobilization run are summarized in the Control Software and logged in the **Chip Properties** (see Section 2.3.4) when the run is completed.

**Note:** The same information can be accessed under File:Properties in the Evaluation Software.

The summary lists the procedure and method, the name of the ligand and whether the target was reached with **Aim for immobilized level**. Two response values are reported, one directly after the ligand immobilization and one after the deactivation injection. The difference between these values is an indication of the amount of non-covalently bound ligand that is washed from the surface by the deactivation injection.

Note that the **Response bound** value does not include the contribution from activation by EDC/NHS. For low ligand levels, this value can give a better indication of the amount of ligand immobilized.

Result files from immobilization can also be opened in the Evaluation Software if you want to prepare other sensorgram displays or plots (see Chapter 7).
4.6 Regeneration scouting

Regeneration scouting

The Regeneration Scouting wizard guides you through the process of finding suitable regeneration conditions for your sensor surface. The principles of regeneration scouting are described in the Biacore Sensor Surface Handbook. Briefly, regeneration scouting is performed by testing repeated cycles of analyte injection and regeneration over a range of regeneration conditions, and assessing the results on the basis of trends in analyte response and baseline levels. The analyte concentration should be relatively high for best results. The analyte response reflects the binding capacity (ligand activity) of the surface, while the baseline level indicates the extent of regeneration. Each condition should be tested for at least 3 cycles in sequence (recommended number 5) in order to detect trends in the regeneration behavior with the given condition. When testing multiple conditions, start with the mildest conditions to minimize the risk of losing ligand activity at the beginning of the scouting series.

**Step 1. Injection sequence**

Choose the detection settings, chip type and injection sequence for the regeneration scouting (see Section 4.2.1). One sample injection and one or two regeneration injections are required. Most sensor surfaces can be adequately regenerated with a single injection, but some situations may benefit from using multiple injections.

**Step 2. Setup**

Specify a conditioning cycle if required.

Regeneration scouting always includes one startup cycle with the same injection sequence as the scouting cycles but with injection of buffer for all injections. Note that this differs from the startup cycle construction in the other application wizards.

**Step 3. Injection parameters**

Specify the injection parameters for each injection in the cycle (see Section 4.2.3). The same sample will be used for all cycles.
Step 4. Experimental parameters

This dialog box determines the design of your regeneration scouting. Set the number of conditions to test and the number of cycles for each condition and specify the conditions in the Settings frame. The default number of cycles for each condition is five. You may use fewer cycles to shorten the total run time for exploratory work, but five cycles are recommended for fine-tuning conditions in order to reveal trends in the regeneration performance.

Use variants of the same kind of regeneration conditions (e.g. different pH values or different concentrations of ethylene glycol) within the same run. Results are most easily interpreted if you use a separate wizard run with a new flow cell or sensor chip for each kind of regeneration condition that you test, so that the outcome with one kind of condition is not affected by the history of exposing the ligand to another condition.

You may choose to lock the solutions or the contact time used for regeneration tests, so that all conditions will use the same setting for the locked parameter. Do not vary both the solution and the contact time at the same time: the results may be difficult to interpret clearly.

Check High viscosity solution if any of the regeneration solutions tested has a relative viscosity higher than about 3 (corresponding to about 35% glycerol or 40% ethylene glycol at 20°C). This will modify the injection procedure for better handling of viscous solutions.

Step 5. System preparations

Check the System preparations options as required (see Section 4.2.5).
Step 6. Rack positions
Accept or change the rack positions for the various solutions required (see Section 4.2.6).

Step 7. Prepare run protocol
Edit the Prepare Run Protocol text if desired (Section 4.2.7). This text will be displayed at the start of a run when the wizard template is used. Save the wizard template if required and start the run.

Regeneration scouting results
Regeneration scouting results are presented in the Control Software when the run is completed.

Note: This result presentation is not shown if the run is opened in the Evaluation Software.

The Trend chart tab shows the results as a plot of baseline and sample response for each cycle in the run, grouped by regeneration conditions. Conditions are identified in tool tips for the data points (place the cursor on a point for a couple of seconds to display the tool tip).

Note: Report points are set before the sample injection for baseline and shortly after the sample injection for sample response. Thus the points for the first cycle indicate the starting values, while those for subsequent cycles each indicate the effect of the previous cycle.
Check **1st cycle** to include the starting values derived from the first sample cycle in the plot. (This cycle is shown as cycle number 2 or 3: conditioning and startup cycles are numbered but not shown.)

Select which conditions to display in the **Conditions** box. Use Shift-click to make multiple adjacent selections, Ctrl-click to make multiple non-adjacent selections. The scale of the display will be adjusted according to the number of cycles displayed.

Select which curves to display in the **Sensorgram** box.

The **Sensorgrams** tab shows the sensorgrams for regeneration scouting. Select the conditions and cycles to display in the respective boxes. Check **Zoom lock** to keep the scale fixed when the choice of sensorgrams is changed.

Result files from regeneration scouting can also be opened in the Evaluation Software if you want to prepare other sensorgram displays or plots (see Chapter 7). See the Biacore Sensor Surface Handbook for a discussion of how to interpret the results of regeneration scouting.
4 Application wizards
4.7 Buffer scouting

4.7 Buffer scouting

The Buffer Scouting wizard helps you to test the effect of up to four different buffers on your assay, using the buffer selector valve to switch running buffers.

**Step 1. Injection sequence**

Choose the detection settings, chip type and injection sequence for the buffer scouting (see Section 4.2.1). The flow path can either be chosen explicitly (in which case the same flow path will be used for each buffer), or set to vary with the buffer (in which case a single flow cell will be used for each buffer, with flow cells 1, 2, 3 and 4 for buffers A, B, C and D respectively).

**Step 2. Setup**

Specify the conditioning and start-up cycles as required. If start-up cycles are chosen, separate rack positions will be created for the start-up sample solution in each buffer tested, and the start-up cycles will be run at the start of each buffer test. Conditioning if used will only be run once at the start of the experiment.

Specify the buffers you want to test. You can enter up to 4 different buffers. The buffers will be tested in the order given.

**Note:** If you use less than 4 buffers, positions A, B and C will be used in order. If you enter buffer names on non-consecutive rows, the table will be adjusted when you leave the dialog box.

**Step 3. Injection parameters**

Specify the injection parameters for each injection in the cycle (see Section 4.2.3).
Step 4. Samples
Enter the samples to be tested in the buffer scouting. The scouting procedure will work through the sample table for the first buffer before switching to the next buffer.

Ligands for capture, samples and enhancement reagents should be prepared in each of the buffers tested. Separate rack positions will be created for samples in each buffer (for example, running buffer scouting with 4 buffers and 5 samples will require 20 sample positions).

Step 5. System preparations
Check the System preparations options as required (see Section 4.2.5).

Step 6. Rack positions
Accept or change the rack positions for the various solutions required (see Section 4.2.6).

Step 7. Prepare run protocol
Edit the Prepare Run Protocol text if desired (Section 4.2.7). This text will be displayed at the start of a run when the wizard template is used. Save the wizard template if required and start the run.

Buffer scouting results
When the wizard run is completed, the results are opened automatically in the Evaluation Software. In addition to the general predefined plots (Section 6.4), plots of binding and stability against sample are created to visualize the behavior in the different buffers.
4 Application wizards
4.8 Surface performance

4.8 Surface performance

The Surface Performance wizard allows you to test the robustness of your surface by performing repetitions of the same analysis cycle. The number of repetitions is in practice usually limited by the capacity of the sample plate (the software allows up to 100 cycles). The cycle requires one sample injection, and can also include capture and enhancement steps and one or two regeneration steps. Use this wizard for example when you want to confirm that the regeneration conditions that you identified in regeneration scouting hold good for an extended number of cycles.

Step 1. Injection sequence
Choose the detection settings, chip type and injection sequence for the surface performance test cycle (see Section 4.2.1).

Step 2. Setup

Specify conditioning and start-up cycles (see Section 4.2.2).

Set the number of repetitions of the analysis cycle according to the purpose of the surface performance test. As a general guide, the test should run for at least as many cycles as will be used normally in the assay.

Step 3. Injection parameters
Specify the injection parameters for each injection in the cycle (see Section 4.2.3).

Step 4. System preparations
Check the System preparations options as required (see Section 4.2.5).
Step 5. Rack positions
Accept or change the rack positions for the various solutions required (see Section 4.2.6).

Step 6. Prepare run protocol
Edit the Prepare Run Protocol text if desired (Section 4.2.7). This text will be displayed at the start of a run when the wizard template is used. Save the wizard template if required and start the run.

Surface performance results
When the wizard run is completed, the results are opened automatically in the Evaluation Software. Examine the plots of baseline and sample response against cycle number. The response values should ideally be unchanged throughout the run.
4.9 Binding analysis

The Binding Analysis wizard supports injection of up to four samples in series, in addition to ligand capture, enhancement and regeneration steps. This wizard is suitable for analysis of applications like multi-component complex formation and pair-wise epitope mapping, as well as simple applications like screening for binding partners to an immobilized ligand.

Step 1. Injection sequence

Choose the detection settings, chip type and injection sequence for the binding analysis (see Section 4.2.1). Up to four sequential sample injections may be included in each cycle.

Step 2. Setup

Specify the conditioning and start-up cycles (see Section 4.2.2).
Step 3. Injection parameters

Specify the injection parameters for each injection in the cycle (see Section 4.2.3). One sample injection panel will be created for each sample injection in the injection sequence. The sample injection for binding analysis has an additional setting for dissociation time. This is the time for which dissociation will be monitored after the end of the injection without disturbances from flow system washing procedures.

Step 4. Samples

The sample table contains one column for each sample injection in the injection sequence. New rows are created as you enter data in the table. (The illustration above shows how the wizard could be used to set up a pair-wise epitope mapping experiment.)

Click Import to import the sample data from an external file. Import of sample information must be enabled in Tools:Preferences to use this function. See Appendix A for details of import functions and file formats.

Click Control Samples to enter control samples for the run.
Specify the details of control samples and the frequency with which they should be run. If you check only Run control samples, the controls will be run once only at the start of the assay. If you check Repeat control samples every..., the controls will be run at the start of the assay and then at the specified interval, and again at the end of the assay. Use the Cycle Run List option in the System Preparations step to verify where control samples will be run during the assay.

**Step 5. System preparations**
Check the System preparations options as required (see Section 4.2.5).

**Step 6. Rack positions**
Accept or change the rack positions for the various solutions required (see Section 4.2.6).

**Step 7. Prepare run protocol**
Edit the Prepare Run Protocol text if desired (Section 4.2.7). This text will be displayed at the start of a run when the wizard template is used. Save the wizard template if required and start the run.

**Binding analysis results**
When the wizard run is completed, the results are opened automatically in the Evaluation Software. Predefined plots (Section 6.4) are created for each sample.
4.10 Concentration analysis

The *Concentration Analysis* wizard helps you to set up an assay for determining analyte concentration in samples with the help of a calibration curve using known concentrations. Control samples may be included at intervals to monitor the stability of the assay.

**Note:** Analyte concentrations can be determined without reference to a calibration curve using the calibration-free concentration assay approach, supported in Method Builder (see Section B.2).

**Step 1. Injection sequence**

Choose the detection settings, chip type and injection sequence for the concentration analysis (see Section 4.2.1).

**Step 2. Setup**

Specify the conditioning and start-up cycles (see Section 4.2.2). Start-up cycles are recommended for concentration assays to ensure that the initial response drift that may occur with a new chip does not interfere with the first measurements.

**Step 3. Injection parameters**

Specify the injection parameters for each injection in the cycle (see Section 4.2.3).

The sample injection for concentration analysis can be extended to include a mixing function, whereby sample is mixed with a specified proportion of a second fixed solution. This feature enables inhibition assay formats where samples are mixed with a constant proportion of a detecting molecule solution. The value specified for *Fraction* refers to the proportion of the fixed component in the final mixture: for example, a value of 30% will mix 7 parts of sample with 3 parts of the specified solution.

The volumes of sample and mixing solution used are determined automatically so that the final volume of mixed solution is sufficient for the injection.
Notes: Mixing in the autosampler is very reproducible, but high accuracy cannot be guaranteed. If your application requires accurate mixing proportions, mix the samples outside the autosampler.

Mixing is not supported in 384-well microplates. The wells on these plates are too small for reliable mixing in the autosampler.

**Step 4. Calibration curve**

Specify the details of the calibration curve for the concentration measurement. Check **Repeat calibration** and enter a repeat interval to repeat the calibration curve at regular intervals during the assay. The calibration curve will be run at the beginning of the assay (as indicated by the checked option **Run first**, which cannot be changed) and at the specified interval. Thus specifying a repeat every 15 sample cycles and running 35 samples will result in calibration curves at the beginning and after samples 15 and 30.

Check **Run last** to include a calibration curve as the last cycle in the assay. Use this option if you intend to use calibration trends in the evaluation to compensate for drift in the calibration curve (see Section 8.2.2).

Enter the concentrations for the calibration points on the curve. You must enter at least two concentrations for the calibration curve. (Two concentrations are sufficient for a linear calibration curve, but if you intend to use the recommended four-parameter fitting function for the calibration curve, you need at least four points.) To run replicate concentrations, enter the same concentration on multiple rows. Calibration points will be run in the order entered. You can choose a different concentration unit if required from the pull-down list in the table header.
Step 5. Control samples

Specify the details of control samples and the frequency with which they should be run. You may choose not to run control samples at all; however, including control samples is generally recommended as an aid in assessing the performance of the assay.

If you check only Run control samples, the controls will be run once only at the start of the assay. If you check Repeat control samples every..., the controls will be run after the first calibration curve and then at the specified interval for sample cycles (not counting repeated calibration curves), and once at the end of the assay. Use the Cycle Run List option in the System Preparations step to verify where control samples will be run during the assay.

Control samples are specified in terms of sample ID and expected concentration. The expected concentrations should lie within the range covered by the calibration curve.
4 Application wizards

4.10 Concentration analysis

Step 6. Samples

Enter the details of the samples to be analyzed. Each sample is defined by a sample ID and a dilution factor: the dilution factor is used during evaluation to calculate the measured concentration in the original undiluted sample. For undiluted samples (dilution factor 1), this column may be left blank. Samples will be analyzed in the order entered. To analyze replicate samples, enter same sample on multiple rows.

Click Import to import the sample data from an external file. Import of sample information must be enabled in Tools:Preferences to use this function. See Appendix A for details of import functions and file formats.

Step 7. System preparations

Check the System preparations options as required (see Section 4.2.5).

Step 8. Rack positions

Accept or change the rack positions for the various solutions required (see Section 4.2.6).

Step 9. Prepare run protocol

Edit the Prepare Run Protocol text if desired (Section 4.2.7). This text will be displayed at the start of a run when the wizard template is used. Save the wizard template if required and start the run.

Evaluation of concentration assays

Chapter 8 describes how to evaluate concentration assays.
4.11 Kinetics/Affinity

The Kinetics/Affinity wizard guides you through the setup of experiments to determine kinetic constants or affinity constants for an interaction. Wizards for control experiments relevant to kinetic analysis are described in Section 4.13.

Note: The Kinetics/Affinity wizard supports kinetic determinations in multi-cycle format, where each analyte concentration is injected in a separate cycle and analyte is allowed to dissociate fully or is removed by regeneration between cycles. An alternative format that does not require regeneration or full dissociation is single-cycle kinetics, supported in Method Builder (see Section B.10).

Step 1. Injection sequence

Choose the detection settings, chip type and injection sequence for the assay (see Section 4.2.1). Only reference subtracted detection using either Fc2-1 or Fc4-3 is available for kinetic analysis. The Kinetics wizard supports capture but not enhancement injections.

A Carry Over injection is also available in the Kinetics/Affinity wizard. This performs an injection of buffer as the last injection in the sample analysis cycle, to check that the flow system is clean from residual analyte that would otherwise carry over to the sample injection in the next cycle. The Carry Over injection is fixed as a 30-second injection at 40 µl/minute, regardless of other settings in the wizard. Report points are set automatically for the carry-over injection to allow evaluation of potential carry-over problems.
**Step 2. Setup**

Specify the conditioning and start-up cycles (see Section 4.2.2). Start-up cycles are recommended for kinetics experiments to ensure that the initial response drift that may occur with a new chip does not interfere with the first measurements.

Check **Run solvent correction** to include solvent correction cycles in the run. Solvent correction adjusts response values for the effects of varying bulk refractive index contribution between samples, and may improve the quality of the results for analyses using small organic analytes that require dimethyl sulfoxide (DMSO) in the sample buffer to maintain solubility. The principles and application of solvent correction are described in Section 6.7.

Set the required number of injections per cycle and the frequency of solvent correction cycles. The default settings use 8 injections per cycle and repeat the solvent correction cycle every 30 sample cycles.
Step 3. Injection parameters

Specify the injection parameters for each injection in the cycle (see Section 4.2.3). The sample injection for kinetics measurement has an additional setting for dissociation time. This is the time for which dissociation will be monitored after the end of the injection without disturbances from flow system washing procedures. An extra wash after the sample injection with 50% DMSO is recommended for work with low molecular weight analytes. This wash solution does not pass over the sensor surface.

Step 4. Samples
Enter the details of the samples for kinetic or affinity determination. For each analyte, a zero concentration sample and at least four non-zero concentrations, one of which is run in duplicate, are strongly recommended. Concentrations are entered in the left-hand **Concentration** column, either in molar or weight-based units. Choose the concentration units from the pull-down list in the column header. If a weight-based concentration unit is chosen (e.g. µg/ml) a molecular weight must also be specified. When a molecular weight is entered, the right-hand **Concentration** column displays the conversion from molar to weight-based or vice versa. Samples with the same sample name may not be given different molecular weights.

The samples may be analyzed either in the order entered in the table or sorted in increasing concentration. The order displayed in the sample table is not affected by the choice of run order.

If you enter samples with different names, they will be handled as separate concentration series regardless of the order in which they are entered. The samples will be run as separate concentration series even if the order is mixed in this dialog: thus samples entered in the order A, B, A, B, A, B, B... will be run in the order A, A, A, A, B, B, B, B... The **Run order** setting applies within each concentration series.

Click **Import** to import the sample data from an external file. Import of sample information must be enabled in **Tools:Preferences** to use this function. See Appendix A for details of import functions and file formats.

Click **Control Samples** to enter control samples for the run.
Specify the details of control samples and the frequency with which they should be run. If you check only **Run control samples**, the controls will be run once only at the start of the assay. If you check **Repeat control samples every**..., the controls will be run at the start of the assay and then at the specified interval, and again at the end of the assay. Use the **Cycle Run List** option in the **System Preparations** step to verify where control samples will be run during the assay.

**Note:** Control samples do not require a concentration series and will not be evaluated for kinetics and affinity. These samples are intended as check on the consistency of binding levels based on single report point measurements.

**Step 5. System preparations**
Check the **System preparations** options as required (see Section 4.2.5).

**Step 6. Rack positions**
Accept or change the rack positions for the various solutions required (see Section 4.2.6).

**Step 7. Prepare run protocol**
Edit the **Prepare Run Protocol** text if desired (Section 4.2.7). This text will be displayed at the start of a run when the wizard template is used. Save the wizard template if required and start the run.

**Evaluation of kinetics/affinity assays**
Chapter 9 describes how to evaluate kinetics/affinity assays.
4.12 Thermodynamics

The Thermodynamics wizard supports kinetic and affinity determinations over a range of temperatures. The corresponding evaluation software extracts thermodynamic data from the dependence of rate and affinity constants on temperature (see Section 10.1).

**Step 1. Injection sequence**
Choose the detection settings, chip type and injection sequence for the assay (see Section 4.2.1). Options for detection settings are the same as for kinetics/affinity determinations (Section 4.11). The thermodynamics wizard supports capture but not enhancement injections. The same injection sequence will be used at all temperatures.

**Step 2. Setup**

Specify the conditioning and start-up cycles (see Section 4.2.2). Start-up cycles are recommended for thermodynamics experiments to ensure that the response drift that may occur with a new chip or when the temperature is changed does not interfere with the first measurements. Start-up cycles will be run at each temperature.
Check Run solvent correction to include solvent correction cycles in the run. Solvent correction adjusts response values for the effects of varying bulk refractive index contribution between samples, and may improve the quality of the results for analyses using small organic analytes that require dimethyl sulfoxide (DMSO) in the sample buffer to maintain solubility. The principles and application of solvent correction are described in Section 6.7.

Set the required number of injections per cycle and the frequency of solvent correction cycles. The default settings use 8 injections per cycle and repeat the solvent correction cycle every 30 sample cycles. Separate solvent correction cycles will be run at each temperature.

Enter the temperatures at which the measurements are to be performed. For most purposes, 5-7 temperatures will be adequate: fewer points make the determination of thermodynamic parameters uncertain, while more points increase the run time. Distribute the points evenly over the widest temperature interval that the ligand and analyte tolerate. Start from the lowest temperature to minimize the time needed for temperature equilibration between measurements (increasing the analysis temperature takes less time than decreasing it). The system will wait for a stable temperature between each determination. As an additional control, you may want to include a replicate of the first temperature at the end of the run.

Note: For robust determination of standard thermodynamic parameters ($\Delta G^\circ$, $\Delta H^\circ$ and $\Delta S^\circ$), arrange the analysis temperatures so that 25°C is in the middle of the range (e.g. 10–40°C). See Section 10.1 for further details.

Specify the sample compartment temperature, or check Vary with analysis temperature to change the sample compartment temperature automatically when the analysis temperature is changed.

**Step 3. Injection parameters**

Specify the injection parameters for each injection in the cycle (see Section 4.2.3). The sample injection for thermodynamics measurement has an additional setting for dissociation time. This is the time for which dissociation will be monitored after the end of the injection without disturbances from flow system washing procedures.

**Step 4. Samples**

Enter the details of the samples. This dialog box is equivalent to the concentration series dialog for kinetics determination (see Section 4.11, step 4). Separate rack positions will be created for each sample at each temperature.

**Step 5. System preparations**

Check the System preparations options as required (see Section 4.2.5). The settings for analysis temperature and sample compartment temperature
cannot be changed here: the values for the first temperature as specified in step 2 are shown.

**Step 6. Rack positions**
Accept or change the rack positions for the various solutions required (see Section 4.2.6).

**Step 7. Prepare run protocol**
Edit the *Prepare Run Protocol* text if desired (Section 4.2.7). This text will be displayed at the start of a run when the wizard template is used. Save the wizard template if required and start the run.

**Evaluation of thermodynamics assays**
Section 10.2 describes how to evaluate thermodynamics assays.
4.13 Control experiments

Two control experiments are currently supported by wizards, both for kinetic analyses:

- **The Mass transfer** control experiment analyses the interaction of one or more analyte concentrations at three different flow rates, to establish whether the observed binding rate varies with flow rate. A dependence of binding rate on flow rate indicates that the binding is limited to some extent by mass transfer of analyte to the sensor surface. If mass transfer limitations are too significant, reliable kinetic data cannot be obtained (see Section 9.4.2).

- **The Linked reactions** control experiment analyses the interaction of one or more analyte concentrations for different contact times, to identify a particular kind of deviation from a 1:1 binding mechanism. Variation of the dissociation behavior after the end of the injection with the contact time indicates that the observed binding consists of at least two processes, one where the analyte binds to the surface and a second where the surface-attached complex undergoes a stabilizing alteration such as a conformational change.

The control experiment wizards have the same basic structure as the kinetic analysis wizard, but some of the analysis settings are fixed.

4.13.1 Mass transfer control

Each sample entered in the **Samples** step will be analyzed three times, at fixed flow rates of 5, 15 and 75 µl/min, with a contact time of 1 minute and dissociation time of 2 minutes. Use an analyte concentration that gives readily measurable initial binding rates. Mass transfer limitation is not affected by analyte concentration, but dependence of binding rate on flow rate may be difficult to detect if the binding rate is too low or too high.

**Note:** All predefined kinetic evaluation models include a term for mass transfer, and the original purpose of the mass transfer control experiment has largely been superseded by functions in the evaluation software. The control experiment may however still be useful to confirm suspected mass transfer limitations if desired.

4.13.2 Linked reactions control

Each sample entered in the **Samples** step will be analyzed at a flow rate of 10 µl/min with fixed contact times of 0.5, 3 and 10 minutes and a dissociation time of 10 minutes. Use one or more fairly high analyte concentrations, preferably so that steady state is approached or reached within the shortest contact time. The experiment is easiest to interpret if the interaction is
4 Application wizards
4.13 Control experiments

maintained at steady state for varying lengths of time, so that the starting
response for dissociation is constant.

**Note:** The *Injection Parameters* step allows you to enter a stabilization time
after each sample injection, but all other settings in this dialog are fixed in
the wizard.

4.13.3 Evaluation of control experiments

When the wizard run is completed, the results are opened automatically in the
Evaluation Software, with overlay plot of adjusted sensorgrams according to the
control experiment.

**Mass transfer**

The sensorgrams are adjusted to zero response and time at the baseline report
point. Compare the observed binding rates at the different flow rates. If the
observed binding rate during sample injection varies with flow rates, there is
some degree of mass transfer limitation in the data. Note that it may still be
possible to obtain kinetic information from the sensorgrams (see Chapter 9).

This example shows a clear dependence of binding rate on flow rate, indicating
mass transfer limitations in the observed binding. It may however still be possible
to extract kinetic information from the sensorgrams.
**Linked reactions**

The sensorgrams are adjusted to zero response at the baseline report point and to zero time at the end of the sample injection. Compare the observed dissociation rates at the different contact times. Variation of the observed dissociation rate after sample injection with contact time indicates linked reactions in the interaction model.

![Graphic of sensorgrams](image)

*This example shows a clear dependence of dissociation rate on contact time, indicating linked reactions in the interaction model.*
4.14 Immunogenicity

Biacore T200 Control Software includes application wizard support for three areas of immunogenicity studies:

- **Immunogenicity Screening**, designed for detection of immune response to administered drugs. Enhancement reagents can be used to reduce false positive results by confirming that observed responses derive from antibodies. This wizard can also be used to provide an estimate of the binding stability of detected antibodies to the drug.

- **Immunogenicity Confirmation**, designed for confirming the specificity of detected antibodies through inhibition of the observed response by excess drug added to the sample

- **Immunogenicity Isotyping**, designed to assist in identifying the isotype of detected antibodies through the use of isotyping reagents as enhancement reagents.

Both **Confirmation** and **Isotyping** wizards can be used for direct analysis of serum samples or to detect antibodies in the presence of drug interference, where antibodies are complexed with excess drug in the sample and would not be detected by a standard direct analysis.

Details of support for immunogenicity studies are provided in the separate Biacore T200 Immunogenicity Handbook.
Methods

Methods in Biacore T200 offer flexibility in instrument control, providing support for applications that cannot conveniently be handled with wizards. Methods are constructed with the Method Builder tool as described in this chapter. Templates from application wizards can be opened in Method Builder (Section 5.1) to provide a starting point for development of customized applications. Method examples are provided with the software installation.

5.1 Opening methods

To open an existing method or create a new method, choose File: Open/New Method.

Select a method and click Open to open the method, or click New to create a new method. Predefined methods for common applications are provided in the folder Biacore Methods. If you make changes to a predefined method, you must save your changed method under a new name.

Check Show importable wizard templates to display wizard templates that can be opened in Method Builder. Opening a wizard template imports all wizard settings into a method and allows you to add functionality that is not supported in the wizard. Templates from all wizards except immobilization can be imported into Method Builder.

The top-level folder for methods is defined under Tools: Preferences (see Section 2.4). You can navigate between subfolders under the top level in the dialog box, but files outside the top-level folder are not listed in the dialog box. Click Browse to navigate freely in the computer file structure and open methods stored in other locations.
5 Methods
5.2 Method structure

5.2 Method structure
Methods are handled in Method Builder in a series of sections representing different aspects of the method definition.

Overview
The Overview screen summarizes the method definition. Use this information as an aid in checking that the method is correctly built.

General settings
Here you define general parameters such as the concentration unit for samples, sample compartment temperature, data collection rate, detection mode and buffer names.

Assay steps
An assay step represents a specific function in the assay, defined in terms of what the step is intended to achieve. Assay steps may for example be start-up operations, solvent correction, sample analysis or control sample analysis. Steps can be run singly or repeated within the context of other steps: for example, start-up operations are typically performed once at the start of a run, while control samples may be repeated at intervals during the sample analysis. Analysis temperature and buffer selection can be set individually for each assay step.

Cycle types
Cycle types define the details of how assay steps will be performed, in terms of sample and reagent injections. Each assay step is linked to one cycle type, but the same cycle type can be used in multiple assay steps. For example, sample and control sample analysis are two assay steps that will typically use the same cycle type, ensuring that controls are analyzed in exactly the same way as samples.

Parameters for injections in a cycle type definition may be variable, so that they can be assigned a series of different values when the method is used. Sample names will typically be variable. The number of values for variable parameters together with assay step repetition determines the number of cycles that will be run.

Report points can also be defined for each cycle type.

Variable Settings
This section determines how values for variable parameters are specified. You can choose whether values are specified in the method or at run-time: this can be used to restrict the number of parameters that have to be entered when the method is run while at the same time maintaining flexibility for method development purposes.
Verification
Once the method has been defined in full, this section verifies that all aspects are consistent and completely specified. The verification results are reported in the work area. A method that does not pass verification can be saved but cannot be run.

Note: Verification only checks the consistency and completeness of the method. It does not in any way verify that the method is suitable for the intended purpose.

Each aspect of Method Builder is described in detail in the following sections.

5.3 Method overview

This screen provides a summary of the method. The main panel shows the assay steps in the method (see Section 5.5). Click on an assay step to show the settings for the step and the details of the cycle definition (see Section 5.6) in the panels to the right. The cycle definition is listed as a series of injection commands: to see command details, expand individual commands by clicking on the + marking or use the Expand All button to expand all commands in the panel.

This screen is for information only: settings cannot be changed here.
5.4 General settings

The settings that are specified here are:

**Data collection rate**
Choose between 1 and 10 Hz for data collection. The higher setting will provide better resolution for kinetic analysis of fast interaction processes, but will result in larger result files.

**Detection**
Choose the detection setting for the run:

- **Single**: Records data from one flow cell according to the chosen flow path. Data is not recorded from the other flow cells.
- **Dual**: Records data from one flow cell pair (1,2 or 3,4) according to the chosen flow path. Data is not recorded from the other flow cell pair.
- **Multi**: Records data from all four flow cells.

This setting affects the choice of flow path that can be made for each injection command in the cycle types definitions (see Section 5.6.1). Use the setting **Multi** if you are not sure what you need: this will provide maximum flexibility for the injection settings.

**Sample compartment temperature**
This is the temperature in the sample compartment (not the analysis temperature at the flow cell, which is set for each assay step). Check the **Vary with analysis temperature** box to set the sample compartment temperature automatically to the same value as the analysis temperature.
**Concentration unit**
This setting defines the unit for entering sample concentrations. The unit must be specified here, and cannot be changed at any other step in the assay definition. The unit can however be changed in the Evaluation Software when the results of the run are evaluated.

**Buffer settings**
Enter names if desired for the buffers in bottles A to D. Names entered here will be displayed in the *Prepare Run Protocol* (Section 4.2.7). Different buffers may be chosen for different assay steps (Section 5.6).

**Specify analysis temperature after run**
Check this option and enter a temperature to set the analysis temperature when the run is completed. The rack temperature will also be reset if the *Vary with analysis temperature* box is checked. This setting provides automated control of the chip and detector environment after the completion of a run, for example in preparation for another run at a different temperature.

### 5.5 Assay steps

This screen determines the main structure of the method in terms of assay steps. Steps at the top level (i.e. not indented or marked with the symbol ) are executed in the order given. Nested steps (marked with the symbol ) are executed within the context of the level in which they are placed, as specified by the settings for *Recurrence*.

To create a new assay step, click *New Step*. The step will be created with default settings at the end of the current method. Move the step in the method with the
**5 Methods**

**5.5 Assay steps**

Move Up/Down buttons until it is in the required position in the method. You can make a copy of the currently selected step with the **Copy Step** button.

The **Cycle Run List** button beside the method summary panel allows you to check the number and order of cycles for the method as defined.

Enter the number of cycles to be run in each assay step in the left-hand panel. The list of cycles for the run is displayed in the right-hand panel.

### 5.5.1 Base settings

**Name**

This is the name of the assay step. Each step in a method must have a unique name. New steps are by default named **Assay Step n**, where *n* is a serial number: change the name to something that describes the context or intent of the step, to make the method easier to follow.

**Purpose**

Assay steps are assigned a purpose, used to identify cycles in the evaluation software. The choice of purpose can help to document the method structure, and also determines the way the data is treated in evaluation. Choose the purpose from the list.
An assay step may have one of the following purposes:

<table>
<thead>
<tr>
<th>Purpose</th>
<th>Usage and restrictions</th>
</tr>
</thead>
<tbody>
<tr>
<td>Calibration</td>
<td>Used for calibration curves in concentration assays and affinity in solution. This assay step should be connected to the same cycle type as the <strong>Sample</strong> step so that the calibration and sample analyses are performed in the same way.                                                                                                     Set <strong>Calibration</strong> to recurring within <strong>Sample</strong> to repeat the calibration at intervals through the assay.</td>
</tr>
<tr>
<td>Conditioning</td>
<td>Used to condition the sensor surface at the start of an assay.</td>
</tr>
<tr>
<td>Control sample</td>
<td>Used for control samples. This assay step should be connected to the same cycle type as the <strong>Sample</strong> step so that the control sample and sample analyses are performed in the same way.                                                                                                                     Set <strong>Control sample</strong> to recurring within <strong>Sample</strong> to repeat the control sample analysis at intervals through the assay.</td>
</tr>
<tr>
<td>Sample</td>
<td>Used for sample analysis in all applications. At least one sample step is required for application-specific evaluation.</td>
</tr>
<tr>
<td>Solvent correction</td>
<td>Used for solvent correction cycles. This step should be connected to a cycle type designed for solvent correction.                                                                                                                                                                                                                                                                                                                                                                                        Set <strong>Solvent correction</strong> to recurring within <strong>Sample</strong> to repeat the solvent correction at intervals through the assay.</td>
</tr>
<tr>
<td>Startup</td>
<td>Used to condition the flow system at the start of an assay. This assay step will commonly be connected to the same cycle type as the <strong>Sample</strong> step.</td>
</tr>
<tr>
<td>Undefined</td>
<td>Used for assay steps that do not fit the predefined purposes. Assay steps with <strong>Undefined</strong> purpose will not be included in application-specific evaluation.</td>
</tr>
</tbody>
</table>

**Note:** For simple methods, the assay step name and purpose may often be the same (e.g. Solvent Correction, Sample, Control Sample etc.). It is however important to remember that the name is for documentation from the user’s perspective and may be chosen freely, while the purpose has significance for the step properties and for evaluation of the run and must be chosen from the predefined list.
5.5 Assay steps

Connect to cycle type
Each assay step is connected to one cycle type, which determines the detailed operation of the step (see Section 5.6). Choose the cycle type from the list of types available in the method.

5.5.2 Assay step preparations

Temperature
This value determines the analysis temperature for the assay step. The setting will also control the sample compartment temperature if the appropriate option is checked under General Settings (Section 5.4).

If the actual temperature at the start of an assay step does not match the setting for the step, the system will wait until the setting is reached.

Buffer
Select the running buffer to be used for the assay step. The default buffer is A (corresponding to buffer bottle and tubing A on the instrument).

5.5.3 Recurrence
An assay step can be set to recur at a specified interval within the context of another step. The recurrence can be specified as Every ... cycles (so that the number of occurrences will depend on the number of cycles in the assay step) or Distribute ... occurrences evenly (in which case the number of occurrences is fixed and they are distributed as evenly as possible among the cycles in the assay step). In addition, the recurring step can be specified to be executed at the beginning and/or end of the step in which it is set to recur.

Note: The Run assay step first/last options refer to the beginning and end of the context in which the assay step recurs. To ensure that a recurring assay step is run at the beginning and end of a whole assay with several different assay steps, set up a separate copy of the assay step to run once at the beginning and/or end of the entire assay. The illustration below shows an assay step setup that ensures calibration before the first sample, repeated during the samples and at the end of the assay.

![Assay step setup illustration](image-url)
If the step within which another step recurs is run in replicate, the recurring step is distributed among the total number of cycles including replicates. This is illustrated by the calculations below for a recurring step set to *Every 5 cycles*:

<table>
<thead>
<tr>
<th>Top level step</th>
<th>Number of cycles</th>
<th>Number of replicates</th>
<th>Total number of cycles</th>
<th>Number of recurrences for the nested step</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>10</td>
<td>1</td>
<td>10</td>
<td>2</td>
</tr>
<tr>
<td></td>
<td>20</td>
<td>1</td>
<td>20</td>
<td>4</td>
</tr>
<tr>
<td></td>
<td>10</td>
<td>2</td>
<td>20</td>
<td>4</td>
</tr>
</tbody>
</table>

### 5.5.4 Number of replicates

Assay steps can be set to run in replicate, which means that all cycles in the assay step will be repeated the specified number of times. The order in which cycles in the assay step are repeated can be specified:

- **As entered** performs all cycles in the step once, then repeats the step until the number of replicates is reached (this is represented as 1,2,3,1,2,3 to illustrate the order of 3 cycles in a step repeated twice)

- **Order** performs the first cycle in the step for the specified number of replicates, then the second cycle and so on (represented as 1,1,2,2,3,3)

- **Random** randomizes the order of the cycles within the step until all cycles have been executed the specified number of times. The order is randomized each time the method is run.
5.6 Cycle types

Cycle types define the detailed sequence of operations to be performed in each assay step.

The top panel in the work area lists the cycle types currently available in the method. Use the **New** button to create a new cycle type. Mark a cycle type and click **Delete** to remove the cycle type from the method. Use the **Copy** button to make copies of cycle types in the method; this can be useful if a method requires a number of similar cycle types with small variations. Click **Rename** to rename a cycle type.

Enter a description of the cycle type if desired.

Settings for the currently selected cycle type are configured in the lower part of the work area. The settings are divided into **Commands** and **Report Points**, accessed on the respective tabs.
5.6.1 Commands

The commands in a cycle definition correspond to different kinds of injection of liquid over the sensor surface. To add a new command to the cycle definition, choose the command type from the pull-down list and click **Insert**. The command will be inserted with default parameter settings immediately after the currently marked command (or at the end of the cycle definition if no command is marked). Use the **Up** and **Down** buttons to change the position of the selected command in the cycle, and the **Remove** button to remove the selected command from the cycle definition. Commands are executed from top to bottom in the cycle definition.

**General command properties**

Common features of several commands are check-boxes for **Predip**, **Extra Wash After Injection** and **Stabilization period**.

**Predip** Check this box to dip the needle in a separate position before aspirating the solution to be injected. The predip position will normally contain the same solution as is injected, so that the needle is rinsed briefly to minimize carry-over effects. The same predip position is used for all cycles.

**Extra Wash After Injection** Check this box and specify a wash solution to perform an extra wash of the flow system after the injection. The flow system is washed automatically with buffer after each injection, but an extra wash with a different solution can be included if required. This wash solution does not pass over the sensor surface.

**Stabilization period** Check this box and specify a time in seconds to introduce a delay before the next command is started. This can sometimes be necessary (for example after regeneration steps) to allow the response to stabilize before performing the next injection.
Capture command
This command is intended for injection of ligand over a capturing molecule at the beginning of a cycle. The injected solution, contact time and flow rate can be set as variables.

Sample command
This command is intended for injection of sample containing analyte. Only Sample commands are recognized as analyte injections in the Evaluation Software for kinetics, affinity and concentration evaluation. The injected solution, contact time, dissociation time and flow rate can be set as variables. Evaluation variables can also be defined for the Sample command (see Section 5.6.2).

The Sample command offers 5 alternative settings under Type:

- **High performance**
  Optimizes the injection for high performance by using extra segments of air and sample during aspiration to separate the injected solution from running buffer, thereby minimizing dispersion of sample at the beginning and end of the injection at the expense of additional 25 µl sample consumption.

- **Low sample consumption**
  Optimizes the injection for low sample consumption by using fewer air segments than the High performance setting. The Low sample consumption setting however still achieves a performance that is adequate for most applications except analysis of rapid kinetics.

- **Single cycle kinetics**
  Injects a series of sample concentrations in the same cycle, intended for single-cycle kinetics analysis (see Section B.10). The samples are injected in direct sequence, separated only by the time required to prepare the next injection. A dissociation time is included after the last sample injection. With this option checked, an evaluation variable (see Section 5.6.2) will be set up for each of the specified number of concentrations.

- **Merged injection**
  Performs simultaneous injection of acidified sample and neutralization solution so that the sample is neutralized during passage through the flow cells. This injection type is specifically designed for use in immunogenicity studies, for dealing with drug interference by neutralization of acidified samples immediately before analysis. See the Biacore T200 Immunogenicity Handbook, Chapter 4, for details of how to use Merged injection.
**Double mix**
Sample is first acidified then neutralized by sequential addition of acid and neutralization solution to the sample in the microplate well. The neutralized sample is then injected over the sensor surface. This injection type is specifically designed for use in immunogenicity studies, for dealing with drug interference by analysis of acidified-and neutralized samples. See the Biacore T200 Immunogenicity Handbook, Chapter 4, for details of how to use **Double mix**.

**Note:** The injection types Merged injection and Double mix are specifically designed for dealing with drug interference in immunogenicity studies and should only be used in such studies.

The **High performance** and **Low sample consumption** options support a **Mix** function for mixing sample with a defined solution in the autosampler before injection. Check the **Mix** option and enter a mix solution and mixing fraction to use this function. Entering a fraction of e.g. 20% will mix one part of mixing solution with four parts of sample. The sample and mixing solution are taken from respective positions in the autosampler and mixed in a third position. The option **Stabilization period after mix** allows you to specify a wait period between the mixing operation and injection of the mixed solution. **Mix** is not supported for the **Single cycle kinetics** option.

**Enhancement command**
This command is intended for injection of a secondary enhancement reagent following the sample injection. Enhancement reagents are most commonly used to amplify the analyte response and to confirm the identity of the bound analyte. The injected solution, contact time and flow rate can be set as variables.

**Regeneration command**
This command is intended for injection of a regeneration solution following the sample injection. Check **High Viscosity Solution** if the regeneration solution has a relative viscosity higher than about 3 (corresponding to about 35% glycerol or 40% ethylene glycol at 20°C). This will adapt the solution aspiration and injection procedure for higher viscosity. The injected solution, contact time and flow rate can be set as variables.

**Carry-over control command**
This command injects a 30-second pulse of buffer over the surface at a flow rate of 40 µl/min, in order to check that there is no carry-over of analyte or other material from an injection earlier in the cycle. The injection is suitably placed at the end of the cycle, and can be used in a conditional context (see the **if...then** command) to perform additional buffer injections or regeneration steps if carry-over is detected. A plot of the binding response from a carry-over injection
5 Methods
5.6 Cycle types

against cycle number is created automatically for quality control purposes in the evaluation software (see Section 6.4).

**Solvent correction command**

This command injects a 30-second pulse of solvent correction solution over the surface at a flow rate of 30 µl/min. A solvent correction cycle should contain 4-8 **Solvent correction** commands for the different solvent concentrations used to construct the correction curve (see Section 6.7). **Solvent correction** commands will be correctly evaluated only when they are used in cycle types that are connected to assay steps with purpose **Solvent correction**.

**Inject and recover command**

This command recovers analyte that is bound to the sensor surface, and is intended for use in applications where the bound material is analyzed further. Several features of the command are designed specifically for integration of Biacore analysis with mass spectrometry (MS). Most of the parameters for the command can be set as variables. This command can only be used for injection over all four flow cells: normally, the same ligand should be immobilized in each cell.

The command initiates a sequence of operations in the instrument:

1. The specified volume of **Deposition solution** is transferred in the autosampler to a target position in the sample and reagent rack. Target positions are assigned as required in the **Rack Positions** dialog (Section 4.2.6). The deposition solution should be MS-compatible, and may be used for example to neutralize the recovery conditions (which are often acidic) or to add trypsin or another protease to the sample for peptide digestion. The presence of deposition solution also helps to collect the small recovered volume reliably from the autosampler needle.

2. The **Sample solution** is injected over the sensor surface with the **Contact time** and **Flow rate** as specified. The **Flow path** is fixed so that sample
passes through all four flow cells to maximize the amount of analyte that binds to the surface.

3. The flow system is washed with the specified **Wash solution**. Distilled water or an MS-compatible buffer should be used as washing solution.

4. A small volume (approximately 2 µl) of **Recovery solution**, separated from the surrounding buffer by air segments, is injected into the flow cells. The flow is stopped for the specified **Incubation time** while the recovery solution is in contact with the sensor surface, to allow the bound analyte to dissociate into the recovery solution.

5. The flow direction over the sensor surface is reversed and the recovery solution containing recovered analyte is deposited in the target position where it mixes passively with the deposition solution.

6. Steps 2-5 are repeated for the specified **Number of repetitions**. This increases the yield of recovered analyte without requiring additional commands. The same target position is used for recovered analyte from all repetitions. Note that only one aliquot of deposition solution is used, regardless of the number of repetitions.

**Notes:** Methods that include the **InjectAndRecover** command require a sample and reagent rack and cannot be used with microplates (see the Biacore T200 Instrument Handbook for rack details).

The contact time for sample, flow rate and number of repetitions determine the total injected volume for both sample and recovery solution. You may need to adjust one or more of these parameters if the method does not pass verification.

**General command**

This command is a general-purpose injection that supports the following options under Type:

- **Dual**: Injects two solutions in direct succession, with no intervening automatic wash routines. A dissociation time may be set for the second injection but not for the first.

- **High performance**: Prioritizes high injection performance over sample consumption (see description under **Sample** command).

- **Low sample consumption**: Prioritizes low sample consumption over injection performance (see description under **Sample** command).

**General** commands are not recognized as sample injections for evaluation of concentration, kinetics or affinity evaluation and may therefore also be used to “hide” injections from the predefined evaluation facilities. The injected solution,
5 Methods
5.6 Cycle types

contact time and flow rate can be set as variables. Evaluation variables can also be defined for **General** commands (see Section 5.6.2).
**If...then command**

This command allows construction of conditional methods, where commands are executed or skipped depending on the outcome of certain conditions. The illustration below shows a cycle which will perform an additional regeneration if the relative response after the first regeneration exceeds a specified value:

To set up a conditional command:

1. Insert an **If...then** command at the appropriate place in the cycle definition.

2. Specify the condition. This is defined as the outcome of a comparison between a report point value (absolute response, relative response or slope) and a constant or another report point value with an added or subtracted constant value. Only report points that have already been set in the cycle definition may be used in the condition.

   Check **Use additional condition** to combine two conditions, using either AND (both conditions must be fulfilled) or OR (fulfillment of one condition is sufficient) as a logical operator.

   The available comparison conditions are **Greater than** and **Less than**. The conditions do not include **Equal to** since exact equality is an unpredictable condition in view of noise in the SPR response. To construct an equality condition, combine one **Greater than** and one **Less than** condition so that a window of tolerance is created. For example, the combined condition A greater than B-1 AND A less than B+1 is equivalent to A equals B with a tolerance of ±1.

3. Choose the actions to be taken when the condition is met and when it is not met. You may choose to execute a command sequence, terminate the cycle or the method, or introduce a stabilization period.

   If you choose a command sequence for either the **True** or **False** outcome, click on the appropriate branch of the command (**Then** or **Else** respectively) and insert the commands you wish to be executed. If you leave the branch empty, the cycle will simply continue with the next command following the **If...then** construction.
5 Methods
5.6 Cycle types

You can use the Move up and Move down buttons to rearrange the order of commands within a branch, but you cannot move commands outside the branch in which they are placed.

If you have chosen a command sequence for an action and have entered commands, you must delete the commands before you can change to a different action.

5.6.2 Variables

Parameters for many of the commands in a cycle definition may be set as variables. Values for variables are entered in either the Variable Settings in the method or the Setup Run step when the method is run (Sections 5.7 and 5.9.2), and determine the number of cycles that will be performed in the run. Variables fall into two broad classes:

- **Method variables** such as sample name or contact time control the way the cycle will be performed. Parameter values that are not set as variables are defined in the main command panel.

- **Evaluation variables** such as concentration or analyte molecular weight are used in evaluation of the data. Some evaluation variables are required for correct functioning of application-specific evaluation procedures (for example, kinetic evaluation requires a variable called Conc which holds the analyte concentration). These are selected from a predefined list. Other evaluation variables may be freely defined by the user, to hold information that is relevant to the assay but not required by an application-specific evaluation procedure (an example might be the sample batch number). Evaluation variables may only be defined for Sample and General commands.

Variables are set in the list at the right in the command panel. For method variables, check a parameter to set it as variable. Sample solution is checked by default for Sample commands. For evaluation variables, choose the purpose of the assay to display an appropriate list of predefined variables and check the variables you want to use. Click Add to set up user-defined variables.

**Note:** For specific assay purposes, you should generally check all suggested variables. If you leave some variables unchecked, the assay-specific evaluation may not work.
Predefined evaluation variables for different assay purposes are described in the table below (see also Section 5.10).

**Evaluation purpose: General, Kinetics/Affinity, Thermodynamics**

<table>
<thead>
<tr>
<th>Variable</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Conc</td>
<td>Analyte concentration. Multi-cycle kinetics requires a single concentration variable. Single-cycle kinetics requires one concentration variable for each sample injection in the cycle.</td>
</tr>
<tr>
<td>MW</td>
<td>Analyte molecular weight: required for molecular weight adjustment of report points, and for kinetic evaluation when the concentration is entered in weight-based units.</td>
</tr>
</tbody>
</table>

**Evaluation purpose: Kinetics (heterogeneous analyte)**

<table>
<thead>
<tr>
<th>Variable</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Conc1, Conc2</td>
<td>Analyte concentrations for the two analytes.</td>
</tr>
<tr>
<td>MW1, MW2</td>
<td>Molecular weight for the two analytes: these variables are required even if concentrations are entered in molar units, to determine the relative contributions of the two analytes to the observed response.</td>
</tr>
</tbody>
</table>

**Evaluation purpose: Concentration (using calibration curve)**

<table>
<thead>
<tr>
<th>Variable</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Conc</td>
<td>Analyte concentration: required for calibration and control samples, left blank for unknowns.</td>
</tr>
<tr>
<td>Dilution</td>
<td>Dilution factor: used for unknown samples to calculate original concentrations.</td>
</tr>
</tbody>
</table>
5 Methods
5.6 Cycle types

**Evaluation purpose: Calibration-free concentration analysis**

<table>
<thead>
<tr>
<th>MW</th>
<th>Analyte molecular weight: used in calculation of concentration from binding rates (empty for blank injections).</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>D</em>(20°C)</td>
<td>Diffusion coefficient of the analyte at 20°C (empty for blank injections).</td>
</tr>
<tr>
<td>Blank</td>
<td>Identifies blank cycles for blank subtraction purposes. Blank cycles have the value <em>y</em> or <em>yes</em> (upper or lower case); any other value identifies the cycle as non-blank.</td>
</tr>
</tbody>
</table>

**Evaluation purpose: Affinity in solution***

<table>
<thead>
<tr>
<th>ConcB-calibration</th>
<th>Concentration of interactant B used to construct a calibration curve.</th>
</tr>
</thead>
<tbody>
<tr>
<td>ConcB-fix</td>
<td>Concentration of interactant B in the sample mixture (the concentration of B is kept constant).</td>
</tr>
<tr>
<td>ConcA-variable</td>
<td>Concentration of interactant A in the sample mixture (the concentration of A is varied).</td>
</tr>
</tbody>
</table>

*See Section 11.1.1 for details of how this assay is set up.

5.6.3 Report points

The Report points tab lists the report points in the cycle type definition, ordered as far as possible in the order they will appear in the cycle. Several injection commands have a predefined set of report points that are added to the list when the command is included in the cycle type. You can add your own report points by filling in the details in the empty row at the bottom of the table. A new empty row is added whenever you create a report point.
Report points are set at defined times in relation to the start or end of injections in the cycle. Report points that are set outside the time range for the cycle (i.e. a significant time before the start of the first injection or after the end of the last injection) will not be created.

*Note:* Do not position report points far away from events, so that they lose their relevance to the event, or so close to an event so that the report point window overlaps the event itself.

Enter the report point details as follows:

- **Name**: The report point name must be unique within the cycle type. Choose a name that reflects the function of the report point.

- **Sec**: Enter the time in seconds between the report point and the event.

- **Before/After**: Choose whether the report point is to be placed before or after the event.

- **Start of/End of**: Choose whether the report point is to be placed relative to the start or end of the injection.

- **Inject**: Choose the injection to which the report point will be related.

- **Window**: Set the window for the report point calculation. The report point will be placed at the center of the window, and the reported response will be an average of the response values within the window. A window of 5 seconds is adequate for most purposes.

- **Baseline**: Choose whether the report point will be defined as a baseline or not. Response values for report points that are not defined as a baseline will be calculated relative to the closest preceding baseline value.
5.7 Variable settings

This screen determines whether variable parameter values are specified in the method or at run time. You may choose to specify all values in the method, all at run time, or a mixture of the two. Values that are specified in the method are saved with the method and cannot be changed at run time. You can change the level at which values are specified without changing the cycle type definition, so that the same cycle type can be used for different assay steps with different sets of run-time variables. An example of this may be found in the predefined method for affinity in solution (Section B.1).

Variables are configured independently for the different assay steps (even if the assay steps use the same cycle type).

If you choose to specify values for all variables in the method, the values are entered in this screen. For each assay step, one row of variable values represents one analysis cycle (the cycle may be repeated if the Repeat property is set in the Assay Step screen). Each row in the variables table corresponds to a cycle in the run. A new empty row (marked with an asterisk) is created automatically at the bottom of the table as soon as data is entered. Columns in the table correspond to variables for the cycle type used in the assay step, and are grouped according to commands in the cycle type definition. Use the right mouse button in the variables table to access functions for copying and pasting cell contents and for inserting and removing rows. When all variables are specified at run time, variables are handled in the same way in the Setup Run step.

To specify that some variables are specified in the method and others at run time, check the appropriate option and then distribute the variables as required.
between the method and run-time lists. This mode can be used to hide variables at run time that are not relevant for the assay step. Fill in values for variables that are specified in the method: those specified at run time are filled in the Setup Run step. Note that in this mode only one value can be given for each variable that is specified in the method. These values will be used for all cycles: the number of cycles is determined by the number of rows of variable values in Setup Run.

Depending on how the method is defined, there may be variable tables for several assay steps. Variable handling must be defined for all steps before the method will pass verification.

5.8 Verification

This step checks that the method is correctly and completely defined. A method that does not pass the verification step can be saved but cannot be run. Verification may fail because parameters are missing (e.g. variables that are specified in the method have not been assigned values) or because the method construction is invalid (e.g. an assay step is not connected to a cycle type).

Note: The verification step does not check whether the run fulfils the requirements for any assay-specific evaluation (see Section 5.10).

5.9 Setup Run

5.9.1 Detection

Set the flow path for the method in the Detection dialog.

You can only choose a flow path that is consistent with the Detection setting for the method (see Section 5.4).

5.9.2 Variables

In this step you assign values to variables that are to be defined at run time (typically a sample table, see Section 5.7). Each row in the variables table corresponds to a cycle in the run. A new empty row (marked with an asterisk) is created automatically at the bottom of the table as soon as data is entered.
Columns in the table correspond to variables for the cycle type used in the assay step, and are grouped according to commands in the cycle type definition.

Use the right mouse button in the variables table to access functions for copying and pasting cell contents and for inserting and removing rows. The columns in the table are listed in the order they are defined in the method (see Section 5.6.2).

Click **Import** to import the variables table from an external file. See Appendix A for details of the import format.

Depending on how the method is defined, there may be variable tables for several assay steps. Method variable values must be entered in all tables before you can continue to the next step. Evaluation variables may be left blank if desired at this step and values entered in the Evaluation Software.
5.9.3 Cycle run list

When you have completed the variables table, click **Next** to view a complete summary of the cycles that will be performed in the run. This view is for information only and cannot be edited. Check through the cycle list to confirm that the variable tables are correctly filled in.

![Method Builder - Cycle run list](image)

Click **Overview** to display the method overview (Section 5.3).

5.9.4 System preparations

Choose which preparation steps should be executed before the method starts.

![Method Builder - System Preparations](image)
5 Methods
5.10 Requirements for assay-specific evaluation

System preparations are equivalent to those for wizard-based runs (see Section 4.2.5). The temperature settings are taken from the first assay step in the method and cannot be changed here: they are shown for information only.

5.9.5 Rack positions
See Section 4.2.6 for a description of the Rack positions dialog box.

5.9.6 Prepare Run Protocol
See Section 4.2.7 for a description of the Prepare Run Protocol dialog box. Save the method at this step to include all settings including the Prepare Run protocol in the saved method.

5.9.7 Starting the run
When the positions are finalized, the sample plate is prepared and loaded into the instrument and a chip is docked, choose Start Run to start the run. You will be asked to specify a file name for the results.

5.10 Requirements for assay-specific evaluation
This section describes the requirements and recommendations if assay-specific evaluation is to be applied to method-based runs.

5.10.1 Concentration analysis
See Chapter 8 for a description of concentration evaluation.

Using calibration
- At least one assay step is required with purpose Calibration and one with purpose Sample. An assay step with purpose Control Sample is also required in order to create trend plots for control samples.
- Assay steps Calibration, Sample and Control Sample must be connected to a cycle type that includes one Sample command. The three assay steps will normally be connected to the same cycle type.
- Samples in the Calibration step must have concentrations specified in the variable Conc. At least two different concentrations are required for linear calibration curves and at least four for 4-parameter fitting.
- Full use of the Calibration trends function (Section 8.2.2) requires calibration curves run before the first sample cycle and after the last.
- Samples in the Sample step will normally not have specified concentrations. If concentrations are specified, they will be ignored.
**Calibration-free**

- At least one assay step is required with purpose **Sample**, connected to a cycle type that includes one **Sample** injection with flow rate set to **Variable**.
- Each sample must be injected at least twice using different flow rates.
- Evaluation variables **D**(20°C), **MW** and **Blank** must be included for the **Sample** injection.

**5.10.2 Kinetics/Affinity**

See Chapter 9 for a description of kinetics and affinity evaluation.

- At least one assay step is required with purpose **Sample**, connected to a cycle type that includes one **Sample** command.
- Sample concentration must be specified in the variable **Conc**. If weight-based units are used, a molecular weight for the analyte must be specified in the variable **MW**.
- As a recommendation for multi-cycle kinetics and affinity, there should be a concentration series with at least four non-zero analyte concentrations and one zero concentration. At least one of the non-zero concentrations should be measured in duplicate. Although kinetic and affinity evaluation can be applied to runs with fewer sensorgrams, the results will generally be less reliable if these recommendations are not followed.
- Five sample concentrations are recommended for single-cycle kinetics and affinity, injected in order of increasing concentration. Duplicate cycles are recommended to ensure robust evaluation. For each determination, there should be a blank cycle corresponding to the sample cycle, with buffer replacing the sample for each injection.

**5.10.3 Thermodynamics**

See Chapter 10 for a description of thermodynamics evaluation.

Thermodynamics evaluation requires that kinetics or affinity (see Section 5.10.2) is determined at two or more (recommended 5) temperatures.

**5.10.4 Affinity in solution**

See Chapter 11 for a description of affinity in solution evaluation.

- At least one assay step is required with purpose **Calibration** and one with purpose **Sample**. Both assay steps must be connected to a cycle type that includes one **Sample** command. The two assay steps will normally be connected to the same cycle type.
5 Methods
5.10 Requirements for assay-specific evaluation

- Samples in the Calibration step must have concentrations specified in the variable ConcB-calibration. At least two different concentrations are required for linear calibration curves and at least four for 4-parameter fitting. These samples should contain only component B.

- Samples in the Sample step must have concentrations specified in the variables ConcB-fix and ConcA-variable. At least 3 samples with the same concentration of component B mixed with different concentrations of component A are required.

5.10.5 Immunogenicity

Evaluation of immunogenicity experiments is described in the separate Biacore T200 Immunogenicity Handbook.

- For evaluation of specificity confirmation by addition of excess drug, sample analysis cycles must use the option to mix samples with a specified solution in the autosampler. For correct evaluation, each sample should be analyzed twice, once mixed with drug and once with buffer.

- For evaluation of isotyping, isotyping reagents must be injected with the Enhancement command.

5.10.6 Other requirements

Application of solvent correction (see Section 6.7) requires an assay step with purpose Solvent Correction, connected to a cycle type that includes at least four Solvent Correction commands for different solvent concentrations.
Evaluation Software
Evaluation software – general features

Biacore T200 Evaluation Software offers general functions for presentation of results as sensorgrams, report point plots and bar charts, and for evaluation functions for specific applications such as concentration, kinetics and affinity, thermodynamics and immunogenicity. There is also a function that corrects for solvent effects that can sometimes distort the results from analyses with low molecular weight analytes that give low response levels and require organic solvents (e.g. dimethyl sulfoxide (DMSO)) to maintain solubility.

This chapter describes the organization of the evaluation software. The various evaluation functions are described in detail in the following chapters.

6.1 User interface

6.1.1 Organization

The Biacore T200 Evaluation Software screen is divided into four main regions:

- The **menu and toolbar** provide access to the evaluation functions.
- The **Evaluation Explorer** at the left of the screen lists the evaluation items (sensorgrams, plots, bar charts and other result displays) that have been created in the current session. See below for a description of how to use the Evaluation Explorer.
6 Evaluation software – general features
6.2 Basic operations

- The work area displays the currently open items. Each item is shown in a separate window that can be moved, resized or closed independently of the other items.

- The status bar indicates the progress of current operations such as application of evaluation methods.

6.1.2 The Evaluation Explorer

The Evaluation Explorer lists the sensorgram windows, plots and other evaluation items in the current evaluation session. Items are organized in folders according to type. Double-click on a folder to expand or collapse the folder. Click on an item to display it in the work area. Right-click in the explorer area for options for adding new items: right-click on an item for additional options relating to that item. You can also remove or edit selected items using the Remove and Edit buttons at the top of the explorer panel.

You can hide the Evaluation Explorer panel to increase the useful size of the work area by clicking on the pin icon in the explorer title bar. If the panel is hidden, it will automatically reappear when you move the mouse over the left-hand edge of the evaluation window.

6.2 Basic operations

6.2.1 Opening files

To open a file in the evaluation software, use the File:Open menu option. You can open result files from the control software (file extension .blr) and saved evaluation sessions (file extension .bme). If a file is already open in the software, opening a new file with File:Open will automatically close the first file. You will be asked if you want to save the file if any changes have been made.

Opening a file automatically creates and opens an All sensorgrams item, and creates a number of default plots according to the content of the file.

To open multiple result files in the same session, use the File:Append Result File option. You can only append result files, not evaluation sessions. Appending a file to a session will delete all the user-defined evaluation items in the session.

Note: The software does not check that appended result files contain compatible data. Beware of appending files from different types of run (e.g. kinetic and concentration analyses) or from the same type of assay run under different conditions (e.g. concentration assays using different sensor chips).
6.2.2 Printing evaluation results

Choose **File:Print** to print a hard-copy of the results. Select the printer to use and choose the items you wish to print.

![Print dialog box](image)

6.3 Common display functions

6.3.1 Zooming the display

To zoom a display window, drag with the mouse over the area you want to enlarge. To restore the previous zoom level, double-click anywhere in the display window except on the axes or legend, or select **Unzoom** from the right-click menu.

Displays are normally rescaled automatically whenever you change the displayed data. To keep the current zoom setting when data is changed, check **Zoom lock** in the display window.

You can also set the display scale with the **Scale** option from the right-click menu. The display is not rescaled automatically if the scale has been set to specified values.

6.3.2 Right-click menus

Right-click in display windows for options relating to the display. The available options vary according to the type of window, and also depending on whether you right-click on a point, a curve or elsewhere in the window.
Labels
Displays a label on each point in a plot window, showing cycle number, flow cell and sample name. (Labels may overlap and be difficult to read if the points in a plot are closely spaced.)

Caption
Displays a caption in the item window. The displayed caption can have system defined and user-defined components.

Show sensorgram
Displays the sensorgram relating to a point in the plot. This option is only available when you right-click on a point: the sensorgram is displayed in a separate window that must be closed before you can continue with the evaluation.

Exclude cycle/curve/point
Excludes data from the evaluation session. The data that can be excluded differs according to the type of window. Excluding cycles in sensorgram, plot or bar chart windows affects all other sensorgram, plot and chart windows correspondingly; however, application-specific evaluation items that have already been created are not affected until they are edited and updated. If you have applied solvent correction and then exclude solvent correction cycles, the correction remains applied, but the excluded cycles will not be available if you edit the solvent correction item.
**Scale**

Sets the scale for the display. You can also access this function by double-clicking on either the x- or y-axis in the display.

![Scale Settings](image)

Uncheck *Auto* and enter minimum and maximum values to specify a scale.

**Copy graph**

Copies the current display as it appears on the screen to the Windows clipboard as a graphic object, from where it can be pasted into third-party software such as word-processing or presentation programs.

**Copy table**

Copies the table data in the current display to the Windows clipboard as tab-separated text. All rows in the table are copied (including any header row) regardless of how many rows are visible in the current display. However, in tables where the columns to be displayed can be selected by the user, only columns that are currently shown are copied.

**Export curves**

Exports the curves in the current display to a tab-separated text file, for import to third-party software. Complete data is exported regardless of the scale setting of the screen display.

**Unzoom**

Restores the previous zoom setting.

**Gridlines**

Shows or hides major and minor gridlines in the display window.

**Legend**

Shows or hides a legend for the display window. Choose the legend placing from the dialog box. In sensorgram and plot windows, the legend corresponds to the *Color by* setting for the display.
6 Evaluation software – general features
6.4 Predefined evaluation items

6.4 Predefined evaluation items

When a result file is opened, a number of evaluation items are created automatically if the results contain the appropriate cycles and report points. This section describes the items created for all result files: special items are also created for certain wizard runs (see Chapter 4).

6.4.1 Sensorgram
An overlay plot of all sensorgrams is created and opened. The sensorgrams are not aligned, and are colored by assay step.

6.4.2 Plots
Plots are created for most wizards if the appropriate report point is present in the results. The settings for predefined plots are locked and cannot be edited. Common predefined plots are listed below. Separate plots will be created if there are multiple injections with similar report points (for example baseline for capture and sample injections). The Plot Settings cannot be changed, but the plot can be modified using the selectors and the Tools menu (see Section 7.2). Note that changing the selector settings can sometimes defeat the purpose of the plot.

| Baseline: Sample | Absolute response for report point baseline against cycle number. |
| Baseline: Capture | Absolute response for report point capture_baseline against cycle number. |
| Baseline: General | Absolute response for report point general_baseline against cycle number. |
| Binding to reference | Relative response for report point stability against cycle number for the reference flow cell. |
| Capture | Relative response for report point capture_level against cycle number for the capture injection. |
| Carry-over | Relative response for the report point co_binding against cycle number for the carry-over injection (only for reference-subtracted curves). |
| Controls, binding | Relative response for the report point binding against cycle number for control samples (only for reference-subtracted curves). |
6.5 Custom report points

Report points are automatically created for all wizard- and method-based runs, placed at strategic positions in relation to injections. These report points, and any other report points that have been created in the Control Software (Section 2.3.3), are not listed in the Custom Report Points dialog and cannot be edited or deleted in the Evaluation Software.

Choose **Tools:**Custom Report Points to add and edit custom report points.

<table>
<thead>
<tr>
<th>Controls, stability</th>
<th>Relative response for the report point <em>stability</em> against cycle number for control samples.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Binding levels</td>
<td>Relative response for the report point <em>binding</em> against cycle number for samples (only for reference-subtracted curves).</td>
</tr>
<tr>
<td>Binding stability</td>
<td>Relative response for the report point <em>stability</em> against cycle number for samples.</td>
</tr>
<tr>
<td>Binding to reference, enhancement</td>
<td>Relative response for report point <em>enhance_level</em> against cycle number for enhancement injections on the reference surface.</td>
</tr>
<tr>
<td>Enhancement</td>
<td>Relative response for the report point <em>enhance_level</em> against cycle number for enhancement injections on the active surface.</td>
</tr>
</tbody>
</table>
6.5 Custom report points

6.5.1 Adding report points

Click **Add** to add a new report point.

Enter a name for the report point (maximum 30 characters) in the **Id** field. The report point name must be unique within an evaluation session.

Enter a value between 1 and 35 for the **Window**. Preset values of 5, 10 and 15 seconds are provided for convenience; a window of 5 seconds is recommended for most purposes. The response value for the report point is the average response over the window, with the report point time at the center of the window.

Use the settings under **Position the report point** to define where the report point will be placed. Report points are placed a specified number of seconds before or after the beginning or end of injections or the beginning or end of the cycle.

**Notes:** Do not position report points far away from events so that they lose their relevance to the event, or so close to an event so that the report point window overlaps the event itself.

You cannot define a report point with settings that would place part of the window outside the time limits of the sensorgram.

Check **Calculate response relative to report point** and select the required report point if you want to calculate the responses at the custom report point relative to another report point. If the box is not checked, the closest preceding baseline report point will be used for calculating relative response values.
You can apply custom report points either to cycles with selected assay step purposes or to cycles selected by cycle number. Choose the appropriate option and check the assay step purpose(s) or cycles to which the report point should apply.

**Note:** If you append a result file to the evaluation session after creating custom report points, the custom report points are retained but they are not applied to sensorgrams in the appended file.

### 6.5.2 Editing and deleting report points

Select a report point in the list in the **Custom Report Points** dialog and click **Edit** to edit the report point definition or **Delete** to delete the report point.

**Notes:** All user-defined evaluation items are deleted from the session when custom report points are edited or deleted, since changes to the report point definitions may affect existing evaluation items. You will be warned when this occurs. To avoid losing work, save your evaluation session before editing or deleting custom report points.

*If you delete a custom report point that is used as baseline for other report points, the relative response can no longer be calculated for the latter points. You will be warned if this situation arises.*

### 6.6 Keywords

Keywords are assigned to cycles when the run is performed, and are then used for identification and evaluation purposes. Keywords are created automatically for wizard-based runs and may be defined in the method for method-based runs, and include:

- automatically generated identifiers such as cycle number or assay step purpose,
- method variables and predefined evaluation variables such as sample name, concentration and molecular weight,
- user-defined variables (see Section 5.6.2).

You can add and remove user-defined keywords in the evaluation software, and edit the contents of certain keywords. Choose **Tools:Keyword Table** to open the keyword table. When you save changes to the keyword table, all user-defined items in the evaluation session will be deleted. Save the session before editing the keyword table if you do not want to lose your work. Click **Cancel** in the **Keyword Table** dialog to close the dialog without applying changes and deleting user-defined evaluation items.
To simplify management of the keyword table, you can sort and filter the table display:

- Click on a column header to sort the table by the contents of that column.
- Click in the filter row (directly below the column header) and select a value to display only rows with that value for the chosen column. Click **Reset All Filters** to restore all filters to the [All] setting.

To change a keyword value, simply enter the new value in the appropriate cell. Values for some system-generated keywords (such as **Assay Step Purpose**) are chosen from a predefined list of values: the list is displayed when you click in such a cell.

To change the units for concentration keywords, choose a new unit from the **Concentration Unit** list. This changes the unit but not the numerical value of the keyword. For example, a concentration entered as 10 µM will become 10 mM if you change the concentration unit to mM. If the evaluation session includes data from multiple files, a table of concentration units for the different files is displayed. Make sure that the unit is the same for all files if data are to be evaluated together.

**Note:** The concentration unit affects only predefined concentration keywords. Numerical user-defined keywords are simply numbers, and will not be re-interpreted when you change the concentration unit even if they are intended to hold concentration information.

Click **Add Keyword** to create a new keyword in the table. You can choose between predefined keywords and user-defined keywords (see Section 5.6.2). If there are multiple **Sample** or **General** commands in the method from which the data is obtained (see Section 5.6.1), specify the command to which the new keyword should apply.
Enter the required keyword values in the empty column that is created for the new keyword.

To rename or delete a keyword, click the appropriate button, then select the keyword in the dialog box. You cannot remove system-generated keywords such as file number or cycle number, or keywords derived from method variables such as sample name.

6.7 Solvent correction

6.7.1 Background

Solvent correction adjusts reference-subtracted responses for small artefacts that can be introduced by variations in the bulk refractive index between samples. The correction is only relevant when variations in the bulk refractive index are of the same order of magnitude as the response: this situation arises commonly in work with small organic analytes that give intrinsically low response values and that often require organic solvents such as dimethyl sulfoxide (DMSO) to maintain solubility.

The need for solvent correction arises because subtraction of the reference response does not exactly eliminate the contribution of the bulk solution to the measured response. Bulk solution is excluded from the volume occupied by ligand on the active surface, so that the bulk contribution to the response on the active surface is slightly smaller than that on the reference surface (Figure 6-1).
Solvent correction

As long as the refractive index of the samples is constant, this excluded volume effect introduces a constant error in reference subtraction which may be ignored for practical purposes. However, if the refractive index of the samples varies, the magnitude of the excluded volume effect will also vary.

Organic solvents like DMSO often give a high bulk response (addition of 1% DMSO gives a bulk response of about 1200 RU), so that small variations in the DMSO content lead to significant variations in the bulk response between samples. Such variations are unavoidable in the preparation of diverse samples such as drug candidates for screening applications. The solvent correction procedure corrects for the variations arising from the excluded volume effect in these cases.

A more detailed description of solvent correction background and procedures may be obtained from GE Healthcare.

6.7.2 When solvent correction should be used

It is important to bear in mind that solvent correction is only relevant when

- the expected analyte responses are low,
- the ligand is a macromolecule immobilized at a high density (typically 5,000 RU or more – lower ligand densities lead to excluded volume effects that are too small to merit correction),
- the bulk response is subject to variations between samples of at least the same order of magnitude as the measured binding response.

Solvent correction should not be applied in situations that do not meet all three of these criteria. Attempts to use solvent correction in other circumstances may introduce errors that are larger than the solvent effects that the procedure is intended to correct.
6.7.3 How solvent correction works

Solvent correction factors are determined by injecting a series of blank samples containing a range of solvent concentrations over the active and reference surfaces, and plotting the difference in relative response between the surfaces as a function of the relative response on the reference surface. Each sample measurement is then corrected by a factor obtained by measuring the relative response on the reference surface and reading the corresponding difference between active and reference surfaces from the correction curve (Figure 6-2).

![Figures 6-2](image)

**Figure 6-2. The principle of solvent correction.** 1. The sensorgram from the reference flow cell shows a bulk displacement (-150 RU in the illustration) during sample injection because the sample and running buffer are not exactly matched. 2. From the solvent correction curve, a displacement of -150 RU in the reference sensorgram corresponds to a solvent error of +5 RU in the reference-subtracted sensorgram. 3. The reference-subtracted sensorgram is corrected by subtracting the solvent error. This procedure is applied to every point during sample injection.

Solvent correction is applied only to response levels during sample injection, since the correction adjusts for differences in the bulk refractive index of the samples compared with running buffer. Solvent correction is meaningless when buffer is flowing over the surface before and after the sample injection.

6.7.4 Applying solvent correction

To apply solvent correction, choose *Add solvent correction* from the *Evaluation* menu. In order to apply solvent correction, the run must include solvent correction cycles (see Section 5.6.1).
6.7 Solvent correction

The left-hand panel of this dialog lists the solvent correction curves in the run, and the corresponding curves are shown in the right-hand panel. For each solvent correction cycle, there is one correction curve for each set of reference-subtracted sensorgrams. All cycles are shown by default in an overlay plot. Select specific rows in the cycle list to display the corresponding solvent correction curves. Clear the checkmark in the Included box to exclude cycles from the correction calculation. You must include at least one solvent correction cycle for each curve. Sample responses are corrected according to the curve obtained from the nearest preceding correction cycle in the run. If there is no preceding correction cycle, the nearest following cycle is used.

Examine the curves for fitting to the experimental points. Right-click on outlying points to exclude either the single point from the curve fit or the whole correction cycle from the correction process. Statistical fitting parameters (chi-squared) are shown for each correction curve in the cycles list. Right-click on a point or curve in the right panel and choose Show sensorgrams if you want to examine the sensorgrams from solvent correction cycles as an aid in judging the quality of the data.

The solvent correction curve is fitted to the experimental points using a second-degree equation. Beware of applying solvent correction if the correction curve does not fit the experimental points closely. Scatter in the correction points indicates that the measurements are not reliable, and applying correction derived from such curves can distort the measured responses unnecessarily. For reliable solvent correction, the chi-squared value should be less than 2 RU.

**Note:** In judging the quality of the solvent correction data, take note of the y-axis scale in the display. The curves are automatically scaled to fit the window.
If the range of solvent correction is small (as in the illustration above),
points may appear to scatter around the fitted curve without necessarily
indicating poor curve quality.

The shape and slope of the solvent correction curve (even the direction of slope)
may vary between measurements on different occasions. This is normal and the
shape of the curve should not be taken as an indicator of curve quality.

The range of report point values that are candidates for solvent correction in the
assay data is indicated by vertical red lines in the window. If report points lie
outside the correction range, these values cannot be properly corrected. Some
small extrapolation of the correction plots may be permissible. Use the
Extrapolate button to extend the correction range. The shape of the solvent
correction plots is however not fully predictable, and extrapolation over more
than about 10% of the range of the reference values is dangerous.

Click OK to apply the solvent correction. Correction will be applied to the sample
and carry-over injection phase(s) of all sensorgrams. Any data points that lie
outside the correction range will be discarded and the corresponding
sensorgram will contain gaps corresponding to the invalid data.

Note: In assays where the temperature is varied during the run, such as
thermodynamics assays, you should make sure that there is a solvent
correction curve included at the start of each temperature series.
Correction will only be applied to sample measurements made at the
same temperature as the correction curve.

6.8 Evaluation methods

Evaluation methods allow you to save the definitions of all evaluation items
(except Thermodynamics) in an evaluation session, and apply them
automatically to the same or different result files or evaluation sessions. Use this
feature to apply standardized evaluation procedures to different result files, or
to re-create an evaluation session after operations that delete user-defined
items such as changes made in the custom report points or keyword table (see
Sections 6.5 and 6.6).

6.8.1 Creating evaluation methods

To create an evaluation method, choose File:Save Evaluation Method As and
specify a file name and location. The method is saved with file extension
.evalmethod.

An evaluation method may only contain one definition of a kinetics/affinity
evaluation item. If there are more than one item in the evaluation session, or if
an item contains more than one fitting, you will be asked to choose which item
should be represented in the method.
6 Evaluation software – general features
6.8 Evaluation methods

Click **Skip** to create the method with no kinetics/affinity item included.

Calibration-free concentration analysis items that have different fitting ranges for different samples (Section 8.3) and kinetic/affinity items that use data sets with multiple $R_{\text{max}}$ (Section 9.2.2) cannot be saved in evaluation methods.

### 6.8.2 Applying evaluation methods

To apply an evaluation method to the contents of an evaluation session, choose **File:Apply Evaluation Method** and choose the method. A preview of the method is shown so that you can check the method contents.

Click **Apply** to apply the method. Items in the method will be created where possible. Items that cannot be created will be reported in a dialog box.
7 Data presentation tools

This chapter describes the tools available for presentation and examination of the data in a result set. These tools comprise:

- Sensorgram display, with facilities for aligning sensorgrams in overlay plots.
- Plot tool for displaying and ranking response values.
- Bar chart for displaying response values.
- A report point table for listing numerical values associated with report points.

7.1 Sensorgram items

Sensorgram items display sensorgrams from one or more cycles in the results. A sensorgram item showing all sensorgrams for the first reference-subtracted curve in the results (or first active curve if reference subtraction is not used) is created automatically when the result file is opened. You can change the display settings in this item, or create additional sensorgram items if required. Click Sensorgram in the toolbar or choose Add sensorgram from the Evaluation menu to add a new sensorgram item.

Hold the cursor over a sensorgram to display a tool tip identifying that particular curve. The sensorgram coloring can be changed if desired with the Tools:Color by option.

The following sections describe display functions specific to sensorgram items. General display functions are described in Section 6.3.
7 Data presentation tools
7.1 Sensorgram items

7.1.1 Selecting sensorgrams for display
The selector bar at the top of the window controls which sensorgrams will be displayed.

- **Curve name** lists the type of sensorgram (active, reference, reference subtracted and solvent corrected where applicable).
- **Assay Step Purpose** filters the sensorgram according to the assay step purpose.
- **Cycle** lists all the cycles in the result set. When multiple files are open, cycles are identified with two numbers, one for the file in the result set and one for the cycle within the file (thus cycle 1-10 is the 10th cycle in the first file added to the result set, cycle 2-4 is the 4th cycle in the second file and so on).

For each display controller, click the browse buttons to browse backwards or forwards through the list, one item at a time.

Click the selector button to open the list for selecting one or more items. Drag with the mouse or use shift-click to select contiguous multiple items. Use control-click to select non-contiguous multiple items. To accept a selection, click anywhere outside the list or press Enter.

7.1.2 Removing data
To remove data from the display, mark the section to be removed by dragging with the right mouse button, then choose **Cut** from the right-click menu. The data will be removed from the current sensorgram display item only; no other windows or evaluation items will be affected. This function can be useful for removing injections with high bulk contributions (such as regeneration injections) or other visual disturbances from the display.

Choose **Undo Cut** from the right-click menu to restore the removed data.
7.1.3  Sensorgram adjustment

Choose **Sensorgram adjustment** under the **Tools** button for options for aligning and adjusting the sensorgram display. For curve alignment, sensorgrams that do not include the chosen reference point for alignment will not be shown. Sensorgram adjustment only affects the display in the current sensorgram item.

**X-adjustment**

Choose to set the zero time point to either a report point or an injection event. If this setting is **Off**, the zero time point will be at the beginning of the cycle.

**Y-adjustment**

Choose to set the zero response point to either a report point or an injection event. If this setting is **Off**, the actual response values will be shown.

If you check **Enable Second Y-Adjustment**, you can select a report point or injection event where the response value will be set to 100. Each sensorgram will then be normalized separately to the first and second adjustment point, so that all sensorgrams will have values of 0 and 100 at these points regardless of the original response levels. This can help in comparing the shapes of sensorgrams independently of their response levels, or in adjusting response levels that are dependent on others (e.g. adjusting analyte response for varying capture levels, by adjusting the baseline to 0 and the capture level to 100).
**Blank subtraction**

Check *Enable Blank Subtraction* and choose a curve to be used as the blank to subtract one sensorgram from all others in the display. Use this feature to eliminate systematic disturbances in sensorgrams that are not removed by reference subtraction. Blank subtraction only affects the current sensorgram window: other evaluation items are not affected.

**Note:** Subtracting a blank sensorgram is not the same as using reference-subtracted data. Reference subtraction gives the difference between active and reference values for each cycle separately for each curve, whereas blank subtraction subtracts one curve from all others in the result set.

### 7.1.4 Markers

You can choose to display markers and/or labels for report points and events in the cycle with the **Report points** and **Event markers** options respectively under the **Tools** button. Report points are displayed on the curve and event markers on the x-axis.

![Diagram](image-url)
7.2 Plot items

Plot items display report point values plotted against either variables or other report point values in the same cycle. Curves can be fitted to points using linear or 4-parameter fitting functions. Ranking borders can be added to plot items to classify response levels into groups such as high, medium and low responses.

To create a plot item, click Plot in the toolbar or choose Add Plot from the Evaluation menu. Enter a name for the plot, choose the parameters that define the plot and click Finish. Cycles that do not contain the selected report point will not be represented in the plot.

Response type may be response (absolute or relative response, or relative response adjusted for molecular weight if the keyword MW is defined) or sensorgram slope at the selected report point. Adjustment for molecular weight is performed by dividing the response in RU by the molecular weight in Da. Points for which the molecular weight value is zero or missing are omitted from the plot.

Variables may be numerical (e.g. molecular weight or concentration) or non-numerical (e.g. sample name or assay step purpose).

The plot will be created with default display settings, with a graphical representation at the left and a table of x-and y-values at the right. Tool tips identify the data points (place the cursor on a point for a couple of seconds to display the tool tip).
Right-click on the plot item in the evaluation explorer and choose *Rename* to change the plot name.

### 7.2.1 Selector functions

The selector bar at the top of the window controls which points will be displayed.

- **Curve name** lists the type of sensorgram from which the points are taken (active, reference, reference subtracted and solvent corrected where applicable).
- **Assay Step Purpose** filters the points according to the assay step purpose.
- The third selector lists the variable values represented on the x-axis. (This option is not available for plots of report point against report point.)

Selection operates in the same way as in the sensorgram window (Section 7.1.1).

Other general display functions are described in Section 6.3.
7.2.2 Table functions

The table to the right of the plot area lists values for the points in the plot. Excluded points are shown struck out in red text. You can exclude or include cycles from the right-click menu in the table area, in the same way as from the right-click menu in the plot. The table also allows you to exclude or include multiple cycles in a single operation.

Select rows in the table to highlight the corresponding points in the plot. If you select a single row, the highlight is augmented with lines drawn to the plot axes:

By default, the table shows x- and y-values and is sorted in ascending order of x-values. Click on the header row to select the sort value and to change the sort order. Sorting the table does not have any effect on the plot display.

Choose Tools: Table columns to select columns that will be displayed in the table. You can also change the order in which columns will be displayed using the Move up and Move down buttons (the top of the column list represents the left-hand column in the table).
7 Data presentation tools
7.2 Plot items

7.2.3 Sorting the plot
Plots of report point values against variables can be sorted in order of ascending or descending y-axis value, regardless of the variable chosen for the x-axis. A sorted plot can be useful for example in visualizing the frequency of different levels of response, which may be more difficult to see if the levels are scattered more or less randomly with respect to the variable parameter defined for the x-axis. Choose Sort:Ascending or Descending under the Tools button to sort the plot. Sorting the plot also sorts the rows in the table, although the table can be sorted independently of the plot by clicking in the table column header (Section 7.2.2).

Note: By default, the table associated with a sorted plot retains a column labeled X-Value. This is the value of the variable originally defined for the plot, and does not correspond to the x-axis as displayed in the sorted plot.

Choose Sort:As Defined to restore the x-axis to the originally defined variable value.

Plots of one report point against another cannot be sorted.

7.2.4 Fitting curves to points
Choose Curve Fitting under the Tools button to fit lines to the points in the plot. using either linear or curved (4-parameter) fitting functions. If Fit by color is checked, each color will be fitted to an independent line. If this option is not checked, all points derived from the same curve will be fitted to a single line. The numerical fitting results are displayed in an extra panel below the plot. Choose Curve Fitting:Curve parameters under the Tools button to toggle display of this extra panel.
For linear fitting, the points are fitted to the equation

\[ y = \text{slope} \times x + \text{intercept} \]

The equation for a 4-parameter fit is

\[ y = \frac{R_{hi} - R_{lo}}{1 + \left( \frac{x}{A_2} \right)^{A_1}} \]

where \( y \) and \( x \) are the plot coordinates

\( R_{hi} \) and \( R_{lo} \) are fitting parameters that correspond to the maximum and minimum response levels respectively

\( A_1 \) and \( A_2 \) are additional fitting parameters

The closeness of fit is reported for linear fitting as the coefficient of determination \( R^2 \) and for 4-parameter fitting as the chi-squared value (see Section 9.7.1).

Curve fitting cannot be applied to plots that have been sorted.
7.2.5 Adjusting plots for controls

Plots can be adjusted for systematic changes in response during the course of the assay, such as progressive loss of binding capacity resulting from harsh regeneration conditions. Adjustment is calculated from the response values obtained for control samples analyzed at intervals during the assay.

Adjustment for controls cannot be applied to plots of one report point against another.

To apply this adjustment, choose Adjustment for controls from the Tools menu in the plot window. Check Use adjustment for controls and select the sample to use as a positive control. You can also select a sample for the negative control. Select whether the adjustment should be made using a linear or polynomial (second degree) fitting function. The display panels in the dialog show a plot of the response against cycle number before and after adjustment. Click OK to apply the adjustment.

Adjustment normalizes the sample responses relative to the positive and negative control levels as follows:

- Curves are fitted to the control sample responses for positive and (if used) negative controls. The Linear option fits the points to a function with the form \( y = ax + b \) (where \( a \) and \( b \) are constants). Polynomial fits the points to a second-degree function with the form \( y = ax^2 + bx + c \) (where \( a \), \( b \) and \( c \) are constants). If no sample is chosen as a negative control, 0 RU is used as the negative control response.

- The fitted line(s) are transformed to straight horizontal lines with values 100 for the positive control and 0 for the negative control.
• The transformation used to create straight horizontal lines for the control points is applied to all points in the plot (including the actual control sample responses) so that each point retains the same position relative to the positive and negative controls before and after adjustment.

Adjustment for controls cannot be applied in regions where the positive control curve lies below the negative control curve (or below the x-axis if no negative control is selected). Any points that lie in such regions will be excluded from the adjusted plot.

Adjustment for controls applies only to the current plot and does not affect any other evaluation item.

Note: Beware of using a polynomial fitting function with less than 4 control samples. The parabolic curve created by the function can deviate greatly from the points, leading to adjustment that does not reflect the drift in the control responses (Figure 7-1).

Figure 7-1. Polynomial function fitted to 3 control points (highlighted) with the resultant adjustment for controls.
7 Data presentation tools
7.2 Plot items

7.2.6 Ranking

Choose **Ranking** under the **Tools** button to add ranking boundaries to the plot. You can add one or two boundaries, classifying the plot points as **Low/High** or **Low/Medium/High**. The boundary values are specified in RU.

Boundaries are shown as horizontal red lines in the plot, labeled with the boundary value. The classification of the points is recorded in the table. Ranking results are independent for each plot item.

**Note:** Editing the definition of a plot does not affect ranking boundaries. If you for example change the y-axis parameter of a ranked plot from relative response to absolute response, you will need to revise the placing of the ranking boundaries if the ranking is to remain meaningful.
7.3 Bar chart items

Bar charts display report point data plotted against cycle number. Unlike plots, bar charts can display multiple report points in the same chart, and can group the display in various ways.

To create a bar chart, click Bar Chart in the toolbar or choose Add Bar Chart from the Evaluation menu. The chart is created directly. Numerical values on which the chart is based are listed on the Table tab.

7.3.1 Selector functions

Select the curves to display in the lower left hand panel, cycles in the center panel and report points in the right-hand panel. In each panel you can select multiple rows by dragging with the mouse or using Shift-click for contiguous rows or Control-click for non-contiguous rows. Click on the column header in the table of cycles in the center panel to sort the table by the contents of the column, to simplify selection of rows according to the required criteria.

Select whether to display relative or absolute response values in the Response field. This selection applies to all report points shown in the chart. Points for which the selection is invalid are not shown (for example, the report point baseline will normally not be shown on a bar chart of relative response).
7 Data presentation tools
7.4 Report point table

7.3.2 Display options
Drag with the mouse to zoom the bar chart display. Double-click to restore the original scaling.

Use the Tools menu to determine how the bar chart is organized and displayed:

Group by
You can group bars in the chart by various parameters such as curve, cycle, report point or sample name. Groups are separated by vertical lines in the chart. Grouping is most useful when you have multiple selections in more than one selection panel: the examples below illustrate bar charts showing two report points, grouped by cycle (left) and by report point (right).

Color by
Select a parameter in the Color by list to color the bars in the chart according to the parameter value. A Color key will be added to the table in the appropriate panel.

Show column labels
Use this option to display labels identifying the bars in the chart. The bars are also identified by tool tips, regardless of whether labels are displayed or not.

7.4 Report point table
Report points are automatically created for all wizard- and method-based runs, placed at strategic positions in relation to injections. See Section 6.5 for details of how to add custom report points.

7.4.1 Displaying the report point table
The report point table lists numerical values for all report points in the current result set. The report point table is created automatically as an evaluation item. Each evaluation session can only have one report point table item: the table is updated automatically if you add custom report points or apply solvent correction.
The contents of the report point table cannot be edited.

The following columns can be represented in the report point table. Click the Table Columns button to choose which columns to display.

<table>
<thead>
<tr>
<th>Column</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>File</td>
<td>File number. This column is only shown when the evaluation session includes more than one file, the cycle number is prefixed with a file number. Choose File:Properties to display the mapping of source files to file numbers.</td>
</tr>
<tr>
<td>Cycle</td>
<td>Cycle number within the file.</td>
</tr>
<tr>
<td>Fc</td>
<td>The curve to which the report point applies, identified as the flow cell.</td>
</tr>
<tr>
<td>Report point</td>
<td>Report point id.</td>
</tr>
<tr>
<td>Time (s)</td>
<td>Report point time in seconds from the start of the cycle.</td>
</tr>
<tr>
<td>Window (s)</td>
<td>Report point time window in seconds.</td>
</tr>
<tr>
<td>AbsResp (RU)</td>
<td>Absolute response in RU, calculated as the mean value over the time window.</td>
</tr>
</tbody>
</table>
### 7 Data presentation tools

#### 7.4 Report point table

| **SD** | Standard deviation of data points in the time window, calculated as  
\[
SD = \sqrt{\frac{1}{n-1} \sum (y - \bar{y})^2}
\]
where \( n \) = number of points and \( y \) = response in RU |
| **Slope (RU/s)** | Slope during time window in RU s\(^{-2}\), calculated as  
\[
slope = \frac{\sum (y - \bar{y})(x - \bar{x})}{\sum (x - \bar{x})^2}
\] |
| **LRSD** | Alignment of slope to a straight line (regression coefficient), calculated as  
\[
LRSD = \sqrt{\frac{Q_0}{n(n-2)}}
\]
where  
\[
Q_0 = \frac{\sum (y - \bar{y})^2 - \left( \sum (y - \bar{y})(x - \bar{x}) \right)^2}{\sum (x - \bar{x})^2}
\] |
| **Baseline** | Yes for report points defined as baseline. Otherwise No. |
| **RelResp (RU)** | Relative response (difference in absolute response from the baseline) in RU. #N/A if no baseline has been set. |
| **AssayStep Purpose** | Identifiers for the cycles, set explicitly in methods (see Chapter 5) and automatically in wizards. |
| **CycleType** | |
| **Keywords** | One column is created for each keyword in the data. |

#### Sorting and filtering the report point table

The report point table can be sorted by any column in ascending or descending order for any column. Click in the header for a column to sort the table by that column. Each click in the same header toggles the sort order.
The second row in the report point table contains a filter setting for each column. All values will be included if the filter setting is blank. To apply a filter, click on the setting and select a value from the list. The value will be shown in the filter setting and only rows in the table that contain the value in the selected column will be displayed. You can apply multiple filters to the table at the same time. To remove a filter, choose All from the list of column values in the filter setting.

**Copying report point table contents**

To copy the selected contents of the report point table, select cells by dragging with the mouse and press Ctrl-C or choose Copy from the right-click menu. Choose Copy Table to copy the whole table. The contents will be copied in tab-separated text format to the Windows clipboard, and can be pasted from there into other programs. All selected cells will be copied, including header cells and filter settings.
7 Data presentation tools
7.4 Report point table
8 Concentration analysis

Biacore T200 software supports two approaches to concentration analysis:

- Using a calibration curve determined by measurement of known samples (referred to as calibrated measurements).
- Calibration-free measurements that rely on the relationship between the analyte concentration and the rate of mass transfer of analyte to the sensor surface.

Principles and experimental practice for concentration measurements are described in the Biacore Concentration Analysis Handbook.

8.1 Requirements for concentration evaluation

8.1.1 Calibrated measurements

In order to evaluate concentration analysis using a calibration curve, the run must include at least one calibration curve and unknown sample injections must have the appropriate properties and keywords. For wizard-based runs, the conditions are met automatically when the Concentration Analysis wizard is used. For method-based runs, the method must be constructed as described in Section 5.10.1: if necessary, the keyword table can be edited so that the conditions are met in full (see Section 6.6). Note however that the command name for the sample injection cannot be edited in the keyword table. Refer to Chapter 5 for details of how to construct methods in Method Builder.

Unknown samples may be evaluated in three ways:

- Based on the nearest preceding measured calibration curve, or the nearest following curve if there is none preceding.
- Based on a single calibration curve constructed as an average of all measured curves.
- Based on individual calibration curves constructed for each separate cycle, using a calibration trend plot to interpolate calibration data between the measured points. This option is useful if the calibration responses show a steady drift during the course of the assay.

8.1.2 Calibration-free measurements

Runs for calibration-free concentration analysis require a minimum of two cycles for each sample, run at different flow rates. Blank cycles for each flow rate are recommended. Both samples and blanks must be injected with a command of type Sample (see Section 5.6.1). No calibration standards are required.
Correct values for the diffusion coefficient $D$ at 20°C and the molecular weight $MW$ must be provided. These values are normally provided when the run is set up in the Control Software, but can be added or edited in the keyword table in the Evaluation Software.

### 8.2 Evaluating calibrated concentration analyses

To evaluate a concentration analysis run, open the run and choose *Concentration Analysis: Using calibration* from the evaluation toolbar, or choose *Add Concentration Analysis: Using calibration* from the *Evaluation* menu. The evaluation dialog is divided into three tabs, for calibration curves, control samples and unknown samples respectively. Choose the appropriate settings for the calibration curves and click *Finish* to complete the evaluation.

#### 8.2.1 Calibration curves

A calibration curve is constructed from the cycles in each calibration step. If two calibration steps are run in direct succession so that there are no other cycles between the steps, they will be combined in a single calibration curve.

Settings on the *Calibration* tab define the report point and fitting function that are used to create the calibration curve. The settings apply to all calibration curves in the evaluation. Choose the settings appropriate for your analysis:
**Flow cell**  Concentration analysis may be performed without a reference cell, since the unknown samples are determined by direct reference to a calibration curve obtained under the same conditions. If a reference cell is included in the flow path for the run, the reference-subtracted curve may however be selected if desired.

**Report point**  Response levels for concentration analysis (relative response) are normally taken from a report point shortly after the end of the sample injection, to avoid contributions from the bulk refractive index of the sample. For analyses based on the rate of binding (report point slope), a report point early in the sample injection is normally used.

**Response type**  Choose between relative response and slope.

Check **Use average calibration curve** to use one average curve constructed from all calibration points in the run.

Check **Use calibration trends** to create individual calibration curves for each cycle in the assay by interpolation between the actual measured calibration curves. See Section 8.2.2 for details of calibration trends.

The left-hand panel lists the details of the calibration curve data, with concentration and response or slope values for all calibration points. The **Calc. Conc.** column lists the concentration corresponding to the actual response value as determined from the fitted calibration curve. Select a row in the table to highlight the corresponding point on the calibration curve or vice versa. Fitting parameters are shown at the bottom of the left-hand panel when a single calibration curve is displayed.

The calibration curves are shown in the right-hand panel. Each calibration curve is constructed from the cycles in an assay step with the purpose **Calibration**. Right-click on a curve or calibration point for options for excluding calibration curves or single cycles from the evaluation, and for displaying sensorgrams corresponding to individual calibration cycles. Excluded cycles are shown as open symbols and are marked in the left-hand panel with red strikethrough text. If you exclude a calibration curve and then re-include sufficient points to allow a new curve to be fitted, the curve will still be excluded but will be shown as a broken line.
8 Concentration analysis
8.2 Evaluating calibrated concentration analyses

You can choose the curve to display from the list at the top of the panel. The options differ according to how the calibration curves are used:

- For **Use preceding calibration curve**, the panel shows one calibration curve by default. Individual curves multiple curves can be selected from the list.
- For **Use average calibration curves**, the panel shows all calibration points with a single average curve.
- For **Use calibration trends**, the panel shows all measured calibration curves in an overlay plot.

**Note:** The sample name entered for the calibration samples is used only as a title for the calibration curve. For wizard-based runs, the sample name will be the same for all calibration cycles, but different names can be introduced either in method-based runs or by editing the keyword table. If the points in a calibration curve have different sample names, all points will still be used and the title will be shown as **Mixed analytes**.

Choose the **Fitting Function** for the calibration curve from the option above the curve panel. Linear and 4-parameter functions and provided with the software, and custom models can be defined if required (Section 8.2.5). The 4-parameter function is a general fitting function for continuous curves, and is recommended for most purposes. Use a linear function only if you have good reason to expect the calibration curve to be a straight line. See Section 7.2.4 for the equations for fitting functions.

8.2.2 Calibration trends

Calibration trends represent the stability or otherwise of the calibration curves during the course of the assay, and can be used to compensate for drift in the calibration responses by constructing an interpolated calibration curve for each individual cycle.

**Examining calibration trends**

Open the **Calibration trends** tab to display the trends as a plot of calibration points against cycle number. Trend lines are fitted through each set of points with the same calibrant concentration, using a linear function for trends with two points and second-degree polynomial function for trends with three points or more. Actual measured points are listed in the table at the left.
If calibration curves are run in replicate (i.e. using the same concentration in two successive calibration curves with no intervening cycles of another type), the trends are fitted to points representing the average of replicate measurements.

Right-click on points in the plot or rows in the table for options relating to the specific calibration sample.

**Using calibration trends**

Check *Use calibration trends* on the *Calibration* tab to create an individual calibration curve for each cycle in the assay by using interpolated calibration points from the calibration trend plot. The illustration below shows how the calibration curve is constructed for cycle 25.

When calibration trends are used, the *Calibration* tab shows an overlay plot of the measured calibration curves. Note however that none of these measured curves is used directly for calibration: an individual (virtual) calibration curve is constructed for every cycle including the calibration cycles themselves.

The trend lines are extrapolated to cover the cycles in the first and last calibration cycles so that every cycle has a complete virtual curve. However, if
you exclude calibration points at the ends of the trend lines, the trend lines will not be extrapolated and calibration curves in the regions outside the excluded range may be incomplete (see illustration below).

![Calibration Curve](image)

**Note:** When *Use calibration trends* is selected, results are recalculated whenever points are excluded from or included in the trend lines, which can take some time for large runs. For best performance, establish the trend lines before choosing *Use calibration trends*.

### 8.2.3 Control samples

The *Control Samples* tab displays the measured concentration for control samples as a plot of response against cycle number.

![Control Sample Plot](image)

Numerical results are presented in the table at the left, and plotted as calculated concentration against cycle number on the right. Select a row in the table to highlight the corresponding point on the plot.
The table lists the expected concentration as entered for the control samples, the response and calculated concentration, the calculated concentration as a percentage of expected and the calibration curve used to calculate the concentration. Replicate control samples are summarized with average values and coefficient of variation (CV%) for the response and calculated concentration.

Right-click on a sample row in the table or a sample point in the plot and choose **Exclude Cycle** to exclude that sample from the sample evaluation. Excluded cycles are shown as open symbols and are marked in the table with red strikethrough text.

### 8.2.4 Samples

The **Samples** tab displays the measured concentrations for samples.

The left-hand panel lists the results sorted by sample ID, with averages and CV values for each sample ID. The column **Calc.Conc.** gives the concentration calculated for the original sample, obtained as the measured concentration multiplied by the dilution factor. Concentrations for samples that give a response outside the range of the calibration curve are listed as below or above the limits of the calibration.

**Note:** The limits of calibration are defined by the concentrations corresponding to the response values on the fitted curve for the highest and lowest calibration samples. Depending on how well the curve fits the experimental points, these limits may not coincide exactly with the actual concentrations in the highest and lowest samples.
8 Concentration analysis
8.2 Evaluating calibrated concentration analyses

The right-hand panel shows the calibration curve for the currently selected sample. All sample points calculated from the curve are shown in black; calibration points are shown in orange. Select a row in the table to highlight the corresponding point on the plot and vice versa.

**Note:** If *Use calibration trends* is selected in the Calibration tab, each sample is calculated from its own individual calibration curve, shown in the right-hand panel with a single sample point.

Right-click on a sample row in the table or a sample point on the curve and choose *Exclude Cycle* to exclude that sample from the average sample calculation. Excluded cycles are shown as open symbols and are marked in the table with red strikethrough text.

8.2.5 Custom models for calibration curves

You can define your own fitting models for calibration curves in concentration analysis, for example to support validated assay procedures that do not use linear or 4-parameter calibration fitting. Choose *Tools:Models:Concentration* to define or edit fitting models. Models for calibration curves are defined using similar principles as equation models for kinetics and affinity (see Section 9.9.3).
8.2.6 Evaluating combined result sets

If you use the Append File function to combine result sets from separate runs, concentration analysis can be evaluated provided that the conditions specified in Section 5.10 are fulfilled in the combined set. The software does not check the validity of any evaluation applied to a combined result set, so that it is your responsibility to determine that the evaluation results are meaningful. It is for example in principle possible to append a kinetic analysis result file to a concentration analysis, and then apply concentration analysis evaluation. Provided that the report point used for the calibration curve also exists in the cycles from the kinetic run, calculated concentrations will be reported for these samples.

In order to ensure that concentration analysis is correctly evaluated in combined result sets, make sure that all files that contribute to the combined result set are derived from concentration analysis runs. Provided that each file starts with a calibration curve that is not excluded from the evaluation, the results will be calculated within the respective files even in the combined result set. However, if calibration curves at the beginning of files are missing or excluded, there will be overlap between the individual file sets and some samples from one file will be evaluated on the basis of calibration curves from another file. In such cases, it is important to ensure that the calibration curves and sample analyses in the different files refer to the same analyte and are performed under as far as possible identical condition.

Note: Beware of using calibration trends with appended result files. If the appended files use the same calibration concentrations, trends will be fitted over the whole data set and will be valid only if the response levels from the different files form a continuous function. The trends will differ according to the order in which the result files are appended.
8.3 Evaluating calibration-free measurements

Calibration-free concentration analysis calculates the analyte concentration from the measured mass transport properties and values for the diffusion coefficient and molecular weight, provided as evaluation variables when the assay is run (Section B.2). The evaluation is based on fitting the sensorgram data to a model of 1:1 interaction kinetics, with mass transport parameters calculated from the supplied diffusion coefficient and molecular weight, and with the analyte concentration set as a globally fitted variable.

To evaluate calibration-free concentration analysis, choose **Concentration Analysis:Calibration-free** from the evaluation toolbar, or choose **Add Concentration Analysis:Calibration-free** from the **Evaluation** menu.

8.3.1 Selecting samples

Choose the samples you want to evaluate in the first dialog:

*Sample and cycle list*

The table in the top panel lists the samples available for analysis. By default, the table shows a full list of cycles including details of the flow rate and subtracted blank cycle (see below) for each sample cycle. Remove the checkmark from **Expand all cycles** to collapse the cycle details and show only samples in the table.
Concentration analysis

Notes: Selecting a row in the cycle list does not display the corresponding cycle in the sensorgram panel. Use the selector bar in the sensorgram panel to view different samples.

The value shown for the diffusion coefficient \( D \) is adjusted to the analysis temperature, and may not be the same as the value for 20°C as entered when the assay was run.

The option **Use reference subtracted data** is checked by default if a reference surface is included in the run setup. You should generally perform the evaluation using reference-subtracted data, but you can uncheck this option to use unsubtracted data if you wish.

Remove the check-mark from the **Include** column to exclude a sample from the evaluation, or from the **Cycle#** column in the expanded list to exclude individual cycles. Remember that each sample must be represented by at least two cycles with different flow rates for the evaluation to be possible. The initial binding rate should be clearly higher at the higher flow rate, and should be reliably measurable (in practice above about 0.2–0.3 RU/s) at the lower flow rate.

It is important that the sensorgrams at the lowest and highest flow rates (recommended 5 and 100 µl/min respectively) are sufficiently separated. If the curves are close together or coincide, this indicates that there is not sufficient mass transport limitation in the binding for reliable concentration measurement. The table column **QC ratio prel** provides a guidance value for this assessment: sufficient mass transport limitation is indicated by a value for the QC ratio of about 0.2 or higher. You may exclude samples with low QC ratios before continuing with the evaluation, or include such samples in the data processing and judge the final results for acceptability.

<table>
<thead>
<tr>
<th>Include</th>
<th>Cycle</th>
<th>Sample</th>
<th>Assay</th>
<th>Dilution factor</th>
<th>F (µl/min)</th>
<th>Temp (°C)</th>
<th>Time (min)</th>
<th>QC ratio prel</th>
</tr>
</thead>
<tbody>
<tr>
<td>✓</td>
<td>1</td>
<td>E100</td>
<td>1</td>
<td>1</td>
<td>25</td>
<td>10000</td>
<td>1</td>
<td>0.965</td>
</tr>
<tr>
<td>✓</td>
<td>2</td>
<td>E100</td>
<td>2</td>
<td>1</td>
<td>25</td>
<td>10000</td>
<td>1</td>
<td>0.965</td>
</tr>
<tr>
<td>✓</td>
<td>3</td>
<td>E100</td>
<td>3</td>
<td>1</td>
<td>25</td>
<td>10000</td>
<td>1</td>
<td>0.965</td>
</tr>
<tr>
<td>✓</td>
<td>4</td>
<td>E100</td>
<td>4</td>
<td>1</td>
<td>25</td>
<td>10000</td>
<td>1</td>
<td>0.965</td>
</tr>
<tr>
<td>✓</td>
<td>5</td>
<td>E100</td>
<td>5</td>
<td>1</td>
<td>25</td>
<td>10000</td>
<td>1</td>
<td>0.965</td>
</tr>
<tr>
<td>✓</td>
<td>6</td>
<td>E100</td>
<td>6</td>
<td>1</td>
<td>25</td>
<td>10000</td>
<td>1</td>
<td>0.965</td>
</tr>
<tr>
<td>✓</td>
<td>7</td>
<td>E100</td>
<td>7</td>
<td>1</td>
<td>25</td>
<td>10000</td>
<td>1</td>
<td>0.965</td>
</tr>
</tbody>
</table>
The QC ratio is calculated as follows from the quotient Q which reflects the degree of mass transport limitation:

\[
\text{quotient } Q = \frac{\text{initial rate at high flow rate}}{\text{initial rate at low flow rate}} \times 3^{\frac{\log \text{low flow rate}}{\log \text{high flow rate}}}
\]

Under conditions of complete mass transport limitation, the binding rate is proportional to the cube root of the flow rate, so the quotient Q has a value of 1. When there is no mass transport limitation, the binding rate is independent of the flow rate so Q has a value equal to the cube root of the flow rate ratio. The range of possible theoretical values for Q will thus depend on the flow rates used (for flow rates of 5 and 100 µl/min, the value is 0.37). The QC ratio is calculated from the measured value for Q normalized to a scale of 0–1:

\[
Q_{\text{max}} = 1
\]

\[
Q_{\text{min}} = 3^{\frac{\log \text{low flow rate}}{\log \text{high flow rate}}}
\]

\[
\text{QC ratio} = \frac{Q_{\text{measured}} - Q_{\text{min}}}{Q_{\text{max}} - Q_{\text{min}}}
\]

The **Initial rate** and **QC ratio** reported in the Select Samples step are preliminary values based on the sensorgram data from 7.5 to 12.5 seconds after the start of the injection. This value may be misleading if the sensorgrams are disturbed. If data is removed from this region (see below) so that initial rates cannot be calculated, the **QC ratio prel** is listed as nd (not determined). A final value, determined from the fitted curves rather than the experimental data, is presented when the evaluation is complete.

Occasionally, the measured binding rate may be lower at the higher flow rate, leading to a negative value for the QC ratio. This can occur if the sensorgrams are disturbed or as a result of experimental variation when the binding rate is not affected by flow rate (so the binding rates should in theory be equal and the QC ratio should be 0).

**Sensorgram display**

The lower panel shows the set of sensorgrams for a selected sample. Select the sample to display with the selector bar. By default, this panel shows data after subtraction of blank cycles (see below). Uncheck **Show blank subtracted data** to see the unsubtractected data and the corresponding blank sensorgrams (shown in gray).

**Note:** The evaluation will always use blank-subtracted data where possible, even if this first dialog shows unsubtractected data.
With the option *Show original sensorgrams* checked, the sensorgram panel shows the full sensorgrams for the sample injection. Uncheck this option to show the data that will be used in the fitting. The range of data used is indicated by blue vertical lines, and any data removed from the sensorgrams is not shown.

**Blank subtraction**

Blank cycles are identified by the value *y* or *yes* in the *Blank* keyword (Section B.2), and must have the same flow rate and injection time as the sample cycles from which they are subtracted. If there is no preceding blank cycle, the nearest following one is used. If there is no blank cycle available for one or more of the cycles in a sample series, the evaluation will be performed using data without blank subtraction for that series.

Click *Included blanks* to display an overlay plot of blank cycles in the run.
8 Concentration analysis
8.3 Evaluating calibration-free measurements

Remove the check-mark from the **Include** column to exclude a blank cycle from the evaluation (for example if the sensorgram shows unreasonable response levels or unacceptable disturbances).

**Setting fitting ranges and removing disturbed data**

Click Advanced settings if you want to adjust the range of data that will be used in the evaluation and/or remove disturbances from the data.

Choose whether you want to make the changes on all sample series in one operation or on selected single sample series. You cannot separate the individual sensorgrams in a sample series.

To adjust the fitting range, drag the blue vertical lines that mark the limits of the range. The default range is from 10 seconds before the start of the injection to 5 seconds before the end. Including baseline data before the start of the injection helps to ensure robust evaluation of the data.
8.3.2 Performing the evaluation

When you are satisfied with the data selection, click Next to start the evaluation. Results appear in the table as each sample series is completed. The evaluation progress is shown in the sensorgram panel.

The fitting procedure is normally fast. If fitting appears to hang for a particular sample series, you can click Accept to accept the current fitting results for that series or Abort current to abandon the series. Click Abort remaining if you want to abandon all remaining series in the evaluation. The fit status is recorded in the table of results.

Click Expand all cycles in the finished result display to display the details of the sample series in the cycle list.

The results are reported in terms of Measured Conc, which is the value calculated from the fitting, and Calc Conc which is obtained by multiplying the Measured Conc by the dilution factor to give the concentration in the original sample. You can change the units for the reported concentration in the header for the Measured Conc column.

**Note:** If you adjust the fitting range differently for different sample series, the range indicators will not be shown when you display All sample series. Evaluation items with different ranges for different sample series cannot be saved in evaluation methods (Section 6.8).

To remove disturbed data, drag with the right mouse button to select the data, then click Remove Selection. Click Undo to restore deleted data.
8 Concentration analysis
8.3 Evaluating calibration-free measurements

**Notes:** The diffusion coefficient $D$ listed in the evaluation results is the value at the analysis temperature, calculated automatically in the software from the value at 20°C provided when the assay is run.

The Initial rate and QC ratio in the finished evaluation are calculated from the fitted curves and are therefore not affected by sensorgram disturbances or data removal.

*If the fitting fails*
If the sensorgrams appear to be satisfactory on visual inspection and quality assessment but the evaluation fails to fit the experimental data, it may help to adjust the initial values for fitting parameters. To do this, step back to the Select samples dialog, open the Advanced settings and click Parameters. Try in the first place changing the starting value for Conc by several orders of magnitude, either up or down. A typical example of a failed fitting is shown below (this was obtained using an initial value for Conc of $1e^{-3}$ where a value of $1e^{-6}$ is more appropriate).

8.3.3 Interpreting the results
Consider the following aspects of calibration-free concentration analysis when interpreting the results:

- Check the appearance of the sensorgrams and fitted curves. Reject samples where the fit is poor at one or both of the flow rates. Use the Chi2 (chi-squared) value as a guideline for fitting quality; this value should be low in relation to the maximum measured response. As a rough guideline, acceptable chi-squared values should be less than about 5% of the response reached at the end of the sample injection at the lowest flow rate (disregarding the difference in units between chi-squared and response).

- Check the QC ratio. Treat the results with caution if the value is lower than about 0.2.
- Check the value for **SE (Conc)** or **T (Conc)**. This value represents the statistical significance of the calculated concentration, and is shown as standard error (SE) or T-value (see Section 9.4.2) according to the setting in Tools:Preferences. Reject samples where the standard error is more than about 20% of the calculated concentration (or correspondingly the T-value is lower than about 5).

- The optimal concentration range that can be measured with calibration-free concentration analysis is 0.5-50 nM. Values for **Meas. Conc** outside this range should be treated with caution.

### 8.3.4 Fitting model

Evaluation of calibration-free concentration relies on fitting the data to a model for 1:1 interaction (see Section 9.8.1) where the mass transport coefficient is provided (through calculation from the diffusion coefficient, and flow cell characteristics) and the analyte concentration is evaluated as a global variable. Although this model is equivalent to the model for evaluation of 1:1 kinetics in terms of the interaction description, the model for calibration-free concentration analysis cannot be accessed through the model editor.

**Notes:** The diffusion coefficient for the analyte at the analysis temperature is calculated automatically from the value at 20°C. The value at 20°C is provided as an evaluation variable when the assay is run (Section B.2).

*Parameters for calculating the mass transport coefficient from the diffusion coefficient are fixed in the software.*
8 Concentration analysis
8.3 Evaluating calibration-free measurements
Biacore T200 offers three main functions for analysis of interaction kinetics and affinity:

- Kinetics and affinity measurements on the sensor surface, which determine the interaction characteristics between ligand and analyte. Kinetic parameters are evaluated from the association and dissociation phases of the sensogram, and affinity either from the kinetic parameters or from plots of steady-state analyte binding levels ($R_{eq}$) against concentration.

- Thermodynamic analysis, which relies on measurement of either kinetics or affinity at different temperatures.

- Affinity in solution, where the interactants are mixed in known concentrations in solution and allowed to reach equilibrium. Biacore T200 is then used to determine the concentration of free interactant in equilibrium with the complex.

Kinetic constants derived from surface interaction measurements in multiple separate experiments can be collated and summarized in the separate Biacore T200 Kinetic Summary software (Section 9.6).

This chapter describes how to evaluate surface-bound kinetics and affinity. If there are multiple sample series in the same data set, evaluation can be performed either in single mode (where each sample series is evaluated separately with interactive control over several aspects of the evaluation), or in batch mode (where multiple series are evaluated automatically using default settings).

Thermodynamic analysis and measurement of affinity in solution are described in Chapters 10 and 11 respectively.

Kinetics and affinity are normally determined from the binding characteristics of a series of analyte concentrations. These concentrations may be injected in separate cycles with surface regeneration between the cycles (multi-cycle analysis) or sequentially in a single cycle with no regeneration between injections (single-cycle analysis), as illustrated in Figure 9-1. Results from these two approaches are evaluated in the same way, using the same tools and fitting models, and may even be evaluated together in a single evaluation. Significant differences in evaluation appearance between the approaches are illustrated where relevant in this chapter.
9 Kinetics and affinity analysis

9.1 Requirements for kinetics and affinity evaluation

The minimum requirements for evaluation of kinetics or affinity are one cycle with a Sample injection in an assay step with purpose Sample, and with the sample concentration in the keyword Conc. If the concentration is not given in molar-based units, the keyword MW must also be included with a value for the molecular weight. For method-based runs, the method must be correctly constructed as described in Section 5.10.2: if necessary, the keyword table can be edited so that the conditions are met in full (see Section 6.6). Note however that the command type cannot be edited in the keyword table. Refer to Chapter 5 for details of how to construct methods in Method Builder.

The recommended minimum conditions for kinetic and affinity evaluation are:

- a concentration series of analyte with at least four non-zero concentrations
- at least one blank cycle consisting of zero concentration sample (for single-cycle kinetics the blank cycle must replicate the sequence of injections in the analysis cycle)
- for multi-cycle kinetics, duplicate determinations for at least one non-zero concentration.

These conditions are recommended but not mandatory in the Kinetics/Affinity wizard.

Figure 9-1. In multi-cycle kinetics and affinity determinations, each sample is injected in a separate cycle. The concentration series is presented as an overlay plot aligned at the start of the injection in the evaluation software. In single-cycle determinations, the samples are injected sequentially in the same cycle. Arrows in the illustrations mark the start of sample injections.
9.2 Evaluating kinetics and affinity in single mode

9.2.1 Basic procedure

To start a kinetics or affinity evaluation in single mode, choose Surface-bound kinetics/Affinity from the Kinetics/Affinity button on the toolbar or the Add Kinetics/Affinity in the Evaluation menu. Check the Single mode option in the first dialog.

1. The first dialog presents the concentration series available in the current result set and allows you to choose the curves included in the evaluation.

A concentration series is defined by a set of curves with the same sample name, analysis temperature and curve identity. Select the concentration series you want to work with in the respective pull-down lists.

**Note:** For runs with immobilized ligand, there will be only one choice for Ligand name. Multiple choices may be available if the ligand is captured and varies between cycles.

For single-cycle kinetics, the cycles table lists all concentrations injected in each cycle. You can however only include or exclude whole cycles at this stage.
9 Kinetics and affinity analysis
9.2 Evaluating kinetics and affinity in single mode

Note: Sample names are case-sensitive, so that “Sample” and “sample” belong to different concentration series. Edit the sample names in the keyword table if you have unintentionally mixed upper- and lower-case letters.

If the result set contains data from more than one file, curves with the same ligand, sample name, temperature and curve identity are grouped together in a single concentration series.

Use the include column in the table of curves to choose which curves should be included in the data set to be evaluated. You can select several curves and use the right-click menu to exclude or include multiple curves in one operation. By default, all curves for the sample are included.

Sensorgrams for non-zero concentrations are shown in color, and those for blanks (zero concentrations) in light gray. The sensorgrams are adjusted to zero at the start of the sample injection on both the response and time axes. The average of the blank sensorgrams will be automatically subtracted from the other curves when you proceed to the next step. If you do not want to perform blank subtraction, exclude the zero concentration sensorgrams from the data set. You can also choose to use blanks from other concentration series for blank subtraction; these are listed at the bottom of the table, and are excluded by default. Only blank sensorgrams with the same contact and dissociation times as the samples are used.

The three check-boxes below the sensorgram panel control the type of curves shown in the display. You can use these check-boxes to examine the sample and blank curves without interference from each other, and to show the average blank that will be used for subtraction. Bear in mind however that these boxes control the display only and do not affect the data set that will be evaluated.

If you have multiple ligand densities represented in the result set, click Multiple Rmax to assign curves to the different sets (see Section 9.2.2).

Click Adjust Injection Events if you want to adjust the injection start and end positions for the evaluation. These positions are set automatically from the event markers in the run, but may need slight adjustment for best fitting to fast interaction processes. The adjustment compensates for small systematic discrepancies in the interval between the event as recorded in
the event log and the time that the sample actually reaches the detection spot on the sensor surface. The difference is most apparent at low flow rates.

The event markers for injection start and end in the result file are shown on the x-axis. Drag the vertical reference lines to adjust the injection start or end point for evaluation. You can adjust the events by \( \pm 10 \) s from the original position. The same adjustment is applied to all curves in the data set, whether they are currently included for evaluation or not. This function cannot be used for single-cycle kinetics, for evaluations with mixed single- and multi-cycle analyses, or for multi-cycle analyses with different contact times in different cycles.

**Note:** If you adjust the injection start, the time axis in the evaluated data is adjusted correspondingly so that the start of the injection is always at time zero. The display in the *Adjust injection events* dialog however always shows the original time axis, with zero at the event marker in the run data.
The second dialog shows the blank subtracted curve set and allows you to delete selected regions from all or selected curves, for example to eliminate spikes or other disturbances.

To delete a selected region from all curves, drag with the right mouse button over the region to be deleted and click **Remove Selection**. Click **Undo** to restore the deleted data.

If you want to delete a region from only selected curves, remove the checkmark from the **Edit** column in the table for the curves that are to be left unchanged. All curves are selected by default and are shown in dark color. Curves that are not selected for editing are shown in light color. Note that all curves will be evaluated, whether they are selected for editing or not: removing the **Edit** checkmark does not exclude a curve from the data set for evaluation.

**Note:** For best resolution of fast kinetics, you should delete any disturbances in connection with injection start and stop (see illustration below). These are commonly caused by small misalignments in reference and blank subtraction, leading to spikes at the beginning and end of the injection. The effect of subtraction spikes on the calculated kinetic constants is usually negligible except for very fast interactions.
Click **Affinity** for steady state affinity evaluation or **Kinetics** for kinetics evaluation when you are satisfied with the curves.

3 **Affinity** only. If you choose to evaluate steady state affinity, the next dialog gives a preview of the plot of steady state response against concentration, with the option to adjust the selection of data used to calculate response values.

The top panel shows the plot of $R_{eq}$ against $C$, based on average response values over the region marked on the sensorgrams. Click **Settings** to adjust the region used for calculation of $R_{eq}$ values.
For single-cycle affinity runs, the same settings are applied to each injection in the cycle.

Click **Next** when you are satisfied with the data selection.

4. In the next dialog (applicable to both **Kinetics** and **Affinity**), you select the fitting model and perform the fit. The same data can be fitted repeatedly to different models or to the same model with different settings.

Select the model from the pull-down list. Available models are described in Section 9.8. Click **Parameters** if you want to change the starting values or scope of any of the parameters (see Section 9.7.2 for details), then click **Fit** to perform the fitting.

During the fitting procedure, the fitted curves are shown in black overlaid on the experimental data. Fitting progress is indicated in the sensorgram window by display of the iteration number, the current chi-squared value and the relative change in the parameter that was changed most from the previous iteration. You can use the **Abort** or **Accept** buttons to cancel the fitting or accept the fitting after the current iteration. You may want to cancel the fitting if it is clear that a fit cannot be found, or to accept the fitting if the chi-squared value and/or maximum relative change indicate that an acceptable fit has been achieved. Clicking on **Accept** will stop the fitting at the end of the current iteration, which may take a few moments to
complete.

You can enter a short description for the fit in the **Description** box. This can be useful for example to distinguish different fits for thermodynamic analysis (Section 10.2).

To perform additional fits on the same data, choose a new model or new parameter settings and click **Fit**. To remove a fit from the evaluation item, select the fit in the list of current fits and click **Delete**.

Click **Finish** to complete the evaluation and place the item in the evaluation explorer panel. You can click **Back** to review the choice of data for the evaluation: however, if you make any changes to the data (e.g. remove additional sections from a curve or exclude a curve from the set), all current fits will be deleted. Current fits are also deleted if you switch between kinetics and affinity evaluation.

**Kinetics results**

When the fit is completed, the results for kinetics are displayed as fitted curves overlaid in black on the experimental data, with details in the table below the curves:

The detailed results are presented on four tabs:

- **Quality control** summarizes important aspects of the quality of the fitting, as an aid to judging the reliability of the reported results. The quality control criteria are discussed in Section 9.4.1. This tab is only shown for kinetic evaluations using the predefined 1:1 model.
9 Kinetics and affinity analysis
9.2 Evaluating kinetics and affinity in single mode

- **Report** shows selected parameters and calculated values. The contents of the Report tab are defined in the model. Global parameters are listed on a single row at the top of the table, and local parameters are listed on one row for each curve.

- **Residuals** plots the difference between the experimental and fitted curves for each point in the curves. Use this display as an aid in judging how closely the results fit the experimental data.

- **Parameters** shows the values for all parameters in the fitting equations.

**Affinity results**
For affinity determination, the reported $K_D$ value is marked on the plot as a vertical line (for a 1:1 interaction, $K_D$ is the same as the analyte concentration at a response equal to half $R_{max}$). If the reported value is higher than half the highest concentration used, this line will be shown broken in red as a warning that the value may be unreliable because the plot does not flatten out sufficiently.

The detailed results are presented on two tabs:
- **Report** shows selected parameters and calculated values. The contents of the Report tab are defined in the model. Global parameters are listed on a single row at the top of the table, and local parameters are listed on one row for each curve.

- **Parameters** shows the values for all parameters in the fitting equations.
9.2.2 Multiple ligand densities

Analysis of the same analyte concentration series over multiple ligand densities can provide more robust fitting than a single ligand density. The kinetic and steady-state affinity fitting functions support simultaneous analysis of up to five sets of data with independent values for $R_{\text{max}}$, returning a single set of rate constants for the whole combined data set. Analyses over multiple ligand densities may be performed in separate runs that are combined with the Append file function, but should preferably be performed on multiple flow cells in the same run to ensure that the experimental conditions are comparable as far as possible.

To set up evaluation of the same sample series over multiple ligand densities, click Multiple $R_{\text{max}}$ in the first dialog box for kinetics and affinity evaluation. A panel for data subsets representing different $R_{\text{max}}$ values opens to the left of the curve table.

Click Add to add a new data subset. If you have multiple sets of the currently chosen curve type in the evaluation session (e.g. multiple reference subtracted curves), the next curve in the list is assigned to the new data subset. If there are no more curves of the same type available, the new set will be a copy of the most recently created subset. You can add up to five data subsets, representing five ligand densities. The same sample name and analysis temperature apply to all subsets.

Click on a subset to manage the contents of the subset. You cannot mix subsets that use different samples or different temperatures.
9 Kinetics and affinity analysis
9.3 Batch mode evaluation

**IMPORTANT!** Make sure that the same curve numbers are not assigned to more than one data subset. If a curve is assigned to two data sets, the software will try to evaluate the same curve with two different $R_{\text{max}}$ values, and the fitting may be distorted.

Do not evaluate multiple copies of the same subset for multiple ligand densities. Subsets that are duplicated will be weighted more than those that are not duplicated in the fitting procedure.

Do not attempt to use subsets for any purpose other than multiple ligand densities. The data will be evaluated in terms of multiple ligand densities regardless of how you have assigned curves to the subsets.

The subsets will be evaluated together, with rate or affinity constants that are global for the whole data set and $R_{\text{max}}$ values that are global within each subset but can differ between subsets.

**Note:** The Parameters tab in evaluation of kinetics with a global $R_{\text{max}}$ parameter within subsets lists a parameter $i_n$, which is set to 1 for the $n^{\text{th}}$ subset and 0 for all others. This is used to control $R_{\text{max}}$ values for each subset: values are fitted as $R_{\text{max}} \times i_n$ which returns a non-zero value for the $n^{\text{th}}$ subset only. For affinity evaluations, each subset is represented by one curve and $R_{\text{max}}$ is simply evaluated as a local parameter.

9.3 Batch mode evaluation

Batch mode allows you to evaluate multiple sample series (e.g. different analytes interacting with a common ligand, or measurement of the same interaction at different temperatures) automatically. You select the series to be evaluated and the fitting model to be used, and evaluation is carried to completion without further intervention.

To perform evaluation in batch mode, start the kinetics/affinity evaluation (Section 9.2.1) and check the **Batch mode** option in the first dialog.
Choose the **Evaluation purpose** as kinetics or affinity, and choose the fitting model. Click the **Parameters** button if you want to make adjustments to the fitting parameters (see Section 9.2.1). The same model and parameter settings will be used for all evaluations in the batch.

Choose the curve type to use (you should normally evaluate reference-subtracted curves) and check the data sets you want to evaluate in the table. Click **Finish** to perform the evaluation. One evaluation item will be created for each sample series.

**Notes:** A data set for batch evaluation is defined as a set of cycles with the same ligand, sample and analysis temperature. Blank cycles will be subtracted within each data set.

*Evaluation in batch mode does not support multiple ligand densities, adjustment of injection start, selection of blank cycles or removal of selected data. Individual evaluation items created during batch evaluation can however be edited separately.*
9.4 Quality assessment for kinetics evaluation

9.4.1 The Quality Control tab

The Quality Control tab in the kinetic evaluation results (for evaluations using the predefined 1:1 model only) gives a brief overview of the reliability of the results. If you prefer, you can hide the quality control tab by setting the appropriate option in Tools:Preferences on the main menu.

The symbols used on this tab have the following meanings:

- **(Yellow)** Warning: quality assessment close to the limits of acceptability
- **(Red)** Fail: quality assessment unacceptable
- **(Blue)** User assessment recommendations

The quality of the fitting is assessed in five areas:

**Magnitude of kinetic constants**

If either association or dissociation rate constants are close to or outside the limits that can be determined in the instrument, this will be reported. For values close to the limit, judge the validity of the results on other assessment criteria as described in this chapter.

**Parameter uniqueness**

In some situations, it may be possible to determine a value for a combined function of two or more parameters without being able to determine unique values for the individual parameters. Such parameters are said to be correlated. One example is the pair of kinetic rate constants $k_a$ and $k_d$, that are related through the affinity constant $K_D (K_D = k_d/k_a)$; it may be possible to determine the affinity constant reliably without being able to resolve the individual rate constants.
Parameter uniqueness is assessed by testing correlation between pairs of the parameters $k_a$, $k_d$, and $R_{max}$. If significant correlation is found, this will be reported as a warning that parameters cannot be uniquely determined.

**Note:** This test does not explore all possible parameter correlations. A Pass status for this test is not a fail-safe indication that parameters are uniquely determined.

The *Check Kinetic Data* tool (Section 9.4.4) provides a visualization of potential correlation between $k_a$ and $k_d$.

**Bulk refractive index**

After reference subtraction and blank subtraction, sensorgrams for kinetic evaluation should not in principle contain any bulk refractive index shifts (parameter $RI$ in predefined models). However, there may be some circumstances where small bulk refractive index shifts may remain in reference- and blank-subtracted data. On the other hand, the fitting algorithm tends to interpret rapid interaction events (incorrectly) as bulk shifts. If the fitting returns significant values for $RI$, a warning will be issued in the quality control tab.

Examine the sensorgrams and fitted curves to determine whether bulk shifts as reported by the fitting are true or false. In cases where reported bulk shifts are unreasonably large, you may want to set $RI$ to a constant value of zero in the *Parameters* setting for the fitting. If you do this, the bulk contributions component in the quality control tab will be reported as neutral since the $RI$ parameter was not evaluated.

**Sensorgram curvature**

You should check that the sensorgrams have sufficient curvature for kinetic determination. Ideally, the sensorgrams for at least the one or two highest concentrations should show measurable binding rates at the beginning of the sample injection and approach a steady state towards the end of the injection. Sensorgrams that approximate to “square-wave” pulses (indicating rapid association and dissociation) and those that do not flatten out during the injection generally do not contain sufficient kinetic information for reliable evaluation. Ideally, the dissociation phase should be long enough to monitor a fall in response of at least 10-15% of the starting value.
9 Kinetics and affinity analysis

9.4 Quality assessment for kinetics evaluation

Figure 9-2. Examples of sufficient and insufficient sensorgram curvature.
Top: Ideal sensorgrams approaching steady state during sample injection and returning to baseline during dissociation.
Bottom left: Rapid interaction approaching “square wave” appearance. These sensorgrams return rate constants close to the limit of measurement for the instrument.
Bottom right: Slow association and dissociation, giving insufficient curvature in both association and dissociation phases. Evaluation is possible but will not be very reliable.

If the interaction is too fast to provide kinetic information, you may only be able to determine affinity constants. Interactions that do flatten out sufficiently during the injection or dissociate sufficiently during the dissociation phase may sometimes be analyzed by prolonging the association or dissociation phase respectively.

Residuals
You should check that the residuals (the difference between experimental and fitted value for each data point in the sensorgrams) lie within reasonable limits. For a perfect fit, the residuals reflect the short-term noise in the sensorgrams and scatter around zero (typically ±1-2 RU). Systematic deviations, seen as a definite shape in the residual plot, indicate that the interaction model is to a greater or lesser extent unsuitable for the interaction. As an aid in judging the residuals, guidelines are drawn on the residual plot to indicate the range of acceptability. Most of the residuals should be within the inner (green) limits.

Figure 9-3. The residuals for a good fit (left) scatter around 0, ideally in a random distribution representing the noise in the sensorgrams. For a poor fit (right) the residual curves show a definite shape and deviate farther from 0.
The guideline positions are calculated in relation to the response range of the sensorgrams. The guidelines are only shown for evaluations using the predefined 1:1 model (i.e. when the quality control tab is included).

**IMPORTANT!** Use the Quality Control assessment as a help in making your own judgement of the results. Pass status in the quality control parameters does not necessarily indicate that the fit is acceptable or that the results are biologically relevant. On the other hand, Fail status in any of the parameters is a reliable warning indicator.

Base your assessment on the overall quality of the results and the fitting, taking all quality control parameters into account.

### 9.4.2 Statistical parameters

Two statistical parameters are provided to help in assessing the results:

- **Chi-squared** is an indicator of how closely the fitted curves agree with the experimental data. One chi-squared value is reported for the whole fitting.
- **Standard error or T-value** is an indicator of parameter significance, and is reported separately for each fitted parameter.

#### Chi-squared

Chi-squared is a measure of the average squared residual (the difference between the experimental data and the fitted curve), calculated as:

$$\text{chi-squared} = \sum_{i=1}^{n} \frac{(r_f - r_x)^2}{n - p}$$

where  
- $r_f$ is the fitted value at a given point  
- $r_x$ is the experimental value at the same point  
- $n$ is the number of data points  
- $p$ is the number of fitted parameters

For sensorgram data, the number of data points is very much larger than the number of fitted parameters in the model, so

$$n - p \approx n$$

and chi-squared reduces to the average squared residual per data point. If the model fits the experimental data precisely, chi-squared represents the mean square of the signal noise.
Chi-squared is listed on the Report tab.

**Standard error and T-value**

The significance of parameter values is indicated by the standard error (SE) or T-value listed on the Parameters tab in the fitting results. This is a statistical indication of the significance of a fitted parameter. Lower standard error values indicate higher significance: if the standard error represents less than 10% of the parameter value, the parameter is significant for the experimental data.

For ease of comparison between parameters with widely different absolute values (e.g. $k_a$ and $k_d$), the standard error may be expressed as a **T-value**, which is obtained by dividing the value of the parameter by the standard error. A high T-value corresponds to a low standard error. As a general guideline, parameters with a T-value greater than about 10 should be regarded as significant.

The choice of whether to display parameter significance as standard error or T-value is made on the Fit tab of the Tools:Preferences dialog.

The significance of a parameter is a measure of how much a change in the parameter value affects the closeness of fit. A parameter with low significance can have a wide range of values without affecting the fit. Typically (but not always), parameters with a low significance have unreasonable values: for example typical values for the mass transfer constant for proteins are around $10^8$ RU·M$^{-1}$s$^{-1}$, but evaluation of data with no mass transfer limitation might return a value of $10^{12}$ or higher. Similarly, rate constants that lack significance are often assigned values outside the reasonable range for biomolecular interactions, or outside the range that can be measured with Biacore.

**Notes:** The standard error and the Check Kinetic Data tool assess parameter significance in different ways, even if the results of the assessment may sometimes be related. Check Kinetic Data tests the contribution of a group of parameters (rate constants for the interaction and mass transport processes) to the closeness of fit by examining the results of correlated changes, whereas the standard error is a mathematical assessment of the significance of a single parameter. If the Check Kinetic Data tool indicates that the rate constants are not significant, the standard error for the constants may be expected to be high. However, the converse is not always true (a high standard error will not always be reflected in the behavior of the Check Kinetic Data tool).

Even if parameters with low significance can have a wide range of values without affecting the fit, repeated evaluation of the same data set will always return the same value for all parameters. Consistency of a value between repeated evaluations is **not** a test of significance.
**U-value**

The U-value is an estimate of the uniqueness of the calculated values for rate constants and $R_{\text{max}}$. If parameters are correlated (see Section 9.4.1), the fitting procedure can determine their relative magnitudes but not absolute values (for example, knowing the affinity gives the ratio but not the values for rate constants). The U-value is determined by testing the dependence of the fit on correlated variations in pairs of parameters, and is reported on the Report tab as a single value for the whole fitting. U-values above about 25 indicate that absolute values for two or more of the parameters (rate constants and $R_{\text{max}}$) are correlated and cannot be determined. If the U-value is below about 15 the parameter values are not significantly correlated.

### 9.4.3 Components of the fit

Choose **Tools:Components** from the fitted results window to display a plot showing the contribution of components in the interaction model to the fitted curve. Choose which cycle to display in the selector bar.

![Components](image)

The example illustrated here is taken from a fitting to a bivalent analyte model (see Section 9.8.2), and shows clearly how the component AB2 (analyte attached to the surface through both binding sites) is displaced by AB as the interaction progresses.

### 9.4.4 Check kinetic data

Kinetic constants obtained from the fitting procedure are only significant if the observed binding is not seriously limited by mass transport of analyte to the surface (see Section 9.8). For 1:1 fitting results, you can check whether mass transport is limited or not using the **Check Kinetic Data** function.
Choose **Tools:Check Kinetic Data** to open a dialog that displays simulated sensorgrams based on the fitting results, with the interaction rate constants $k_a$ and $k_d$ varied in parallel (so that the affinity constant, remains unchanged). If curves do not shift as values for $k_a$ and $k_d$ are changed, this means that the actual values are not important for the fitting, and the curves do not contain kinetic information. Conversely, if the simulated curve shape changes as the values of $k_a$ and $k_d$ are varied, the fitting is dependent on the actual values and the curves do contain kinetic information.

This tool is only available for results obtained with the 1:1 fitting model.

(This example illustrates a fitting with only three concentrations for clarity.)

To use the tool, drag the slider for the modification factor $M$ and observe the behavior of the curve display. The original curves (which remain unchanged as you drag the slider) are shown in black: blue curves show the simulation for $k_a$ and $k_d$ multiplied by $M$, while red curves show the simulation for $k_a$ and $k_d$ divided by $M$. If the red (reduced rate constants) and blue (increased rate constants) curves clearly diverge from the original curves, the fitting is sensitive to changes in the rate constants and the curves probably contain significant
kinetic information. If on the other hand the divergence is negligible, the values of the rate constants do not matter because the binding is fully limited by mass transfer. Mass transfer places an upper limit on the rate constants that can be measured: on the borderline, the fitting is sensitive to a reduction in rate constants but not to an increase.

Choose the Residuals option in the Compare to frame to examine the effect of varying the modification factor on the difference between the original and modified curves in relation to the experimental residuals. The tool display allows the simulated difference curves to be compared to the experimental residuals or to residuals averaged over a moving time window. The latter option smooths the experimental residual display, making it easier to observe the general shape of the residual curves. Movable horizontal Limit guides can be displayed to mark the extent of the residual variation and aid visual interpretation. (Note that the limit guides are not related to the guidelines shown on the residual tab for QC purposes (Section 9.4.1), and do not in any way imply acceptance limits.)

Figure 9-4. Kinetic data check comparison to residuals (left) and averaged residuals (right).

9.5 Quality assessment for affinity evaluation

Steady state affinity evaluations are performed by fitting a plot of $R_{eq}$ against concentration $C$ to a model representing equilibrium 1:1 binding. The closeness of fit is reported as a chi-squared value, calculated in the same way as for kinetics. Note however that the number of points in the steady state affinity plot is very much lower than for kinetic evaluation, so that chi-squared is a more sensitive indicator of fitting quality.

The plot of $R_{eq}$ against $C$ approaches a limiting value (equivalent to $R_{max}$) at very high concentrations. Robust evaluation of the data requires that the plot shows sufficient curvature for reliable estimation of $R_{max}$. As a rule of thumb, the evaluation is acceptable only if the calculated $K_D$ value is less than half the highest analyte concentration used. (For a 1:1 interaction, the $K_D$ value is equal to the analyte concentration that gives 50% saturation of the binding sites, so that $R_{eq}=0.5R_{max}$. In other words, reliable evaluation is only obtained if the surface is more than 50% saturated at the highest analyte concentration.)
To help in this assessment, the calculated $K_D$ value is indicated as a vertical line at the corresponding analyte concentration. The line is red and broken if the value is greater than half the highest concentration.

**Figure 9-5.** For reliable evaluation, the calculated $K_D$ value should be less than half the highest analyte concentration used. When this condition is met, the $K_D$ value is indicated with a full black line in the result plot (left). When the condition is not met, the $K_D$ value is indicated with a broken red line.

### 9.6 Summarizing kinetics and affinity results

You can summarize the kinetics and affinity evaluation results from several runs or from several evaluation items for the same run using the separate Biacore T200 Kinetics Summary software (this is installed automatically together with the Evaluation Software).

#### 9.6.1 Creating kinetic summaries

To create a kinetic summary of the results in a single evaluation file, simply open the evaluation file (file type `.bme`) in the Kinetics Summary software. The Kinetics Summary software can also be started from the *Tools* menu in the Evaluation Software.

To create a summary of the results in multiple files, you can either select multiple files within the same folder in the *Open* dialog box using *Ctrl-click* and/or *Shift-click*, or use *File:Append File* to add results to an existing summary. You can open saved summary files (file type `.bks`) in addition to evaluation files.

#### 9.6.2 Basic summary presentation

The summarized kinetic and affinity results are presented as thumbnail plots of the kinetics and evaluation items and as a table of result data. Multiple fits in the same item in the Evaluation Software are presented as separate fits in the summary.
Thumbnail types

Thumbnails can be displayed as small, standard or extended diagrams, selected from the View menu or from the options under the Thumbnails button.

Small Provides an overview of many thumbnails for general comparison. Details will not be legible in the thumbnails. Individual thumbnails are identified in a tool tip.

Standard Shows the thumbnails with identification, legible axis, and a summary of fitting model and kinetic or affinity constants.

Extended Shows more detailed identification of each thumbnail than the standard view.

Thumbnail display settings

Choose Display Settings from the View menu or the right-click menu for display settings for the thumbnails.
If you choose to use the same x- and/or y-axis scales, the scale(s) will be chosen to include the widest range in the currently included set of thumbnails. Scales are chosen independently for steady-state affinity and kinetic items. If the **Use same scales** options are not checked, each individual thumbnail will be scaled according to the range of data in the item.

Check **Show curve fits in thumbnails** to display the fitted curves overlaid on the experimental data in the thumbnails. This may not have any significant effect on the appearance of small thumbnails for kinetic evaluation items, but will in general be readily visible for affinity items.

**Sorting items**

Choose **Arrange by** from the right-click menu in the thumbnail display to sort the items. The list of sorting parameters corresponds to the columns on the **Table** tab.

Sorting the thumbnail display also sorts the rows in the **Table** tab and vice versa. You can sort the rows directly in the table according to the contents of a column by clicking on the column header. Clicking repeatedly on the column header toggles the sort order.

**Note:** The **Arrange by** option on the ** Thumbnails** tab always sorts the thumbnails and table rows in ascending order. Sort the rows on the **Table** tab if you want to use a descending sort order.

**Excluding items**

To exclude an evaluation from the summary, right click on the item thumbnail and choose **Exclude thumbnail**, or remove the checkmark from the Included column on the **Table** tab. You can also exclude kinetic evaluation items using the right-click menu in the on-off rate map (see Section 9.6.3).

**Evaluation item details**

Double-click on a thumbnail or table row or choose **Evaluation details** from the right-click menu to display the evaluation item as it appears in the Evaluation Software (Section 9.2). You can also display evaluation details using the right-click menu in the on-off rate map (see Section 9.6.3).

**Note:** You can only display the evaluation details for one item at a time.

**Copying and exporting summaries**

From the right-click menu on the **Thumbnail** tab, you can copy the single or all graphs or thumbnails to the Windows clipboard as graphic objects for pasting into other programs. Copying a graph copies only the graph itself; copying a thumbnail copies the graph together with the additional information as shown on the screen. (Copying a graph and a thumbnail is equivalent when small thumbnails are displayed, since there is no additional information in this mode.) Copying all items copies all the included items in the summary (regardless of
how many are currently visible on the tab) as a single graphic object, arranged as in the display.

You can also export the curve data from the currently selected item to a tab-separated text file for import into third-party software. The exported file includes data for the fitted curves if they are displayed in the summary.

From the Table tab, you can copy the table data for single rows or the complete table as text to the Windows clipboard.

**Note:** The header row is not included when single rows are copied.

### 9.6.3 On-off rate maps

On-off rate maps (also called $k_a/k_d$ plots) provide an overview of kinetic and affinity properties for the interactions in the summary by plotting the association rate constant $k_a$ against the dissociation rate constant $k_d$, both on logarithmic scales. Since the affinity constant $K_D$ is the ratio of $k_d$ to $k_a$, interactions that have the same affinity will appear on diagonal lines representing the $K_D$ value. The diagonals are shown as broken lines on the plot with the $K_D$ value indicated. Points that lie widely separated on the same diagonal represent interactions with the same affinity but different kinetics.

[Image of an on-off rate map showing association and dissociation rates on a log-log scale, with diagonal lines indicating constant affinity and dots representing different sets of rate constants.]

Choose the **On-Off Rate Map** tab to view the rate map for the current kinetics summary. Only items with single sets of values for kinetic rate constants are represented in the plot. The plot does not therefore include steady state affinity items or kinetic evaluation items with multiple sets of kinetic constants, such as evaluation with heterogeneous models.

Click on a point in the on-off rate map or select a row in the table below the map to display the thumbnail for the evaluation.
9 Kinetics and affinity analysis
9.7 Curve fitting principles

Right-click in the on-off rate map for display options and copy and export functions.

9.7 Curve fitting principles

With all kinetic and affinity analysis, it is important to remember that the results obtained represent the results of fitting the experimental data to a mathematical model, and that obtaining a good fit is not in itself evidence that the model describes the physical reality of the interaction. The fitting procedure does not have any “knowledge” of the biological significance of parameters in the model equations, and it is wise always to examine the results obtained for reasonableness of the values obtained. In addition, any mechanistic conclusions drawn for the interaction from fitting results (e.g. concerning multiple interaction sites or conformational changes) should ideally be tested using independent techniques.

9.7.1 Fitting procedure

Kinetic parameters are extracted from experimental data by an iterative process that finds the best fit for a set of equations describing the interaction. The equations are created automatically from the definition of the interaction model. The fitting process begins with initial values for the parameters in the equation set, and optimizes the parameter values according to an algorithm that minimizes the chi-squared value (Section 9.4.2) for the fitting.
In some situations, the fitting algorithm may be unable to find a fit for the experimental data with the initial parameter values as specified in the model. This may happen typically if the concentration unit is incorrect: for example if the unit is set to mM instead of nM in the keyword table. On occasion, however, it can be necessary to adjust the starting values for fitting parameters, accessed through the Parameters button in the fitting dialog (step 4 in Section 9.2).

### 9.7.2 Local and global parameters

Parameters in the fitting equations are treated as either **local** or **global** variables or **constants**:

- **Local parameters** are assigned an independent value for each curve in the data set. Typical local parameters are concentration (which is different for different curves) and bulk refractive index contribution (which may be expected to vary between curves).

- **Global parameters** have one single value that applies to the whole data set. Typical global parameters are the rate constants for the interaction, which should in principle have the same value for all curves in the data set.

- **Constants** have a fixed value that is not changed in the fitting procedure. An example is the analyte concentration. Constants may also be local (separate values for each curve) or global (one value for the whole data set).

The local/global status of parameters can be changed through the Parameters button in the fitting dialog (step 4 in Section 9.2), without making any changes to the model.

Evaluating kinetics or affinity with global rate constants gives a more robust value for the rate constants, although the curves may fit the experimental data more closely if all parameters are fitted locally. This is because local fitting allows variation between the constants obtained from different curves; when the constants are fitted globally, this variation appears in the closeness of fit rather than the reported values. Rate constants are always global in predefined kinetic models.

In general, kinetic constants should be fitted as global parameters and bulk refractive index contribution as a local parameter. The analyte binding capacity of the surface $R_{\text{max}}$ is a global parameter by default in the predefined models (this assumes that the ligand activity is unchanged between cycles in the assay). It is however justified to use a local $R_{\text{max}}$ if there is reason to believe that the ligand activity may vary between cycles (e.g. in a capture assay, if the capture level varies between cycles).
9.8 Predefined models

A set of predefined models for kinetics and steady state affinity is provided with Biacore T200 Evaluation Software. These models are marked in the model selection list (see Section 9.2) with a red dot, and cannot be removed or modified.

**Mass transfer parameters**

All kinetic models include a term for mass transfer of analyte to the surface. If transport is slow compared with binding of analyte to the ligand, the transport process will limit the observed binding rate, at least partially. All models take account of this potential limitation and can extract rate constants from the data provided that mass transfer is not totally limiting (see Section 9.8).

The rate of mass transfer of analyte to the surface under the conditions of non-turbulent laminar flow that prevail in the Biacore flow cell is characterized by the mass transfer coefficient $k_m$ (units m·s$^{-1}$):

$$k_m = 0.98 \left( \frac{D^2 \cdot f}{0.3 \cdot h^2 \cdot w \cdot l} \right)^{1/3}$$

where $D$ is the diffusion coefficient of the analyte (m$^2$·s$^{-1}$)

$f$ is the volume flow rate of solution through the flow cell (m$^3$·s$^{-1}$)

$h, w, l$ are the flow cell dimensions (height, width, length in m)

One form used in fitting models in Biacore T200 is referred to as the mass transfer constant $k_t$ (units RU·M$^{-1}$·s$^{-1}$), obtained by adjusting the mass transfer coefficient approximately for the molecular weight of the analyte and for the conversion of surface concentration to RU:

$$k_t = k_m \times \text{MW} \times \text{G}$$

where $G$ is the conversion factor from surface concentration to RU. The value of $G$ is approximately $10^9$ for proteins on Sensor Chip CM5.

A further modification of this expression gives the flow rate-independent component of the mass transfer constant (units RU·M$^{-1}$·s$^{-2/3}$·m$^{-1}$), referred to as $tc$ in the models:

$$t_c = \frac{k_t}{\sqrt[3]{f}}$$
9.8.1 Kinetics – 1:1 binding

This is the simplest model for kinetic evaluation, and is recommended as default unless there is good experimental reason to choose a different model. The model describes a 1:1 interaction at the surface:

$$A + B = AB$$

### Model parameters

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Description</th>
<th>Obtained from</th>
</tr>
</thead>
<tbody>
<tr>
<td>$ka$</td>
<td>Association rate constant ($M^{-1}s^{-1}$)</td>
<td>Fitted</td>
</tr>
<tr>
<td>$kd$</td>
<td>Dissociation rate constant ($s^{-1}$)</td>
<td>Fitted</td>
</tr>
<tr>
<td>$R_{max}$</td>
<td>Analyte binding capacity of the surface (RU)</td>
<td>Fitted</td>
</tr>
<tr>
<td>$Conc$</td>
<td>Analyte concentration (M)</td>
<td>Provided as input</td>
</tr>
<tr>
<td>$tc$</td>
<td>Flow rate-independent component of the mass transfer constant</td>
<td>Fitted</td>
</tr>
<tr>
<td>$f$</td>
<td>Flow rate (µl/min)</td>
<td>Provided as input</td>
</tr>
<tr>
<td>$t_{On}$</td>
<td>Sample injection start time (s)</td>
<td>Provided as input</td>
</tr>
<tr>
<td>$t_{Off}$</td>
<td>Sample injection end time (s)</td>
<td>Provided as input</td>
</tr>
<tr>
<td>$RI$</td>
<td>Bulk refractive index contribution in the sample</td>
<td>Fitted</td>
</tr>
</tbody>
</table>

### Report parameters

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Description</th>
<th>Calculated as</th>
</tr>
</thead>
<tbody>
<tr>
<td>$ka$</td>
<td>Association rate constant ($M^{-1}s^{-1}$)</td>
<td>$ka$</td>
</tr>
<tr>
<td>$kd$</td>
<td>Dissociation rate constant ($s^{-1}$)</td>
<td>$kd$</td>
</tr>
<tr>
<td>$KD$</td>
<td>Equilibrium dissociation constant (M)</td>
<td>$kd/ka$</td>
</tr>
<tr>
<td>$R_{max}$</td>
<td>Analyte binding capacity of the surface (RU)</td>
<td>$R_{max}$</td>
</tr>
<tr>
<td>$Conc$</td>
<td>Analyte concentration (M)</td>
<td>$Conc$</td>
</tr>
<tr>
<td>$tc$</td>
<td>Flow rate-independent component of the mass transfer constant</td>
<td>$tc$</td>
</tr>
<tr>
<td>$f$</td>
<td>Flow rate (µl/min)</td>
<td>$f$</td>
</tr>
<tr>
<td>$kt$</td>
<td>Mass transfer constant</td>
<td>$tc \times f^{1/3}$</td>
</tr>
<tr>
<td>$RI$</td>
<td>Bulk refractive index contribution in the sample</td>
<td>$RI$</td>
</tr>
</tbody>
</table>
9.8.2 Kinetics – Bivalent Analyte

This model describes the binding of a bivalent analyte to immobilized ligand, where one analyte molecule can bind to one or two ligand molecules. The two analyte sites are assumed to be equivalent. The model may be relevant to studies among others with signaling molecules binding to immobilized cell surface receptors (where dimerization of the receptor is common) and to studies using intact antibodies binding to immobilized antigen. As a result of binding of one analyte molecule to two ligand sites, the overall binding is strengthened compared with 1:1 binding. This effect is often referred to as avidity.

\[
A + B = AB \\
AB + B = AB_2
\]

Note: Once analyte is attached to the ligand through binding at the first site, interaction at the second site does not contribute to the SPR response. For this reason, the association rate constant for the second interaction is reported in units of RU\(^{-1}\)s\(^{-1}\), and can only be obtained in M\(^{-1}\)s\(^{-1}\) if a conversion factor between RU and M is available. Similarly, a value for the overall affinity or avidity constant is not reported.

<table>
<thead>
<tr>
<th>Model parameters</th>
<th>Obtained from</th>
</tr>
</thead>
<tbody>
<tr>
<td>ka1 Association rate constant for the first site (M(^{-1})s(^{-1}))</td>
<td>Fitted</td>
</tr>
<tr>
<td>kd1 Dissociation rate constant for the first site (s(^{-1}))</td>
<td>Fitted</td>
</tr>
<tr>
<td>ka2 Association rate constant for the second site (RU(^{-1})s(^{-1}))</td>
<td>Fitted</td>
</tr>
<tr>
<td>kd2 Dissociation rate constant for the second site (s(^{-1}))</td>
<td>Fitted</td>
</tr>
<tr>
<td>Rmax Analyte binding capacity of the surface (RU)</td>
<td>Fitted</td>
</tr>
<tr>
<td>Conc Analyte concentration (RU)</td>
<td>Provided as input</td>
</tr>
<tr>
<td>tc Flow rate-independent component of the mass transfer constant</td>
<td>Fitted</td>
</tr>
<tr>
<td>f Flow rate (μl/min)</td>
<td>Provided as input</td>
</tr>
<tr>
<td>tOn Sample injection start time (s)</td>
<td>Provided as input</td>
</tr>
<tr>
<td>tOff Sample injection end time (s)</td>
<td>Provided as input</td>
</tr>
<tr>
<td>RI Bulk refractive index contribution in the sample</td>
<td>Fitted</td>
</tr>
</tbody>
</table>
9.8.3 Kinetics – Heterogeneous Analyte

This model is intended for analysis of the kinetics of interaction of mixtures of two analytes that compete for the same ligand site. Experiments of this kind can be used to deduce kinetic parameters for a low molecular weight analyte that gives a small response from measurements of binding of a competing high molecular weight analyte. Response contributions from both analytes are taken into account, although the high molecular weight analyte is responsible for the dominant component in the observed sensorgrams.

Concentrations and molecular weights are required for both analytes. If absolute molecular weights are not known, relative values can be entered without affecting the outcome of the fitting. The model cannot evaluate interactions where the proportions and relative sizes of the analytes are unknown.

\[
\begin{align*}
A_1 + B &= A_1B \\
A_2 + B &= A_2B
\end{align*}
\]
### Model parameters

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Description</th>
<th>Obtained from</th>
</tr>
</thead>
<tbody>
<tr>
<td>ka1, ka2</td>
<td>Association rate constant for the first and second analytes (M⁻¹s⁻¹)</td>
<td>Fitted</td>
</tr>
<tr>
<td>kd1, kd2</td>
<td>Dissociation rate constant for the first and second analytes (s⁻¹)</td>
<td>Fitted</td>
</tr>
<tr>
<td>Conc1, Conc2</td>
<td>Concentration of the first and second analytes (M)</td>
<td>Provided as input</td>
</tr>
<tr>
<td>mw1, mw2</td>
<td>Molecular weights of the first and second analytes</td>
<td>Provided as input</td>
</tr>
<tr>
<td>tc1, tc2</td>
<td>Flow rate-independent component of the mass transfer constant for the first and second analytes</td>
<td>Fitted</td>
</tr>
<tr>
<td>Rmax1</td>
<td>Analyte binding capacity of the surface for the first analyte (RU)</td>
<td>Fitted</td>
</tr>
<tr>
<td>Rmax2</td>
<td>Analyte binding capacity of the surface for the second analyte (RU)</td>
<td>Fitted</td>
</tr>
<tr>
<td>rcf</td>
<td>Response correction factor, allowing for different refractive index contributions for the two analytes. This factor is defined as (Rmax1/Rmax2) / (MW1/MW2).</td>
<td>Fitted</td>
</tr>
<tr>
<td>f</td>
<td>Flow rate (µl/min)</td>
<td>Provided as input</td>
</tr>
<tr>
<td>tOn</td>
<td>Sample injection start time (s)</td>
<td>Provided as input</td>
</tr>
<tr>
<td>tOff</td>
<td>Sample injection end time (s)</td>
<td>Provided as input</td>
</tr>
<tr>
<td>RI</td>
<td>Bulk refractive index contribution in the sample</td>
<td>Fitted</td>
</tr>
</tbody>
</table>
## 9.8.4 Kinetics – Heterogeneous Ligand

This model describes an interaction between one analyte and two independent ligands. The binding curve obtained is simply the sum of the two independent reactions. Unlike the case of heterogeneous analyte, the relative amounts of the two ligands does not have to be known in advance.

Heterogeneous ligand situations frequently arise in practice through heterogeneous immobilization of ligand (e.g. amine coupling of proteins, where the ligand has multiple attachment points), as well as through heterogeneity in the ligand preparation itself. In cases where the heterogeneous ligand model is found to give the best fit to the observed sensorgrams, further experimental efforts to reduce the heterogeneity are recommended where possible.

\[
A + B1 = AB1 \\
A + B2 = AB2
\]

<table>
<thead>
<tr>
<th>Report parameters</th>
<th>Calculated as</th>
</tr>
</thead>
<tbody>
<tr>
<td>(ka1) (ka2)</td>
<td>(ka1) (ka2)</td>
</tr>
<tr>
<td>(kd1) (kd2)</td>
<td>(kd1) (kd2)</td>
</tr>
<tr>
<td>(KD1) (KD2)</td>
<td>(kd1/ka1) (kd2/ka2)</td>
</tr>
<tr>
<td>(Rmax1)</td>
<td>(Rmax2 \times rcf)</td>
</tr>
<tr>
<td>(Rmax2)</td>
<td>(Rmax2)</td>
</tr>
<tr>
<td>(Conc1) (Conc2)</td>
<td>(Conc1) (Conc2)</td>
</tr>
<tr>
<td>(tc1) (tc2)</td>
<td>(tc1) (tc2)</td>
</tr>
<tr>
<td>Flow</td>
<td>(f)</td>
</tr>
<tr>
<td>(kt1) (kt2)</td>
<td>(tc1 \times f^{1/3}) (tc2 \times f^{1/3})</td>
</tr>
<tr>
<td>RI</td>
<td>(RI)</td>
</tr>
</tbody>
</table>
Note: The model is limited to two ligands because the fitting algorithm tends to become unstable with more components, and three or more ligand species cannot be reliably resolved.

<table>
<thead>
<tr>
<th>Model parameters</th>
<th>Obtained from</th>
</tr>
</thead>
<tbody>
<tr>
<td>$k_a1$</td>
<td>Fitted</td>
</tr>
<tr>
<td>$k_a2$</td>
<td>Fitted</td>
</tr>
<tr>
<td>$k_d1$</td>
<td>Fitted</td>
</tr>
<tr>
<td>$k_d2$</td>
<td>Fitted</td>
</tr>
<tr>
<td>$R_{max1}$</td>
<td>Fitted</td>
</tr>
<tr>
<td>$R_{max2}$</td>
<td>Fitted</td>
</tr>
<tr>
<td>$C_{on}$</td>
<td>Provided as input</td>
</tr>
<tr>
<td>$C_{off}$</td>
<td>Provided as input</td>
</tr>
<tr>
<td>$C_{r_i}$</td>
<td>Fitted</td>
</tr>
</tbody>
</table>

Report parameters

<table>
<thead>
<tr>
<th>Report parameters</th>
<th>Calculated as</th>
</tr>
</thead>
<tbody>
<tr>
<td>$k_a1$</td>
<td>$k_a1$</td>
</tr>
<tr>
<td>$k_a2$</td>
<td>$k_a2$</td>
</tr>
<tr>
<td>$k_d1$</td>
<td>$k_d1$</td>
</tr>
<tr>
<td>$k_d2$</td>
<td>$k_d2$</td>
</tr>
<tr>
<td>$K_{D1}$</td>
<td>$k_d1/k_a1$</td>
</tr>
<tr>
<td>$K_{D2}$</td>
<td>$k_d2/k_a2$</td>
</tr>
<tr>
<td>$R_{max1}$</td>
<td>$R_{max1}$</td>
</tr>
</tbody>
</table>
9.8.5 Kinetics – Two State Reaction

This model describes a 1:1 binding of analyte to immobilized ligand followed by a conformational change that stabilizes the complex. To keep the model simple, it is assumed that the conformationally changed complex can only dissociate through the reverse of the conformational change:

\[
A + B = AB = AB^* \]

Note that conformational changes in ligand or complex do not normally give a response in Biacore. A good fit of experimental data to the two-state model should be taken as an indication that conformational properties should be investigated using other techniques (e.g. spectroscopy or NMR), rather than direct evidence that a conformational change is taking place.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Description</th>
<th>Obtained from</th>
</tr>
</thead>
<tbody>
<tr>
<td>( R_{\text{max}}^2 )</td>
<td>Analyte binding capacity of the second ligand (RU)</td>
<td>( R_{\text{max}}^2 )</td>
</tr>
<tr>
<td>Conc</td>
<td>Analyte concentration (M)</td>
<td>Conc</td>
</tr>
<tr>
<td>( t_c )</td>
<td>Flow rate-independent component of the mass transfer constant</td>
<td>( t_c )</td>
</tr>
<tr>
<td>Flow</td>
<td>Flow rate (( \mu \text{l/min} ))</td>
<td>( f )</td>
</tr>
<tr>
<td>( k_t )</td>
<td>Mass transfer constant ( t_c \times f^{1/3} )</td>
<td>( t_c \times f^{1/3} )</td>
</tr>
<tr>
<td>( R_I )</td>
<td>Bulk refractive index contribution in the sample</td>
<td>( R_I )</td>
</tr>
<tr>
<td>( R_{\text{max}} )</td>
<td>Analyte binding capacity of the surface (RU)</td>
<td>Fitted</td>
</tr>
<tr>
<td>Conc</td>
<td>Analyte concentration (M)</td>
<td>Provided as input</td>
</tr>
<tr>
<td>( t_c )</td>
<td>Flow rate-independent component of the mass transfer constant</td>
<td>Fitted</td>
</tr>
</tbody>
</table>

Model parameters

- **\( k_a^1 \)**: Association rate constant for analyte binding (M\(^{-1}\)s\(^{-1}\))
  - Obtained from: Fitted
- **\( k_d^1 \)**: Dissociation rate constant for analyte from the complex (s\(^{-1}\))
  - Obtained from: Fitted
- **\( k_a^2 \)**: Forward rate constant for the conformational change (s\(^{-1}\))
  - Obtained from: Fitted
- **\( k_d^2 \)**: Reverse rate constant for the conformational change (s\(^{-1}\))
  - Obtained from: Fitted
- **\( R_{\text{max}} \)**: Analyte binding capacity of the surface (RU)
  - Obtained from: Fitted
- **Conc**:
  - Obtained from: Provided as input
- **\( t_c \)**:
  - Obtained from: Fitted
### Kinetics and affinity analysis

#### 9.8 Predefined models

<table>
<thead>
<tr>
<th>Symbol</th>
<th>Description</th>
<th>Notes</th>
</tr>
</thead>
<tbody>
<tr>
<td>f</td>
<td>Flow rate (µl/min)</td>
<td>Provided as input</td>
</tr>
<tr>
<td>tOn</td>
<td>Sample injection start time (s)</td>
<td>Provided as input</td>
</tr>
<tr>
<td>tOff</td>
<td>Sample injection end time (s)</td>
<td>Provided as input</td>
</tr>
<tr>
<td>RI</td>
<td>Bulk refractive index contribution in the sample</td>
<td>Fitted</td>
</tr>
</tbody>
</table>

**Report parameters**

<table>
<thead>
<tr>
<th>Symbol</th>
<th>Description</th>
<th>Formula</th>
</tr>
</thead>
<tbody>
<tr>
<td>ka1</td>
<td>Association rate constant for analyte binding (M⁻¹s⁻¹)</td>
<td>ka1</td>
</tr>
<tr>
<td>kd1</td>
<td>Dissociation rate constant for analyte from the complex (s⁻¹)</td>
<td>kd1</td>
</tr>
<tr>
<td>ka2</td>
<td>Forward rate constant for the conformational change (s⁻¹)</td>
<td>ka2</td>
</tr>
<tr>
<td>kd2</td>
<td>Reverse rate constant for the conformational change (s⁻¹)</td>
<td>kd2</td>
</tr>
<tr>
<td>KD</td>
<td>Overall equilibrium dissociation constant (M)</td>
<td>( \frac{kd1}{ka1} \times \ \left( \frac{kd2}{kd2+ka2} \right) )</td>
</tr>
<tr>
<td>Rmax</td>
<td>Analyte binding capacity of the surface (RU)</td>
<td>Rmax</td>
</tr>
<tr>
<td>Conc</td>
<td>Analyte concentration (M)</td>
<td>Conc</td>
</tr>
<tr>
<td>tc</td>
<td>Flow rate-independent component of the mass transfer constant</td>
<td>tc</td>
</tr>
<tr>
<td>Flow</td>
<td>Flow rate (µl/min)</td>
<td>f</td>
</tr>
<tr>
<td>kt</td>
<td>Mass transfer constant</td>
<td>( tc \times f^{1/3} )</td>
</tr>
<tr>
<td>RI</td>
<td>Bulk refractive index contribution in the sample</td>
<td>RI</td>
</tr>
</tbody>
</table>
9.8.6 Affinity – Steady State 1:1

This model calculates the equilibrium dissociation constant $K_D$ for a 1:1 interaction from a plot of steady state binding levels ($R_{eq}$) against analyte concentration ($C$). The equation includes a term for the bulk refractive index contribution $RI$, which is assumed to be the same for all samples. This term simply serves as an offset on the $R_{eq}$-axis.

$$R_{eq} = \frac{C R_{max}}{K_D + C} + RI$$

Model parameters and reported results are:

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Obtained from</th>
</tr>
</thead>
<tbody>
<tr>
<td>$KD$</td>
<td>Equilibrium dissociation constant [M]</td>
</tr>
<tr>
<td>$R_{max}$</td>
<td>Analyte binding capacity of the surface [RU]</td>
</tr>
<tr>
<td>$RI$</td>
<td>Bulk refractive index contribution in the sample</td>
</tr>
</tbody>
</table>

**Note:** Reported $K_D$ values that are higher than half the highest analyte concentration used should be treated with caution. If the response against concentration plot does not flatten out sufficiently because the concentrations are not high enough in relation to the $K_D$ value, the reported value may be unreliable. The reported $K_D$ value is marked as a vertical line on the fitting plot (see Section 9.2).

9.9 Creating and editing models

To create your own models for kinetics of affinity evaluation, choose **Tools:Models** from the main menu and select the type of model you want to work with. You can use existing models as templates. Choose an existing model from the list and click **New**: answer **Yes** in the following dialog to create a new model based on the chosen template or **No** to create a blank model. For kinetic models, you can define a new model either as a reaction scheme describing the interaction or as an equation defining response as a function of time. Interaction models are described in Section 9.9.1 and equation models in Section 9.9.2.

Predefined models cannot be edited or removed. If you want to modify a predefined model, create a new model using the predefined model as a template.
9.9.1 Interaction models for kinetics

The reaction scheme for an interaction model supports up to five component reactions. Follow the steps below to define a new model or edit an existing definition.

1. On the Interaction tab, click **New** to add new reactants. For each reactant, choose whether it is analyte, ligand or complex (see below) and enter an identifier for the reactant. Enter parameter names or expressions for the reactant properties.

   **Note:** Numbers are used as part of the identifier, not in the conventional chemical sense of stoichiometry. Thus a complex named AB2 does not imply two molecules of B binding to one of A.
### Analyte

The analyte is injected in solution at a constant concentration, and has the properties listed below. Analyte is usually denoted by the letter A.

<table>
<thead>
<tr>
<th><strong>Concentration</strong></th>
<th>Injected concentration in molar units.</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Mass transfer</strong></td>
<td>Check this box to include a mass transfer term in the fitting, and enter a parameter name or expression for the mass transfer constant.</td>
</tr>
<tr>
<td><strong>Molecular weight</strong></td>
<td>Check this box and enter a molecular weight if required. This information is used to calculate relative response contributions for heterogeneous analyte models (it is not used for conversion of weight-based to molar concentration units: this conversion is performed if necessary in the sample table).</td>
</tr>
</tbody>
</table>

### Ligand

The ligand is immobilized or captured on the surface, and has the properties listed below. Ligand is usually denoted by the letter B.

<table>
<thead>
<tr>
<th><strong>Binding capacity</strong></th>
<th>Maximum analyte binding capacity of the surface in RU.</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>At molecular weight</strong></td>
<td>This parameter is only used in heterogeneous analyte models. Check the box and enter the molecular weight parameter for the analyte to which the binding capacity parameter refers. Binding capacity for the other analyte will be calculated using the molecular weight values.</td>
</tr>
</tbody>
</table>

### Complex

The complex is formed on the surface and generates response and has the properties listed below.

<table>
<thead>
<tr>
<th><strong>Generates response</strong></th>
<th>Uncheck this box for complexes that form in solution and that do not contribute to the response.</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Molecular weight</strong></td>
<td>Check this box and specify a parameter for complexes that form in solution and then bind to the surface. Do not check this box if <strong>Generates response</strong> is also checked.</td>
</tr>
</tbody>
</table>
9 Kinetics and affinity analysis
9.9 Creating and editing models

In the *Bulk and Drift* panel, enter details for bulk refractive index contribution. Normally, there will be one bulk refractive index term applicable during association (from the start to the end of the injection). A second term can be used if necessary during dissociation (after the end of the injection), for example to accommodate a permanent shift in baseline as a result of the sample injection:

Check the *Drift* box and enter an expression describing the drift (most commonly a linear function of time) to account for baseline drift.

2 Enter the reaction scheme in the *Reaction* panel using the pull-down list for each reactant. Enter parameter names for the forward and backward rate constants for each line in the reaction scheme. (The terms *k-forward* and *k-backward* apply to the reaction as entered in the scheme, reading from left to right). You can also enter mathematical expressions or constant values for the rate constants.

3 Click the *Parameters* tab and define the parameters used in the reaction scheme. Click *Add* to add a new parameter, and define the parameter properties in the dialog:
Choose a default type for the parameter (Fit global, Fit local or Constant).

For the Initial value, enter a numerical value or select a value expression from the pull-down list. The expression represent functions evaluated within the current data set (e.g. \(Y_{\text{max}}\) is the maximum y-value in the data set). Alternatively, choose Attach to and select a parameter from the list. If you attach a parameter to Keyword, the initial parameter value will be set to the value of the keyword with the same name as the parameter.

Check Allow negative value if the parameter can be below zero. Enter a description of the parameter for ease of identification.

If you have only used single parameter names (as opposed to expressions) for the rate constants and properties, you can click Rate equations or OK as a shortcut to defining parameters. The software will then enter suggested definitions for all undefined parameters. This shortcut cannot be used if you have entered expressions.

In the Report panel, define the parameters you want to appear in the Report tab of the results. Report parameters are defined by a name that may be chosen freely and a value that is entered as a parameter or expression containing parameters.
9 Kinetics and affinity analysis
9.9 Creating and editing models

4 Click **Rate Equations** to display the equations generated by the software.

![Rate Equations Window]

You can select the equations in the display and click **Copy** to copy the equations to the Windows clipboard. Use this function and paste the equations in to e.g. Wordpad to print a copy of the rate equations.

### 9.9.2 Equation models for kinetics

Models for kinetic evaluation can also be entered as an expression defining response as a function of time t. To create an equation model, choose **New** in the kinetics models dialog, then choose to create the new model without using the currently selected model as a template. Select **Equation model** in the subsequent dialog.

![Editing Mode Window]
The example below shows a model for evaluation of the dissociation phase only.

Parameters and report parameters are defined in the same way as for kinetic models.
9.9.3 Models for steady state affinity

Models for steady state affinity evaluation are entered as an expression defining $R_{eq}$ as a function of concentration $Conc$. The example below shows a model for two-site affinity evaluation.

Parameters and report parameters are defined in the same way as for kinetic models.

**Note:** Beware of trying to define and use complex models for steady state affinity. Because of the relatively few points available for fitting to steady state affinity models (typically about five concentrations in duplicate), complex models tend to give unstable fitting behavior.
10 Thermodynamic analysis

Biacore T200 supports automated measurement of kinetics or affinity at a series of temperatures using the Thermodynamics wizard (Section 4.12). In addition to displaying the variation of kinetic and affinity constants with temperature, the evaluation software extracts standard thermodynamic parameters from the data.

10.1 Background

10.1.1 Equilibrium thermodynamics

For equilibrium thermodynamics, the van’t Hoff equation states:

\[ \Delta G^\circ = -RT \ln \frac{1}{K_D} = RT \ln K_D \]

where \( \Delta G^\circ \) is the standard Gibbs free energy change

\( R \) is the universal gas constant

\( T \) is the absolute temperature (K)

\( K_D \) is the equilibrium dissociation constant

Substituting in the expression

\[ \Delta G^\circ = \Delta H^\circ - T \Delta S^\circ \]

and rearranging gives:

\[ \ln K_D = \frac{\Delta H^\circ}{RT} - \frac{\Delta S^\circ}{R} \]

where \( \Delta H^\circ \) is the standard enthalpy change

\( \Delta S^\circ \) is the standard entropy change

A plot of \( \ln K_D \) against \( 1/T \) should thus be a straight line, with slope \( \Delta H^\circ/R \) and intercept on the y-axis \( \Delta S^\circ/R \).

This simplified relationship does not hold if the heat capacities of reagents and products differ, since different amounts of energy will be required to raise the temperature by the same amount on the two sides of the reaction. In such cases, the plot of \( \ln K_D \) against \( 1/T \) is not linear, and the relationship becomes
10 Thermodynamic analysis

10.1 Background

A value for the standard heat capacity change $\Delta C_p^\circ$ can thus be obtained in addition to $\Delta H^\circ$ and $\Delta S^\circ$ from non-linear fitting of the data to this extended equation.

10.1.2 Transition state thermodynamics

Transition state theory holds that the equilibrium constant for formation of the transition state in a reaction can be related to the rate constant for the overall reaction by the Eyring equation:

$$
K^\ddagger \equiv \frac{k \hbar}{k_B T}
$$

where $K^\ddagger$ is the equilibrium constant for formation of the transition state for the forward or back reaction

- $k$ is the kinetic rate constant for the interaction in the corresponding direction ($k_a$ or $k_d$)
- $\hbar$ is Planck’s constant
- $k_B$ is Boltzmann’s constant

Applying a similar rearrangement of the thermodynamic equations for the transition state gives:

$$
\ln \frac{k \hbar}{k_B T} = -\frac{\Delta H^\ddagger}{RT} + \frac{\Delta S^\ddagger}{R}
$$

so that the thermodynamic transition state constants for the forward and backward reactions can be obtained from plots of $\ln(k_o/T)$ and $\ln(k_a/T)$ respectively against $1/T$.

Note that the Eyring equation does not have a corresponding non-linear form that takes account of the heat capacity change for transition state formation. Non-linear fitting to obtain values for $\Delta C_p^\circ$ can only be applied to equilibrium thermodynamic analysis.
### 10.2 Performing thermodynamic analysis

Before thermodynamic analysis can be performed for a set of data, the kinetics and/or affinity must be evaluated at each temperature. Create a separate Kinetics/Affinity evaluation item at each temperature used in the run, using the same fitting model for each item (see Section 9.2).

When all required kinetic and affinity evaluation items have been created, click Thermodynamics on the toolbar.

Choose the sample and the fitting model in the pull-down lists. You may only choose one sample, and you should only choose one fitting model. Options for the model are **1:1 kinetics/steady state affinity** (recommended) or **All**. If you choose **All** it is possible to combine data from different fitting models in the same evaluation; however, values for thermodynamic constants are in all likelihood meaningless if the data is obtained from a mixture of different models.

Check the rows for data that you want to use in the thermodynamic evaluation. Use the **Check All** and **Uncheck All** as quick options to select and deselect the whole list.

**Note:** If you use data from fitting models that include multiple rate or affinity constants, be sure to select the correct rows so that equivalent constants are included from each fit. In some cases it may be necessary to examine the kinetic or affinity evaluation items to determine which constants belong together.
Click **Next**> when you have selected the data to be included. The results are displayed first as plots of affinity and rate constants against temperature.

Click **Next**> to display the van't Hoff and Eyring plots together with a table of thermodynamic constants for the equilibrium and transition state formation. In any of the plots, right-click on a point to exclude the point from the line fitting.
Choose whether to use a linear or non-linear fitting function for the van’t Hoff plot (see Section 10.1.1). If you choose non-linear fitting, a value for $\Delta C_p$ will be included in the reported parameters. Energies of activation ($E_a$), derived from the Eyring plots, are also listed for the transition states. All thermodynamic parameters are calculated for a temperature of 25°C.

In any of the plots, right click on a point to exclude it from the evaluation.

**Notes:** Regardless of the setting for the van’t Hoff plot, the Eyring plots are always fitted to a linear function. Calculation of $\Delta C_p$ by non-linear fitting is not valid for transition state data (see Section 10.1.2).

If you have combined kinetic and steady state affinity data in the thermodynamic evaluation, the van’t Hoff plot will show all affinity values, but the Eyring plots will be empty because the steady state data lacks values for the rate constants.

Plots of kinetic and affinity constants against temperature show temperature values in °C, while van’t Hoff and Eyring plots use absolute temperature values (K).

Click on **Finish** to finalize the thermodynamic analysis.

When assessing the validity of thermodynamic constants reported by this analysis procedure, pay particular attention to the kinetic analysis at different temperatures. With complex interactions involving macromolecules, there is a significant possibility that the characteristics of the interaction (including the role of mass transport limitations in the observed interaction) change with temperature, resulting in different fitting quality at different temperatures. This may be evident from direct comparison of the kinetic fits, but will not be immediately apparent in the thermodynamic analysis.
10 Thermodynamic analysis
10.2 Performing thermodynamic analysis
11 Affinity in solution

Determination of affinity in solution provides an alternative to steady state affinity measurements (see Chapter 9) for interactions that take a long time to reach equilibrium or for any other reason are difficult to determine with a direct binding assay. In principle, the affinity in solution approach uses Biacore to determine the free concentration of one interactant in equilibrium mixtures containing known total interactant concentrations.

11.1 Conventions and background

11.1.1 Experimental setup

The interactants in affinity in solution determination are denoted A and B:

\[ A + B = AB \]

Experiments are set up so that a fixed concentration of B is mixed with variable concentrations of A and allowed to reach equilibrium. The free concentration of B is then determined by injecting the sample over a ligand that binds B but not A or the complex AB (the interactant A or a derivative thereof is usually suitable as ligand). It is assumed that the measurement itself does not significantly disturb the equilibrium in the sample.

The experimental setup requires a calibration curve with known concentrations of B determined over the same sensor surface, in order to calculate the free B concentrations in the samples.

11.1.2 Evaluation principles

The equilibrium constant for a 1:1 interaction is given by

\[ K_D = \frac{A_{\text{free}} \cdot B_{\text{free}}}{AB} \]

or

\[ K_D = \frac{(A_{\text{tot}} - AB)(B_{\text{tot}} - AB)}{AB} \]
11 Affinity in solution

11.2 Requirements for affinity in solution

Rearranging gives

\[ K_D \cdot AB = A_{tot} \cdot B_{tot} - AB(A_{tot} + B_{tot}) + AB^2 \]

or

\[ AB^2 - AB(A_{tot} + B_{tot} + K_D) + A_{tot} \cdot B_{tot} = 0 \]

Solving for AB:

\[ AB = \frac{(A_{tot} + B_{tot} + K_D)}{2} \pm \sqrt{\left(\frac{A_{tot} + B_{tot} + K_D}{4}\right)^2 - A_{tot} \cdot B_{tot}} \]

Substituting in the relationship \( B_{free} = B_{tot} - AB \) gives

\[ B_{free} = \frac{(B_{tot} - A_{tot} - K_D)}{2} \pm \sqrt{\left(\frac{A_{tot} + B_{tot} + K_D}{4}\right)^2 - A_{tot} \cdot B_{tot}} \]

This equation can be fitted to a plot of \( B_{free} \) against \( A_{tot} \) to calculate a value for \( K_D \). (Formally, the equation has two solutions, but one is always negative and is not meaningful in the context of an affinity determination.)

11.2 Requirements for affinity in solution

Affinity in solution experiments are run using a method. The method must be correctly constructed as described in Section 5.10.4: if necessary, the keyword table can be edited so that the conditions are met in full (see Section 6.6). Note however that the command type cannot be edited in the keyword table. Refer to Chapter 5 for details of how to construct methods in Method Builder. Determination of affinity in solution is not supported by a wizard.
11.3 Evaluation of affinity in solution

To evaluate affinity in solution measurements, open the result file and click **Affinity in Solution** on the toolbar. The first step displays the calibration curve for measurement of free B:

Choose the sensorgram, report point, response type and fitting function from the pull-down lists at the top of the dialog. See Section 8.2.1 for more details of these choices.

If you have run multiple sample series in the experiment, choose the sample to evaluate in the **Sample** list. A sample series is defined as all cycles with the same sample name in the assay step(s) with purpose **Sample**.

If you have run multiple calibration curves in the experiment, choose the curve to use in the **Calibration curve** list. A calibration curve is defined as measurements from assay step(s) with purpose **Calibration**, regardless of the sample name. If two or more **Calibration** assay steps are run contiguously with no intervening steps with a different purpose, they will be combined into a single calibration curve.

**Note:** All samples in a series are evaluated against the chosen calibration curve. You cannot use different calibration curves for different samples in the same series.

The table lists the data for the calibration curve. The plot panel shows the curve with calibration points as black inverted triangles and sample points as red squares. Samples that lie outside the range of the calibration curve are not shown. Right click on calibration points to exclude the points from the curve.
The table shows the numerical results for the sample series. Samples that lie outside the range of the calibration curve are marked as \textit{N/A} (not applicable) in the column for \textit{Calc. Conc. Beq}.

The plot panel shows the sample measurements with a line fitted according to the equation for 1:1 affinity (see Section 11.1.2). Right-click on a point to exclude it from the fitting.

\textbf{Notes:} The plot of free B against total A is presented by default with a logarithmic scale on the x-axis.

Zero values cannot be plotted on a logarithmic scale. If you have included a sample with zero concentration of A in the sample series and want to display this point on the plot, choose \textit{Scale} from the right-click menu in the plot panel and set a linear scale for the x-axis.

The intercept of the fitted curve on the y-axis represents a fitted value for the parameter \textit{ConcB}. This value should be the same as or close to the value entered for the variable \textit{ConcB} in the method.

The calculated $K_D$ value is shown in the panel below the table.
Evaluation of immunogenicity experiments is supported by 4 application-specific tools in the evaluation software:

- **Screening**, based on ranking plots for detection of antibody responses.
- **Confirmation**, for evaluating antibody specificity as tested by inhibition of responses by added drug. Confirmation evaluation can be based either on the degree of inhibition by added drug or on a cut-off boundary to classify inhibition in yes/no terms.
- **Isototyping**, for identifying positive responses from isotyping reagents. Isotyping results are presented as bar charts with cut-off boundaries to exclude negative responses from the chart.
- **Stability**, which assesses the rate dissociation of detected antibody from the antigen using a two-site dissociation model.

Evaluation of immunogenicity studies is described in detail in the separate Biacore T200 Immunogenicity Handbook.
12 Immunogenicity
Appendices
Appendix A  Data import and export

This appendix describes the functions and data format for data import and export.

A.1  Exporting data

A.1.1  Export functions

Data can be exported from both the Control and Evaluation Software in Microsoft Excel or Extended Markup Language (XML) format. The report point table can be exported to a tab-separated text file. Rack positions may also be exported from the Rack Positions dialog box.

Export to Excel

To export data to an Excel file, choose File:Export:Results to Excel.

Export from the Control Software creates an Excel spreadsheet file (extension .xls) containing separate worksheets for the file properties and report point table. The audit trail is also exported to a separate worksheet if it is present when the GxP module is installed (see the separate Biacore T200 GxP Handbook).

Export from the Evaluation software an Excel spreadsheet file (extension .xls) containing separate worksheets for the file properties and for tabulated data for all evaluation items where appropriate (i.e. plot data and evaluation results). The worksheets for each item are identified with the item name. For plots, only the columns shown in the plot window table are exported. Data from sensorgram items is not exported. The audit trail is also exported to a separate worksheet if it is present when the GxP module is installed (see the separate Biacore T200 GxP Handbook).

Export to XML

To export data to an XML file, choose File:Export:Results to XML.

This option exports the same data as Results to Excel but creates a text file in XML format (file extension .xml). Details of the XML format may be determined by exporting data from the Control or Evaluation Software and opening the exported file in an XML-compatible editor.
Appendix A  Data import and export
A.2  Importing data

**Report point table**
To export the report point table to a tab-separated text file, choose
*File:Export:Report Point Table*. The exported file has the extension *.rpt*.

**Note:** If you open an exported report point table in Microsoft Excel, make sure that the format for the *Fc* column is set to *Text* in the Excel import file wizard. The default setting of *General for text file import* may interpret the flow cell identification for reference-subtracted data as a date instead of a text string.

**Rack positions**
Rack positions can be exported from the Control Software to a tab-separated text file in either ASCII or Unicode format using the *Menu:Export Positions* function in the *Rack Positions* dialog (Section 4.2.6). The file contains two lines identifying the microplate and reagent rack settings followed by the contents of the rack positions table with the columns separated by tabs.

A.2  Importing data

**A.2.1  Control Software**
The Control Software supports data import to sample tables in assay wizard templates and in the *Setup Run* step of methods and to position information to the *Rack Positions* step in all runs. In order to use the import function, the option must be activated in *Tools:Preferences* and valid import programs must be specified. Import programs and data files for use with import are the responsibility of the user.

For each import function, a check box allows the import program to be started automatically without user intervention. If the respective box is not checked, the
program will only be started when the user actively requests data import. If the box is checked:

- Data is imported automatically to the sample table only if the table is empty. The program is not started if the table already contains data.

- Data is imported to the Rack Positions dialog whenever automatic positioning is invoked. This happens when the dialog is first opened with new or modified sample data, and also when the user requests Automatic positioning from the dialog menu (Section 4.2.6). The program is not started when the user makes manual changes to the rack positioning or when the dialog is opened with no changes in previously positioned samples (for example when the user clicks Back and Next in the dialog sequence without changing sample information).

Sample table import

When the sample import function is invoked, the contents of the sample table are first exported in Extended Markup Language (XML) format to a temporary file that is submitted to the specified import program. The import program may append new sample data to the file or overwrite the file contents with new data as required. The modified file is then imported back into the sample table and the temporary file is deleted.

Development or choice of a suitable import program is the responsibility of the user. To document the detailed XML format of the import file, specify an XML-compatible text editor as the import program and save a copy of the import file from a suitable table.

```xml
<xml version="1.0" encoding="iso8859-1" ?>
<!DOCTYPE MethodBuilderImport [ View source for full doctype ... ]>
  <MethodBuilderImport importFileVersion="1.0">
    <AssayStep name="Startup">
      <DataTable row="1">
        <Data cmd="Sample 1" grp="" fid="Solution" val="Sample 1/2" />
        <Data cmd="Sample 1" grp="" fid="Conc1" val="0" />
        <Data cmd="Sample 1" grp="" fid="Conc2" val="125" />
        <Data cmd="Sample 1" grp="" fid="MW1" val="200" />
        <Data cmd="Sample 1" grp="" fid="MW2" val="10000" />
      </DataTable>
    </AssayStep>
    <AssayStep name="Sample">
      <DataTable row="1">
        <Data cmd="Sample 1" grp="" fid="Solution" val="Sample 1/2" />
        <Data cmd="Sample 1" grp="" fid="Conc1" val="0" />
        <Data cmd="Sample 1" grp="" fid="Conc2" val="0" />
        <Data cmd="Sample 1" grp="" fid="MW1" val="200" />
        <Data cmd="Sample 1" grp="" fid="MW2" val="10000" />
      </DataTable>
    </AssayStep>
    <DataTable row="2">
      <Data cmd="Sample 1" grp="" fid="Solution" val="Sample 1/2" />
      <Data cmd="Sample 1" grp="" fid="Conc1" val="0" />
      <Data cmd="Sample 1" grp="" fid="Conc2" val="195" />
      <Data cmd="Sample 1" grp="" fid="MW1" val="200" />
      <Data cmd="Sample 1" grp="" fid="MW2" val="10000" />
    </DataTable>
  </MethodBuilderImport>
</xml>
```

Figure A-1. Part of an XML import file from a method for heterogeneous analyte kinetics displayed in an XML-compatible editor.
A.2 Importing data

**Rack positions import**

The **Menu:Custom Position Import** and **Simple Position Import** functions in the **Rack Positions** dialog (Section 4.2.6) import rack position data from an external file such as one from a laboratory robot used to prepare sample microplates. If you choose **Custom Position Import**, the external file is first processed by the import program as specified in **Files:Preferences**. Output from this program must be tab-separated text in either ASCII or Unicode format conforming to the specification below. The **Simple Position Import** option imports data from a file conforming to the specifications with no intervention from an external program.

- Two lines in the file specify the microplate and reagent rack settings, in the format:
  
  Rack1=<microplate specification>
  Rack2=<reagent rack specification>

- Specifications are not case-sensitive, but microplate and reagent rack specifications must be given otherwise exactly as they appear in the selection lists in the **Rack Positions** dialog. If either specification is invalid, the corresponding definition will not be imported. The position of these two lines in the file does not matter.

- One line specifies the headers for table columns to be imported, separated by tabs. The headers should correspond to the column headers as they appear in the **Rack Positions** table, with the exception of the **Volume** column in the table which can be omitted from the import file (and is ignored if it is present). This line may not be preceded by any line other than the microplate and reagent rack specifications.

- A set of lines hold the content of the table columns separated by tabs. Each line must contain the same number of tab characters as the header line.

When import is requested, the contents of each table line in the import file are matched as far as possible to the contents of the **Rack Positions** table, with the exception of the **Position** and **Volume** column. For matched rows, the **Position** in the table is replaced by the value in the **Position** column from the import file. Rows for which a match cannot be found are not imported. Any rows in the **Rack positions** table which do not have a matching row in the import file are left without a **Position** specification and must be placed in the microplate or reagent rack before the run can be started.

Details of the required import file format can be investigated further by examining a file created with the **Menu:Export Positions** command.
Figure A-2. Example of an exported file for rack positions, opened in Microsoft Excel.

A.2.2 Evaluation Software

The Evaluation Software supports import of model definitions for kinetics and affinity evaluation. Model files for import should be obtained from GE Healthcare or created by exporting models from another installation.
Appendix A  Data import and export
A.2  Importing data
Appendix B  Method examples and recommendations

A selection of predefined methods covering common applications is provided in the **Biacore Methods** folder (see Section 5.1). Use these methods either directly or as starting points for your own method development. This section describes the essential features in each method that are not supported in wizards. Refer to these methods as guidelines in constructing your own methods that exploit similar features.

### B.1 Affinity in solution

This method is designed for measurement of affinity in solution as described in Chapter 11. The method includes a **Calibration** assay step for measurement of component B and a **Sample** step for measurement of mixtures of components A and B. Both these assay steps are connected to the same cycle type.

Predefined evaluation variables are included for the evaluation purpose **Affinity in solution**.

The **Variable Settings** are different for the two assay steps, so that only the relevant variables are entered at run-time for each assay step.
Appendix B  Method examples and recommendations
B.2  Calibration-free concentration analysis

B.2  **Calibration-free concentration analysis**

Requirements and recommendations for running calibration-free concentration analysis are provided in this method.

B.2.1  **Assay steps and general settings**

Blank cycles are run before the samples and are repeated every 12 sample cycles. Assay steps for samples and blanks are connected to the same cycle types:

Analysis is performed at a data collection rate of 10 Hz (set in the General workspace). Do not change this setting.

B.2.2  **Cycle types**

The cycle type for sample analysis performs a *High performance* injection with a contact time of 36 seconds and a dissociation time of 5 seconds. The dissociation time is not critical: evaluation uses only data from the association phase. Do not use shorter contact times than 36 seconds, since the actual contact time (which is determined by the injected volume, rounded to the nearest µl) may differ significantly at the lowest flow rate.

Sample solution and flow rate are set as method variables. For evaluation variables, the evaluation purpose is set to *Calibration-free conc* and predefined variables *MW, D(20°C), Blank* and *Dilution* are required.
B.2.3 Variable settings

Sample cycles

All the variables for assay step Sample are set at run time, except for the method variable Blank which is set to No in the method.
Blank cycles
For assay step **Blank**, only the flow rate is set at run time. The method variable **Blank** is set to **Yes** in the method and **Sample solution** to **Buffer**. The remaining variables are not used and may be left blank.

B.2.4 Setup Run
At run time, variables need to be assigned for assay steps **Sample** and **Blank**. Each sample should be run at two or more flow rates: values of 5 and 100 µl/min are recommended and are entered for one sample. Use the same set of flow rates for each sample. Values are also required for analyte molecular weight (**MW**), diffusion coefficient at 20°C (**D(20°C)**) and dilution factor (**Dilution**).

Only flow rates are required for assay step **Blank**: these values must be the same as the flow rates used for samples.
B.3  **CAP single-cycle kinetics**

This method implements single-cycle kinetics using reversible capture of biotinylated ligand on Sensor Chip CAP (supported by the Biotin CAPture Kit from GE Healthcare). This section focuses on the method aspects required for using the Biotin CAPture Kit. If you want to create a method for other applications using the Biotin CAPture Kit, modify the **Sample** cycle definition in the CAP single-cycle kinetics method. See Section B.10 for aspects of the method directly related to single-cycle kinetics.

Follow the Instructions for Use supplied with the Biotin CAPture Kit before using a new Sensor Chip CAP:

### B.3.1 Assay steps

The method includes a **Conditioning** step run once before the start-up cycles.
This step injects three 1-minute pulses of regeneration solution, and should always be included at the start of an assay using a new sensor chip. The step can be omitted if the sensor chip has been used previously.

B.3.2 Sample analysis cycle for Sensor Chip CAP

The cycle definition for sample analysis with Sensor Chip CAP includes 4 injection commands:

- **General** for injection of the Biotin CAPture reagent
- **Capture** for injection of the biotinylated ligand
- **Sample** for injection of analyte
- **Regeneration** to remove the Biotin CAPture reagent, ligand and bound analyte
An extra wash with buffer after the injection is included in the *Regeneration* command.
B.4 GST kinetics

The recommended method for using a GST-tagged ligand is set up with Dual detection in General Settings. Detection settings Multi and Single are not recommended. The method involves 5 conditioning cycles with injection of recombinant GST (provided in the GST Capture Kit from GE Healthcare) over both flow cells followed by regeneration:

The sample analysis cycle uses two capture injections, one to capture recombinant GST on the reference surface only (flow path setting First) and the second to capture GST-tagged ligand on the active surface only (flow path setting Second). A stabilization time of 180 seconds is included after capturing the ligand on the active surface, to allow for dissociation of any weakly bound components. The sample is then injected over both surfaces. This approach provides a reference surface that mimics the active surface with respect to occupancy of the anti-GST capturing sites.
This method is set up for kinetic analysis but can be adapted for other assays.

**Note:** If you change the detection setting to **Multi**, you will need to include additional capture injections for the active surfaces in flow cells 3 and 4. Do not use detection setting **Single** without removing the first capture injection.
Appendix B  Method examples and recommendations  
B.5  Inject and Recover

B.5  Inject and Recover

The example method provided for using the InjectAndRecover command is targeted to recovery for mass spectrometry, and contains two assay steps. The first conditions the surface by washing three times with 0.5% trifluoroacetic acid, while the second performs the sample injection and recovery operation.

The cycle for recovery of bound analyte contains just one single InjectAndRecover command. Additional commands are usually not needed, since the recovery component of the InjectAndRecover command serves as a regeneration step (see Section 5.6.1).
All parameters in the **InjectAndRecover** command are fixed: check the appropriate boxes in the **Method Variables** list if you want to use variable parameters. There are no evaluation variables for this command.
Appendix B  Method examples and recommendations

B.6  Kinetics heterogeneous analyte

This method is a straightforward kinetics determination with evaluation variables included for the evaluation purpose *Kinetics – heterogeneous analyte*, providing separate concentration and molecular weight variables for two analytes.

---

B.7  L1 liposome capture

The recommended method for liposome capture on Sensor Chip L1 uses one conditioning cycle consisting of two injections of 40 mM octylglucoside, followed by three start-up cycles using a dummy sample before the actual samples are run.
B.8  **LMW kinetics and LMW Screen**

Methods are provided for both kinetics and screening of low molecular weight compounds. The essential addition to wizard-based counterparts is the inclusion of a *Solvent Correction* assay step.

This assay step is connected to a cycle type that includes 8 *Solvent Correction* commands for injection of 8 different solvent concentrations (see Section 6.7).
Screening applications suitably include a carry-over control injection, to identify potential carry-over problems from “sticky” compounds that may affect the response in subsequent injections. The Biacore method is designed for screening of low molecular weight compounds and includes a solvent correction step as described in Section B.8. Low molecular weight compounds frequently dissociate readily from their targets, and the example method does not include regeneration. If you require a regeneration step, it is advisable to include regeneration after both the sample and the carry-over injections.
B.9 **LMW single-cycle kinetics**

This method combines functions for working with low molecular weight analytes (Section B.8) with the Sample command for single-cycle kinetics (Section B.10).

B.10 **Single-cycle kinetics**

The Biacore method for single-cycle kinetics uses the sample command type *Single cycle kinetics* with 5 sample concentrations per cycle. Note that with this command type, the evaluation purpose is fixed as *Kinetics/Affinity* and the number of predefined variables for sample concentration corresponds automatically to the number of sample concentrations per cycle. The predefined variables cannot be deselected.
Index

Numerics
1:1 binding 187

A
aborting fitting procedure 166
absolute response 9, 127, 137
accepting fits 166
active surface 9
adaptive methods 92
adding keywords 115, 116
adding report points 114
adjust injection start 162
adjustment for controls 132
adjustment for molecular weight 127
affinity 159, 165, 168
application wizard 65
affinity in solution 95, 159
evaluation 209
experimental design 209
method example 223
requirements for evaluation 102, 210
variables 95
aim for immobilized level 41, 46, 48
aligning sensorgrams 125
alignment 125
analysis temperature 7, 20, 36, 81, 84
thermodynamics 71
analysis temperature after run 81
analyte 8, 197
analyte concentration 94
analyte molecular weight 68, 94
adjusting response 127
anchor 39
append file 108, 149, 169
application wizards 13, 29
applying solvent correction 119
assay methods 77
assay setup 32
assay step name 82
assay step purpose 82, 115
assay steps 78, 81
analysis temperature 84
buffer change 84
connecting to cycle type 84
recurrence 84
replicates 85
run order 85
auto scale 18
automatic positioning 39
autosampler 62
average calibration curve 141, 143
avidity 188

B
bar charts 135
display options 136
baseline 96
for report points 96, 114, 138
in regeneration scouting 52
in sensorgram display 18
plot 112
batch mode 159, 170
Biacore methods 77, 223
Biacore terminology 8
BIAnormalizing solution 36
binding analysis 58
results 60
binding capacity 197
binding levels 113
binding partners 58
binding to reference 112
biological significance 184
bivalent analyte 188
blank cycles 95, 141, 152, 160, 224
blank immobilization 46
blank method 47
blank subtraction 126, 153, 162
boundaries 134
boundaries for ranking 134
browse buttons 124
buffer 84
buffer bottles 81
buffer change 84
buffer names 81
buffer requirement 41
buffer scouting 54
results 55
buffer selector 7
bulk contribution 117
bulk refractive index 66, 173, 198

C
calc conc 147, 155
calculated concentration 146
calculated concentrations 143
calculated parameters 199
calibration curve 61, 62, 83, 141, 142
calibration curves 211
calibration trends 141, 144
calibration-free concentration assays 61, 95, 102, 141, 150, 224
caption 110
capture 31, 56, 88
  flow path 31
  plot 112
capturing molecule 8
carry-over control 89, 236
  plot 112
carry-over injection 65
changing rack positions 38
check kinetic data 176, 178
chip properties 16, 17, 49
chip type 31
chi-squared 131, 175
clear positions 38
coefficient of determination 131
coefficient of variation 147
color by 111, 123, 136
color-coding 37
column labels 136
combined result sets 149
command queue 23, 25
commands
  in manual run 25
  in Method Builder 87
competing analytes 189
complex 197
components 177
concentration analysis
  application wizard 61
  combined result sets 149
  evaluation 141
  requirements for evaluation 101, 141
concentration series 68, 160, 161
concentration unit 81, 116
conditional command 92
conditional statements 41
conditioning 32
conditioning cycle 83
conformational change 193
connect to cycle type 84
constants 185
contact time 33, 46
  at low flow rates 33
control experiments 73
  results 74
control samples 34, 60, 83, 132
  binding analysis 59
  concentration analysis 63, 146
  kinetics/affinity 68
  plot 112
control software

Edit menu 17
File menu 15
Run menu 19
screen regions 14
Tools menu 19
View menu 18
copy graph 20, 111
copy table 111
correlated parameters 172, 177
create
  assay step 81
  cycle type 86
  evaluation method 121
  fitting model 195
  method 77
  wizard template 29
curve 9, 19
curve fitting 130, 184
curve name 124, 128
curve type 171
custom methods for immobilization 47
custom position import 220
custom report points 14, 95, 113
CV 147
cycle 124
  manual run 27
cycle run list 36, 100
cycle types 78, 84, 86
D
data collection rate 80
data import 20
data presentation 123
delete
  keywords 115
  report points 115
deposition solution 90
detecting molecule 61
detection 80, 98
  in wizards 31
diffusion coefficient 95, 142, 156, 186
diluting ligand for immobilization 46, 48
dilution factor 64, 94, 147, 155
dimethyl sulfoxide 66, 71, 117
dissociation time 59, 67, 88
disturbed data 154
DMSO 66, 71, 117
drift 198
dual detection 80
dual inject 91
dummy sample 32
E
EDC/NHS 49
Edit menu
control software 17
editing keywords 115
editing report points 115
in control software 17
in evaluation software 113
eject chip 20, 27
eject rack 20, 24, 27
manual run 26
ending a manual run 27
enhancement 31, 56, 89
plot 113
enthalpy 203
entropy 203
epitope mapping 58
estimated buffer requirement 41
estimated run time 41
evaluating concentration analysis 141
Evaluation Explorer 108
evaluation items 108
evaluation methods 121
evaluation of kinetics and affinity 159
evaluation procedures 121
evaluation purpose 93, 171
evaluation requirements 101
evaluation software
general features 107
screen regions 107
evaluation variables 93, 115
event log 14, 19
event markers 19, 126
Excel 217
excluded cycles 143
excluded volume effect 118
excluding data
from concentration analysis 143, 147, 148
from evaluation 110
from kinetics/affinity
evaluation 162
from plots 129
expand all cycles 150
expected concentration 63, 147
export 15
export curves 20, 111
export rack positions 40
export to Excel 217
export to XML 217
exporting data 217
exporting plots 111
exporting rack positions 40, 218
exporting the report point table 218
extra wash after injection 87
extrapolating solvent correction
curves 121
Eyring equation 204
F
File menu
control software 15
file name extensions 21
file storage 21
filter keywords 116
filtering report point table 138
fit by color 130
fitting curves to plots 130
fitting function 142, 144
fitting models 148, 157, 166, 171
fitting parameters 166, 168
flow cell 7, 9, 45, 80
flow path 80, 98
for buffer scouting 54
for capture 31
in wizards 30
manual run 24, 26
method runs 98
flow rate 33, 46, 141, 151, 186
effect on binding rate 73
manual run 24, 26
range 33
folder for methods 77
folder for wizard templates 29
four-parameter fit 62, 131, 144
free energy 203
G
general command 91
general settings 80
global parameters 168, 185
gridlines 21, 111
in wizards 136
GST Capture Kit 230
GST-tagged ligand 230
GxP compliance 8
H
heat capacity 204
heterogeneous analyte 94, 189
method example 234
heterogeneous ligand 191
high performance injection 88, 91
I
if...then command 92
immobilization 45
methods 45
results 49
immobilization pH scouting 42
import program 219
import rack positions 40
importable wizard templates 77
importing rack positions 40, 220
importing sample tables 59, 64, 68, 99, 218
importing wizard templates into Method Builder 77
included blanks 153
initial values 199
inject and recover 90, 232
inject sample
    manual run 26
injection parameters 33
injection sequence 30
injection volumes 33
injections 31, 87
insert chip 20
instrument preparation 23
instrument preparations for manual run 23
interaction kinetics 159
interaction models 196
interpolated calibration 141
K
keyword table 14, 115
kinetic constants 172
kinetics 159, 165
    predefined models 186
kinetics control experiments 73
kinetics/affinity
    application wizard 65
    calculated values 168
    evaluation 159
    requirements for evaluation 102, 160
    subsets 169
kinetics/affinity report 168
$k_m$ 186
$k_i$ 186
L
labels in plot windows 110
legend 111
ligand 8, 197
ligand capture 31
ligand immobilization 45
limit guides 179
linear fit 131, 144
linked reactions 73
control experiment 73
calculated values 168
control software 14
evaluation software 107
maintenance tools 20
manual run 13, 23
    initial settings 24
markers 126
mass spectrometry 7, 90
mass transfer 73, 179, 197
    control experiment 73
    control results 74
mass transfer coefficient 186
mass transfer constant 176, 186
mass transfer parameters 186
mass transport limitation 151
measured conc 155
menus 14
Method Builder 13, 77
    commands 87
    importing wizard templates 77
method examples 223
method files 21
method overview 78, 79
method structure 78
method variables 93
methods 77
    assay 13
    evaluation 121
    folder 21, 77
microplate 23, 38, 39
    in manual run 24
Microsoft Excel 217
mix 61, 89
    in concentration analysis 61
mix and inject 48
mixing sample before injection 89
modification factor 178
molar concentrations 68
molecular weight 142
    of analytes 68
multi detection 80
multi-component complex 58
multi-cycle kinetics 65, 159
multiple data sets 162
multiple ligand densities 169
evaluation 170
multiple result files 108
multiple $R_{\text{max}}$ 162, 169

N
nested assay steps 81
new cycle in manual run 27
normalize 36
normalizing sensorgram display 125
notebook 19, 21

O
octylglucoside 234
opening files for evaluation 108
optimum buffer pH 44
optimum immobilization pH 44
orientation 39
overview 79

P
parameter significance 176
parameters button 171
parameters for kinetics/affinity 166
pausing a manual run 27
pH range 43
pH scouting 42
results 44
plot windows 127
fitted curves 130
table columns 129
polynomial fitting 132
pooling 39
preconcentration 42, 43, 46, 48
predefined
evaluation items 112
evaluation variables 93
immobilization methods 45
methods 77, 223
models 186
plots 112
pre dip 87
preferences 40, 77
prepare run protocol 41
prime 19, 36
print layout 16
print range 16
printing rack positions 40
printing results 16, 109
printing wizard templates 40

Q
QC ratio 151
acceptance 151
quality control 167, 172
queue 23, 25

R
rack 39
rack illumination 20
rack positions 37
automatic import 40
default positions 38
export 40, 218
import 40, 220
rack settings in manual run 24
ranking 134
rate equations 199, 200
reactants 196
reagent rack 23, 38, 39
recovering bound analyte 90
recovery solution 91
recurrent assay steps 84
reference line 18
reference subtraction 118, 126
reference-subtraction 9
regeneration 9, 31, 34, 89
manual run 26
regeneration conditions 51
regeneration scouting 50
results 52
regeneration solutions
viscosity 26, 34, 51, 89
regions 39
regression coefficient 138
regulated environments 8
relative response 9, 114, 127, 138
reliability of $K_D$ values 195
remove selection 164
removing data from sensorgram display 124
renaming keywords 117
repeat calibration 62
replicate assay steps 85
replicate concentrations 62
report point id 114, 137
report point markers 19
report point range 121
report point table 14, 21, 136
export 218
report point window 114, 137
report points 9, 95, 113, 136
adjustment for molecular weight 127
editing in control software 17
in regeneration scouting 52
markers 126
$R_{\text{eq}}$ against C 165
requirements for evaluation 101
requirements for kinetics and affinity evaluation 160
residual analyte 65
residuals 168, 174
resonance unit 9
response bound 49
response correction factor 190
response type 127
result files 21
right-click menu 20, 25, 99, 109
Run menu
control software 19
run order 68, 85
run protocol 41
run time 97
running buffer 84
running wizards 30
sample capacity 7
sample command 88
sample compartment
  illumination 20
  temperature 7, 36, 71, 80, 84
sample concentrations 68
sample injection 31, 34
  binding analysis 58
sample mixing in concentration analysis 61
sample name in calibration curve 144
sample names 162
sample parameters
  binding analysis 59
sample response
  in regeneration scouting 52
samples 34
  for calibration curve 62
  for concentration analysis 64
  for kinetics/affinity 68
scale 18, 20, 109, 111
screen regions
control software 14
evaluation software 107
screening 58
  method example 236
SD 138
selecting data for thermodynamic evaluation 205
selector bar 15, 124, 128
selector button 124
Sensor Chip L1 234
Sensor Chip SA 48
sensor chip type 31
sensorgram 9
adjustment 125
baseline 18
copy graph 20
curvature 173
display 123
export curves 20, 111
in evaluation 123
markers 19
normalizing 125
printing 16
reference line 18
removing data for display 124
window 14
service tools 20
settings for regeneration scouting 51
settings for $R_{eq}$ 165
setup run 98
short-term noise 174
show sensorogram 110
shutdown 19
significance 176
simple position import 220
simulate cycle run list 82
single detection 80
single-cycle kinetics 65, 88, 159, 161, 237
slope 138
sodium acetate buffers 43
software help 15
solvent correction 66, 71, 83, 90, 117
  applying 119
  extrapolation 121
  method example 235
  principle 119
  requirements for evaluation 103
solvent correction curves 120
  fitting 120
  shape 121
  slope 121
sort keywords 116
sorting
  plots 130
  report point table 138
specifying variable values 97
SPR technology 7
stabilization period 87, 89
stabilization time 34
standard deviation 138
standard error 176
standby 19
start-up cycles 32, 83
  in regeneration scouting 50
statistical parameters 175
status bar 14, 108
steady state affinity 165, 195, 202
stop command in manual run 27
stop run 27
subset 169
surface performance 56
results 57
system caption 110
system overview 7
system preparations 35, 100

T

table 129
in plot windows 129
table columns 129
target immobilization level 46
tc 186
temperature 20, 23, 36
temperature control 7
temperature dependence of kinetics and affinity 206
temperature equilibration 41, 71
template types 29
templates 29
folder 21, 29
importing into Method Builder 77
terminology 8
test tools 20
thermodynamic parameters 203
thermodynamics 159
application wizard 70
evaluation 203
requirements for evaluation 102
tool tips 123
toolbar 14
Tools menu
control software 19
transition state 204
T-value 176
two-state reaction 193

U

uniqueness 172, 177
unknown samples 141
unzoom 18, 109, 111
user-defined caption 110
user-defined variables 93
using calibration trends 145
U-value 177

V

van’t Hoff equation 203
variable settings 97
variables 78, 93, 97, 115, 127
entering values at run time 98
entering values in a method 97
verification 79, 98
vial size 39
view curve 19
View menu
control software 18
viscosity 26, 34, 51, 89

W

wait 26
weight-based concentrations 68
window 96, 114
wizard components 30
wizard results 19
wizard templates 29, 77
wizards 13, 29
importing into Method Builder 77
work area 108

X

XML 217, 219

Z

zero concentration samples 68, 160, 162
zero response point 125
zero time point 125
zoom 18, 109
zoom lock 53, 109