

CAINTEGRATOR2 v.1.0

User's Guide



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TABLE OF CONTENTS

| | |
|---|-----------|
| Credits and Resources | i |
| Using the caIntegrator2 v.1.0 User's Guide | 1 |
| Introduction to the caIntegrator2 User's Guide | 1 |
| Organization of this Guide | 1 |
| User's Guide Text Conventions | 2 |
| Chapter 1 | |
| Getting Started with caIntegrator2 | 5 |
| About caIntegrator2 | 5 |
| Registering as a New caIntegrator2 User | 6 |
| Logging In | 8 |
| Using the caIntegrator2 Workspace | 8 |
| caIntegrator2 Functions | 9 |
| Using Online Help | 10 |
| Logging Out | 10 |
| Application Support | 11 |
| Chapter 2 | |
| Creating a New Study | 13 |
| Creating a Study – Overview | 13 |
| Configuring and Deploying a Study | 14 |
| Creating/Editing a Study | 15 |
| Adding Clinical Data | 16 |
| Adding/Editing Genomic Data | 24 |
| Adding Imaging Data | 29 |
| Deploying the Study | 31 |
| Managing a Study | 31 |
| Managing Platforms | 32 |

Chapter 3

Searching a caIntegrator2 Study35

| | |
|-------------------------------|----|
| Search Overview | 35 |
| Searching a Study | 36 |
| Results Type Tab | 40 |
| Sorting Tab | 42 |
| Managing Queries | 43 |
| Saving a Query | 43 |
| Editing a Query | 43 |
| Exporting Query Results | 43 |

Chapter 4

Viewing Query Results45

| | |
|--|----|
| Query Results Overview | 45 |
| Browsing Query Results | 46 |
| Clinical and Imaging Data | 46 |
| Genomic Data | 46 |
| Expanding Imaging Data Results | 48 |
| Relationship of Patient to Study to Series to Images | 51 |
| Exporting Data | 52 |

Chapter 5

Analyzing Studies53

| | |
|---|----|
| Data Analysis Overview | 53 |
| Creating Kaplan-Meier Plots | 54 |
| K-M Plot for Annotations | 54 |
| K-M Plot for Gene Expression | 56 |
| K-M Plot for Queries | 58 |
| Creating Gene Expression Plots | 60 |
| Gene Expression Value Plot for Annotation | 61 |
| Gene Expression Value Plot for Genomic Queries | 64 |
| Gene Expression Value Plot for Clinical Queries | 66 |
| Understanding a Gene Expression Plot | 70 |
| Analyzing Data with GenePattern | 73 |
| GenePattern Modules | 75 |
| Comparative Marker Selection (CMS) Analysis | 76 |
| Principal Component Analysis (PCA) | 78 |
| GISTIC-Supported Analysis | 81 |

Chapter 6

Administering User Accounts85

| | |
|---|----|
| Administering caIntegrator2 User Accounts Using UPT | 85 |
| Steps for Creating User Access to caIntegrator2 | 86 |
| Creating a New caIntegrator2 User | 86 |
| Creating a New User Group | 88 |
| Creating a New Protection Group | 89 |
| Assigning a User Group to a Protection Group | 90 |
| Adding a User to a User Group | 93 |
| Changing a User Password | 95 |

Appendix A

Data Import Configurations97

| | |
|---|----|
| Subject Clinical Data Configuration | 97 |
| Delimited-Text Annotation Import | 97 |
| Annotation Field Configuration | 98 |
| Sample Data Configuration | 98 |
| Genomic Data Configuration | 99 |
| Imaging Data Configuration | 99 |

Index101

USING THE CAINTEGRATOR2 v.1.0 USER'S GUIDE

This chapter introduces you to the *calIntegrator2 v.1.0 User's Guide* and suggests ways you can maximize its use.

Topics in this chapter include:

- [Introduction to the calIntegrator2 User's Guide](#) on this page
- [Organization of this Guide](#) on this page
- [User's Guide Text Conventions](#) on page 2

Introduction to the calIntegrator2 User's Guide

The *calIntegrator2 v.1.0 User's Guide* is the companion documentation to the calIntegrator2 software application. The user's guide includes information and instructions for the end user about using calIntegrator2.

Organization of this Guide

The *calIntegrator2 v.1.0 User's Guide* contains the following chapters and appendices:

Using the calIntegrator2 User's Guide — This chapter introduces you to the *calIntegrator2 v.1.0 User's Guide* and suggests ways you can maximize its use.

Chapter 1 Getting Started in calIntegrator2 — This chapter introduces general calIntegrator2 procedures and how to obtain help to use calIntegrator2. .

Chapter 2 Creating a Study — This chapter describes the processes for creating and managing studies in calIntegrator2.

Chapter 3 Searching a calIntegrator2 Study — This chapter describes the processes for searching studies within calIntegrator2 using the search and browse tools.

Chapter 4 Viewing Search Results — This chapter describes search results that calIntegrator2 returns after queries.

Chapter 5 Analyzing Studies — This chapter describes how to use calIntegrator2 tools to analyze data in clinical or genomic studies that have been deployed in calIntegrator2.

Chapter 6 Administering User Accounts —This chapter describes the process for creating and managing user accounts in caIntegrator2.

Appendix A Exporting Data — This appendix describes how MAGE-TAB documents are parsed, validated and imported into caIntegrator2. It also provides examples of the types of MAGE-TAB documents that are expected by caIntegrator2.

Index—This section of the guide provides a complete index.

User's Guide Text Conventions

Table 2.1 illustrates how text conventions are represented in this guide. The various typefaces differentiate between regular text and menu commands, keyboard keys, toolbar buttons, dialog box options and text that you type.


| Convention | Description | Example |
|--|--|---|
| Bold & Capitalized Command Capitalized command > Capitalized command | Indicates a Menu command Indicates Sequential Menu commands | Admin > Refresh |
| TEXT IN SMALL CAPS | Keyboard key that you press | Press ENTER |
| TEXT IN SMALL CAPS + TEXT IN SMALL CAPS | Keyboard keys that you press simultaneously | Press SHIFT + CTRL and then release both. |
| Monospace type | Used for filenames, directory names, commands, file listings, and anything that would appear in a Java program, such as methods, variables, and classes. | URL_definition ::= url_string |
| Icon | A toolbar button that you click | Click the Paste button () to paste the copied text. |
| Boldface type | Options that you select in dialog boxes or drop-down menus. Buttons or icons that you click. | In the Open dialog box, select the file and click the Open button. |
| <i>Italics</i> | Used to reference other documents, sections, figures, and tables. | <i>caCORE Software Development Kit 1.0 Programmer's Guide</i> |
| <i>Italic boldface monospace type</i> | Text that you type | In the New Subset text box, enter <i>Proprietary Proteins.</i> |
| Note: | Highlights a concept of particular interest | Note: This concept is used throughout the installation manual. |

Table 2.1 caIntegrator2 User's Guide Text Conventions

| Convention | Description | Example |
|-------------------|---|--|
| Warning! | Highlights information of which you should be particularly aware. | Warning! Deleting an object will permanently delete it from the database. |
| { } | Curly brackets are used for replaceable items. | Replace {root directory} with its proper value, such as c:\cabio |

Table 2.1 caIntegrator2 User's Guide Text Conventions (Continued)

CHAPTER

1

GETTING STARTED WITH CAINTEGRATOR2

This chapter introduces general calIntegrator2 procedures and how to obtain help to use calIntegrator2.

Topics in this chapter include:

- [About calIntegrator2](#) on this page
- [Registering as a New calIntegrator2 User](#) on page 6
- [Logging In](#) on page 8
- [Using the calIntegrator2 Workspace](#) on page 8
- [Using Online Help](#) on page 10
- [Logging Out](#) on page 10
- [Application Support](#) on page 11

About calIntegrator2

NCI, Center for Biomedical informatics and Information Technology (CBIIT) is developing a novel translational informatics platform called calIntegrator that allows researchers and bioinformaticians to access and analyze clinical and experimental data across multiple clinical trials and studies. The calIntegrator framework provides a mechanism for integrating and aggregating biomedical research data and provides access to a variety of data types (e.g. Immunohistochemistry (IHC), microarray-based gene expression, SNPs, clinical trials data, etc.) in a cohesive fashion.

calIntegrator2 is a web based or locally installed portal that allows researchers and study managers to access the biomedical informatics infrastructure and data analysis tools established by calIntegrator from one common software platform. As a calIntegrator2 user, you can perform the following tasks:

- Integrate clinical data with genomic and/or imaging data
- Import data of various types in a predefined flat format, and create new studies with multiple study data
- Update an existing study to add new attributes or to add/modify data
- Perform analyses on study data.

Registering as a New caIntegrator2 User

To request a caIntegrator2 user account, you must register as a new user, completing the following steps:

1. Go to the CBIIT caIntegrator2 login page <http://caintegrator2.nci.nih.gov> or use the URL provided by your System Administrator for the caIntegrator2 instance at your institution.
2. Click the **Register Now** hypertext link, under the caIntegrator2 login section in the upper left of the page. This opens the account registration form (*Figure 1.1*).

Register

Figure 1.1 New user account registration form

3. In the Register form, enter the appropriate information¹.
 - **Security Information**
 - **Do you have an LDAP account** [a user profile with your institution] at [NCICB or your institution]?

If **Yes**, enter your username and case-sensitive password for the purposes of verifying that it is correct. After you submit your request, you can continue to use caIntegrator2 without an account to browse and

1. Items with an asterisk or highlight are required.

search available experiments and download data while your account is verified and activated.

–**Username***

–**LDAP Password***

–**Requested role(s)*** – Select one or more of the roles. Roles are described in [Table 1.1](#).

If your LDAP profile is not validated, caIntegrator2 indicates that the LDAP credentials do not check out. You are asked to reenter them, but you can choose to answer no, and the System Administrator will manually ensure you don't get a duplicate LDAP account during provisioning. You can **Cancel** or talk with your System Administrator about the problem.

If you select **No** [you do not have an LDAP account], the text boxes for entering the LDAP account information disappear. You must indicate the role you would like to be assigned in caIntegrator2, and continue entering the appropriate information in the **Account Details** section.

| Role | Description | Permissible 1.0 Actions |
|---------------------------|---|--|
| Study Manager | Creates, owns and manages studies | Create studies Assign annotations to studies Edit studies Search studies Perform analyses on study data |
| Study Investigator | Investigates and queries the study data | Query study data Save queries Analyze using K-M Plot Analyze using Gene Expression Plots Analyze using GenePattern |
| Platform Manager | Uploads and maintains the array platform annotation | Uploads array platform annotation files Ensures that if study requires a certain platform, then it is available within the installation |

Table 1.1 caIntegrator2 role descriptions

◦ **Account Details**

— **First Name***

— **Last Name***

— **Email [address]***

— **Organization***

— **Address [Lines 1* and 2]**

- City*
- State*
- Country*
- Postal [or Zip] Code*
- Phone*
- Fax

4. Click **Submit Registration Request** to execute the request, or click **Cancel** to abort the registration.

After registration is sent, the screen displays a confirmation message.

At this point, an email containing all of the information you specified in the new user request form is sent to the caIntegrator2 system administrator and an account request confirmation email is also sent to you, the prospective user, at your specified email address. In response, the caIntegrator2 system administrator uses UPT to create your user account and assign the requested roles (in predefined groups like Study Investigator). When your account is created, the system administrator sends you an email to alert you, after which you can login to caIntegrator2.

When your account is registered, the UserID and password you are assigned determines your access rights for the software.

Logging In

To log into caIntegrator2, follow these steps:

1. On the login page, enter your **username** and **password**.
2. Click the **Login** button. If your login is successful, the Welcome to Browse/ Study page appears (*Figure 1.2*).

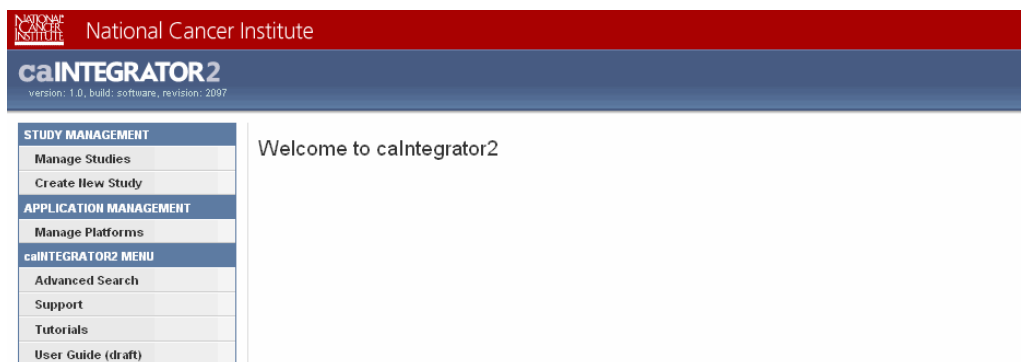


Figure 1.2 The caIntegrator2 workspace before any studies have been deployed

Using the caIntegrator2 Workspace

The caIntegrator2 workspace enables quick access to all caIntegrator2 functions and information. To access caIntegrator2 functions, use the options listed on the left sidebar of the workspace.

caIntegrator2 Functions

When you log into caIntegrator2, before any studies have been created the workspace opens with a Welcome page, as shown in ([Figure 1.2](#)). Once a study is created, its name is listed at the top of the left sidebar.

[Table 1.2](#) describes each caIntegrator2 option in the workspace ([Figure 1.2](#)).

| Sidebar Option | Function |
|-------------------------------|--|
| [Study Name] | When you log in, one study displays in the left sidebar by default. Any study that you select in the My Studies drop-down list in the upper right of the page replaces this default selection. |
| Home | Click this to return to the home page for the selected study. |
| Search [study name] | Click this option to open the Search [study name] page from which you can launch queries into your selected study. For more information, see Searching a caIntegrator2 Study . |
| Study Data | Click Queries > My Queries to open the list of previous queries you saved. Click any item in the list to open the saved query, which displays on the Criteria, Columns and Sorting tabs. From those tabs, you can modify criteria and/or launch the query again. For more information, see Saving a Query on page 43. |
| Analysis Tools | Click either of the listed options to open a page where you can launch an analysis of the data in the selected study. <ul style="list-style-type: none"> • Generate a K-M Plot. See Creating Kaplan-Meier Plots on page 54. • Generate a Gene Expression Plots. See Creating Gene Expression Plots on page 60. • Launch GenePattern Analysis. Analyzing Data with GenePattern on page 73. |
| Study Management | Click either of the listed options to manage the selected study through editing or deleting it or by creating a new study. <ul style="list-style-type: none"> • Click Manage Studies. See Managing a Study on page 31. • Click Create a New Study. See Configuring and Deploying a Study on page 14. |
| Application Management | Click Manage Platforms to identify, add or remove platforms that caIntegrator2 supports . For more information, see Managing Platforms on page 32. |
| caIntegrator2 Menu | <ul style="list-style-type: none"> • Click Support to view contact information for Application Support. • Click Tutorials to view a tutorial to help you get started using caIntegrator2. • Click User Guide to open the caIntegrator2 v.1.0 User's Guide in PDF format. |

Table 1.2 caIntegrator2 tabs

In the **My Studies** drop-down list in the upper right of the page, select the study you want to use for your current session. (The list includes all studies to which you are subscribed.) As you do so, the following left sidebar contents change to reflect options relevant to your study selection:

- the logo for the selected study (if it exists)

- the name for the selected study
- the list of saved queries for that study

Using Online Help

The online help explains how to use all of the features.

To access online help, click the help icon at the top of each page to open a context-sensitive topic. Context-sensitive help displays information that corresponds to the page from which help was opened.

Help opens displaying the table of contents in the left panel.

Once you are in online help, several buttons and/or options help you locate topics of interest.


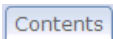
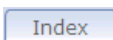
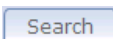
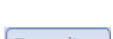



| <i>Icon or Button</i> | <i>Description</i> |
|---|--|
|  | Locates and highlights your current topic in the table of contents. |
|  | Select a topic from the complete online help table of contents. |
|  | Select a topic from the online help index. |
|  | Perform word searches of Help by entering query text in the search text box. |
|  | Create a list of your frequently-accessed topics. |
|  Related Topics  | Opens other closely related topics. |
|  | Prints the current topic. |
| Topic Name > Topic Name | The breadcrumb trail shows the relative location of the current help topic relative to neighboring topics. Click a breadcrumb link to display that help topic. |
| Back Forward | Navigates through previously viewed topics. |

Table 1.3 Online help tips

Logging Out

To log out of calIntegrator2, click the **logout** link in the upper right-hand corner of the page.

Application Support

For any general information about the application, application support or to report a bug, contact NCICB Application Support.

| | |
|---|--|
| Email: ncicb@pop.nci.nih.gov | When submitting support requests via email, please include: <ul style="list-style-type: none">• Your contact information, including your telephone number.• The name of the application/tool you are using• The URL if it is a Web-based application• A description of the problem and steps to recreate it.• The text of any error messages you have received |
| Application Support URL | http://ncicb.nci.nih.gov/NCICB/support |
| Telephone: 301-451-4384 Toll free: 888-478-4423 | Telephone support is available: Monday to Friday, 8 am – 8 pm Eastern Time, excluding government holidays. |

CHAPTER 2

CREATING A NEW STUDY

This chapter describes the processes for creating and managing studies in calIntegrator2.

Topics in this chapter include:

- [*Creating a Study – Overview*](#) on this page
- [*Configuring and Deploying a Study*](#) on page 14
- [*Managing a Study*](#) on page 31

Creating a Study – Overview

You can create a calIntegrator2 study by importing clinical study data, genomics data and imaging data, using a combination of spreadsheet/files and existing caGrid applications as source data. Each instance of calIntegrator2 can support multiple studies. As the manager creating a study, it is important that you understand the study well and that the data you wish to aggregate has been submitted to the applications whose data can be integrated in calIntegrator2.

- **Clinical** – The clinical data should be available in CSV files, with a unique patient identifier in one column, one patient per row. Other relevant data can be supplied in other columns to be identified as annotations in the file from within calIntegrator2. You, as the study creator, must have access to the clinical data file, as the file does not come from a caBIG[®] repository.
- **Genomic** – To use calIntegrator2 to integrate array data, the data should be imported into caArray, either locally or the CBIIT installation, using that system's data file import functionality. You must also have a mapping file in CSV format. This file indicates correlations between array files and the clinical subjects in the clinical data files. A mapping file consists of two columns: one with the patient ID, and one with the sample ID.

- **Imaging** – Imaging data should have been submitted to the NBIA grid node as public data, either locally or as part of the CBIIT installation. Image annotations, which includes information about images provided by radiologists or other researchers can include such information as tumor size, tumor location, etc. It must be in CSV format, with unique image series IDs in one column and annotation IDs in the second column. You must also have an image mapping file in CSV format. This file indicates correlations between clinical subjects or images in NBIA and clinical subjects in the clinical data files. A mapping file consists of two columns: one with the patient ID, and one with the NBIA image series ID in the other column.

As you create the study, you define its structure in the process, identifying the data sources and mapping the data between different source data. After the study has been created and deployed, the study can then be used to perform analyses.

Configuring and Deploying a Study

Note: Only a user with a Study Manager role can create a study.

When you create a study, you must specify different data-types (clinical, array, image, etc), data sources (caGrid applications – caArray and NBIA) and map the data, (patient to sample, image series, etc.).

To create a new study, follow these steps:

1. In the Study Management section of the left sidebar, click **Create New Study**.
2. In the Create New Study dialog box that opens, provide a name and description for the study you are creating ([Figure 2.1](#)).

Create New Study

Figure 2.1 Create Study page

3. Click **Save**.

This opens an Edit Study page where you can add identify data files for your study.

Creating/Editing a Study

The Edit Study page displays the Name and Description that you entered for a new study, or for an existing study that you are editing ([Figure 2.2](#)).

Figure 2.2 Edit Study page

To continue creating a study or to modify a study, on the Edit Study page complete these steps:

1. Change the name and or description, if you so choose. Click **Save**.
Note: You can save the study at any point in the process of creating it. You can resume the definition and deployment process later.
2. If you choose to add a logo for the study, click the **Browse** button corresponding to **Logo File**. Navigate for the file, then click **Upload Now**. Once you save the study (or its edit), the logo displays in the center of the page ([Figure 2.3](#)). On the home page for the study, the logo displays in the upper left, above the sidebar.



Figure 2.3 Example of a logo added to the caIntegrator2 browser on the Edit Study page

To continue, you can add clinical data sources, genomic data or imaging data.

Adding Clinical Data

The Edit Study page opens after you save a new study or click to edit an existing study.

Note: To edit information for an existing study, follow the same basic directions in this section. Instead of entering new information, you will modify existing information.

To add or edit clinical metadata in this page, follow these steps:

1. On the Edit Study page, click the **Browse** button in the Clinical Data Sources section. Navigate to locate the file. Files must be in CSV file format.

In the Clinical Data Sources section, if a file has already been selected, its information displays in the varying fields.

2. Click **Add Clinical Data Source**. This opens the Define Fields for Clinical Data page ([Figure 2.4](#)).

Define Fields for Clinical Data

Assign annotation definitions to data fields and click **Done**.

| Field Definition | Field Header from File | Data from File |
|---|------------------------|--------------------|
| Identifier Change Assignment | PATIENT_ID | 151 |
| DC_STUDY_ID Change Assignment | DC_STUDY_ID | B-NCI_U133A_1L.CHP |
| MICROARRAY Change Assignment | MICROARRAY | NCI_U133A_1L |
| SITE Change Assignment | SITE | MSKCC |
| IN_DC_STUDY Change Assignment | IN_DC_STUDY | 1 |
| GENDER Change Assignment | GENDER | Male |
| AGE_AT_DIAGNOSIS Change Assignment | AGE_AT_DIAGNOSIS | 64 |
| RACE Change Assignment | RACE | White(01) |
| ADJUVANT_CHEMO Change Assignment | ADJUVANT_CHEMO | Yes |
| ADJUVANT_HORMONE Change Assignment | ADJUVANT_HORMONE | No |

Figure 2.4 Define Fields for Clinical Data page

The Field Header from File column on the Define Fields... page displays column headers taken from the source *CSV file. The page also displays data values in the file you have designated. You must map each column name to an existing

column name in the caIntegrator2 database or in caDSR. If it doesn't yet exist, you can create a custom column name ([Figure 2.5](#)).

| | A | B | C | D | E | F | G | H |
|----|--------|-------|--------|----------|-------------|-------|-----------|---|
| 1 | Pa | Age | Gender | Survival | Disease | Grade | Race | |
| 2 | ASP221 | 50-54 | M | | ASTROCYTOMA | | WHITE | |
| 3 | ASP308 | 50-54 | M | | GBM | | WHITE | |
| 4 | FPH113 | 20-24 | M | | UNKNOWN | | WHITE | |
| 5 | FPH114 | 40-44 | M | | UNKNOWN | | WHITE | |
| 6 | FPH118 | 55-59 | M | | GBM | | WHITE | |
| 7 | FPH309 | 50-54 | M | | GBM | | WHITE | |
| 8 | E09238 | 45-49 | M | 18-24M | GBM | | WHITE | |
| 9 | E09239 | 25-29 | M | | UNKNOWN | | WHITE | |
| 10 | E09262 | 35-39 | M | | ASTROCYTOMA | | WHITE | |
| 11 | E09278 | 30-34 | M | | UNKNOWN | | WHITE | |
| 12 | E09331 | 35-39 | M | | UNKNOWN | | ASIAN NOS | |
| 13 | E09332 | 55-59 | M | | GBM | | WHITE | |
| 14 | E09336 | 30-34 | M | | GBM | | WHITE | |
| 15 | E09348 | 60-64 | M | | GBM | | WHITE | |
| 16 | E09378 | 45-49 | M | | UNKNOWN | | WHITE | |
| 17 | E09449 | 50-54 | M | | UNKNOWN | | OTHER | |
| 18 | E09454 | 0-4 | M | | UNKNOWN | | WHITE | |
| 19 | E09489 | 55-59 | M | | GBM | | WHITE | |
| 20 | E09515 | 35-39 | M | | UNKNOWN | | WHITE | |
| 21 | E09569 | 45-49 | M | | UNKNOWN | | WHITE | |
| 22 | E09587 | 35-39 | M | | UNKNOWN | | OTHER | |
| 23 | E09601 | 40-44 | M | | GBM | | WHITE | |
| 24 | E09610 | 55-59 | M | | GBM | | WHITE | |
| 25 | E09611 | 60-64 | M | | UNKNOWN | | ASIAN NOS | |
| 26 | E09615 | 45-49 | M | | UNKNOWN | | WHITE | |
| 27 | E09624 | 35-39 | M | | GBM | | WHITE | |
| 28 | E09645 | 45-49 | M | | UNKNOWN | | WHITE | |
| 29 | E09657 | 50-54 | M | | UNKNOWN | | WHITE | |
| 30 | E09730 | 40-44 | M | | UNKNOWN | | WHITE | |

Figure 2.5 Example of a source CSV file whose data you are mapping in caIntegrator2

The MOST important steps in creating a new study in caIntegrator2:

- You MUST designate one column in the file as a unique “identifier” column type.
- You MUST review and define column annotation definitions for each column header in the file.

If caIntegrator2 “recognizes” the same column header in other files already in the system, a term, for example “age” or “survival”, which is the current definition appears in the **Field Definition** column above the blue **Change Assignment** link. If the area above the blue **Change Assignment** link is blank, no correlating term exists in the database; you must specify the field type, and then the term will populate the space.

3. To indicate the unique identifier of choice, on the row showing the column header (PatientID in the figure, but other examples are subject identifier, sample identifier, etc), click **Change Assignment** in the **Field Definition** column.

Assigning An Identifier or Annotation

When you click **Change Assignment** on the Define Fields... page, the Assign Annotation Definition for Column dialog box opens ([Figure 2.6](#)). On this page you can change the column type and the field definition for the specific data field you selected.

Note: When you change an assignment, you must make sure the data types match--numeric, etc.

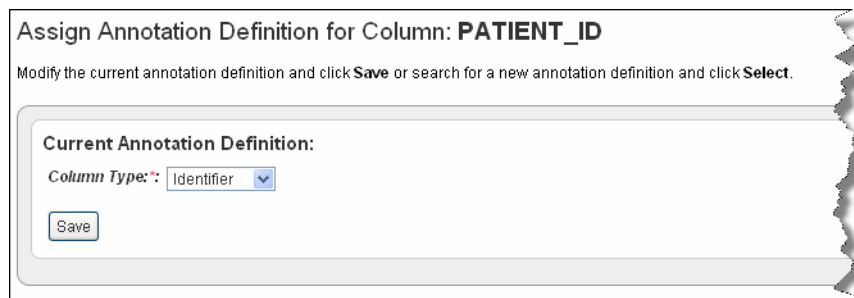


Figure 2.6 The Assign Annotation Definition dialog box

1. For the column (PatientID) that you choose to be the one and only Identifier column, in the **Column Type** drop-down list, select **Identifier**.
2. Click **Save** to save the identifier. This returns you to the Define Fields for Clinical Data page where the Identifier is noted in the Field Definition column.
3. After you have defined which field is the Identifier, you must ensure that ALL other fields also have a field definition assignment. For those fields without a Field Definition assignment or for those whose Field Definition you want to review, click **Change Assignment**.
4. In the Assign Annotation Definition for Column: [column header] dialog box, select **Annotation** in the drop-down list. The page extends, displaying additional fields for fleshing out the annotation description.

As you select the column type, you can work with column headers in one of four ways in this dialog box.

- You can accept existing default definitions (those that are inherent in the data file you selected). See Step 5.
 - You can create your own definitions manually. See Step 6.
 - You can search for and use definitions in other caIntegrator2 studies. See [Searching for Annotation Definitions](#) on page 20.
 - You can search for and use definitions found in caDSR. See [Searching for Annotation Definitions](#) on page 20.
5. If there is anything you want to change about an existing annotation definition of the field such as its name, or if you want to view or edit its definition, click the **Change Assignment** link on the Define Fields... page. The Assign Definition page opens, expanded now to include a Current Annotation Definition section above a section where you can still initiate a search for an annotation definition ([Figure 2.7](#)).

Note: If the column header you are working with already has a designated Field Definition, the Current Annotation Definition section of the Assign

Annotation Definition for Column page is already visible when you open this dialog box.

Assign Annotation Definition for Column: **MICROARRAY**

Modify the current annotation definition and click **Save** or search for a new annotation definition and click **Select**.

Current Annotation Definition:

Column Type: Annotation

Name: MICROARRAY

Definition: Created via selenium for DC Lung Full on 09/14/09 15:11:17.

Keywords: MICROARRAY

Data Type: string

Non-Permissible

Permissible

Permissible Values:

Add >

< Remove

New

Save

Search for an Annotation Definition:

Search

Search existing studies and caDSR for definitions.

Figure 2.7 Current Annotation Definition

- To enter a new name annotation, or any other information about the annotation definition, click the **New** button and enter the information described in [Table 2.1](#)

| Annotation Field | Field Description |
|-------------------|---|
| Name | Enter the name for the annotation. |
| Definition | Enter the term(s) that define the annotation. |
| Keywords | Insert keyword(s) that can be used to find the annotation in a search, separated by commas. |
| Data Type | Enter a string (default), numeric, or date |

Table 2.1 Annotation fields for new definitions

| Annotation Field | Field Description |
|---|--|
| Permissible/Non-permissible Values | <p>Note: The first time you load a file, before you assign annotation definitions (step 3 on page 17), these panels may be blank. If the column header for the data is already “recognizable” by calIntegrator2, the system makes a “guess” about the data type and assigns the values to the data type in the newly uploaded file. They will display in the Non-permissible values sections initially. Use the Add and Remove buttons to move the values shown from one list to the other, as appropriate.</p> <p>When you select or change annotation definitions by selecting matching definitions (described in Searching for Annotation Definitions on page 20), this may add (or change) the list of non-permissible values in this section.</p> <p>If you leave all values for a field in the Non-permissible panel, then when you do a study search, you can enter free text in the query criteria for this field.</p> <p>If there are items in the Permissible values list, then the values for this annotation are restricted to only those values. When you perform a study search, you will select from a list of these values when querying this field. If there are no items in the permissible values list then the field is considered free to contain any value.</p> <p>To edit a field's permissible values, you must change the annotation definition. You can do this even after a study has been deployed.</p> |

Table 2.1 Annotation fields for new definitions

Searching for Annotation Definitions

An alternative to creating a new definition is to search for annotation definitions already present in calIntegrator2 studies or in caDSR.

1. Enter search keyword(s) in the **Search** text box on the Assign Annotation Definition page. Click **Search**. After a few moments, the search results display on the page (*Figure 2.8*).

Search for an Annotation Definition:

microarray Search existing studies and caDSR for definitions.

| Matching Annotation Definitions from caIntegrator2 | | | | |
|--|------------------------|---------------|-----------|--|
| Name | Actions | CDE Public ID | Data Type | Definition |
| MICROARRAY | Select | | string | Created via selenium for DC Lung Full on C |
| MICROARRAY | Select | | string | Created via selenium for DC Lung Full on C |

| Matching Annotation Definitions from caDSR | | | | | |
|---|---|---------------|---------|----------|--|
| Name | Actions | CDE Public ID | Context | Status | Definition |
| Microarray Microarray Analysis Data float One Dimensional Array | Select View | 2658378 | caBIG | RELEASED | A microarray is a piece of glass or plastic on which different samples have been affix are usually DNA fragments but may also be antibodies, other proteins, or tissues. _Ana microarrays to profile the pattern of proteins). A collection or single item of factual infor drawn. _Generic value domain for a single dimensional array with floating numbers as i |
| Microarray Identifier java.lang.Long | Select View | 2223905 | caCORE | RELEASED | A microarray is a piece of glass or plastic on which different samples have been affix are usually DNA fragments but may also be antibodies, other proteins, or tissues. _One |

Figure 2.8 Results for annotation definition search

2. To view the definitions corresponding to any of the “Matching Annotation Definitions”, which are those currently found in other caIntegrator2 studies, click the [term], such as “age”, hypertext link. The definition then appears in the Current Annotation Definition segment of the page just above.

In summary, when you click the link, that assigns the definition to the Define Fields for Clinical Data page, and it also closes the Annotation Definition page.

You can modify any portion of the definition, as described in [step 6](#) on page 19.

3. The matches from caDSR display some of the details of the search results. To view more details of a match, such as permissible values, click **View**, which opens caDSR to the term. If you click **Select**, the caDSR definition automatically replaces the annotation definition for this field with which you are working.

Caution: Take care before you add a caDSR definition that it says exactly what you want. caDSR definitions can have minor nuances that require specific and limited applications of their use.

4. Once you have settled on an appropriate field definition for the annotation, click **Save**. This returns you to the Define Field for Clinical Data page.

Note: If you have not clicked **Select** for alternate definitions in this dialog box, then click **Save** to return to the Define Field...dialog box without making any definition changes.

5. From the Define Fields for Clinical Data page, be sure and designate the annotations for each field in the file. Click **Save** on each page to save your entries or click **New** to clear the fields and start again. You will not be able to proceed until every Field Definition entry on the Fields for Clinical Data screen has a unique entry, one as an Identifier and the remainder as annotations.

The Data From File columns on the page display the column header *values* of the first three rows you designated as “annotations”.

6. Click **Done**. This saves the study by name and description, but does not deploy the study. See [Deploying the Study](#) on page 31.

Saving the study returns you to the Edit Study page where a “Not Loaded” status now appears for the file whose annotations (column headers) you have defined ([Figure 2.9](#)).

Study Overview

Study Name: test jbh

Study Description:

Status: Not Deployed

Status Description:

Study Logo: Study Logo

Logo File: Browse...

JPEG/GIF, 200x72 maximum

Upload Now

Clinical Data Sources

Add New Edit Survival Values

| Type | Description | Status | Action |
|----------------|---------------------------|------------|--------|
| DELIMITED_TEXT | dc_lung_clinical_data.csv | Not Loaded | |

Figure 2.9 Example file whose annotations have been defined

7. Click the **Load Clinical** link in the Action section to load the data file you configured. At this point, the Status changes to “Loaded”.

Note: You can add as many files as are necessary for a study. Patients 1-20 in first file, 21-40 in second file, or many patients in first file and annotations in second file, etc. As long as IDs are defined correctly, it works.

8. Once you have assigned data types to every column header in the data file and have loaded the clinical data, click **Save and Deploy**. At that point, calIntegrator2 loads data from the file to the calIntegrator2 database.

Note: You can change assignments even after the study is deployed, using the Edit feature. For more information, see [Creating/Editing a Study](#) on page 15.

The Manage Studies page opens when the study is deployed. The **Deployed** status is indicated on the Manage Studies page as well as the Edit Study page. For more information, see [Managing a Study](#) on page 31.

You can continue to perform other tasks in calIntegrator2 while deployment is in process.

See also [Deploying the Study](#) on page 31.

Note: You can repeatedly upload additional or updated subject annotations, samples, image data, array data to the study at later intervals. These later imports do not remove any existing data; they instead insert any new subjects or update annotations for existing subjects.

Defining Survival Values

Survival value is the length of time a patient lived. If you plan to analyze your data in calIntegrator2 to create a Kaplan-Meier (K-M) Plot, then during the Annotation Definition process described above, you must make sure that you have defined at least three fields set to the “date” Data Type. These will be matched to the following three properties during Survival Value definition.

- **Survival Start Date**
- **Death Date**
- **Last Followup Date**

Note: Setting survival values is optional if you do not plan to use the K-M plot analysis feature or if you do not have this kind of data (survival values) in the file.

For some applications, such as REMBRANDT and I-SPY, survival values are pre-defined in the databases when you load the data. In calIntegrator2, however, you can review and define survival value ranges in a data set you are uploading to a study. To be able to do so, you need to understand the kind of data that can comprise the survival values.

To set up survival values, follow these steps:

1. On the Edit Study page, click **Edit Survival Values**. This opens the Survival Value Definitions dialog box ([Figure 2.10](#)).

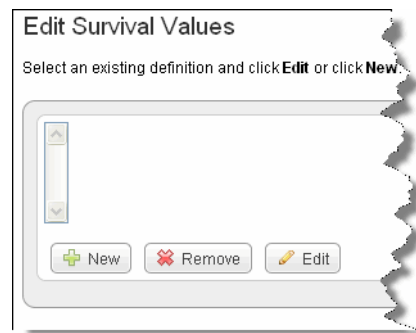


Figure 2.10 Survival Value Definition dialog box

2. Click **New** to enter new survival value definitions.

- OR -

Click **Edit** to edit existing survival value definitions.

- The dialog box extends, now displaying three drop-down lists that show column headers for date metadata in the spreadsheet you have uploaded. [Figure 2.11](#) displays survival value ranges that have already been added to a study.

Survival Value Definitions for 'Demo Study based on 101 Long-Full data'

Survival from enrollment

New

Remove

Edit

Survival Value Definition Properties for 'Survival from enrollment'

| | |
|----------------------|--------------------------|
| Name: | Survival from enrollment |
| Survival Start Date: | ENROLLMENT_DATE |
| Death Date: | DEATH_DATE |
| Last Followup Date: | LAST_CONTACT_DATE |

Save

Figure 2.11 Survival Definitions example

In the drop-down lists, select the appropriate survival value definitions for each field listed. You might want to refer to the column headers in the data file itself. Dates covered by the definitions are already in the data set. You cannot enter specific dates.

- **Name** – Enter a unique name that adequately describes the survival values you are defining here. *Example:* Survival from Enrollment Date or Survival from Treatment Start. The name you enter displays later when you are selecting survivals to create the K-M plot.
- **Survival Start Date** – Select the column header for this data.
- **Death Date** – Select the column header for this data.
- **Last Followup Date** – Select the column header for this data.

See also [Creating Kaplan-Meier Plots](#) on page 54.

Adding/Editing Genomic Data

Note: Genomic data must be parsed and stored in caArray to be able to use it in caIntegrator2.

Once you have loaded clinical data and identified patient IDs, you can add either array genomic sample data from caArray, which caIntegrator2 maps by sample IDs to the patient IDs in the clinical data, covered in this section, or you can load imaging files from NBIA, also mapped by IDs to the patient data, covered in [Adding Imaging Data](#) on page 29. You can also edit genomic data information that you have already added to the study. Genomic sample data and imaging data are independent of each other, so neither is required before loading the other.

It is essential that you are well acquainted with the data you are working with--the clinical data, and the corresponding array data in caArray.

caIntegrator2 supports a limited number of array platforms. For more information, see [Managing Platforms](#) on page 32.

To add genomic data to your caIntegrator2 study, follow these steps:

1. On the Edit Study page where you have selected and added the clinical data, click the **Add** button under Genomic Data Sources. You can upload genomic data only from caArray.

This opens the Edit Genomic Data Source dialog box (Enter the appropriate information in the fields (*Figure 2.12*).

Edit Genomic Data Source

| | |
|-------------------------------------|-------------------|
| caArray Server Hostname: | array.nci.nih.gov |
| caArray server JNDI Port: | 0 |
| caArrayUsername: | |
| caArrayPassword: | |
| caArray Experiment Id: | |
| Vendor: | Affymetrix |
| Data Type: | Expression |
| Platform (only needed for Agilent): | |

Cancel Save

Figure 2.12 Edit Genomic Source dialog box

- **caArray Host Name** – Enter the hostname for your local installation or for the CBIIT installation of caArray, array.nci.nih.gov. If you misspell it, you will receive an error message.
 - **caArray JNDI Port** – Enter the appropriate server port. See your administrator for more information. *Example:* For the CBIIT installation of caArray, enter **8080**.
 - **caArray Username** and **caArray Password** – If the data is private, you must enter your caArray account user name and password; you must have been given permissions in caArray for the experiment. If the data is public, you can leave these fields blank.
 - **caArray Experiment ID** – Enter the caArray Experiment ID which you know corresponds with the clinical data you uploaded. *Example:* Public experiment “beer-00196” on the CBIIT installation of caArray (array.nci.nih.gov). If you misspell your entry, you will receive an error message.
 - **Vendor** – Select either **Agilent** or **Affymetrix**
 - **Data Type** – Select **Expression** or **Copy Number**.
 - **Platform (needed only for Agilent)** – If appropriate, select the **Agilent** platform.
2. Click **Save**.

caIntegrator2 goes to caArray, validates the information you have entered here, finds the experiment and retrieves all the sample IDs in the experiment. Once

this finishes, the experiment information displays on the Edit Study page under the Genomic Data Sources section ([Figure 2.13](#)).

| Host Name | Experiment Identifier | File Description | Data Type | Status | Action |
|--------------------------|-----------------------|--|------------|--------|---|
| ncies-d227-v.nci.nih.gov | admin-00001 | Mapping File(s): nci_sample_mapping.csv Control Sample Mapping File(s): page_0034_control_samples.csv | Expression | Loaded | Edit Map Samples Delete |

Figure 2.13 Genomic Data Sources section of the Edit Study page

3. If you want to redefine the caArray experiment information, you can edit it. Click the **Edit** link corresponding to the Experiment ID. The Edit Genomic Data Source dialog box reopens, allowing you to edit the information.

Mapping Genomic Data to Clinical Data

Because the goal of caIntegrator2 is to integrate data from clinical, genomic and imaging data sources, data from uploaded source files must be mapped to each other.

To map the samples from the caArray experiment to the patients in the clinical data you uploaded, follow these steps:

1. On the Edit Study page, click the **Map Samples** link. This opens the Edit Sample Mappings page ([Figure 2.14](#)).

Edit Sample Mappings
Upload mapping files and click **Map Samples**.

Data Source

caArray Server Hostname: array-stage.nci.nih.gov
caArray server JNDI Port: 8080
caArrayUsername:
caArrayPassword:
caArray Experiment Id: rembr-00037
Subject to Sample Mapping File: [Browse...](#)
Control Sample Set Name:
Control Samples File: [Browse...](#)
[Map Samples](#)

Control Sample Sets

| Set Name | Sample Name |
|----------|-------------|
| | |

Sample Mappings

| Unmapped Samples |
|---|
| Sample Name |
| E09907B_U133P2 |
| GeneratedSample.ASTROCYTOMA_L_E07733_U133P2 |
| GeneratedSample.ASTROCYTOMA_L_E09137_U133P2 |
| GeneratedSample.ASTROCYTOMA_L_E09629_U133P2 |
| GeneratedSample.ASTROCYTOMA_L_E09920_U133P2_2 |
| GeneratedSample.ASTROCYTOMA_L_E09921_U133P2 |
| GeneratedSample.ASTROCYTOMA_L_E09948_U133P2 |
| GeneratedSample.ASTROCYTOMA_L_E09971_U133P2 |

Figure 2.14 Edit Sample Mappings page

When you first open this page, all of the samples in the caArray experiment you selected are listed as unmapped, because caIntegrator2 does not know how

these sample names correlate to the patient data in the clinical file until you upload the subject to sample mapping file.

2. At the top of the page, click **Browse** to navigate for the CSV file that identifies the mapping information. Click the **Upload Mapping File** button.

The mapping file has only two columns (typically without headers)—one that shows the subject ID (designated in calIntegrator2 as the “Identifier”) and one that has “Sample name” field from the linked caArray experiment, with one subject per row (*Figure 2.15*). This provides calIntegrator2 with the information for mapping patients to caArray samples.

| | A | B | C | D | E | F | G | H |
|---|--------|---|---|---|---|---|---|---|
| 1 | E10216 | GeneratedSample.UNKNOWN_DISEASE_L_E10216_U133P2 | | | | | | |
| 2 | E10144 | GeneratedSample.UNKNOWN_DISEASE_L_E10144_U133P2 | | | | | | |
| 3 | E09212 | GeneratedSample.UNKNOWN_L_20070227_16-22-37-238_E09212_U133P2 | | | | | | |
| 4 | E09369 | GeneratedSample.UNKNOWN_L_20070227_16-22-37-238_E09369_U133P2 | | | | | | |
| 5 | E10162 | GeneratedSample.UNKNOWN_DISEASE_L_E10162_U133P2 | | | | | | |
| 6 | E10318 | GeneratedSample.UNKNOWN_DISEASE_L_E10318_U133P2 | | | | | | |
| 7 | E09264 | GeneratedSample.OLIGO_L_20070227_11-27-27-881_E09264_U133P2 | | | | | | |
| 8 | E10252 | GeneratedSample.UNKNOWN_DISEASE_L_E10252B_U133P2 | | | | | | |

Figure 2.15 Example sample mapping file, in CSV format

Note: When you open the mapping file, make sure that the patient ID is used for mapping.

Unmapped samples continue to show at the top of the calIntegrator2 page. They were loaded from caArray, but they are not in the mapping file. These are not used for integration.

3. Scroll down the page to see samples that are mapped to the patients in the clinical data (*Figure 2.16*).

| 1336 | GeneratedSample.OLIGO_L_20070227_11-49-51-876_HF0599_U133P2 | |
|-----------------------------------|---|--------------------|
| 1338 | GeneratedSample.UNKNOWN_DISEASE_L_E10029_U133P2 | |
| 1339 | GeneratedSample.GBM_L_20070226_14-05-29-569_HF1356_U133P2 | |
| 1340 | GeneratedSample.OLIGODENDROGLIOMA_L_HF0599_U133P2 | |
| 1342 | GeneratedSample.GBM_L_20070226_13-30-40-39_HF0142_U133P2 | |
| 1345 | GeneratedSample.GBM_L_20070226_14-31-20-427_HF1400_U133P2 | |
| Samples Mapped to Subjects | | |
| Sample ID | Sample Name | Subject Identifier |
| 901 | GeneratedSample.UNKNOWN_DISEASE_L_E10216_U133P2 | E10216 |
| 911 | GeneratedSample.UNKNOWN_DISEASE_L_E10144_U133P2 | E10144 |
| 914 | GeneratedSample.UNKNOWN_L_20070227_16-22-37-238_E09212_U133P2 | E09212 |
| 918 | GeneratedSample.UNKNOWN_L_20070227_16-22-37-238_E09369_U133P2 | E09369 |
| 922 | GeneratedSample.UNKNOWN_DISEASE_L_E10162_U133P2 | E10162 |
| 925 | GeneratedSample.UNKNOWN_DISEASE_L_E10318_U133P2 | E10318 |
| 930 | GeneratedSample.OLIGO_L_20070227_11-27-27-881_E09264_U133P2 | E09264 |
| 940 | GeneratedSample.UNKNOWN_DISEASE_L_E10252B_U133P2 | E10252 |
| 954 | GeneratedSample.GBM_L_20070226_13-14-06-57_E09624_U133P2 | E09624 |
| 957 | GeneratedSample.ASTROCYTOMA_L_E09137_U133P2 | E09137 |
| 958 | GeneratedSample.UNKNOWN_DISEASE_L_E09890_U133P2 | E09890 |
| 968 | GeneratedSample.UNKNOWN_L_20070227_16-57-07-283_E09515_U133P2 | E09515 |
| 1004 | GeneratedSample.UNKNOWN_L_20070227_17-26-09-910_E09722_U133P2 | E09722 |

Figure 2.16 Example of samples mapped to patients' data

Uploading Control Samples

A Control Samples file is used to calculate fold change data, which compares “tumor” sample gene expression in the caArray experiment to the control samples to identify those that exhibit up or down gene regulation. Control samples can be the “normal” samples, but that is not necessarily the case.

To upload the control samples, follow these steps:

1. On the Edit Sample Mappings page, click the **Map Samples** link.
2. Click **Browse** to navigate for the control samples file, and click the **Upload Control Samples** File button. The control sets display at the top of the page once they have been uploaded ([Figure 2.17](#)).

Figure 2.17 Example list of control samples

The control samples now display toward the bottom of the page.

3. This information will be used when performing other tasks in caIntegrator2, to be described in other sections.

Configuring Copy Number Data

You can add copy number data for a genomic data source by uploading the mapping file. This allows you to configure parameters to be used when segmentation data is being configured.

The name specified in the third column of the mapping file is specific for each array manufacturer as follows:

- Affymetrix – The third column of the mapping file must contain filenames that end in cnchp. The corresponding experiment in caArray must have these files and the extensions must match .cnchp.
- Agilent – The third column must name a file which contains level 2 copy number data. Level one copy number will not work. This file name is repeated for each line in the mapping file.

To add copy number data relating to the genomic data you are adding, follow these steps:

1. In the Genomic Data Sources section, for the data you have already added, click **Configure Copy Number Data** hypertext link.

Note: This link is available only if you have uploaded copy number data and you are configuring a Copy Number data type (as indicated by the Data Type column on the Edit Study page).

The Edit Copy Number page opens ([Figure 2.18](#)).

Figure 2.18 Edit Copy Number page

2. Browse for and enter appropriate information to identify the copy number mapping file. The fields are described in [Table 2.2](#). An asterisk* indicates a required field..

| Field | Description |
|---|---|
| Subject and Sample to Copy Number Mapping File | Browse for the appropriate CN mapping file |
| caDNACopy Service URL* | Control for selecting the URL which hosts the caDNACopy grid service |
| Change Point Significance Level | Significance levels for the test to accept change-points |
| Early Stopping Criteria | The sequential boundary used to stop and declare a change |
| Permutation Replicates | The number of permutations used for p-value computation |
| Random Number Seed | The segmentation procedure uses a permutation reference distribution. This should be used if you plan to reproduce the results. |

Table 2.2 Fields for retrieving a copy number mapping file.

3. Click **Configure copy number data** for a genomic data source. On the screen upload a copy number mapping file (format: subject id, sample id, file name) and configure the parameters to be sent when computing segmentation data.

Adding Imaging Data

Once you have loaded clinical data and identified patient IDs, you can add either array genomic sample data from caArray which caIntegrator2 maps by sample IDs to the patient IDs in the clinical data, or you can load imaging files from NBIA, also mapped by IDs to the patient data, covered in this section. Genomic sample data and imaging data are independent of each other, so neither is required before loading the other.

It is essential that you are well acquainted with the data you are working with--the clinical data, and the corresponding imaging data in NBIA.

Any data in NBIA can be uploaded to caIntegrator2. Imaging data consist of images and or annotations for images.

To add imaging data to the study you are creating or are editing, follow these steps:

1. On the Edit Study page, click the **Add** button under Imaging Data Sources section. Imaging data can be NBIA images or image annotations, which are uploaded in spreadsheet format.

This opens the Edit Imaging Data Source dialog box. Enter the appropriate information in the fields (*Figure 2.19*). Asterisks indicate required fields..

Edit Imaging Data Source

Enter a NBIA Data Source and Image Mapping Data from a file and click **Save**.

Figure 2.19 Edit Image Data Source dialog box

- **NBIA Server Grid URL*** – Enter the URL for the grid connection to NBIA
 - **NBIA Username and NBIA Password.** This information is not required, as currently all data in the NBIA grid is Public data.
 - **Collection Name*** – Enter the name/source for the collection.
 - **Current Mapping** – If a mapping file has already been uploaded to the study to map imaging data, the file name displays here.
 - **Select Mapping File Type*** – Click to select the file type:
 - **Auto** – No file is required. Selecting this takes all clinical subject IDs and attempts to map them to the corresponding ID in the collection in NBIA. If the ID does not exist in NBIA, then no mapping is made for that ID.
 - **By Subject** – Requires a file to be uploaded. The “clinical to imaging mapping file” must be a two column mapping (CSV) from the caIntegrator2 clinical subject ID to the NBIA subject ID.
 - **By Image Series** – Requires a file to be uploaded. The clinical to imaging mapping file needs to be a two column mapping (CSV) from the caIntegrator2 clinical subject ID to the NBIA study instance UID.
 - **Clinical to Imaging Mapping File** – Click **Browse** to navigate to the appropriate clinical to imaging mapping file. See **Select Mapping File Type*** field description.
2. Click **Add** to upload the data to caIntegrator2.

The imaging data information displays on the Edit Study page under the Imaging Data Sources section ([Figure 2.20](#)).

| Host Name | Collection Name | File Description | Status | Action |
|---------------------|-----------------|---|--------|--|
| imaging.nci.nih.gov | NCRI | Annotation File: nci_image_annotations.csv Mapping File: nci_image_mapping.csv | Loaded | Edit Edit Annotations Delete |

Figure 2.20 Imaging Data Sources section of the Edit Study page

- Once the data is uploaded, you must assign identifiers and annotations to the data in the same way you did with the clinical data. For more information, see [Assigning An Identifier or Annotation](#) on page 17 and [Searching for Annotation Definitions](#) on page 20.
- To deploy the study, see [Deploying the Study](#).

Deploying the Study

When you are ready to deploy the study, click the **Deploy Study** button on the Edit Study page. caIntegrator2 retrieves the selected data from the data service(s) you defined and makes the study available to a study manager or to anyone else who may want to analyze the study's data. Using the Manage Studies feature, you can then configure and share data queries and data lists with all investigators who access the study.

Note that you can continue to work in caIntegrator2 while study is being deployed.

Managing a Study

Note: A user without management privileges has no access to this section of caIntegrator2.

Once you have started to create a study or have deployed it, you can update an existing study in the following ways:

- Add new attributes (annotations) and upload relevant data to an existing study.
- Delete a study
- Modify existing annotation definitions
- Reload subset of study data and re-deploy the study and perform new analyses
- Re-deploy the entire study with new set of data and mappings.

To update, edit or delete a study, follow these steps:

1. On the left sidebar, click **Manage Studies**. The Manage Studies page appears (*Figure 2.21*).

Manage Studies

View studies and click **Edit** to modify or click **Delete**.

| Name | Description | Status | Deployment Start Date | Deployment Finish Date | Action |
|--|---|--------------|-----------------------|------------------------|---|
| Demo Study based on DC Lung Full data. | DC Lung Full. Study created via selenium. | Deployed | 2009/09/14 15:18:32 | 2009/09/14 15:41:10 | Edit Delete |
| Demo Study based on Rembrandt with NCRI data. | Rembrandt with NCRI. Study created via selenium. | Deployed | 2009/09/14 14:45:58 | 2009/09/14 14:57:55 | Edit Delete |
| Demo Study based on Rembrandt with no images data. | Rembrandt with no images. Study created via selenium. | Deployed | 2009/09/14 15:54:55 | 2009/09/14 16:40:56 | Edit Delete |
| Demo Study based on Small Copy Number data. | Small Copy Number. Study created via selenium. | Not Deployed | | | Edit Delete |
| Demo Study based on TCGA Agilent data | TCGA Agilent. Study created via selenium. | Processing | 2009/09/14 16:28:57 | | |
| NCRI by TJ | | Not Deployed | | | Edit Delete |

Figure 2.21 Manage Studies page

All of the “in process” or “completed” studies display on this page.

2. Click the **Edit** link corresponding to your study of choice to open the Edit Studies page.

On this page you can edit any details such as adding or deleting files, survival values, and so forth. For information about working in the Edit Study, see *Creating/Editing a Study* on page 15.

3. Click the **Delete** link to delete the corresponding study.

Managing Platforms

caIntegrator2 supports a limited number of array platforms, all of which originate from Agilent or Affymetrix. While they do not represent all of the platforms supported by caArray, caIntegrator2 must have array definitions loaded for the platforms it supports, and be able to properly load the data from caArray and parse it.

You can create a study without genomic data, but you cannot add genomic data to a caIntegrator2 study without a corresponding supported array platform.

On the Manage Platforms page, you can identify, add or remove supported platforms.

To manage platforms in caIntegrator2, follow these steps:

1. Click **Manage Platforms** on the left sidebar.

The Manage Platforms page that opens lists the platforms caIntegrator2 currently supports, those that the system can pull from caArray (*Figure 2.22*).

You can also add a new platform by entering information in the fields at the top of the page.

Manage Platforms (draft)

Platform Type: Affymetrix Gene Expression ▼

Platform Name (For NON-GEML.xml file):

Annotation File: [Browse...](#)

[Add Annotation File](#)

Annotation File(s) Selected:

[Create Platform](#)

| Name | Vendor | Reporter List | Action |
|---------------------------------|------------|--|------------------------|
| AgilentG4502A_07_01 | AGILENT | AgilentG4502A_07_01, AgilentG4502A_07_01 | Delete |
| GeneChip Human Mapping 100K Set | AFFYMETRIX | Mapping50K_Hind240, Mapping50K_Xba240 | None |
| HG-U133A | AFFYMETRIX | HG-U133A, HG-U133A | Delete |
| HG-U133A | AFFYMETRIX | HG-U133A, HG-U133A | None |

Figure 2.22 Manage Platforms page

- To add a platform, in the Platform Type field, select the appropriate platform type from the drop down list. Click **Browse** to navigate for the Affymetrix or Agilent file you want to add.
- Enter a **Platform Name** if the file is a NON-GEML.xml file.
Depending on what Platform Type is selected, there may be other parameters to provide here as well. Once all parameters have been provided, click **Create Platform**.
- Click the **Browse** button to browse for the appropriate annotation file. When you have located it, click **Add Annotation**. The system displays annotation files you select in the Associated File(s) Selected box.
- Click the **Add** button.

CHAPTER 3

SEARCHING A CAINTEGRATOR2 STUDY

This chapter describes the processes for searching studies within calIntegrator2.

Topics in this chapter include:

- [Search Overview](#) on this page
- [Searching a Study](#) on page 36
- [Managing Queries](#) on page 43

Search Overview

The search and browse functions in calIntegrator2 allow you to search for clinical data, genomic or imaging data that were uploaded into the application as part of a study. When gene expression and imaging data are uploaded into a calIntegrator2 study, mapping files that correlate the data in those files to patient IDs in the clinical data file must also be uploaded. When you launch a search, calIntegrator2 finds and integrates the clinical, genomic and imaging data based on the mapping files and the criteria that you define in the search query.

In a search query, you can specify criteria for just one of the data types, or configure complex search criteria that join two or three data types. The available criteria for the query were defined when the study was deployed.

The basic workflow for a study search follows these steps:

1. Select the study to be searched.
2. Select one data type:
 - **Clinical** – searches one or more uploaded CSV files for data identifiers or annotations (column headers) specified when the study was created
 - **Genomic** – Searches caArray experiments samples uploaded in the study for gene expression data by gene name or reporter ID.

- **Image Series** – Searches NBIA files uploaded in the study for image annotations or links to images, identified by subject identifiers or image series IDs.
3. Define criteria for the search in the selected data type and run the search.
 4. For a more complex search, select multiple criteria from more than one data type.
 5. Specify whether you want clinical/imaging annotations to display or genomic data to display.
 6. Review search results.
 7. Configure results column and sorting display settings. You can do this before or after you run a search. If you choose to do it after, you must re-run the search.
 8. Download annotation search results as a CSV file. The CSV file contains only the data you specified in the annotation and display configurations.
 9. Follows links to NBIA in the search results to view or download images located in the search.

Searching a Study

To initiate a search of all annotations and/or other data in a study, follow these steps:

1. In calIntegrator2, in the upper right hand corner, select the study you want to browse or perform a simple search.
2. On the left sidebar, under the first section that displays the study name, click **Search [Study Name]**. This opens a simple search query page with five tabs (*Figure 3.1*).

Search Demo Study based on Rembrandt with NCRI data.

The screenshot shows the 'Criteria' tab of the search interface. It includes a dropdown menu for selecting criteria (currently set to 'Clinical'), an 'Add' button, and a message indicating no criteria have been added yet. At the bottom, there are options to combine results using 'or' or 'and' logic, and a 'Run Query' button.

Figure 3.1 Search page

3. On the Criteria tab, in the drop-down list, select the type of data you want to search. The listed options reflect the type of data that have been uploaded to the study.

Note: You can perform a search using one or more criteria you set in one of the data types, or you can define criteria in more than one data type per query, creating a more complex search.

- **Clinical**
- **Gene Expression**
- **Image Series**

Note: NBIA submissions are organized in the following hierarchy, which illustrates the relationship of an image series to its parent study and patient, as well as to the images in the series.:

Clinical trial > Patient (Subject) > Study > Series > Images

4. Click the **Add** button to define annotation elements for the search.

Continue with:

[Clinical and Image Series](#) on page 37

[Gene Expression](#) on page 38

5. To add additional criteria for the search, repeat steps 3 and 4, as appropriate. You can set more than one data type or more than one criterion for a data type. The criteria become cumulative, thus refining the search.
 6. Once you have configured the query criteria, select the Boolean **Or** or **And** search operator at the bottom of the page.
 - **Or** finds a data subset with at least one of the search criteria
 - **And** finds a data subset with both/or all search criteria.
 7. Click the **Remove** button to clear any data elements you have defined.
 8. You can launch the search from this tab. Click the **Run Search** button. For information about the search results, see [Chapter 4 Viewing Query Results](#). You may want to run the search first to see what kind of results you get before you configure the data display, described in step 9.
- or –**
9. On the Results Type tab, you can specify the columns you want to display in the search results data. On the Sorting tab, you can specify how the data is to be sorted. For more information, see [Results Type Tab](#) on page 40.

Note: As long as you are still in the current query session, you can return to the Criteria, Columns and Sorting tabs to add, modify or remove data and display criteria and re-run the search. If you configure another query without saving the first, the first query will be lost. If you save the query, your current search criteria are saved.

Clinical and Image Series

- If you select Clinical or Image Series data types, an additional drop-down list displays data elements that are annotation definitions specified when

the data was uploaded into the study ([Figure 3.2](#)). Select a search criterion from among the options. You can make only one selection at a time.

Figure 3.2 Additional clinical search criteria

- ° Each choice opens other fields relevant to the selection where you can further define your search query.
 - If permissible values were added when the annotation was defined, you must select among the values in a drop-list that displays on the right side of the page.
 - If no permissible values were defined as part of the annotation, you have the option to enter descriptive text in a text box on the right side of the page ([Figure 3.3](#)).


Figure 3.3 You may be able to further define search criteria when you select a specific clinical or imaging annotation element

Note: When working with image data, if only an Imaging Mapping file was uploaded when the study was created and not an Image Series Annotation file, you cannot enter image search criteria. The search results will, however, display a link that allows you to view the associated images in NBIA.

Continue with step 5 in [Searching a Study](#) on page 36.

Gene Expression

1. For the Gene Expression selection, select **Gene Name** or **Fold Change**.
2. For Gene Name or Fold Change, enter one or more gene symbols (up to 100 characters, separated by commas) in the text box or click the icons to locate genes in the following databases. If entering more than one gene in the text box, separate entries by commas.

- **CGAP** – Use this directory to identify genes. Before clicking this link you must enter gene symbols in the text box. This link does not pull anything into calIntegrator2 but does provide information about the gene(s) whose names you entered.
- **caBio** – This link searches caBIO, then pulls identified genes into calIntegrator2 for analysis. Click the caBIO icon (), enter **Keyword(s)** in the text box that opens and click **Search**. The search results display (Figure 3.4).

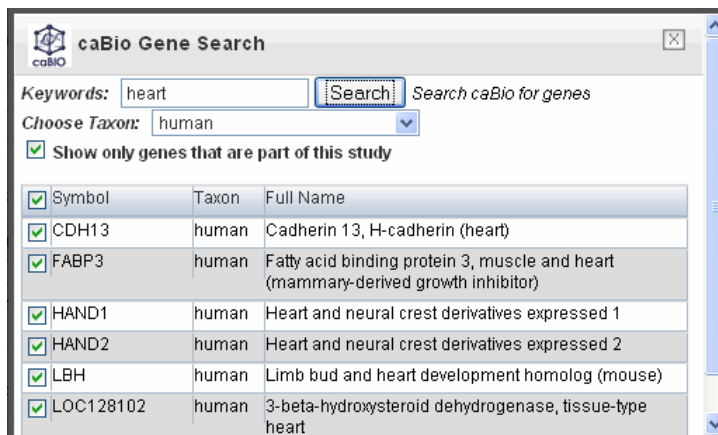


Figure 3.4 Example caBIO gene search results

3. Use the check boxes to identify the genes whose symbols you want to use in the gene expression analysis.
4. Click the **Use Genes** button at the bottom of the page. This pulls the checked genes into the Criteria tab (Figure 3.5).



Figure 3.5 Genes pulled in from caBIO display on the Criteria tab

Additional fields display for the Fold Change selection.

The fold change option appears only if genomic control samples have been uploaded to the study. Fold change identifies genes with expression differences compared to control samples, as defined when the study was deployed in calIntegrator2. You can enter query values in greater/lesser-than-or-equal-to arguments.

5. Select or enter data for the Fold change fields shown in [Figure 3.6](#) :

Figure 3.6 Fields for identifying fold change search criteria

- **Control Sample Set** – Select from the drop down list the name of the uploaded control sample set to serve as the fold change reference.
- **Regulation Type** – Select the term that describes the gene expression in comparison with the control samples: **Up** is increased expression; **Down** is decreased expression; **Up or Down** is increased or decreased; **Unchanged** means no change in expression.
- **Up-Regulation Folds** – The options here are dependent upon the Regulation Type you selected.
 - **Up** = Up Regulation Folds – Samples with a fold change greater than this value, when compared to the control samples, will be returned.
 - **Down** = Down Regulation Folds – Samples with a fold change less than this value, when compared to the control samples, will be returned.
 - **Up or Down** = Down Regulations Folds, Up Regulation Folds – Samples with a fold change either up or down, when compared to the control samples, will be returned.
 - **Unchanged** = Samples with a fold change between the two specified values will be returned.

Continue with step 5 in [Searching a Study](#) on page 36.

Results Type Tab

You can specify columns for the way you want the search results to display either before or after you run the search. If you run the search directly from the Criteria tab before setting the results type/sorting features, by default only the Subject Identifiers display on the Search Results tab. You can then come back to the Results Type tab and [Sorting Tab](#) to expand the display options and re-run the search, having set the display parameters.

The selection you make on the Results Type tab determines whether caIntegrator2 displays search results for clinical or genomic data. It filters the search based on the criteria you set on the Criteria tab, whether it is clinical, gene expression or image series data type(s). In other words, if you select clinical criteria on the Criteria tab, but select Genomic on the Results Type tab, the data subset that displays on the Search Results tab is genomic data that is filtered by the clinical criteria you defined on the Criteria tab.

1. On the Results Type tab, select the **Clinical** or **Genomic** radio button to search clinical data (*Figure 3.7*).

Search Demo Study based on Rembrandt with NCRI data.

Criteria Results Type Sorting Query Results Save as...

Select Results Type: ☐ Genomic ☒ Clinical

Genomic result type - will display a gene expression data matrix.
Clinical result type - will display tabular data, including column selection.

Select Columns for Results

| Subject Annotations | Image Annotations |
|--|--|
| <input checked="" type="checkbox"/> Age | <input type="checkbox"/> Comments |
| <input checked="" type="checkbox"/> Death Date | <input type="checkbox"/> Eloquent Cortex |
| <input type="checkbox"/> Disease | <input type="checkbox"/> Exam Number |
| <input type="checkbox"/> Followup Month | <input type="checkbox"/> Radiologist |
| <input checked="" type="checkbox"/> Gender | <input type="checkbox"/> Side of Tumor Epicenter |
| <input type="checkbox"/> ... | <input checked="" type="checkbox"/> Tumor Location |

Select All Unselect All Select All Unselect All

Run Query

Figure 3.7 Results Type tab

Clinical – Select the annotation elements that you want to display in the search results. All elements listed are column headers in the data uploaded to the study. You can make multiple selections on this list.

Note: For Clinical Annotations, the Patient or Subject Identifier display by default in the search results.

Results display as tabular data.

Genomic – Select the Reporter Type:

- **Gene Name** – Finds and summarizes at the gene level all reporters that match criteria for the gene you defined on the Criteria tab.
- **Reporter ID** – Finds all reporters that map to the gene(s) you identified on the Criteria tab

Results display in a gene expression data matrix.

Imaging – If imaging annotations have been added to the study, annotation elements also display on the lower right section of this page when you select **Clinical**. All elements listed are column headers in the image annotation data uploaded to the study. You can make multiple selections on this list.

Note: If you select even one Image Annotation on the Results Type tab, the Image Series IDs display by default in the search results. If you select no Image Annotations on the Results Type tab, however, even if you have selected image series criteria on the Criteria tab, no image series IDs display in the search results. The fact that images can be located, however, in NBIA is indicated by two image-related buttons at the bottom of the Query Results page. You can open the images in NBIA, but

they will be at StudyInstance UID level. See [Relationship of Patient to Study to Series to Images](#) on page 51.

Results display as tabular data.

2. Use the **Select All** or **Unselect All** buttons to aid you in making your selections.

The column selection is saved as part of the query if you save it. See [Saving a Query](#) on page 43.

Sorting Tab

On the Sorting tab, you can set the sort order for data columns in the query results. You can also indicate whether column contents are sorted in ascending or descending order.

The columns that display on the Sorting tab are those criteria that you selected on the [Results Type Tab](#) for a Clinical Results type search.

1. Select the Sorting tab and indicate the left to right column order of the Search Results by changing one or more numbers in the Column Order column in this table ([Figure 3.8](#)).

| Column | Column Order (L-R) | Row Order |
|---|--------------------|---|
| Death Date | 1 | <input type="radio"/> Ascending <input type="radio"/> Descending <input checked="" type="radio"/> No Sort |
| Prior Therapy Radiation Fraction Number | 2 | <input type="radio"/> Ascending <input type="radio"/> Descending <input checked="" type="radio"/> No Sort |
| Prior Therapy Surgery Procedure Title | 3 | <input type="radio"/> Ascending <input type="radio"/> Descending <input checked="" type="radio"/> No Sort |
| Institution | 4 | <input type="radio"/> Ascending <input type="radio"/> Descending <input checked="" type="radio"/> No Sort |
| Gender | 5 | <input type="radio"/> Ascending <input type="radio"/> Descending <input checked="" type="radio"/> No Sort |
| Prior Therapy Surgery Tumor Histology | 6 | <input type="radio"/> Ascending <input type="radio"/> Descending <input checked="" type="radio"/> No Sort |
| Prior Therapy Radiation Type | 7 | <input type="radio"/> Ascending <input type="radio"/> Descending <input checked="" type="radio"/> No Sort |
| Last Followup Date | 8 | <input type="radio"/> Ascending <input type="radio"/> Descending <input checked="" type="radio"/> No Sort |
| Age | 9 | <input type="radio"/> Ascending <input type="radio"/> Descending <input checked="" type="radio"/> No Sort |
| Race | 10 | <input type="radio"/> Ascending <input type="radio"/> Descending <input checked="" type="radio"/> No Sort |
| Prior Therapy Surgery Outcome | 11 | <input type="radio"/> Ascending <input type="radio"/> Descending <input checked="" type="radio"/> No Sort |
| Karnofsky | 12 | <input type="radio"/> Ascending <input type="radio"/> Descending <input checked="" type="radio"/> No Sort |

Figure 3.8 Sorting tab

2. In the **Row Order** column, indicate how you want columns sorted, **Ascending** or **Descending**, or leave the default, **No Sort**, if you choose.
3. Click **Run Query** at the bottom of the page to execute your sorting changes in the search results. When you do so, the change in column order is visible on the Query Results tab, as well as on the Sorting tab. For example, any column that you have indicated to be number 1 now appears in Query Results immediately after the Subject Identifier column and at the top of the Set Sort Order table on the Sorting tab.

Sorting parameters are saved as part of the query if you choose to save it using the Save Query feature. See [Saving a Query](#) on page 43.

4. If you click the **Reset** button before running the query from the Sorting tab, the original column settings are restored.

For information about the search results, see [Chapter 4 Viewing Query Results](#).

Managing Queries

When you create a search query in calIntegrator2, you can save the query for later use or edit it.

[Exporting Query Results](#)

Saving a Query


To save a query, follow these steps:

1. Click the **Save As** tab and enter a **Search Name** and **Search Description**, unique to the search. *Example: **Batch ID 6 and female***
2. Click **Save**.

Once the query is saved, it is listed by its name under the **Study Data > Queries > My Queries** in the left sidebar, whenever the study to which the query applies is selected. Click on the saved query in this list to either edit or re-run the query. Click on the query name to retrieve query results. If you hover over the Name text for the query, a popup displays the query description.

Editing a Query

To edit a query, follow these steps:

1. To edit a query, select it in the left sidebar under the **Study Data > Queries > My Queries**.
2. Click the **Edit** icon () corresponding to the study.
3. Change the query and display criteria on the Criteria, Columns and Sorting tabs.
4. On the Save As tab, check the appropriate options and click **Save As**. You can use the same name as the original query or modify the name as needed.

Exporting Query Results

After running a search, you can export the result set or a subset as a tab-delimited text file. For more information, see [Exporting Data](#) on page 52.

CHAPTER 4

VIEWING QUERY RESULTS

This chapter describes search results that calIntegrator2 returns after queries.

Topics in this chapter include the following:

- [Query Results Overview](#) on this page
- [Browsing Query Results](#) on page 46

Query Results Overview

After you launch a search of a calIntegrator2 study, the system automatically opens the Query Results tab showing the results of your search.

If you have not configured column and sort display parameters before launching the search, by default the tab shows only the subject identifiers and a column that allows you to select each row of the data subset.

To display and/or sort additional data, you must return to the Columns and/or Sorting tabs to set display parameters, then re-run the search. The new search results will display the additional information, with the columns and data sorted as you specified. See [Results Type Tab](#) on page 40.

calIntegrator2 paginates search results into pages of configurable size (default 20) with standard paginated navigation controls. To sort columns by ascending or descending parameters for on any displayed field, click on the underlined column header.

You can download search results as a CSV file. The file contains the annotations, columns and data sort configurations you specified in the search query. See [Exporting Query Results](#) on page 43.

Browsing Query Results

The query results that can display depend upon the criteria you established for the search. Follow the links below for more information about the category of data you searched.

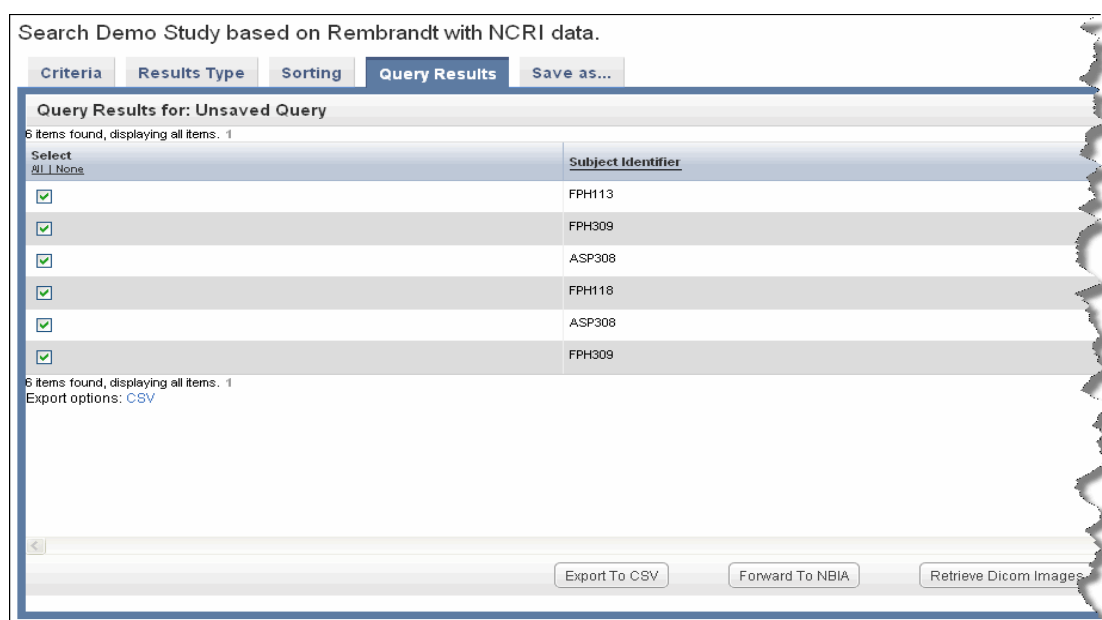
[Clinical and Imaging Data](#) on page 46

[Genomic Data](#) on page 46

[Expanding Imaging Data Results](#) on page 48

Clinical and Imaging Data

If you run the search before configuring column and sort display parameters, only the [subject] ID that meet the criteria and a column allowing you to select each row appear on the table ([Figure 4.1](#)). .



Search Demo Study based on Rembrandt with NCRI data.

Criteria Results Type Sorting **Query Results** Save as...

Query Results for: Unsaved Query

6 items found, displaying all items. 1

| Select | Subject Identifier |
|-------------------------------------|--------------------|
| <input checked="" type="checkbox"/> | FPH113 |
| <input checked="" type="checkbox"/> | FPH309 |
| <input checked="" type="checkbox"/> | ASP308 |
| <input checked="" type="checkbox"/> | FPH118 |
| <input checked="" type="checkbox"/> | ASP308 |
| <input checked="" type="checkbox"/> | FPH309 |

6 items found, displaying all items. 1
Export options: [CSV](#)

Export To CSV Forward To NBIA Retrieve Dicom Images

Figure 4.1 Query Results page

You can add details for one or more subjects by configuring them on the Results Type tab. Annotations listed there are the column headers in the CSV file(s) that were uploaded to the study. For information about using the Results Type tab, see [Results Type Tab](#) on page 40.

Genomic Data

If you select the Genomic result type on the Results Type tab, genomic data search results display in a gene expression data matrix. Because the data was downloaded from caArray, the data permissions granted there still apply. In other words, if you have been given access to the data in caArray, you can see it in caIntegrator2.

In the matrix, samples in the experiment form the column headings. If the rows display Gene Name, the cells display the median gene expression value for each gene. If the

rows display Probe Set, the cells display the normalized signal-based value for a given reporter for a given sample (*Figure 4.2, Figure 4.3*).

Search Demo Study based on DC Lung Full data

Criteria

Results Type

Sorting

Query Results

Save as...

Query Results for: Unsaved Query

| | | | | | | | | | | | | | | | | | | |
|------------|--------|--------|--------|-------|-------|--------|-------|-------|--------|--------|--------|-------|--------|-------|--------|--------|-------|-----|
| Subject ID | 321 | 326 | 158 | 53 | 180 | 350 | 545 | 54 | 142 | 199 | 539 | 89 | 193 | 107 | 305 | 244 | 74 | 361 |
| Sample ID | 321 | 326 | 158 | 53 | 180 | 350 | 545 | 54 | 142 | 199 | 539 | 89 | 193 | 107 | 305 | 244 | 74 | 361 |
| Gene Name | | | | | | | | | | | | | | | | | | |
| EGFR | 151.48 | 395.43 | 179.63 | 70.39 | 91.81 | 175.26 | 97.61 | 80.96 | 111.84 | 115.25 | 309.05 | 81.36 | 136.61 | 84.75 | 145.52 | 208.45 | 66.68 | 14 |

Export options: CSV

Figure 4.2 Gene Name search EGFR, Reporter Type: Gene

Search Demo Study based on DC Lung Full data

Criteria

Results Type

Sorting

Query Results

Save as...

Query Results for: Unsaved Query

| | | | | | | | | | | | | | |
|-----------|-------------|---------|--------|---------|---------|--------|--------|--------|---------|---------|---------|---------|--------|
| | Subject ID | 21 | 362 | 369 | 332 | 480 | 135 | 93 | 345 | 255 | 486 | 333 | 115 |
| | Sample ID | 21 | 362 | 369 | 332 | 480 | 135 | 93 | 345 | 255 | 486 | 333 | 115 |
| Gene Name | Reporter ID | | | | | | | | | | | | |
| EGFR | 201983_s_at | 7071.93 | 957.15 | 1747.74 | 2614.96 | 875.01 | 646.42 | 503.99 | 4350.84 | 1297.73 | 2425.24 | 1275.78 | 365.28 |
| EGFR | 201984_s_at | 2625.52 | 402.21 | 1225.3 | 1629.62 | 569.71 | 477.81 | 311.24 | 673.6 | 611.8 | 637.53 | 347.0 | 376.42 |
| EGFR | 210984_x_at | 420.18 | 44.28 | 56.72 | 214.84 | 133.44 | 24.43 | 20.37 | 36.96 | 40.0 | 31.66 | 45.54 | 40.23 |
| EGFR | 211550_at | 9.43 | 18.68 | 33.64 | 11.73 | 23.74 | 15.93 | 12.4 | 31.44 | 29.33 | 20.11 | 33.91 | 14.49 |
| EGFR | 211551_at | 180.43 | 157.4 | 349.1 | 208.0 | 161.24 | 173.67 | 59.78 | 213.68 | 319.14 | 189.92 | 250.89 | 131.74 |
| EGFR | 211607_x_at | 378.93 | 96.02 | 48.57 | 108.57 | 209.22 | 13.32 | 29.6 | 61.36 | 62.75 | 52.88 | 87.53 | 47.32 |

Export options: CSV

Figure 4.3 Gene Name search EGFR, Reporter Type: Reporter ID

- Genomic data does not display in tandem with clinical and imaging data; it only displays when you select the Genomic result type on the Results Type tab. Genomic data is however, filtered by clinical and imaging query criteria configured on the Criteria tab.
- Click the Export Options CSV link to download the CSV file whose data displays on the Search Results tab. When you do so, the CSV file opens automatically in MS Excel or similar applications for working with spreadsheets, showing the columns and sorting as you defined them in calIntegrator2 on the appropriate tabs.

Expanding Imaging Data Results

In reviewing imaging search results, it is important to understand the hierarchy of submissions in NBIA. For more information, see [Relationship of Patient to Study to Series to Images](#) on page 51.

If you run the search before configuring column and sort display parameters, only the Subject Identifiers for the patients/images that meet the criteria and a column containing one check box per row display by default ([Figure 4.4](#)).

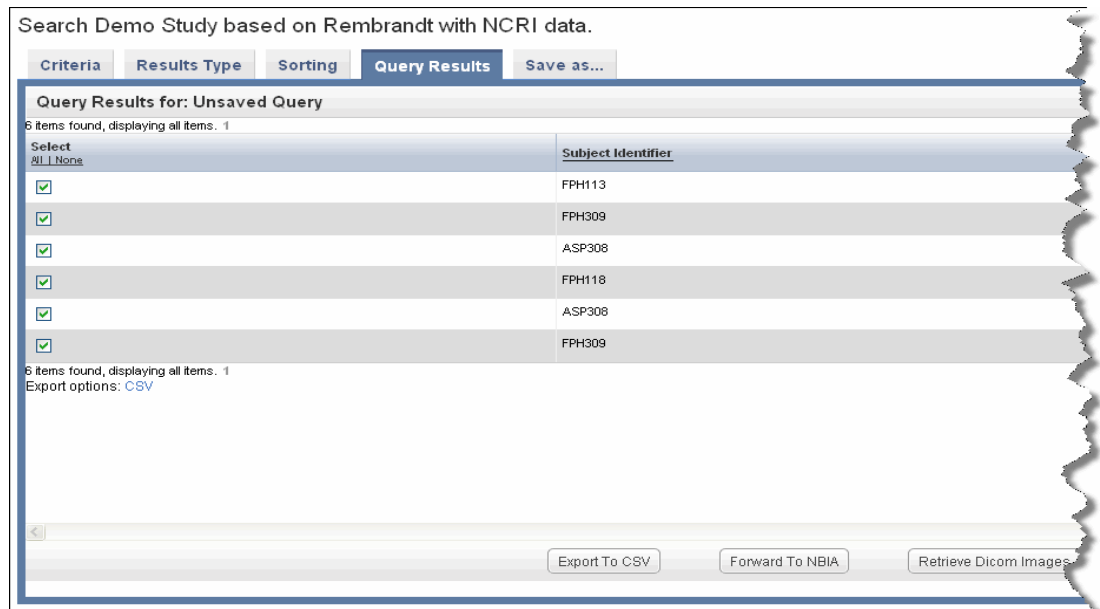


Figure 4.4 With imaging criteria only and no column definition, only Subject IDs display

If your annotation choice on the Columns page identifies annotations such as tumor size or tumor location, the search results display image series subsets that have those annotations. The check boxes work in conjunction with buttons at the bottom of the results page ([Figure 4.5](#)).

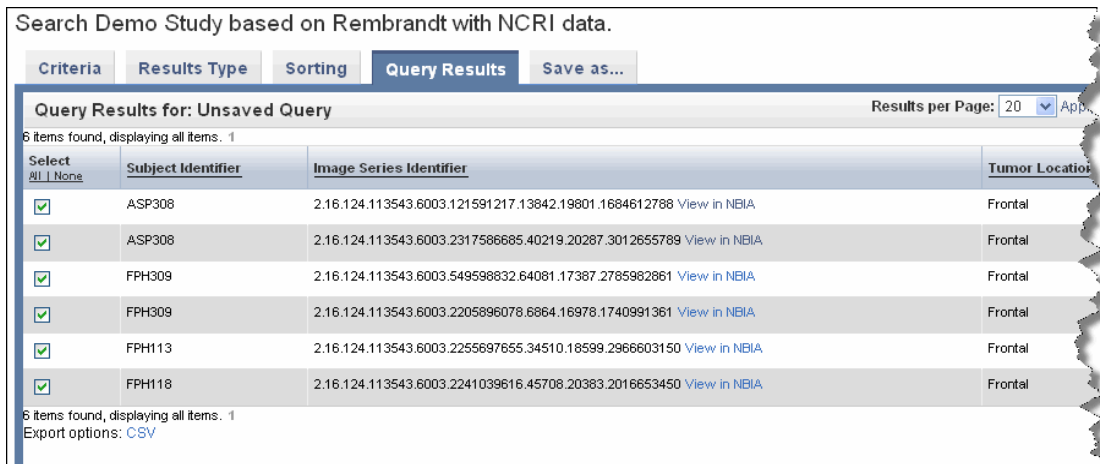


Figure 4.5 By expanding display parameters, you can view complete details for image search results

You can add more details for images by configuring image annotations on the Results Type tab. Annotations listed there are the column headers in the image series CSV file(s) that were uploaded to the study. Examples of image details include the following:

- All image details (name, size, etc.)
- The series that the image belongs to
- Image feature attributes
- The subject ID. Click the subject ID under Clinical Annotations on the Results Type tab to display this.

You can set display parameters for the results on the Columns and Sorting tabs. For more information, see [Results Type Tab](#) on page 40.

See also [caIntegrator2](#) and [NBIA](#) and

caIntegrator2 and NBIA

Images can be accessed in NBIA if you see buttons on the Search Results page. See the Imaging Note in [Results Type Tab](#) on page 40. You can click links on the Search Results tab to view or download image data.

- **View in NBIA** – This link corresponds to each Image Series listed in the results table. If you click the link, NBIA opens to the login page. After you log in, NBIA brings up the first image in the corresponding image series ([Figure 4.6](#)). You must log into NBIA to see the data. On the NBIA page that opens, you can opt to view the entire series containing this image, or you can display the image as a large JPEG-formatted image. You can also add the image to the NBIA basket. For more information, see the NBIA online help or user's guide accessible from NBIA.



Figure 4.6 An example of displaying the first image in image series

- **Forward to NBIA** – This button is linked to results you have selected by row. Click the button to open NBIA, where the image series you select are loaded in the NBIA image basket. In the event that the caIntegrator2 study was NOT

configured with image annotation for an image series, calIntegrator2 sends NBIA a list of Study Instance UIDs, for which NBIA will add all corresponding image series to the basket. In the event that the calIntegrator2 study was configured with annotations for an image series, the system sends NBIA a list of Image Series IDs, for which NBIA adds all corresponding image series to the basket.

Retrieving Dicom Images

On the Imaging data Search Results page, you can click the **Retrieve DICOM Images** button which is linked to results you have selected by row. calIntegrator2 retrieves the corresponding image(s) from NBIA through the grid. NBIA organizes the download file by patientID, StudyInstanceUID, and ImageSeriesUID, and compresses it into a zip file. When calIntegrator2 notifies you that the file is retrieved, the DICOM Retrieval page indicates whether the retrieved files are Study Instance UIDs or Image Series UIDs (*Figure 4.7*). For more information, see the note below.



Figure 4.7 DICOM Retrieval result

Click the **Download DICOM** link to download and save the file. calIntegrator2 unzips the file and displays the list of images in the file. To open the DICOM images, you must have a DICOM image viewer application installed on your computer. For more information, see <http://dicom.online.fr/fr/download.htm>.

In the search results, not all of the patients in the data subset may be mapped to image series IDs. If you select a mixture of patients that have image annotations as indicated by an image series ID and patients that do not have image annotations (no image series ID), when you click the **Retrieve DICOM Images** button, NBIA retrieves the images for the entire *NBIA study instance UID* that includes the image seriesIDs you checked.

If on the Search Results tab you select only patients that have image annotations as indicated by an image series ID, when you click the **Retrieve DICOM Images** button, NBIA retrieves images for the *NBIA image series* that were matched in the search. If the results are a mixture, but you select one specific row with a valid image annotation, calIntegrator2 aggregates to the *images series*. If results are a mixture and you select multiple rows, calIntegrator2 aggregates to the NBIA study in which multiple image series you have selected in the search results are found.

If your query does not have image annotations and all check boxes are selected, results will go up to image series UID and gives all image series in it. Search results

may ultimately depend on how the study was created. For example, if no image series display in query results, it means they were not mapped in the study. In that case, the results “move” up to Study Instance UIDs.

To best understand this, it is important to review the hierarchy of submissions in NBIA. For more information, see [Relationship of Patient to Study to Series to Images](#) on page 51.

Example of Retrieving Images:

You are searching a study that has image data and image annotation(s) for at least one image series.

1. Open a study that has imaging data associated with it that points to the production NBIA server.
2. Make a query that will have image series or patients who are associated to Image Studies and select a few of those patients in the check box.
3. Click the **Retrieve Dicom Images** button.

Note that it aggregates to the image study.

4. Now go back to Results Type tab, select all image annotations and run the query again.
5. Select an image series type column and click the **Retrieve Dicom Images** button.

calIntegrator2 now aggregates to the Image Series that were selected and not the Image Study.

6. Select a row that doesn't have image series data, and a row that does, and push the button.

This should aggregate to the study for the rows selected.

7. Click **Forward to NBIA**. You should see the same types of aggregation for these tests.

When the image Study is in the checked boxes (regardless of image series being there or not), the system aggregates up to the Image Study level.

Relationship of Patient to Study to Series to Images

This flowchart illustrates the relationship of patient to study to series and lastly to images.

Clinical trial > Patient (Subject) > Study > Series > Images

For example, the Study Instance UID is the set of images resulting from one patient office visit. When you upload a spreadsheet of an image series, the hierarchy of images in an image series might look like this:

Study Instance UID (one office visit):

Brain (image series)

- Brain image 1
- Brain image 2
- Brain image 3

Leg (image series)

- Leg image 1
- Leg image 2
- Leg image 3

You can add details for images by configuring image annotations on the Results Type tab. Annotations listed there are the column headers in the image series CSV file(s) that were uploaded to the study. Examples of image details include the following:

- All image details (name, size, etc.)
- The series that the image belongs to
- Image feature attributes
- The subject ID. Click the subject ID under Clinical Annotations on the Results Type tab to display this.

Exporting Data

You can choose to download tabular search results as a CSV file. Click the **Export .csv** link at the bottom of the page. You may need to scroll the page to see it. The file contains the annotations, columns and data sort configurations you specified in the search query.

Note: You will not see the Export option when genomic data displays as query results.

CHAPTER 5

ANALYZING STUDIES

This chapter describes how to use calIntegrator2 tools to analyze data in clinical or genomic studies that have been deployed in calIntegrator2.

Topics in this chapter include the following:

- [Data Analysis Overview](#) on this page
- [Creating Kaplan-Meier Plots](#) on page 54
- [Creating Gene Expression Plots](#) on page 60
- [Analyzing Data with GenePattern](#) on page 73

Data Analysis Overview

Once a study has been deployed, you can analyze the data using calIntegrator2 analysis tools.

You can verify that the study is in “Deployed” status by selecting the study name in the My Studies dropdown selector. After selecting the study name, click **Home** in the left sidebar of the calIntegrator2 Menu. A study summary should appear, including a status field. If the status is not deployed, or if the study summary does not appear, then the study is not deployed and available for analysis.

If the study is ready for analysis, you will see an **Analysis Tools** menu in the left sidebar with the following options:

- **K-M Plot:** This tool analyzes clinical data, generating a Kaplan-Meier (K-M) plot based on survival data sets. See [Creating Kaplan-Meier Plots](#) on page 54.
- **Gene Expression Plot:** This tool analyzes annotation, clinical or genomic data based on gene expression values. See [Creating Gene Expression Plots](#) on page 60.

- **GenePattern:** This feature provides an express link to GenePattern where you can perform analyses on selected calIntegrator2 studies, or it enables you to perform several GenePattern analyses on the grid. See [Analyzing Data with GenePattern](#) on page 73 .

After defining or running the analysis on selected data sets, analysis results display on the same page, allowing you to review the analysis method parameters you defined.

Creating Kaplan-Meier Plots

The Kaplan-Maier method analyzes comparative groups of patients or samples. In calIntegrator2, the K-M method compares survival statistics among comparative groups. You can configure the survival data in the application. For example, you might identify a group of patients with smoking history and compare survival rates with a group of non-smoking patients, or compare the survival data for two groups of patients with a specific disease type and based on Karnofsky scores . You could compare groups of patients with varying gene expression levels. You can also identify data sets using the query feature in the application, saving the queries, then configuring the K-M to compare groups identified by the queries.

The key is to first identify subsets of patients or samples that meet criteria you want to establish, thus filtering the data you want to compare. Next, generate a K-M plot based on their survival probability as a function of time. Survival differences are analyzed by the log-rank test.

Note: To perform a K-M plot analysis, survival data must have been identified for the study you want to analyze. For more information, see [Defining Survival Values](#) on page 23.

K-M Plot for Annotations

The groups identified for this K-M plot generation are based on clinical annotations.

1. Select the study whose data you want to analyze in the upper right portion of the calIntegrator2 page.
2. Under Analysis Tools on the left sidebar, select **K-M Plot**.
3. Select the **For Annotation** tab at the top of the page ([Figure 5.1](#)).

Kaplan-Meier Survival Plots (draft)

For Annotation
For Gene Expression
For Queries

Annotation Based Kaplan-Meier Survival Plots

| | Annotation Type | Annotation | Values |
|--------------------------------------|---|--|----------------------|
| 1.) Patient Groups: | <input type="text" value="Select Annotation Type"/> | <input type="text" value="Select Annotation"/> | <input type="text"/> |
| Survival Value | | | |
| 2.) Select Survival Measure: | <input type="text" value="Survival from enrollment"/> | | |
| <input type="button" value="Reset"/> | | | |

Figure 5.1 Fields for defining annotation data for a K-M plot

4. Select fields as described in [Table 5.1](#).

| Field | Description |
|-----------------------|---|
| Patient Groups | <p>The groups to be compared in the K-M plot originate from one patient group. Varying data sets are based upon multiple values corresponding to the selected annotation.</p> <p>Annotation Type – Select the annotation type that identifies the patient group. Selections are based on the data in the chosen study.</p> <p>Annotation – Select an annotation. Fields are based on the annotation type you select. For example, if you choose Subject, then you could select Gender or Radiation Type or any field that would distinguish the patients into groups based upon their values.</p> <p>Values – Using conventional selection techniques, select two or more values which will be the basis for the K-M plot. Permissible (available) values or “No Values” correspond to the selected annotation.</p> |
| Survival Value | <p>Survival value is the length of time the patient lived. Select the survival measure, which is the unit of measurement for the survival value to be used for the K-M plot.</p> |

Table 5.1 Fields for selecting K-M annotations plot values

5. Click the **Create Plot** button.

Note: The Create Plot button displays only after you have selected appropriate criteria.

calIntegrator2 generates the plot which then displays below the K-M plot criteria ([Figure 5.2](#)).

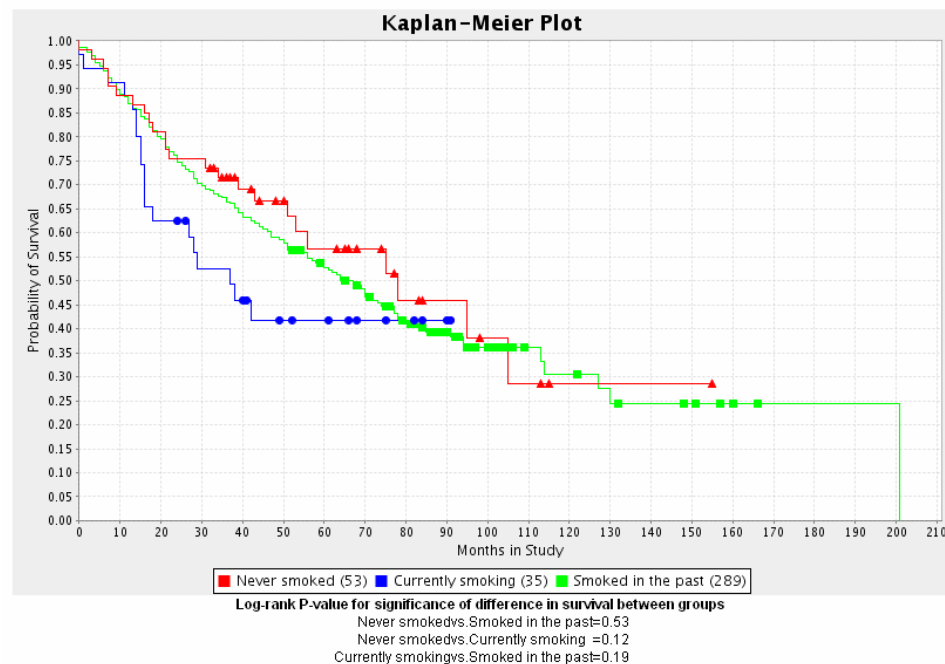


Figure 5.2 A K-M plot generated for groups based on clinical annotations

The number of subjects for each group appears embedded in the legend of the graph below the plot.

calIntegrator2 generates a P-value for the selected groups; it displays at the bottom of the page. A low P-value generally has more significance than a high P-value.

K-M Plot for Gene Expression

calIntegrator2 allows you to compare expression levels for one given gene at a time. The relative expression level is referred to as “fold change” and the numeric value for a given sample and reporter combination is the ratio of the expression value for that particular reporter for the given sample to a reference value calculated for that reporter across all control samples. The reference value is calculated by taking the mean of the \log_2 of the expression values for all control samples for the reporter in question. The \log_2 mean value (n) is then converted back to a comparable expression signal by returning 2 to the exponent n .



To create a K-M plot illustrating gene expression values, follow these steps:

1. Select the study whose data you want to analyze in the upper right portion of the calIntegrator page. You must select a study with gene expression data.
2. Under Analysis Tools on the left sidebar, select **K-M Plot**.
3. Select the **For Gene Expression** tab ([Figure 5.3](#)).

Kaplan-Meier Survival Plots


For Annotation | **For Gene Expression** | For Queries

Gene Expression Based Kaplan-Meier Survival Plots

1.) Gene Symbol:  

2.) Overexpressed >= fold

3.) Underexpressed >= fold

4.) Select Survival Value: 


5.) Select Control Sample Set: 

Figure 5.3 Fields for defining gene expression data for a K-M plot

4. Enter or select fields as described in [Table 5.2](#).

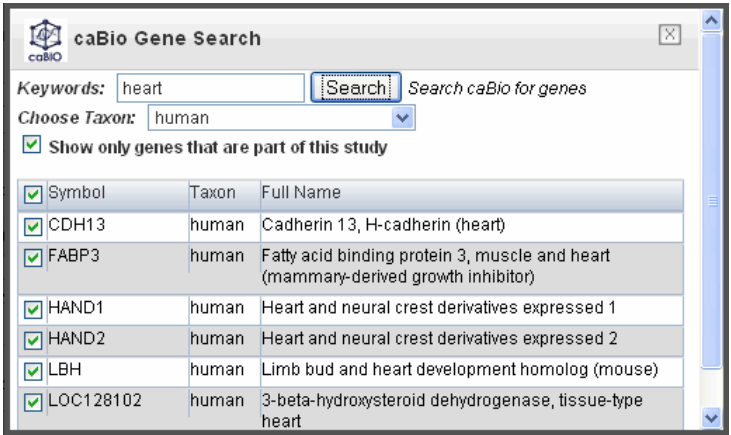


| Field | Description |
|------------------------------------|--|
| Gene Symbol | <p>In the text box, specify a single gene whose expression values can be used to split the subjects into three categories: high, low and intermediate expression or click the icons to locate genes in the following databases. If entering more than one gene in the text box, separate entries by commas.</p> <p>CGAP – Use this directory to identify genes. Before clicking this link you must enter gene symbols in the text box. This link does not pull anything into calIntegrator2 but does provide information about the gene(s).</p> <p>caBio – This link pulls identified genes into calIntegrator2 for analysis. Click the icon, enter Keyword(s) in the text box that opens and click Search.</p>  <p>Use the check boxes to identify the genes whose symbols you want to use in the gene expression analysis.</p> <p>Click the Use Genes button at the bottom of the page. This pulls the checked genes into the Gene Expression ... tab.</p> <p>Annotation Based Gene Expression Plots</p> <p>1.) Gene Symbol(s) (comma separated list): CDH13,FABP3,HAND1,HA  </p> |
| Over-expressed/ Under-expressed | Define the over- and under-expression criteria, expressed in terms of fold-change. Fold change is the ratio of the measured gene expression value for an experimental sample to the expression value for the control sample. |
| Survival Value | Survival value is the length of time the patient lived. Select the survival measure which is the unit of measurement for the survival value to be used for the K-M plot. |
| Control Sample Set | One or more control sample sets are created by the study manager when the study is deployed. Select the control sample set you would like to use to calculate fold-change. |

Table 5.2 Fields for selecting K-M gene expression plot values

- Click the **Create Plot** button. calIntegrator2 generates the plot which then displays below the K-M plot criteria ([Figure 5.4](#)).

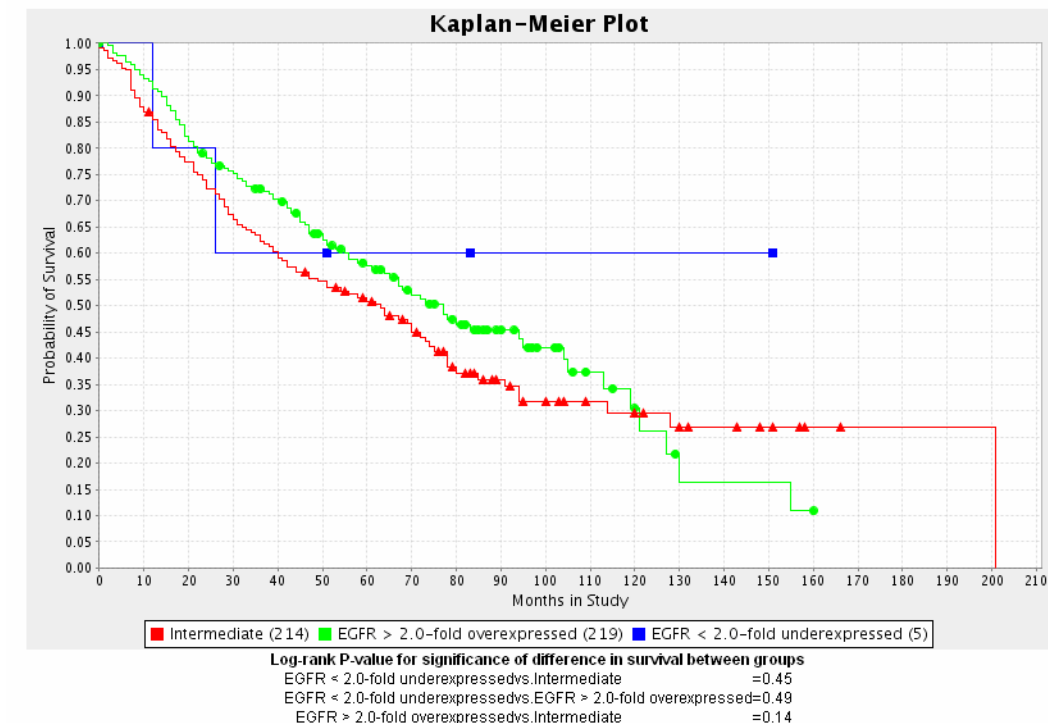


Figure 5.4 K-M plot generated from gene expression data.

The number of subjects for each group appears embedded in the legend of the graph below the plot. Note the appearance of an intermediate group (red entries), which is a group with gene expression values that are not up-regulated nor down-regulated.

In queries that include a fold change criterion and that are configured to return genomic data, the raw expression values are replaced with the calculated fold change value.

A P-value is also generated for the selected groups; it displays at the bottom of the page. A low P-value generally has more significance than a high P-value.

K-M Plot for Queries

You can identify data sets using the query feature in the application. You can manipulate the queries to find the groups you want to compare, save the queries, then configure the K-M to compare the query groups. This is one method of limiting the data considered in the K-M plot calculation.

- Select the study whose data you want to analyze in the upper right portion of the calIntegrator page. You must select a study for which the queries you will identify for the K-M plot have been saved.
- Under Analysis Tools on the left sidebar, select **K-M Plot**.

3. Select the **For Queries** tab (*Figure 5.5*).

Kaplan-Meier Survival Plots (draft)

For Annotation For Gene Expression **For Queries**

Query Based Kaplan-Meier Survival Plots

1.) Select Queries:

All Available Queries

- gender female
- equal to or > 60
- equal to or > 70
- Never smoke
- equal to or > 40
- equal to or > 30

Add >

< Remove

Selected Queries

v ^

2.) ☐ Exclusive Subjects in Queries (Subjects in upper queries are removed from subsequent queries)

3.) ☐ Add additional group containing all other subjects not found in selected queries.

4.) Select Survival Value: Survival from enrollment

Reset Create Plot

Figure 5.5 Fields for defining K-M plot parameters based on saved queries in caIntegrator2

4. Enter or select fields as described in *Table 5.3*.

| Field | Description |
|---|---|
| Queries | Select the queries whose data you want to analyze from the All Available Queries panel and move them to the Selected Queries panel using the Add >> button. Note: Genomic queries do not appear in the lists; they cannot be selected for this type of K-M plot. |
| Exclusive Subject in Queries | Check the box if you want to exclude any subjects that appear in both (or all) queries selected for the plot, thus eliminating overlap. |
| Add additional group...other subjects... | Check the box to create an additional group of subjects that are not in your other selected query groups. |
| Survival Value | Survival value is the length of time the patient lived. Select the survival measure, which is the unit of measurement for the survival value to be used for the K-M plot. |

Table 5.3 Fields for selecting K-M plot values based upon caIntegrator2 queries

5. Click the **Create Plot** button. caIntegrator2 generates the plot which then displays below the K-M plot criteria (*Figure 5.6*).

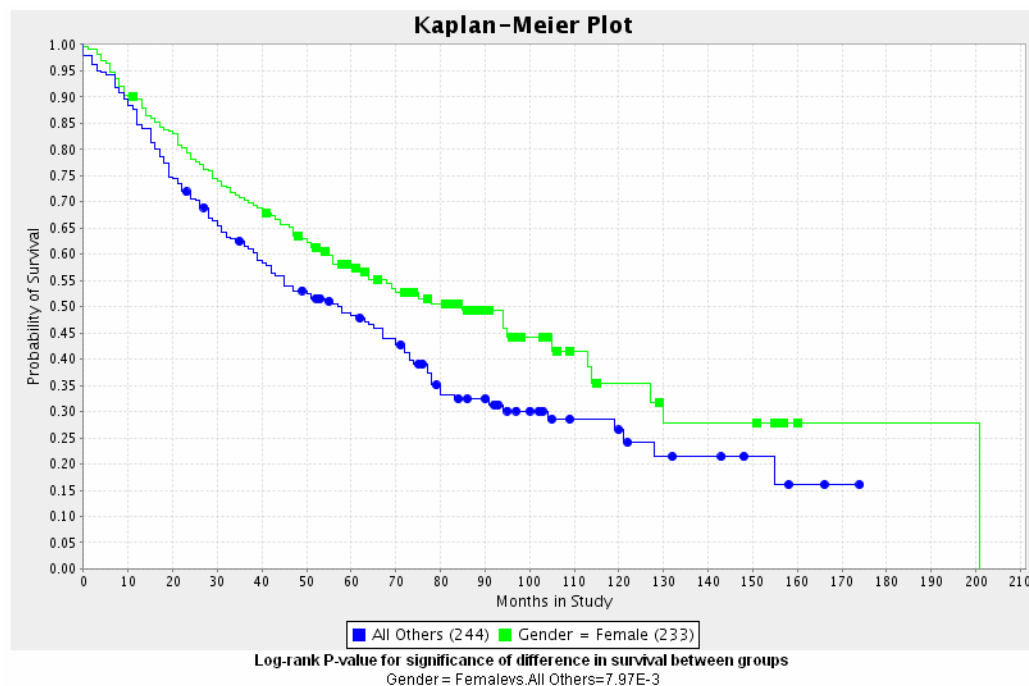


Figure 5.6 K-M Plot comparing statistics between subjects in two queries

The number of subjects for each group appears embedded in the legend of the graph below the plot.

A P-value is also generated for the selected groups; it displays at the bottom of the page. A low P-value generally has more significance than a high P-value.

Creating Gene Expression Plots

Gene expression plots compare signal values from reporters or genes. This statistical tool allows you to compare values for multiple genes at a time, but it does not require only two sets of data to be compared. It also allows you to compare expression levels for selected genes against expression levels for a set of control samples designated at the time of study definition.

calIntegrator2 provides three ways to generate meaningful gene expression plots, indicated by tabs on the page. The tabs are independent of each other and allow you to select the genes, reporters and sample groups to be analyzed on the plot.

- [Gene Expression Value Plot for Annotation](#) – You can locate genes in the CGAP and caBio directories and criteria can be defined using clinical and image annotations.
- [Gene Expression Value Plot for Genomic Queries](#) – You can select data based on saved genomic queries.
- [Gene Expression Value Plot for Clinical Queries](#) – You can select data based on saved clinical queries.

See also [Understanding a Gene Expression Plot](#) on page 70.

Gene Expression Value Plot for Annotation

To generate a gene expression plot, follow these steps:

1. Select the study whose data you want to analyze in the upper right portion of the calIntegrator2 page. (You must select a study which has genomic data.)
2. Under Analysis Tools on the left sidebar, select **Gene Expression Plot**. This opens a page with three tabs
3. Select the **For Annotation** tab ([Figure 5.7](#)).

Gene Expression Value Plots

For Annotation For Genomic Queries For Clinical Queries

Annotation Based Gene Expression Plots

1.) Gene Symbol(s) (comma separated list): CxAP CASIO

2.) Select Reporter Type: ☒ Reporter Id ☐ Gene

| Annotation Type | Annotation | Values |
|--|--|----------------------|
| 3.) Sample Groups: <input type="text" value="Select Annotation Type"/> | <input type="text" value="Select Annotation"/> | <input type="text"/> |

4.) ☐ Add additional group containing all other subjects not found in selected queries.

5.) ☐ Add additional group containing all control samples for this study.

Reset

Figure 5.7 Gene expression value tab for configuring gene expression annotation value plot

4. Enter or select fields as described in [Table 5.4](#).

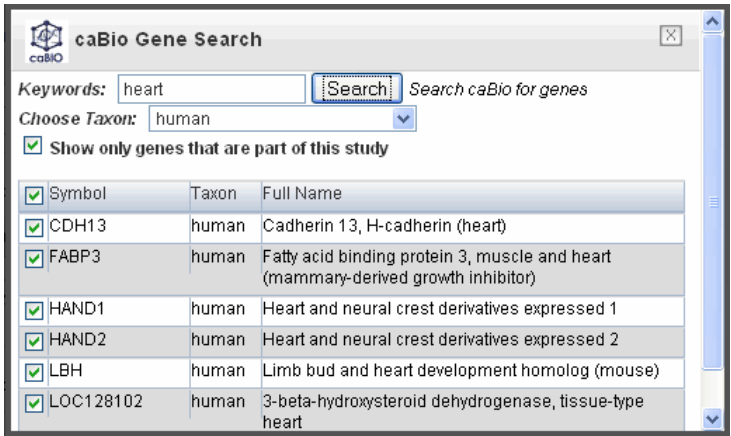


| Field | Description |
|---------------|---|
| Gene Symbol | <p>Enter one or more gene symbols in the text box or click the icons to locate genes in the following databases. If entering more than one gene in the text box, separate entries by commas.</p> <p>CGAP – Use this directory to identify genes. Before clicking this link you must enter gene symbols in the text box. This link does not pull anything into caIntegrator2 but does provide information about the gene(s).</p> <p>caBio – This link pulls identified genes into caIntegrator2 for analysis. Click the icon, enter Keyword(s) in the text box that opens and click Search.</p>  <p>Use the check boxes to identify the genes whose symbols you want to use in the gene expression analysis.</p> <p>Click the Use Genes button at the bottom of the page. This pulls the checked genes into the Gene Expression ... tab.</p> <p>Annotation Based Gene Expression Plots</p> <p>1.) Gene Symbol(s) (comma separated list): CDH13,FABP3,HAND1,HA  </p> |
| Reporter Type | <p>Select the radio button that describes the reporter type:</p> <p>Reporter ID – Summarizes expression levels for all reporters you specify.</p> <p>Gene Name – Summarizes expression levels at the gene level.</p> |

Table 5.4 Fields for selecting gene expression plot values based upon annotations

| Field | Description |
|--|---|
| Sample Groups | <p>Annotation Type – Select the annotation type. Selections are based on the data in the chosen study</p> <p>Annotation – Select an annotation. Fields are based on the annotation type you select. For example, if you choose Subject, then you could select Gender or Radiation Type or any field that would distinguish the patients into groups based upon study values.</p> <p>Values – Using conventional selection techniques, select one or more values which will be the basis for the plot. Permissible (available) values or “No Values” correspond to the selected annotation.</p> |
| Add additional group...all other subjects | Check the box to create an additional group of all other subjects that are not in selected query groups. |
| Add additional group...control group | Check the box to display an additional group of control samples for this study. |

Table 5.4 Fields for selecting gene expression plot values based upon annotations

- Click the **Create Plot** button. calIntegrator2 generates the plot which then displays below the Gene Expression Plot criteria in bar graph format (*Figure 5.8*). Legends below the plot indicate the plot input.

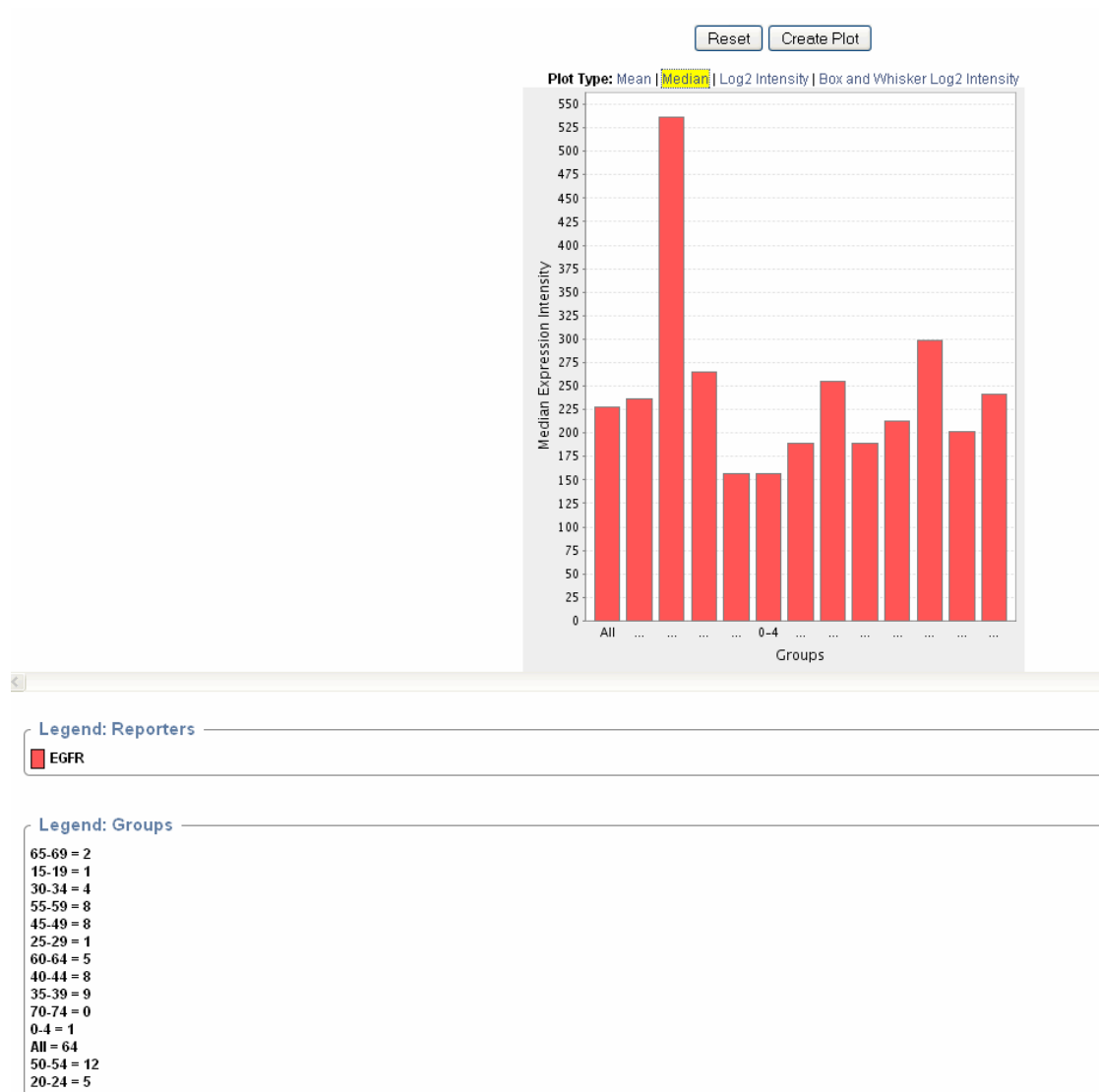


Figure 5.8 Gene expression plot based on selected annotations

6. By default, calIntegrator2 displays a plot showing the mean of the data. You can recalculate the data display by clicking the **Plot Type** above the graph. See [Understanding a Gene Expression Plot](#) on page 70

Figure 5.8 displays a plot with gene expression median calculation summaries. Legends below the plot indicate the plot input.

7. You can modify the plot parameters and click the **Reset** button to recalculate the plot.

Gene Expression Value Plot for Genomic Queries

Data to be analyzed on this tab must have been saved as a genomic query. For more information, see [Saving a Query](#) on page 43.

To generate a gene expression plot using a genomic query, follow these steps:

1. Select the study whose data you want to analyze in the upper right portion of the caIntegrator page. (You must select a study which has genomic data.)
2. Under Analysis Tools on the left sidebar, select **Gene Expression Plot**.
3. Select the **For Genomic Queries** tab ([Figure 5.9](#)).

Gene Expression Value Plots

Figure 5.9 Gene expression value tab for configuring gene expression genomic queries plot

4. Enter or select fields as described in [Table 5.5](#).

| Field | Description |
|----------------------|---|
| Genomic Query | Click on the genomic query upon which the plot is to be based. |
| Reporter Type | Select the radio button that describes the reporter type: Reporter ID – Summarizes expression levels for all reporters you specify. Gene Name – Summarizes expression levels at the gene level. |

Table 5.5 Fields for selecting gene expression plot values based upon genomic queries

- Click the **Create Plot** button. calIntegrator2 generates the plot which then displays below the Gene Expression Plot criteria. Legends below the plot indicate the plot input (*Figure 5.10*).



Figure 5.10 A gene expression plot (Mean) based on a genomic query.

- You can recalculate the data display by clicking the **Plot Type** above the graph. See *Understanding a Gene Expression Plot* on page 70. Legends below the plot indicate the plot input.
- You can modify the plot parameters and click the **Reset** button to recalculate the plot.

Gene Expression Value Plot for Clinical Queries



Data to be analyzed on this tab must have been saved as a clinical query, but it must have genomic data identified in the query. For more information, see *Adding/Editing Genomic Data* on page 24.

To generate a gene expression plot using a clinical query, follow these steps:

- Select the study whose data you want to analyze in the upper right portion of the calIntegrator page. You must select a study saved as a clinical study, but which has genomic data.
- Under Analysis Tools on the left sidebar, select **Gene Expression Plot**.

3. Select the **For Clinical Queries** tab (*Figure 5.11*).

Clinical Query Based Gene Expression Plots (draft)

1.) Gene Symbol(s) (comma separated list):  

2.) Select Reporter Type: ☒ Reporter Id ☐ Gene

3.) Select Queries:


All Available Queries

Selected Queries

Add > < Remove

4.) ☐ Exclusive Subjects in Queries (Subjects in upper queries are removed from subsequent queries)

5.) ☐ Add additional group containing all other subjects not found in selected queries.

6.) ☐ Add additional group containing all control samples for this study. 

Reset Create Plot

Figure 5.11 Gene expression value tab for configuring gene expression clinical queries plot

4. Enter or select fields as described in [Table 5.6](#).

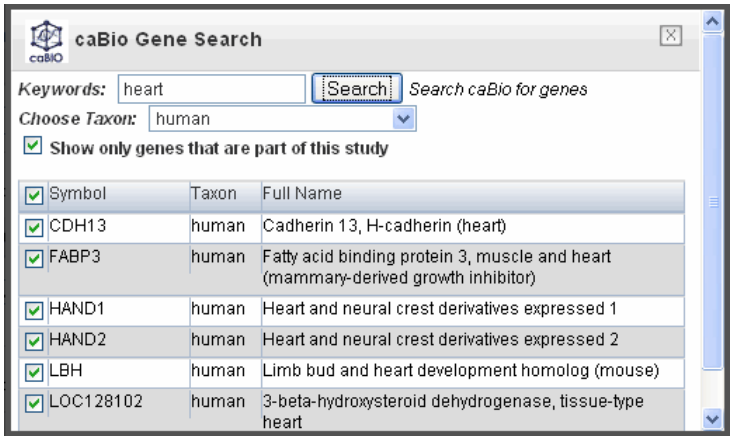


| Field | Description |
|---------------|--|
| Gene Symbol | <p>Enter one or more gene symbols in the text box or click the icons to locate genes in the following databases. If entering more than one gene in the text box, separate entries by commas.</p> <p>CGAP – Use this directory to identify genes. Before clicking this link you must enter gene symbols in the text box. This link does not pull anything into caIntegrator2 but does provide information about the gene(s).</p> <p>caBio – This link pulls identified genes into caIntegrator2 for analysis. Click the icon, enter Keyword(s) in the text box that opens and click Search.</p>  <p>Use the check boxes to identify the genes whose symbols you want to use in the gene expression analysis.</p> <p>Click the Use Genes button at the bottom of the page. This pulls the checked genes into the Gene Expression ... tab.</p> <p>Annotation Based Gene Expression Plots</p> <p>1.) Gene Symbol(s) (comma separated list): <input type="text" value="CDH13,FABP3,HAND1,HA"/>  </p> |
| Reporter Type | <p>Select the radio button that describes the reporter type:</p> <p>Reporter ID – Summarizes expression levels for all reporters you specify.</p> <p>Gene Name – Summarizes expression levels at the gene level.</p> |

Table 5.6 Fields for selecting gene expression plot values based upon clinical queries

| Field | Description |
|--|---|
| Sample Groups | <p>Annotation Type – Select the annotation type. Selections are based on the data in the chosen study</p> <p>Annotation – Select an annotation. Fields are based on the annotation type you select. For example, if you choose Subject, then you could select Gender or Radiation Type or any field that would distinguish the patients into groups based upon its values.</p> <p>Values – Using conventional selection techniques, select two or more values which will be the basis for the K-M plot. Permissible (available) values or “No Values” correspond to the selected annotation.</p> |
| Add additional group...all other subjects | Check the box to create an additional group of all other subjects that are not in selected query groups. |
| Add additional group...control group | Check the box to create an additional group of control samples for this study.. |

Table 5.6 Fields for selecting gene expression plot values based upon clinical queries

- Click the **Create Plot** button. By default, calIntegrator2 generates the plot which displays the mean of the data below the Gene Expression Plot criteria. Legends below the plot indicate the plot input.

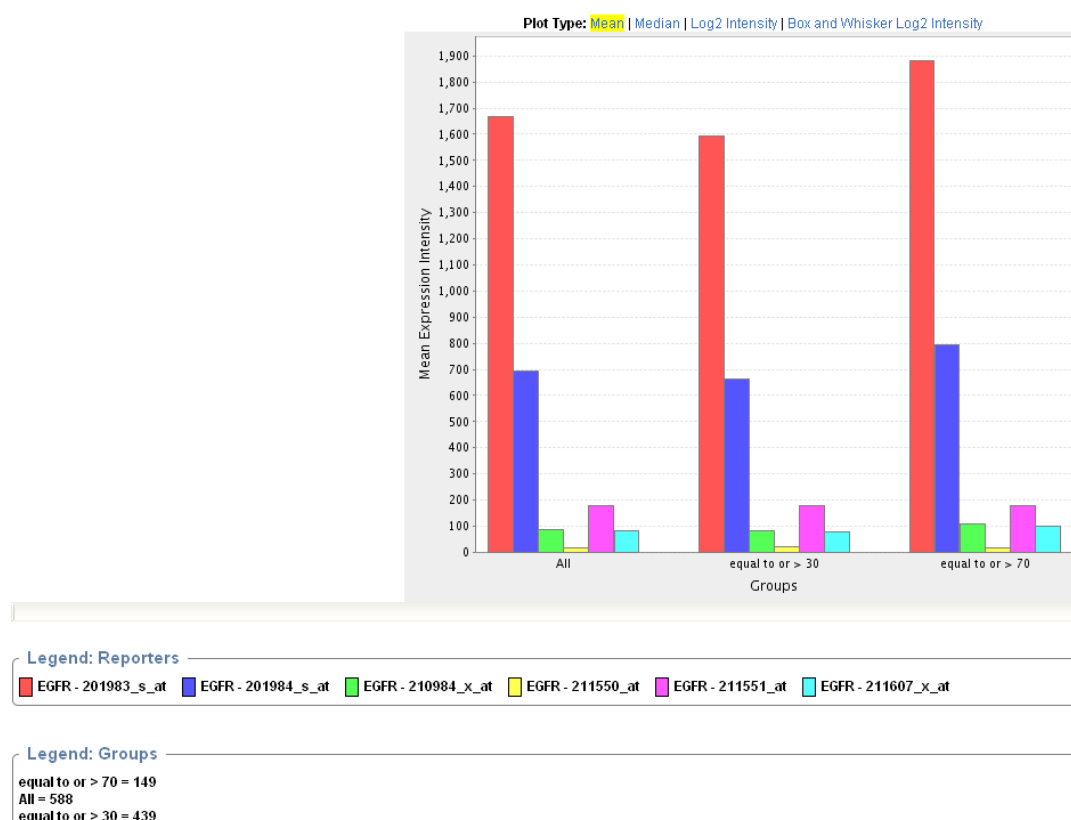


Figure 5.12 Gene expression plot based on clinical queries

6. You can recalculate the data display by clicking the **Plot Type** above the graph. See [Understanding a Gene Expression Plot](#) on page 70.
7. You can modify the plot parameters and click the **Reset** button to recalculate the plot.

Understanding a Gene Expression Plot

Above the plot, you can select various plot types. When you do so, the plot is recalculated. Although all of the plots in this section appear similar, note the differences in calculation results and legends between the Y axis on each of the plots.

When you perform a Gene Expression simple search, by default the **Mean** Gene Expression Plot ([Figure 5.13](#)) appears.

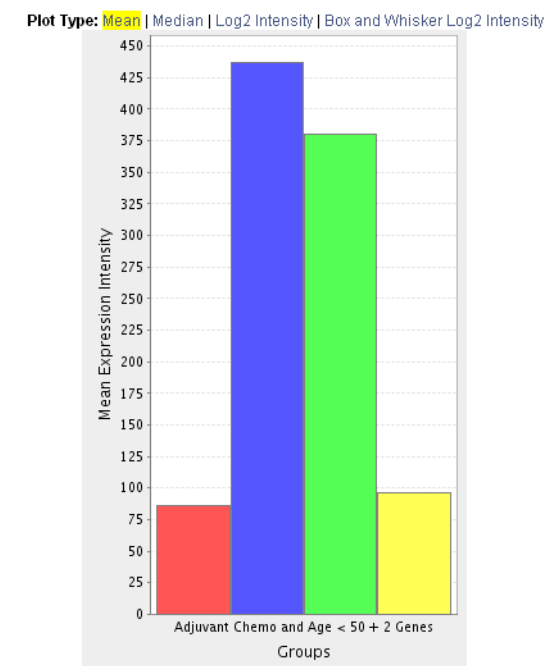


Figure 5.13 Gene expression plot calculating the mean

The **Mean** Gene Expression Plot ([Figure 5.13](#)) displays mean expression intensity (Geometric mean) versus Groups.

The **Median** Gene Expression Plot ([Figure 5.14](#)) displays the median expression intensity versus Groups..

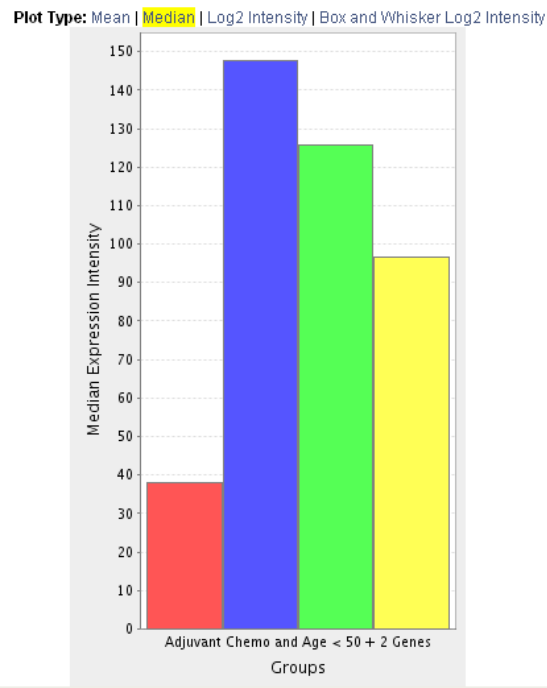


Figure 5.14 Gene expression plot calculating the median

The **Log2 Intensity** Gene Expression Plot ([Figure 5.15](#)) displays average expression intensities for the gene of interest based on Affymetrix GeneChip arrays (U133 Plus 2.0 arrays).

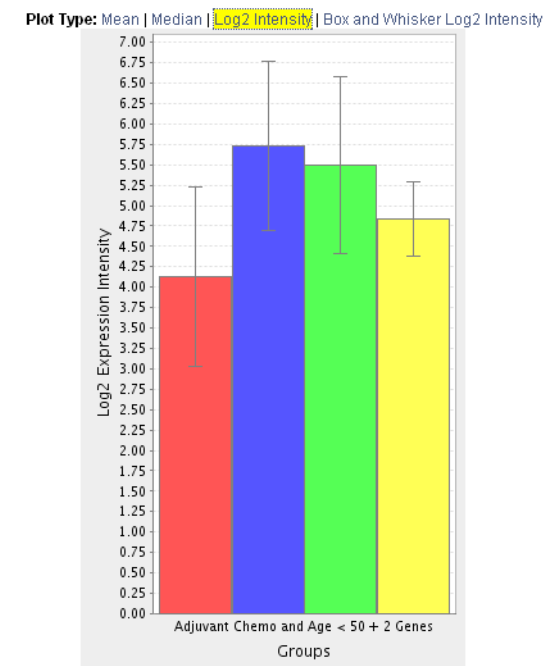


Figure 5.15 Gene expression plot displaying log2 intensity values

The box and whisker log2 expression intensity plot displays a box plot ([Figure 5.16](#), [Figure 5.17](#)). Example uses of box and whisker plots include the following:

- Indicate whether a distribution is skewed and whether there are potential unusual observations (outliers) in the data set.
- Perform a large number of observations.
- Compare two or more data sets.
- Compare distributions because the center, spread, and overall range are immediately apparent.

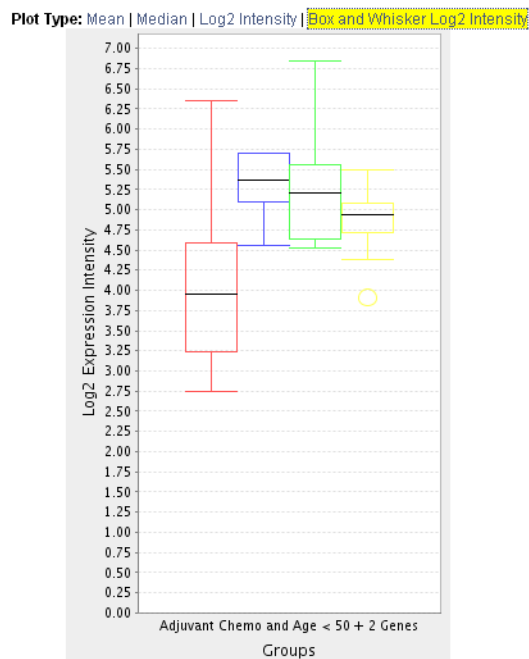


Figure 5.16 Box and whisker plot based on the same data set as represented in [Figure 5.13](#), [Figure 5.14](#), [Figure 5.15](#)

In descriptive statistics, a box plot or boxplot, also known as a box-and-whisker diagram or plot, is a convenient way of graphically depicting groups of numerical data through their five-number summaries (the smallest observation excluding outliers, lower quartile [Q1], median [Q2], upper quartile [Q3], and largest observation excluding outliers).

The box is defined by Q1 and Q3 with a line in the middle for Q2. The interquartile range, or IQR, is defined as $Q3 - Q1$. The lines above and below the box, or 'whiskers', are at the largest and smallest non-outliers. Outliers are defined as values that are

more than $1.5 \times \text{IQR}$ greater than Q3 and less than $1.5 \times \text{IQR}$ than Q1. Outliers, if present, are shown as open circles (*Figure 5.17*).

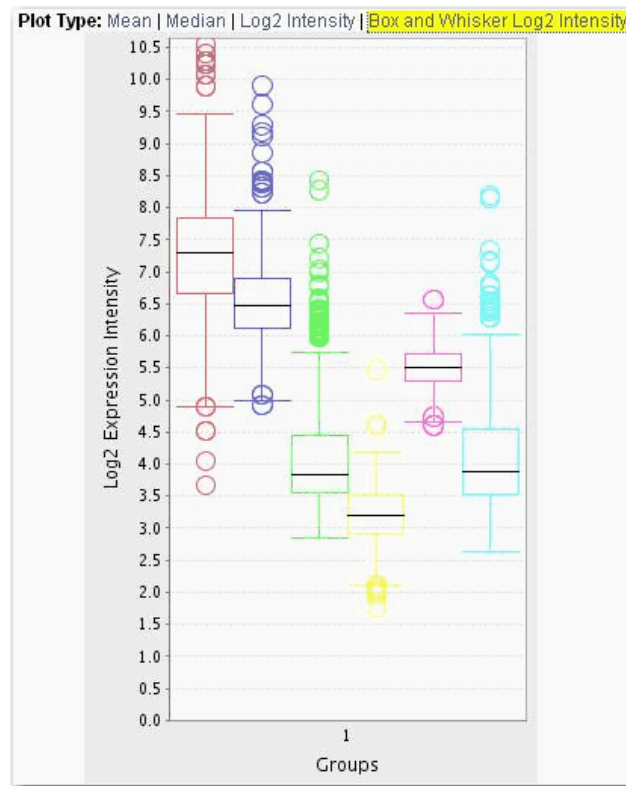


Figure 5.17 Box and whisker plot showing outliers

Boxplots can be useful to display differences between populations without making any assumptions of the underlying statistical distribution: they are non-parametric. The spacings between the different parts of the box help indicate the degree of dispersion (spread) and skewness in the data.

Analyzing Data with GenePattern

GenePattern is an application developed at the Broad Institute that enables researchers to access various methods to analyze genomic data. caIntegrator2 provides an express link to GenePattern where you can analyze data in any caIntegrator2 study.

Information is included in this section for connecting to GenePattern from caIntegrator2. Specifics for launching GenePattern tools from caIntegrator2 are included as well, but you may want to refer to additional GenePattern documentation, available at this website: http://www.broadinstitute.org/cancer/software/genepattern/tutorial/gp_concepts.html.

You have two options for using GenePattern from calIntegrator2:

- Option 1 – Use the web-interface of any available GenePattern instances.
 - a. To use the public instance from Broad, first register for an account at <http://genepattern.broad.mit.edu/gp/pages/login.jsf>
 - b. In calIntegrator2, enter the URL for connecting: <http://genepattern.broad.mit.edu/gp/services/>, then enter your userId and password.
- Option 2 – Use GenePattern on the grid.

The GenePattern feature in calIntegrator2 currently supports three analyses on the grid: Comparative Marker Selection (CMS), Principal Component Analysis (PCA) and GISTIC-supported analysis.

Tip: If you are using the web interface to access GenePattern (option #1 listed above), then you can run other GenePattern tools in addition to CMS, PCA and GISTIC.

1. Select the study whose data you want to analyze in the upper right portion of the calIntegrator2 page.
2. Click **GenePattern Analysis** in the left sidebar of calIntegrator2. This opens the GenePattern Analysis Status page (*Figure 5.18*).

GenePattern Analysis Status

| Gene Pattern Modules | | New Analysis Job | | |
|----------------------|------------------------------|--------------------------------------|---------------------|---------|
| Job Name | Job Type | Status | Creation Date | Status |
| JP - CMS - 2 | Comparative Marker Selection | Completed - Download | 2009/08/26 21:38:03 | 2009/08 |
| CMS 1 | Comparative Marker Selection | Completed - Download | 2009/08/26 11:43:44 | 2009/08 |
| PCA1 | Principal Component Analysis | Completed - Download | 2009/08/26 11:38:41 | 2009/08 |

Figure 5.18 GenePattern Analysis Status page

3. Select from the drop-down list the type of GenePattern analysis you want to run on the data.
 - **GenePattern Modules** – This option launches a session within GenePattern from which you can launch analyses. See [GenePattern Modules](#) on page 75.
 - **Comparative Marker Selection (Grid Service)**. This option enables you to run this GenePattern analysis on the grid. See [Comparative Marker Selection \(CMS\) Analysis](#) on page 76.
 - **Principal Component Analysis (Grid Service)**. This option enables you to run this GenePattern analysis on the grid. See [Principal Component Analysis \(PCA\)](#) on page 78.
 - **GISTIC (Grid Service)**. This option enables you to run this GenePattern analysis on the grid. See [GISTIC-Supported Analysis](#) on page 81.

- Click the **New Analysis Job** button to open a corresponding page where you can configure the analysis parameters.

GenePattern Modules

Note: To launch the analyses described in this section, you must have a registered GenePattern account. For more information, see <http://genepattern.broad.mit.edu/gp/pages/login.jsf>.

To configure the link for accessing GenePattern from caIntegrator2, open the appropriate page as described in *Analyzing Data with GenePattern* on page 73.

- Select the study whose data you want to analyze in the upper right portion of the caIntegrator2 page.
- Click **GenePattern Analysis** in the left sidebar of caIntegrator2. This opens the GenePattern Analysis Status page.
- Make sure **GenePattern Modules** is selected in the drop down list. Click **New Analysis Job**.
- In the GenePattern Analysis dialog box (*Figure 5.19*), specify connection information, , described *Table 5.7*.

GenePattern Analysis

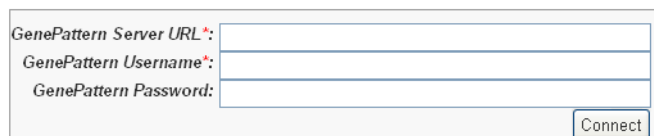


Figure 5.19 Dialog box for configuring the link to GenePattern

| Fields | Description |
|-----------------------------|---|
| Server URL | Enter any GenePattern publicly available URL, such as http://genepattern.broad.mit.edu/gp/services/Analysis . |
| GenePattern Username | Enter your GenePattern user name. |
| GenePattern Password | Enter your GenePattern password. |

Table 5.7 Fields for selecting GenePattern configurations

If you choose to access GenePattern in this way, you can continue to use GenePattern tools from within that application. See GenePattern user documentation for more information.

Tip: If you run these analysis within GenePattern itself, you may be able to view results in the GenePattern visualization module. If you run them on the grid from caIntegrator2, your results will be available only in spreadsheet and XML format.

You can run GenePattern analyses for Comparative Marker Selection, Principal Component Analysis and GISTIC-based analysis on the grid if you choose.

Comparative Marker Selection (CMS) Analysis

The Comparative Marker Selection (CMS) module implements several methods to look for expression values that correlate with the differences between classes of samples. Given two classes of samples, CMS finds expression values that correlate with the difference between those two classes. If there are more than two classes, CMS can perform one-vs-all or all-pairs comparisons, depending on which option is chosen.

For more information, see the GenePattern website: http://www.broad.mit.edu/cgi-bin/cancer/software/genepattern/modules/gp_modules.cgi.

To perform a CMS analysis, follow these steps:

1. Select the study whose data you want to analyze in the upper right portion of the calIntegrator2 page. You must select a study saved as a clinical study, but which has genomic data.
2. Click **GenePattern Analysis** in the left sidebar of calIntegrator2. This opens the GenePattern Analysis Status page.
3. In the GenePattern Analysis Status page, select **Comparative Marker Selection (Grid Service)** from the drop down list and click **New Analysis Job**. This opens the Comparative Marker Selection Analysis page (*Figure 5.20*).

Comparative Marker Selection Analysis

Job Name*:

Preprocess Server*: Default Broad service - <http://node255.broad.mit.edu:6060/wsrf/services/cagrid/PreprocessDatasetMAGEService> ▼

Comparative Server*: Default Broad service - <http://node255.broadinstitute.org:11010/wsrf/services/cagrid/ComparativeMarkerSelMAGESvc> ▼

Clinical Queries*: Must select two clinical queries, which are used to group the samples into two separate classifications to run against ComparativeMarkerSelection. The queries selected here have been previously saved by the user. Selected queries will result in the processing of only those samples which are mapped to patients in the saved query result.

All Available Queries

Selected Queries

Add > < Remove

Filter flag: ☐

Preprocessing Flag*: no-disc-or-norm ▼

Min Change*:

Min Delta*:

Threshold*:

Ceiling*:

Max Sigma Binning*:

Probability Threshold*:

Num Exclude*:

Log Base Two: ☐

Number Of Columns Above Threshold*:

Test Direction*: two-sided ▼

Test Statistic*: T-test ▼

Min Std*:

Number Of Permutations*:

Complete: ☐

Balanced: ☐

Random Seed*:

Smooth Pvalues: ☐

Phenotype Test*: one-versus-all ▼

Perform Analysis

Figure 5.20 Comparative Marker Selection analysis parameters

4. Select or define CMS analysis parameters, described in [Table 5.8](#). An asterisk indicates required fields. The default settings are valid; they should provide valid results.

| CMS Parameter | Description |
|---|--|
| Job Name* | Assign a unique name to the analysis you are configuring. |
| Preprocess Server* | A server which hosts the grid-enabled data GenePattern PreProcess Dataset module. Select one from the list and calIntegrator2 will use the selected server for this portion of the processing. |
| Comparative Server* | A server which hosts the grid-enabled data GenePattern Comparative Marker Selection module. Select one from the list and calIntegrator2 will use the selected server for this portion of the processing. |
| Clinical Queries* | All clinical queries with appropriate data for the analysis are listed. Select and move 2 or more queries from the All Available Queries panel to the Selected Queries panel. Note: If a query has a genomic component (e.g. gene criteria), it does not display in the queries field. |
| Filter Flag | Variation filter and thresholding flag |
| Preprocessing Flag* | Discretization and normalization flag |
| Min Change* | Minimum fold change for filter |
| Min Delta* | Minimum delta for filter |
| Threshold* | Value for threshold |
| Ceiling* | Value for ceiling |
| Max Sigma Binning* | Maximum sigma for binning |
| Probability Threshold* | Value for uniform probability threshold filter |
| Num Exclude* | Number of experiments to exclude (max & min) before applying variation filter |
| Log Base Two | Whether to take the log base two after thresholding |
| Number of Columns Above Threshold* | Remove row if n columns no \geq than the given threshold |
| Test Direction* | The test to perform (up-regulated for class0; up-regulated for class1, two sided). By default, Comparative Marker Selection performs the two-sided test. |
| Test Statistic* | Select the statistic to use. |
| Min Std* | The minimum standard deviation if test statistic includes the min std option. Used only if test statistic includes the min std option. |

Table 5.8 Comparative Marker Selection analysis options

| CMS Parameter | Description |
|--------------------------------|---|
| Number of Permutations* | <p>The number of permutations to perform. (Use 0 to calculate asymptotic P-values.) The number of permutations you specify depends on the number of hypotheses being tested and the significance level that you want to achieve (3). The greater the number of permutations, the more accurate the P-value.</p> <p>Complete – Perform all possible permutations. By default, complete is set to No and Number of Permutations determines the number of permutations performed. If you have a small number of samples, you might want to perform all possible permutations.</p> <p>Balanced – Perform balanced permutations</p> |
| Random Seed* | The seed for the random number generator. |
| Smooth Pvalues | Whether to smooth P-values by using the Laplace's Rule of Succession. By default, Smooth Pvalues is set to Yes , which means P-values are always less than 1.0 and greater than 0.0. |
| Phenotype Test* | <p>Tests to perform when class membership has more than 2 classes: one versus-all, all pairs.</p> <p>Note: The P-values obtained from the one-versus-all comparison are not fully corrected for multiple hypothesis testing.</p> |

Table 5.8 Comparative Marker Selection analysis options

- When you have completed the form, click **Perform Analysis**.

calIntegrator2 takes you to the JobStatus/Launch page where you will see the job and its status in the Status column of the list ([Figure 5.21](#)).

GenePattern Analysis Status (draft)

Gene Pattern Modules


| Job Name | Job Type | Status | Creation Date | Status Update Date |
|----------------------------------|------------------------------|--|---------------------|---------------------|
| Well-diff vs adjuvant chemo | Comparative Marker Selection |  Processing Locally | 2009/08/14 11:48:35 | 2009/08/14 11:48:35 |
| Filter out non-interesting genes | Gene Pattern | Completed - View 122444 | 2009/08/14 10:16:29 | 2009/08/14 10:19:47 |

Figure 5.21 The progress of a GenePattern analysis that has been launched displays in the status column of page

- When the job is complete, the system displays a completion date on the GenePattern Analysis status page. Click the **Download** link. This downloads zipped result files to your local work station. The number of files and their file type will vary according to the processing. The results format is compatible with GenePattern visualizers and can be uploaded within GenePattern.

Principal Component Analysis (PCA)

Principal Component Analysis is typically used to transform a collection of correlated variables into a smaller number of uncorrelated variables, or components. Those components are typically sorted so that the first one captures most of the underlying variability and each succeeding component captures as much of the remaining variability as possible.

You can configure GenePattern grid parameters for preprocessing the dataset in addition to PCA module parameters. For more information, see the GenePattern website: http://www.broad.mit.edu/cgi-bin/cancer/software/genepattern/modules/gp_modules.cgi.

To perform a PCA analysis, follow these steps:

1. Select the study whose data you want to analyze in the upper right portion of the calIntegrator page. You must select a study with gene expression data.
2. Click **GenePattern Analysis** in the left sidebar of calIntegrator2. This opens the GenePattern Analysis Status page.
3. In the GenePattern Analysis Status page, select **Principal Component Analysis (Grid Service)** from the drop down list and click **New Analysis Job**. This opens the Principal Component Analysis page (*Figure 5.22*).

Principal Component Analysis

(draft)

This form submits a job which analyzes samples using the GenePattern Principal Component Analysis module.

Job Name - Please enter a job name.
Principal Component Analysis Server - Select a PCA grid service from the dropdown.
Clinical Queries - Select saved Clinical queries to specify which samples will be processed.
Enable Preprocess Dataset - (Optional) Check this to display and configure preprocessing parameters.

* Job Name:

* Principal Component Analysis Server: Default Broad service - <http://node255.broad.mit.edu:8060/awstf/services/cagrid/PCA>

* Clinical Queries: Clinical Queries enable the user to specify which samples will be processed using PCA. The queries selected here have been previously saved by the user. Selected queries will result in the processing of only those samples which are mapped to subjects in the saved query result. If multiple queries are selected, all of the sample from each saved query are processed PLUS the results set will be classified according to those queries. (One class per selected query.)

All Available Queries

- gender female
- Never smoke

Selected Queries

Add >

< Remove

Enable Preprocess Dataset: ☐
(check to display preprocess parameters)

Perform Analysis

Figure 5.22 Principal Component Analysis parameters

4. Select or define PCA analysis parameters, described in *Table 5.9*. You must enter a job name and select a clinical query, but you can accept the other default settings..

| PCA Parameters | Description |
|---|---|
| Job Name* | Assign a unique name to the analysis you are configuring. |
| Principal Component Analysis Server* | A server which hosts the grid-enabled data GenePattern Principal Component Analysis module. Select one from the list and calIntegrator2 will use the selected server for this portion of the processing. |
| Clinical Queries* | All clinical queries display in this list. Select one or more of these queries to define which samples are analyzed using PCA. If you select more than one query, then the union of the samples returned by the multiple queries is analyzed. |

Table 5.9 PCA analysis options

| PCA Parameters | Description |
|----------------|--|
| Cluster By* | Selecting rows looks for principal components across all expression values, and selecting columns looks for principal components across all samples. |

Table 5.9 PCA analysis options

- If you want to preprocess the data set, click **Enable the Preprocess Dataset**. This opens an additional set of parameters (Figure 5.23), discussed in Table 5.10. The preprocessing is executed prior to running the PCA.

Figure 5.23 Parameters for pre-processing parameters for PCA

| PCA Preprocessing Parameters | Description |
|------------------------------|---|
| Preprocess Server* | A server which hosts the grid-enabled data GenePattern PreProcess Dataset module. Select one from the list and caIntegrator2 will use the selected server for this portion of the processing. |
| Filter Flag | Variation filter and thresholding flag |
| Preprocessing Flag | Discretization and normalization flag |
| Min Change | Minimum fold change for filter |
| Min Delta | Minimum delta for filter |
| Threshold | Value for threshold |
| Ceiling | Value for ceiling |
| Max Sigma Binning | Maximum sigma for binning |
| Probability Threshold | Value for uniform probability threshold filter |

Table 5.10 Parameters for preprocessing data sets for PCA

| PCA Preprocessing Parameters | Description |
|--|---|
| Num Exclude | Number of experiments to exclude (max & min) before applying variation filter |
| Log Base Two | Whether to take the log base two after thresholding |
| Number of Columns Above Threshold | Remove row if n columns no \geq than the given threshold |

Table 5.10 Parameters for preprocessing data sets for PCA

- When you have completed the form, click **Perform Analysis**.
- When the job is complete, the system displays a completion date on the GenePattern Analysis status page. Click the **Download** link. This downloads zipped result files to your local work station. The number of files and their file type will vary according to the processing. The results format is compatible with GenePattern visualizers and can be uploaded within GenePattern.

GISTIC-Supported Analysis

The GISTIC Module is a GenePattern tool that identifies regions of the genome that are significantly amplified or deleted across a set of samples. For more information, see http://www.broad.mit.edu/cgi-bin/cancer/software/genepattern/modules/gp_modules.cgi.

To perform a GISTIC-supported analysis, follow these steps:

- Select the study whose data you want to analyze in the upper right portion of the calIntegrator2 page. You must select a study with copy number (either Affymetrix SNP or Agilent Copy Number) data.
- Click **GenePattern Analysis** in the left sidebar of calIntegrator2. This opens the GenePattern Analysis Status page.

3. In the GenePattern Analysis Status page, select **GISTIC (Grid Service)** from the drop down list and click **New Analysis Job**. This opens the GISTIC Analysis page (*Figure 5.24*).

GISTIC Analysis

This form submits a job which analyzes samples using the GenePattern GISTIC module.

Job Name - Please enter a job name.

GenePattern Server URL / GISTIC Server - Select whether to use the GISTIC web service or grid service and provide or select the service address. If the web service is selected, authentication information is also required.

Clinical Queries - (Optional) Select a saved Clinical query to specify which samples will be processed.

Exclude Sample Control Set - (Optional) Select a Control Sample Set to be excluded from the Clinical Query.

* Job Name:

* GenePattern Server URL:

* GenePattern Username:

GenePattern Password:

* GISTIC Server:

Use GenePattern GISTIC Web Service ☒

Use GISTIC Grid Service ☐

For the Clinical query parameter below, choose either "All Samples" or a clinical query. If "All Samples" is selected, then all samples will be used. If a clinical query is selected, only those samples which map to the subjects in the clinical query results will be used. The clinical queries in this list have been previously saved by the user. Control samples can be excluded from this processing by selecting a control set name in the Exclude Sample Control Set dropdown.

Clinical query:

* Exclude Sample Control Set:

* Amplifications Threshold:

* Deletions Threshold:

* Join Segment Size:

* QV Threshold:

* Remove X:

cnv File:

Figure 5.24 GISTIC analysis criteria

4. Select or define GISTIC analysis parameters, as described in *Table 5.8*. You must indicate a Job Name, but you can accept the other defaults settings, which are valid and should produce valid results.

| GISTIC Parameters | Description |
|----------------------------------|--|
| Job Name* | Assign a unique name to the analysis you are configuring. |
| GISTIC Server* | A server which hosts the grid-enabled data GISTIC-based analysis module. Select one from the list and caIntegrator2 will use the selected server for this portion of the processing. |
| Refgene File* | Enter or select the cytoband file to use in the analysis. Allowed values: {Human Hg18, Human Hg17, Human Hg16}. Default = Human Hg16. |
| Clinical Query | All clinical queries display in this list as well as an option to select all non-control samples. Select a clinical query if you wish to run GISTIC on a subset of the data and select all non-control samples if wish to include all samples. |
| Amplifications Threshold* | Threshold for copy number amplifications. Regions with a log2 ratio above this value are considered amplified. Default = 0.1. |
| Deletions Threshold* | Threshold for copy number deletions. Regions with a log2 ratio below the negative of this value are considered deletions. Default = 0.1. |

Table 5.11 GISTIC analysis parameters

| GISTIC Parameters | Description |
|---------------------------|---|
| Join Segment Size* | Smallest number of markers to allow in segments from the segmented data. Segments that contain fewer than this number of markers are joined to the neighboring segment that is closest in copy number. Default = 4. |
| QV Thresh[hold]* | Threshold for q-values. Regions with q-values below this number are considered significant. Default = 0.25. |
| Remove X* | Flag indicating whether to remove data from the X-chromosome before analysis. Allowed values = {1,0}. Default = 1(yes). |
| cnv File | <p>This selection is optional.</p> <p>Browse for the file. There are two options for the cnv file.</p> <p>Option #1 enables you to identify CNVs by marker name. Permissible file format is described as follows:</p> <p>A two column, tab-delimited file with an optional header row. The marker names given in this file must match the marker names given in the markers_file. The CNV identifiers are for user use and can be arbitrary. The column headers are:</p> <ol style="list-style-type: none"> 1. Marker Name 2. CNV Identifier <p>Option #2 enables you to identify CNVs by genomic location. Permissible file format is described as follows:</p> <p>A 6 column, tab-delimited file with an optional header row. The 'CNV Identifier', 'Narrow Region Start' and 'Narrow Region End' are for user use and can be arbitrary. The column headers are:</p> <ol style="list-style-type: none"> 1. CNV Identifier 2. Chromosome 3. Narrow Region Start 4. Narrow Region End 5. Wide Region Start 6. Wide Region End |

Table 5.11 GISTIC analysis parameters

5. When you have completed the form, click **Perform Analysis**.
6. When the job is complete, the system displays a completion date on the GenePattern Analysis status page. Click the **Download** link. This downloads zipped result files to your local work station. The number of files and their file type will vary according to the processing. The results format is compatible with GenePattern visualizers and can be uploaded within GenePattern.

CHAPTER 6

ADMINISTERING USER ACCOUNTS

This chapter describes the process for creating and managing user accounts in calIntegrator2. It also discusses the processes for managing ownership and access to studies in calIntegrator2.

Note: The options for performing user management tasks are visible in calIntegrator2 on the left sidebar of the browser only if you have these Admin privileges.

Administering calIntegrator2 User Accounts Using UPT

Note: If you are interested in registering an account in calIntegrator2, see *Registering as a New calIntegrator2 User* on page 6.

In calIntegrator2, all tasks related to creating and managing user accounts can be performed only by a calIntegrator2 administrator using the CBIIT User Provisioning Tool (UPT) v. 4.2. The following sections discuss the use of the UPT for performing these tasks. For further information about UPT, see Chapter 3 of the CSM 4.2 Programmer's Guide located here: https://gforge.nci.nih.gov/docman/view.php/12/18945/caCORE_CSM_v42_ProgrammersGuide.pdf

The UPT is a separately installed application which serves as the user management interface for all National Cancer Institute CBIIT Life Sciences Distribution (LSD) applications, including calIntegrator2. The UPT application is the central point for all user management functionality within calIntegrator2. You can use UPT to add new users and to apply user group assignments to the calIntegrator2 database directly. The UPT groups can refer to predefined groups such as Study Manager or Study Investigator, which determine what roles the user has.

The following terms are used both in this chapter and in the UPT to define user-related roles:

- **User** – a person who is accessing calIntegrator2. The user has an associated account and user id.

- **User Group** – a group of users, typically grouped by organization and role, for example, “Columbia University Study Managers”
- **Protection Group** – a group of studies given a secure status and typically grouped by organization, for example, “Columbia University Protected Studies”.

Steps for Creating User Access to caIntegrator2

The following steps summarize the process for establishing user access to caIntegrator2:

1. A potential user requests a user account in caIntegrator2. See [Registering as a New caIntegrator2 User](#) on page 6.
1. You, as a caIntegrator2 administrator, check if the **User** already exists in caIntegrator2. If not, create the new user. See [Creating a New caIntegrator2 User](#) on page 86.
2. Check if the requestor's **User Group** already exists in caIntegrator2. If not, create a new **User Group**. See [Creating a New User Group](#) on page 88.
3. Check if the **Protection Group** (e.g. “Columbia University Protected Studies”), containing the studies to which this user wants access currently exists. If not, create a new **Protection Group**. See [Creating a New Protection Group](#) on page 89.

Note: If the Protection Group already exists, contact the Organizational Contact person to confirm that it is OK to give this person access to this Protection Group.

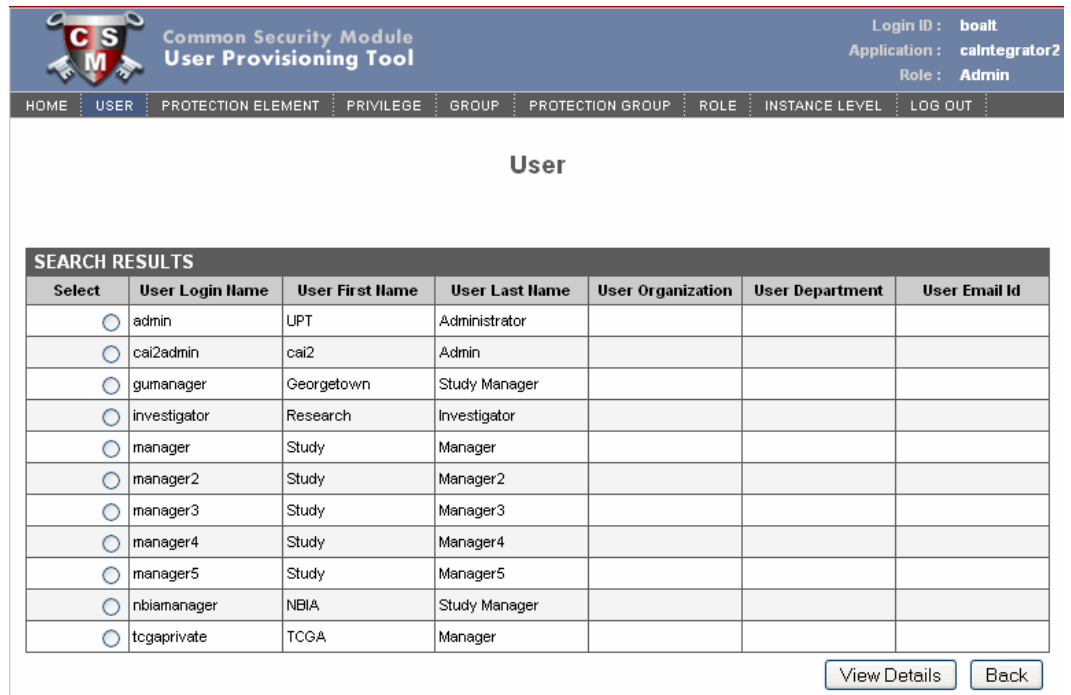
4. Give the requestor's **User Group** access to the **Protection Group**. See [Assigning a User Group to a Protection Group](#) on page 90.
5. Add the **User** to the **User Group**. See [Adding a User to a User Group](#) on page 93

Creating a New caIntegrator2 User

To create a new User in caIntegrator2, follow these steps:

1. Login to UPT as a caIntegrator2 Admin.
2. First, search to see if the user already exists. Click the **User** menu option.
3. On the User page that opens, click **Select an Existing User**.

- Use the form and search for the user. If you define no criteria, UPT returns a list of all caIntegrator2 users currently in the system (*Figure 6.1*).



The screenshot shows the UPT interface with a top navigation bar and a main content area. The top bar includes the CS M logo, the title 'Common Security Module User Provisioning Tool', and user information: Login ID: boalt, Application: caIntegrator2, and Role: Admin. Below this is a navigation menu with options: HOME, USER, PROTECTION ELEMENT, PRIVILEGE, GROUP, PROTECTION GROUP, ROLE, INSTANCE LEVEL, and LOG OUT. The main content area is titled 'User' and displays a 'SEARCH RESULTS' table. The table has columns for Select, User Login Name, User First Name, User Last Name, User Organization, User Department, and User Email Id. It lists 12 users, each with a radio button in the Select column. At the bottom right of the table are two buttons: 'View Details' and 'Back'.

| Select | User Login Name | User First Name | User Last Name | User Organization | User Department | User Email Id |
|-----------------------|-----------------|-----------------|----------------|-------------------|-----------------|---------------|
| <input type="radio"/> | admin | UPT | Administrator | | | |
| <input type="radio"/> | cai2admin | cai2 | Admin | | | |
| <input type="radio"/> | gumanager | Georgetown | Study Manager | | | |
| <input type="radio"/> | investigator | Research | Investigator | | | |
| <input type="radio"/> | manager | Study | Manager | | | |
| <input type="radio"/> | manager2 | Study | Manager2 | | | |
| <input type="radio"/> | manager3 | Study | Manager3 | | | |
| <input type="radio"/> | manager4 | Study | Manager4 | | | |
| <input type="radio"/> | manager5 | Study | Manager5 | | | |
| <input type="radio"/> | nbiamanager | NBIA | Study Manager | | | |
| <input type="radio"/> | tcgaprivate | TCGA | Manager | | | |

View Details Back

Figure 6.1 A list of current caIntegrator2 users displays in UPT after a user search

- If the user does not already exist (is not listed in the search results), then create a new user. To do so, select the **User** menu option again, then click **Create a New User**.

This opens the page for creating a new caIntegrator2 user (*Figure 6.2*).

Common Security Module
User Provisioning Tool

Login ID : boalt
Application : caIntegrator2
Role : Admin

HOME USER PROTECTION ELEMENT PRIVILEGE GROUP PROTECTION GROUP ROLE INSTANCE LEVEL LOG OUT

Enter the details to add a new User. The **User Login Name** uniquely identifies the User and is a required field. The **User First Name** and **User Last Name** identifies the User. The **User Organization**, **User Department** and **User Title** provides his work details. The **User Phone Number** and **User Email Id** provides the contact details for the User. The **User Password** can be entered if the same schema is also going to be used for Authentication. The **User Start Date** and **User End Date** determine the period for which the User is a valid User.

* indicates a required field

| ENTER THE NEW USER DETAILS | |
|----------------------------|---|
| * | User Login Name <input type="text"/> |
| * | User First Name <input type="text"/> |
| * | User Last Name <input type="text"/> |
| | User Organization <input type="text"/> |
| | User Department <input type="text"/> |
| | User Title <input type="text"/> |
| | User Phone Number <input type="text"/> |
| | User Password <input type="text"/> |
| | Confirm Password <input type="text"/> |
| | User Email Id <input type="text"/> |
| | User Start Date <input type="text"/> (MM/DD/YYYY) |
| | User End Date <input type="text"/> (MM/DD/YYYY) |

Add Reset Back

Figure 6.2 UPT page for creating new user details

6. Enter details for the following required fields:

- **User Login Name**
- **User First Name**
- **User Last Name**
- **User Password**

Caution: If the requestor is an LDAP user, then the User Login Name must match the LDAP login id AND the User Password field must be left blank. If the requestor is not an LDAP user, then provide a password.

- **User Organization**
- **User Department**

7. Click **Add** to confirm the new user.

Creating a New User Group

You can assign a user group to a protection group. The advantage of working with a user group is that you do not have to assign roles to each user individually. You can assign users to a user group to which you assign a role, and then assign that user group to the protection group, or you can assign a role collectively to a protection group after it is created.

To create a new User Group in calIntegrator2, follow these steps:

1. Login to UPT as calIntegrator2 Admin.
2. First search for an existing group that the user wishes to join. Click the **Group** menu option.
3. On the Group page that opens, click **Select an Existing Group**.
4. Use the form and search for the group. If you define no criteria, UPT returns a list of all calIntegrator2 groups currently in the system
5. If a user group does not already exist, then create a new user group. Click the **Group** menu option, then click **Create a new Group**.
6. On the form that opens (*Figure 6.3*), enter a unique **Group Name** and a description, if appropriate. Click **Add**.

Note: The recommended naming convention for a new User Group is *[insert organization name] Study [insert role]s* Example: "Columbia University Study Managers".

Figure 6.3 UPT page for creating a new group

Creating a New Protection Group

: If you prefer that a study or group of studies have limited access, you can assign a user to a particular protection group and assign roles which allow the users in the protection group study access. : A protection group provides security or limited access for studies listed there.

To create a new Protection Group in calIntegrator2, follow these steps:

1. Login to UPT as calIntegrator2 Admin.
2. Click the **Protection Group** menu option.

- On the page that opens, click **Create a New Protection Group**. The page opens for defining PG Group details (*Figure 6.4*).

Common Security Module
User Provisioning Tool

Login ID : boalt
Application : calintegrator2
Role : Admin

HOME USER PROTECTION ELEMENT PRIVILEGE GROUP PROTECTION GROUP ROLE INSTANCE LEVEL LOG OUT

Enter the details to add a new Protection Group. The **Protection Group Name** uniquely identifies the Protection Group and is a required field. The **Protection Group Description** is a brief summary about the Protection Group. The **Protection Group Large Count Flag** is used to indicate if the Protection Group has a large number of associated Protection Elements.

* indicates a required field

ENTER THE NEW PROTECTION GROUP DETAILS

* **Protection Group Name**

Protection Group Description

Protection Group Large Count Flag ☐ Yes ☒ No

Add Reset Back

Figure 6.4 UPT page for creating a new protection group

- Enter a unique **Protection Group Name** and Description, if appropriate. Click **Add**.

Note: The recommended naming convention is *[insert organization name here] Protected Studies*. Example: "Columbia University Protected Studies".

Assigning a User Group to a Protection Group

To give a User Group access to a Protection Group (a group of protected studies), follow these steps:

- Login to UPT as calIntegrator2 Admin.
- Find the user group that you want to assign. Click the **Group** menu option and click **Select an Existing Group**. In the page that opens, click **Search**. If you define no criteria, UPT returns a list of all calIntegrator2 groups currently in the system (*Figure 6.5*).

Group

| SEARCH RESULTS | | |
|-----------------------|-------------------------|---|
| Select | Group Name | Group Description |
| <input type="radio"/> | Study Managers Group 3 | Study Managers who can create/modify any Group 3 studies. |
| <input type="radio"/> | Study Managers Group 4 | Study Managers who can create/modify any Group 4 studies. |
| <input type="radio"/> | Study Managers Group 5 | Study Managers who can create/modify any Group 5 studies. |
| <input type="radio"/> | NCI Study Investigators | Study investigators for the NCI studies. |
| <input type="radio"/> | NCI Study Managers | Study Managers who can create/modify any NCI studies. |
| <input type="radio"/> | Platform Manager Group | The platform manager group. |
| <input type="radio"/> | TCGA Study Managers | Study Managers who can create/modify any TCGA studies. |

View Details Back

Figure 6.5 UPT page showing Group search results

3. Select the radio button next to the group name you want to assign to the Protection Group. Click **View Details**. This opens the Group Details page (Figure 6.6).

Common Security Module
User Provisioning Tool

Login ID : boalt
Application : calintegrator2
Role : Admin

HOME USER PROTECTION ELEMENT PRIVILEGE GROUP PROTECTION GROUP ROLE INSTANCE LEVEL LOG OUT

Update the details of the displayed Group. The **Group Name** uniquely identifies the Group and is a required field. The **Group Description** is a brief summary about the Group. The **Update Date** indicates the date when this Group's Details were last updated.

| GROUP DETAILS | |
|-------------------|---|
| * Group Name | NCI Study Managers |
| Group Description | Study Managers who can create/modify any NCI studies. |
| Group Update Date | 09/24/2009 (MM/DD/YYYY) |

Update Delete Back

Associated Users Associated PE & Privileges Associated PG & Roles Assign PG & Roles

Figure 6.6 UPT page showing details for a selected group

4. Below the group details, click **Associated PG & Roles**. The page that opens displays any PG to which the user group is already assigned (Figure 6.7).

Group, Protection Group and Roles

SELECTED GROUP

| Group Name |
|--------------------|
| NCI Study Managers |

Select the **Protection Group** association which to be removed for the selected **Group** or whose **Roles** Association needs to be updated.

| SEARCH RESULTS | | |
|-----------------------|----------------------------------|----------------------|
| Select | Associated Protection Group Name | Associated Role Name |
| <input type="radio"/> | NCI Protected Studies | STUDY_MANAGER_ROLE |

Remove PG & Roles Associated Roles Back

Figure 6.7 UPT page that shows any PGs to which the select user group is assigned

5. Below the group name, examine if the Protection Group of your choice is already listed there. If so, this means your user group is already assigned to the protection group of choice, and you can skip the remainder of the steps in this section. If the Protection Group is not listed there, then click **Back**.

6. Back on the User Group details page, click **Assign PG & Roles**. This opens the Group, Protection Group and Roles Association page ([Figure 6.8](#)).

Group, Protection Group and Roles Association

| SELECTED GROUP | |
|-------------------|--------------------|
| Group Name | NCI Study Managers |

Select a single **Protection Group** to associate with the selected **Group**.

| AVAILABLE PROTECTION GROUPS |
|-------------------------------|
| Platforms |
| Protected Studies for Group 3 |
| Protected Studies for Group 4 |
| Protected Studies for Group 5 |
| TCGA Protected Studies |

| ASSIGNED PROTECTION GROUP |
|---------------------------|
| |

Select **Roles** which are to be associated with the selected **Group**.

| AVAILABLE ROLES |
|-------------------------|
| PLATFORM_MANAGER_ROLE |
| STUDY_INVESTIGATOR_ROLE |
| STUDY_MANAGER_ROLE |

| ASSIGNED ROLES |
|----------------|
| |

Figure 6.8 UPT page for assigning user group to a protection group and selected roles

7. From the list of Available Protection Groups, highlight your PG of choice and click **Assign**.

Now you can assign a role to the user. The caIntegrator2 Roles are defined in [Table 6.1](#):

| Role Name | Role Definition |
|--------------------|--|
| STUDY_MANAGER_ROLE | Assigning this role allows the user to modify existing studies, create new studies, and deploy existing studies. |

Table 6.1 Names and definitions for caIntegrator2 roles

| Role Name | Role Definition |
|-------------------------|--|
| STUDY_INVESTIGATOR_ROLE | Assigning this role allows the user to search the study, save queries about the study and perform analyses. |
| PLATFORM_MANAGER_ROLE | Assigning this role allows the user to create and delete array platforms for the entire calIntegrator2 installation. Caution: Array platforms are shared by all users and studies in the calIntegrator2 installation. A user with this role can affect the platforms that are used by all users and studies in the calIntegrator2 installation. |

Table 6.1 Names and definitions for calIntegrator2 roles

8. If this user group is a group of study managers, then select STUDY_MANAGER_ROLE. If this user group is a group of study investigators, then select STUDY_INVESTIGATOR_ROLE. Click **Assign**.
9. Click **Update Association** at the bottom of the page. This completes the assigning of the user group to the protection group you chose.

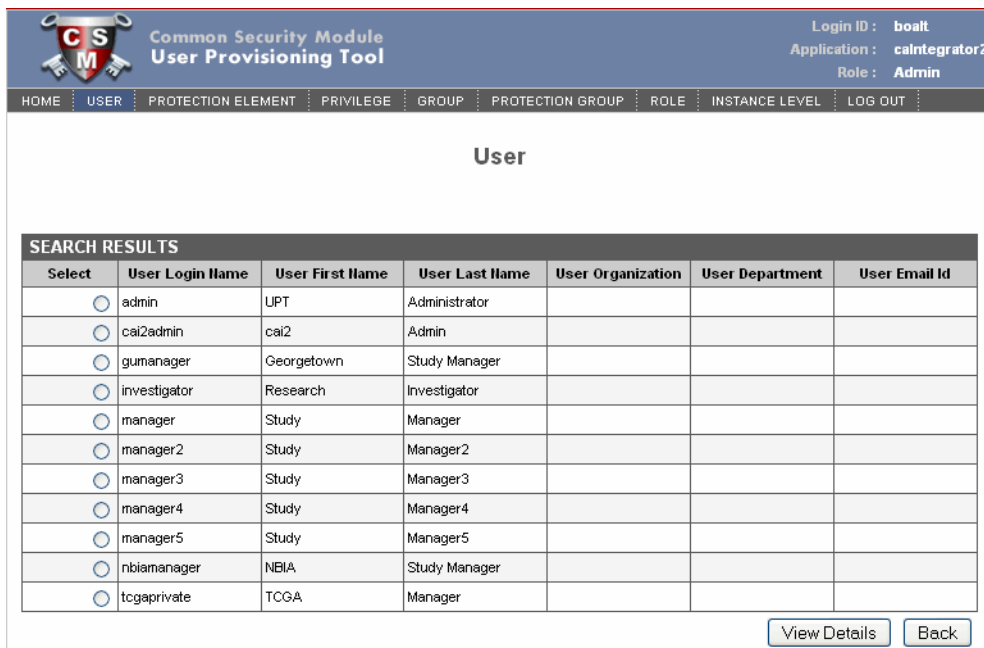
Note: If a **User** has the STUDY_MANAGER_ROLE role for more than one **Protection Group**, then any study that the **User** creates will be assign to each of those **Protection Groups**.

Adding a User to a User Group

To add a user to an existing user group, follow these steps:

1. Login to UPT as calIntegrator2 Admin.
2. Find the user that you want to assign to a user group. Click the **User** menu option, then click **Select an Existing User**.

- Enter the name of the user you are looking for and click **Search**. If you define no criteria, UPT returns a list of all calIntegrator2 users currently in the system (Figure 6.10).



The screenshot shows the 'Common Security Module User Provisioning Tool' interface. The top navigation bar includes links: HOME, USER, PROTECTION ELEMENT, PRIVILEGE, GROUP, PROTECTION GROUP, ROLE, INSTANCE LEVEL, and LOG OUT. The user's session information is displayed as: Login ID: boalt, Application: calIntegrator2, Role: Admin.


The main section is titled 'User' and displays 'SEARCH RESULTS' in a table. The table has the following columns: Select, User Login Name, User First Name, User Last Name, User Organization, User Department, and User Email Id.

| Select | User Login Name | User First Name | User Last Name | User Organization | User Department | User Email Id |
|-----------------------|-----------------|-----------------|----------------|-------------------|-----------------|---------------|
| <input type="radio"/> | admin | UPT | Administrator | | | |
| <input type="radio"/> | cal2admin | cal2 | Admin | | | |
| <input type="radio"/> | gumanager | Georgetown | Study Manager | | | |
| <input type="radio"/> | investigator | Research | Investigator | | | |
| <input type="radio"/> | manager | Study | Manager | | | |
| <input type="radio"/> | manager2 | Study | Manager2 | | | |
| <input type="radio"/> | manager3 | Study | Manager3 | | | |
| <input type="radio"/> | manager4 | Study | Manager4 | | | |
| <input type="radio"/> | manager5 | Study | Manager5 | | | |
| <input type="radio"/> | nbiamanager | NBIA | Study Manager | | | |
| <input type="radio"/> | tcgaprivate | TCGA | Manager | | | |

At the bottom right of the table, there are two buttons: 'View Details' and 'Back'.

Figure 6.9 UPT page showing a list of calIntegrator2 users

- Select the radio button next to the name and click **View Details** (Figure 6.10).



The screenshot shows the 'Common Security Module User Provisioning Tool' interface for editing a user. The top navigation bar and session information are the same as in Figure 6.9.

The main section contains instructions: 'Update the details of the displayed User. The **User Login Name** uniquely identifies the User and is a required field. The **User First Name** and **User Last Name** identifies the User. The **User Organization**, **User Department** and **User Title** provides his work details. The **User Phone Number** and **User Email Id** provides the contact details for the User. The **User Password** can be entered if the same schema is also going to be used for Authentication. The **User Start Date** and **User End Date** determine the period for which the User is a valid User. The **Update Date** indicates the date when this User's Details were last updated.'

The 'USER DETAILS' section contains the following fields:

| | | |
|---|-------------------|-------------------------|
| * | User Login Name | manager |
| * | User First Name | Study |
| * | User Last Name | Manager |
| | User Organization | |
| | User Department | |
| | User Title | |
| | User Phone Number | |
| | User Password | •••••• |
| | Confirm Password | •••••• |
| | User Email Id | |
| | User Start Date | (MM/DD/YYYY) |
| | User End Date | (MM/DD/YYYY) |
| | User Update Date | 09/24/2009 (MM/DD/YYYY) |

At the bottom right, there are three buttons: 'Update', 'Delete', and 'Back'.

At the bottom, there are four links: 'Associated Groups', 'Associated PE & Privileges', 'Associated PG & Roles', and 'Assign PG & Roles'.

Figure 6.10 UPT page showing details for a selected user

- Click the **Associated Groups** button at the bottom of the page. This opens the page where you can assign a user to a group (*Figure 6.11*).

User and Groups Association

| SELECTED USER | |
|-----------------|---------|
| User Login Name | manager |

Assign or Deassign multiple **Groups** for the selected **User**. To remove the complete association Deassign all the **Groups**.

| AVAILABLE GROUPS |
|-------------------------|
| Study Managers Group 3 |
| Study Managers Group 4 |
| Study Managers Group 5 |
| NCI Study Investigators |
| TCGA Study Managers |

| ASSIGNED GROUPS |
|------------------------|
| Platform Manager Group |
| NCI Study Managers |

Figure 6.11 UPT page for assigning a user to user groups

- Select the group(s) that you want the user to be in and click **Assign**.
- At the bottom of the page click **Update Association**. This completes the assigning of the user to the user group. Now the user will have access to any studies to which the user group has been given access.

Note: You can add a user to more than one user group. For example, a user could be assigned to “Columbia University Study Managers” as well as to “Columbia University Study Investigators”.

Changing a User Password

To change a password for a User, follow these steps:

- Confirm if the User is an LDAP user or not. If the User is an LDAP user, then this person must change their password using the NCI password change utility. Skip the rest of these steps.
- If the User is not an LDAP user, then continue with the rest of these steps.
- Login to UPT as calIntegrator2 Admin.
- Find the User that you want to change. Click the **User** menu option, then **Select an Existing User**.
- Enter the name of the user you are looking for and click **Search**. If you define not criteria, UPT returns a list of all calIntegrator users.
- Select the radio button next to the name and click **View Details**

7. Replace the **User Password** and **Confirm Password** fields with the new password (*Figure 6.12*).

Common Security Module
User Provisioning Tool

Login ID : boalt
Application : calIntegrator2
Role : Admin

HOME USER PROTECTION ELEMENT PRIVILEGE GROUP PROTECTION GROUP ROLE INSTANCE LEVEL LOG OUT

Update the details of the displayed User. The **User Login Name** uniquely identifies the User and is a required field. The **User First Name** and **User Last Name** identifies the User. The **User Organization**, **User Department** and **User Title** provides his work details. The **User Phone Number** and **User Email Id** provides the contact details for the User. The **User Password** can be entered if the same schema is also going to be used for Authentication. The **User Start Date** and **User End Date** determine the period for which the User is a valid User. The **Update Date** indicates the date when this User's Details were last updated.

| USER DETAILS | |
|--------------------------|-------------------------|
| * User Login Name | manager |
| * User First Name | Study |
| * User Last Name | Manager |
| User Organization | |
| User Department | |
| User Title | |
| User Phone Number | |
| User Password | •••••• |
| Confirm Password | •••••• |
| User Email Id | |
| User Start Date | (MM/DD/YYYY) |
| User End Date | (MM/DD/YYYY) |
| User Update Date | 09/24/2009 (MM/DD/YYYY) |

Update Delete Back

Associated Groups Associated PE & Privileges Associated PG & Roles Assign PG & Roles

Figure 6.12 UPT page where you can edit user details, such as a password

8. At the bottom of the page click **Update**.

DATA IMPORT CONFIGURATIONS

This appendix describes configurations for importing data into a study.

Topics in this appendix include the following:

- [Subject Clinical Data Configuration](#) on this page
- [Delimited-Text Annotation Import](#) on page 97
- [Annotation Field Configuration](#) on page 98
- [Sample Data Configuration](#) on page 98
- [Genomic Data Configuration](#) on page 99
- [Imaging Data Configuration](#) on page 99

Subject Clinical Data Configuration

The following clinical data configuration information is collected:

- Clinical Data Source (delimited text)
- Protocol Id (of study to import)
- For delimited text, see [Delimited-Text Annotation Import](#). For subject annotation files, one field must be identified as the subject identifier.
- See [Annotation Field Configuration](#) for details on specification of visibility and browse configuration.

Delimited-Text Annotation Import

Delimited-text annotation files must be in standard comma-separated value format. The file must include a header line that specifies the name for each field. Each row of data must contain the same number of values as the header row. The file must include a column that will be designated as the identifier (e.g. subject identifier, sample identifier,

etc.) for each row. Optionally a file may include a single column that will be designated as a time-point indicator. Each row must contain a unique combination of identifier and time-point indicator of a unique identifier if no time-point is included. An example of the content of a file including a time-point is shown below.

```
"patientId", "timepoint", "bloodPressure", "weight"  
"1234", "T1", "120/80", "180"  
"1234", "T2", "125/80", "190"  
"5678", "T1", "120/85", "200"
```

After upload of the file, the Study Manager must indicate for each field:

- Field type (identifier, timepoint indicator, text, integer, float or boolean)
- After specification of these types, the file will be validated to ensure that the values are valid for the types selected and that the file conforms to the requirements given above.

Annotation Field Configuration

For each annotation field (regardless of the source), the Study Manager must specify the following information:

- Annotation semantics: each annotation field (whether associated with a subject, image series, image or sample) must either:
 - be associated with an existing annotation definition known to the system,
 - be associated to an existing CDE in caDSR or
 - have sufficient semantic metadata recorded so that the field may be submitted for registration as a CDE in caDSR.
- Field authorization: Each field must be either declared publicly visible or restricted to a list of groups. The default will be the visibility settings given at the study level.
- Whether the field is to be included in the results list for a given entity type (i.e. Subject, Sample, ImageSeries or Array Data) when browsing data (See Use Case: Browse Study Data).
- Whether the field is to be included in simple single-input searches when browsing data (See Use Case: Browse Study Data).

Sample Data Configuration

Sample data may be uploaded from either caArray 2 or from delimited-text import. Samples imported from caArray 2 may have annotation updated by use of the delimited-text import functionality if sample annotation is required. Import from caArray 2 requires specification of the following information:

- caArray server hostname
- caArray server JNDI port
- caArray username

- caArray password
- Either the experiment identifier (to import all samples in the experiment) or a file containing a comma-separated list of samples in the format “experiment identifier”, “sample name”.
- Mapping of samples to subjects. This may be specified by a comma-separated list in the format “subject identifier”, “sample identifier” or by a regular-expression based mapping formula.

When samples are imported via delimited-text import, the time-point is associated to the sample itself. This means that each sample may be associated with only one time-point (i.e. multiple time-points for the same sample are invalid).

Genomic Data Configuration

All genomic data (i.e. array data) is imported from caArray 2. First the Study Manager must specify sufficient information to map study samples to caArray 2 samples. If all samples were imported directly from caArray 2 as described in Special Requirement: Sample Data Configuration, no further information is required for this step. If samples were imported via delimited-text, the Study Manager must specify

- caArray server hostname
- caArray server JNDI port
- caArray username
- caArray password
- A mapping of calIntegrator2 sample identifiers to caArray 2 samples, specified as a comma-separated list in the format “calIntegrator2 sample identifier”, “caArray 2 experiment identifier”, “caArray 2 sample name”.

The system will enable the Study Manager to navigate easily to the selected caArray 2 instance.

Next, the system will indicate the available platforms and array data types available for the study samples. The Study Manager will indicate which platforms and data types to import and for each platform/data type combination will indicate:

- Whether to import the data
- The visibility of the data; either public or restricted to a set of groups. Low-level genotyping data (raw data and normalized) will always have restricted visibility.

Imaging Data Configuration

The following imaging data configuration information is collected:

- NBIA grid server hostname (defaults to NCICB instance)
- NBIA grid server port (defaults to NCICB instance port)
- Protocol Id

- Mapping of NBIA Patients to subjects imported from clinical data source. This may be specified by a comma-separated list in the format “subject identifier”, “NBIA patient identifier” or by a regular-expression based mapping formula.
- Which annotation fields to import from NBIA.
- The system will enable the Study Manager to navigate easily to the selected caArray 2 instance.

Additional annotation for either images or image series may be imported using the delimited-text import functionality.

INDEX

A

account, requesting new user 6

adding

clinical data 16

genomic data 24

imaging data 29

new user to user group 93

annotation

assigning identifier 17

configuring field 98

importing delimited text 97

K-M plot for 54

searching for definitions 20

Application Support i, 11

assigning, annotation identifier 17

B

box and whisker plot

interpretation 72

uses for 72

C

caIntegrator2

logging in 8

logging out 10

requesting user account 6

using workspace 8

workspace 8

caIntegrator2 User's Guide

introduction 1

organization 1

text conventions 2

clinical study

adding data 16

data for 13

columns, defining display 40, 42

Comparative Marker Selection (CMS)

data analysis 76

configuring

annotation fields for import 98

copy number data 28

genomic data for import 99

imaging data for import 99

sample data for import 98

subject clinical data for import 97

control samples, uploading 27

copy number data, configuring 28

creating

K-M plot 54

new user 88

protection group 89

study 14, 15

user account 85

D

data analysis, overview 53

defining survival values 23

delimited text annotation import 97

DICOM, retrieving images 50

E

editing a query 43

exporting

data 52

query results 43

F

fold change, control samples file 27

G

gene expression, K-M plot for 56

gene expression plot

description 53, 60, 70

for clinical queries 66

for genomic queries 64

interpreting 70

plot display, box & whisker 72

plot display, log2 intensity 71

plot display, mean 70

plot display, median [71](#)

GenePattern

analyses, description [73](#)

analyses, in caIntegrator2 [74](#)

analyses, modules [75](#)

CMS analysis [76](#)

GISTIC analysis [81](#)

PCA [78](#)

plot description [54](#)

genomic data

adding copy number data to [28](#)

adding to study [24](#)

configuring for import [99](#)

for study [13](#)

mapping to clinical data [26](#)

GISTIC-based data analysis [81](#)

H

hierarchy of objects, NBIA [37, 51](#)

I

imaging data

adding to study [29](#)

configuring for import [99](#)

for study [14](#)

importing delimited text annotations [97](#)

K

Kaplan_Meier plot see K-M plot

K-M plot

creating [54](#)

description [53, 54](#)

for annotations [54](#)

for gene expression [56](#)

for queries [58](#)

L

logout link [10](#)

M

managing

platforms [32](#)

queries [43](#)

study [31](#)

user accounts [85](#)

mapping genomic to clinical data [26](#)

N

NBIA

forwarding imaging results to [49](#)

viewing imaging results in [49](#)

NCICB Application Support [i, 11](#)

O

objects in NBIA, hierarchy of [37, 51](#)

overview, chapters in guide [1](#)

P

password, changing [95](#)

patient

relationship to study, series, images [51](#)

platforms, managing [32](#)

plot

gene expression, description [53, 60](#)

gene expression description [70](#)

GenePattern description [54](#)

K-M description [53](#)

Principal Component Analysis

data analysis [78](#)

protection group

creating [89](#)

definition [86](#)

Q

query

editing [43](#)

exporting results [43](#)

K-M plot for [58](#)

managing [43](#)

saving [43](#)

R

registering new user [6](#)

Results Type tab [40](#)

S

sample data, configuring for import [98](#)

saving query [43](#)

searching

annotation definitions [20](#)

overview [35](#)

study [36](#)

search results

browsing [46](#)

exporting data [52](#)

forwarding imaging results to NBIA [49](#)

genomic data [46](#)

imaging data [48](#)

overview [45](#)

retrieving DICOM images [50](#)

viewing imaging data in NBIA [49](#)

Sorting tab [42](#)

study

adding clinical data [16](#)

adding genomic data [24](#)

- adding imaging data 29
- clinical data, description 13
- configuring copy number data 28
- creating 13, 14, 15
- deploying 31
- editing 31
- genomic data, description 13
- imaging data, description 14
- managing 31
- mapping genomic data to clinical 26
- relationship patient, study, series, images 51
- searching 36
- uploading control samples to 27
- subject clinical data, configuring for import 97
- survival values, defining 23

T

- Technical Support i
- text conventions in user guide 2

U

UPT

- adding new user to user group 93
- assigning user group to protection group 90
- creating new user 86
- creating protection group 89
- creating user group 88
- description 85
- summary of steps 86

user

- adding to user group 93
- changing password 95
- creating new 86
- creating user access, summary 86
- definition 85

- user's manual conventions 2

- user account, new 85

user group

- assigning to protection group 90
- creating 88
- definition 86

