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**LIST OF FIGURES (continued)**

**LIST OF TABLES**

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# 5.0   INTRODUCTION TO THE SYSTEM SOFTWARE

The BioLogic Duo-Flow system software is based on Microsoft Windows®, a widely used operating system for PC-compatible computers. This chapter discusses how to use the BioLogic Duo-Flow system software.

## 5.1   USING THE SYSTEM INTERFACE

The Manual control screen (Figure 5-1) is the first screen displayed when you turn on the system. This screen, like all BioLogic screens, is grouped into the Main (or System) Menus, the System Control Window, and the System Status Bar. The contents of the System Control Window are discussed in Chapters 6 and 7 and in the separate documentation for the Econo Gradient Pump (EGP) and the QuadTec UV/VIS Detector.
- Browser screen: Chapter 6
- Manual Control screen: Chapter 7, section 7.1
- Setup screen: Chapter 7, section 7.2
- Protocol screen: Chapter 7, section 7.3
- Run screen:  Chapter 7, section 7.4
- Post-Run screen:  Chapter 7, section 7.5
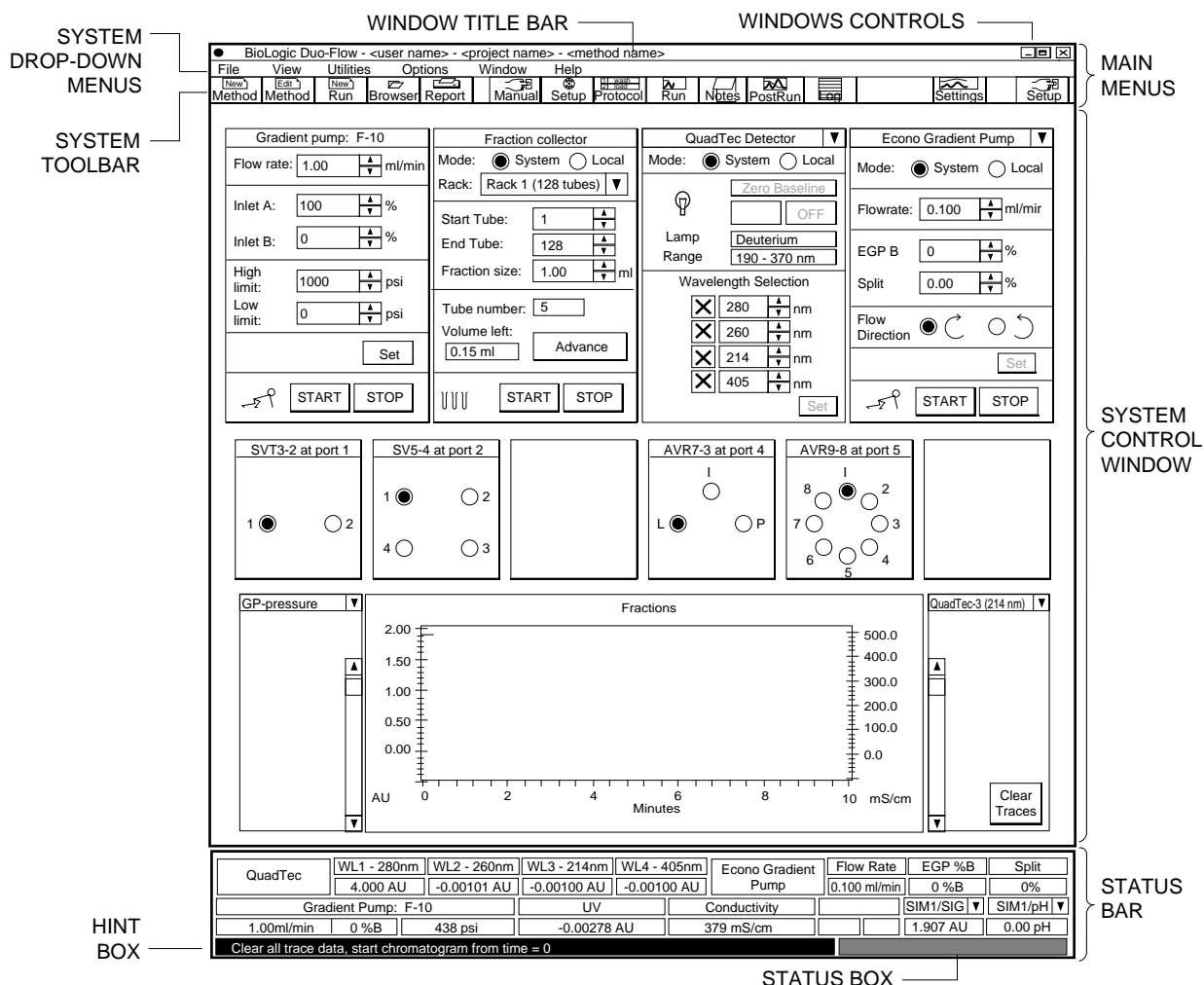


***Figure 5-1.  Layout of the Screen Display, showing the Manual Control Screen***

The BioLogic Duo-Flow system is controlled and monitored using the following:
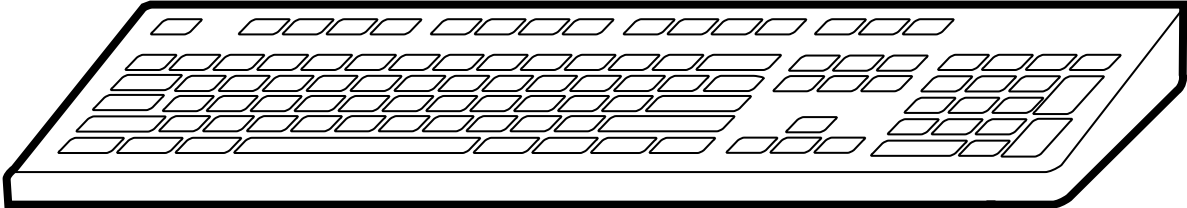
- **System Toolbar:** This toolbar is the primary navigation tool for the system software.

- **System drop-down menus:** These menus are always displayed. Access to advanced functions are found only in the drop-down menus. **Note**: Some functions found on the System Toolbar are duplicated in the drop-down menus.

- **System Status bar**: This area shows information about components in the system, as well as providing information about software icons.

   **Note**:  Items shown in gray are not currently active.

## 5.2    STANDARD MOUSE AND KEYBOARD FUNCTIONS

The BioLogic Duo-Flow system is supplied with a standard PC-compatible mouse and keyboard. The left mouse button is used with system software, except as noted.

**Table 5-1.**
**Special Function Keys**



| Special Function Keys | Description |
|---|---|
| F2 | **Hold until Keypress:**  This is used during a run to advance the Method (i.e., satisfy the Hold) when the Method includes a "Hold until Keypress" step. |
| F1 | **Help:**  Displays the Help menu for the currently displayed screen. |
| Esc | **Esc:**  Functions as an alternative to the Cancel selection in a Dialog box. |
| Alt | **Alt:**  Some system commands can be executed either by selecting them from a drop-down menu or by holding down the **Alt** key and then pressing the appropriate character key. |

## 5.3   SYSTEM MENUS

The system menus consist of both System drop-down menus and the System Toolbar. In some cases, identical functions are found in both areas. Advanced features are located only in the System drop-down menu.

### 5.3.1   System Toolbar Buttons

Most of the System Toolbar buttons are continuously displayed across the screen.  The function of each toolbar button is provided in Table 5-2.

*Table 5-2.*
*Toolbar Buttons*
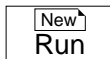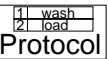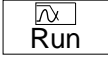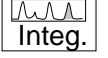
| Button | Description |
|---|---|
| **File Buttons** | **Note**: Similar functions can be found by selecting the File drop-down menu. |
| New Method | Initiates the process of writing a new method. The new method will be saved into the currently open User and Project directories, which are listed in the window title bar. **Note:** To place a new method in a different user and project directory, choose **Select User and Project** under the File menu before clicking on the **New Method** icon in the Toolbar.  Choose the desired user and project from the appropriate drop-down list. |
| Edit Method | This button is available when an open method has runs associated with it.  It allows you to create a new method based on editing the open method. A copy of the open method is made, and the system software prompts you for a name change. |
| New Run | Initiates a new run for the currently open method.  You will be prompted for a new run name prior to launching the run, or the runs will be numbered consecutively. |
| Browser | Opens the database of existing users, projects, methods, and runs for creating new users and projects. See chapter 6 for a complete description of the browser. |
| Report | Allows you to define when to print and what is to be printed in the report for a run. |

*Table 5-2. (continued)*
*Toolbar Buttons*

| Button | Description |
|--------|-------------|
| **View Buttons**<br><br>Manual | **Note:** Similar functions can be found by selecting the View drop-down menu.<br><br>Displays the Manual screen, which allows individual control of instruments and devices in the system. In Manual mode, data can be viewed but it cannot be saved. |
| Setup | Displays the Method Setup screen for specifying the components required for your method. From the Setup screen you can add a descriptive name to the valve positions and buffers. This screen is automatically displayed after the New Method button is selected. |
| Protocol | Displays the Protocol screen for creating and editing a method. |
| Run | Displays the Run screen, from which you can initiate a run of the currently loaded method. You will be prompted for a new run name prior to launching the run. |
| Notes | The notes section is a prompted section which allows you to enter additional information about your method and/or run. This section is always editable, even after a run is executed. |
| PostRun | PostRun allows you to customize your chromatogram prior to printing. |
| Log | As the run progresses, all events are recorded in the Log. This information cannot be edited. This button is not displayed when the EzLogic integration software is installed. To view the Log, select Run Log from the View drop-down menu. |
| Integ. | Appears only when the EZLogic integration software option is installed; this button replaces the Log button. It launches the software. |
| Settings | Enables the selection of up to 8 instrument traces to display on screen and on the printed report. The trace options are : standard UV detector, Conductivity monitor, pH, system back pressure, theoretical %B concentration, four QuadTec wavelengths, and detector traces acquired via the System Interface Module (SIM). |
| Setup | Displays the Manual Setup window, which allows you to enter:<br><br>• Econo Gradient Pump (EGP) Split Time Period when the EGP is to be used.<br><br>• QuadTec UV/VIS Detector Time Constant when the QuadTec Detector is to be used.<br><br>• Pump purge rate.<br><br>• Signal Import Module (SIM) signal parameters. |

### 5.3.2    System Drop-down Menus

The system drop-down menus are discussed in the following tables.

**Table 5-3.**
**File Drop-down Menu**

---

```
● BioLogic Duo-Flow - <user name> - <project name> - <method name>
File    Edit   View    Utilities    Options    Window    Help
New Method
Copy and Edit Method
New Run
Open...
Close
Select User and Project
Change Name, Author...

Save setup ...
Load setup...

Print Report...
Data management...
Export Data...
Export Chromatogram Image...
Integration...

Exit

1 Method 1-
2 Method 2-
3 Method 3-
4 Method 4-
```

---
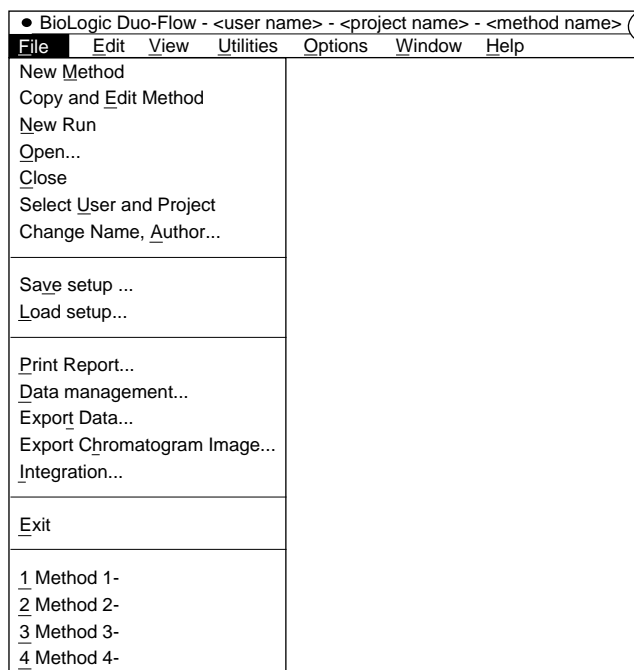
The File menu is always displayed.  It consists of the following:

- **New Method**: Initiates the process of writing a new method.  The new method will be saved into the open User and Project directories. To save to a different User or Project, first select **Select User and Project**.  Selecting **New Method** during a run will automatically open the "BioLogic offline" function, allowing you to create a method while a run is in progress.

- **Copy and Edit Method**:  Allows you to use an existing method as a template for a new method.  A copy of the method is made, and the system software prompts you for a name change.

- **New Run**:  Initiates a run for the currently open method.  You will be prompted for a new run name prior to launching the run.

- **Open**:  Displays the browser.  Refer to Chapter 6, Browser Screen.

- **Close**:  Closes the current method and/or run.

- **Select User and Project...**:  Allows you to select the default user and project of the system when new methods are created.

- **Change Name, Author...**:  Allows you to change the method/run name, author, and description.  This function is inactive once a run has been initiated.

- **Save setup...**:  Allows you to save and name different hardware setups and to set one as the default.

- **Load setup...**:   Recalls and loads the selected setup.

***Table 5-3. (continued)***
***File Drop-down Menu***

- **Print Report**: Allows you to print a report of the currently open method, including its setup and run data, run results, and the run log report.

- **Data Management**: Displays the Browser screen, from which you can copy and move run data. Refer to Chapter 6, Browser Screen.

- **Export Data**: This feature establishes data export parameters, and is available from the PostRun screen.

- **Export Chromatogram Image**: This feature exports a chromatogram image in a Windows Meta File (.WMF) format. This feature is available in the PostRun screen.

- **Integration:** Launches the Bio-Rad EZLogic integration software. Contact Bio-Rad Technical Support at 1-800-4-BIORAD in the USA or your local Bio-Rad representative for more information on EZLogic.

- **Exit**: Exits the BioLogic system software and returns to the Windows desktop.

- **1 through 4**: Display the last four user names and methods

*Table 5-4.*
*Edit Drop-down Menu*

**Edit menu: Setup Screen**

● BioLogic Duo-Flow - <user name> - <project name> - <method name>
File    **Edit**    View    Utilities    Options    Window    Help
　　　　Edit...
　　　　Delete
　
　　　　Select All
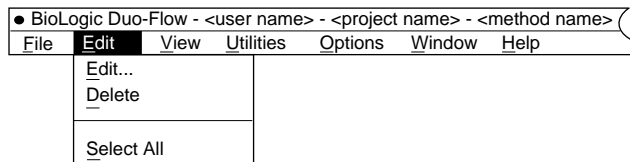
**Edit menu: Protocol Screen**

● BioLogic Duo-Flow - <user name> - <project name> - <method name>
File    **Edit**    View    Utilities    Options    Window    Help
　　　　Edit...
　　　　Cut
　　　　Copy
　　　　Paste
　　　　Delete
　
　　　　Select All:

**Edit menu: Run Screen**

● BioLogic Duo-Flow - <user name> - <project name> - <method name>
File    **Edit**    View    Utilities    Options    Window    Help
　　　　Full view in Run Chromatogram
　　　　Multiple runs...
　　　　Start run
　　　　Pause buffer pump
　　　　Hold gradient
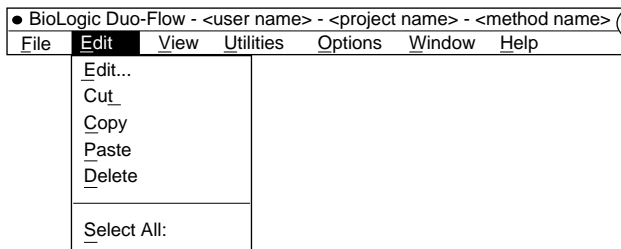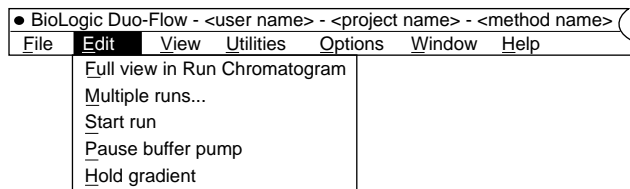
**Edit menu: Post Run Screen**

● BioLogic Duo-Flow - <user name> - <project name> - <method name>
File    **Edit**    View    Utilities    Options    Window    Help
　　　　Edit Activity Trace ...
　　　　Tag and Trace Options ...
　　　　Full view in zoom Chromatogram
　　　　Print zoom Chromatogram
　　　　Trace Legend

　　　　Copy zoom Chromatogram to clipboard

　　　　Delete selected tag
　　　　Delete all tags
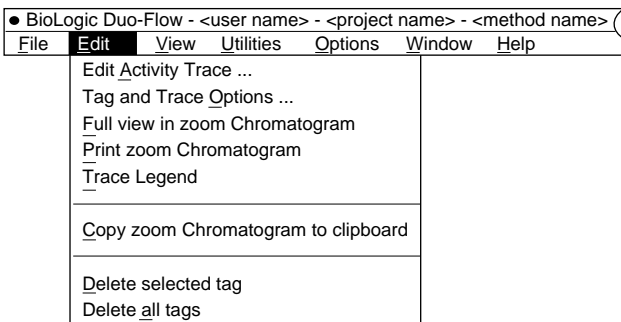
**Note:  The Edit menu is not available in the Manual screen.**

The contents of the Edit menu depends upon the currently displayed screen.  In most instances, the item in the drop-down menu also appears in the System Toolbar; exceptions are noted below.

**Setup Screen**

- **Edit...**:  Allows you to edit the selected device in Setup.

- **Delete**:  Deletes the currently highlighted device in Setup.  To delete all devices, first select Select All, as described below.

- **Select All**:  Highlights all devices in Setup.  (Not available from the System Toolbar.)

**Protocol Screen**

- **Edit...**:  Displays the Edit window for the step selected in the protocol.

- **Cut**:  Cuts (deletes) the currently highlighted step from the protocol.  A cut step may be pasted elsewhere.

- **Copy**:  Copies the currently highlighted step so that it can be pasted elsewhere in the protocol.

- **Paste**:  Pastes the cut or copied step into the protocol.

- **Delete**:  Deletes the currently highlighted step from the protocol.

- **Select All**:  Highlights all protocol steps.  Cut, Copy, and Delete then act on all steps.  To remove the highlighting from individual steps, hold down the **Ctrl** key and click the mouse over the desired step. (Not available from the System Toolbar.)

**Run Screen**

- **Full view in Run Chromatogram**: Zooms out to show the full view for the run.

- **Multiple runs...**: Specifies the number of times the method is to run.

*Table 5-4. (continued)*
*Edit Drop-down Menu*

**Run Screen** (continued)

- **Start run**:  Starts the run.

- **Pause buffer pump**:  Pauses progression of method's protocol.  Stops Gradient Pumps and the method time (volume) does not advance.  From Pause, you can Abort, Continue, or Edit-During-Run.

- **Hold gradient**:  Holds the current %B Gradient pump conditions and halts the advance of the method's protocol (including fraction collection). The method time (volume) does not advance.  From Hold, you can Abort, Pause, or Continue.


**Post Run Screen**

- **Edit Activity Trace...**: Allows you to input post-run sample activity data obtained off-line. (See page 7-31.)

- **Tag and Trace Options...**:  Displays the Post-Run Tags window that allows you to specify which traces are to be displayed, whether or not their tags are to be displayed, the tag style for the active trace, and the values for each tag in the active trace.  (Also available from the **Tag** button in the System Toolbar.)

- **Full View in zoom chromatogram**:  Allows you to view the complete chromatogram.  (Also available from the **Full View** button in the System Toolbar.)

- **Print zoom chromatogram**:  Allows you to print the complete chromatogram.  (Also available from the **Print** button in the System Toolbar.)

- **Copy zoom chromatogram to clipboard**:  Copies to clipboard the complete chromatogram.  From the clipboard, it can then be copied into other applications.  (Not available from the System Toolbar.)

- **Delete selected tag**:  To delete a tag, highlight the selected tag before using this function.  (Also available from the **Del. Tag** button in the System Toolbar.)

- **Delete all tags**:  Deletes all tags in the chromatogram.  (Not available from the System Toolbar.)

- **Trace Legend**:  Displays different line formats for each of the chromatogram traces. This is useful for distinguishing traces when printing to a black and white printer.

***Table 5-5.***
***View Drop-down Menu***

```
● BioLogic Duo-Flow - <user name> - <project name> - <method name>
  File    Edit    View    Utilities    Options    Window    Help
                  Manual
                  Setup
                  Protocol
                  Run
                  Post-Run
                  Run Notes
                  Run Log
```
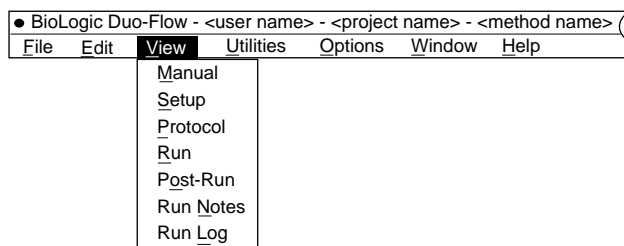
The View menu selections below remain the same in each toolbar menu selection.

- **Manual**:  Displays the Manual screen for individual control of installed instruments and devices in the system.

- **Setup**:  Displays the Setup screen, which allows you specify the components required for your method.

- **Protocol**:  Displays the Protocol screen for creating and editing chromatography steps in the method.

- **Run**:  Displays the Run screen, from which you can initiate the running of the currently open method.

- **Post-Run**:  When a run is completed, select **Post-Run** to annotate the chromatogram (for example, to apply tags to UV, Conductivity, and/or %B values).

- **Run Notes**:  To keep notes regarding the run, select **Run Notes** and enter them into the Notes screen. There are fields for description of the sample, the column, the operator, the buffer(s), flow rates, gradients, chart speed, fraction size, and general text entry of variable length.  These notes are printed with the report.

- **Run Log**:  Displays a log of all events occurring during the run.  This information cannot be edited. The Run Log may be disabled by selecting **Edit User Preferences** from the Option drop-down menu. It is recommended, however, that the Run Log be active for assistance in troubleshooting.

  **Note** :  If the Bio-Rad EzLogic Integration software option is installed, the **Log** toolbar button is replaced by the Integ toolbar button. To display the Log window, select **Run Log** from the View drop-down menu.

**Table 5-6.**
**Utilities Drop-down Menu**

```
● BioLogic Duo-Flow - <user name> - <project name> - <method name>
  File    Edit    View    Utilities    Options    Window    Help
                          Validate Method
                          Check Hardware connections

                          System Information
                          Diagnostics Information

                          Gradient Pump Calibration
                          Conductivity Flow Cell Constant Calibration
                          pH Probe Calibration
                          EGP Calibration
```
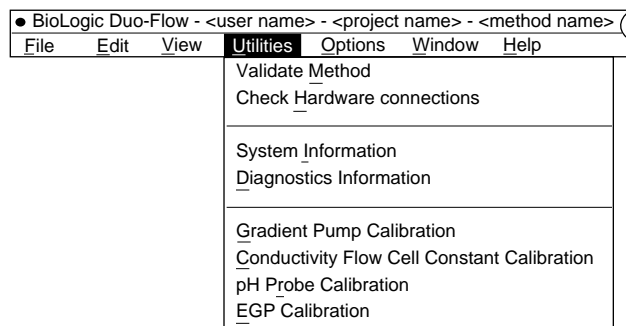
The Utilities menu selections relate to system options, and remain the same regardless of which screen is displayed. The Utilities menu consists of the following:

•   **Validate Method**: This verifies that the devices required by the Protocol have been defined in the Setup screen. Validation is automatically run at the start of a new run.

•   **Check Hardware connections**: This checks that all devices appearing in the Setup screen and required for the run are connected to the system. Check Hardware does not detect a V7-3 Manual Inject valve, chart recorder, Model 2110 or generic fraction collector, or Auxiliary pump, and it does not verify plumbing connections.

•   **System Information**: Displays the current system configuration, including each instrument in the system and its firmware version number, the Windows® version number, available hard disk space, and the number of methods and runs in the database.

•   **Diagnostics Information**: Displays information for service diagnostic purposes only.

•   **Calibrate Gradient Pump**: This selection is available only from the Manual screen, and it is typically used only after servicing of the pump. It allows the user to calibrate the gradient pump flow rate and zero the system pressure gauge.
    **Note**: You must exit the Calibration screen in order for the system to accept the calibration values.

•   **Conductivity Flow Cell Constant Calibration**: This utility allows the user to calibrate the conductivity flow cell. The conductivity flow cell constant is printed on a tag attached to the flow cell cable. When a new conductivity flow cell is installed, use this value to enter the flow cell constant.
    **Note**: You must exit the Calibration screen in order for the system to accept the calibration values.

•   **pH Probe Calibration**: This utility allows the user to calibrate the pH probe. It is typically used at the start of each day's use of the system.
    **Note**: You must exit the Calibration screen in order for the system to accept the calibration values.

•   **EGP Calibration**: This is informational only. It tells the user to calibrate the pump through the Pump software. Refer to the EGP Instruction Manual.

**Table 5-7.**
**Options Drop-down Menu**

**Options menu: Manual Screen**

● BioLogic Duo-Flow - <user name> - <project name> - <method>
File    Edit    View    Utilities    Options    Window    Help

| Edit User Preferences |
| Chromatogram Settings |
| Status lines |
| Browser Settings |
| Manual Setup |

**Options menu: Setup, Protocol, Post-Run Screen**

● BioLogic Duo-Flow - <user name> - <project name> - <method>
File    Edit    View    Utilities    Options    Window    Help

| Edit User Preferences |
| Chromatogram Settings |
| Status lines |
| Browser Settings |

---

These selections are to change screen options.

- **Edit User Preferences**: This lets you specify the following:

  **Run Options**: These determine whether or not the protocol's steps are listed in the Run Log and whether or not the valves are automatically returned to position 1 at the end of a run.

  **Protocol Editor Mode**: This determines whether the protocol is to be based on time or volume.

- **Chromatogram settings**: Allows you to select which device traces will be visible on the chromatograms. Up to 8 instrument traces are selectable from among the following: standard UV detector, Conductivity monitor, pH, system back pressure, theoretical %B concentration, four QuadTec wavelengths, and detector traces acquired via the System Interface Module (SIM). Both the X-axis (time) and the Y-axis (AU) ranges can be set.

- **Status Lines**: Allows the status information at the bottom of the screen to be toggled on or off.

- **Browser settings**: This enables you to select how the browser is to display users/projects/methods/ runs. For example, select Projects to display a top level folder called "Projects" which contains all projects in the database.

In addition, the following selection is available from the Manual screen:

- **Manual Setup**: This enables you to set the pump's purge flow rate. Additionally, you can set an EGP split time period, select a QuadTec time constant, and set the SIM-HR parameters. For the F-10 pumps, you can specify a maximum purge rate of 10 ml/min; for the F-40 pumps, you can specify a maximum purge rate of 40 ml/min.

**Table 5-8.**
**Window Drop-down Menu**

---

```
● BioLogic Duo-Flow - <user name> - <project name> - <method name>
   File    Edit    View    Utilities    Options    Window    Help
                                                   1  BioLogic Online - <method name>
                                                   2  BioLogic Offline - <method name>
```

---
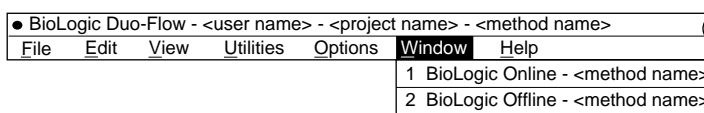
The BioLogic Duo-Flow software allows you to continue working while a run is in progress (Offline). A complete discussion of this function is provided in section 7.4.2, Working Offline.

•   **BioLogic Online**:  Allows you to observe and control the run in progress.

•   **BioLogic Offline**:  Allows you to write a protocol, analyze data, and export data while a separate run is in progress.

---

### 5.4   BIOLOGIC CONFIGURATION UTILITY SOFTWARE FOR USE WHEN CHANGING PUMP HEADS

Whenever the pump heads are changed, the BioLogic Configuration Utility software must be run before the system is used. The BioLogic Configuration software updates the flow rate and pressure ranges in the BioLogic Duo-Flow software. To run this software:

1.   Exit the BioLogic Duo-Flow software by selecting **Exit** from the File drop-down menu.

2.   Double-click on the BioLogic Configuration icon.

3.   In the BioLogic Configuration Utility window, indicate the pump that will be used with the system.

4.   Exit the BioLogic Configuration Utility and double-click on the BioLogic Duo-Flow icon to start the software.
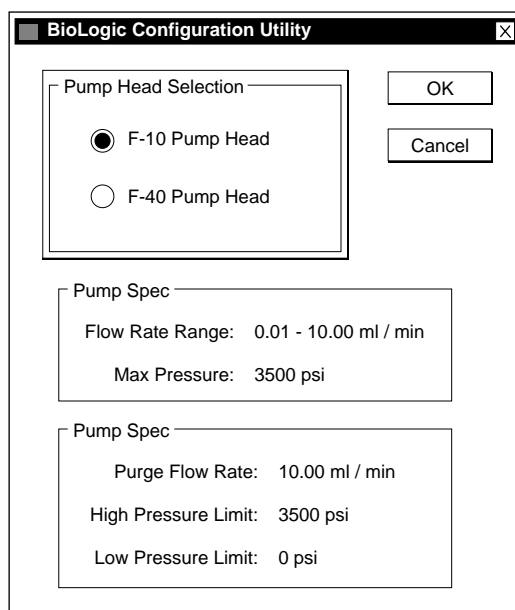


*Figure 5-2.  BioLogic Configuration Utility Software Screen*

# 6.0   INTRODUCTION TO THE BROWSER SCREEN

The browser is the organizational tool for chromatography data. It is a database that is displayed as a tree hierarchy which can be sorted by Users, Projects, Methods, and Runs. Figure 6-1 shows the layout of the Browser screen.
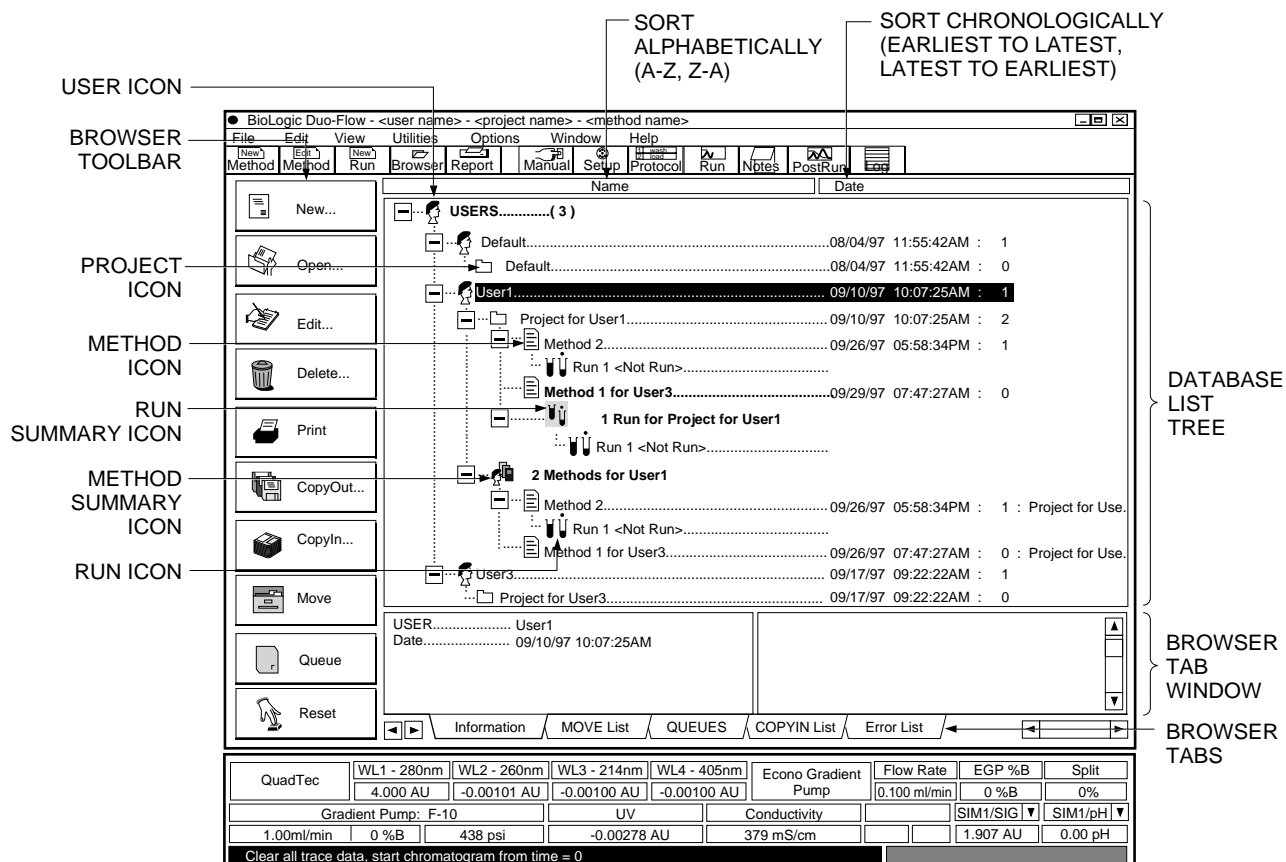


*Figure 6-1.  The Browser Screen*

## 6.1   OVERVIEW

The Browser screen displays the following information and controls:

- **Collapsing/expanding the tree hierarchy:**

    □   Click on this icon in the database tree to collapse the listing for a user, project, method, or run folder.

    ⊞   Click on this icon in the database tree to expand the listing for a user, project, method, or run folder.

    RESET   Updates and refreshes the Browser screen by collapsing all folders to the single user icon.

- **Icon colors in the database tree indicate the following:**

    Green:  Currently open in the active window.  The active window may be the on-line window or the off-line window. For more detailed information about on-line and off-line, refer to the discussion in Chapter 7, section 7.4.2, Working Off-Line.

    Red:  Currently open in the window that is not active.  This condition is displayed during off-line use: it applies to the method and run data for the window that is **not** active (on-line or off-line).  Items with a red check mark indicate that item is in the Move List or is marked for deletion.

    Blue (profile): User folder

    Yellow: Project folder

    Blue (text page):  Method folder

    Blue (test tubes):  Runs

- **Bolded text in the database tree applies to the following:**

    Top level folders:  Typically this applies to the USERS folder, which lists all users of the system.  By selecting **Browser Settings** from the Options menu, you can use the "Starting Browser Selections" to move Projects, Methods, and Runs to the top level of the tree hierarchy where they will be bolded.

    Summary information:  The total number of Methods are listed for each user, and the total number of Runs are provided for each user project folder.

    **Note:**  You cannot delete summary information.

**Browser screen controls and information**

- **Browser toolbar**:  These buttons, located down the left side of the screen, control the different uses of the Browser screen.  They are discussed in greater detail on the following page.

- **Name and Date bars**:  These are located across the top of the database tree and allow you to sort the tree alphabetically or chronologically.  These are toggle buttons.  For example, if you click on the Name button to sort from a to z, then clicking again sorts from z to a. If you click on the Date button to sort from the latest listing, then clicking again sorts from the earliest listing.

- **Browser Tabs and Browser Tab window**:  The Browser Tabs are at the bottom of the Browser screen, and they control what is displayed in the Browser Tab window.  The following tabs are available:

    Information tab:  Clicking on this tab displays two panels in the Browser Tab window: The left panel shows details about the specific item selected.

    MOVE List tab:  Displays the list of all items selected to be moved from one place to another in the database list.  Items selected to be moved are indicated by a red checkmark in the database tree.  Projects can be moved from one user to another; Methods can be moved from one Project to another. When a Project is moved, it takes all Methods and Runs associated with it; when a Method is moved, it takes all Runs associated with it.  The procedure for moving an item is discussed later in this chapter.

    Queue tab:  Displays the methods in the Queue selected or highlighted in the Browser. The sequence of methods in a queue can be changed by dragging and dropping.

    COPYIN List tab:  Displays the list of all items selected to be copied in (restored) to the database from an archived data file or disk.  The procedure for copying in items is discussed on page 6-4.

    Error List tab:  Displays discrepancies that exist between methods in queue. These discrepancies must be corrected to run the queue.

**Browser toolbar buttons**

**New...**      The **New...** button creates a new User, Project, Method, Run, or Queue depending on which icon within the Browser is highlighted.

**Open...**      The **Open...** button allows you to open a selected method or run from the database tree. Opening a Run allows you to view, analyze, and print the run data.

**Note**: Opening a Method or Run while a run is in progress will result in the Run-in-progress remaining in the On-Line window and the selected method or run being opened in the Off-Line window. For more on On-Line and Off-Line, refer to the discussion in Chapter 7, section 7.4.2, Working Off-Line.

**Edit...**      There are multiple functions of the **Edit...** button, depending upon what is highlighted when the button is pressed.

Selecting a user or project name allows you to change that user name or project name and/or description.

Selecting a Method or Run copies the methods to a new Setup and Protocol. The new method will be numbered sequentially and placed in the Browser. When a method is selected, you create a new method when you enter the method name, author, and description. When a run is selected, you can change the run name, operator, and description.

**Note**: Summary items are not editable.

**Delete**      The **Delete** button eliminates Users, Projects, Methods and Runs from the Browser. This is useful after using the Copy Out function to archive files to floppy disk, as discussed below. To delete from the Browser, you must first delete associated data. For example, to delete a Method, you must first delete all of the Runs associated with the Method.

**Note**: You cannot delete Methods or Runs from method and run summaries.

**Print**      To generate and print the report for a run. "**Print**" on the browser tool bar allows multiple runs to be printed with one command; highlight the desired group and press "**Print**".

**CopyOut...**      This function lets you copy methods and runs to floppy disk or to another location on your computer's hard disk. The copy out function is used for backup and archiving purposes and to transfer data to another BioLogic Duo-Flow system.

Copied out files are stored in a file with the default name biologic.zib, which is a compressed file format to conserve disk space. The copy out function copies the User, Project, and Method names associated with the Run.

**Note:** Initiating a CopyOut to floppy disk completely erases the contents of the floppy disk. Do not use a floppy disk containing data that you want to keep.

To copy out, highlight the Runs you wish to copy out. Press the **CopyOut...** button on the Browser Toolbar. A dialog box appears which allows you to select the destination of the .ZIB file (either the hard drive or a floppy disk). Choose the destination and press **OK**.
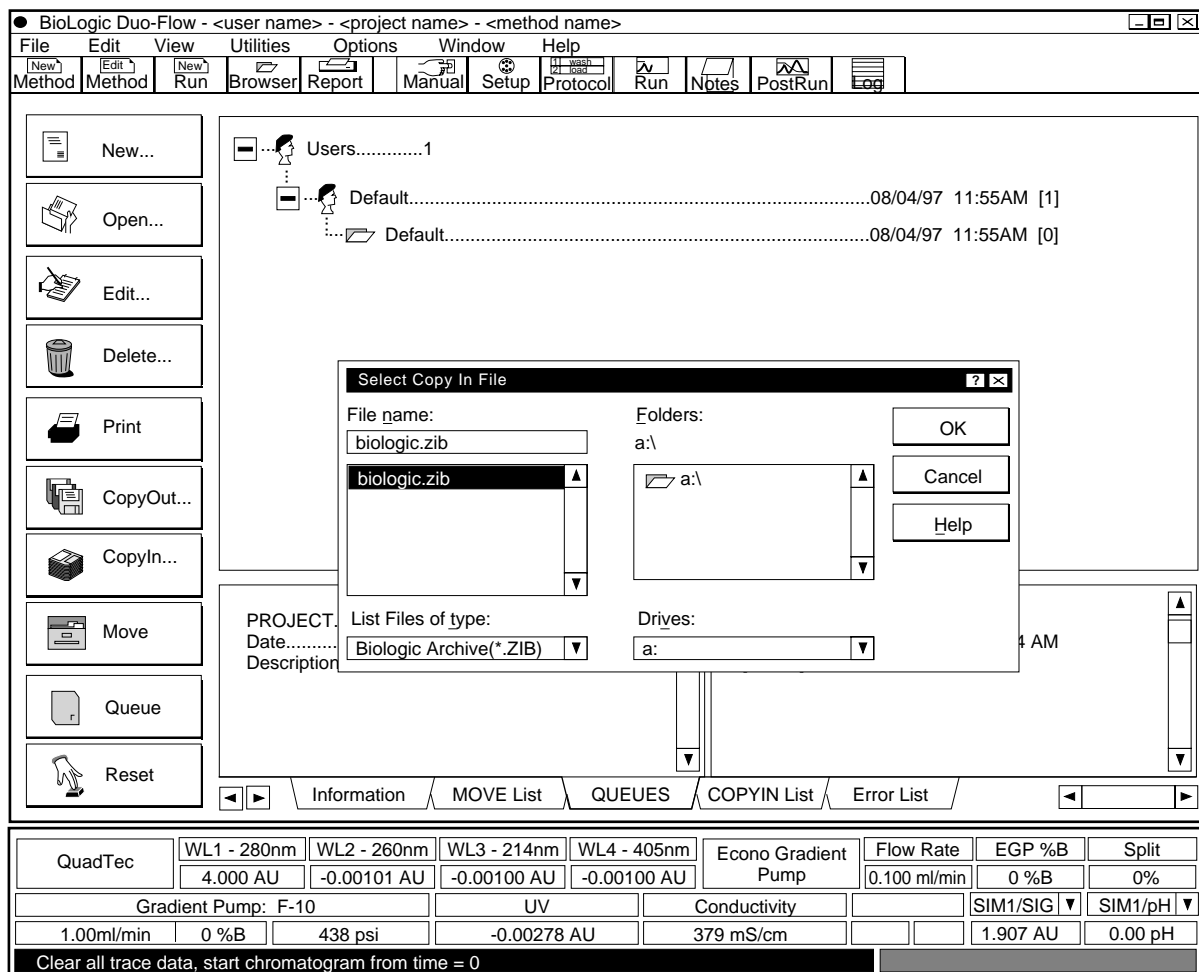
**Figure 6-2.  The CopyIn Window**

**CopyIn**...    This function allows you to copy in methods and runs from a backed-up or archived .ZIB file. To copy in,

    a.  Press the **CopyIn...** button and select the .ZIB file you wish to CopyIn.  (See Figure 6-2.)

    b.  From the COPYIN List in the Browser Tab window, select the desired methods and runs.

       All methods and runs in the file are displayed in the COPYIN List.  (See Figure 6-3.)

       From the database tree window, click the left mouse button to select the Project to which you want the methods and runs to be copied.

    c.  Again, click on the  **CopyIn...** button (or click on the right mouse button) and select **Copy to <project name>**.

       **Note**:  Clicking the right mouse button displays all the available Browser toolbar buttons for the *highlighted* selection.

    d.  Once you finish the copy in procedure, use the **Clear List** button on the Browser toolbar to delete the remaining methods and runs in the COPYIN List.
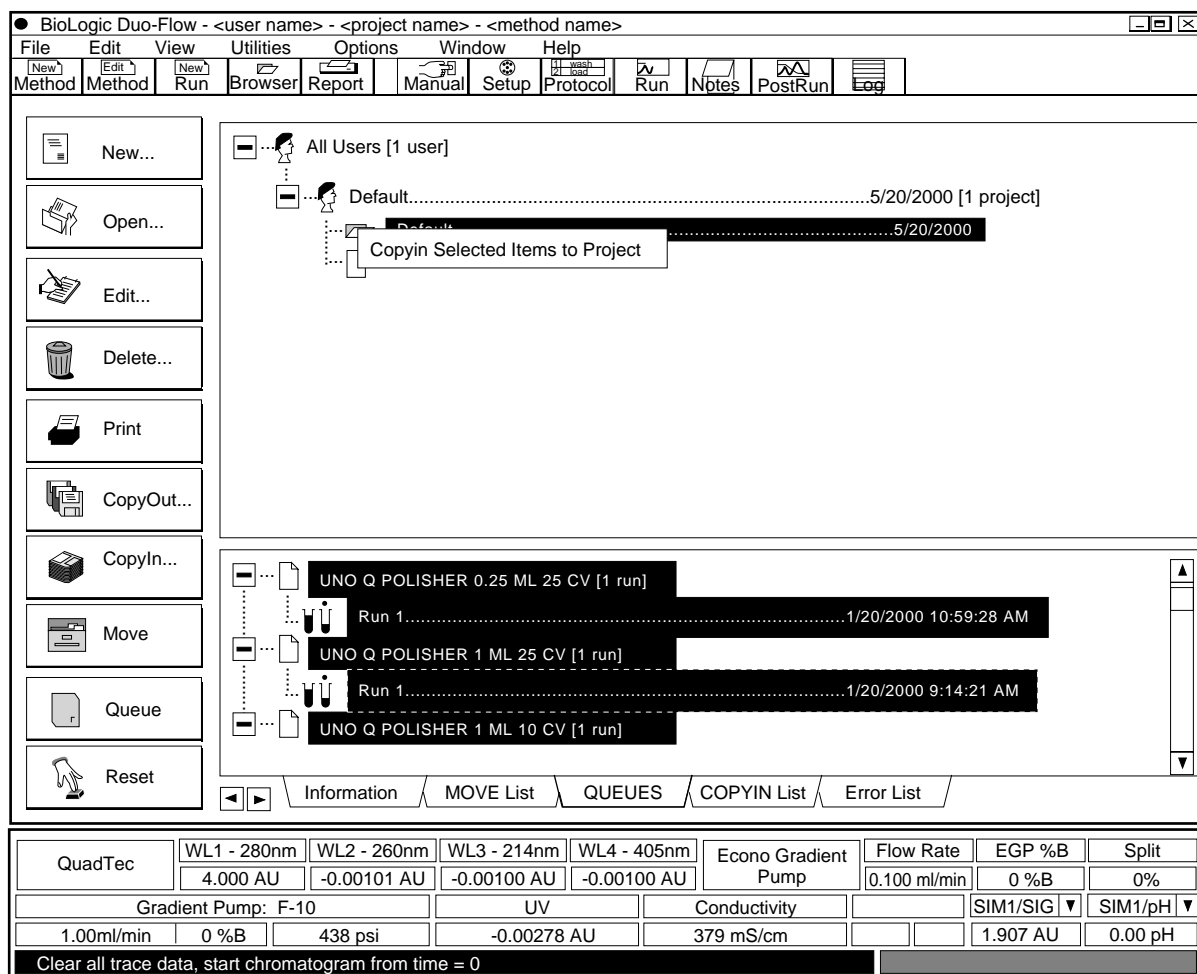
**Figure 6-3.  CopyIn with Information in Browser Tab Window**

**Move**          Move allows you to transfer a selected item from one location to another.  For example, a
                 Project can be moved from one user to another, and a Method (and its Runs) can be moved
                 from one Project to another. Individual Runs cannot be moved.  To move an item,

                 a.   In the database tree, highlight the item you want to move. You may highlight a Project or
                      a Method.

                 b.   Click once on the **Move** button on the Browser Toolbar. The items to be moved will
                      appear in the MOVE List in the Browser Tab window.

                 c.   From the MOVE List, select what you want to move.

                 d.   From the menu tree, select the new destination for the items you are moving.

                 e.   Again click on the **Move** button. The items will automatically move to the destination.
                      **Hint:** Alternatively, highlight the destination icon, right-mouse click and select the "Move
                      Selected" message with a left-mouse click.

                 f.   Once you finish the move procedure, use the **Clear List** button on the Browser toolbar
                      to delete the remaining methods and runs in the MOVE List.

**Queue**         Places methods in a queue.  Highlight the method(s) you wish to place in a queue and
                 select **Queue**. The method will appear in the Tab screen.  See section 6.2 for detailed
                 description of queuing.

**Reset**         Updates and refreshes the Browser screen by collapsing all folders to the single user icon.

There are several options available for how information is displayed in the Browser screen. Select **Browser Settings** from the Options menu, discussed in Table 5-7 Options Drop-down Menu.  The Set Browser Options window provides the following options:

- **Enable Projects**:  A summary of all Projects, regardless of User, will be listed in the Browser.

- **Enable Methods**:  A summary of all Methods, regardless of User or Projects, will be listed in the Browser.

- **Enable Runs**:  A summary of all Runs will be listed in the Browser.

- **Enable User Methods**:  A summary of all Methods for the specified user will be listed in the Browser.

- **Enable Project Runs**:  A summary of all Project Runs will be listed in the Browser.
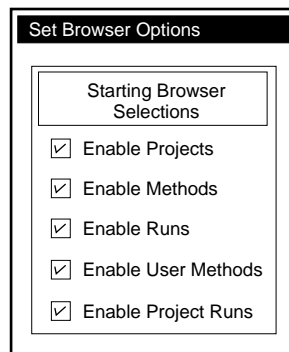


*Figure 6-4.  Set Browser Options Window*

**Procedures for creating a run in the Browser**

**1. Define a New User.**

Define a new user to allow you to group your methods and runs within the Browser.

a. Click once on the **Users** icon in the Browser.

b. Click once on the **New...** button and select New User. You will be prompted for a new User Name. Enter the name you would like to use.

c. Press **OK** to accept the new name.

**Hint:** Alternatively, highlight a User icon, right-mouse click, and select **New User** with a left-mouse click.

**Note:** If you define a new user, you must also assign a project to the user before writing a new method.

**2. Define a New Project.**

The Browser is further segmented into Projects within a User's domain. To define a new project:

a. With the new User Name highlighted, click once on the **New...** button.

b. Select **New Project**. Enter the project name and description.

c. Press **OK** to accept the new name.

**Hint:** Alternatively, highlight the User icon, right-mouse click, and select **New Project** with a left-mouse click.

**3. Write a New Method.**

To begin writing a new method from the Browser:

a. Select the Project folder or a Method in the Project folder where you want the new Method to be located.

b. Click once on the **New...** button.

c. Select **New Method**. Enter the method name and press **OK**. This transfers you to the hardware Setup screen.

**Hints: 1.** Alternatively, highlight the Project icon, right-mouse click, and select **New Method** with a left-mouse click.

**2.** You can also use the **New Method** button in the System Toolbar. By default the system assumes that the current user and project shown in the window title bar is the path for the new method.

**4. Name a New Run in a previously defined Method.**

To start a new run from the Browser:

a. Select the Method folder or a Run in the Method folder where you want the new Run to be located.

b. Click once on the **New...** button.

c. Select **New Run**. Enter the run name and press **OK**. The Run screen appears. Pressing the **Start** button in the toolbar will begin your sample run.

**Hints: 1.** Alternatively, highlight the Method icon, right-mouse click, and select **New Run** with a left-mouse click.

**2.** You can also use the **New Run** button in the System Toolbar. By default the system assumes that the current user, project, and method shown in the window title bar is the path for the new run.

## 6.2    CREATING AND RUNNING A QUEUE

Queue allows you to run multiple methods in sequence. **Methods placed in a queue must have identical hardware Setups or the system will not permit them to run.**
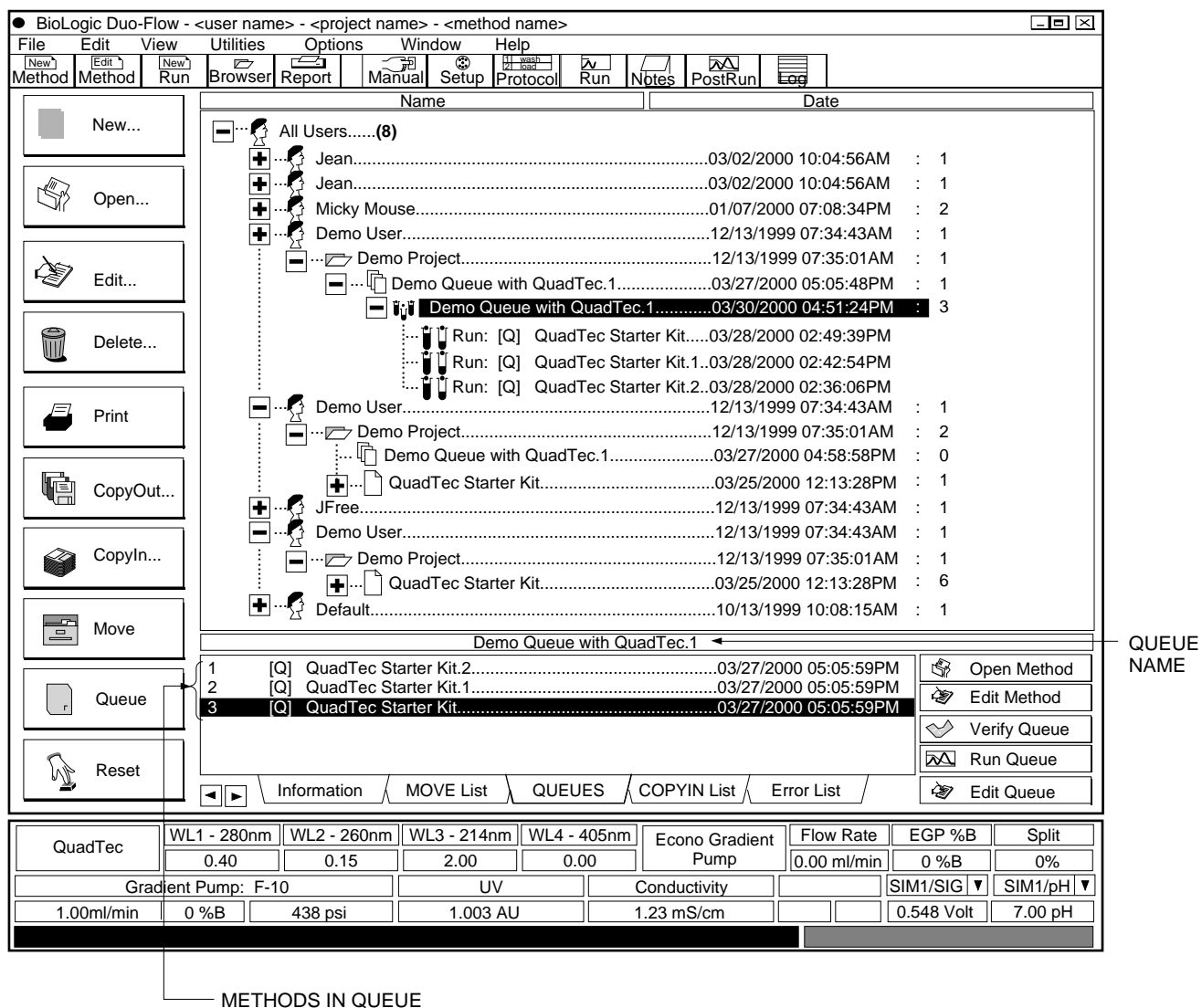


*Figure 6-5.  Queues Displayed in the Browser Window*

To create a queue,

**1.** In the Browser screen select a current user name or enter a new user name.

**2.** Select a current project name or enter a new project name.

**3.** Select the **NEW** icon from the left screen sidebar, and from the displayed menu select **New Queue.**
Enter a name and description for your queue and click **OK**.  Your queue name will be listed under your
user and project name, and the Queue tab will appear at the bottom of the screen. (See Figure 6-5.)

**4.** To place methods in a queue, highlight the method in the Browser and click the **QUEUE** icon on the sidebar. Repeat until all desired methods are in the Project Queue. This places each method into your Project Queue and the Queue Tab window at the bottom of the screen.

- Whenever you highlight the Queue icon under the Project Queue, the list of methods will appear in the Tab window.

- Multiple methods may be selected in the Browser by either shift/click or control/click.

- Methods will run in the sequence they are listed in the Tab window. The run order of the methods can be changed by dragging and dropping methods in the queue in the Tab window.

**5.** After all methods are listed in the Project Queue, click the **Verify Queue** button in the right sidebar of the Tab window to confirm that all methods meet the necessary criteria to be run.

- If the methods meet the necessary run criteria to run correctly, "Methods were successfully verified" will appear.

- If methods do not meet the necessary run criteria, a list of errors will appear. Errors will appear if methods are not compatible to run in sequence or the hardware Setup screens are not identical.

- To run your queue, click the **Run Queue** icon in the right sidebar of the Tab window.

**Description of the Queue Icons**
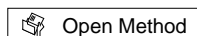
| | |
|---|---|
| Open Method | **Open Method**: Opens a highlighted method in the Protocol screen. |
| Delete Method | **Delete Method**: Deletes a highlighted method from the queue. Note: This is displayed only prior to the queue running. |
| Edit Method | **Edit Method**: Opens a highlighted method in the Protocol screen to be edited Note: This is displayed only when the queue has completed its run. Queued methods cannot be edited after the run has started. |
| Verify Queue | **Verify Queue**: Checks methods in a queue to assure there is not a conflict in the methods to prevent running.<br><br>If there is a conflict a list of errors will appear.<br><br>If methods are compatible the message "Methods were successfully verified," appears. |
| Run Queue | **Run Queue**: Starts a run queue. |
| Edit Queue | **Edit Queue**: Permits editing a queue. "**Edit Queue**" is a "**Copy and Edit**" function; i.e. it makes a duplicate. |
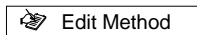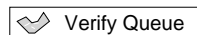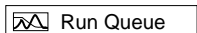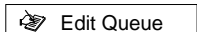
## 7.0   MODES OF OPERATION

There are two primary modes of system operation: the user can operate the system manually (from the Manual control screen) or through the use of a user-defined Method.  Each of these modes of operation is discussed below, as well as the screens involved in setting up, defining, and running a method.

- Operating the system using the Manual control screen:  This allows you to individually control the devices and instruments connected to the Workstation. Examples of its use include purging the pumps and equilibrating the columns.  For further discussion, refer to section 7.1.

- Operating the system using a User-Defined Method:  This involves the following:

    Creating a new method or selecting an existing method.

    Setup screen:  Allows you to define the instruments and devices to be used for any method.

    Protocol screen:  To define each of the steps to be run during a Method.  Most steps are programmed using a dialog box to enter the parameter values.  Note that the number of the step being programmed is displayed in the upper left corner of the dialog box.

    Run screen:  To run a method or view a run in progress.

    Start:  Starts a selected run.

    **Note**: You may move through the four screens by activating buttons on the system toolbar as long as a run is not in progress.  Those functions available during a run are discussed in section 7.4, Run Screen.

The relationship between the the two modes of operation, as well as the screens involved in defining and running a method, is illustrated in Figure 7-1 below.
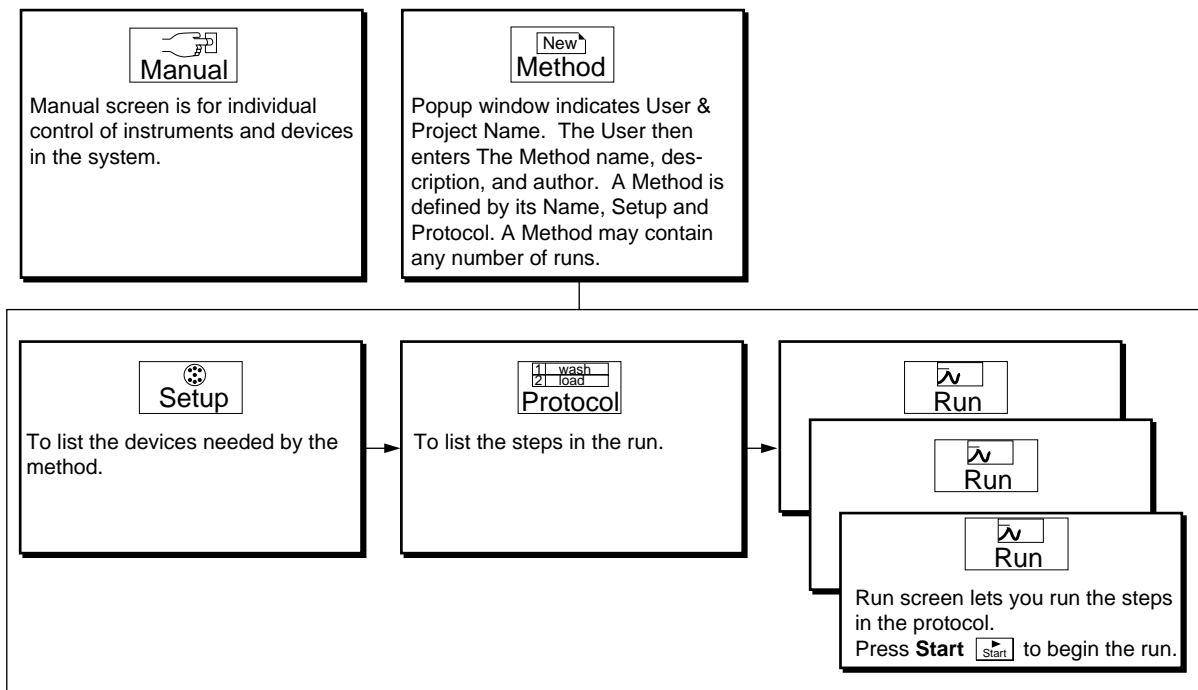


**Figure 7-1.  Relationships between Modes of Operation**

## 7.1    MANUAL CONTROL SCREEN

The Manual Control screen allows you to control different devices and instruments on an individual basis.

**Note**:  If an instrument is connected via the instrument bus but does not appear in the Manual screen, check that its power switch is On and that the instrument bus cable connection is secure.



**Figure 7-2.  Manual Control Screen, showing a Model 2128 Fraction Collector, QuadTec Detector, Econo Gradient Pump, and Four Valves**

From the Manual control screen you can control the following:

- Gradient Pump:  This control panel allows you to control the composition of the mobile phase being pumped, set the flow rate and pressure limits, and start/stop the pump. (User-defined pressure limit settings is discussed in the section below.)  If the pressure limit is exceeded, the pumps stop.

    **Note**: Changing pressure limits requires pressing either the **Start** or **Set** buttons.  When the pump is running, changing any parameters requires pressing the **Set** button.

The fraction collector, detectors, and Econo Gradient Pump have two modes of operation: **System**, in which the full functionality of the unit is available from the BioLogic Duo-Flow system software, and **Local**, in which control of the device is from its front panel faceplate.

- Fraction Collector:  When a Bio-Rad Model 2128 Fraction Collector is connected, the control panel displays the "Tube number," which is the tube currently under the drop-head (assuming the run started at tube 1), and the "Volume left," which is the volume remaining to be delivered.  There are also toggle buttons for **Start** and **Stop** and a button for **Advance.**  Note that the Tube number is a display field only.

   **Note**: Fraction collection is always by **Volume**. Collection by Drop or Time is not supported by the BioLogic Duo-Flow software.

- Econo Gradient Pump:  For this optional peristaltic pump the control panel allows you to select a flow rate, determine percentage of flow divided by the EGP splitter valve, select a flow direction and start and stop the pump.  Specific details for operation will be in the EGP user instruction manual.

- QuadTec Detector:  For this optional UV/Vis detector the control panel allows you to select 4 wavelengths and to zero them.  The QuadTec down arrow in the upper right corner of the control panel toggles between the QuadTec and the standard UV detector.  Specific details for operation of the QuadTec detector can be found in its instruction manual.

- Standard UV Detector:  The control panel provides access to the standard UV detector, Chart Recorder and indicates Signal Import Module Status. It allows you to turn the UV Lamp on and off and to zero the baseline.

- Chart Recorder:  If you are using the standard UV detector and Conductivity monitor, you can set the chart recorder full scale ranges, start/stop the recorder, and send an event mark.  The full scale ranges set here apply to the chart recorder and are not reflected in chromatograms displayed on the computer screen.

- Signal Import Module (SIM 1 and 2):  This panel indicates the presence of up to two Signal Import Modules (SIM-HR).  The signal from an analog instrument connected to the SIM is digitized so that the instrument can be monitored from the BioLogic Duo-Flow system.  Instruments that may be connected include any detection source, such as a variable wavelength UV detector, a pH meter, a refractive index detector, or a fluorescence detector.  For more information, refer to Chapter 2.4.1 and to the separate instruction sheet for the SIM-HR module.

- Valves:  Allows you to control the valve position of any of the BioLogic Duo-Flow solenoid or automated valves connected to the system.

In addition, the Manual control screen chromatogram displays the system output for up to 8 instrument traces from any of the following: standard UV detector, conductivity, pH, pressure, theoretical %B concentration, four QuadTec wavelengths, and detector traces acquired via the System Interface Module (SIM). Use the scroll bars, the **Settings** button in the toolbar, or the **Chromatogram Settings** selection under the **Options** drop-down menu to change the x- and y-axes settings. The drop-down menus on either side of the chromatogram permit selection of a trace for adjustment to its sensitivity reading.

### 7.1.1 Pressure Limit Settings

The Duo-Flow system is designed for pressures to 3500 psi, with the standard F-10 pump heads installed. The optional F-40 pump heads are rated to 1000 psi at 40 ml/min.  If you will be using a column designed for lower pressures, you can set a lower pressure limit for the system.  From the Gradient Pump panel, set the high limit to the appropriate value for the column. Set a low limit that is 6 psi or greater (the system does not allow minimum pressure below 6 psi) to protect the system from pumping air if it runs out of buffers or mobile phase. If the pressure limits are exceeded, the pumps stop.

**Note**: Changing pressure limits requires pressing either the **Start** or **Set** buttons.

## 7.2    SETUP SCREEN

The Setup screen is used to select instruments/devices to be used in the Protocol for a particular user defined method.  Each Method has its own instruments/devices Setup.  For example, if you plan to use a Model 2128 Fraction Collector for different methods then you must select this particular collector in each Setup screen for each method.

If you routinely use the same Setup(s), it is recommended to select **Save setup...** from the File drop-down menu.  This allows you to save and name different hardware setups and to set one as the default.  The default or selected setup is then loaded each time you create a New Method, thus eliminating the need to build a new Setup each time.

In figure 7-3, Setup screen devices are chosen from the left hand "Available Devices" box. A dialog box will appear allowing you to label the device in the Method. For example, an AVR7-3 may be selected and then defined as a Sample Inject Valve.  Use the **Delete** button in the System Toolbar to remove a device from the Devices in Setup window.  In order to *run* your Method, these devices must be connected to the rear of the Workstation and they must be powered on.

**Note**:  Devices may not be properly connected to the system but may be included in the Setup.  The run will not start, however, until the devices are connected.  An error message will appear on the Run screen.
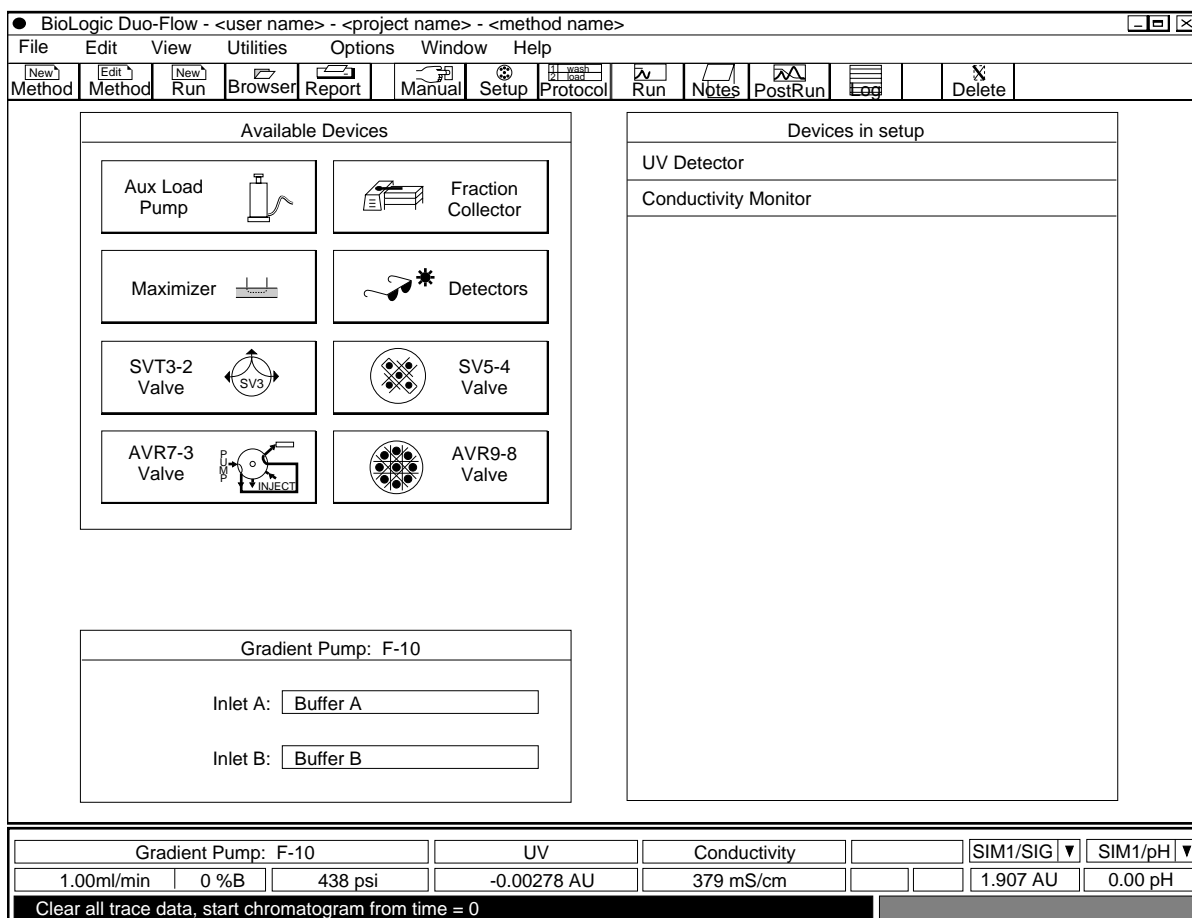


**Figure 7-3.  Setup Screen**

**Valve Functioning and Naming**

One of the most powerful features of the BioLogic Duo-Flow Software is the ability to define the specific purpose of the automated valves and to name each valve position. This greatly simplifies programming individual steps in the Protocol screen.

For example;

- The most common use of an AVR7-3 valve is as a Sample Inject Valve.  Once an AVR7-3 valve has been assigned to this function, the software controls all of its valve changes based on the programmed sample injection step.

- The most common use of a SVT3-2 valve will be as a Fraction Collector Diverter valve with a Model 2110 or generic collector. Once an SVT3-2 valve has been assigned to this function, the software controls eluant diversion based on the programmed fraction collection scheme.

In summary, you can identify the following as applying to all valves:

- **Valve Name/Function**.  What is the valve to be used for? Select from the drop-down list box or input a unique function that you will recognize when the particular valve is selected in the Protocol screen.

- **Connector**.  Which connector at the rear of the workstation is the valve connected to?

- **Position Names**.  These fields let you name each valve position.

The following devices are listed in the Setup screen.  Because the standard UV detector, Conductivity monitor, and AVR7-3 valve are standard with the BioLogic Duo-Flow system, they are always listed in the Devices in Setup panel.  To eliminate any of these, highlight it and click the **Delete** button.

- **Auxiliary Load Pump**:  Allows you to use an Auxiliary Load Pump, such as the Model EP-1 Econo Pump or the Econo Gradient Pump (EGP).  See Section 4, Advanced System Applications, for ways to incorporate an auxiliary load pump into a method.

- **Fraction Collector**:  Select this button if you are using a fraction collector with the system.  A dialog box appears, allowing you to select a Model 2110, Model 2128, or generic (non-Bio-Rad) fraction collector. If you use threshold detection to initiate fraction collection, select the appropriate detector.

   Select a delay volume or leave at default of zero.  The Delay function precisely synchronizes fraction collection with the UV signal on the chart recorder.  It is defined as the tubing volume between the UV detector and the fraction collector drop-head. If you are using the recommended 1/16" (1.6 mm) OD, 0.020" (0.51 mm) ID Tefzel tubing, 1 cm of tubing has a volume of 2 µl. The volume of the supplied backpressure regulator is 146 µl.

   **Note**:  To enable the use of collection windows or threshold collection for a Model 2110 or generic fraction collector, it is necessary to assign an SVT3-2 valve as the fraction collector diverter valve.

- **Detectors**:  The QuadTec multiwavelength detector is available as an option from Bio-Rad; it is discussed in greater detail in its instruction manual.  The standard UV detector and Conductivity monitor are standard with the Duo-Flow system. The SIM/pH is for use with Bio-Rad's optional pH monitor and SIM (Signal Import Module).  SIM Signal is for use with non-Bio-Rad detectors, such as refractive index and fluorescence detectors.

   The **SIM Signal** button requires that you set the **Units** and the minimum and maximum for the **Units Range** and **Device Output Range (volts)**.  Select the device number on the SIM-HR to match the number displayed.  Refer to the analog device's documentation for information on output voltages.

- **Valves (SVT3-2, SV5-4, AVR7-3, and AVR9-8)**:  The BioLogic Duo-Flow system's automatic valves can be configured in a number of different ways depending on their function in a method.  Table 7-1 discusses how each of the valves can be set up.

- **Gradient Pump**: Inlets A and B may be renamed by overwriting "Buffer A" and "Buffer B" with the description of the buffer.

**Table 7-1.**
**Valve Setup Information**

| Valve Type | Valve Name/ Function | Position Names | Notes |
|---|---|---|---|
| **SVT3-2** Low pressure solenoid valve | Fraction Collector Diverter | 1 Waste 2 Collect | Functions as a fraction collection diverter, determined by the actual fraction collection parameters chosen. |
| | Inlet A Inlet B | Named by user Named by user | When used before the inlet to Pump A or B, the valve enables buffer selection. The buffer name specified for each position will appear in the Protocol screen's "Isocratic Flow", "Linear Gradient", and "Change Valve" dialog box. Refer to Chapter 8, Sample Loading, for examples. |
| | Aux Pump Inlet | Named by user | Used for auxiliary pump load selection to select one of two solutions. Refer to Chapter 8, Sample Loading, for examples. |
| | User-assigned name | Named by user | When used for a purpose other than described above. The name specified for each position will appear in the Protocol screen's "Change Valve" dialog box. |
| **SV5-4** Low pressure solenoid valve | Inlet A Inlet B | Named by user Named by user | When used before the inlet to Pump A or B, the valve provides preparative sample loading or buffer selection. The buffer or sample name specified for each position will appear in the Protocol screen's "Isocratic Flow", "Linear Gradient", and "Change Valve" dialog box. Refer to Chapter 8, Sample Loading, for examples. |
| | Aux Pump Inlet | Named by user | Used for Auxiliary Pump load selection to select one of four solutions. Refer to Chapter 8, Sample Loading, for examples. |
| | User-assigned name | Named by user | When used for a purpose other than that described above. The name specified for each position will appear in the Protocol screen's "Change Valve" dialog box. |
| **AVR7-3** High pressure valve | Sample Inject | 1 Load Sample 2 Inject Sample 3 Purge | For automatically loading a sample. |

**Table 7-1. (continued)**
**Valve Setup Information**

| Valve Type | Valve Name/ Function | Position Names | Notes |
|---|---|---|---|
| AVR7-3 (cont'd) | User-assigned name | Named by user | When used for a purpose other than described above. The name specified for each position will appear in the Protocol screen's "Change Valve" dialog box. Refer to Section 4, Advanced System Applications, Chapters 8 through 10. |
| **AVR9-8** High pressure valve | Aux Pump Inlet | Named by user | Used for Auxiliary Pump load selection to select from up to eight samples, buffers, or rinse solution. |
| | Inlet A | Named by user | When used before the inlet to Pump A or B, the valve provides preparative sample loading or buffer selection. The buffer or sample name specified for each position will appear in the Protocol screen's "Isocratic Flow", "Linear Gradient", and "Change Valve" dialog box. Refer to Section 4, Advanced System Applications, Chapters 8 and 9. |
| | Inlet B | Named by user | |
| | Fraction Collector | 1 Waste 2 Collect 3-8 Named by user | Useful for collecting large volume samples. Collects up to 8 samples. The name specified for each position will appear in the Protocol screen's "Change Valve" dialog box. |
| | User-assigned name | Named by user | When used for purposes other than described above. The name specified for each position will appear in the Protocol screen's "Change Valve" dialog box. Refer to Chapters 8 and 9. |
| | Column Switching | Named by user | Allows you to install up to 8 columns and assign valve numbers and names to each column. Requires two AVR9-8 valves: one is an inlet valve and the other functions as an outlet valve. |

## 7.3    PROTOCOL SCREEN

The Protocol screen allows you to program step-by-step instructions for running a method.

**Note**: Methods may be programmed either by Volume (ml) or Time (min).  The default (time or volume) is set by selecting **Edit User Preferences** from the Options drop-down menu.

The Protocol screen contains the following types of icon buttons, which are shown in Figure 7-4 and described in detail in this section.

- Add Step buttons: These buttons are used to define the steps in the protocol.  They are located in a vertical box on the left side of the screen.  To insert a step, highlight the step *below* where the new step is to be added, and then press the appropriate button to begin defining that step.  For example, selecting Isocratic Flow displays a dialog box allowing you to define the buffer and volume (or time) and flow rate.

- Fraction Collector button:  This button is used to define the fraction collection scheme.  It is located below the Add Step buttons.

- Edit buttons:  These buttons, located in the System Toolbar, are used to edit, cut, copy, paste, and delete steps from the method.  To cut, copy, or delete a step, select that step and then press the appropriate button.  To paste, select the step below where the step is to be pasted and then paste.

- File Buttons:  Use the New Method, Edit Method, and Browser System Toolbar buttons to create a new method or to open an existing Method.

The system checks the Setup information before allowing you to add steps or run the Method.  This ensures that each device you need is present and available for the required function.
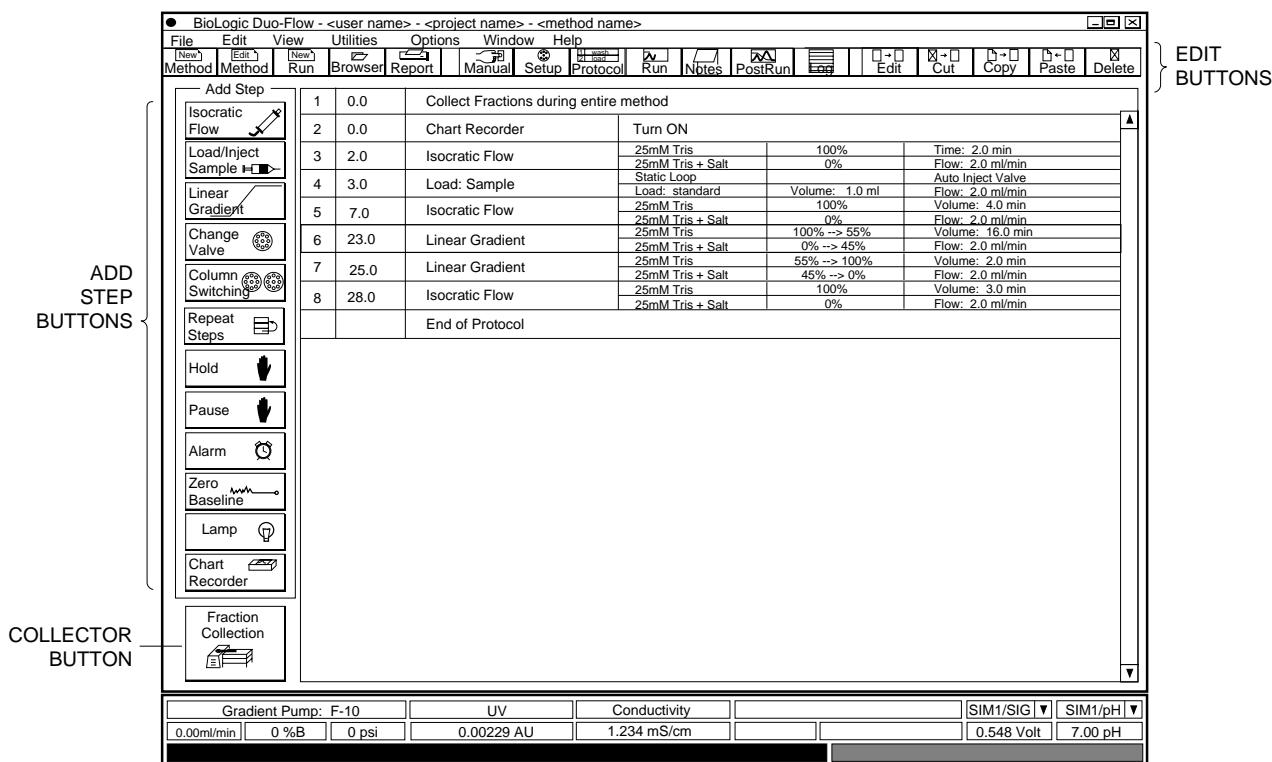


*Figure 7-4.  Protocol Screen*

The following tables describe each button in the "Add Step" column of the Protocol screen.

**Table 7-2.**
**Isocratic Flow**

---

```
┌─────────────────────────────────────────────────────────────────┐
│ Edit Isocratic Flow                                         ☒    │
│ ┌─────────┐                                                      │
│ │Isocratic│                                                      │
│ │ Flow    └──────────────────────────────────────────────────┐  │
│ │                                                             │  │
│ │    Buffers       % Composition    ┌─ Volume (ml.) ──────┐   │  │
│ │  ┌────────────┐               ┌─┐ │                  ┌─┐ │   │  │
│ │  │A-Buffer 1 ▼│ A │ 100     │ │▲│ │ │ 1.0     │      │▲│ │   │  │
│ │  └────────────┘             │ │▼│ │            │▼│ │   │  │
│ │                                   │ Flow (ml./min)      │   │  │
│ │  ┌────────────┐             ┌─┐   │              ┌─┐    │   │  │
│ │  │Buffer B   ▼│ B │ 0      │ │▲│  │ │ 1.00    │  │▲│    │   │  │
│ │  └────────────┘            │ │▼│  │             │▼│    │   │  │
│ │                                   └─────────────────────┘   │  │
│ │                                                             │  │
│ │ Step 2, Volume = 0.00 ml          ┌────────┐  ┌────────┐    │  │
│ │                                   │   OK   │  │ Cancel │    │  │
│ │                                   └────────┘  └────────┘    │  │
│ └─────────────────────────────────────────────────────────────┘  │
└─────────────────────────────────────────────────────────────────┘
```

---
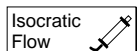
Isocratic
Flow ✐

To deliver a buffer under isocratic conditions. The composition may be chosen by altering either Inlet A or Inlet B percent values which increase or decrease in tandem. Use the up/down arrows or input values from the keyboard when this field is highlighted.

• **Buffers**: Displays all available buffers for separation. Buffer A and Buffer B can be labeled to identify the buffer in use by editing the Gradient Pump fields in the Setup screen. If pump inlet valves are assigned in the Setup screen and buffers are defined, all additional buffers will be displayed in the drop-down boxes.

• **Composition**: Allows you to choose the buffer composition (e.g., 75% A  and 25% B).

• **Volume/Time**: Allows you to choose the duration of the step.

• **Flow rate (ml/min)**: Allows you to choose the flow rate of the step.

• **OK**: Adds the step to the protocol.  This is the same as pressing the Enter key on the keyboard.

• **Cancel**: Cancels all input.  This is the same as pressing the Esc key on the keyboard.

• **Step, Time** or **Volume**: Identifies current step number, and calculates the elapsed time or volume from all previous steps.  This is not user editable.

**Table 7-3.**
**Load/Inject Sample**

```
┌─────────────────────────────────────────────────────────────────────────┐
│ Edit Load/Inject Sample                                              [×]  │
│ ┌─────────────────┐                                                       │
│  Load/Inject Sample  Fill Before Inject  Rinse After Inject               │
│ ┌─ Type ──────────┐  ┌──────────────────────────────┐  ┌───────────────┐ │
│ │                 │  │ Injection Buffers    % Comp.  │  │ Volume (ml.)  │ │
│ │  ● Static Loop   │  │ ┌──────────────┐ ┌─────┐▲    │  │ ┌─────┐ ▲     │ │
│ │                 │  │ │ A-Buffer 1  ▼│ A│ 100 │▼    │  │ │ 1.0 │ ▼     │ │
│ │  ○ Dynamic Loop │  │ └──────────────┘ └─────┘      │  │ └─────┘       │ │
│ │                 │  │                               │  │ Flow (ml./min)│ │
│ │  ○ Direct Inject│  │ ┌──────────────┐ ┌─────┐▲    │  │ ┌──────┐ ▲    │ │
│ │                 │  │ │ Buffer B    ▼│ B│  0  │▼    │  │ │ 1.00 │ ▼    │ │
│ └─────────────────┘  │ └──────────────┘ └─────┘      │  │ └──────┘      │ │
│ Alarm will sound and Protocol will HOLD for Start of Injection, then CONTINUE at end of this step. │
│                                                                           │
│ Step 3, Volume = 1.00 ml                        ┌────────┐   ┌──────────┐ │
│                                                 │   OK   │   │  Cancel  │ │
│                                                 └────────┘   └──────────┘ │
└─────────────────────────────────────────────────────────────────────────┘
```

Load/Inject
Sample ⊢■▷~
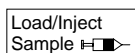
To program the sample injection. The Load Sample dialog box allows you to specify the device that will load sample onto the column, as well as the volume of sample to be loaded. (If programming in time mode, the system calculates the time required for this step.)

- **Load/Inject Sample**: Explains the action of the valve. The AVR7-3 auto inject valve will move to the Inject position at the start of the step and return to the Load position at the end of the step.

- **Type**: There are three options for sample loading. Refer to Section 4, Advanced System Applications, for examples.

  **Static Loop**: This is a standard fixed volume loop for sample loading. On the AVR7-3 valve, the loop is connected at ports 3 and 6, and sample can be loaded with a syringe through port 2. By adding an Auxiliary (AUX) pump and an auxiliary pump valve (such as the SVT3-2, SV5-4, or AVR9-8), the system can fill the sample into the static loop prior to injecting it onto the column, and it can rinse the sample loop after the sample has been injected. Fill Before Inject and Rinse After Inject are active when an Aux pump is used to load sample. Volumes for pre- and post-rinse can be selected in these two tab windows.

  **Dynamic Loop**: This uses Bio-Rad's DynaLoop or other sliding-piston sample loop. Both partial loop and full loop injections can be programmed. By adding an Auxiliary (AUX) pump and an auxiliary pump valve (such as the SVT3-2, SV5-4, or AVR9-8), the dynamic loop can be filled prior to injection onto the column. The Rinse function is not available. The dynamic loop can also be filled manually through Inject valve port #2.

  **Direct Inject**: Allows direct injection of sample through either the Workstation pump or auxiliary load pump. Pre-pump valves (such as the SVT3-2, SV5-4, or AVR9-8) automate direct injection. Use of an auxiliary pump (such as the Bio-Rad Model EP-1 Econo Pump or EGP peristaltic pump) allows direct injection of sample onto a low pressure column (≤15psi). Use of an Auxiliary (AUX) pump and an auxiliary pump valve allows multiple direct injections onto a low pressure column.

- **Injection Buffers and % Composition**: These fields allow you to choose the buffer and composition which will push the sample through the sample loop.

- **Volume**: Allows you to specify the volume of the sample to be injected. The smallest volume selectable is 0.1 ml; the largest volume selectable is 9999 ml (for larger volumes, use another Load/Inject Sample step). Actual load volume is defined by the sample loop size.

- **Flow Rate**: Allows you to specify the flow rate for the sample injection step.

- **OK**: Adds the step to the protocol. This is the same as pressing the **Enter** key on the keyboard.

- **Cancel**: Does *not* add the step to the protocol. This is the same as pressing the **Esc** key on the keyboard.

- **Step, Time** or **Volume**: Identifies current step number, and calculates the elapsed time or volume from all previous steps. This is not user editable.

*Table 7-4.*
*Linear Gradient*

---

```
┌─────────────────────────────────────────────────────────────────────┐
│ Edit Linear Gradient                                              [✕] │
├─────────────────────────────────────────────────────────────────────┤
│ ┌─Linear Gradient──────────────────────────────────────────────────  │
│ │                                                                     │
│ │       Buffers          % Composition                               │
│ │                       Initial      Final          Volume (ml.)     │
│ │   ┌──────────────┐┌─┐      ┌────┐      ┌────┐▲    ┌────┐▲           │
│ │   │ A-Buffer 1   ││▼│  A   │100 │▲     │100 │     │1.0 │           │
│ │   └──────────────┘└─┘      └────┘▼     └────┘▼    └────┘▼           │
│ │                                                   Flow (ml./min)   │
│ │   ┌──────────────┐┌─┐      ┌────┐▲     ┌────┐▲    ┌────┐▲           │
│ │   │ B-Buffer 1   ││▼│  B   │0   │      │0   │     │1.00│           │
│ │   └──────────────┘└─┘      └────┘▼     └────┘▼    └────┘▼           │
│ │                                                                     │
│ ├─────────────────────────────────────────────────────────────────  │
│  Step 8, Volume = 2.00 ml              ┌────────┐   ┌────────┐        │
│                                        │   OK   │   │ Cancel │        │
│                                        └────────┘   └────────┘        │
└─────────────────────────────────────────────────────────────────────┘
```

---

**Linear Gradient**

To deliver buffer gradients. The Linear Gradient dialog box sets the initial and final composition of the buffers and the period over which the change in composition is to occur. There is no limit to the number of gradient steps in a protocol.

- **Buffers**: Allows you to choose two buffers to make a binary gradient. Drop-down menus display all buffers that are set up using BioLogic Workstation pump Inlet valves.

- **% Composition**: Allows you to choose initial and final values that will form a linear binary gradient. Notice that changing one value in the "Initial" column affects the other "initial" value. The "Final" column behaves identically.

- **Volume**: Allows you to choose the duration of this step.

- **Flow**: Allows you to choose the flow rate of this step.

- **OK**: Adds the step to the protocol. This is the same as pressing the **Enter** key on the keyboard.

- **Cancel**: Does *not* add the step to the protocol. This is the same as pressing the **Esc** key on the keyboard.

- **Step, Time** or **Volume**: Identifies current step number, and calculates the elapsed time or volume from all previous steps. This is not user editable.

**Table 7-5.**
**Change Valve**

---

```
Edit Change Valve                                          ☒

 Change Valve

   Valve                                         Position
   ┌────────────────────────────────┬──┐        ┌──────────────┐
   │ SVT3-2 Valve - Inlet A Port 1  │▼ │        │ A-Buffer 1   │
   ├────────────────────────────────┴──┤        │ A-Buffer 2   │
   │ SVT3-2 Valve - Inlet A Port 1      │        │              │
   │ SVT3-2 Valve - Inlet B Port 2      │        │              │
   │ AVR7-3 Valve - User Assigned Name Port 4 │  └──────────────┘
   └────────────────────────────────────┘


 Step 8, Volume = 2.00 ml                   ┌──────┐   ┌────────┐
                                            │  OK  │   │ Cancel │
                                            └──────┘   └────────┘
```

---

Change
Valve  ◉       To select any valve and change its position.


• **Change Valve name and position**:  These fields allow you to choose a valve and to make a change in valve position.  Note that certain valve functions defined in the Setup screen (such as Fraction Collector Diverter, Aux Pump Inlet, Sample Inject, Inlet A, Inlet B)  are "tied" to other Protocol steps, so valve position changes will be made automatically. An example would be an AVR7-3 valve defined as a Sample Inject valve in the Setup screen, which is then tied to the Load/Inject Sample step in the Protocol screen.

  Valves assigned a "User-defined name" during Setup require the "Change Valve" step at the desired point in the Protocol.

• **OK**:  Adds the step to the protocol.  This is the same as pressing the **Enter** key on the keyboard.

• **Cancel**:  Does *not* add the step to the protocol.  This is the same as pressing the **Esc** key on the keyboard.

• **Step, Time** or **Volume**:  Identifies current step number, and calculates the elapsed time or volume from all previous steps.  This is not user editable.
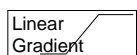
---

***Table 7-6.***
***Column Switching***

---

```
┌─────────────────────────────────────────────────────────┐
│ Edit Switch Columns                                    ☒ │
│  ┌──────────────┐                                         │
│  │ Switch Columns│                                        │
│  │              └──────────────────────────────────────┐ │
│  │ Position                                             │ │
│  │ ┌──────────────────────┬─┐                           │ │
│  │ │ Column 1             │▲│                           │ │
│  │ │ Column 2             │ │                           │ │
│  │ │ Column 3             │ │                           │ │
│  │ │ Column 4             │ │                           │ │
│  │ │ Column 5             │ │                           │ │
│  │ │ Column 6             │ │                           │ │
│  │ │ Column 7             │▼│                           │ │
│  │ └──────────────────────┴─┘                           │ │
│  │                                                      │ │
│  └──────────────────────────────────────────────────────┘ │
│  Step 8, Volume = 5.00 ml          ┌────────┐ ┌────────┐ │
│                                     │   OK   │ │ Cancel │ │
│                                     └────────┘ └────────┘ │
└─────────────────────────────────────────────────────────┘
```

---

Column Switching  The BioLogic Duo-Flow system supports the use of more than one column during a run. (Chapter 9 provides two examples of column switching during a run.)  Use this button to specify which column to load sample onto.

Column switching is supported by installing two AVR9-8 valves that permit the use of 8 different analytical and separation columns.  The AVR9-8 valves are defined by selecting Column Switching in the Setup screen.  This identifies the two valves as installed and synchronizes the switching of the valves without further user input.  Column switching permits identification of the electrical connection port of each valve and user naming for each column.  Installation of the AVR9-8 valve in Setup automatically activates Column Switching in the Protocol screen.

- **Position**:  Select the column to switch to in a run.

- **OK**:  Adds the step to the protocol.  This is the same as pressing the **Enter** key on the keyboard.

- **Cancel**:  Does *not* add the step to the protocol.  This is the same as pressing the **Esc** key on the keyboard.

- **Step, Time** or **Volume**:  Identifies current step number, and calculates the elapsed time or volume from all previous steps.  This is not user editable.
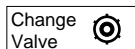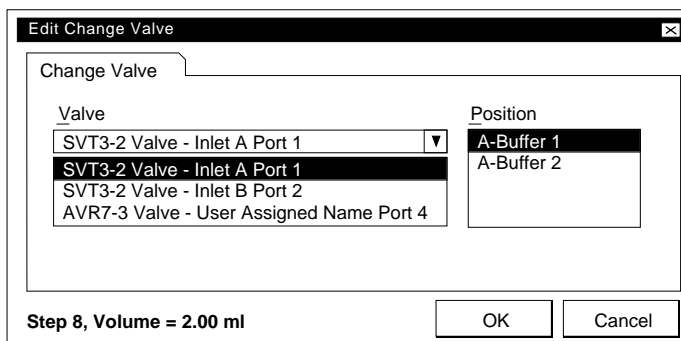
---

**Table 7-7.**
**Hold Until**

---

```
┌─────────────────────────────────────────────────────────────────────┐
│ Edit Hold Event                                                   [X] │
│  ┌─ Hold Until Event ─────────────────────────────────────────────┐  │
│  │ ┌─ Hold Until ──┐  ┌──────────────────┐  ┌─ Threshold Detector ─┐│  │
│  │ │ ● Key Pressed │  │ ☐ Time Out Reqd. │  │ UV Detector      [▼] ││  │
│  │ │               │  │                  │  │                      ││  │
│  │ │ ○ Start of    │  │ Time Out (min)   │  │ Threshold            ││  │
│  │ │   Inject      │  │ ┌──────┐ [▲]     │  │ ┌──────┐ [▲]         ││  │
│  │ │ ○ End of      │  │ └──────┘ [▼]     │  │ │1.000 │ [▼]         ││  │
│  │ │   Inject      │  │                  │  └──────────────────────┘│  │
│  │ │ ○ Above       │  │ The method will  │                          │  │
│  │ │   Threshold   │  │ automatically    │   ☐ Sound Alarm          │  │
│  │ │ ○ Below       │  │ continue after   │                          │  │
│  │ │   Threshold   │  │ the selected     │                          │  │
│  │ │               │  │ time-out duration│                          │  │
│  │ └───────────────┘  └──────────────────┘                          │  │
│  └────────────────────────────────────────────────────────────────┘  │
│  Step 1, Volume = 0.00 ml                      [   OK   ]  [ Cancel ]  │
└─────────────────────────────────────────────────────────────────────┘
```

---

| Hold ✋ | Inserts a Hold step into the method. The Hold step stops the progression of the method; the pumps continue pumping at the Hold step %B composition. The method time (or volume) advances. When the run resumes, the current fraction collection condition is maintained. The |

hold condition is maintained until a specified activity occurs (e.g., "Hold until Start of Inject" or "Hold until Keypress"). For example, if the hold starts at time 2 (minutes), then the method resumes at time 2, no matter how long the Hold was in effect.

**Hold Until** and **Threshold Detector**: Choose one of five events to discontinue the hold:

- **Key Pressed**: Press the **F2** key on the keyboard to end the programmed hold.

- **Start of Inject**: The run will continue when the manually-controlled device (which must be connected to the Workstation AUX connector at pin 1, Inject) is moved to its desired position.

- **End of Inject**: The run will continue when the manually-controlled device (which must be connected to the Workstation AUX connector at pin 1, Inject) is returned to its original position.

- **Above Threshold**: The run will continue when threshold exceeds that specified by the following fields in this window:

    **Threshold Detector**: Select the desired threshold detector absorbance, conductivity, RI, etc.

    **Threshold**: Specify a threshold value.

- **Below Threshold**: The run will continue when threshold falls below that specified by the following fields in this window:

    **Threshold Detector**: Select the desired threshold detector absorbance, conductivity, RI, etc.

    **Threshold**: Specify a threshold value.

- **Time Out Reqd** and **Time Out (min)**: To hold for a specified length of time.

- **Sound Alarm**: When this box is checked, an alarm will sound at the beginning of the step to remind you that the system requires an action on your part to allow the method to advance.

- **OK**: Adds the step to the protocol. This is the same as pressing the **Enter** key on the keyboard.

- **Cancel**: Does *not* add the step to the protocol. This is the same as pressing the **Esc** key on the keyboard.

- **Step, Time** or **Volume**: Identifies current step number, and calculates the elapsed time or volume from all previous steps. This is not user editable.

*Table 7-8.*
*Miscellaneous*
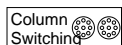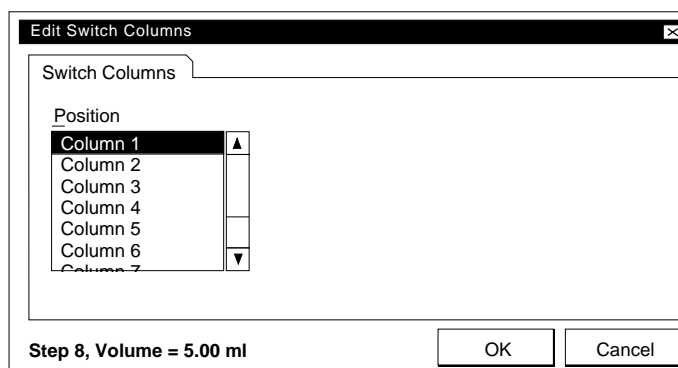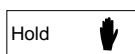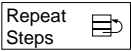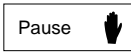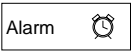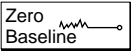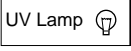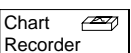
| Button | Description |
|--------|-------------|
| Repeat Steps | To repeat the highlighted step(s) a specified number of times. To highlight more than one step, hold down the Ctrl key while selecting steps with the mouse. |
| Pause | To pause the method at a specific step during the run. Pause stops the progression of the method and time, holds the %B composition, and stops the pumps. **Time Out Req'd** permits a pause time to be entered. The step will be paused for the specified length of time, after which it will automatically resume. An audible alarm may be programmed to sound when the step is paused and when it automatically resumes. |
| Alarm | To sound a 10 second audible alarm at the programmed time. The progress of the run will not be affected. |
| Zero Baseline | To zero the UV baseline at the programmed time. |
| UV Lamp | At system start-up, the default condition is Lamp ON. If the lamp is OFF, it takes about 20 minutes for the lamp to warm up prior to use. |
| Chart Recorder | To turn the Chart Recorder on/off at the programmed time. Inserts a step that tells the Chart Recorder to start/stop the paper feed and lower/lift the pen. |

**Table 7-9.**
**Fraction Collection**

| Button | Description: Collect All |
|---|---|
| Fraction Collection | To set up fraction collection.  Fraction collection schemes are discussed below. |

**IMPORTANT NOTE:**

The BioLogic Duo-Flow system is designed to control the Model 2128 Fraction Collector, which is the only collector that provides a choice of racks and the ability to overlay fractions from consecutive runs.

The BioLogic Duo-Flow system will also accommodate the Bio-Rad Model 2110 Fraction Collector as well as generic collectors.  To use a fraction collection scheme other than "Collect All"  with these collectors, it is **essential** to assign a SVT3-2 valve as a "Fraction Collector Diverter" in the Setup screen.  Rack choice and Overlay features are **not** available with these collectors.



**Collect All**:  The eluant from the entire run will be collected by the fraction collector.

- **Fraction Size**:  Enter the fraction size as volume (ml).

- **Rack**:  This applies only to the Model 2128 Fraction Collector.  The following racks may be specified:
    1    128 tubes; 12-13 mm tube diameter
    2    78 tubes; 16-18 mm tube diameter
    3    Microplates, 3 plates total for 288 wells (96 wells/plate, arranged 8x12)
    4    Microtubes (128 capless microtubes)
    5    Prep-Bottle (10 bottles of any size)
    6    Microplate: 72 wells (24 wells/plate x 3 plates)

- **Tubes Required**:  The displayed value is calculated based on the total volume divided by fraction size.

***Table 7-9. (continued)***
***Fraction Collection***

---

### Description: Collect All (continued)

---

- **Number of Racks**: Indicate the number of racks to be used for fraction collection. The Model 2128 Fraction Collector permits more than one rack to be used. The system will pause and a yellow prompt will tell you to change the rack and continue the method.

- **Start Tube** and **End Tube**: This applies only to the Model 2128 Fraction Collector. Identify the first and last tubes to receive fractions.

---

### Description: Threshold

---



**Threshold**: This method allows the flow to be sent to the collector if the detector signal of the eluant is above a certain signal level. **Note**: You can select any detection source to initiate threshold-based fraction collection. Non-peak eluant can be sent to waste or to the fraction collector.

- **Fraction Size, Rack**, **Tubes Required**, **Number of Racks**, **Start Tube,** and **End Tube**: See description for **Collect All** on page 7-16.

- **Threshold**: Enter the threshold value in absorbance units. Fractions will be collected whenever the detector output goes above the threshold level.

- **Non-peak parameters Destination** and **Fraction Size**: Enter the non-peak parameters. If the buffer stream is diverted to Waste, there is no need to enter a fraction size. But if you want to collect non-peak material, you must enter a fraction size.

*Table 7-9. (continued)*
*Fraction Collection*

**Description: Collection Windows**

```
 ┌──────────────────────────────────────────────────────────────────────────┐
 │ Fraction Collector Scheme │ Rack │ CollectionWindows │                     │
 │                                                                            │
 │  ○ Collect All          Fraction Size      Threshold              ┌──────┐ │
 │                         ┌─────┐▲           ┌─────┐▲               │ Close│ │
 │  ○ Threshold            │2.07 │▼ ml        │0.100│▼ AU            └──────┘ │
 │                         └─────┘            └─────┘                         │
 │  ● Collection Windows                  Non-peak Parameters:                │
 │                                        Destination:                        │
 │  ○ Threshold and Collection Windows     ◉ Waste    Fraction Size           │
 │                                         ○ Tubes    ┌─────┐▲ ml             │
 │                                                    │1.00 │▼                │
 │                                                    └─────┘                 │
 │ Volume                      Description                                    │
 └──────────────────────────────────────────────────────────────────────────┘
```

```
 ┌──────────────────────────────────────────────────────────────────────────┐
 │ Fraction Collector Scheme │ Rack │ CollectionWindows │                     │
 │  Rack                                          Start Tube         ┌──────┐ │
 │  1 (12-13mm tubes)                                                │ Close│ │
 │  2 (16-18 mm tubes        Tubes Required    5   ┌─────┐▲          └──────┘ │
 │  3 (Microplate(96 well plate)                   │1    │▼                   │
 │  4 (Microtubes)                                 └─────┘                    │
 │  5 (Prep Bottle)                                                           │
 │  6 Microplate(24 well plate)   Number of racks  End Tube                   │
 │                                ┌─────┐▲         ┌─────┐▲                   │
 │                                │1    │▼         │288  │▼                   │
 │                                └─────┘          └─────┘                    │
 │ Volume                      Description                                    │
 └──────────────────────────────────────────────────────────────────────────┘
```

```
 ┌──────────────────────────────────────────────────────────────────────────┐
 │ Fraction Collector Scheme │ Rack │ CollectionWindows │                     │
 │       Start      End       Frac. Size                             ┌──────┐ │
 │       (ml)       (ml)      (ml)                                    │ Close│ │
 │      ┌────┐     ┌────┐    ┌─────┐                                  └──────┘ │
 │      │0.0 │     │4.0 │    │0.60 │                                  ADD MODE │
 │      └────┘     └────┘    └─────┘                                           │
 │   ┌───┬──────┬──────┬──────┬─────┬─────┬─────┐    ┌──────────────┐         │
 │   │ 1 │ 4.0  │ 5.0  │ 0.60 │     │     │     │    │ Save Window  │         │
 │   ├───┼──────┼──────┼──────┼─────┼─────┼─────┤    └──────────────┘         │
 │   │ 2 │ 7.0  │ 6.0  │ 1.00 │     │     │     │    ┌──────────────┐         │
 │   ├───┴──────┴──────┴──────┴─────┴─────┴─────┤    │ Delete Window│         │
 │   │                                          │    └──────────────┘         │
 │   └──────────────────────────────────────────┘    ┌──────────────┐        │
 │                                                    │Finished Adding│       │
 │                                                    └──────────────┘        │
 └──────────────────────────────────────────────────────────────────────────┘
```

**Collection Windows**:  This method specifies collection during specified parts of the run (Time or Volume). Each collection window can have a unique fraction size (ml).

**Note**:  Collection windows programming should be done after all other steps have been written.  When using Collection Windows, the Model 2128 will skip a tube between each window collection.  For example, if the first window ends collection at 20 minutes and the second window starts collection starting at 21 minutes, then the fraction collector will leave an empty tube for the period between minutes 20 and 21.

- **Fraction Size, Rack**, **Tubes Required**, **Number of Racks**, **Start Tube,** and **End Tube**:  See description for **Collect All** on page 7-16.

- **Add Mode**: Enter the desired values for **Start**, **End**, and **Frac Size**.
  a.  Select the **Save Window** button and the values will appear in line 1.
  b.  Enter the values for the next window and select **Save Window.** The values will appear in line 2.
  c.  When the collection scheme is complete, select **Finished Adding,** and the Select Mode will appear.

- **Select Mode**: This mode (not shown in illustration above) shows the following two buttons:
  **Add Window**: Inserts a window after the highlighted window.
  **Delete Window**: Deletes that window.

***Table 7-9. (continued)***
***Fraction Collection***

**Description: Threshold and Collection Windows**

---

Fraction Collector Scheme | Rack | CollectionWindows

- ○ Collect All
- ○ Threshold
- ○ Collection Windows
- ● Threshold and Collection Windows

Fraction Size: 2.07 ml   Threshold: 0.100 AU

Non-peak Parameters:
Destination:
- ● Waste
- ○ Tubes   Fraction Size: 1.00 ml

Close

**Volume**                    **Description**

---

Fraction Collector Scheme | Rack | CollectionWindows

Rack
- 1 (12-13mm tubes)
- 2 (16-18 mm tubes)
- 3 (Microplate(96 well plate)
- 4 (Microtubes)
- 5 (Prep Bottle)
- 6 Microplate(24 well plate)

Start Tube: 1

Number of racks: 1

End Tube: 288

Close

**Volume**                    **Description**

---

Fraction Collector Scheme | Rack | CollectionWindows

| | Start (ml) | End (ml) | Frac. Size (ml) | Thresh (AU) | Non-Peak Frac. Size (ml) | |
|---|---|---|---|---|---|---|
| | 0.0 | 5.0 | 1.00 | 1.00 | ● Waste  ○ Tubes  0.01 | |
| 1 | 4.0 | 5.0 | 0.60 | 0.000 | Waste | |
| 2 | 7.0 | 6.0 | 1.00 | 0.010 | Waste | |

Close

ADD MODE

Save Window

Delete Window

Finished Adding

---

Fraction Collector Scheme | Rack | CollectionWindows

| | Start (ml) | End (ml) | Frac. Size (ml) | Thresh. (AU) | Non-Peak Frac. Size (ml) | |
|---|---|---|---|---|---|---|
| 1 | 4.0 | 5.0 | 0.60 | 0.000 | Waste | |
| 2 | 7.0 | 6.0 | 1.00 | 0.010 | Waste | |

Close

SELECT MODE

Add Window

Delete Window

---

**Threshold and Collection Windows**:  Each Collection Window (discussed on page 7-18) can also contain a threshold value to further discriminate when eluant can be collected.  For each collection window, non-peak effluent can be directed to waste or to the fraction collector. The Threshold and Time windows can be combined, so the system collects fractions only during the programmed Time windows and within the Time windows only when UV absorbance is above the threshold value.

*Table 7-9. (continued)*
*Fraction Collection*
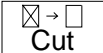
**Description: Threshold and Collection Windows (continued)**

When using the Collection Windows functions, the Model 2128 will skip a tube between each window collection.  For example, if the first window ends collection at 20 minutes and the second window starts collection starting at 21 minutes, then the fraction collector will leave an empty tube for the period between minutes 20 and 21.

- **Fraction Size, Rack**, **Tubes Required**, **Number of Racks**, **Start Tube,** and **End Tube**:  See description for **Collect All**.

- **Start (ml)**, **End (ml)**, **Fraction Size (ml), Thresh (AU),** and **Non-Peak Frac Size (ml)**:  Each window is defined by these parameters.  Use the scroll bar to the right of the window display to scroll through the list of windows.  Enter the fraction size as volume (ml).

- **Add Mode**: Enter the desired values for **Start**, **End**, **Frac Size**, and **Thresh**.

   Select the **Save Window** button and the values will appear in line 1.

   To enter the values for the next window, select **Add Window** and enter the values. Select **Save Window** and the values will appear in line 2.

   When the collection scheme is complete, select **Finished Adding,** and the Select Mode will appear.

- **Select Mode**: This mode shows the following two buttons:

   **Add Window**: Inserts a window after the highlighted window.

   **Delete Window**: Deletes that window.

- **Non-peak parameters Destination** and **Fraction Size**:  Enter the non-peak parameters.  When collecting non-peak material, you must enter a fraction size.

The following Toolbar menu options are available only when creating and editing a protocol.

***Table 7-10.***
***Protocol Screen's Editing Toolbar***

| Button | Description |
|---|---|
| **Edit buttons:**<br><br>☐→☐<br>Edit<br><br>☒→☐<br>Cut<br><br>☐→☐<br>Copy<br><br>☐←☐<br>Paste<br><br>☒<br>Delete | These buttons are available from the System menu.<br><br>To edit the currently highlighted step(s) in the protocol.  If multiple steps are highlighted any changes will be reflected in all steps.<br><br>To cut the currently highlighted step from the protocol.  It places the step in the clipboard, which means it can be placed elsewhere by using the **Paste** button.<br><br>To copy the currently highlighted step in the protocol.<br><br>To paste the cut or copied step before the currently highlighted step.<br><br>To delete the currently highlighted step from the protocol. |

## 7.4    RUN SCREEN

The Run screen displays a run in progress.  All data associated with the run are automatically saved.  An example of a run in progress is shown in Figure 7-5 below.  Table 7-11 discusses the buttons which may be used to control the run.  Note: If the Bio-Rad EzLogic Integration software is installed, the **Integ**. toolbar button appears.
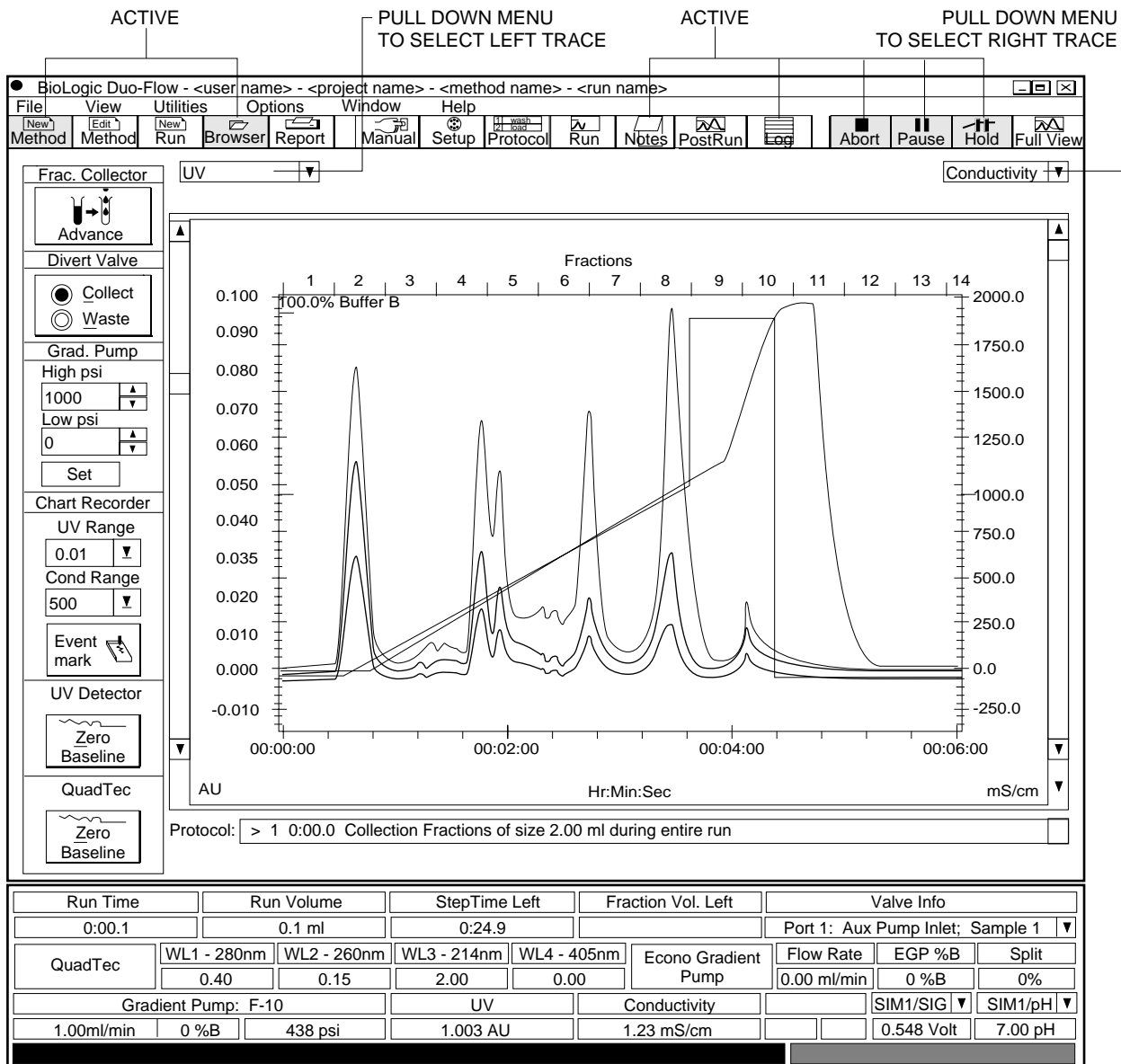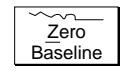


**Figure 7-5.  Run Screen showing a Run in Progress**

**Table 7-11.**
**Run Screen's Control**

| Button | Description |
|---|---|
| **Frac. Collector** Advance | Advances the fraction collector to the next tube. Pressing this button does not modify the method. The event is recorded in the Run Log. |
| **Divert Valve** ○ Collect ● Waste | Immediately changes the position of the diverter valve. Pressing this button does not modify the method. The event is recorded in the Run Log. |
| **Grad. Pump** High psi 400 Low psi 20 Set | To set the pressure limits of the Workstation pump. This is not part of a protocol; it is a system setting from the Manual screen. The pumps will stop when the pressure goes above or below the set values. |
| **Chart Recorder** UV Range 0.2 Cond Range 500 Event mark | To set the **UV Detector** range for the Chart Recorder output. The settings are: 0.001, 0.002, 0.005, 0.01, 0.02, 0.05, 0.1, 0.2, 0.5, 1.0, 2.0 AUFS.  To set the **Conductivity** range on the Chart Recorder output. The settings are: 500, 200, 100, 50, 20, 10, 5 mS/cm.  To set an **Event Mark** on the chart recorder UV trace output only. It will not record event marks on the screen. |
| Zero Baseline | Use the **Zero Baseline** button to reset the UV or QuadTec detector absorbance value to zero. **Note:** Use care during a collection scheme which uses a **Threshold** value. |
| UV ▼ Conductivity ▼ | These drop-down menus, located on either side of the chromatogram, are used to select the active data trace to be scaled during or after a run. As each trace scale is selected, the scroll bars can be used to adjust the scale setting. |
| New Method Browser | To work offline while a run is in progress. (Refer to section 7.4.2, Working Offline.) |
| Notes Log | Notes displays the Run Notebook screen, which allows you to enter notes regarding the run. Log displays the Run Log screen, which provides information about the run and is non-editable. |
| ■ Abort ‖ Pause ⟙ Hold | To stop, pause, or hold a method in progress. (Refer to Figure 7-6.) |
| Settings | To select which traces to display in the chromatograms. (Refer to Table 5-2.) |

### 7.4.1  Pausing/Stopping a Method in Progress

The Run screen provides three options for pausing or stopping a method in progress, as discussed in the figure below.



**Abort**

Stops the current run and saves the data. (A confirmation message is presented to prevent accidents.) The run cannot be continued.

**Pause**

Stops progression of method, method time, and pumps. Valves remain in paused position. From Pause, you can Continue, Abort, or Edit Protocol.

**Hold**

Stops progression of method and method time. Pumps continue pumping at Hold %B concentration, and valves remain in Hold position. From Hold you can Continue, Abort, or Pause. You cannot edit.

**Setup**

Setup screen lets you view the list of devices available to the method. From Setup, you can select Protocol or Run. From Run, select Abort or Continue.

**Protocol**

Protocol screen (View only) lets you view the steps in the run. Select Run or Edit Protocol. From Run, select Abort or Continue. Edit Protocol involves the buttons shown below.

**Continue**

To continue the method run.

**Edit Protocol**

Pressing this button initiates the process of editing the method's protocol.

Method Name:
Method Description:
Method Author:

If there's a run already associated with the method, the method is copied and you must rename it.

You may edit the method's protocol using any of the Add Step buttons and the Cut, Copy, Paste, and Delete buttons.

**Run**

After editing the method's protocol, return to the Run screen. You then can Continue or Abort the run.

***Figure 7-6.  Run Screen's Abort, Pause, and Hold Buttons***

### 7.4.2   Working Offline During a Run

The BioLogic Duo-Flow system software allows you to work offline, creating a new method or editing an existing method, while a run is in progress.  The following actions can be performed:

- Write a new method
- Edit an existing method
- View results from a completed run
- Print a report from a completed run
- Integrate run data from a completed run (using the optional EZLogic software)
- Perform PostRun analysis on a completed run
- Export Data or Export Chromatogram Image of a completed run
- Access the HELP screen

These functions are not available while a run is in progress:

- PostRun analysis of the Run in progress
- Integration of the data from the Run in progress
- Initiation of a new Run
- Utilities functions, including calibrating pH probe, gradient pumps, conductivity
- Manual mode functions

During a run, the following toolbar buttons activate the offline window:

- **Browser** button

The online and offline windows are distinguished as follows:

- Online:  The BioLogic Duo-Flow icon in the upper left of the screen is green, and the title bar does not indicate online.
- Offline:  The BioLogic Duo-Flow icon in the upper left of the screen is yellow, and the title bar indicates offline.

To change between the online and offline windows, you can use any of the following:

- Windows drop down menu:  The online and offline windows are listed.
- Task bar: The Windows® taskbar lists the BioLogic Duo-Flow windows.
- Alt-Tab:  Simultaneously hold down these two keys to list the open windows.

### 7.4.3    Editing A Method During a Run

During a run, you may find you want to change some of the run parameters.  The figure below shows how this is handled by the system and some of the restrictions.
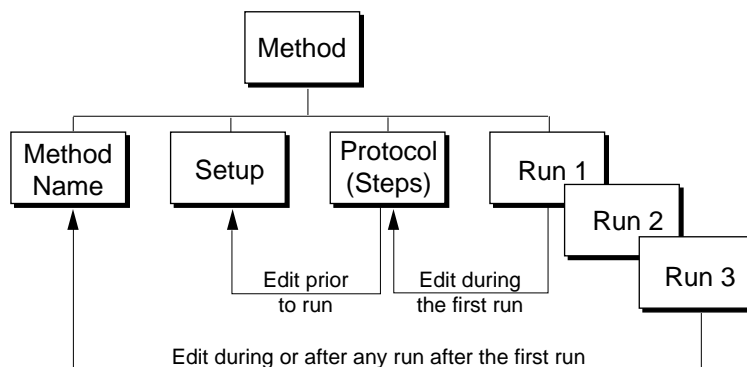


***Figure 7-7.  Editing during a Run***

- During the first run of a method, you can **pause** the method and return to the Protocol screen to edit steps that have not already been started. The method name will not change.

- Any run after the first can be paused and then edited, except runs in queue which cannot be edited. You will first be asked to rename the method (or accept the default new name) when the **Edit Protocol** button is clicked. This is to ensure the integrity of the database in terms of which Method Protocol particular runs are tied to.

   For example, assume you make three runs under a method named "Ion-exchange UNO Q1." During the fourth run, you may decide to pause the run, and enter the Protocol screen to increase the length of the salt gradient. In this case, you will first be prompted to rename the Method (e.g., Ion-exchange UNO Q1 Rev. 1) before proceeding with the edit.

   Upon completion of the run, the database of Methods and Runs will show three runs associated with the method "Ion-exchange UNO Q1" and one run associated with the method "Ion-exchange UNO Q1 Rev. 1."

- A fraction collection scheme is editable, but it cannot be added to a Protocol once a run has been started. In such cases, you should abort the run and s elect **Edit the Method**.

- A currently programmed fraction collection scheme may be edited during a run, but note that changing the fraction size of a "Collect All" scheme will first require selection of  "Collection Windows." This is because the editing process effectively turns the initial "Collect All" scheme into the "first" collection window.

| | | | | | | |
|---|---|---|---|---|---|---|
| 1 | 0.0 | Collect Fractions during entire method | | | | |
| 2 | 0.0 | Chart Recorder | Turn ON | | | |
| 3 | 2.0 | Isocratic Flow | 25mM Tris | 100% | Time: 2.0 min | |
| | | | 25mM Tris + Salt | 0% | Flow: 2.0 ml/min | |
| 4 | 3.0 | Load: Sample | Static Loop | | Auto Inject Valve | |
| | | | Load: standard | Volume: 1.0 ml | Flow: 2.0 ml/min | |
| 5 | 7.0 | Isocratic Flow | 25mM Tris | 100% | Volume: 4.0 min | |
| | | | 25mM Tris + Salt | 0% | Flow: 2.0 ml/min | |
| 6 | 23.0 | Linear Gradient | 25mM Tris | 100% --> 55% | Volume: 16.0 min | |
| | | | 25mM Tris + Salt | 0% --> 45% | Flow: 2.0 ml/min | |
| 7 | 25.0 | Linear Gradient | 25mM Tris | 55% --> 100% | Volume: 2.0 min | |
| | | | 25mM Tris + Salt | 45% --> 0% | Flow: 2.0 ml/min | |
| 8 | 28.0 | Isocratic Flow | 25mM Tris | 100% | Volume: 3.0 min | |
| | | | 25mM Tris + Salt | 0% | Flow: 2.0 ml/min | |
| | | End of Protocol | | | | |

Add Step:

Edit Protocol

View-Only Mode

Fraction Collection

BioLogic Duo-Flow - offline

File   Edit   View   Utilities   Options   Window   Help

New Method   Edit Method   New Run   Browser   Report   Manual   Setup   Protocol   Run   Notes   PostRun   Log   Settings   Edit   Cut   Copy   Paste   Delete

| Gradient Pump: F-10 | | | UV | Conductivity | | SIM1/SIG ▼ | SIM1/pH ▼ |
|---|---|---|---|---|---|---|---|
| 1.00ml/min | 0 %B | 438 psi | 1.003 AU | 1.23 mS/cm | | 0.548 Volt | 7.00 pH |

**Figure 7-8.  Protocol Screen during a Run**

### 7.4.4    Run Notebook

Use the Run Notebook screen to maintain any information you want regarding the run. It contains fields for description of the sample, the column, the operator, the buffer(s), flow rates, gradients, chart speed, fraction size, and any other information you might want to enter.



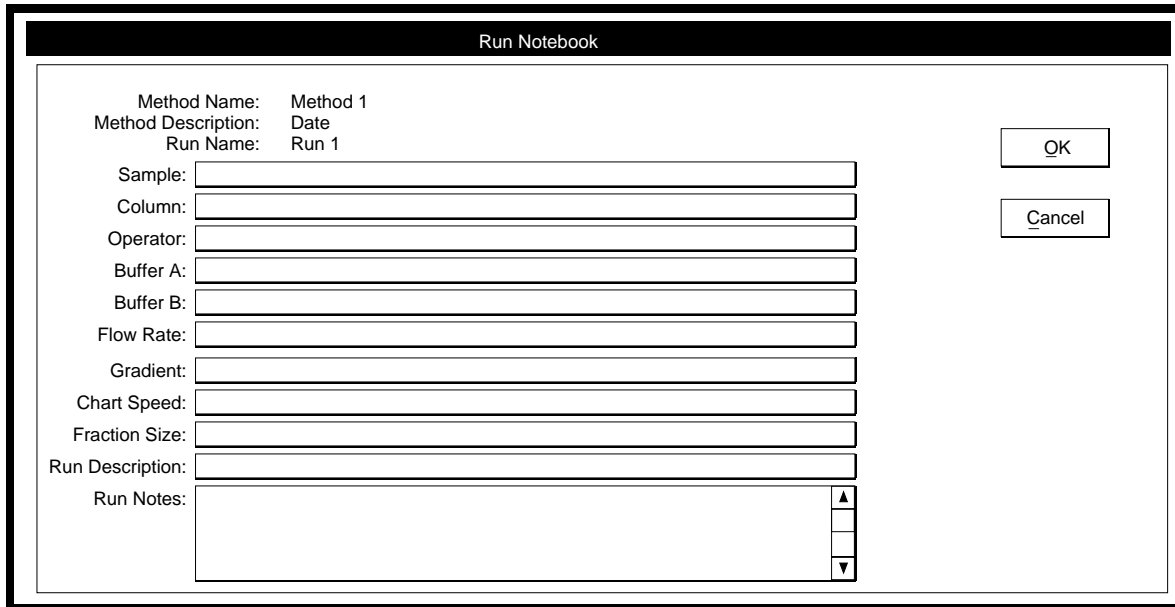*Figure 7-9. Run Notebook Screen*

### 7.4.5    Run Log

The Run Log screen details the time and order of the execution of each step, error, and/or event of the run. This information cannot be edited. It is possible, although not recommended, to turn off the Run Log by deselecting its checkbox in the Edit User Preferences window, available from the **Options** menu.
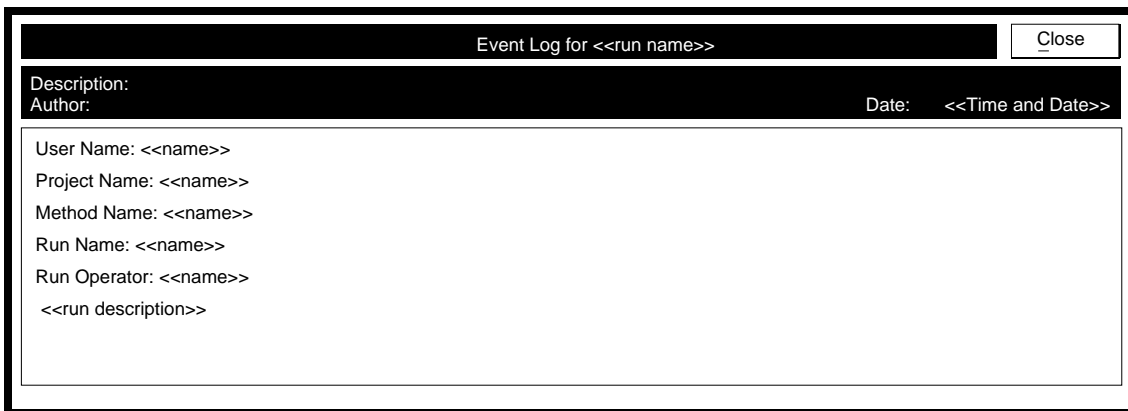


*Figure 7-10. Run Log Screen*

### 7.5    POST-RUN

The Post-Run screen enables peak annotation ("tagging") and data export.  To enter the Post-Run screen, open a Run file in the Browser screen, or when a run is finished choose **PostRun** from the toolbar.  Two chromatograms are viewed:  the main chromatogram in the lower large window and the reference chromatogram in the upper right window.  In the upper reference chromatogram, use the sliding sidebar to make scaling adjustment.

**Note**: The Values at Cursor window in the upper left corner will list fewer traces if you are not using the QuadTec multiwavelength detector.
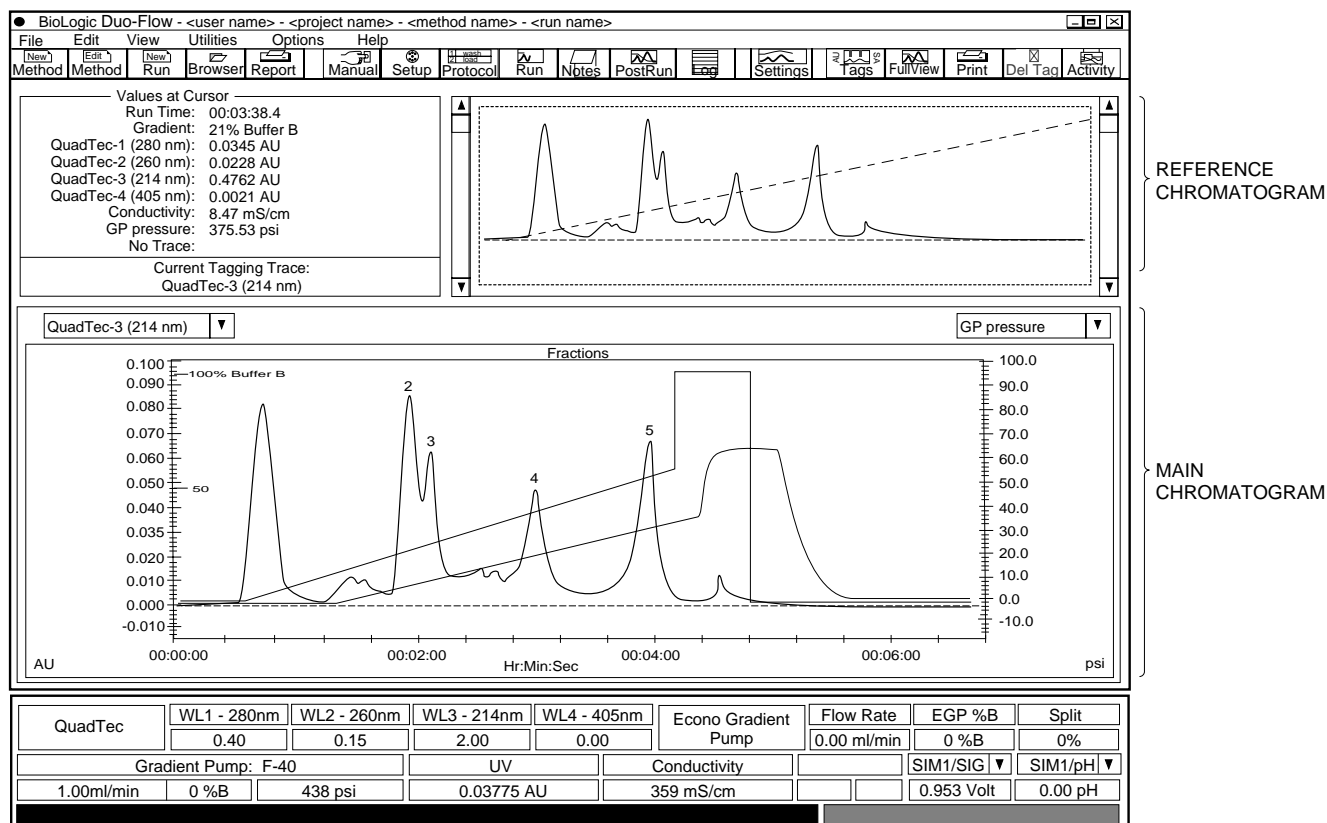


*Figure 7-11.  Post-Run Screen*

### 7.5.1    Resizing

The main chromatogram shows the current "Zoom" region.  To resize the image (or "zoom in" to a particular region of the chromatogram), either change the axes scale in the reference chromatogram, or use the "rubber band" controls on either chromatogram.  To use the rubber band controls, click and hold the left mouse button. Drag it across the chromatogram region.  Release the mouse button and the main chromatogram will be resized.  Click on the **Full View** button in the toolbar to return to normal view.

### 7.5.2    Chromatogram Information (Values at Cursor)

The top left corner of the PostRun screen contains information about the chromatogram.  The position of the vertical bar within the main chromatogram dictates the displayed information.  Run time, the absorbance (AU) and conductivity (mS/cm) values, gradient progression (%B), and the current tagging trace are displayed for the peak selected by panning the vertical bar across the chromatogram.  To view the vertical bar, move the cursor inside the main chromatogram.

### 7.5.3    Annotating ("tagging") the Chromatogram

Tags can be placed on any trace. To assign tags to a trace, and to view all assigned tags and their data, use the Post-Run Tags window, shown below. To display the Post-Run Tags window, click once on the **Tags** button in the toolbar or from the Edit drop down menu, select **Tag and Trace Options.**

In the Post-Run Tags window, select a trace by clicking on an Active Trace radio button. Trace visibility and tag visibility can also be selected from this box. Tag styles are specified in the upper right box. Peaks can be annotated with a sequential tag number (1,2...), a user-defined name (user tag name), with the trace value, or the run time.

| Tag # | Tag name | Retention time (h:m:s) | Volume (ml) | Buffer B (%B) | UV (AU) | Cond. (mS/cm) | (no trace) | (no trace) | Tube # |
|---|---|---|---|---|---|---|---|---|---|
| 1 | Myoglobin | 00:00:32 | 2.2 | 0 | 0.07992 | 1.496 | N/A | N/A | 0.07992 |
| 2 | Conalbumin, peak 1 | 00:01:41 | 6.7 | 18 | 0.08617 | 7.999 | N/A | N/A | 0.08617 |
| 3 | Conalbumin, peak 2 | 00:01:48 | 7.2 | 20 | 0.05921 | 9.025 | N/A | N/A | 0.05921 |
| 4 | Ovalbumin | 00:02:44 | 11.0 | 34 | 0.03799 | 16.472 | N/A | N/A | 0.03799 |
| 5 | Soybean Trypsin Inhibitor | 00:03:30 | 14.0 | 46 | 0.06844 | 22.308 | N/A | N/A | 22.308 |
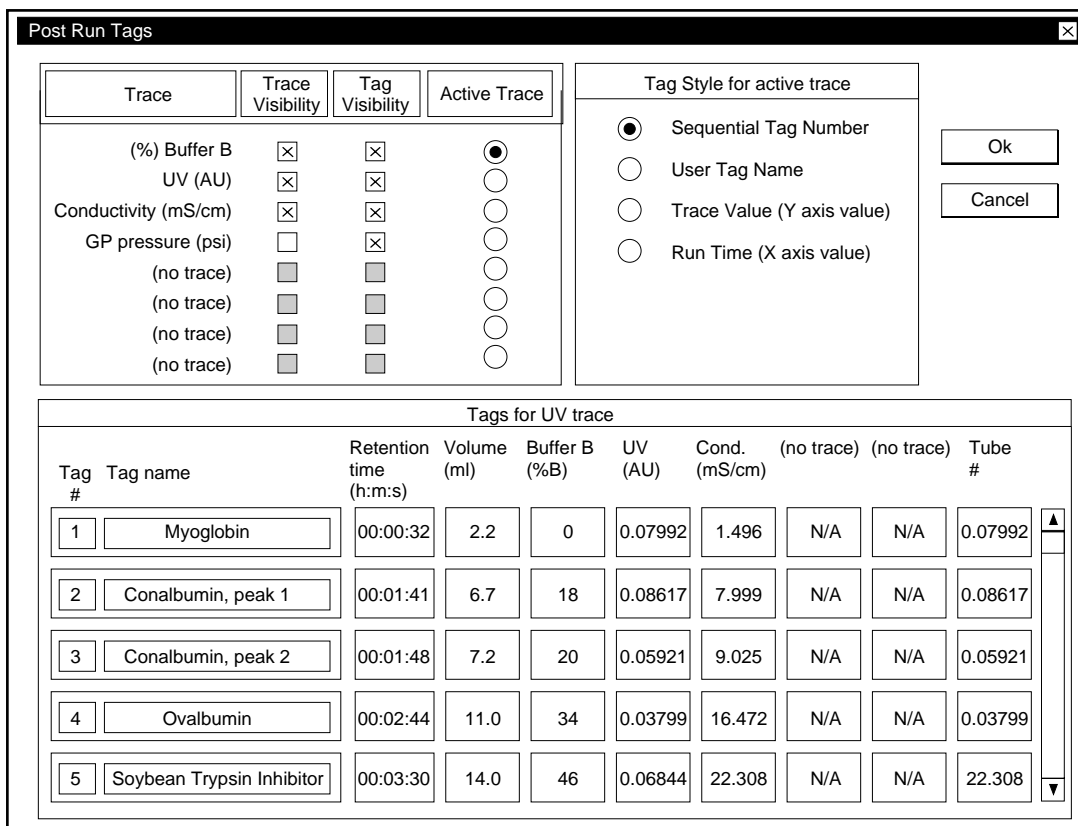
**Figure 7-12.  Post-Run Tags for Standard UV Detector**
**(Note: Additional Traces will be shown when the QuadTec Detector is being used.)**

To change a tag name in the Post-Run screen chromatogram, move the cursor over the tag and note that the cursor changes. Double click on the tag and in the window that appears, enter the tag name. To deselect a tag, highlight the undesired tag and press DelTag from the button bar. To remove all tags, select "Delete all tags" from the **Edit** drop-down menu.

### 7.5.4    Entering Activity Data

The Activity Trace Editor permits data collected by a separate offline method to be included with data collected by the Duo-Flow method.  For example, if fractions collected by the Duo-Flow are also analyzed by an ELISA method or for radioactivity, CPM, or DPM, the data collected by these assays can be entered into the Activity Trace Editor and a trace will appear on screen reflecting the activity for each fraction.

From the Browser screen, select the run for which you wish to enter the post-run activity data. From the post-run screen that is displayed:

1.  Select the **Activity** icon from the toolbar.  A box will appear listing each fraction collected in the opened run.  (See Figure 7-13.)

2.  Enter values for each fraction and enter the desired units.

3.  The activity values entered will appear along with the chromatogram data that appears in the upper left hand corner as the cross hatch is moved across each peak and fraction. (See Figure 7-14.)
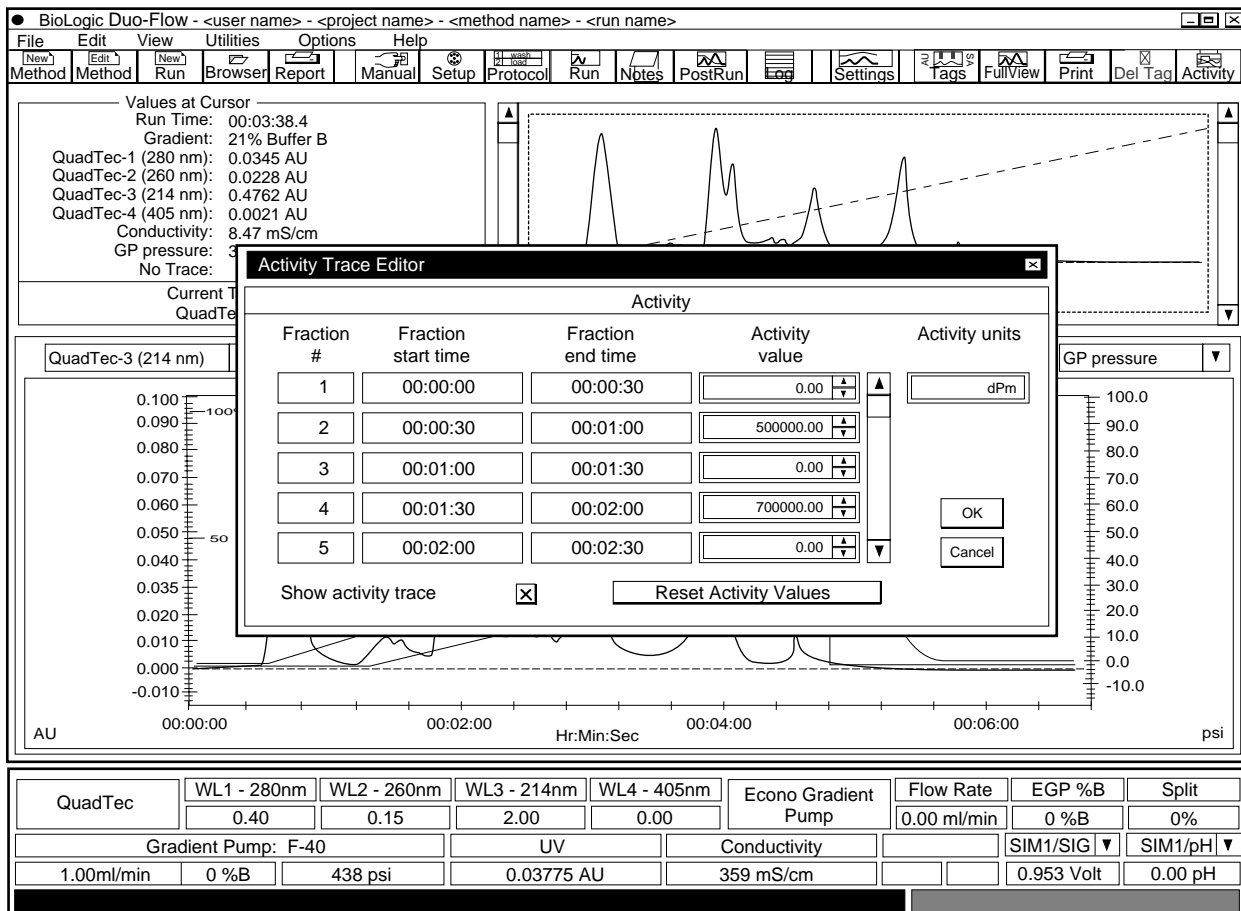


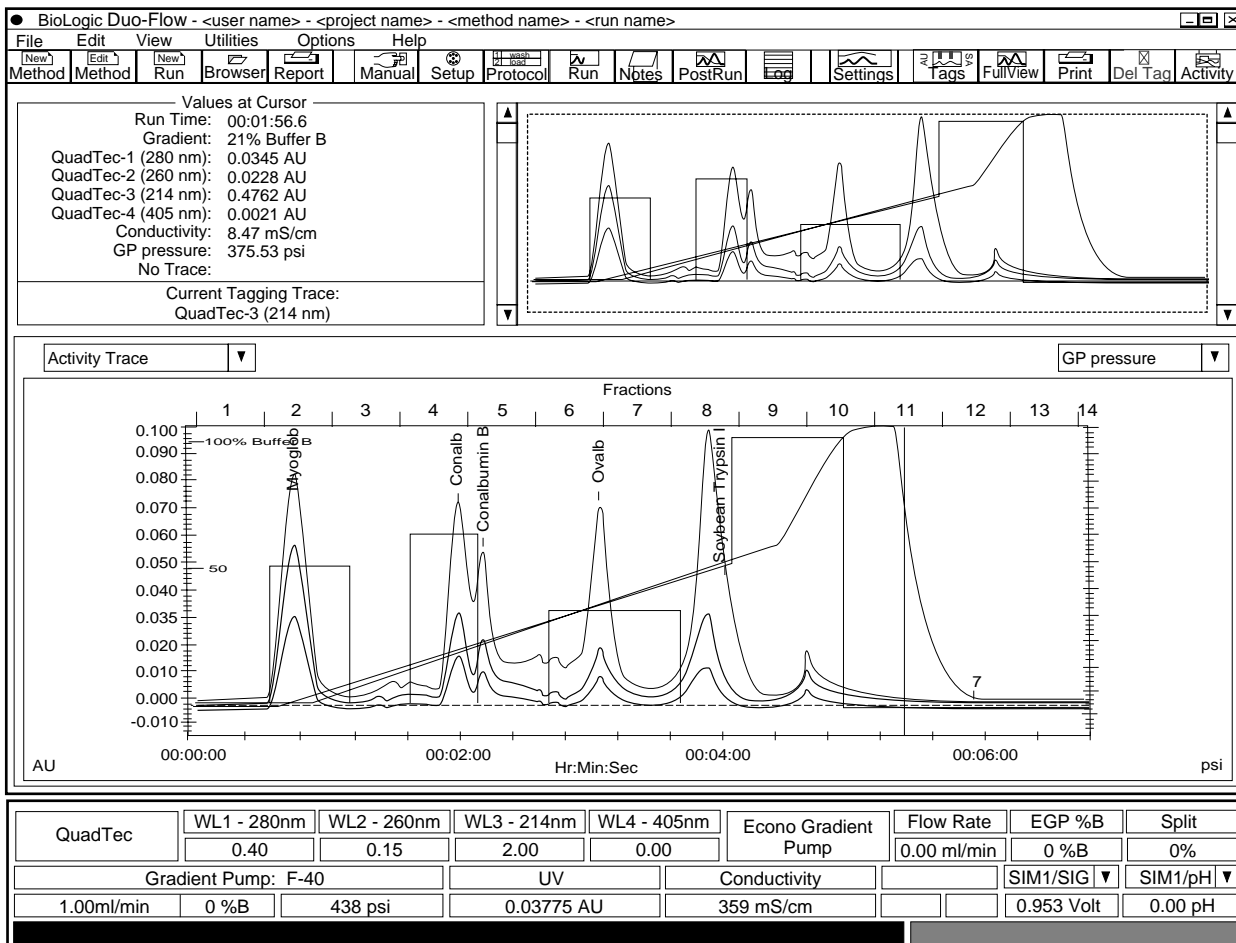**Figure 7-13.  Activity Trace Editor**

**Figure 7-14. Activity Trace**

### 7.5.5    Exporting Chromatogram Data to other Software Applications

To transfer data, go to the Post-Run screen of the desired run.  Select **Export Data** from the File drop-down menu (See Figure 7-15, Export Data Setup screen).  From this screen, choose the run time and the data trace(s) that will be exported.  Although data is collected at a rate of five points per second, it can be exported at a user-defined rate.  Data is exported in an ASCII.TXT file format.  The file name is user-defined and can be saved anywhere on the hard drive.  Note that the default file is called "Export" and is located in the following directory, "Biologic/data/export".  When importing the data into Excel, choose comma delimited.

```
┌─────────────────────────────────────────────────────────────────────────┐
│ Export Data Setup                                                     ⊠   │
│  ┌──────────────────────────────────────────────┐   ┌─────────────────┐  │
│  │          Export Run Time Window              │   │       Ok        │  │
│  │                                              │   └─────────────────┘  │
│  │  From: │0.0 │ ▼ ▲▼  Min   To: │10.5 │ ▲▼  Min │   ┌─────────────────┐  │
│  │                                              │   │     Cancel      │  │
│  └──────────────────────────────────────────────┘   └─────────────────┘  │
│  ┌──────────────────────────────────────────────────────────────────┐    │
│  │          Export Parameters      /      Export Rate               │    │
│  │     ⊠   UV              ┌─────────────┐                           │    │
│  │     ⊠   Conductivity    │ 5.0      ▼  │ points/sec.               │    │
│  │     ▢                   └─────────────┘                           │    │
│  │     ▢                                                             │    │
│  │     ▢                      ┌──────────────────────────────┐       │    │
│  │     ▢                      │        Calculate             │       │    │
│  │     □   GP Pressure        │  ┌──────────┐  ┌──────────┐  │       │    │
│  │     □   Gradient %B        │  │ 135   KB │  │  3150    │  │       │    │
│  │     □   Volume             │    File Size   Number of    │       │    │
│  │     □   Fraction           │                 Records     │       │    │
│  │                            └──────────────────────────────┘       │    │
│  └──────────────────────────────────────────────────────────────────┘    │
└─────────────────────────────────────────────────────────────────────────┘
```

**Figure 7-15.  Export Data Setup Screen**

### 7.5.5    Exporting Chromatogram Images

Chromatographic images can be exported into other applications, such as word processing programs. Before transferring an image, modify the chromatogram in the PostRun screen since changes cannot be made to the file once it is transferred out of the BioLogic Duo-Flow program. The image in the main chromatogram will be the exported image.

Select Export Chromatogram Image from the File drop-down menu (see Figure 7-16). The image is exported as an Enhanced Meta File (*.emf) format. In general, the Enhanced Meta File format is used for transfer into a Windows-based application.

Alternatively, a chromatogram can be copied to the Windows clipboard and pasted into another Windows program. Choose "Copy zoom Chromatogram to clipboard" from the Edit drop-down menu.



**Export Chromatogram Image File**                                        [?][X]

Save in:    [🗀 Export ▼] [↰] [⌨*]    [▦][▦]

File name:    [*.emf]                              [ Save ]
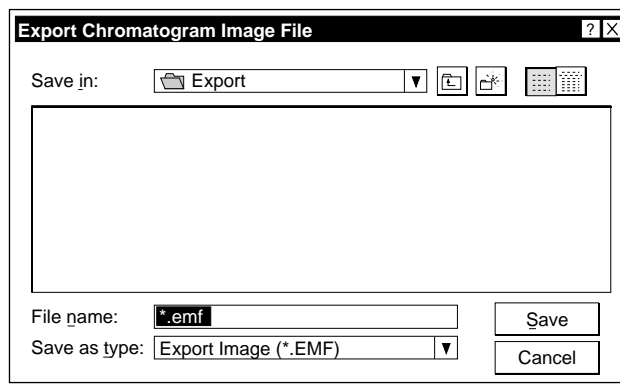Save as type:  [Export Image (*.EMF)  ▼]          [ Cancel ]

*Figure 7-16.  Exporting a Chromatographic Image*