User Manual of Breedbase

Breedbase team

2024-07-22

Contents

1 Basic Website Usage **11**

1.1 Creating a User Account 11

1.1.1 Verifying first that you do not already have an account 11

1.1.2 Creating a user account 11

1.2 Managing your Account 12

1.2.1 Login 12

1.2.2 Editing Account Settings 13

1.2.3 Changing Your Account Status: From "User" to "Submitter" 14

1.2.4 Submitting Feedback on an SGN Database 14

1.3 Menu Layout 15

1.3.1 Menu Options 15

1.4 Working with Lists 17

1.4.1 Creating lists 17

1.4.2 Viewing and editing lists 21

1.5 User Permissions 22

2 Searching the Database **23**

2.1 The Search Wizard 24

2.1.1 How the Search Wizard Works 24

2.1.2 How to use retrieved data 26

2.1.3 Updating the Wizard 29

2.2 Accessions and Plot Search 29

2.3 Trials Search 32

2.4 Trait Search 33

2.5 Ontology Browser 35

2.6 Search Seedlots 36

3 Managing User Roles **39**
3.1 What are User Roles? . . . 39
3.2 The Manage User Roles page . . . 40

4 Managing Breeding Programs **41**

5 Managing Locations **43**

6 Managing Accessions **45**
6.1 Add Accessions Using A List . . . 45
6.2 Uploading Accessions and Accession's Info From A File . . . 48
6.3 Email alert for accession upload . . . 51
6.4 Add Parentage (Pedigree) Information to Accessions . . . 52
6.5 Working with grafts . . . 53
6.6 Bulk renaming of accessions . . . 54

7 Managing Seed Lots **57**
7.1 Add New Seedlot(s) . . . 58
7.2 Seedlot Transactions . . . 59
7.3 Seed Inventory . . . 60
7.4 Find Seedlots For a List of Accessions . . . 61
7.5 Create a seedlot for an Accession or Cross . . . 63
7.6 Add quality data to a seedlot . . . 64
7.7 Seedlot Maintenance Events . . . 64
7.7.1 Setup . . . 64
7.7.2 Adding Events . . . 65
7.7.3 Displaying Events . . . 70
7.7.4 Downloading Events . . . 72
7.8 Deleting Seedlots . . . 72

8 Managing Populations **73**

9 Managing Crosses **75**
9.1 Crossing Experiment . . . 75
9.1.1 Add New Crossing Experiment . . . 76
9.2 Cross . . . 77
9.2.1 Add New Crosses . . . 77
9.2.2 Update Crosses by Uploading . . . 81
9.3 Cross Wishlist . . . 83

9.3.1 Create a Cross Wishlist 83
9.4 Crossing Experiment Detail Page 89
9.5 Cross Detail Page 91

10 Managing Field Trials 95
10.1 Trial Detail Page 96
10.2 Adding Trials 98
10.2.1 Prerequisites 98
10.2.2 Adding a trial by using "Add Trial" form 99
10.2.3 Adding a trial from an uploaded file 106
10.2.4 Multi-location trials 112
10.2.5 Email alert for multiple trial design upload 113
10.2.6 Viewing Plot Layout and Trait HeatMap 113
10.2.7 Adding additional information in the "Trial Detail" page 120
10.2.8 Downloading the Trial Layout from the "Trial Detail" page 129
10.2.9 Adding Plant Entries To Your Trial 131
10.2.10 Adding Tissue Sample Entries To Your Trial 133
10.2.11 Uploading GPS Coordinates For Plots 136
10.2.12 Uploading Additional Files To Trial 137
10.2.13 Deleting Trial Data 138

11 Managing Genotyping Plates 141
11.1 Adding a New Genotyping Plate 143
11.2 Genotyping Plate Detail Page 145

12 Using Field Book App 147
12.1 A typical workflow 148
12.2 Creating Field Layout Files for the Field Book App 148
12.2.1 Creating "Field Layout Files" by using "Field Book Tools" page. 149
12.2.2 Creating "Field Layout Files" by using "Trial Detail" page. 151
12.3 Creating Trait Files for the Field Book App 153
12.3.1 Creating a Trait List 153
12.3.2 Creating a Trait File 153
12.4 Transferring Files from Your Computer to Android Tablet . . 155
12.4.1 Files on your computer 155

12.4.2 Files on your Android tablet 156
12.5 Setting up "Field Book App" for data collection 158
12.6 Exporting Files from Field Book App 167
12.7 Uploading Phenotype Files to an SGN database 170

13 Managing Phenotypic Data 173
13.1 Uploading Fieldbook Phenotypes 173
13.1.1 Export Field Book Database File 173
13.1.2 Upload Field Book Database File 174
13.2 Uploading Spreadsheet Phenotypes 174
13.2.1 Generating Spreadsheet File 175
13.2.2 Uploading Spreadsheet File 177

14 Managing Barcodes 179

15 Using the Label Designer 183
15.0.1 First Select a Datasource 183
15.0.2 Set Page and Label Size 184
15.0.3 Design Your Label 185
15.0.4 Adjust Formatting, Save, and Download 186

16 Managing Downloads 189

17 Managing ODK Data Collection 191
17.1 ONA Crossing Information 192
17.1.1 Managing ONA Crossing Information 192
17.1.2 Reviewing Plant Status 193
17.1.3 Graphical Summary For Performed Crosses 193
17.1.4 Summary Information For Performed Crosses 194

18 Managing Tissue Samples 195
18.1 Tissue samples from field trials 195
18.2 Genotyping Plate Tissue Samples (96 or 384 well plates) . . . 198

19 Managing Observation Variables 201
19.1 Managing Observation Variables with Traits, Methods, and Scales 201

20 Managing Image Data 207

20.1 Image-Phenotyping Dashboard . . . 207
20.2 Image Input . . . 207
20.3 Standard Process . . . 212
20.4 Ground Control Points . . . 220

21 Managing VCF Data **221**
21.1 Uploading VCF Data . . . 221
21.2 Searching and Downloading VCF Data . . . 224
21.3 Searching Protocols . . . 226
21.4 Detail Pages and Deletion . . . 228

22 Managing Spectral Data **231**
22.1 Upload Spectral Data . . . 232
22.2 Evaluate and Remove Outliers . . . 233
22.3 Plot Spectra . . . 233
22.4 Aggregate Spectra . . . 234
22.5 References . . . 235

23 Managing Sequence Metadata **237**
23.1 What is Sequence Metadata? . . . 238
23.2 Loading Sequence Metadata . . . 238
23.3 Searching Sequence Metadata . . . 239
23.3.1 Basic Search . . . 239
23.3.2 Advanced Search . . . 241
23.4 Marker Integration . . . 241
23.5 Sequence Metadata API . . . 241

24 Managing Outliers in Dataset **243**
24.1 What is Outliers Functionality in Dataset ? . . . 243
24.2 Accessing Trait Visualization . . . 244
24.3 Interpreting Visual Elements . . . 244
24.4 Choosing Cut-Off Values . . . 245
24.5 Setting Deviation Multiplier . . . 245
24.6 Utilizing Graph Controls . . . 246

25 Data Analysis Tools **247**
25.1 Selection Index . . . 247
25.2 Genomic Selection . . . 251

25.2.1 Building a Model - Method 1: . . . 252
25.2.2 Building a Model - Method 2 . . . 263
25.2.3 Building a Model - Method 3 . . . 271
25.3 Genome Browsing . . . 272
25.3.1 Browsing Genotype data by Accession . . . 272
25.3.2 Browsing Genotype data by Trial . . . 273
25.4 Principal Component Analysis (PCA) . . . 274
25.5 ANOVA . . . 276
25.6 Clustering (K-Means, Hierarchical) . . . 276
25.7 Genetic Gain . . . 279
25.8 Kinship and Inbreeding Coefficients . . . 280
25.9 Creating Crossing Groups . . . 281
25.10 Search Wizard Genomic Relationship Matrix (GRM) Download 284
25.11 Search Wizard Genome Wide Association Study (GWAS) . . . 285
25.12 Spectral Analysis . . . 289
25.12.1 Dataset selection . . . 290
25.12.2 Cross-validation . . . 291
25.12.3 Preprocessing . . . 292
25.12.4 Algorithms . . . 293
25.12.5 Output: common model summary statistics . . . 294
25.12.6 Export model for later use . . . 295
25.12.7 Predict phenotypes from an exported model (routine use) . . . 295
25.12.8 FAQ . . . 296
25.13 General Mixed Model Tool . . . 297
25.14 Genomic Prediction of Cross Performance (GCPC) . . . 299

Introduction

Welcome to the Breedbase manual!

This manual is intended for database users.
If you are a developer looking for software implementation details, please visit the developer wiki instead: https://github.com/solgenomics/sgn/wiki

Chapter 1

Basic Website Usage

1.1 Creating a User Account

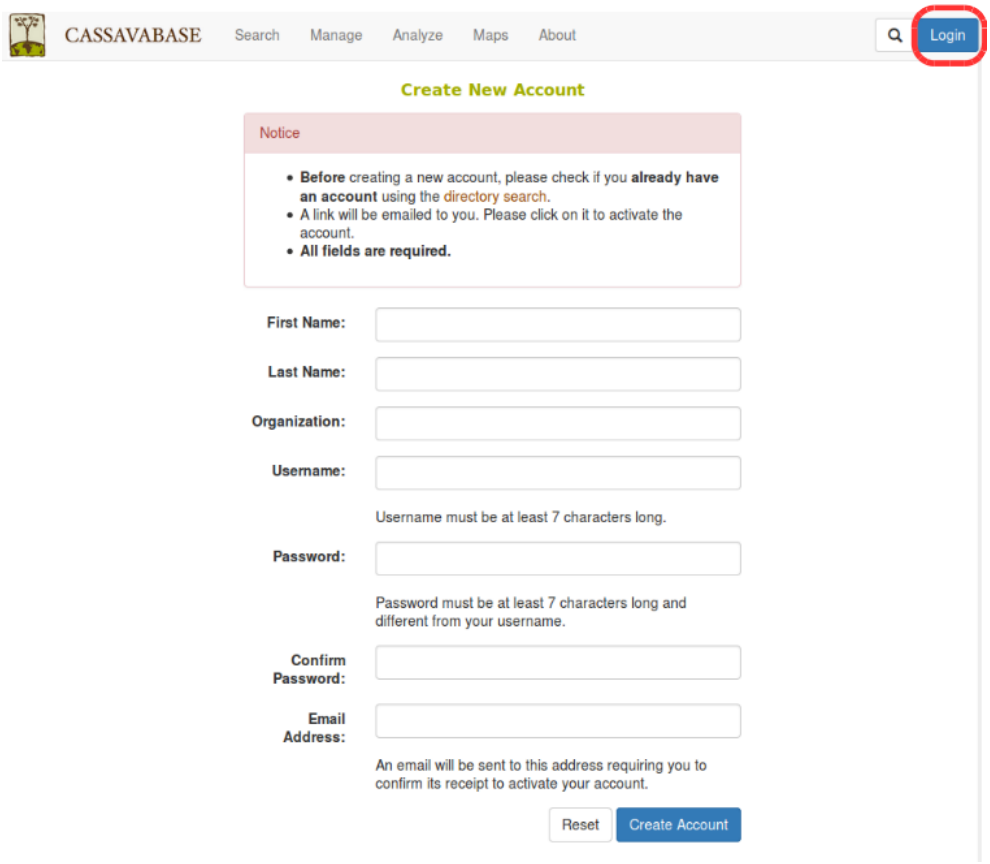
1.1.1 Verifying first that you do not already have an account

Before creating an account, please verify first that you don't already have an account. You can use "Search" menu to check if you already registered as a user.

In the "Search" menu, selecting the "People" tab and search your name. If nothing is found, proceed with the instructions below. Otherwise, clicking the "Login" button. If you have forgotten your password, you can retrieve it by clicking the "Forgot your password?" link on the login page.

1.1.2 Creating a user account

On the right of the toolbar, clicking on "Login." It will take you to the login page. On the login page, clicking on the link "sign up for an account." It will take you to the page below:



Filling in all of the information, then clicking "Create Account."

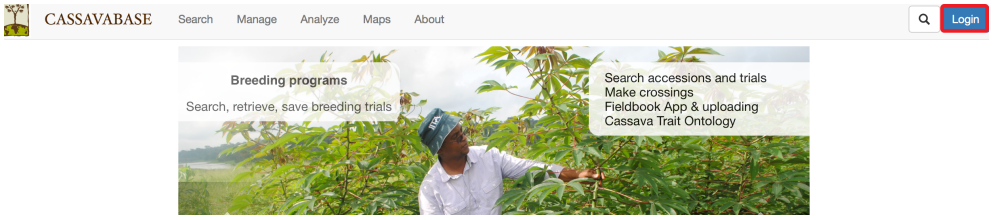
After you submit the information, an email will be sent to the provided email address. Checking your email and clicking on the link to activate your account.

1.2 Managing your Account

1.2.1 Login

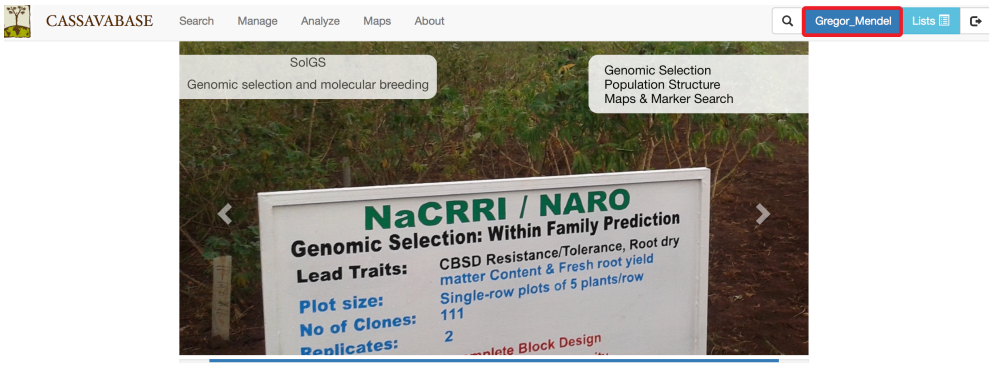
To login, clicking the "Login" link in the toolbar on any page and enter your username and password.

If you have forgotten your password, you can retrieve it by clicking the "Forgot your password?" link on the login page.



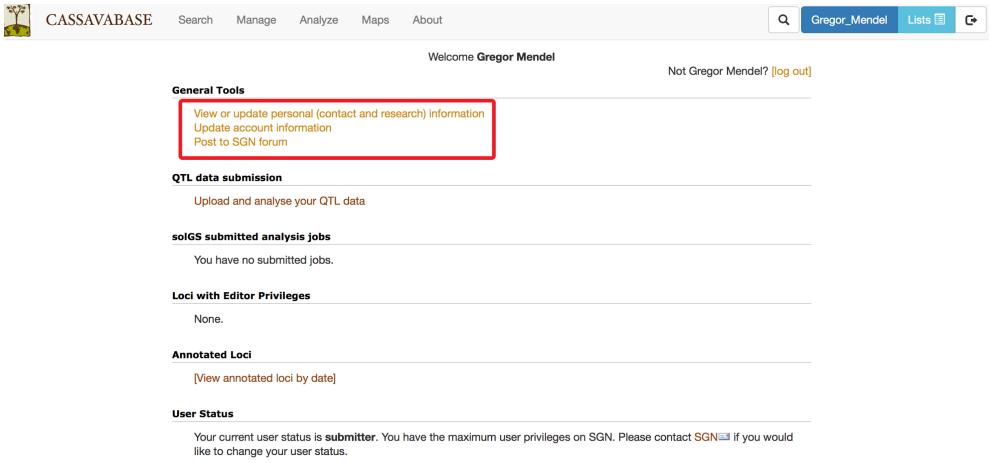
1.2.2 Editing Account Settings

Account settings can be edited by clicking on the "my profile" link displayed as your user name, on the right of the toolbar. You must login, in order to access and change account settings.



You can add personal information to your account using the "View or update personal information" link.

To change your password, username, or your contact email, clicking on "Update account information" link. You must provide your old password before you can make any changes.



1.2.3 Changing Your Account Status: From "User" to "Submitter"

After you create an account, your account has a "user" status. This account has limited privileges.

Accounts with "user" status are able to:

- Change personal information
- Post comments on pages
- Post to the forum

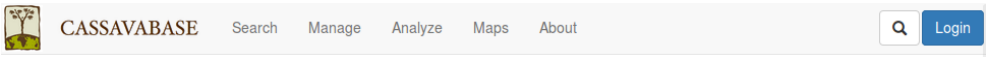
To upgrade your account status to "submitter," contact the database curators using the "contact" link provided at the footer of each page. Submitter accounts can add data, such as new plots, accessions, phenotype data and images.

1.2.4 Submitting Feedback on an SGN Database

We appreciate your feedback! Feel free to submit any questions or suggestions by using the "Feedback" link provided at the footer of each page.

1.3 Menu Layout

SGN Database websites have a toolbar on the top of each page with a number of menus for convenient access of major functions. The menus, as pictured below, are "search," "manage," "analyze," and "maps." The toolbar also provides a quick search, a "log in" button, and a "new user" button.



1.3.1 Menu Options

Search

In the Search menu, the options are:

Tab	Description
Wizard	Search different accessions and plots by location, year, trial, and trait data. Can also be used to create lists of different types.
Accession and plots	Search accessions and plots using a variety of criteria
Trials	Search trials by name, description, breeding program, year, location, and trial type.
Markers	Search different markers
Images	Search images contained in the SGN database
People	Search database users

Manage

In the Manage menu, the options are:

Tab	Description
Breeding Programs	View, add and delete breeding programs
Locations	View, add and delete locations
Accessions	Manage and search different accessions
Seedlots	Manage and search different seedlots

Tab	Description
Crosses	Create new crosses in the database
Field Trials	Manage field trials. Create trials using different field layouts.
Genotyping Plates	Manage genotyping plates. Create 96 or 384 well plates.
Phenotyping	Upload phenotyping files from the Tablet Field Book application
Field Book App	Manage the field book app data (download files to tablet)
Barcodes	Refers to the old barcode system, mainly historical
Download	Download information in the database based on lists

Analyze

Clicking on the "Analyze" link will give a full menu of all analysis functions
In the Analyze menu, the options are:

Tab	Description
Breeder Tools	
Breeder Home	Access breeding functionalities. Lists important and helpful links.
Barcode Tools	Manage, create, and download barcodes. Also access barcode tools.
Genomic Selection	Can search for traits, start building a GS model, and predict values based on genotypes
Sequence Analysis	
BLAST	Sequence homology search
Other	
Ontology Browser	Browse all recorded ontologies

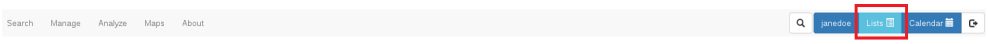
1.4 Working with Lists

Lists are collections of identifiers that are stored in the database. Lists can be composed of accessions, plots, traits, locations, and trials. Lists are attached to the individual user's account, and can only be created and seen by the user while logged in. SGN databases make heavy use of lists in a number of tools on the website. For example, trials are created using lists of accessions.

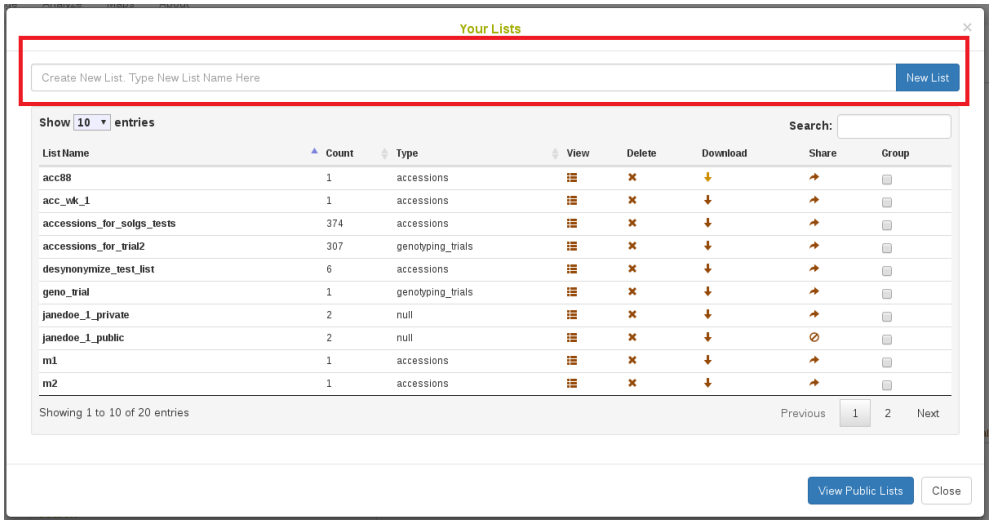
1.4.1 Creating lists

Lists can be generated in various ways:

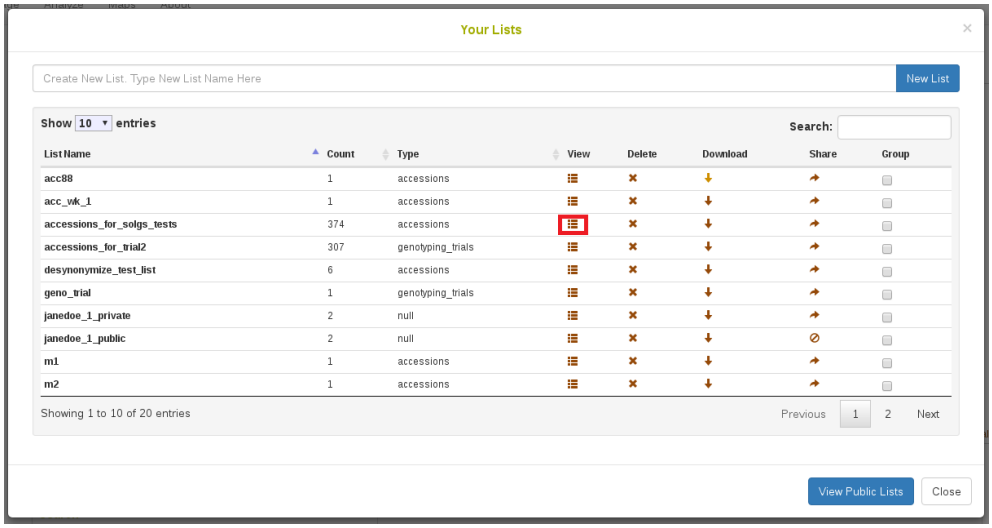
One way to create a list is by clicking on the "Lists" link located on the toolbar.



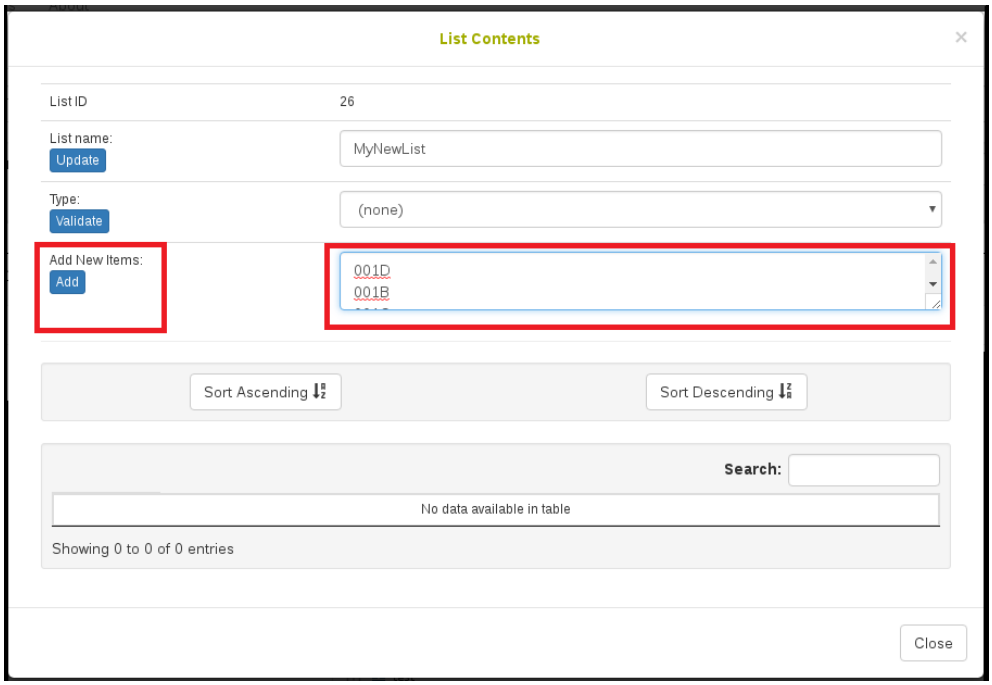
To create a new list, enter the name of your new list and then clicking on the "New List" button. The name of the list can be anything, but should be unique and should be something to help you easily identify.



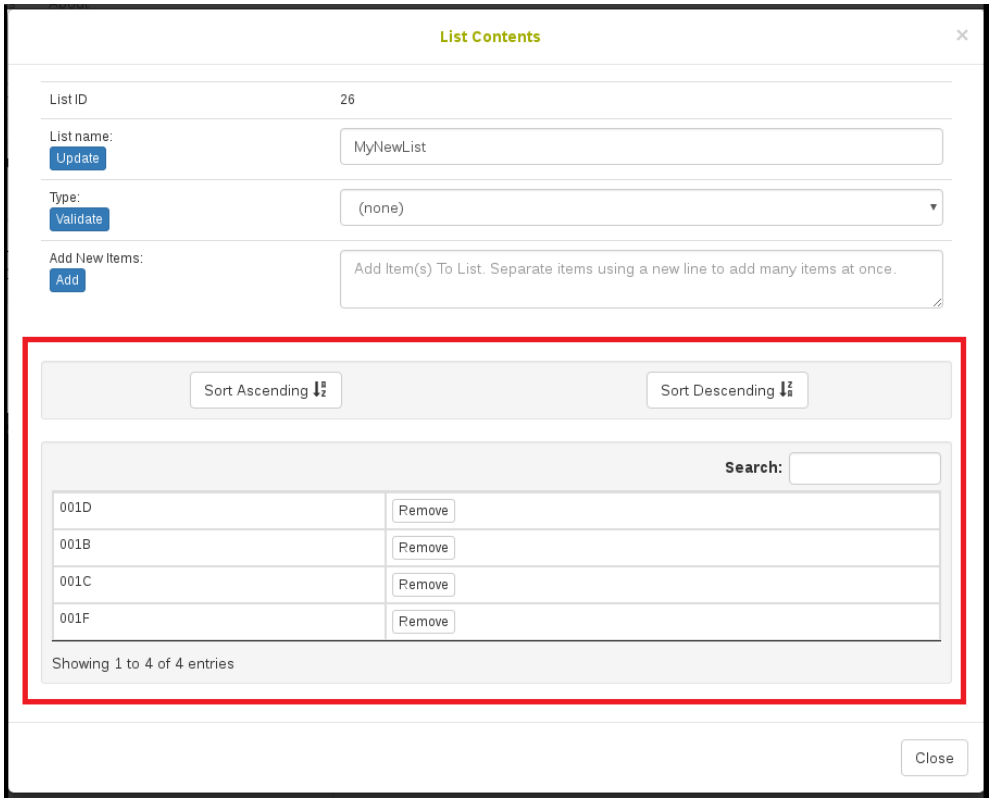
You can find the list that you entered on the "Your Lists" page. To add items to your list, click on the "View" icon to open "List Contents" page.



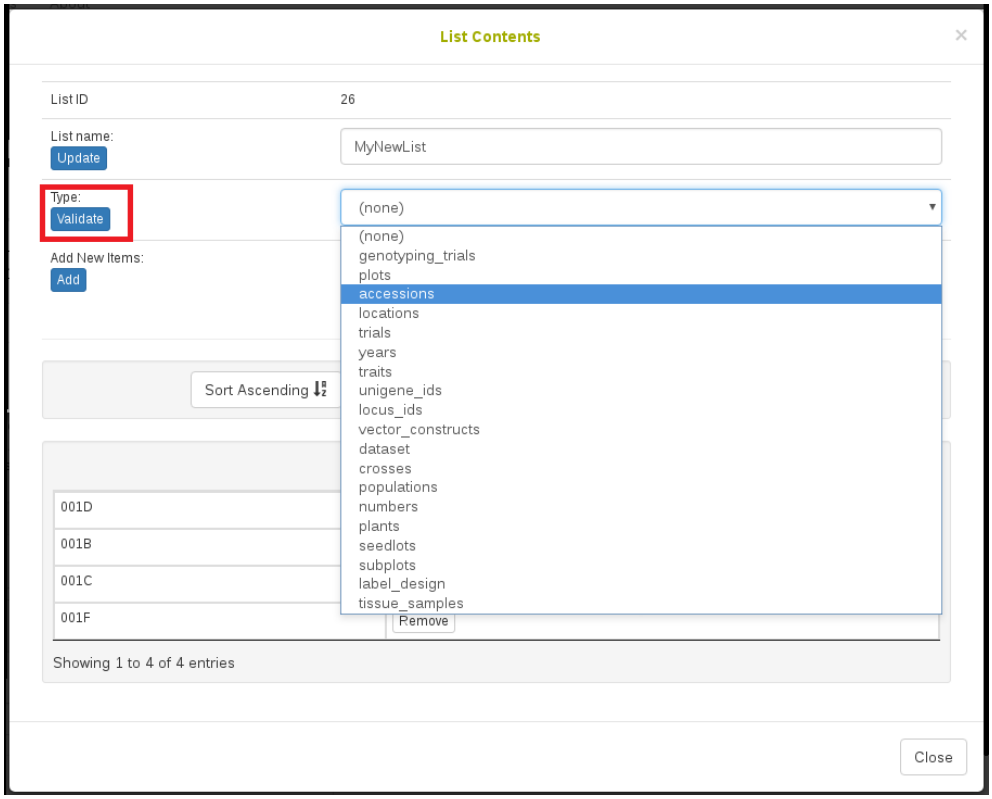
On the "List Contents" page, enter items that you want to add to the list, then click on "Add" button.



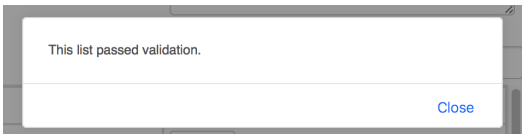
The page will be updated and will display your items in a table at the bottom of the page. It is possible to sort the list if you need.



Select the type of items in your list. To verify that the items that you added to your list are already stored in the database and that you selected a correct type for the items, click on the “Validate” button.



If those items are already in the database, a message will indicate that "This list passed validation"

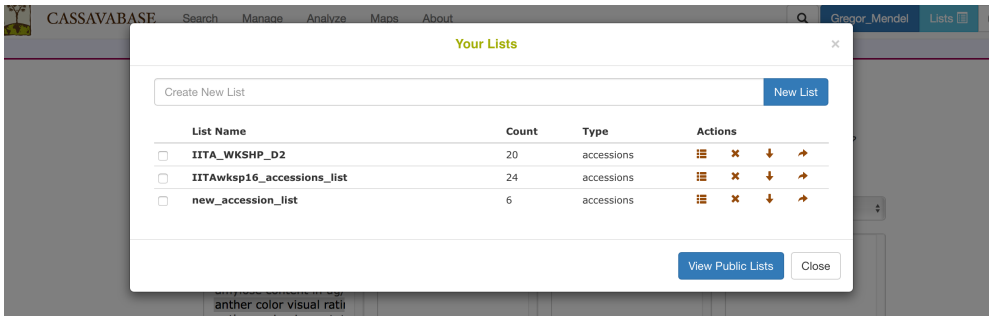


Note that a list cannot contain duplicate elements. If a duplicate item is entered, the list manager will inform the user that the element is already in the list and will not add it again.

Another easy way to create a list is to use 2.1, which can be accessed from the Search menu.

1.4.2 Viewing and editing lists

Lists can be viewed and edited using the "Lists" link on the toolbar. Clicking on the link will open a window that displays all of your lists, as well as an option to create new lists.



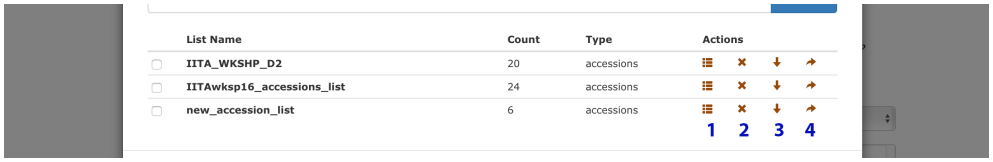
This page shows all lists that have been created, including those created by using the Search Wizard. You can view and edit your lists by using "Actions" buttons.

1. Clicking on the "view" icon will open a new window called "List Contents" that allows you to change the list name, the type of the list, add new items, or delete existing items.

2. Clicking on the "delete" icon will delete your list. **Caution: this action cannot be undone**.

3. Clicking on the "download" icon will download the contents of your list to your computer.

4. Clicking on the "make public" icon will make your list available for other users to view and use your list.



1.5 User Permissions

Breedbase accounts are assigned one or more of four different roles to determine the level of access they have within the database. The possible roles are **User**, **Submitter**, **Sequencer**, and **Curator**. Each role grants specific permissions, and careful management of them helps prevent data from being altered or deleted in error.

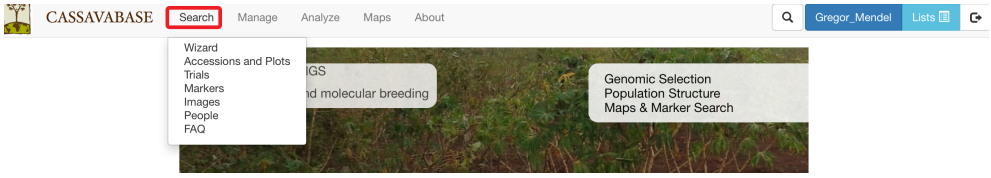
Accounts are also assigned Breeding Program role(s) to grant access to the specfic breeding program(s) they work with.

- The **User** role gives an account permission to view and download data throughout the database.
- The **Submitter** role gives an account permission to design field experiments and to upload and edit data using the tools in the "Manage" section. In order to submit and manage breeding data within a given breeding program, a submitter also must have a matching Breeding Program role.
- The **Sequencer** role gives an account permission to design genotyping experiments and submit plates to a genotyping service.
- The **Curator** role gives an account permission to do all of the above, as well as to delete data within the database. The Curator role also enables the addition or deletion of roles for all database accounts in the 'Manage User Roles' tool.

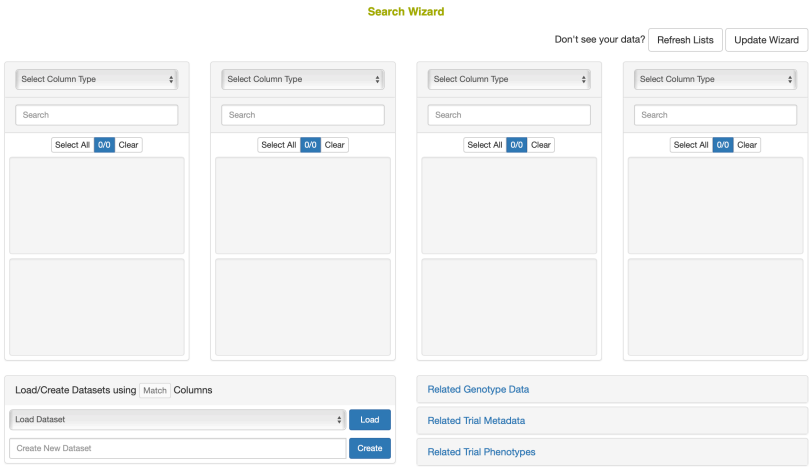
Chapter 2

Searching the Database

You can search for information on the database by using the following search options: Wizard, which uses combined criteria specified by users; Accessions and Plots; Trials; Markers; Images; People; FAQ.



2.1 The Search Wizard

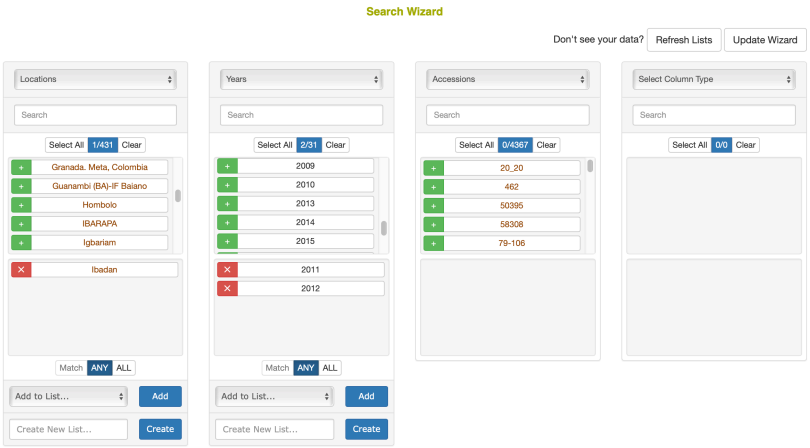


2.1.1 How the Search Wizard Works

The search wizard presents a number of select boxes, which are initially empty. You start searching by picking a category of data from the dropdown above the left-most select box.

Once a category has been picked, the database will retrieve all the options within this category and display them within the first select box. You then select one or more options from the first select box, which activates the second dropdown.

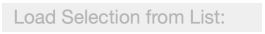
You can then select a category from the second dropdown, and repeat this same search process through all four dropdowns and select boxes.



- In the example above, the "locations" category was chosen in the first dropdown. The first select box then displayed all the possible locations in the database. The option Ibadan was selected.
- This activated the second dropdown. The category "years" was chosen in the second dropdown. The second select box then displayed all the years that are linked in the database to the location Ibadan. From that list, the options 2011 and 2012 were selected.
- This activated the third dropdown. A final category, "accessions", was chosen in the third dropdown. The third select box was then populated with the 3847 accessions in the database that are linked with the location Ibadan in the years 2011 or 2012.

In addition to the basic search operations demonstrated above, users can take advantage of two more features:

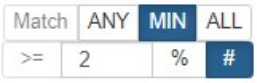
Load Selection from List



- Instead of picking a category in the first dropdown, users can instead populate the first selectbox from a list by scrolling down in the first dropdown to the "Load Selection from List" subheading and selecting a list. This is useful for starting queries with a list of plots, as this category is not among the options in the first dropdown.

ANY/MIN/ALL Toggle

- By default, the search wizard combines options within a category using an OR query. In the example above, in the third panel the wizard retrieved accessions associated with the location 'Ibadan' in **ANY** of the years "2011 **OR** 2012"
- If the user clicked the toggle below the second select box to change it to **ALL** before choosing accessions in the third dropdown, the wizard would instead retrieve accessions associated with the location 'Ibadan' in the years "2011 **AND** 2012". This will be a smaller set of accessions, because any accessions used only in 2011, or only in 2012 will be excluded.
- A more advanced search could use the **MIN** toggle option. This allows the user to make a query in between an ANY or ALL query, where a minimum number of matches from the selected column will be used as a filter for the next column. The minimum can be provided as either a percentage (%) or an actual count of items (#). In the example above, if the years 2011, 2012, and 2013 were selected in the second column, the user could enter '2' in as the minimum and select '#' as the minimum match type. This would select accessions in the third column that were used in 2 or more of the selected years.



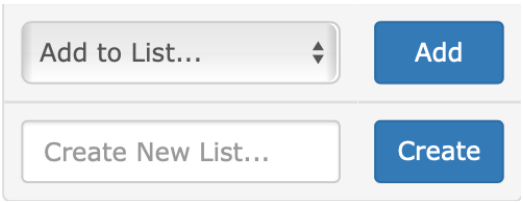
2.1.2 How to use retrieved data

Getting more Info

Any option in the wizard select boxes (except for years) can be clicked to open a page with more details. The new page is opened in a new tab.

Saving to a list

You can store the highlighted items in any selected box to lists. This is done using the inputs and buttons directly below the select box. **Don't forget, you must be logged in to work with lists!**



- To **add items to an existing list**, first pick an existing list using the "Add to List…" dropdown on the left. Then click the "Add" button. A popup window will confirm the action, and display the number of items added to your existing list.

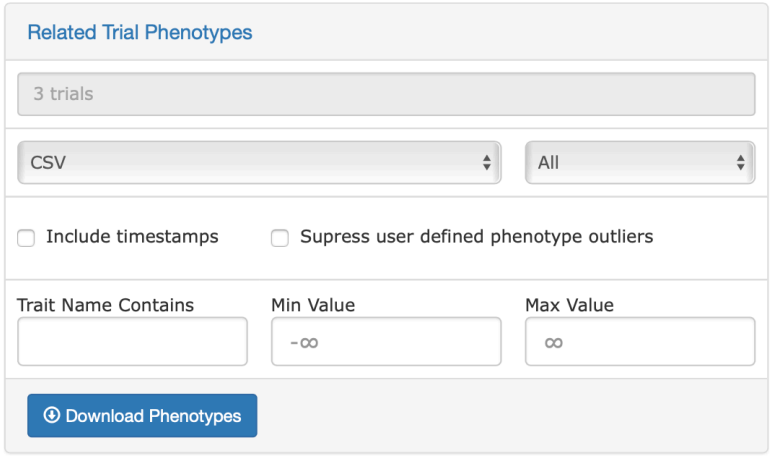
- To **store items to a new list**, first type a new list name in the "Create New List…" text input on the left. Then click on the "Create" button. A popup window will confirm the action, and display the number of items added to your new list.

Downloading Data

You can download trial metadata, phenotypes and genotypes associated with the highlighted items in the wizard select boxes. This is done using the buttons in the download section at the bottom of the page. **Don't forget, you must be logged in to download data!**

Metadata Trial metadata can be downloaded by selecting a subset of trials from the database or based on your search categories. To download, click on "Related Trial Metadata", a dialog will appear. Select download format and click the "Metadata" button to complete your download.

Phenotypes The phenotypes download is quite flexible, and can download a subset of all the trial data in the database based on whichever categories and options you currently have selected. Simply click on the "Related Trial Phenotypes" link, review the options, changing or adding any additional parameters you like, then click 'Download Phenotypes'.



Genotypes The genotype download is more stringent. It requires a minimum of one accession and one genotyping protocol to be selected in the wizard select boxes. The text box in the download section of the page will help track what has been selected. Once clicked, the "Download Genotypes" button will download a genotype file for the selected accessions.

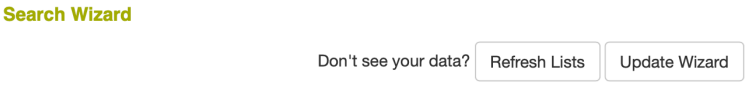
Saving the wizard selections

As discussed above, the selections of the individual select boxes in the wizard can be saved separately to a list. The lists can be used as inputs in other tools on the site. However, sometimes creating a selection is quite time consuming and restoring the selections from four different lists would be cumbersome

too. Therefore, the selections can be saved together in a dataset, and named for later retrieval. This is done in the section "Load/Create Datasets" that is below the first two wizard select boxes. To select an existing dataset, one uses the "Load Dataset" dropdown. A particular dataset can be chosen, and the "Load" button can be clicked to retrieve and display the dataset in the wizard. To create a new dataset using items that are selected in the wizard, one can enter the name of the new dataset in the "Create New Dataset" text box. Once the dataset has been given a name, clicking the "Create" button will save the new dataset.

2.1.3 Updating the Wizard

The search wizard uses a copy of the database, or a cache, to return results quickly. If data appears to be missing, it usually means that the cache needs to be updated. Users with submitter privileges or above can do this using the 'Update Wizard' button. One can also use the 'Refresh Lists' button to update the available lists.

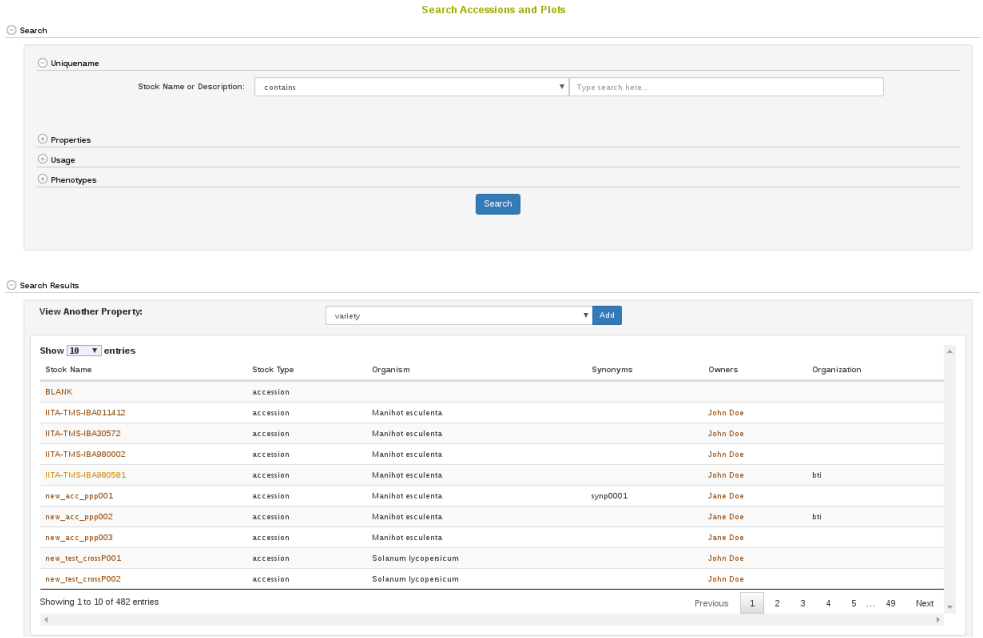


This will take just a few seconds in small databases, but may take a few hours to complete in larger databases.

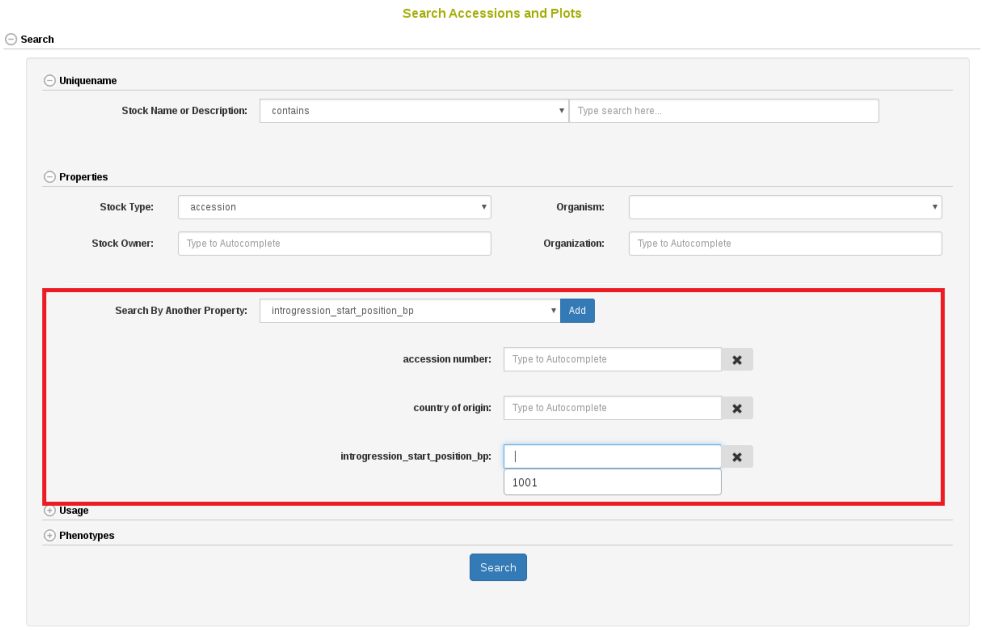
2.2 Accessions and Plot Search

Accessions and their related materials (cross, plant, plot, population, tissue_sample, training population) can be searched by using "Search Accessions and Plots" page. On this page, "accession" is the default stock type; however, you can change stock type by selecting an option from the dropdown list. From this page you can construct detailed queries for stock types.

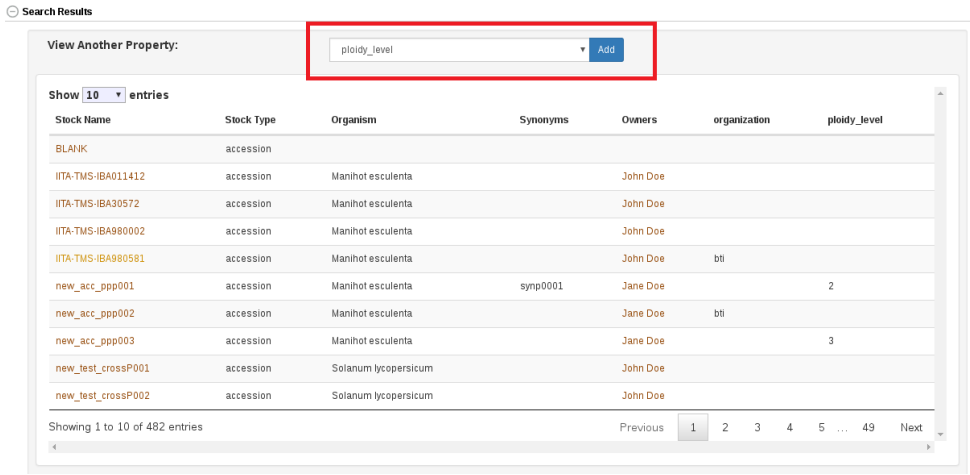
For example, by using the "Usage" section, the "Properties" section, and the "Phenotypes" section you could search for accessions which were diploids used in a specific year and location and were also phenotyped for height. You can also search for accessions based on genetic properties, such as the location of an introgression on a specific chromosome.



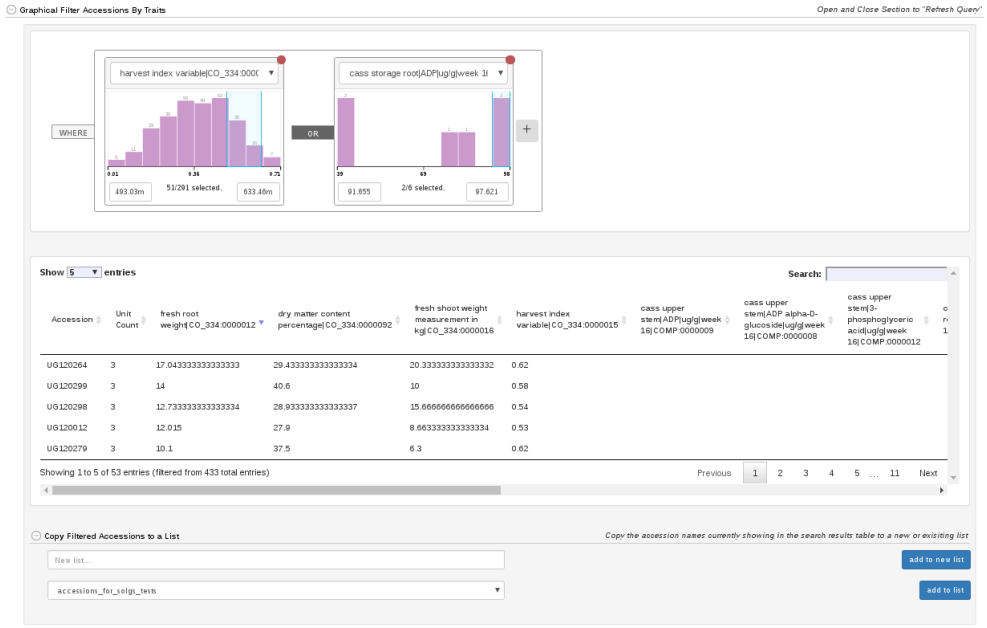
It is possible to query over any of the available properties, such as "ploidy_level", "country of origin", "introgression_chromosome", etc.



In the search result table it is possible to select any of the available properties to view.



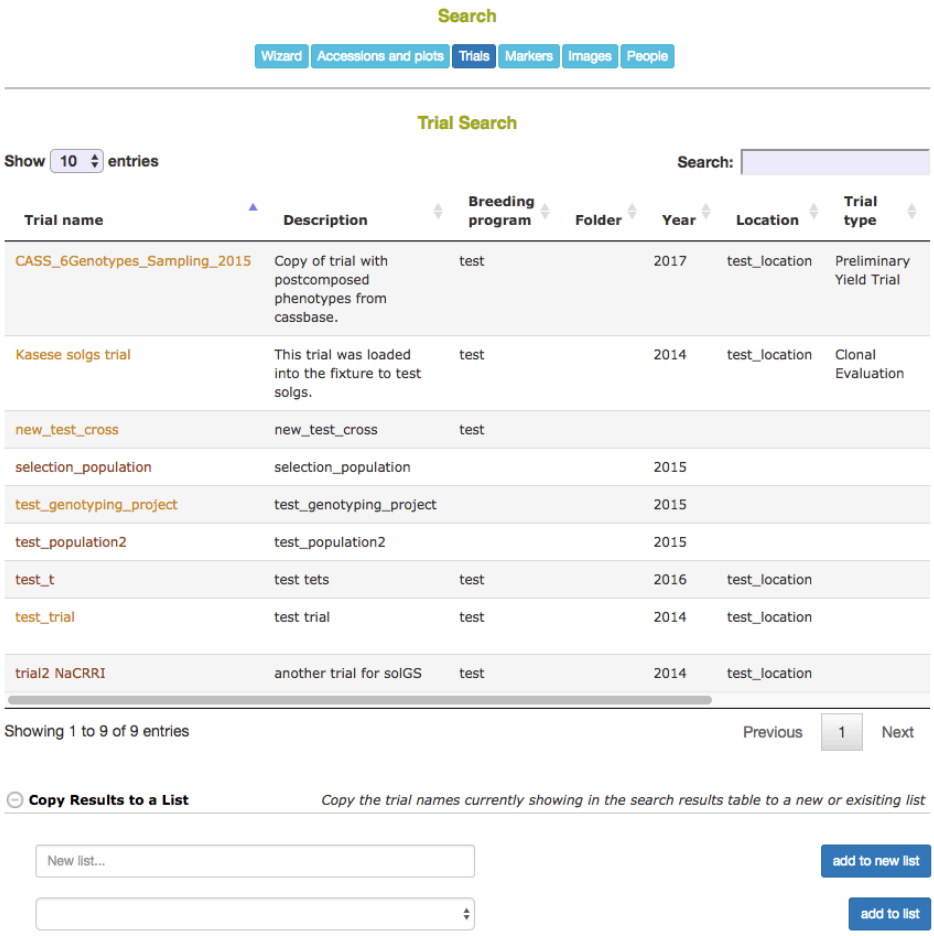
At the bottom of the accession search there is a phenotype graphical filtering tool. Here you can filter down accessions based on combinations of trait performance. The filtered down accessions are then able to be saved to a list.



For information on adding Accessions please see the Managing Accessions help. For information on how field trial plots, plants, tissue samples, and subplots are added to the database, please see the Managing Field Trials help.

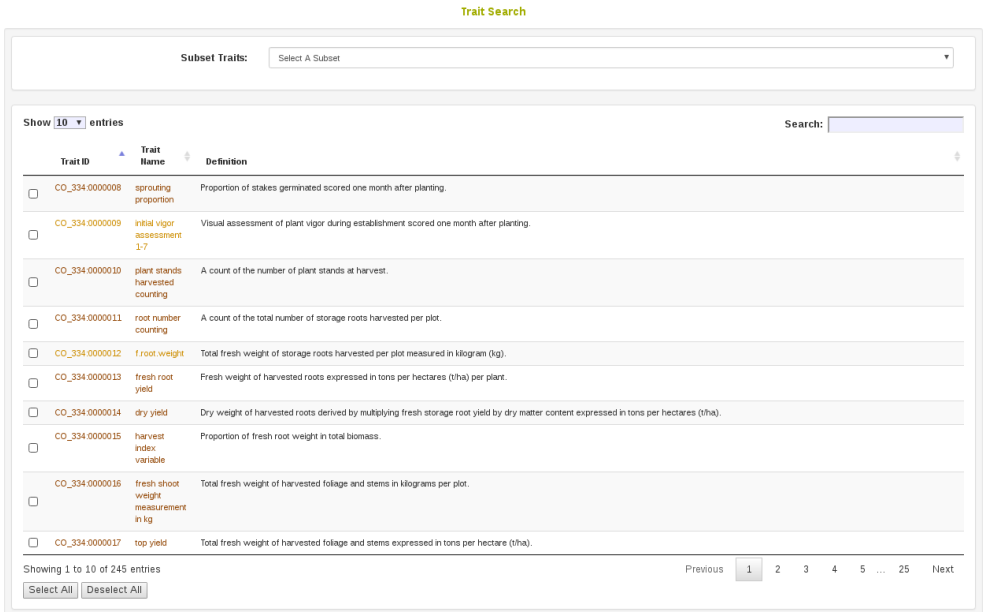
2.3 Trials Search

Trials on the database can be searched based on trial name, description, breeding program, year, location, trial type, design, planting date, and harvest date.

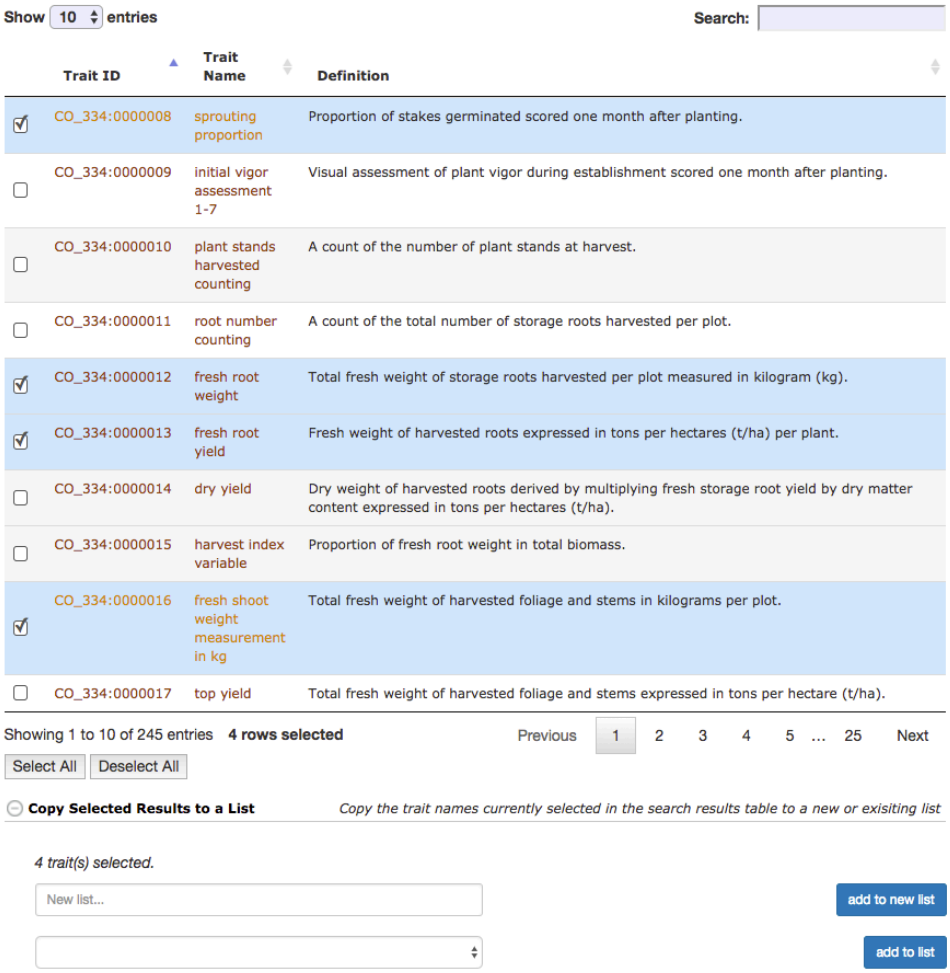


2.4 Trait Search

On the Trait Search page (menu item `Search > Traits`), traits in the database can be searched by ID, name, or descripiton. Optionally, a starting list of traits can be selected to filter down results.



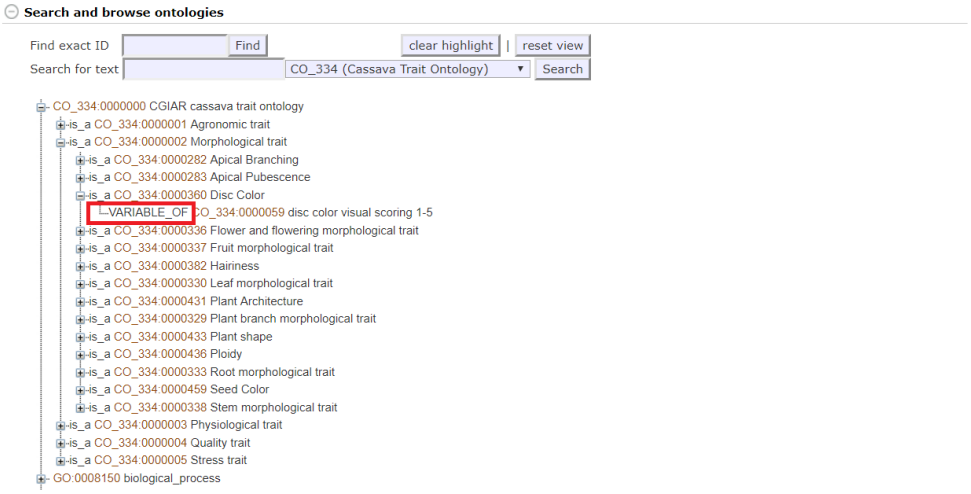
Selecting traits in the results of the search allows one to add the selected results to a trait list, or create a new trait list from the select results.



2.5 Ontology Browser

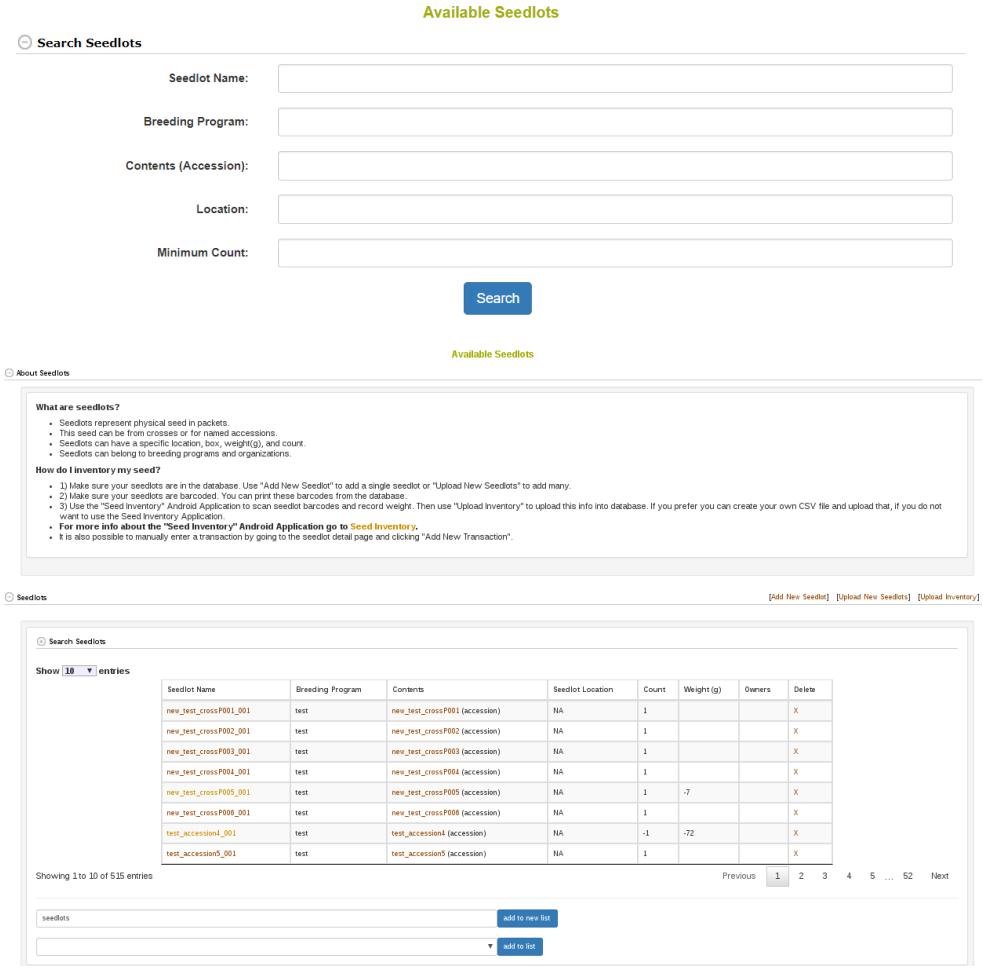
A more advanced tool for searching for Traits is the ontology browser, available by clicking on Analyze and Ontology Browser. From here you can search ontologies and see the various classifications of terms in a tree display.

The terms which appear in the Trait Search in 2.4 are only variable terms. The ontology browser shows these variables as different from their grouping terms by indicating VARIABLE_OF like in the following screenshot.



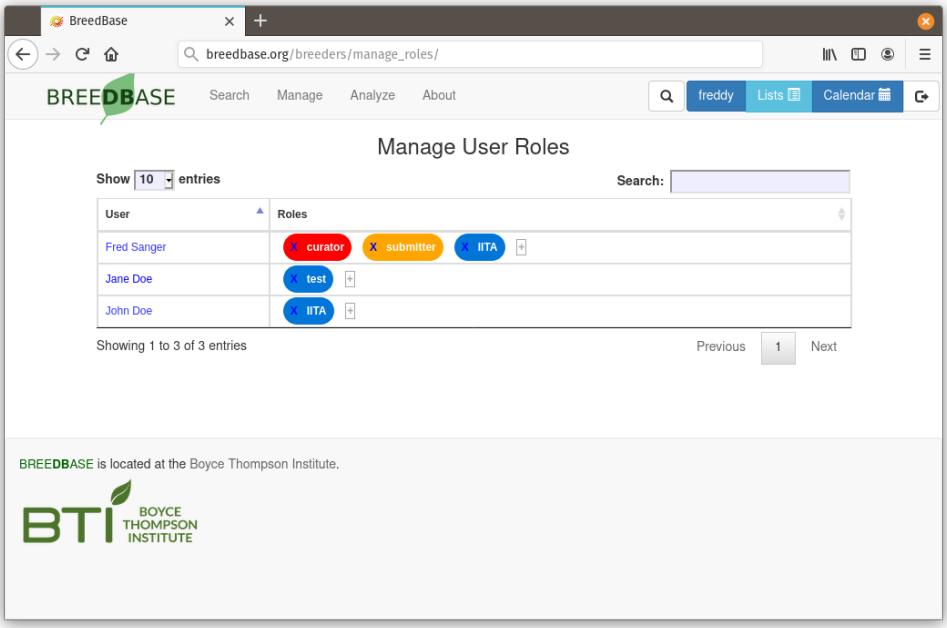
2.6 Search Seedlots

Seedlots are different from Accessions in that they represent the physical seed being evaluated in an experiment. Seedlots have things like physical storage locations and seed quantities, which accessions do not. To search for available seedlots you go to Manage and then click Seed Lots. By clicking Search Seedlots, you can specify query information. The results from your search will be in the table below the search form.


Available Seedlots
Search Seedlots
Seedlot Name:
Breeding Program:
Contents (Accession):
Location:
Minimum Count:
Search
Available Seedlots
About Seedlots
What are seedlots?
Seedlots represent physical seed in packets.
This seed can be from crosses or for named accessions.
Seedlots can have a specific location, box, weight(g), and count.
Seedlots can belong to breeding programs and organizations.
How do I inventory my seed?
1) Make sure your seedlots are in the database. Use "Add New Seedlot" to add a single seedlot or "Upload New Seedlots" to add many.
2) Make sure your seedlots are barcoded. You can print these barcodes from the database.
3) Use the "Seed Inventory" Android Application to scan seedlot barcodes and record weight. Then use "Upload Inventory" to upload this info into database. If you prefer you can create your own CSV file and upload that, if you do not want to use the Seed Inventory Application.
For more info about the "Seed Inventory" Android Application go to Seed Inventory.
It is also possible to manually enter a transaction by going to the seedlot detail page and clicking "Add New Transaction".
Seedlots
[Add New Seedlot] [Upload New Seedlots] [Upload Inventory]
Search Seedlots
Show 10 entries
Seedlot Name
Breeding Program
Contents
Seedlot Location
Count
Weight (g)
Owners
Delete
new_test_crossP001_001 test new_test_crossP001 (accession) NA 1 X
new_test_crossP002_001 test new_test_crossP002 (accession) NA 1 X
new_test_crossP003_001 test new_test_crossP003 (accession) NA 1 X
new_test_crossP004_001 test new_test_crossP004 (accession) NA 1 X
new_test_crossP005_001 test new_test_crossP005 (accession) NA 1 -7 X
new_test_crossP006_001 test new_test_crossP006 (accession) NA 1 X
test_accession4_001 test test_accession4 (accession) NA -1 -72 X
test_accession5_001 test test_accession5 (accession) NA 1 X
Showing 1 to 10 of 515 entries
Previous 1 2 3 4 5 ... 52 Next
seedlots
add to new list
add to list

Chapter 3

Managing User Roles



3.1 What are User Roles?

Every user account in Breedbase has one or more associated "roles" that determine the authorizations (what the user is allowed to do) in the database.

There are three fundamental roles, “curator”, “submitter”, and “user”, which determine basic read/write levels. The “curator” status can read and write everything in the database. The “submitter” status can add information and edit or delete previously submitted information. The “user” type can only read data. Additional roles represent the breeding programs, and are sometimes used to fine-tune write and edit capabilities, as it necessary for multiple users in a breeding program to edit each other’s data.

3.2 The Manage User Roles page

In the “Manage” menu, select the item “User Roles”. This will show the current users in the database with their associated roles. If you are logged in as a curator, the table will show system roles as well as breeding program roles; if you are logged in as a submitter or user, it will show breeding program membership.

If logged in as a “curator”, the roles can be added or deleted.

- To delete a role, click on the X in the role name. A confirm dialog will be displayed to prevent accidental deletion.
- To add a role, click on the plus sign next to the roles. A dialog will pop up with a list of roles. Select the desired role and click “Submit”.
- The new role should be displayed next to the user immediately.
- Role deletions and additions will be effective immediately.

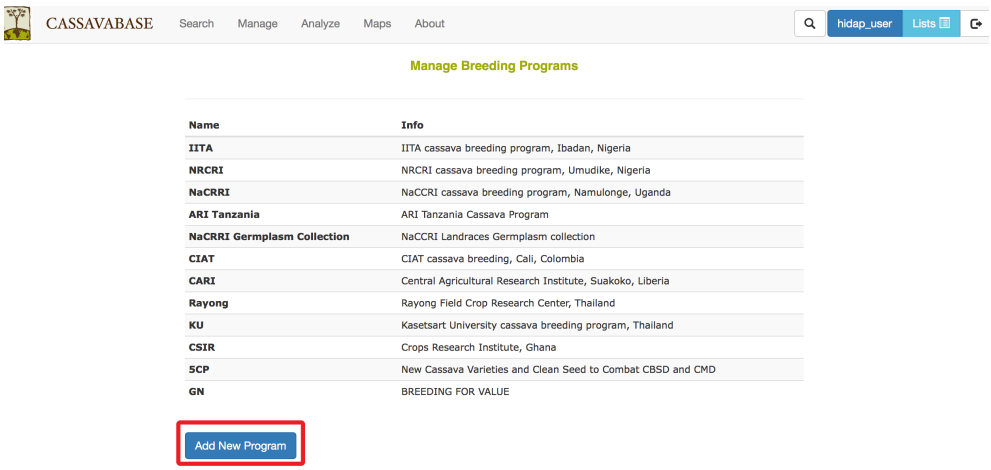
It is recommended that few users be given the “curator” privileges to avoid confusion over data ownership and accidental data overwriting and deletion.

@ref(managing_user_roles)

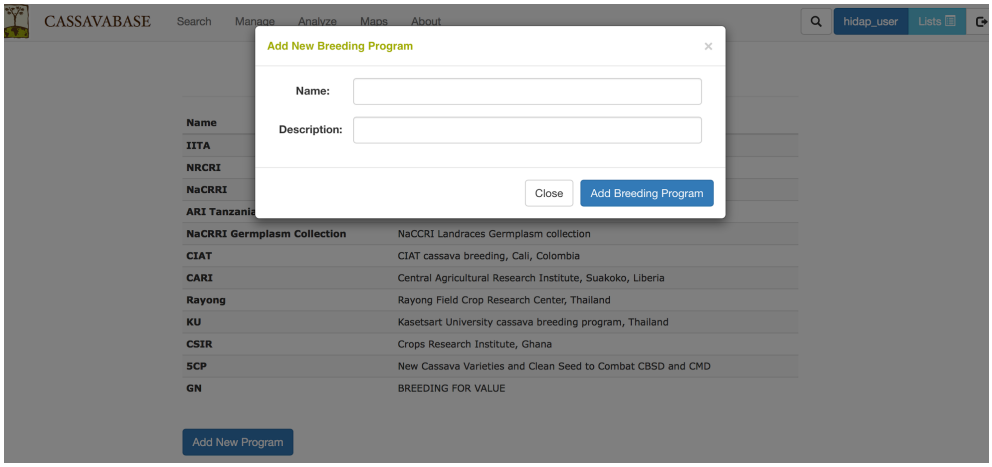
Chapter 4

Managing Breeding Programs

New breeding programs can be added by using "Add New Program" button on the "Manage Breeding Programs" page.



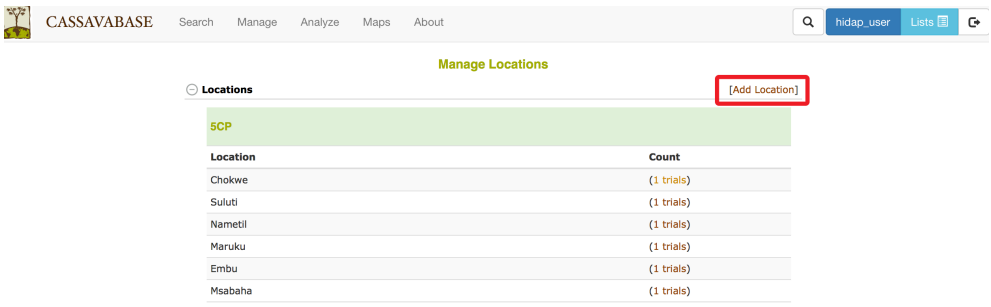
Clicking on the "Add New Program" button will generate a blank form for you to fill out the name and description of the breeding program that you want to add. After completing the form, click on "Add Breeding Program" button to finish the process.


CASSAVABASE
Search
Manage
Analyze
Maps
About
hidap_user
Lists
Add New Breeding Program
Name:
Description:
Close
Add Breeding Program
Name
IITA
NRCRI
NaCRRI
ARI Tanzania
NaCRRI Germplasm Collection
NaCCRI Landraces Germplasm collection
CIAT
CIAT cassava breeding, Cali, Colombia
CARI
Central Agricultural Research Institute, Suakoko, Liberia
Rayong
Rayong Field Crop Research Center, Thailand
KU
Kasetsart University cassava breeding program, Thailand
CSIR
Crops Research Institute, Ghana
5CP
New Cassava Varieties and Clean Seed to Combat CBSD and CMD
GN
BREEDING FOR VALUE
Add New Program

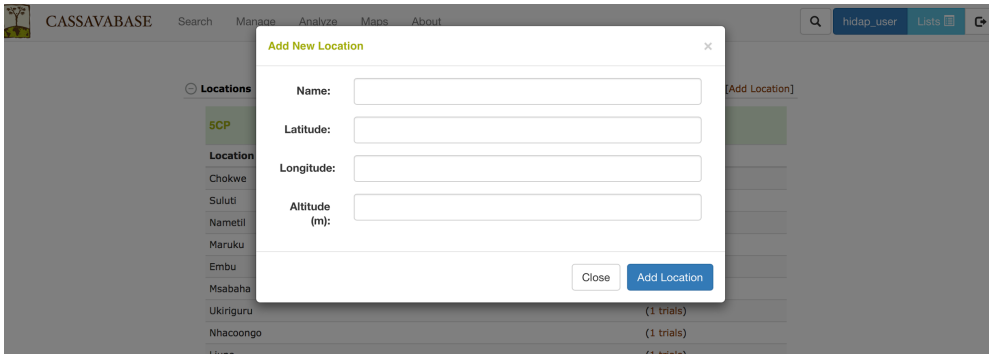
Chapter 5

Managing Locations

Field locations can be managed using the "**Manage Locations**" page. On this page, locations in the database are organized based on their breeding programs. Each location has a link to trials conducted in that location. To add a new location, click on the "Add Location" button that links to the "Add New Location" form.



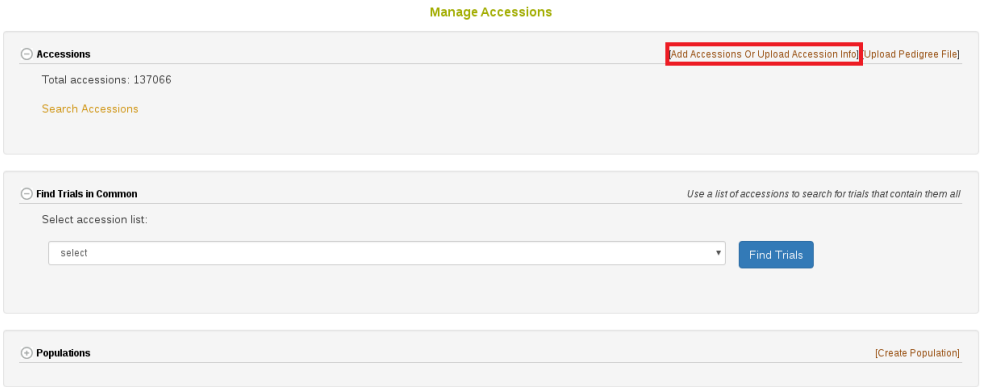
On the "Add New Location" form, fill out the location name that you want to add. Latitude, longitude, and altitude are optional. Submit the new location by clicking on the "Add Location" button at the bottom right of the form.


CASSAVABASE
Search
Manage
Analyze
Maps
About
hidap_user
Lists
Add New Location
Name:
Latitude:
Longitude:
Altitude (m):
Close
Add Location
Locations
[Add Location]
5CP
Location
Chokwe
Suluti
Nametil
Maruku
Embu
Msabaha
Ukiriguru
(1 trials)
Nhacoongo
(1 trials)

Chapter 6

Managing Accessions

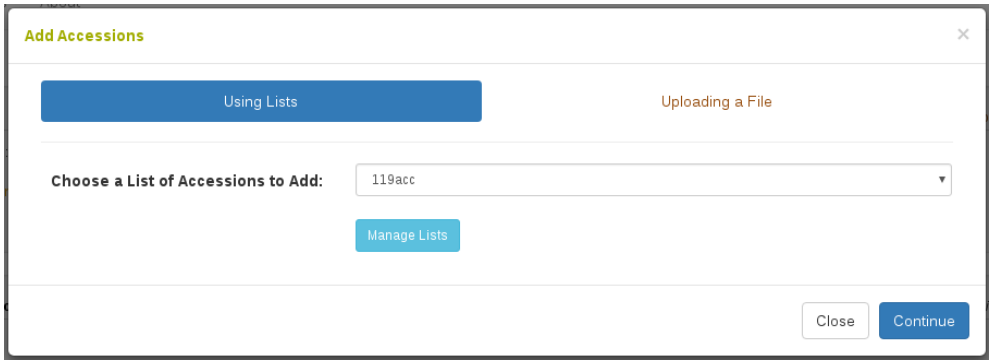
The "Manage Accession" page provides links for adding new accessions. You can choose to add accessions into the database by either using a List you have created or by uploading XLS or XLSX file. Both options will be detailed below. To begin click on the "Add Accessions or Upload Accession Info" link.



This will open a dialog allowing you to select either "Using Lists" or "Uploading a File".

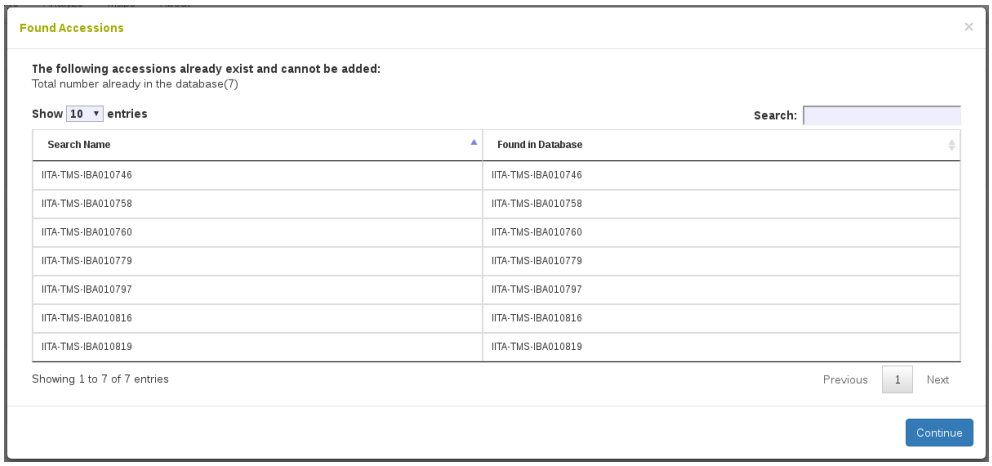
6.1 Add Accessions Using A List

First we will show how to add accessions "Using Lists".

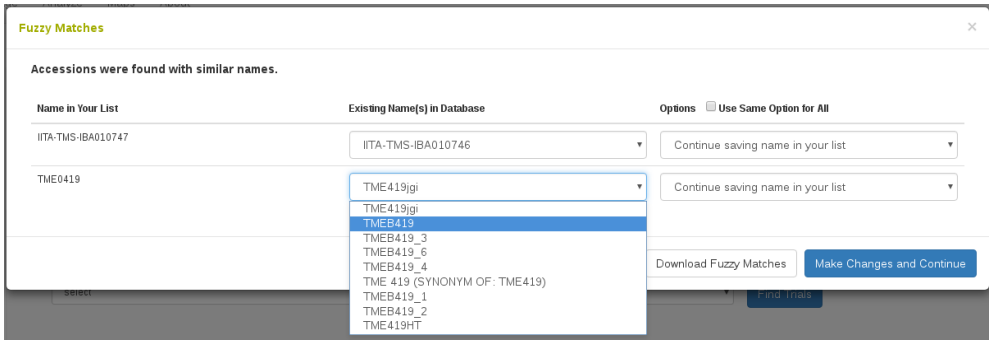


Here you select an accession list which you have previously made. If you need to create or edit your list you can do so now by clicking "Manage Lists". Once you have selected your list you can click "Continue".

The first dialog which can appear will show the accessions which already exist in the database.



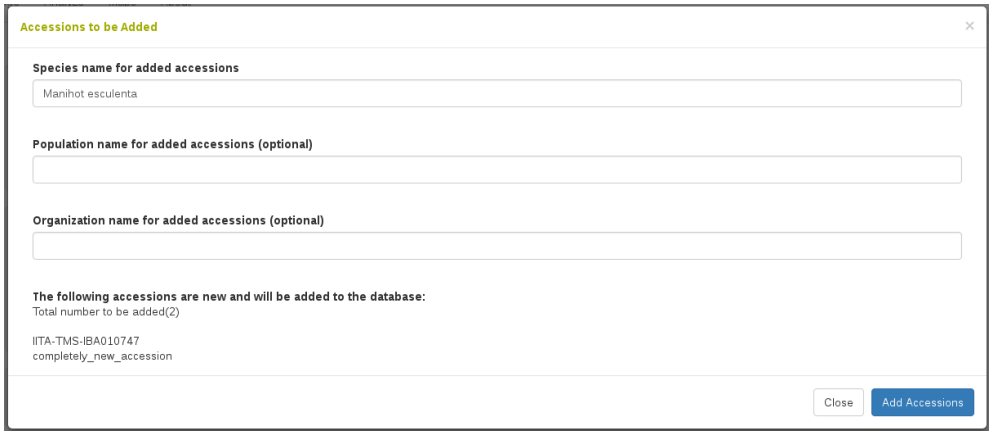
Click "Continue". The next dialog which can appear will show accessions which have very similar matches to the accession names you are adding. In the example below, there are two accession names that are very similar to accession names already in the database. 'TME0419' is very similar to 'TME419', and actually is probably a mistake that should not be added to the database.



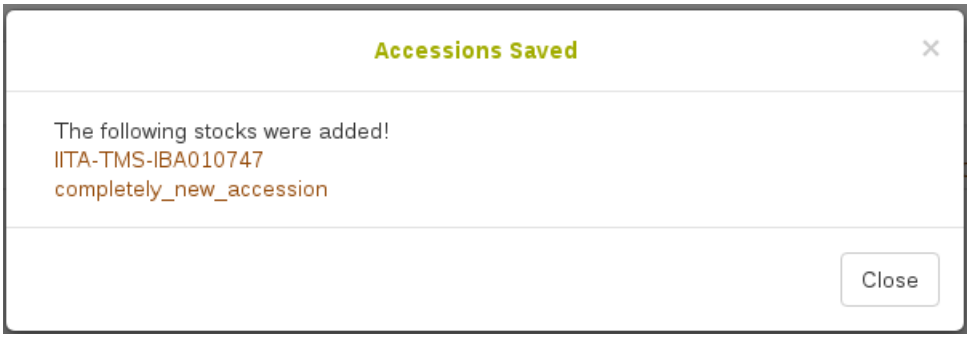
To avoid situations in adding a mistaken duplicate accession, the database gives you options for moving forward with these very similar looking accession names. You can either "continue saving the name in your list", "replace name in your list with selected existing name", "remove name in your list and ignore", or "add name in your list as a synonym to selected existing name".

Clicking "Download Fuzzy Matches" will return a tabular result of the "fuzzy" accession name results shown. Click "Make changes and continue" to move on.

The final dialog shows the accessions that will be added. Here you need to assign the species of these accessions. You can optionally group the accessions into a population and/or add an organization for the accessions.



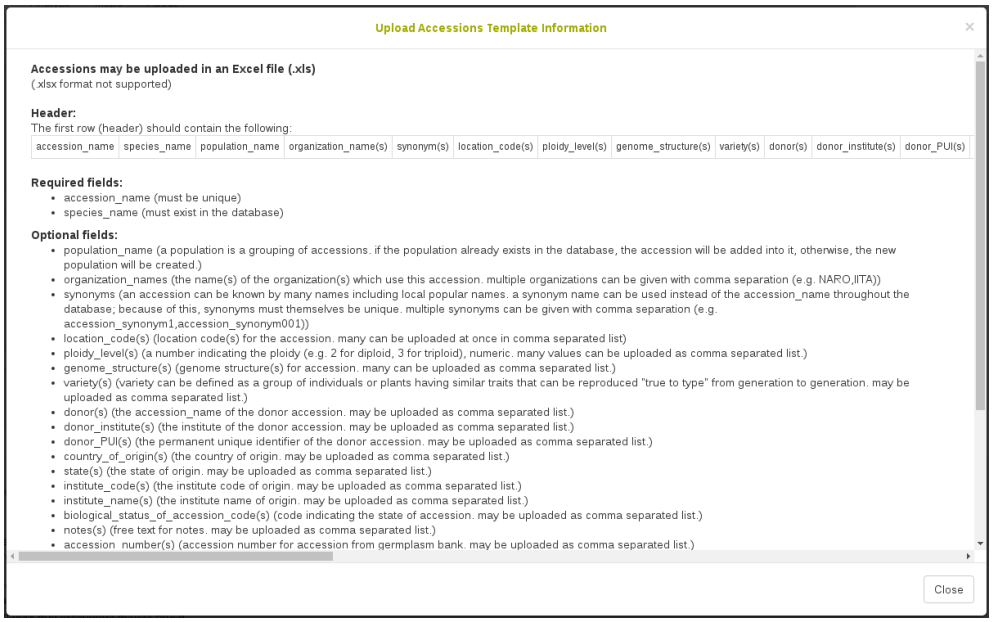
Once you click "Add Accessions", the new accessions will be created in the database and you will see the following confirmation dialog, which includes links to the newly created accessions.



6.2 Uploading Accessions and Accession's Info From A File

The process to upload accessions is very similar to using a list, but enables you to add a variety of properties, such as synonyms, to the accessions in bulk.

Clicking on "Spreadsheet format" will show the following dialog. Here it shows that the file must be XLS or XLSX format and can contain a number of header columns as attributes. It is important that you use exactly the same header column names as listed here. In columns that indicate that many attribute values can be passed at once using (s), such as synonym(s), you can pass a comma separated list of values, such as 'synonym1,synonym2'.



Once you have selected your XLS or XLSX file for upload, click "Continue".

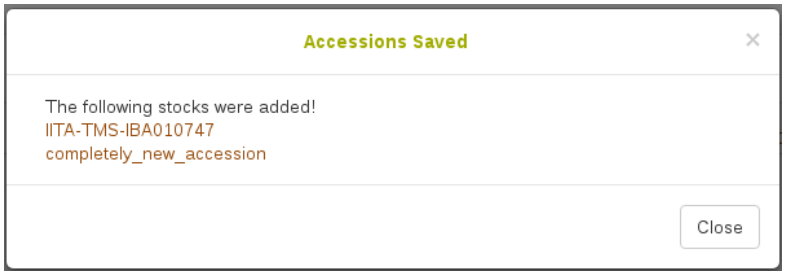
The following process is the same way as with lists:

The first dialog which can appear will show accession names which are already in the database.

Click "Continue" and the next dialog that can appear will show "fuzzy" matches for the accession names you are trying to upload. Here you can choose to prevent adding accession names which look very similar to each other as wrongly duplicated accessions.

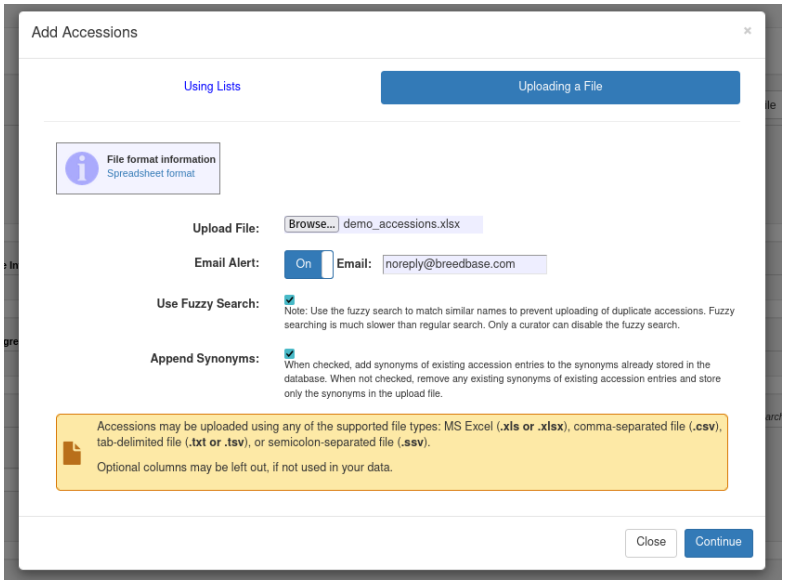
Click "Continue" and the final dialog that will appear will show the information to be added into the database. Here it is divided into accession names that are new and accession names that already exist in the database; however, for the accession names that already exist it will show additional attributes that originated from your file that will be added to these accessions.

Once you click "Add Accessions", the new accessions and information will be created in the database and you will see the following confirmation dialog, which includes links to the created and updated accessions.



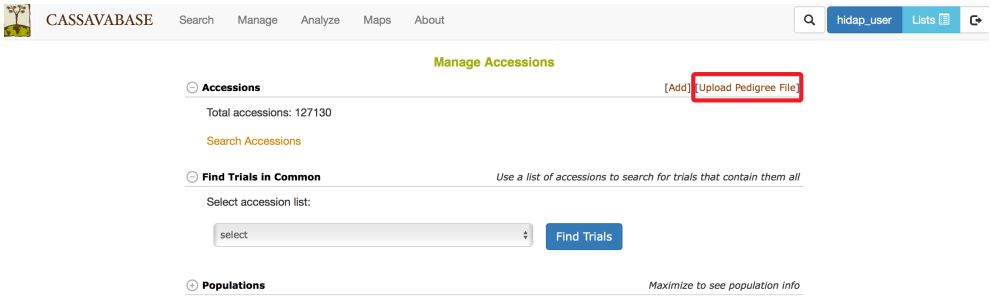
6.3 Email alert for accession upload

When uploading accessions from a file, you have the option to receive email notifications about the status and results of your upload by clicking the "Email Alert" checkbox. By default, the system will use the email address associated with your account, but you have the option to enter a different email address if you prefer. After submitting, the upload process runs in the background, allowing you to continue using the interface without interruptions. Once the process completes, you will receive an email with the upload results, including any warnings or errors that may have occurred during the upload.



6.4 Add Parentage (Pedigree) Information to Accessions

Pedigree data can be uploaded from your computer by clicking on "Upload Pedigree File"



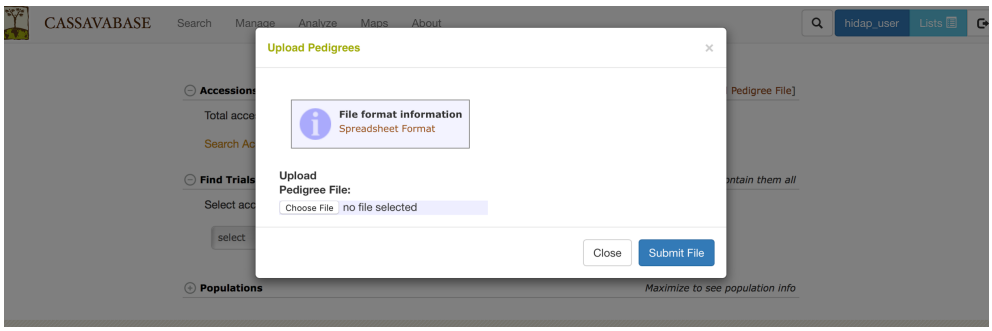
***IMPORTANT!* Please use only tab-delimited text file format (.xls or .xlsx formats are NOT supported).**

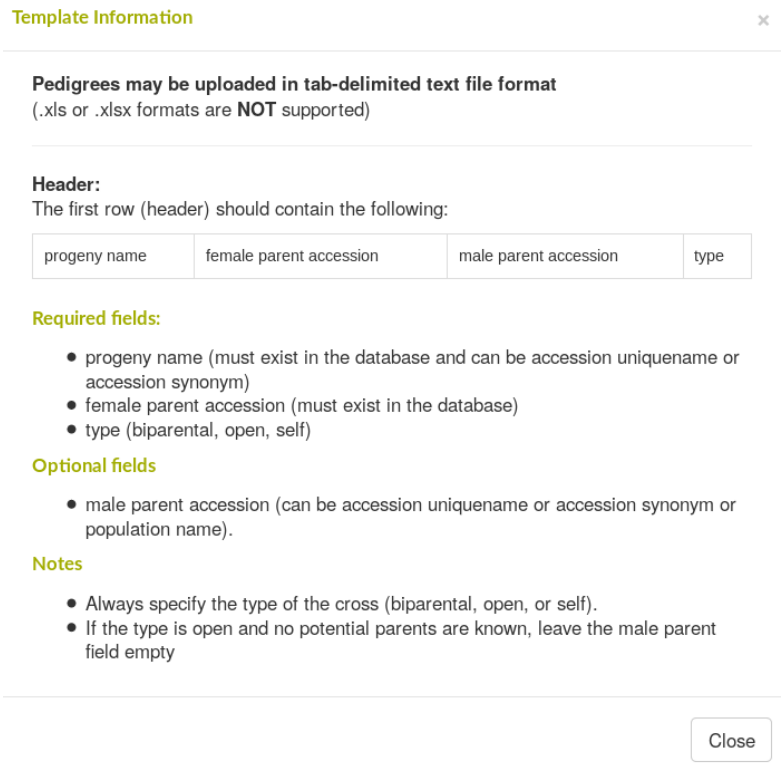
You can find detailed information on how to prepare pedigree file by clicking on "File format information"

The currently supported format has four tab separated columns:

progeny name female parent accession male parent accession type

Type can be biparental, self, backcross, sib, polycross, reselected, or open. In the case of the open type, the male parent accession field can remain blank. For all other types, both columns should be filled, even if they contain the same information as another column (such as self).





6.5 Working with grafts

Grafts are plants that are composed of a rootstock and a scion, which are genetically different and fused together, usually at the stem level.

To work with grafts, the grafts interface needs to be activated by adding a configuration parameter in the sgn_local.conf file. The parameter is show_grafting_interface. It should be set to 1 in sgn_local.conf, the default is 0 (in sgn.conf).

Grafts to be created need to be specified using an Excel file (xlsx format) with two columns. The first column should have the header "scion accession" and should list accession names that will be scions. The second column should have the header "rootstock accession" and should list accession names that will be rootstocks.

In the database, the graft accessions will created as single accessions. The

graft accession will have two relationships, one to the scion accession (scion_of relationship) andone to the rootstock (rootstock_of relationship). These relationships are displayed on the pedigree viewer. The graft accession name is created from the scion accession name and the rootstock accession name, separated by the graft separator character. By default, the graft separator character is the plus sign '+'. The graft separator character can be changed in the sgn_local.conf file, using the parameter graft_separator_string. The graft separator string should not occur in any other accession names that are not grafts.

When the grafting interface is activated, a new button will be shown on the manage accessions page, called "Upload Grafts".

Clicking the button brings up the upload grafts dialog.

Select the Excel file containing the grafting information. The system will validate the file, for example, check whether the accessions are in the database, and if the headers are correct.

The validation result will be presented, and if problems are found, they will be listed. In addition, if there are problems, the Upload button will be grayed out and upload will not be possible. Conversely, if there are no problems, the Upload button will be activated and can be clicked to upload the data.

If the upload completes, a completion message is displayed with a summary what was uploaded.

Grafted accessions can be used like any other accession, for example, they can be used on field layouts. If you create a list of graft accessions, use the list type 'accessions'.

Note that you shouldn't create new grafts based on other grafts. The scion accession and the rootstock accession have to be different, otherwise they will not be created.

6.6 Bulk renaming of accessions

Accessions can be renamed in bulk using the rename accessions feature. To rename accessions, prepare a tab delimited file with two columns: the first column should have the header "old name" and contain the accession names

that need to be changed. The second column should have the header "new name" and contain the names that the accessions in column 1 should be renamed to.

The accession renaming feature is available from the Manage->Accessions page. Click on the "Rename Accessions" button. The first step is the upload of the file with a verification step. The verification step checks whether all the accession names in column 1 exist in the database, and whether all the accession names given in column 2 do NOT exist in the database. Only if both conditions are met, will the "rename" button become active, otherwise an error message is displayed listing the offending accession names.

Optionally, the old name can be automatically added as a synonym to the renamed accession, using the checkbox on the submit form. This option is clicked by default. Unclick the checkbox to NOT save any old names as synonyms.

Note that accession renaming should not be undertaken lightly. This feature is intended for special use cases, such as where accessions are created in a nursery with a name that is different from the accession name in the downstream breeding program.

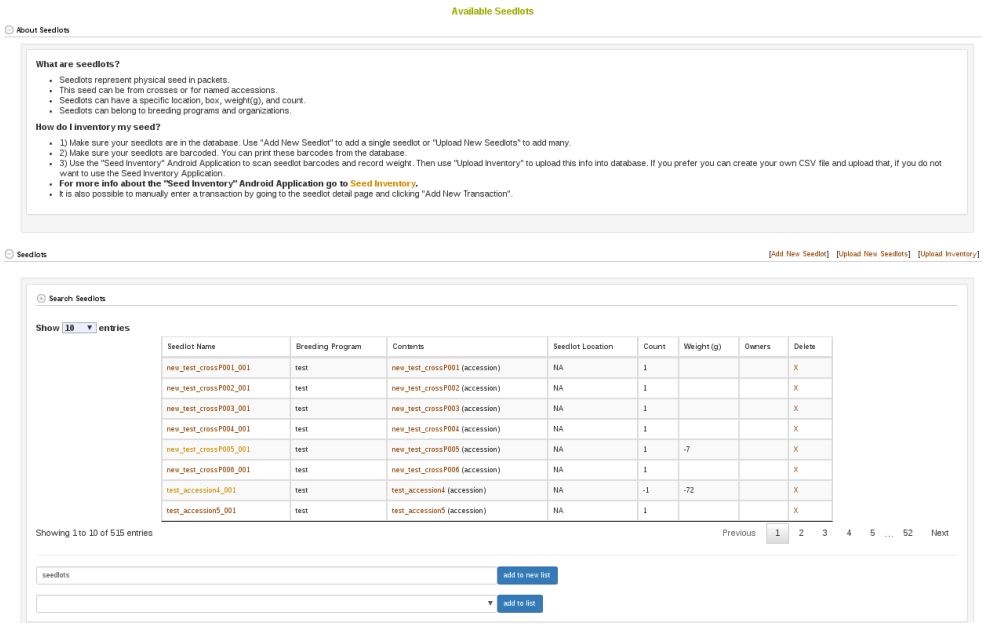
It can also be used to rename accessions in bulk that have spelling mistakes and other issues. Please note however, that the tool does not make any attempt to change the names of associated elements, such a plots, that may have been constructed using accession names.

Because of the many implications of accession renaming, the feature is limited to accounts with the curator role.

Chapter 7

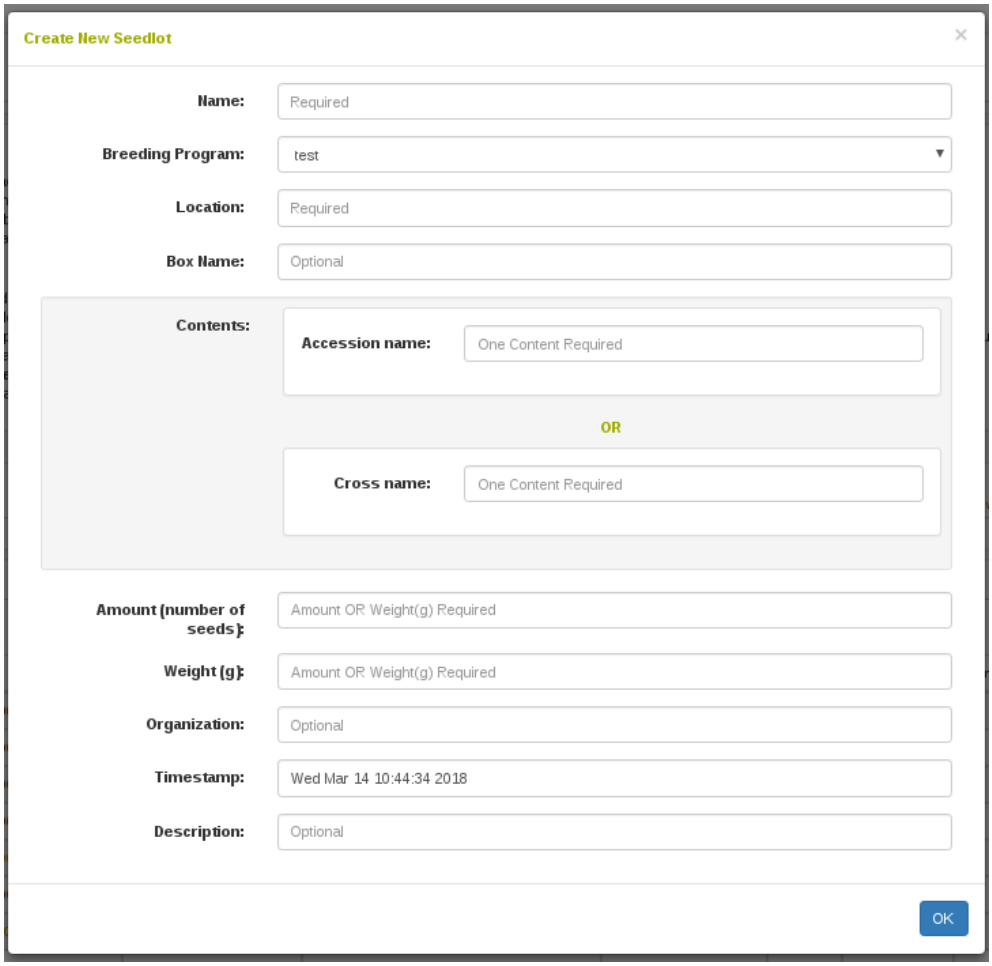
Managing Seed Lots

Seedlots are different from Accessions in that they represent the physical seed being evaluated in an experiment. Seedlots have things like physical storage locations and seed quantities, which accessions do not. The seed in seedlots can be from crosses or can be named accessions. Seedlots from crosses would represent seed harvested. Click Manage and then Seed Lots to begin.



7.1 Add New Seedlot(s)

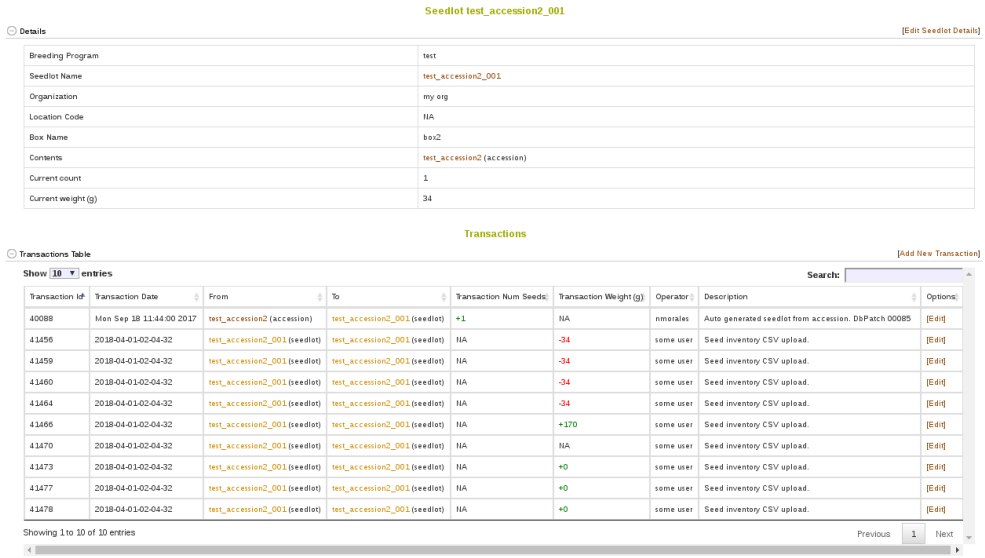
To add a single new seedlot, click on "Add Seedlot". This will bring up the following dialog where you enter information about where the seedlot exists, what accession or cross is contained in it, and how many seeds there are. A seedlot must contain either an accession or a cross, and not both. A seedlot must have a weight in grams or a seed count or both of these.



In the case where you have many seedlots to add to the database, you can upload an excel XLS or XLSX file instead. Click "Upload Seedlots" to see the following dialog.

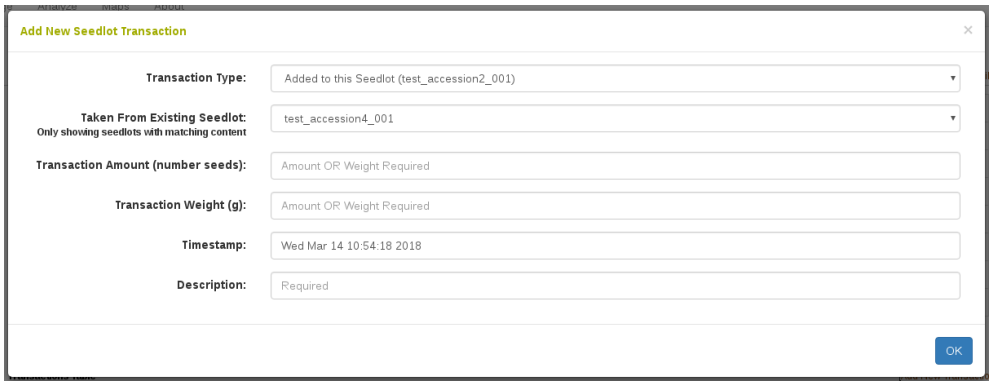
7.2 Seedlot Transactions

Seedlots are capable of tracking where seeds came from, such as from crosses, and to where seeds go, such as to plots in the field. If you navigate to a seedlot detail page you will see the following.



On this page you see and can edit information regarding a single seedlot,

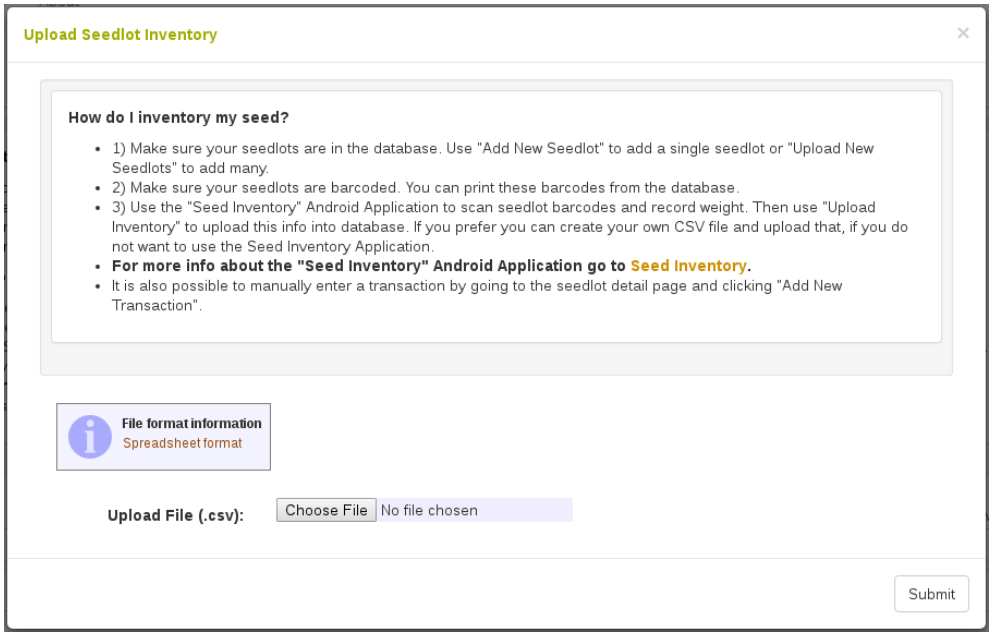
such as its name and location. You will also see a table indicating all t he transactions that a seedlot has been involved in, such as if it was planted in a plot in the field. Transactions to field plots are created when adding or uploading a new trial or from a trial's detail page. Clicking on "Add New Transaction" let you add a transaction from between this seedlot and another seedlot. This kind of transaction is useful for representing if you have distributed seed to different locations.



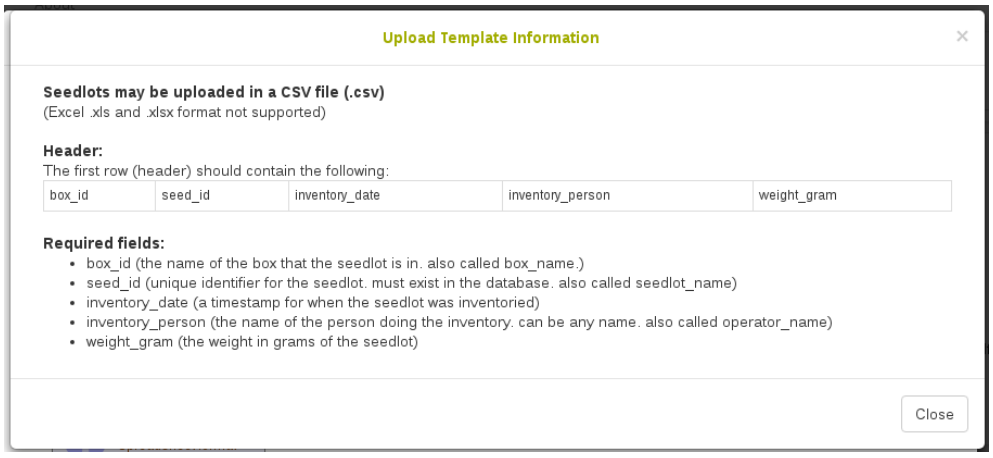
7.3 Seed Inventory

To inventory your seed: 1) Make sure your seedlots are in the database. Use "Add New Seedlot" to add a single seedlot or "Upload New Seedlots" to add many. 2) Make sure your seedlots are barcoded. You can print these barcodes from the database. 3) Use the "Inventory" Android Application to scan seedlot barcodes and record weight. Then use "Upload Inventory" to upload this info into database. If you prefer you can create your own CSV file and upload that, if you do not want to use the Inventory Application. For more info about the "Inventory" Android Application go to Inventory.

Clicking the "Upload Inventory" button will bring the following dialog:



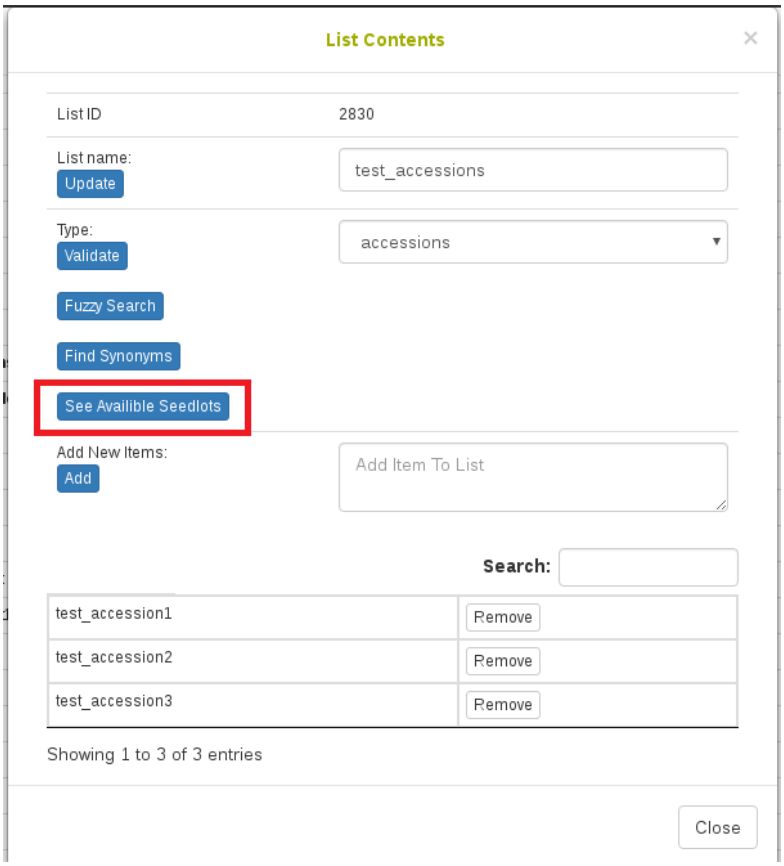
The CSV file that should contain your inventory should meet these Template requirements. The Seed Inventory Android Application exports this exact file.



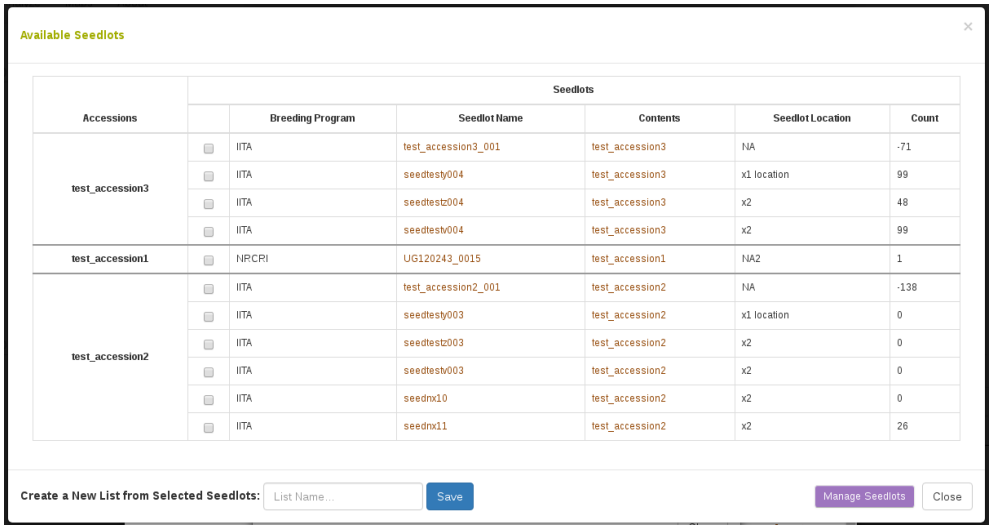
7.4 Find Seedlots For a List of Accessions

A convenient tool for searching available seedlots for a list of accessions is available in the list tool. First open up your list of accessions. For help

opening a list of accessions please see the List section help. There is a button called "See Available Seedlots".

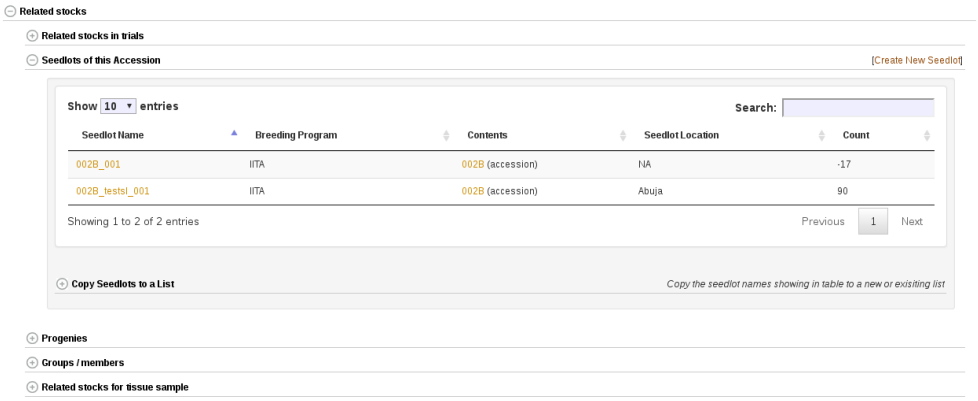


Once you click this, you will see the following table in a dialog. From here you can create a list of seedlots using the checkboxes and the input at the bottom.



7.5 Create a seedlot for an Accession or Cross

Complementary to what we saw above for creating seedlots from the "Manage Seedlots" page, it is possible to create a new seedlot from an accession's detail page or from the cross detail page. On the accession detail page, this is visible in the "Related Stocks" section as seen below. The cross detail page has an identical section. Notice the link for creating a new seedlot, which streamlines adding the seedlot.



7.6 Add quality data to a seedlot

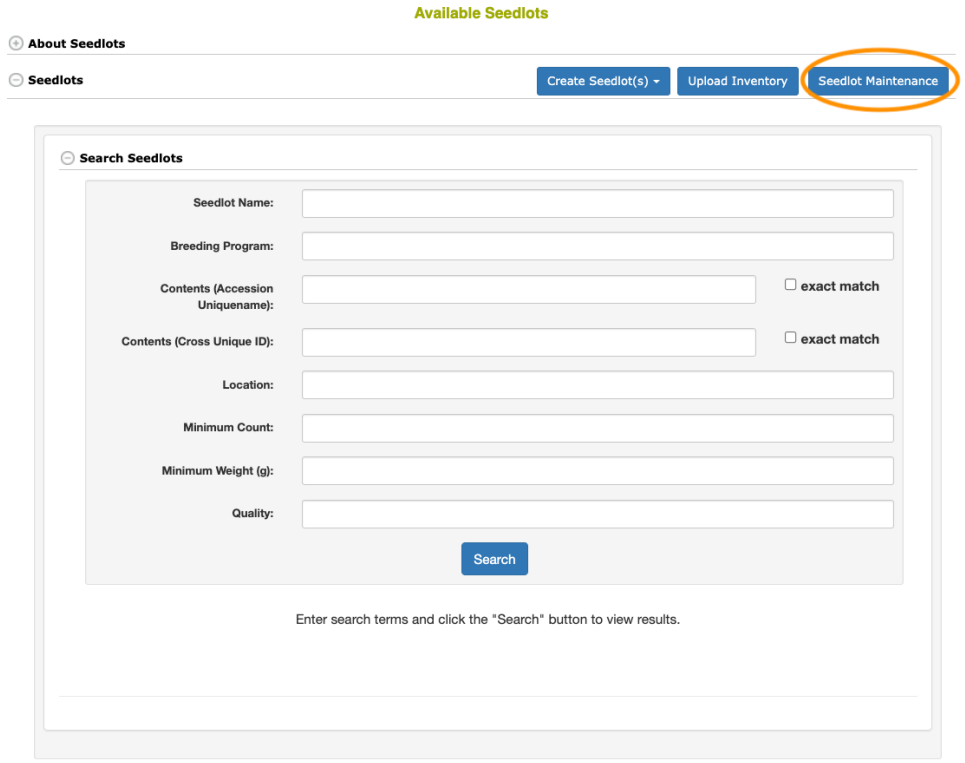
Quality information can be added to a seedlot in the quality field. This is also available as a column in the file upload format. It is recommended to use a controlled vocabulary, defined by the user, for the quality field. For example, good quality seed should be labelled “ok”, whereas other quality descriptors could be “moldy”, “insect damage”, or “low sprouting”, etc.

7.7 Seedlot Maintenance Events

For some crops, such as sugar kelp, a “seedlot” requires routine maintenance for the successful long-term storage of the seedlot. (For example, a Seedlot Maintenance Event for sugar kelp would be the routine change of the water that gametophytes are kept it). Breedbase can now store a record of these Seedlot Maintenance Events associated directly with existing Seedlots. Maintenance Events can be uploaded using a simple Excel template or recorded directly on the website.

7.7.1 Setup

Each Breedbase instance needs to be configured to support the storage of Seedlot Maintenance Events since each crop will have their own distinct set of maintenance events for their seedlots. To check if your Breedbase instance supports this feature, go to the Manage menu and select the Seed Lots page. Make sure you are logged in and look for the **Seedlot Maintenance** button near the top, next to the **Create Seedlot(s)** and **Upload Inventory** buttons. If you don’t see this button, contact the developer(s) supporting your Breedbase instance and ask if they can setup this feature.



The location of the Seedlot Maintenance button on the Manage > Seed Lots page

7.7.2 Adding Events

Seedlot Maintenance Events can be added using two methods: 1) Uploading an Excel template or 2) Recording events directly on the website

Uploading Events with Excel Template

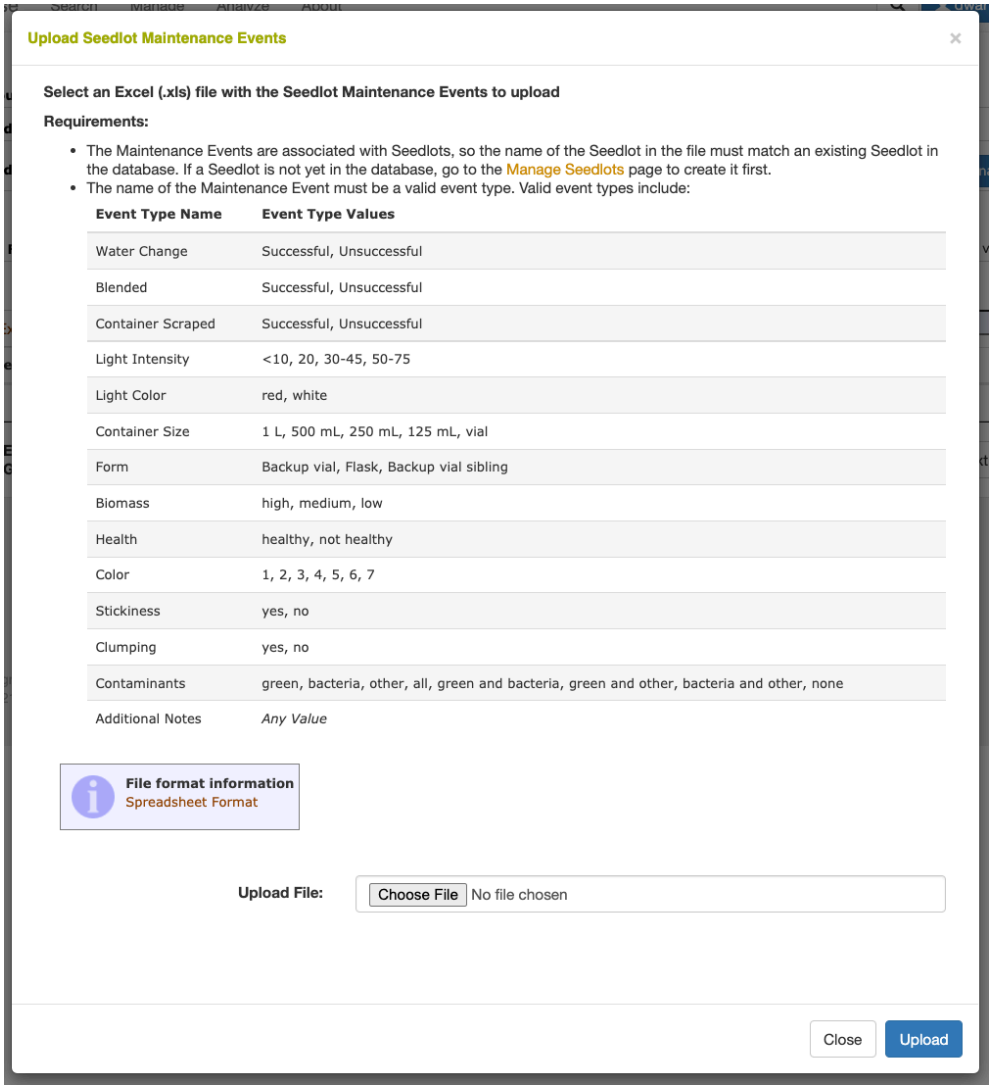
To bulk-upload a file of Seedlot Maintenance Events, first create an Excel (.xls or .xlsx) file with the following headers:

- **seedlot** - the name of the Seedlot to associate the event with (must exactly match an existing Seedlot in the database)
- **type** - the name of the Seedlot Maintenance Event type (these vary between Breedbase instances, a list of supported event types is displayed

on the upload page)
- **value** - the value of the Seedlot Maintenance Event (these may be different for each event type and vary between Breedbase instances, a list of supported event values is displayed on the upload page)
- **notes** - optional, additional notes/comments about the event
- **operator** - the username of the Breedbase user that recorded the event
- **timestamp** - the date/time the event was recorded, in 'YYYY-MM-DD HH:MM:SS' format

Once you have an Excel file with the events filled out, follow these steps to upload the events to the database:

1. Make sure you are logged in to your Breedbase instance
2. Go to the Manage > Seed Lots page
3. Select the **Seedlot Maintenance** button
4. Select the **Upload Maintenance** button
5. Choose your Excel (.xls or .xlsx) file to upload
6. Select the **Upload** button



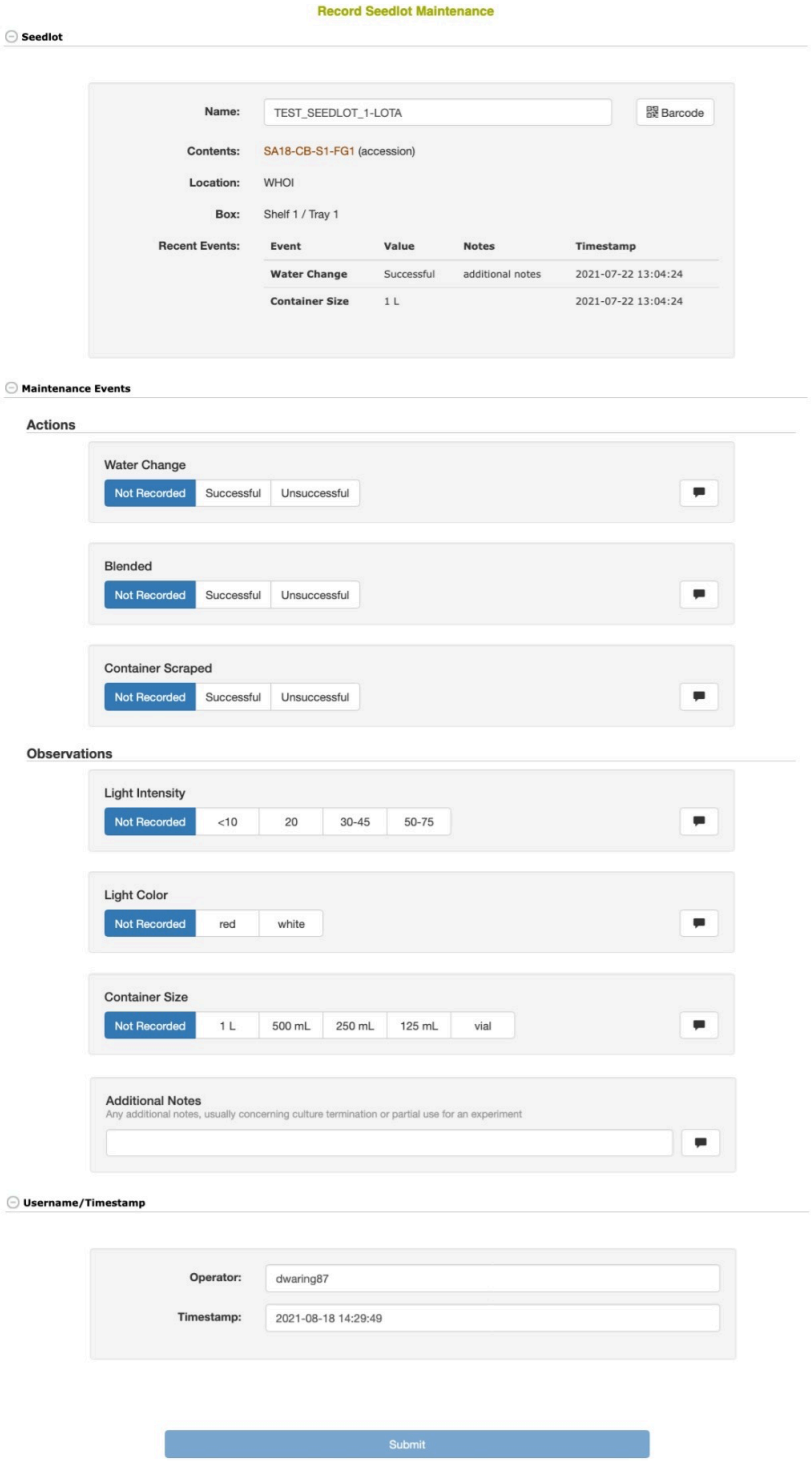
The Seedlot Maintenance upload dialog, showing the supported event types and values (for sugar kelp)

Recording Events on Website

To add individual Seedlot Maintenance Events to the database in real time, as they're being recorded, use the **Record Maintenance** page. Follow these steps to record Seedlot Maintenance Events:

1. Make sure you are logged in to your Breedbase instance

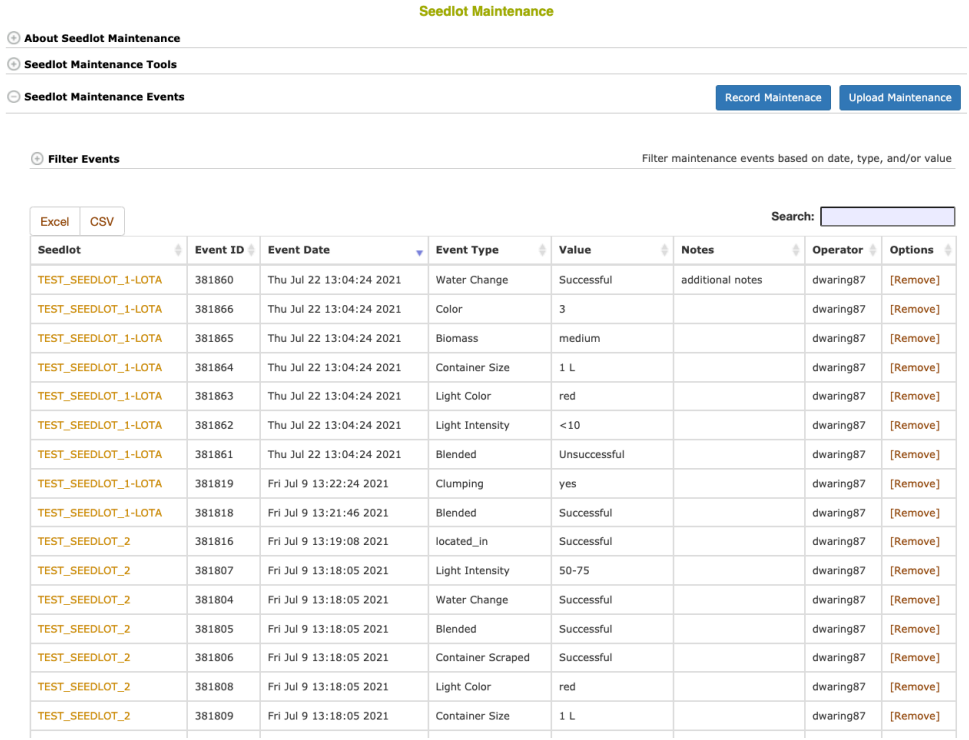
2. Go to the Manage > Seed Lots page
3. Select the **Seedlot Maintenance** button
4. Select the **Record Maintenance** button
5. Enter the **Seedlot Name** or scan a barcode that has the Seedlot Name encoded. Once entered, the box at the top of the page will display basic information about the Seedlot as well its recently recorded events.
6. Select or Enter the values of individual events
7. Optionally, notes button next to each event to add additional notes/comments about that specific event
8. Make sure the operator/username and timestamp are correct
9. Select the **Submit** button to add the recorded events to the database. NOTE: any events that remain selected as "Not Recorded" will not be submitted to the database.


Record Seedlot Maintenance
Seedlot
Name:
TEST_SEEDLOT_1-LOTA
Barcode
Contents:
SA18-CB-S1-FG1 (accession)
Location:
WHOI
Box:
Shelf 1 / Tray 1
Recent Events:
Event
Value
Notes
Timestamp
Water Change
Successful
additional notes
2021-07-22 13:04:24
Container Size
1 L
2021-07-22 13:04:24
Maintenance Events
Actions
Water Change
Not Recorded
Successful
Unsuccessful
Blended
Not Recorded
Successful
Unsuccessful
Container Scraped
Not Recorded
Successful
Unsuccessful
Observations
Light Intensity
Not Recorded
<10
20
30-45
50-75
Light Color
Not Recorded
red
white
Container Size
Not Recorded
1 L
500 mL
250 mL
125 mL
vial
Additional Notes
Any additional notes, usually concerning culture termination or partial use for an experiment
Username/Timestamp
Operator:
dwaring87
Timestamp:
2021-08-18 14:29:49
Submit

The Seedlot Maintenance record page, as configured for sugar kelp

7.7.3 Displaying Events

Recently recorded Seedlot Maintenance Events are displayed in a table from the main Seedlot Maintenance page, as well as the detail page for individual Seedlots.



Seedlot Maintenance

About Seedlot Maintenance

Seedlot Maintenance Tools

Seedlot Maintenance Events — Record Maintenace — Upload Maintenance

Filter Events — Filter maintenance events based on date, type, and/or value

Excel CSV — Search:

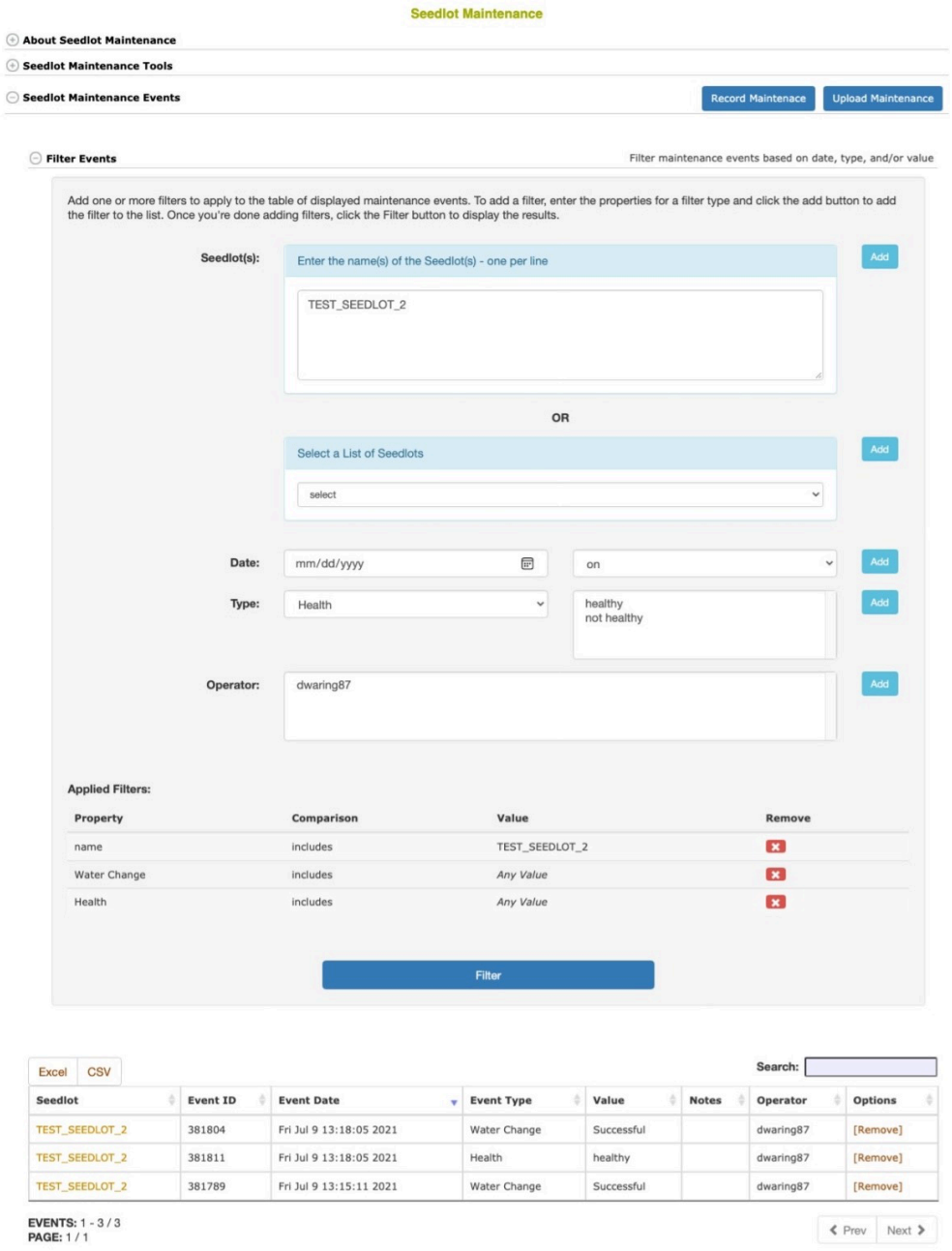
Seedlot	Event ID	Event Date	Event Type	Value	Notes	Operator	Options
TEST_SEEDLOT_1-LOTA	381860	Thu Jul 22 13:04:24 2021	Water Change	Successful	additional notes	dwaring87	[Remove]
TEST_SEEDLOT_1-LOTA	381866	Thu Jul 22 13:04:24 2021	Color	3		dwaring87	[Remove]
TEST_SEEDLOT_1-LOTA	381865	Thu Jul 22 13:04:24 2021	Biomass	medium		dwaring87	[Remove]
TEST_SEEDLOT_1-LOTA	381864	Thu Jul 22 13:04:24 2021	Container Size	1 L		dwaring87	[Remove]
TEST_SEEDLOT_1-LOTA	381863	Thu Jul 22 13:04:24 2021	Light Color	red		dwaring87	[Remove]
TEST_SEEDLOT_1-LOTA	381862	Thu Jul 22 13:04:24 2021	Light Intensity	<10		dwaring87	[Remove]
TEST_SEEDLOT_1-LOTA	381861	Thu Jul 22 13:04:24 2021	Blended	Unsuccessful		dwaring87	[Remove]
TEST_SEEDLOT_1-LOTA	381819	Fri Jul 9 13:22:24 2021	Clumping	yes		dwaring87	[Remove]
TEST_SEEDLOT_1-LOTA	381818	Fri Jul 9 13:21:46 2021	Blended	Successful		dwaring87	[Remove]
TEST_SEEDLOT_2	381816	Fri Jul 9 13:19:08 2021	located_in	Successful		dwaring87	[Remove]
TEST_SEEDLOT_2	381807	Fri Jul 9 13:18:05 2021	Light Intensity	50-75		dwaring87	[Remove]
TEST_SEEDLOT_2	381804	Fri Jul 9 13:18:05 2021	Water Change	Successful		dwaring87	[Remove]
TEST_SEEDLOT_2	381805	Fri Jul 9 13:18:05 2021	Blended	Successful		dwaring87	[Remove]
TEST_SEEDLOT_2	381806	Fri Jul 9 13:18:05 2021	Container Scraped	Successful		dwaring87	[Remove]
TEST_SEEDLOT_2	381808	Fri Jul 9 13:18:05 2021	Light Color	red		dwaring87	[Remove]
TEST_SEEDLOT_2	381809	Fri Jul 9 13:18:05 2021	Container Size	1 L		dwaring87	[Remove]

Unfiltered table of recent Seedlot Maintenance events

The events displayed in these tables are sorted by timestamp, with the most recently recorded events displayed first. The displayed events can be filtered using any number of supported filter criteria, such as: - seedlot names (as entered on the page or using an existing seedlot list), - dates (on, on or before, before, on or after, and/or after the entered dates) - event types - event type values - operator/username

Select the properties of the filter(s) you want to apply, then select the **Add** button next to the button to add the filter to the list of applied filters. Once

you're done adding filters, select the **Filter** button to search the database for the filtered events.



A filtered table of Seedlot Maintenance events

The filtered events can be downloaded directly from the table using the **Excel** or **CSV** buttons at the top of the table. Or Seedlot Maintenance Events can be bulk-downloaded (this includes all events for a Seedlot) using a list of Seedlots from the main downloads page (see below).

7.7.4 Downloading Events

To bulk-download all events for a specific subset of Seedlots:

1. Create a list containing the Seelots you are interested in.
2. Go to the **Download Using Lists** page (Manage > Download)
3. Find the **Download Seedlot Maintenance Events** section
4. Select your list of Seedlots
5. Select the **Download** button to generate the download file

The downloaded file will follow the same format as the upload template and will contain all recorded Seedlot Maintenance Events for each Seedlot in the list.

7.8 Deleting Seedlots

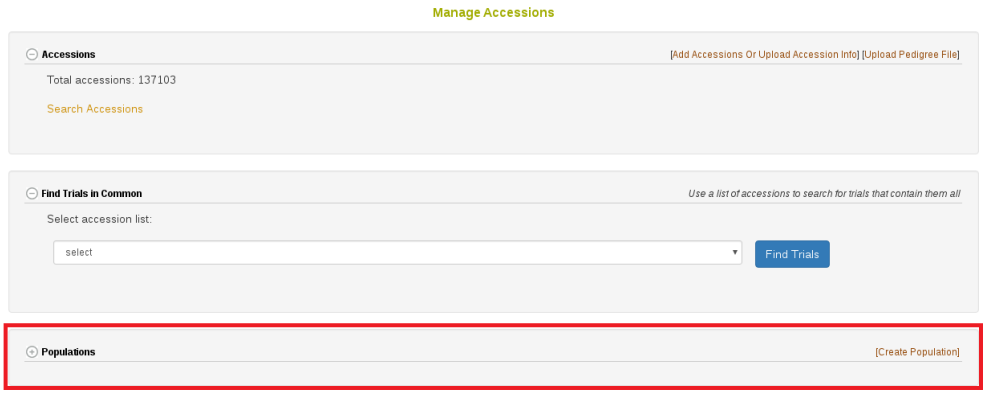
Seedlots can be deleted on the Manage Seedlots page (/breeders/seedlots) by search the seedlot and then clicking the X to delete one seedlot at a time. To delete a seedlot, the logged in user needs the required delete privileges on the seedlot. The seedlot also should not have any transactions associated with it (except for the initial transaction).

To delete seedlots in bulk, generate a list of type seedlot, for example, using the wizard. Open the section “Delete seedlots using a list” on the Manage Seedlots page and select the list. Seedlot deletion using a list is only available to user with curator status.

Chapter 8

Managing Populations

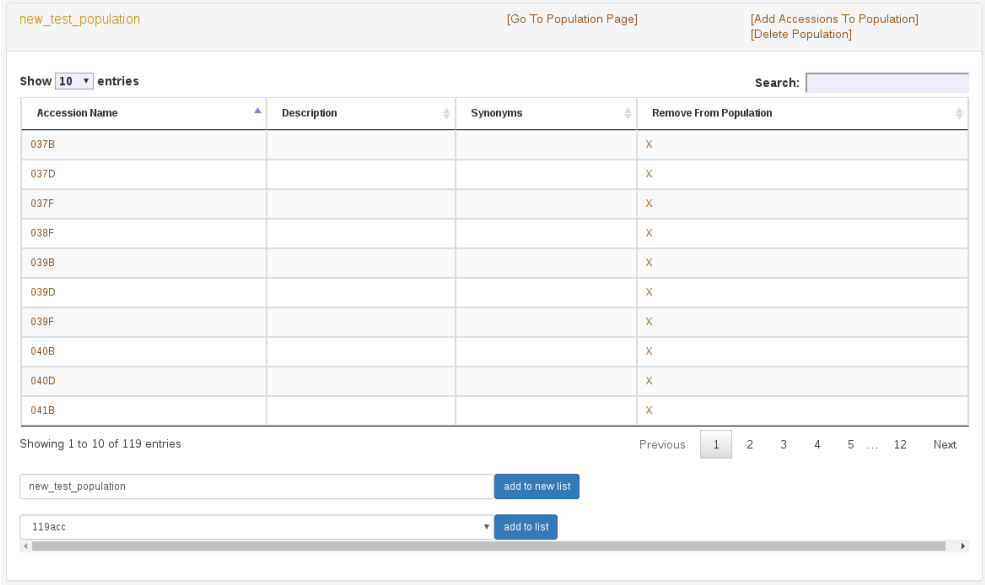
Populations are modeled as groups of accessions. This grouping can be useful in downstream analyses. To manage these populations go to Manage Accessions and scroll tp the bottom.



To add a new population click “Create Population”. The following dialog will appear where you choose a list of accessions and give a name to the new population. Please note it is also possible to create a population when you are uploading new accessions into the database.

Click on the plus (+) button next to Populations to see all the available populations. Click on a population name to see the accessions in the population.

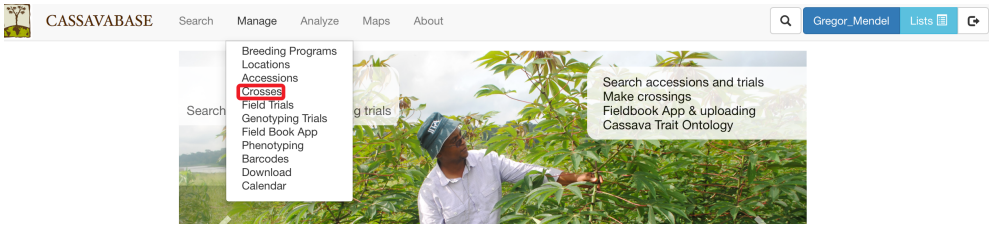
From here you can delete accessions from a population as well as add new accessions to the population.



Chapter 9

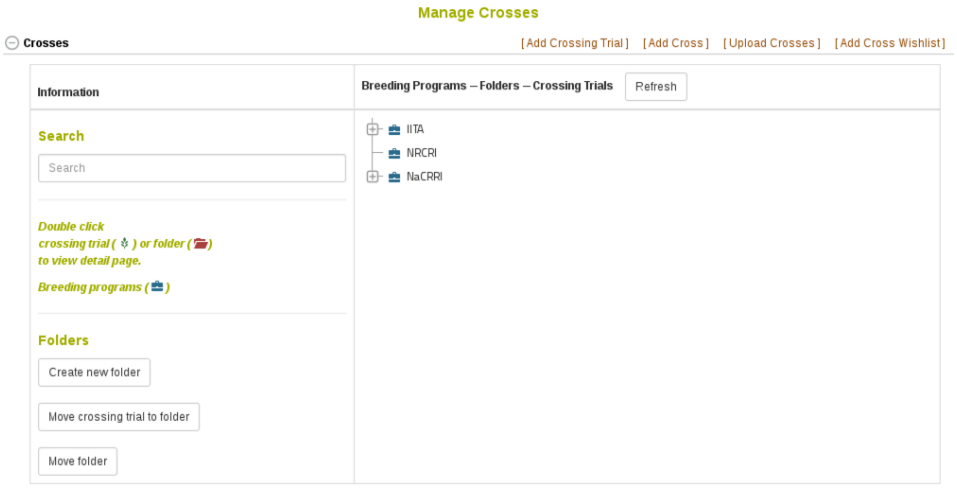
Managing Crosses

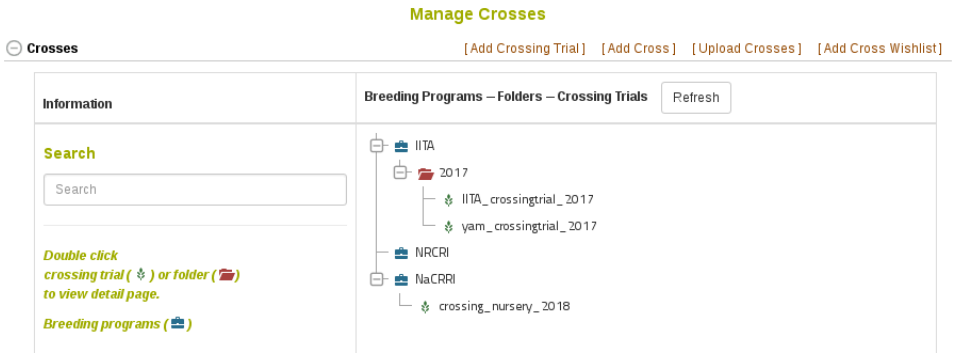
Information for crosses can be managed using the "Crosses" option in the Manage menu.



9.1 Crossing Experiment

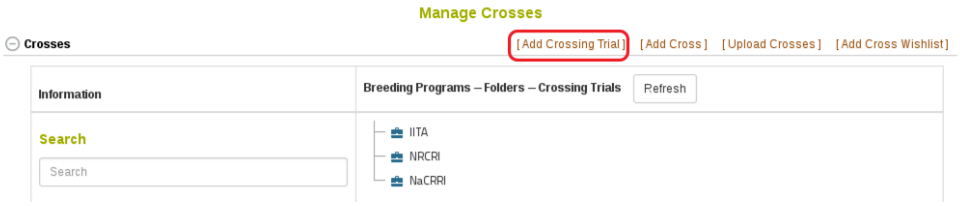
Different crosses in the same trial/nursery/project are grouped via "**crossing experiment**". Crossing experiments are organized based on their breeding programs. To find a crossing experiment, you can either type the crossing experiment name in the "Search" box, or look for the crossing experiment directly in its breeding program by clicking on the "+" icon. In each breeding program, crossing experiments can be placed directly in the breeding program, or organized in folders. The "**Folders**" section allows you to place crossing experiments in folders, move a crossing experiment in a folder to another folder, or rearrange your folders within a breeding program.





9.1.1 Add New Crossing Experiment

To add a new crossing experiment, click on "Add Crossing Experiment" link.



Required Information:

- "**Crossing Experiment Name**": enter a name for the crossing experiment. The crossing experiment name must not already exist in the database.

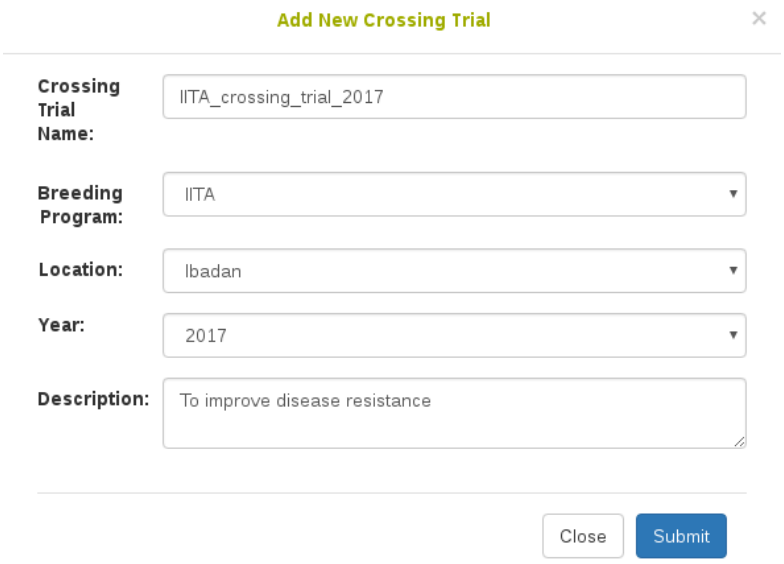
• "**Breeding program**": select a breeding program that is available in the database. New breeding programs can be added on the "Breeding program" page, accessible from the "Manage" menu. *Breeding Program Page*

• "**Location**": select a location for the crossing experiment. New locations can be entered on the "**Locations**" page, accessible from the "**Manage**" menu. *Location Page*

• "**Year**": select a year.

• "**Description**": enter a description for the crossing experiment.

After filling in the information, click "**Submit**" to generate the crossing experiment.

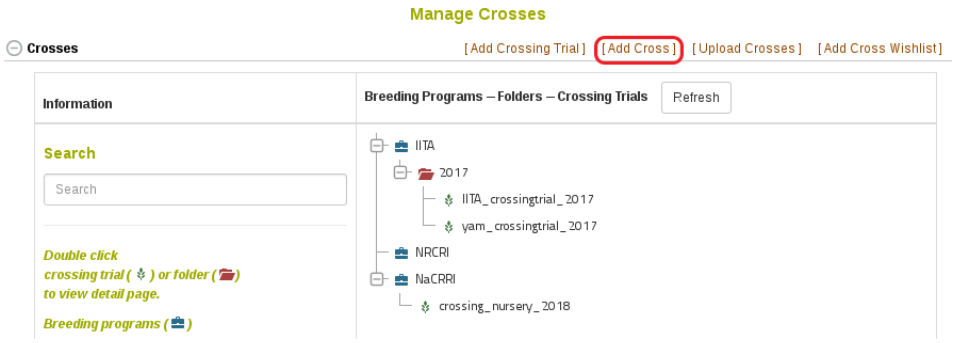


9.2 Cross

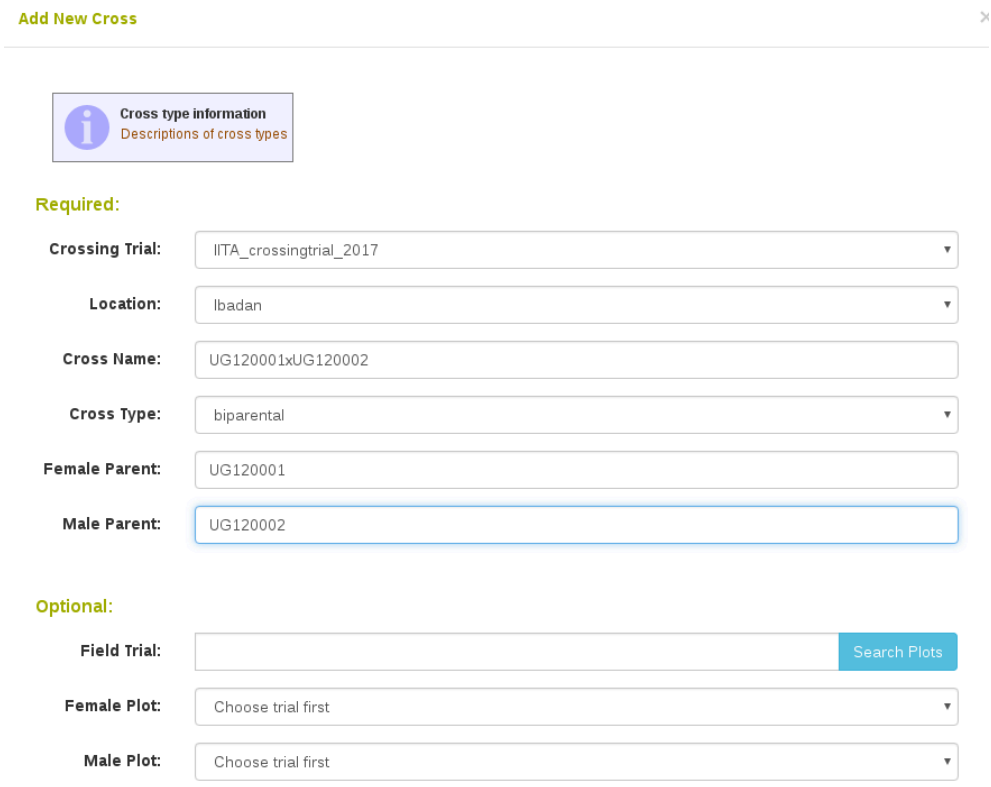
9.2.1 Add New Crosses

Add a cross by using the "Add New Cross" dialog

To add a single new cross, click on "Add Cross" link.



Enter cross information in the popup dialog.

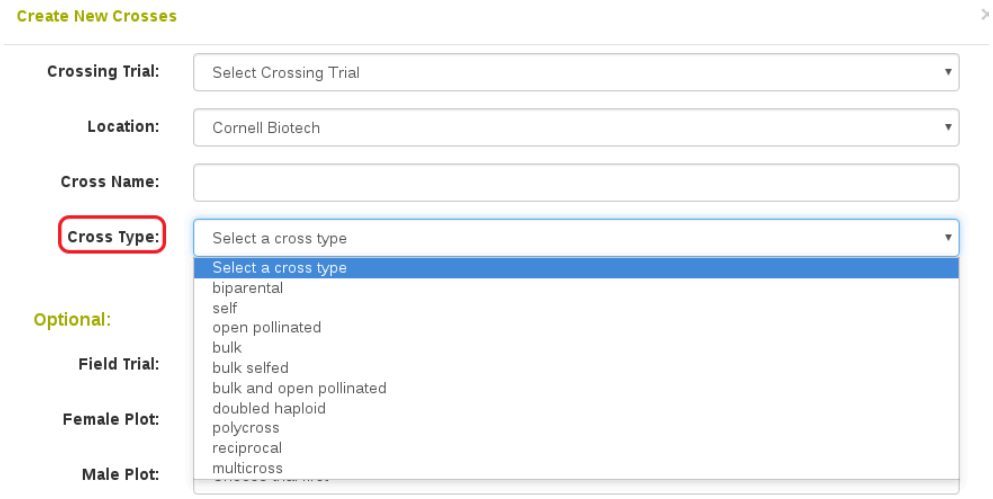


Required Information:

- "**Crossing experiment**": select a crossing experiment available in the database.
- "**Location**": select a location available in the database.

• "**Cross name**": enter a name for the cross. The cross name must not already exist in the database.

• "**Cross type**": the options for cross types are: biparental, self, open pollinated, bulk, bulk selfed, bulk and open pollinated, double haploid, polycross, reciprocal and multicross.



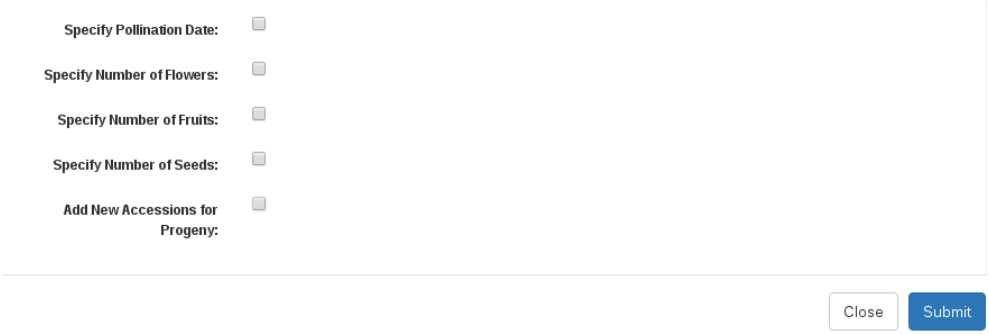
• The "**Female Parent**" and "**Male Parent**" field are auto-complete fields for accessions that are already in the database. The parents specified will be entered in the pedigree of the new accessions generated by this cross.

Optional Information:

• "**Female Plot and/or Male Plot**": In addition to the accession names, specific plots used in the cross can also be added to the database. To retrieve plot names associated with each female/male accession, enter your trial name, then click "**Search Plots**". Plot names of each parental accession in that field trial will be shown in the drop-down list, you can then select the plot used in the cross.

Additional crossing experimental information such as pollination date, number of flowers, number of fruits, number of seeds can be specified during adding new cross. Alternatively, this information can be updated or edited directly on the "**Cross Details**" page.

If you know the number of accessions that are generated from the cross, they can be instantiated immediately in the database by clicking the "**Add accessions for progeny**" checkbox and specifying the number.

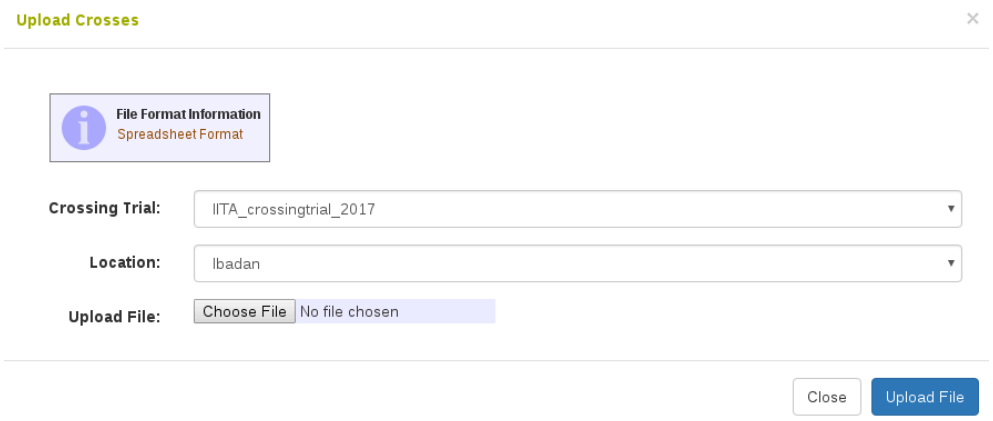


Click "Submit" to generate the cross.

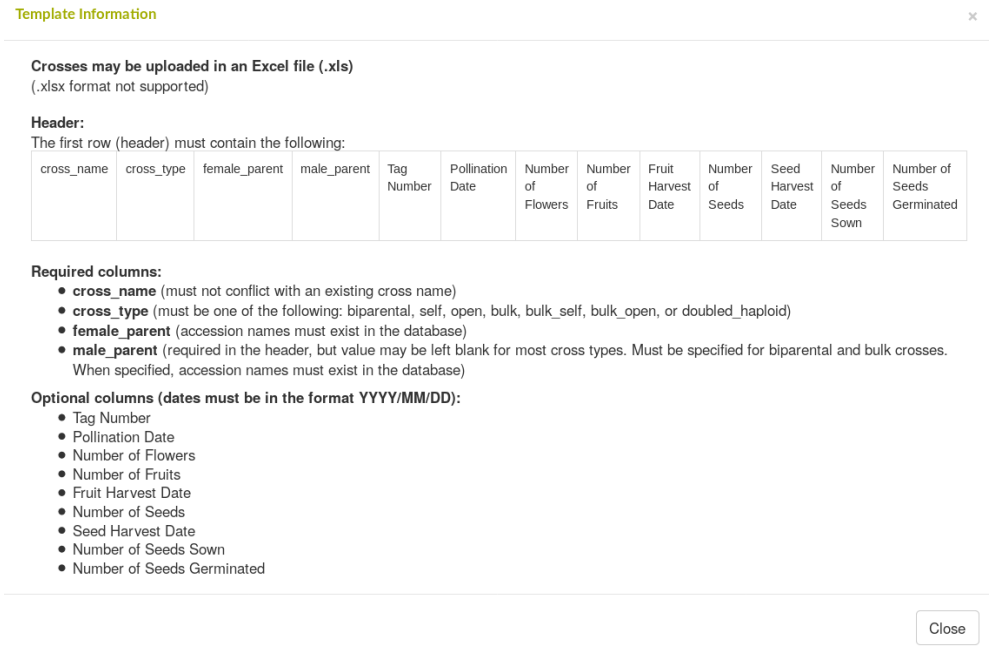
Upload New Crosses

To upload new crosses from an Excel file (.xls or .xlsx), click on "Upload Crosses" link.

Select a crossing experiment and a location available in the database from drop-down lists and choose a file that you want to upload, then click "**Upload File**".



Please check spreadsheet format carefully. The file must be an Excel file (.xls or .xlsx).

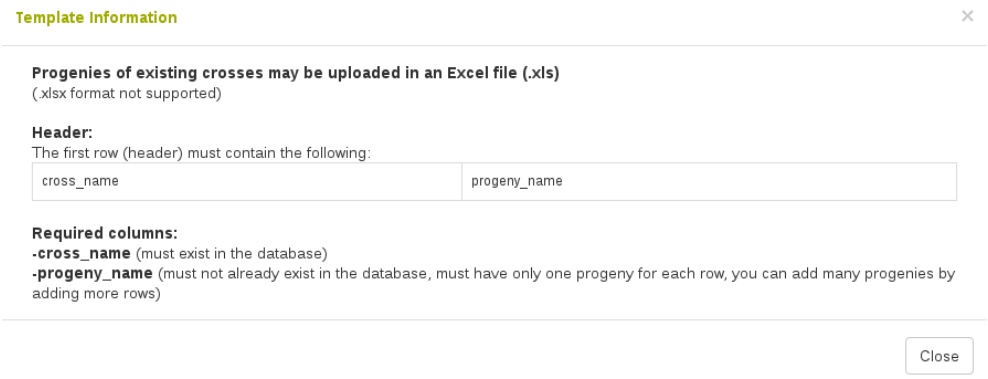


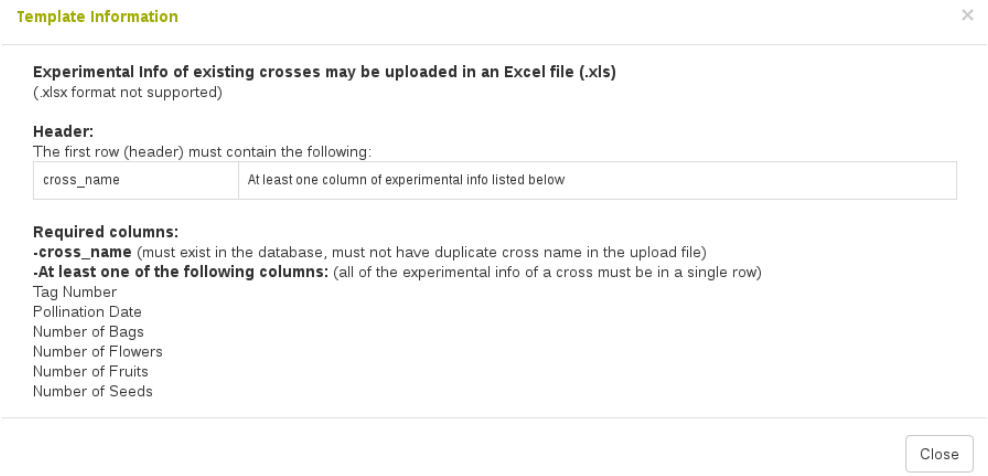
9.2.2 Update Crosses by Uploading

To upload progenies and/or experimental info of crosses already in the database, go to "**Manage-Upload**" page.

In the "**Crosses**" section, there are links for uploading progenies and experimental info.

Please check spreadsheet format in each link carefully. The file must be an Excel file (.xls or .xlsx).





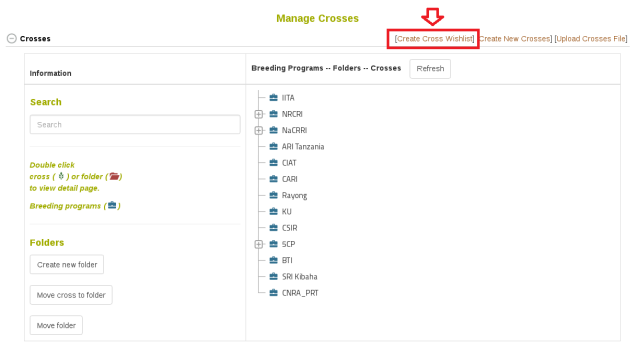
Note: crossing experimental information is customized based on the need for

each crop. As a result, column headers for experimental info in your database may be different from the information shown in this manual.

9.3 Cross Wishlist

An Android ODK application is being developed to record cross information on a mobile device in the field. To link this mobile application with the database, the Cross Wishlist can be used to create a plan for which crosses to perform.

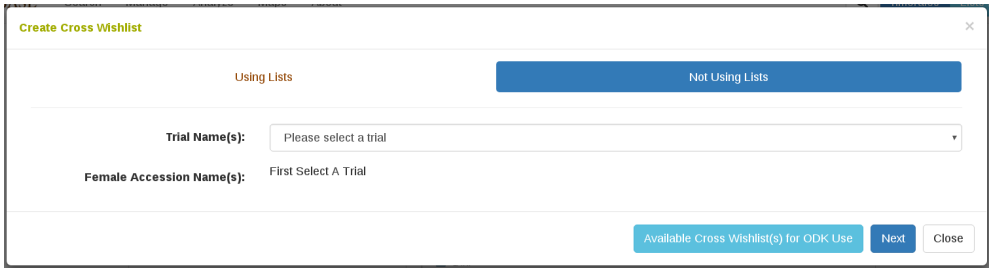
This tool is available on the Manage Cross page. It is currently only available on certain databases, so when you click this link you may see an alert mentioning that the cross wishlist is not available on your database.



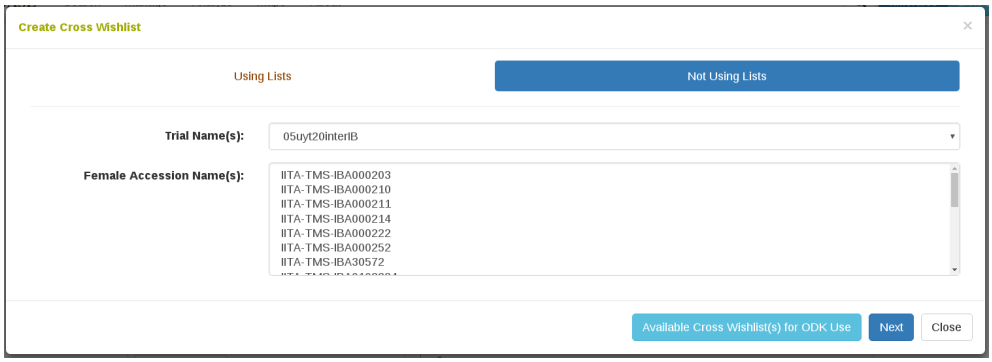
9.3.1 Create a Cross Wishlist

Step 1. Select the accessions to be crossed in your trial

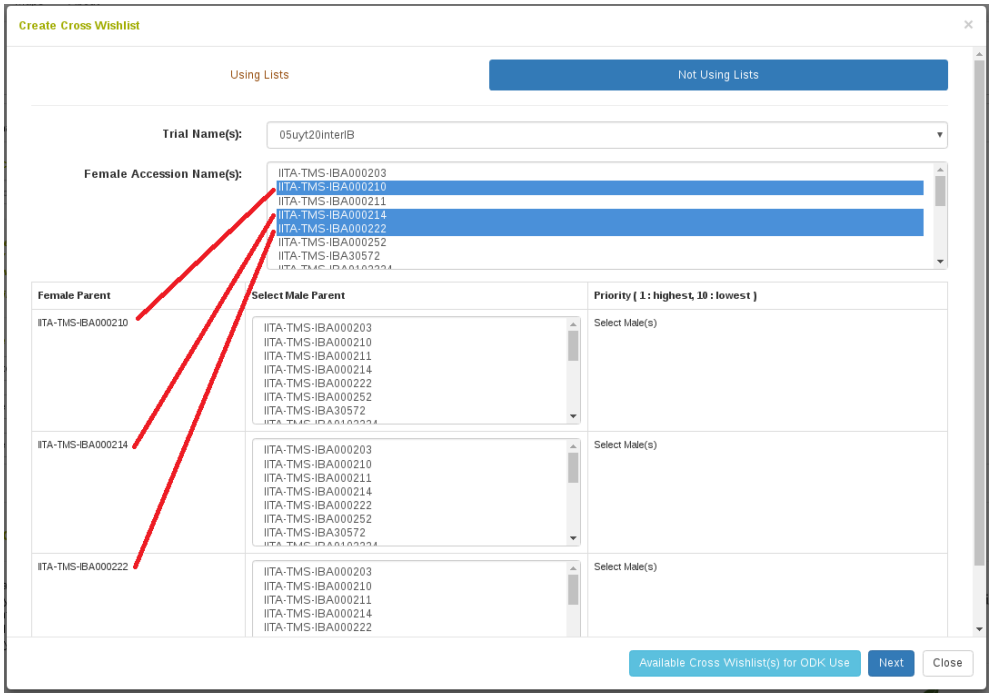
There are two interfaces for this step, either "Not Using Lists" or "Using Lists". Depending on if you already have a list of female and male accessions to use, you can decide on which interface to use. The end result of using either interface is the same.



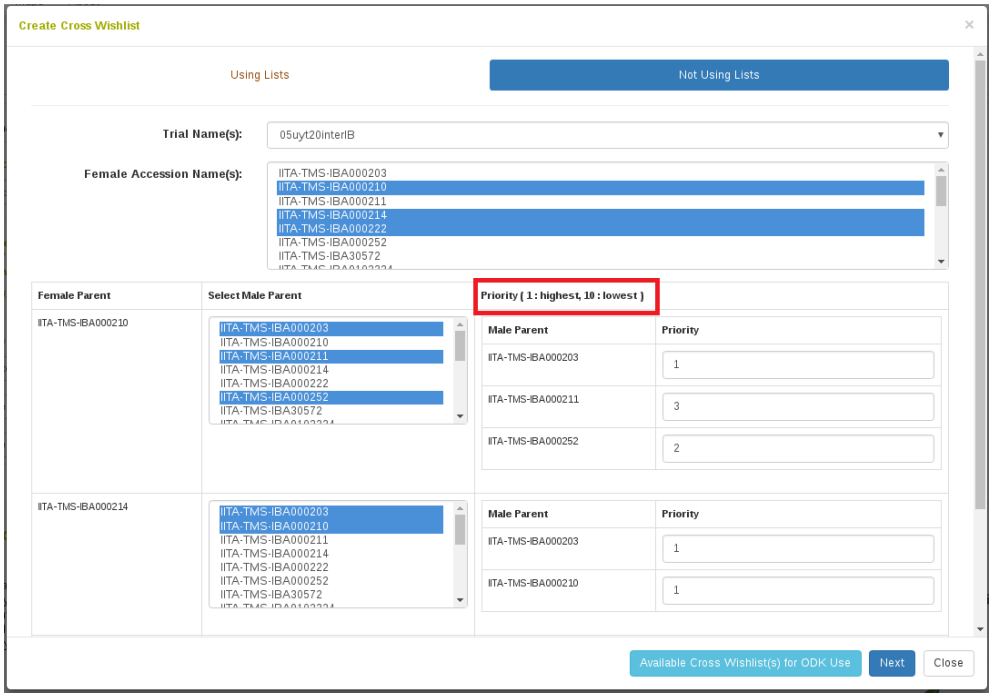
We will start by showing "Not Using Lists". First select the trial in which the crosses are to be performed. This will populate a select box with all the accessions used in that trial. From here, one or many accessions can be selected as the female accession.



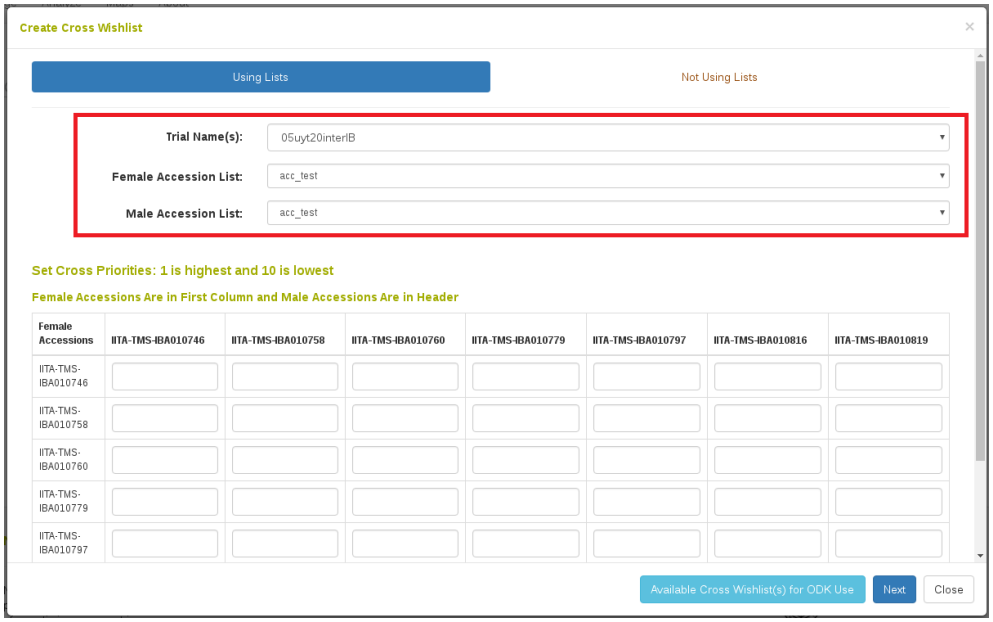
Once the female accessions are selected, a table is populated. Each row in this table begins with the female accession that was selected, followed by a select box with all the accessions used in the trial. From here, one or many accessions can be selected as the male to use in the cross.



Once the male accessions are selected to cross with each female accession, a table indicating priorities appears. Priority is meant to indicate an order in which to attempt the cross; first the highest priority male will be considered, but if this cross is not possible then subsequent males will be considered. An equal priority can be given and this will not indicate a specific order to follow.

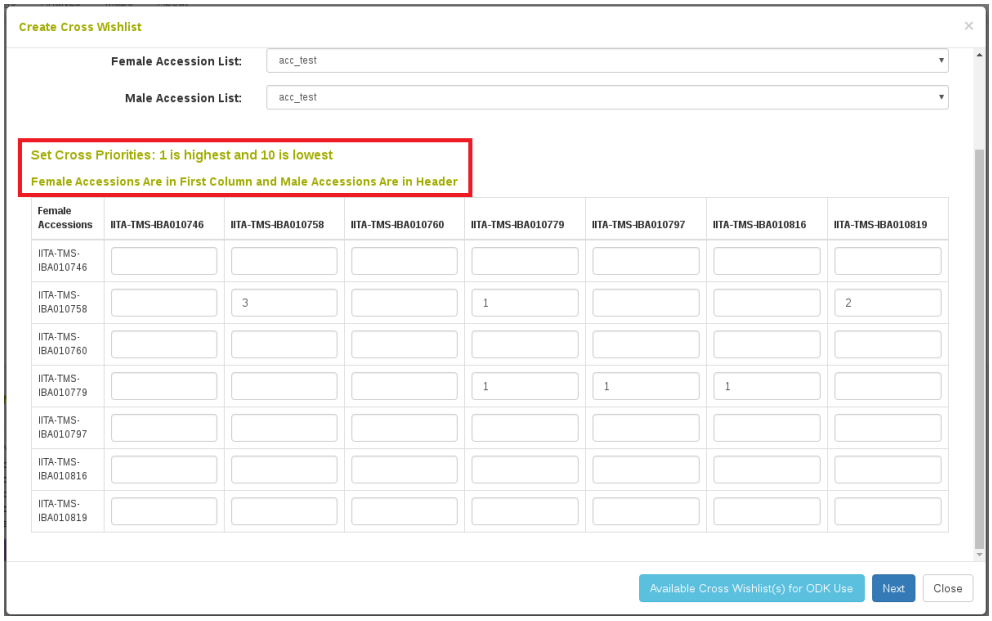


Alternatively, we could have used the "Using List" interface instead. Here we select the trial in which the crosses will be performed and we provide a list of accessions to consider for the females and the males to be crossed.

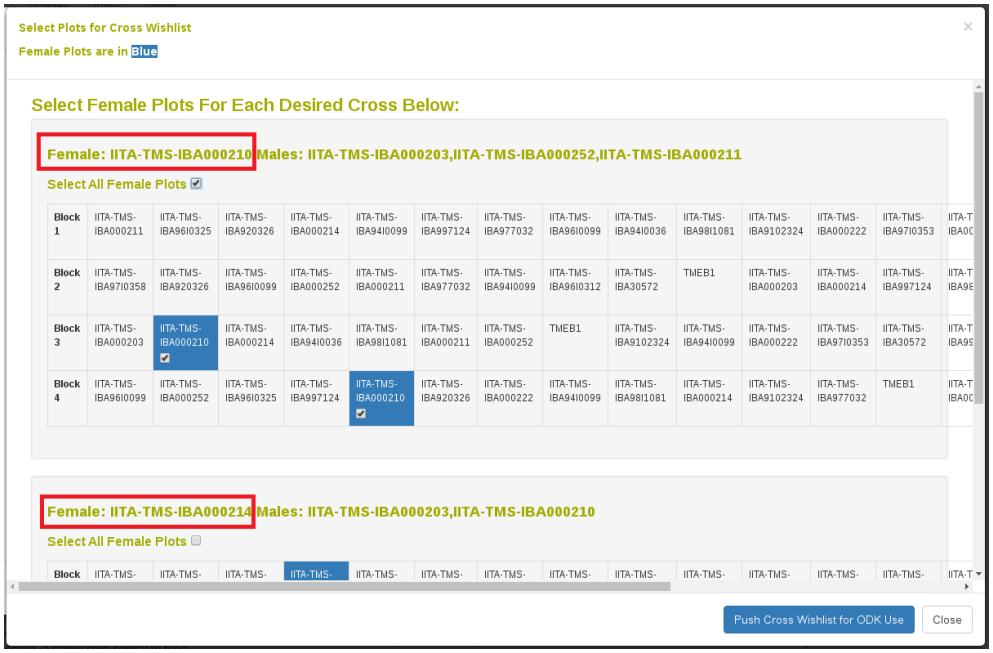


Step 2. Select the female plots to be considered in the crosses

After selecting your lists, the table below is populated. The first column has all the female accessions specified and the header row has all the male accessions specified. The males to consider crossing with each female are indicated with priority.



After female and male accessions are selected to cross, either by the "Nor Using List" or "Using List" interface, click Next. The next dialog will allow selection of specific female plots to use for the cross. Sections for each female accession selected will appear with the field layout displayed. Selecting all plots in which the female is present indicates that the cross should be performed on all plots where that female accession is present.

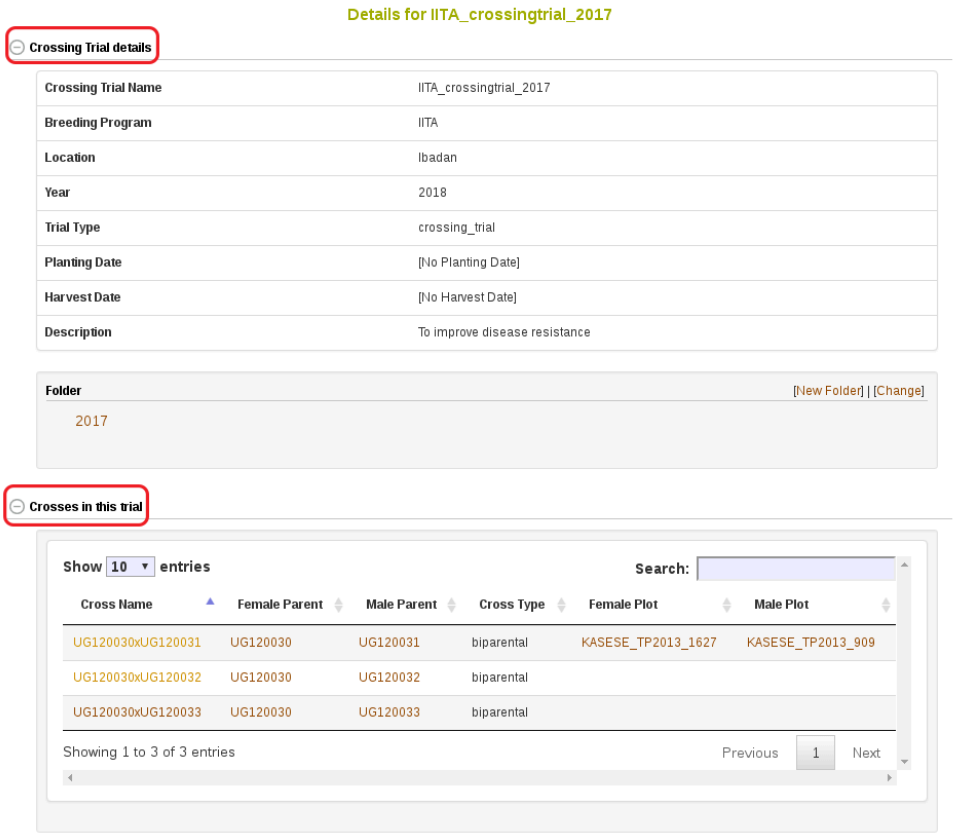


Step 3. Transfer the cross wishlist to your mobile crossing application

Clicking "Push Cross Wishlst for ODK Use" will send the cross wishlist plan to the ONA server for use by the mobile ODK application. Crosses can then be performed and recorded in the field using the mobile application. Afterwards, the crosses are sent back to our database and stored.

9.4 Crossing Experiment Detail Page

Information for crosses in the same crossing experiment is compiled in the crossing experiment detail page.



Details for IITA_crossingtrial_2017

Crossing Trial details

Crossing Trial Name	IITA_crossingtrial_2017
Breeding Program	IITA
Location	Ibadan
Year	2018
Trial Type	crossing_trial
Planting Date	[No Planting Date]
Harvest Date	[No Harvest Date]
Description	To improve disease resistance

Folder [New Folder] | [Change]

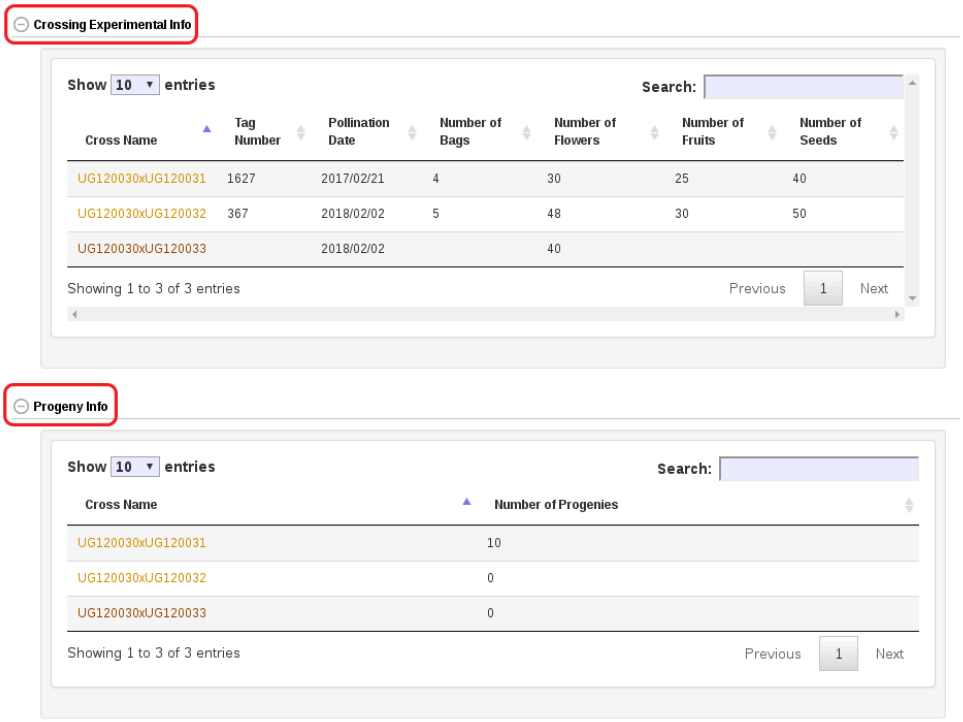
2017

Crosses in this trial

Show 10 entries Search:

Cross Name	Female Parent	Male Parent	Cross Type	Female Plot	Male Plot
UG120030xUG120031	UG120030	UG120031	biparental	KASESE_TP2013_1627	KASESE_TP2013_909
UG120030xUG120032	UG120030	UG120032	biparental		
UG120030xUG120033	UG120030	UG120033	biparental		

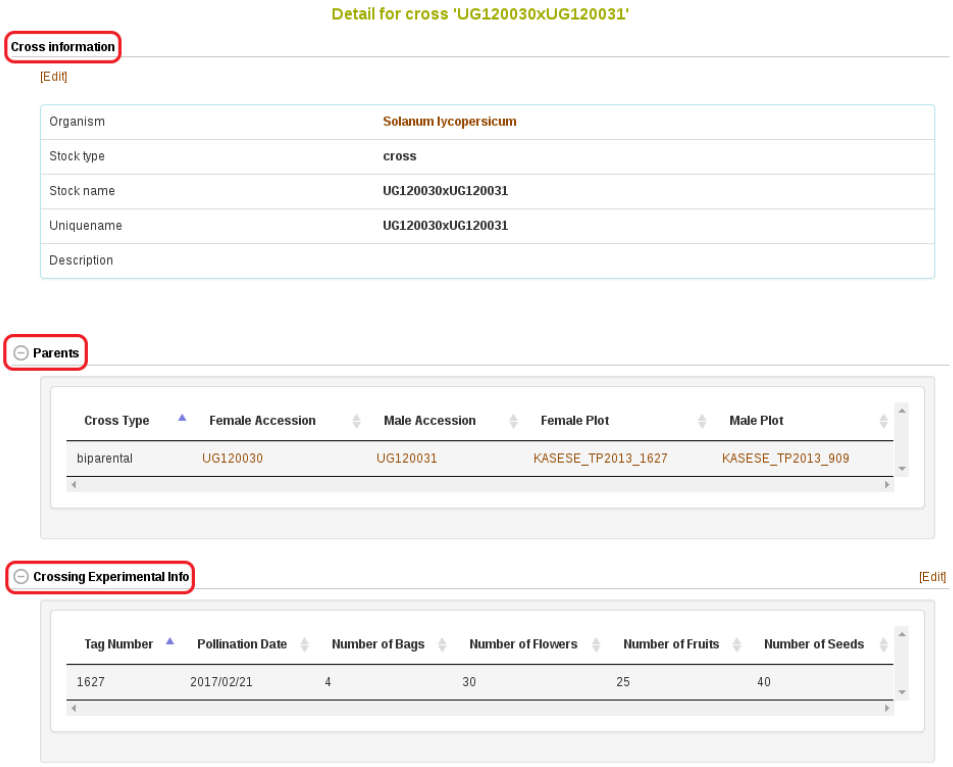
Showing 1 to 3 of 3 entries Previous 1 Next

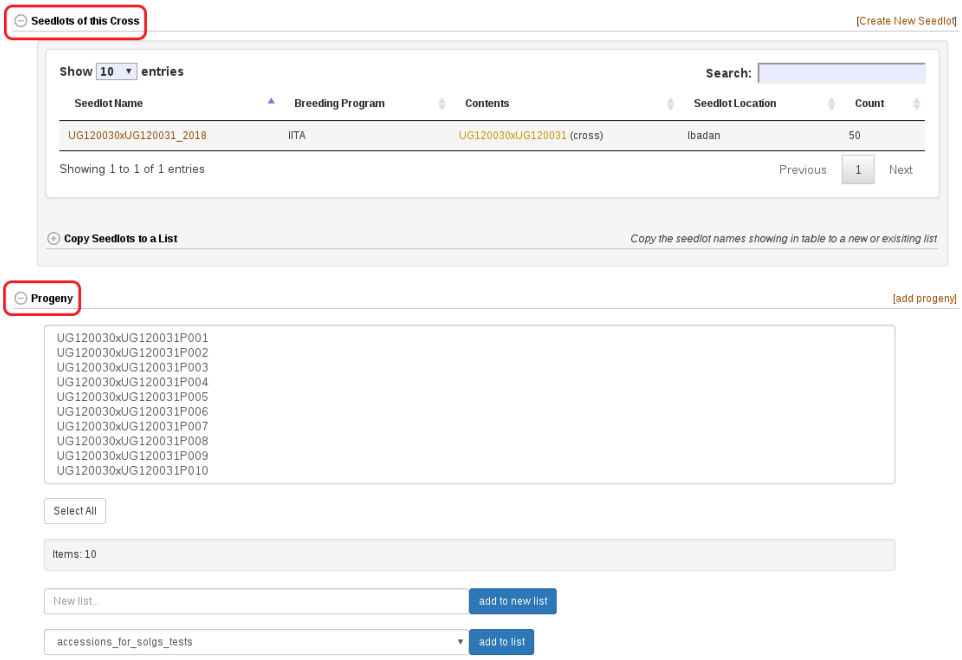


Each cross name, female parent, male parent, female plot and male plot has a link to its own detail page, which contains information specific to each one. Note: crossing experimental information is customized based on the need for each crop. As a result, the details of the information in your database may be different from the information shown in this manual.

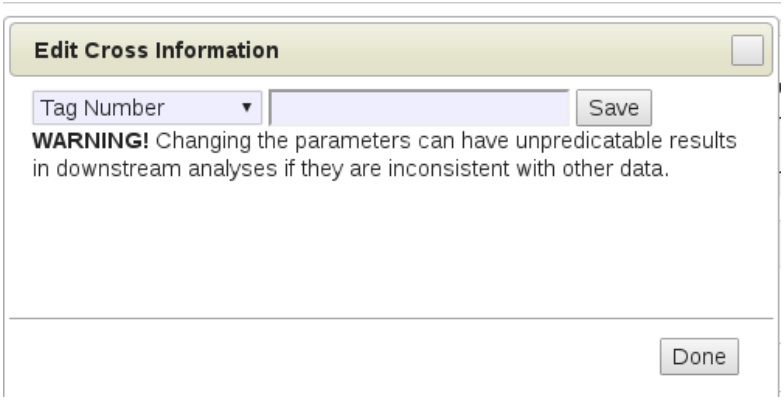
9.5 Cross Detail Page

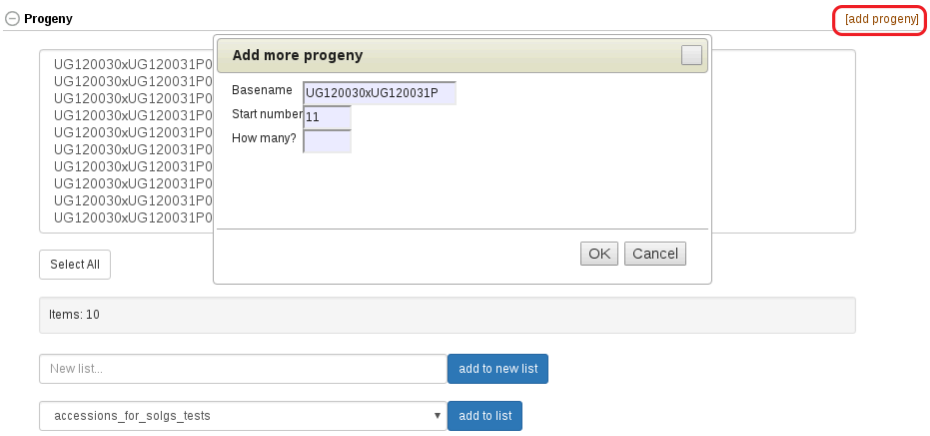
Information of each cross can also be viewed in its detail page.


Detail for cross 'UG120030xUG120031'
Cross information
[Edit]
Organism
Solanum lycopersicum
Stock type
cross
Stock name
UG120030xUG120031
Uniquename
UG120030xUG120031
Description
Parents
Cross Type
Female Accession
Male Accession
Female Plot
Male Plot
biparental
UG120030
UG120031
KASESE_TP2013_1627
KASESE_TP2013_909
Crossing Experimental Info
[Edit]
Tag Number
Pollination Date
Number of Bags
Number of Flowers
Number of Fruits
Number of Seeds
1627
2017/02/21
4
30
25
40



This page allows you to update or edit crossing experimental information and add progenies related to that cross. Note: crossing experimental information is customized based on the need for each crop. As a result, the details of the information in your database may be different from the information shown in this manual.

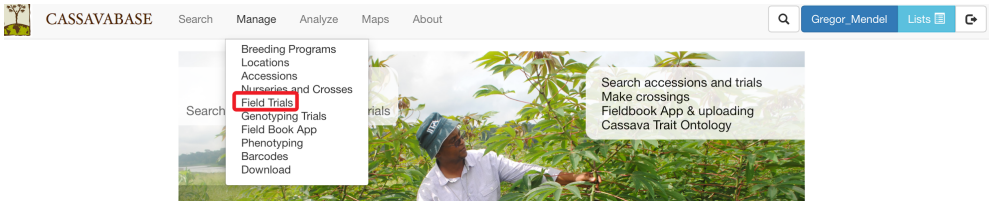



Progeny
[add progeny]
UG120030xUG120031P0
UG120030xUG120031P0
UG120030xUG120031P0
UG120030xUG120031P0
UG120030xUG120031P0
UG120030xUG120031P0
UG120030xUG120031P0
UG120030xUG120031P0
UG120030xUG120031P0
UG120030xUG120031P0
Add more progeny
Basename
UG120030xUG120031P
Start number
11
How many?
OK
Cancel
Select All
Items: 10
New list...
add to new list
accessions_for_solgs_tests
add to list

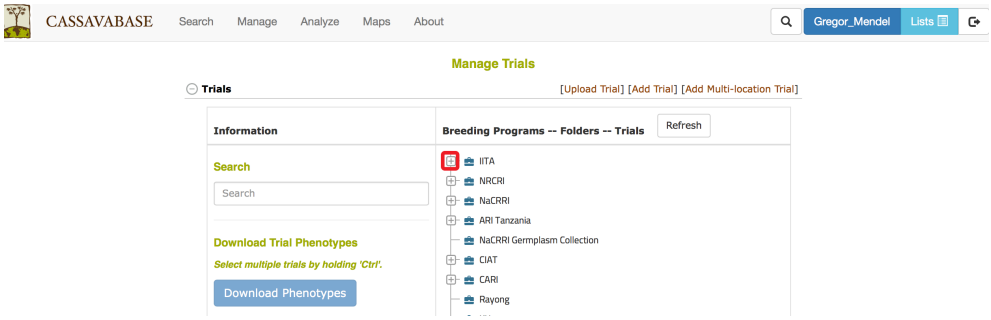
Chapter 10

Managing Field Trials

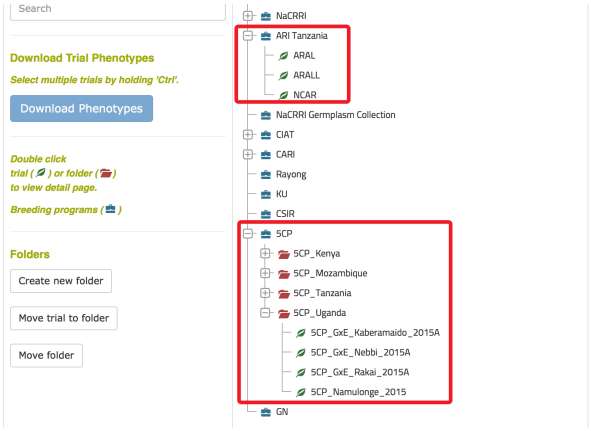
To view trial details on the database, click on the “Field Trials” link under the “manage” menu on the toolbar.



Clicking on the “Field Trials” link will bring you to the “Manage Trials” page. On this page, trials are organized according to their breeding programs. To access trial details, click on the + icon next to your breeding program.

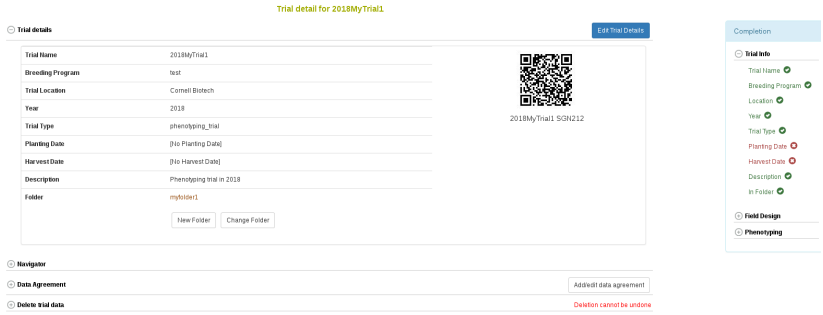


Trials can be placed directly in their breeding program. Alternatively, they can be organized by using folders within each breeding program. Clicking on trial name will take you directly to the trial details page.

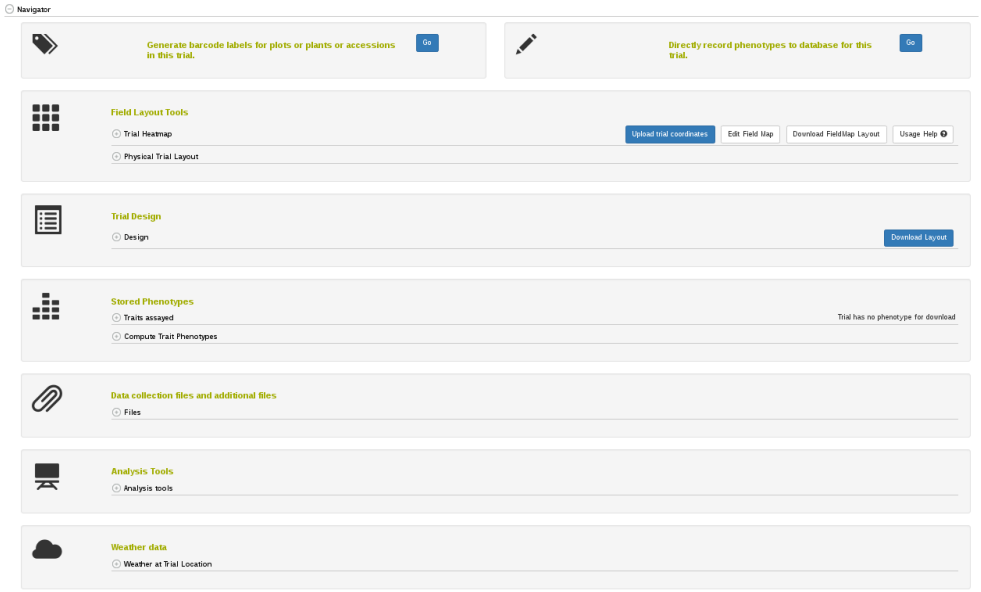


10.1 Trial Detail Page

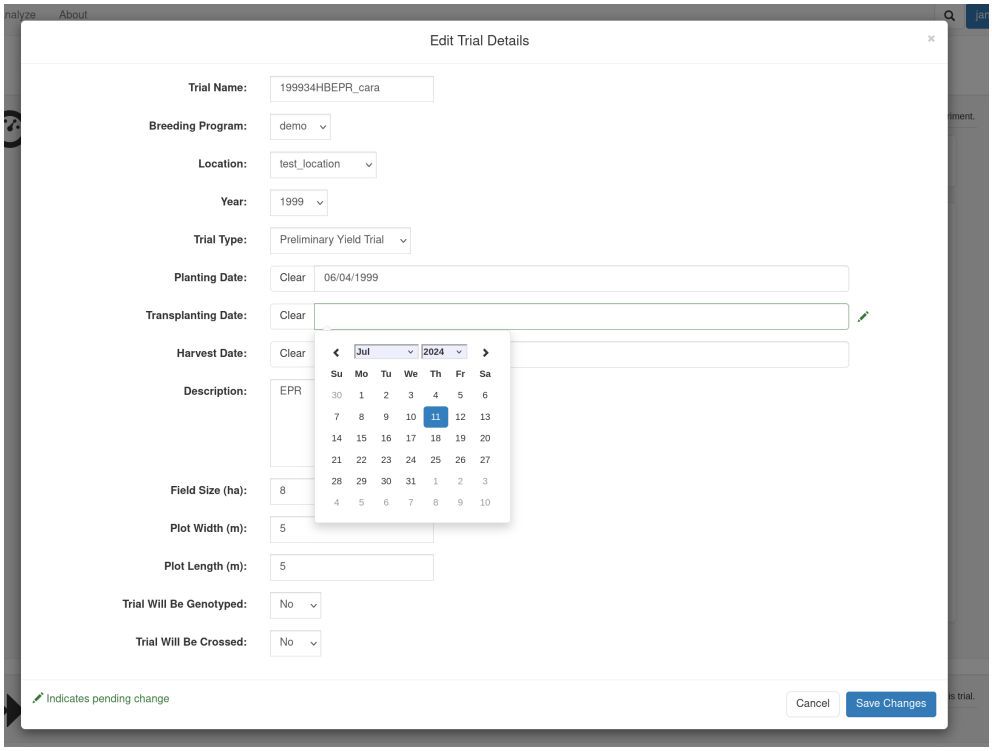
Trial detail page displays important information about individual trial including breeding program, location, year, description of the trial, design, and any files associated with that trial.



The "Navigator" section on the trial detail page allows easy access to all aspects of your trial. This section contains subsections for printing labels for your plots or plants, recording phenotypes, viewing your trial layout or design, viewing phenotypes for this trial, or conducting analyses.



The "transplanting date" field feature will only be shown if it has a value. To add a transplanting date after creating a trial, change the show_transplanting_date parameter from 0 to 1 in the SGN config file. As a result, you will be able to add a date under the transplanting date field by clicking the "Edit Trial Details" on the trial detail page.



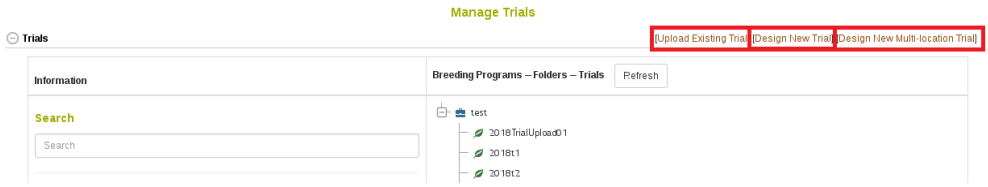
10.2 Adding Trials

Only users with the account status of "submitter" may create trials. To learn how to change your account status from "user" to "submitter" visit the 1.2 page.

10.2.1 Prerequisites

- To add a trial, all of your accessions should already exist in the database before you begin to design a trial. If you have accessions that are not in the database, see the instructions for adding accessions .
- Breeding program and location for your trial should also exist in the database. If you need to add breeding program and/or location to the database, see instructions for adding breeding program and location in the "Managing Breeding Programs" and "Managing locations" respectively.

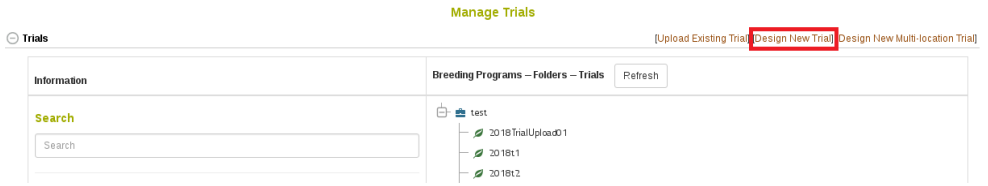
On the "Manage Trials" page, there are three alternative methods for you to add new trials: by using "Add Trial" form, "Upload Trial" form, or "Add Multi-location Trial" form.



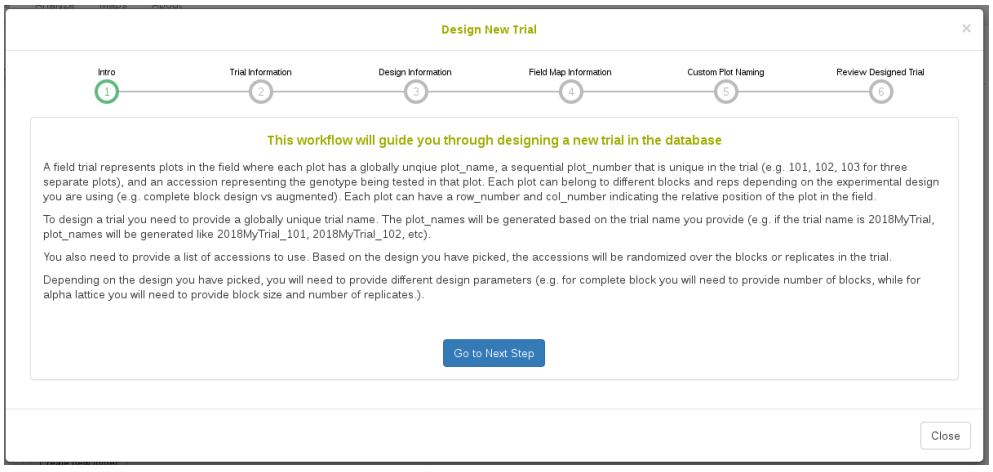
10.2.2 Adding a trial by using "Add Trial" form

Step 1. Begin the "Design new trial" workflow

Click on "Design New Trial" to begin.



The first step in this workflow is an introduction that looks like:

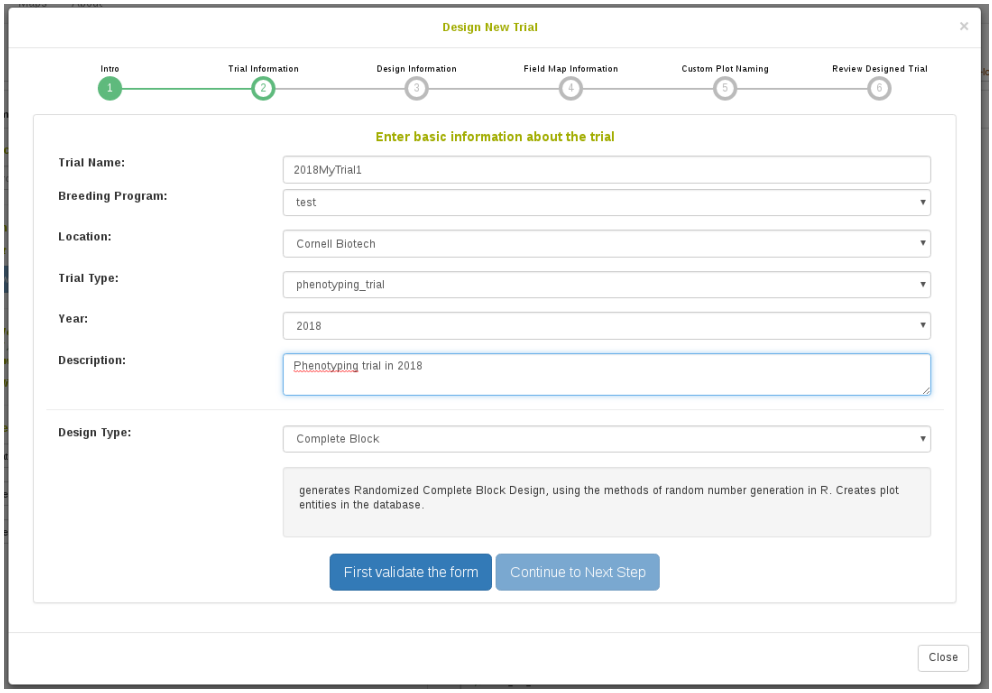


Here it gives information about what is required for a trial, including that to create a new trial, you need to create a list of the accessions that you would like to use in the trial. Lists can be viewed, created, and modified with the

"**lists**" tool at the upper right of the screen. For more information on lists, click *here*.

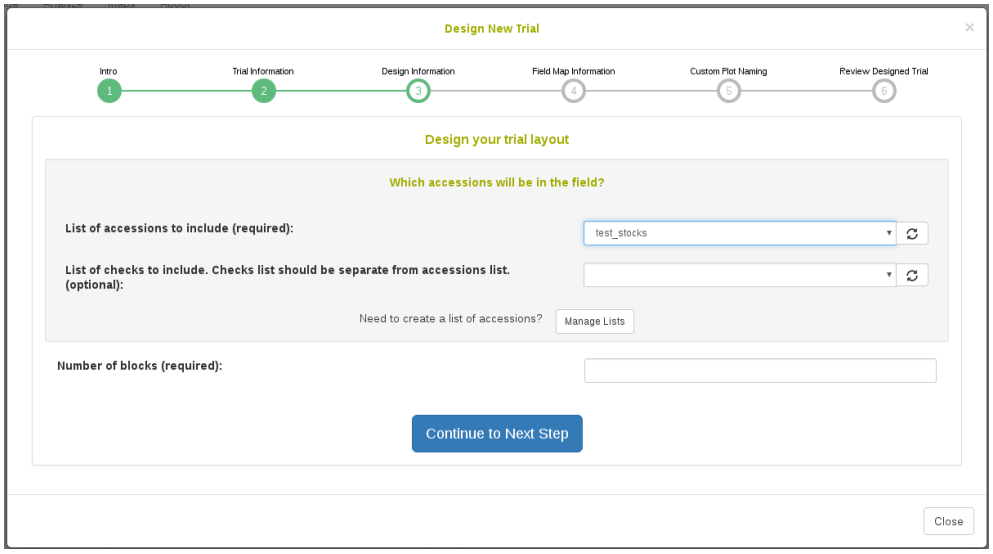
Step 2. Enter "Trial Information"

On this screen you need to enter basic information about the trial, such as breeding program and location(s). You must also select a design type, such as Complete Block Design. The design is important because it influences how your genotypes are distributed and randomized over the trial. You must first click validate before proceeding to the next step.



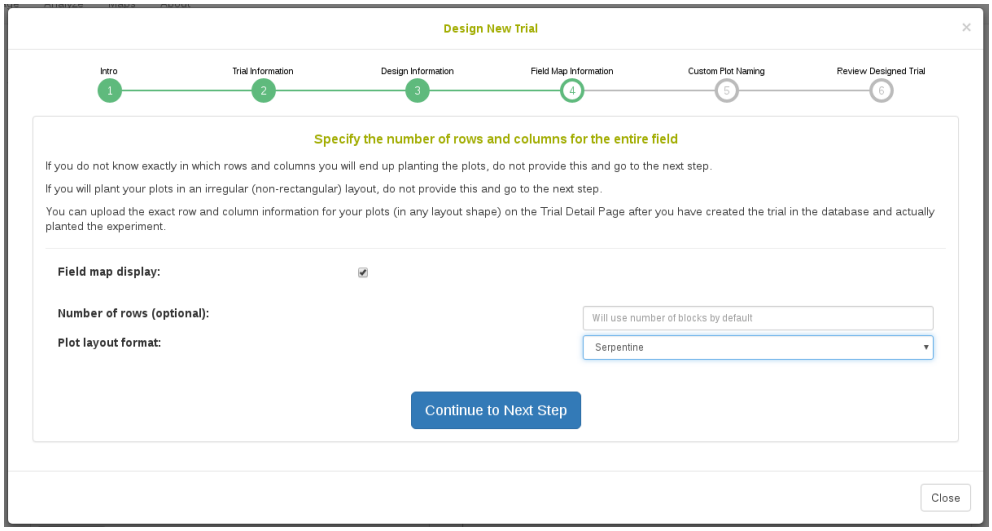
Step 3. Enter "Design Information"

On this screen you need to specify a list of accessions to use in the experiment. This list must be a valid list of accessions. You must also specify all required design information, such as number of blocks in this case.



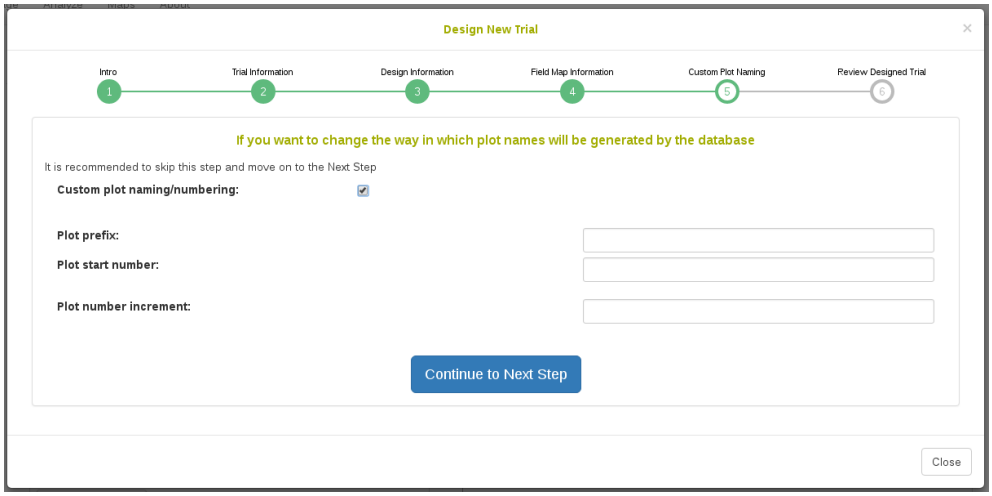
Step 4. Enter "Field Map Information" (Optional)

On this screen you can specify how the row and column numbers will be generated for the plots in the trial. The row and column number represent a relative position of the plot in the field. If you are not exactly sure of how you will plant the plots in the field or you have an irregular (non-rectangular) layout, you can skip this step for now. This information can be added on the Trial Detail Page once the trial is saved in the database in order to reflect exactly how the plots were planted in the field.



Step 5. Custom Plot Naming (Optional)

On this screen it is possible to change the format in which plot names will be generated for your trial. It is recommended to skip this step and just use the format generated by the database by default.

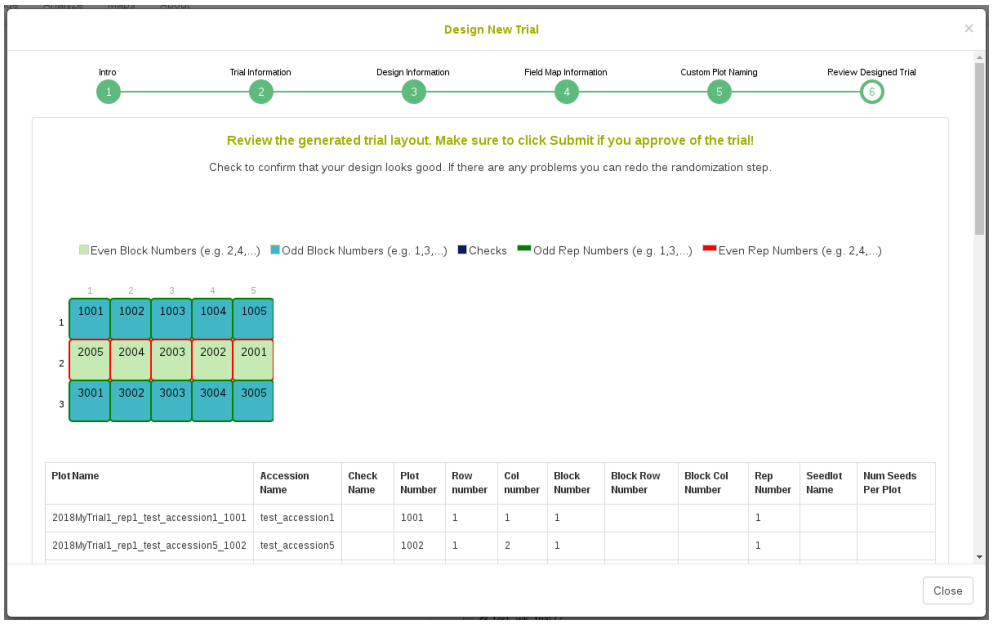


Step 6. Review Designed Trial

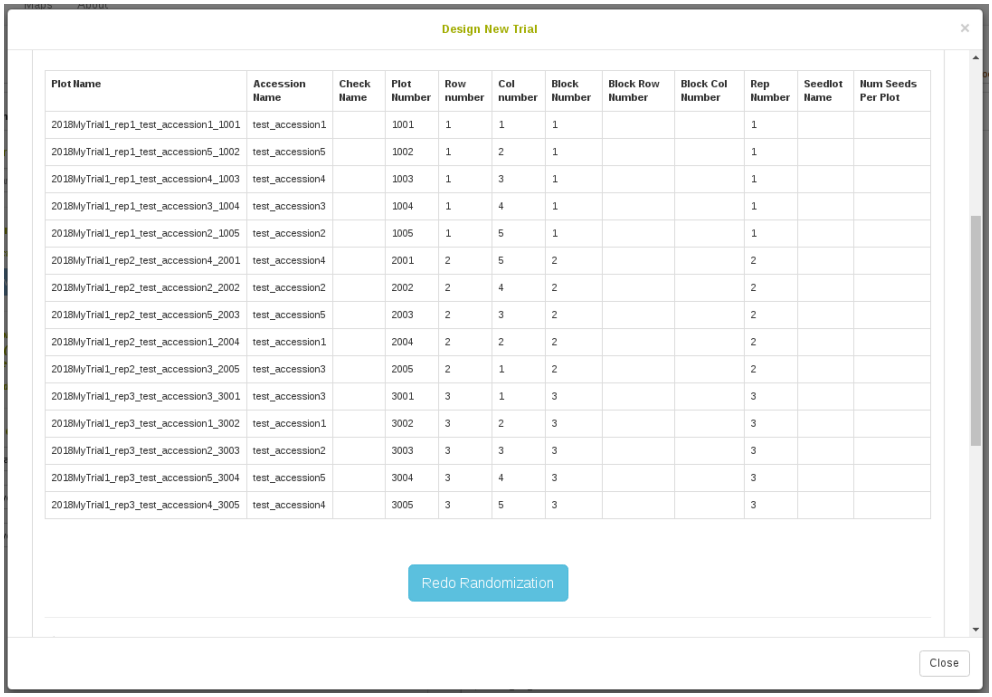
On this screen you can review the trial that the database has generated.

You will see a graphical representation of the trial. The numbers on the

squares represent the plot_number of each plot and on mouse hover you can see further information about the plot.

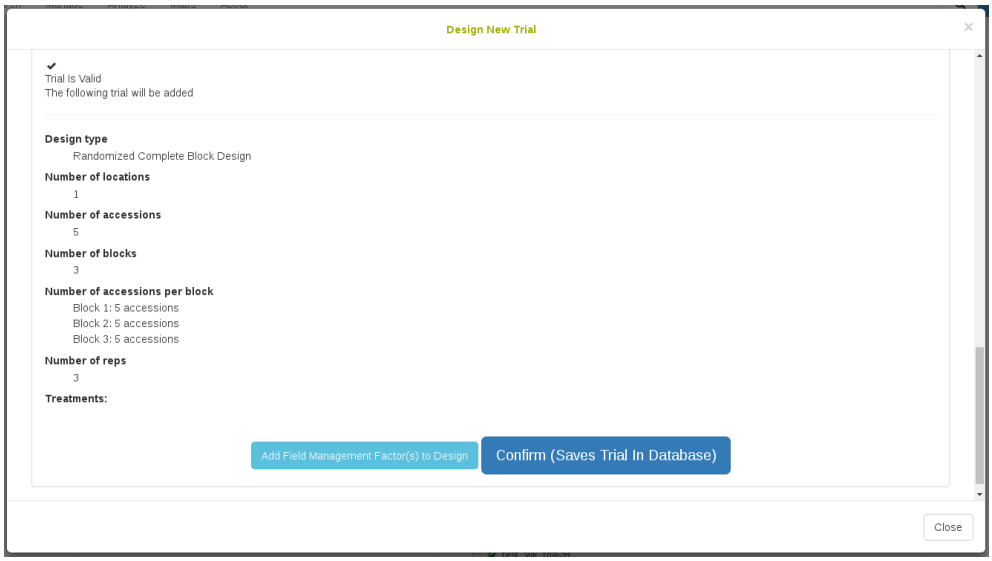


You will also see a table representation of all the plots and their information. If you want to redo the randomization, you can click the “Redo Randomization” button.



Plot Name	Accession Name	Check Name	Plot Number	Row number	Col number	Block Number	Block Row Number	Block Col Number	Rep Number	Seedlot Name	Num Seeds Per Plot
2018MyTrial1_rep1_test_accession1_1001	test_accession1		1001	1	1	1			1		
2018MyTrial1_rep1_test_accession5_1002	test_accession5		1002	1	2	1			1		
2018MyTrial1_rep1_test_accession4_1003	test_accession4		1003	1	3	1			1		
2018MyTrial1_rep1_test_accession3_1004	test_accession3		1004	1	4	1			1		
2018MyTrial1_rep1_test_accession2_1005	test_accession2		1005	1	5	1			1		
2018MyTrial1_rep2_test_accession4_2001	test_accession4		2001	2	5	2			2		
2018MyTrial1_rep2_test_accession2_2002	test_accession2		2002	2	4	2			2		
2018MyTrial1_rep2_test_accession5_2003	test_accession5		2003	2	3	2			2		
2018MyTrial1_rep2_test_accession1_2004	test_accession1		2004	2	2	2			2		
2018MyTrial1_rep2_test_accession3_2005	test_accession3		2005	2	1	2			2		
2018MyTrial1_rep3_test_accession3_3001	test_accession3		3001	3	1	3			3		
2018MyTrial1_rep3_test_accession1_3002	test_accession1		3002	3	2	3			3		
2018MyTrial1_rep3_test_accession2_3003	test_accession2		3003	3	3	3			3		
2018MyTrial1_rep3_test_accession5_3004	test_accession5		3004	3	4	3			3		
2018MyTrial1_rep3_test_accession4_3005	test_accession4		3005	3	5	3			3		

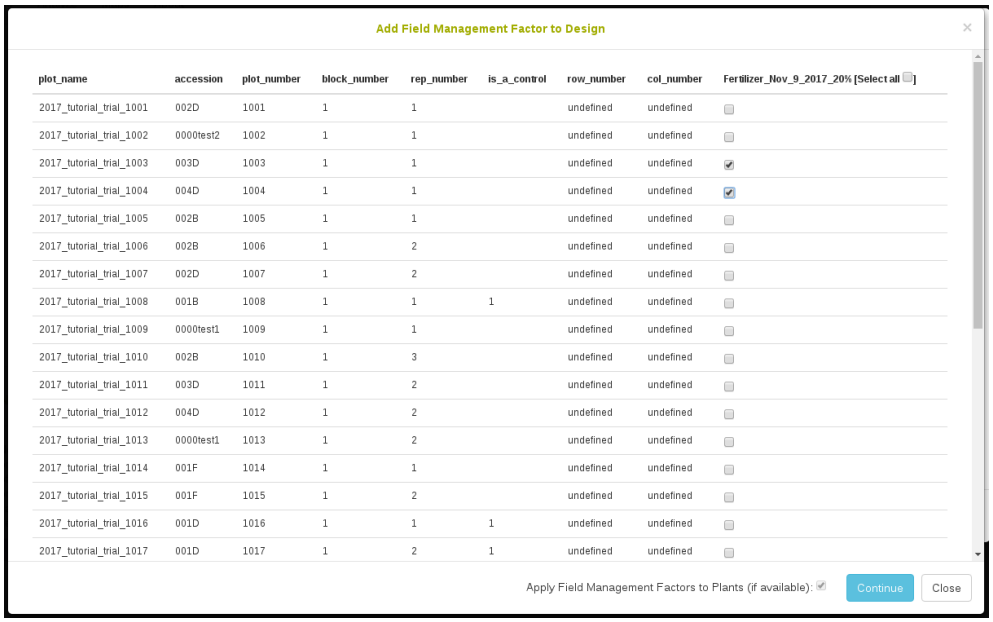
At the bottom there is a brief summary of the trial followed by two buttons.



Step 7. Add Field Management Factors to your design (Optional)

You can add Field Management Factors by clicking "Add Field Management Factor(s) to Design". Clicking this opens a dialog to name your factor. You can name this to account for fertilizer or watering regime or inoculation or anything else. This is optional and can be added from the trial detail page afterwards.

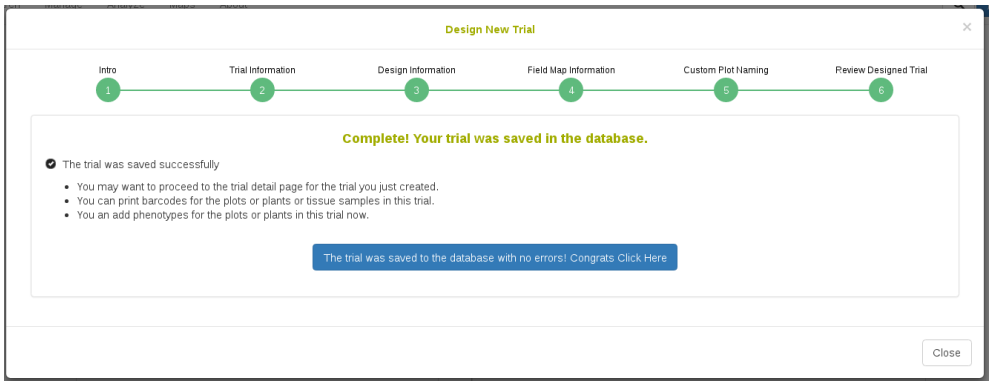
Click "Continue" and a dialog will appear where you can specify plots for which the factor was applied. There is a select all button also.



Step 8. Saving new trial in the database

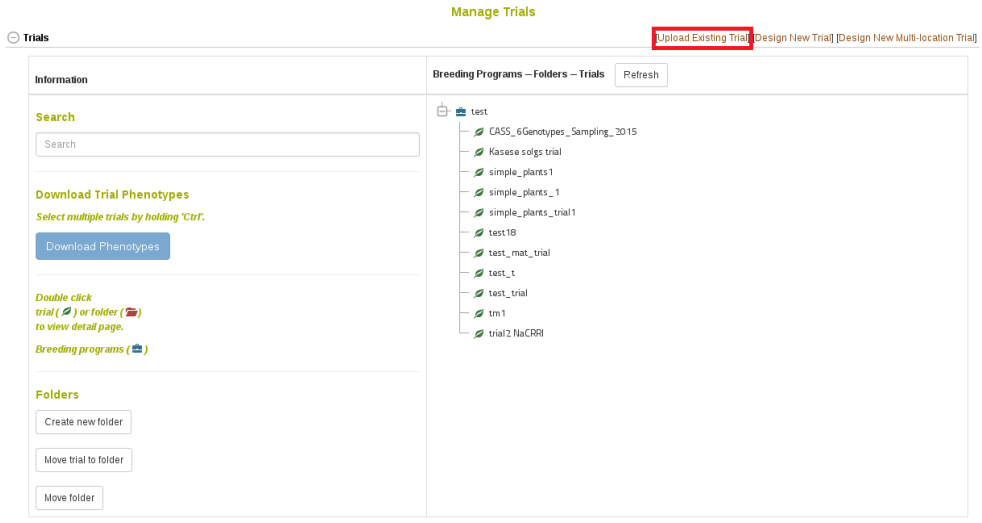
Once you are done reviewing the trial you can click "Confirm" to save the generated trial into the database. Once the trial has saved you will see the

final completion screen:



10.2.3 Adding a trial from an uploaded file

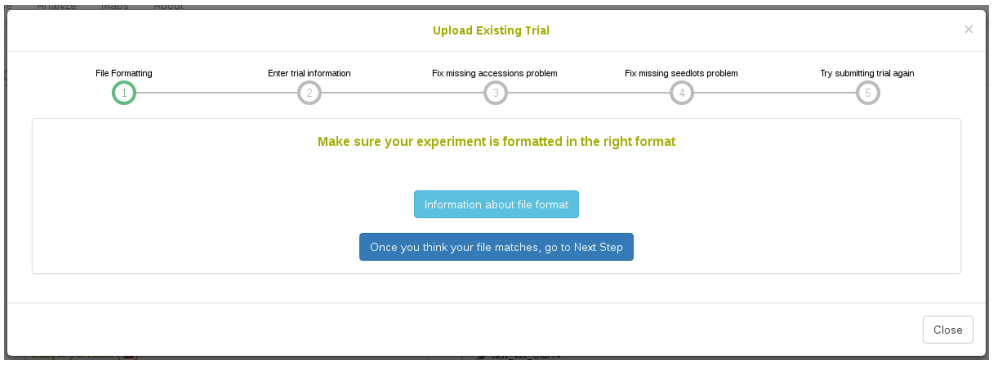
If you already have trial design layout in a spreadsheet, you can add your trial into the database by using "Upload Trial" form. To access "Upload Trial" form, click on "Upload Existing Trial(s)" button on the "Manage Trials" page.



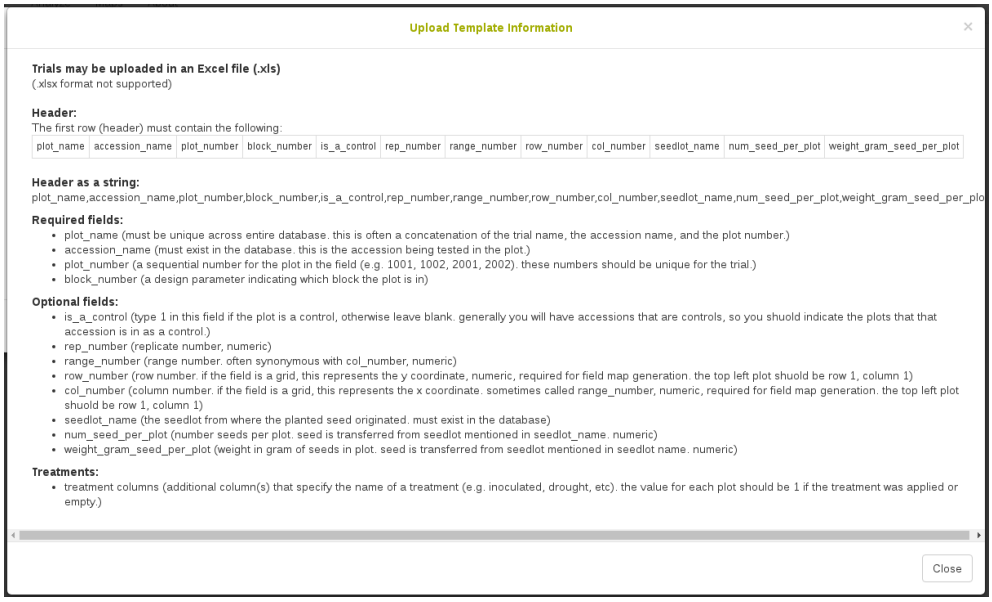
When you click "Upload Existing Trial(s)" you will see the following workflow. Notice that there are 5 numbered sections to the workflow.

Step 1:

The first step is to understand what the format of the trial upload is. It is important to understand that the field layout represents plots in the experiment. Each plot has a globally unique plot_name, a sequential plot_number that is unique in the trial (but not globally unique. e.g. 101, 102, 103 for three separate plots), an accession_name representing what genotype is planted in that plot, and a block_number representing design replication. Each plot can be thought of as having a row_number and a column_number representing the relative position of the plot in a grid (e.g. the top left plot is row 1 column 1 following by row 1 column 2). Each plot can be planted with an amount of seed from a seedlot, where the seedlot_name represents the specific seed packet that was used, and num_seed_per_plot and weight_gram_seed_per_plot represent amount that were transferred from the seedlot_name to the plot_name. Treatments (management factors) can be applied onto plots using additional column names in your file, where a 1 represents if the factor was applied to the plot and an empty cell means it was not applied.



This information and more can be found by clicking “Information about file format”, which shows the following:



Minimum File requirements

- All accession names in the file must exist in the database. See adding accessions for more information.
- The uploaded file should be XLS or XLSX file format (NOT CSV).
- The first row (header) must contain the column names: plot_name accession_name plot_number block_number is_a_control rep_number range_number row_number col_number seedlot_name num_seed_per_plot weight_gram_seed_per_plot

Minimal Example:

plot_name	accession_name	plot_number	block_number	is_a_control	rep_number	range_number	row_number	col_number	seedlot_name	num_seed_per_plot	weight_gram_seed_per_plot	seed_p
2018plot1	accession1	101	1		1							
2018plot2	accession2	201	2		2							
2018plot3	accession1	102	1		2							
2018plot4	accession2	202	2		1							

File validation

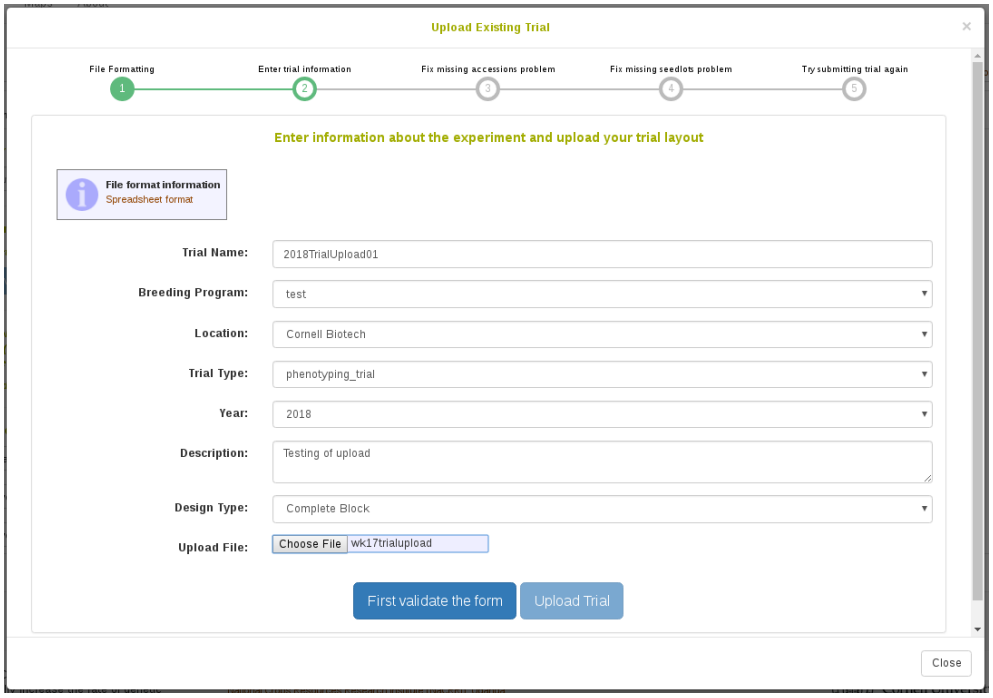
- In case of errors in the uploaded file such as missing or invalid data, a window will appear listing the specific errors in the file that must be corrected before a successful upload.

Uploading a trial with Field Management Factors

- You can upload a trial with field management factor(s) by adding additional column(s). The column header will be the factor e.g. fertilizer, watering regime, inoculation, etc. and the values in these columns will be either 1 or empty, indicating that the factor was applied to the plot or not.

Step 2:

Once you feel that your experiment field layout is in the right format, click on to the Next Step. You will see the following form which must be filled in completely:

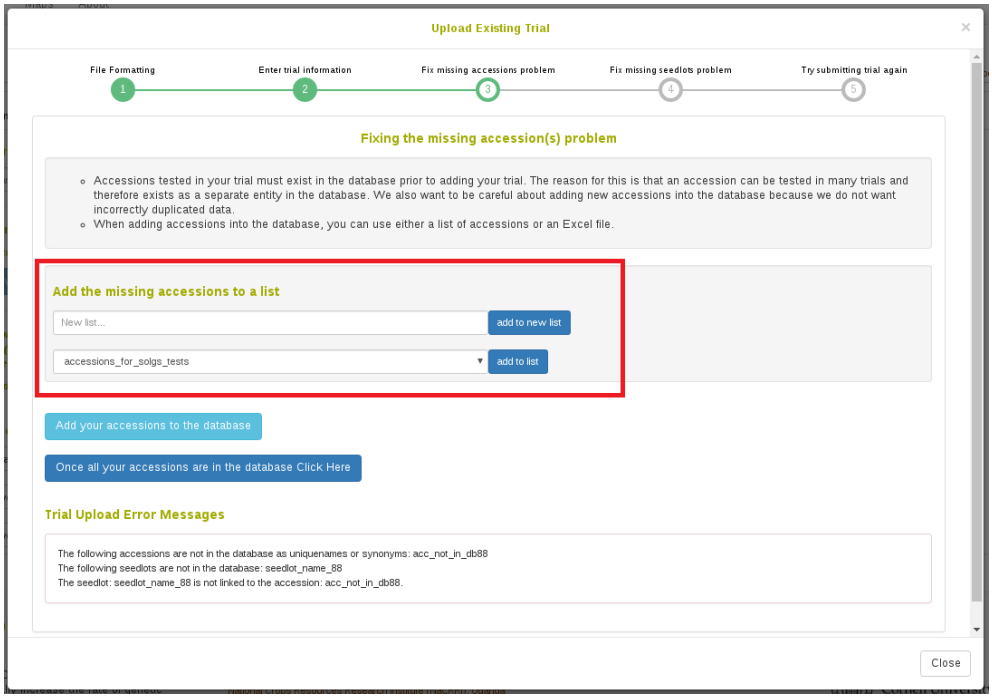


The trial name must be globally unique in the database. Please try to follow standard naming conventions for your group.

First you need to validate the form, and then you can click “Upload Trial”.

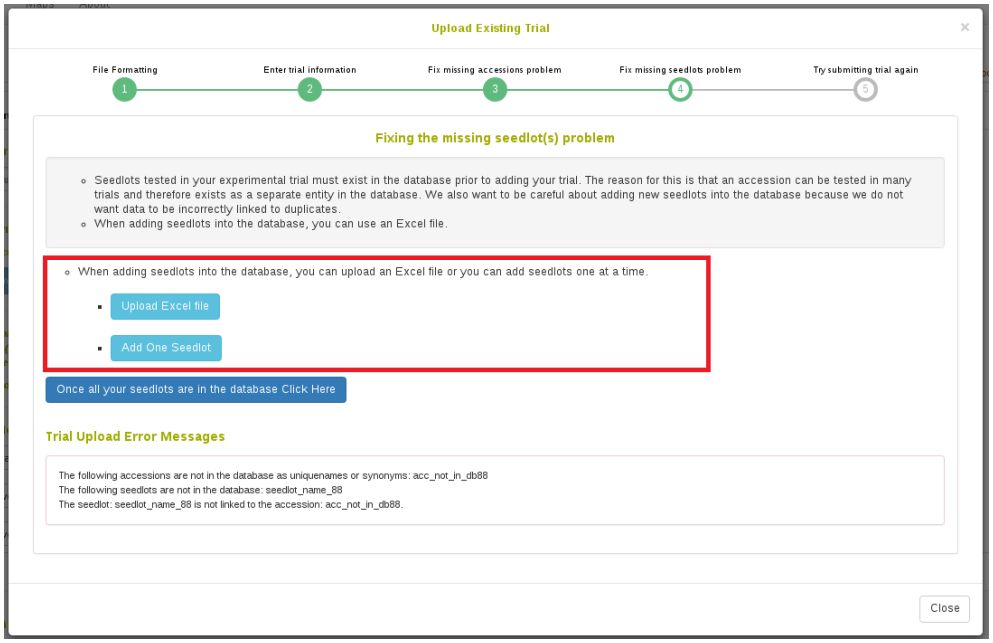
Step 3:

In the case where you have uploaded an experiment using accession_names that are not already present in the database, you will be taken to this screen. If the accession_names in your file are all already in the database, this step will be skipped. The reason it is necessary for your accessions to be in the database before you can add a trial using them is that a single accession can be used among many trials and therefore must exist as a separate entity in the database; because of this it is also very important to be careful about adding wrongly duplicated accession_names into the database. From this screen it is possible to make a new list with the missing accession_names and then click “Add Accessions to the database” to immediately resolve the issue. Once all your accessions are in the database, click to move to the Next Step.



Step 4:

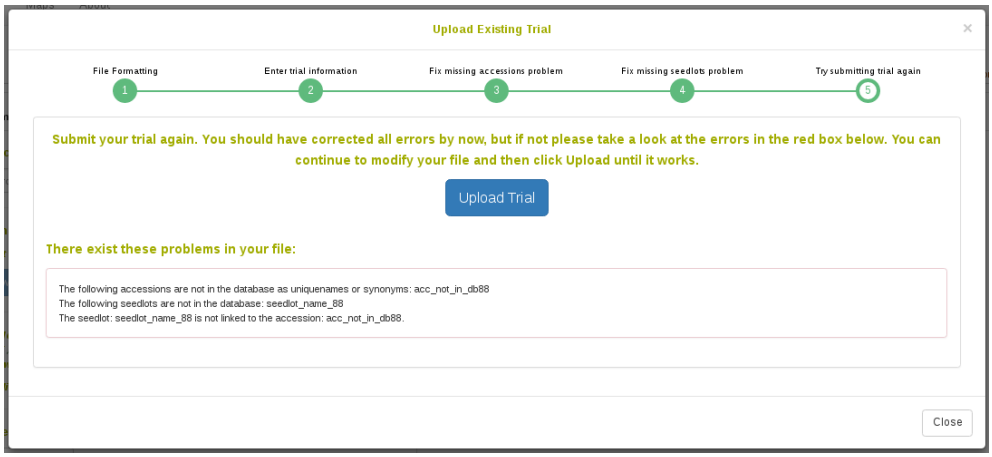
In the case where you have uploaded an experiment using seedlot_names that are not already present in the database, you will be taken to this screen. If the seedlots in your file are all already in teh database, this step will be skipped. The reason it is necessary for your seedlots to be in the database before you can add a trial using them is that a ginel seedlot can be used among many trials and therefore must exist as a separate entity in the database. From this screen it is possible to add the missing seedlots; you can either upload an XLS or XLSX file to add many at once or you can add them one by one. Once all your seedlots are in the database, click to move to the Next Step.



Step 5:

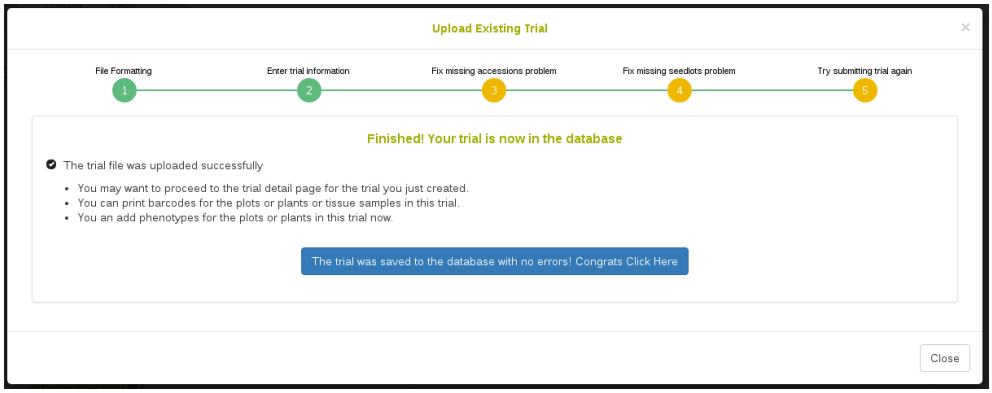
If there are any other errors with your file, such as if the plot_names are not globally unique in the database or your plot_numbers are not unique in your trial or row_number is not an integer or any other error, you will see the errors listed in the red box. It is up to you to correct these errors in your file. Simply open up the file you selected earlier in Excel and correct the issues and then save the file. Then you can click “Submit Trial” and it will resubmit it for you. You can continue to edit your file here and submit

as many times as you need until it is accepted.



Completion screen

Whether you were lucky enough to submit your trial successfully on Step 2 or if you tried many times on Step 5, once your trial has been saved in the database you will see the following screen:



10.2.4 Multi-location trials

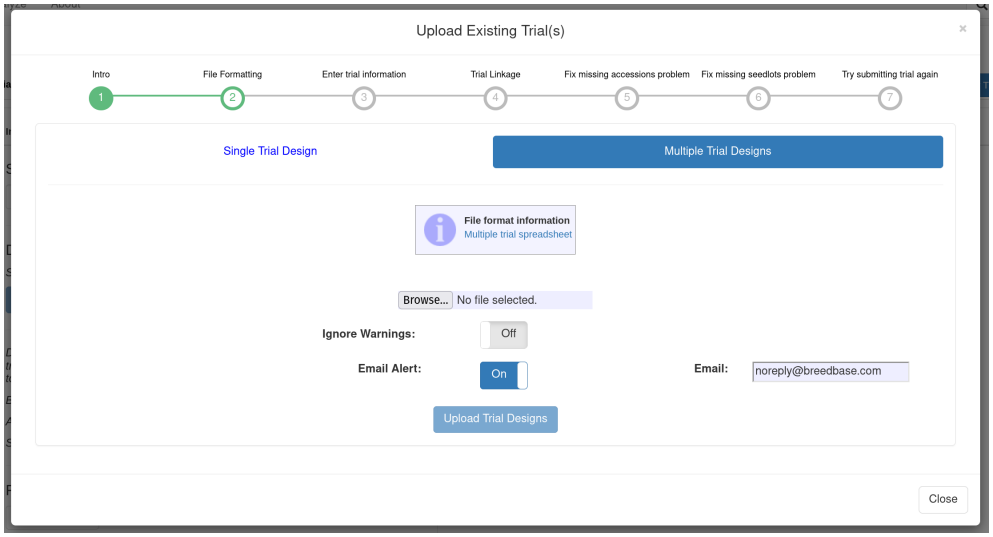
To add multi-location trials, simply select the multiple locations while using the 'Add Trial' form.

This will create a separate trial for each selected location, but they will share the same design and will be grouped in a single folder.

By default each trial design will have a fresh randomization, but if desired you may check the "Use same randomization for all locations" option.

10.2.5 Email alert for multiple trial design upload

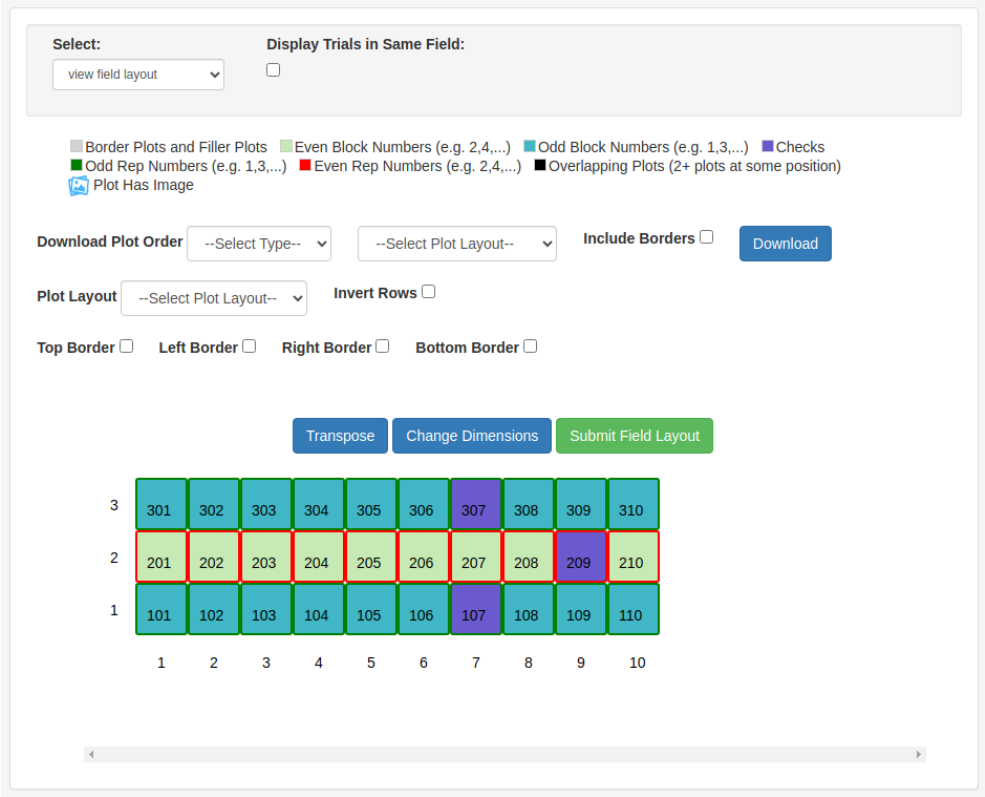
When uploading multiple trials from a file, you have the option to receive email notifications by clicking the "Email Alert" checkbox. By default, the system will use the email address associated with your account, but you have the option to enter a different email address if you prefer. After submitting, the upload process runs in the background, allowing you to continue using the interface without interruptions. Once the process completes, you will receive an email with the upload results.



10.2.6 Viewing Plot Layout and Trait HeatMap

10.2.6.1 Viewing plot layout

In the "Field Layout Tools and Phenotype Heatmap" section of a Trial Detail page, the trial physical layout is displayed by default. The relative position of the plots will be displayed based on the row and column positions given to the plots during the trial creation or upload steps. The plots are color-coded based on the plot's rep and block numbers and whether or not it is used as a check. Hover the mouse over the plot to see details about a specific plot.



If there is more than one trial grown in the same physical field, the trial layouts of all of the trials can be shown together if the trials share these properties:

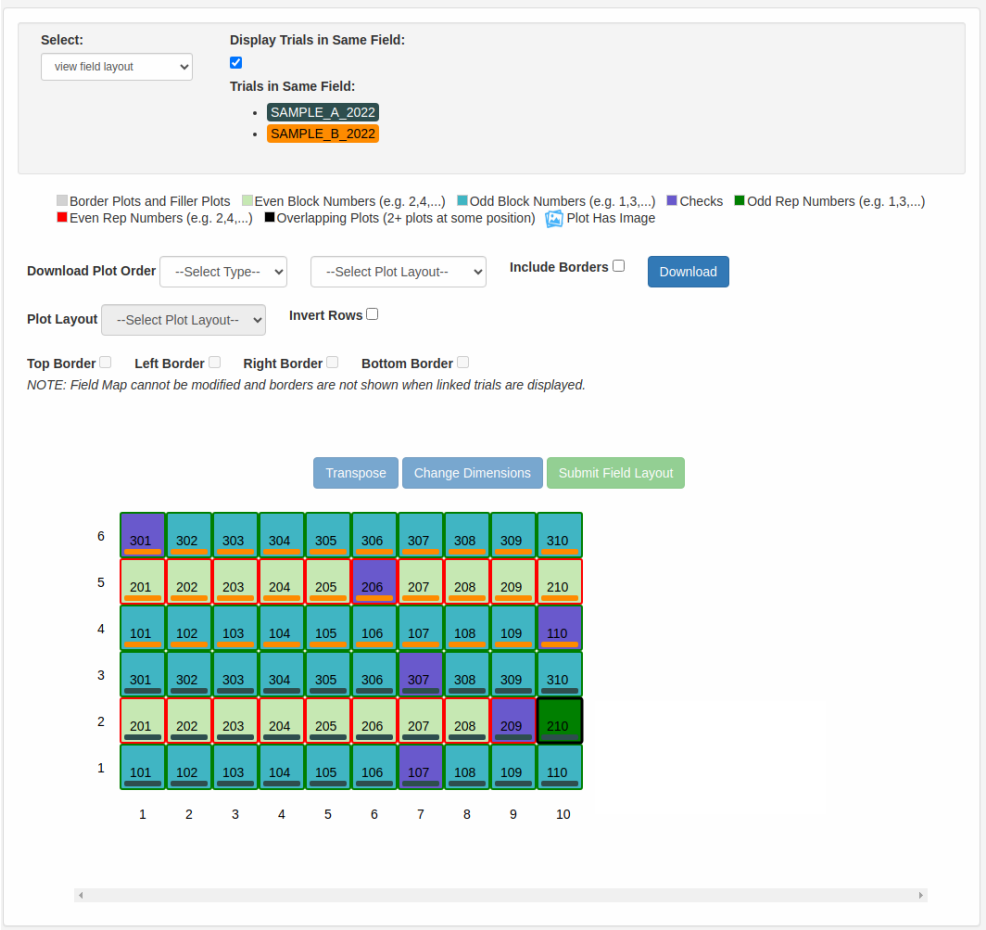
Each trial has the same year

Each trial has the same location

The location type of the trials' location is set to Field

The row and column positions of all of the plots (across the related trials) don't overlap. For example, trial #1 starts at row 1 and trial #2 starts at row 10.

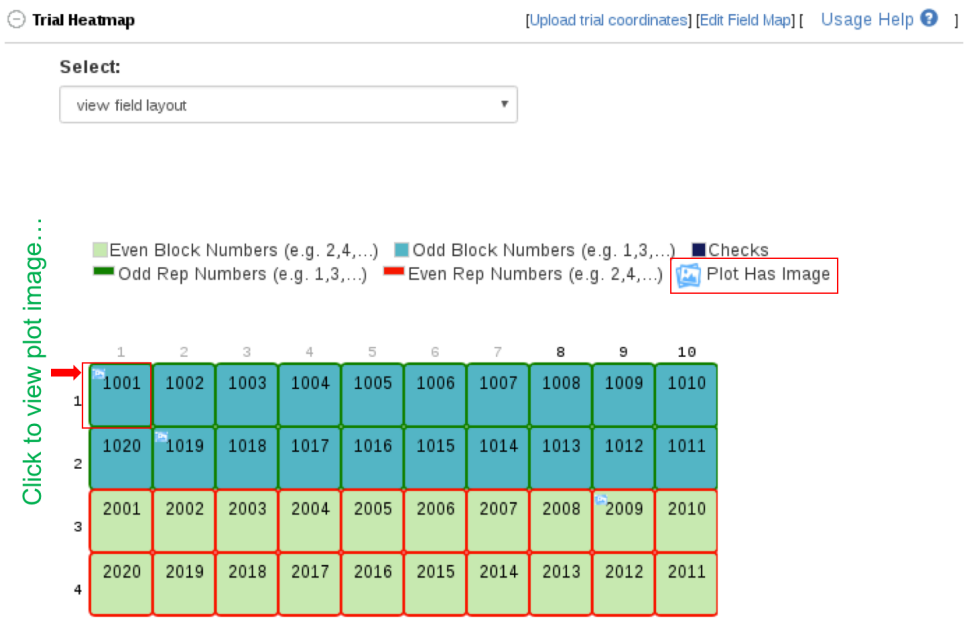
When these conditions are met and you check the "Select Trials in Same Field" checkbox, the plots from all of the related trials will be displayed on the same field layout. The plots will be color-coded by trial. The planting order and harvest order downloads will include the plots from all of the displayed trials in the order in which the plots occur in the field.



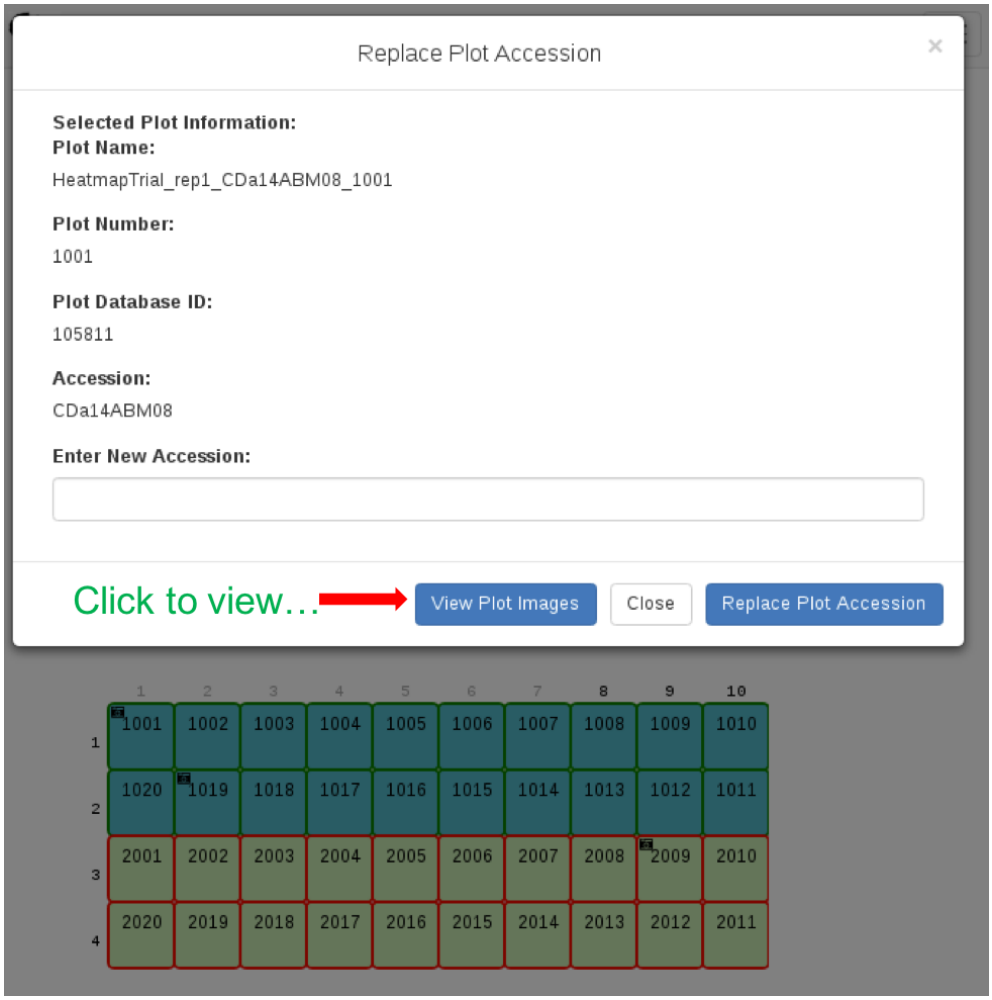
10.2.6.2 Viewing plot layout for multiple trials

Tracking plot images on fieldMap

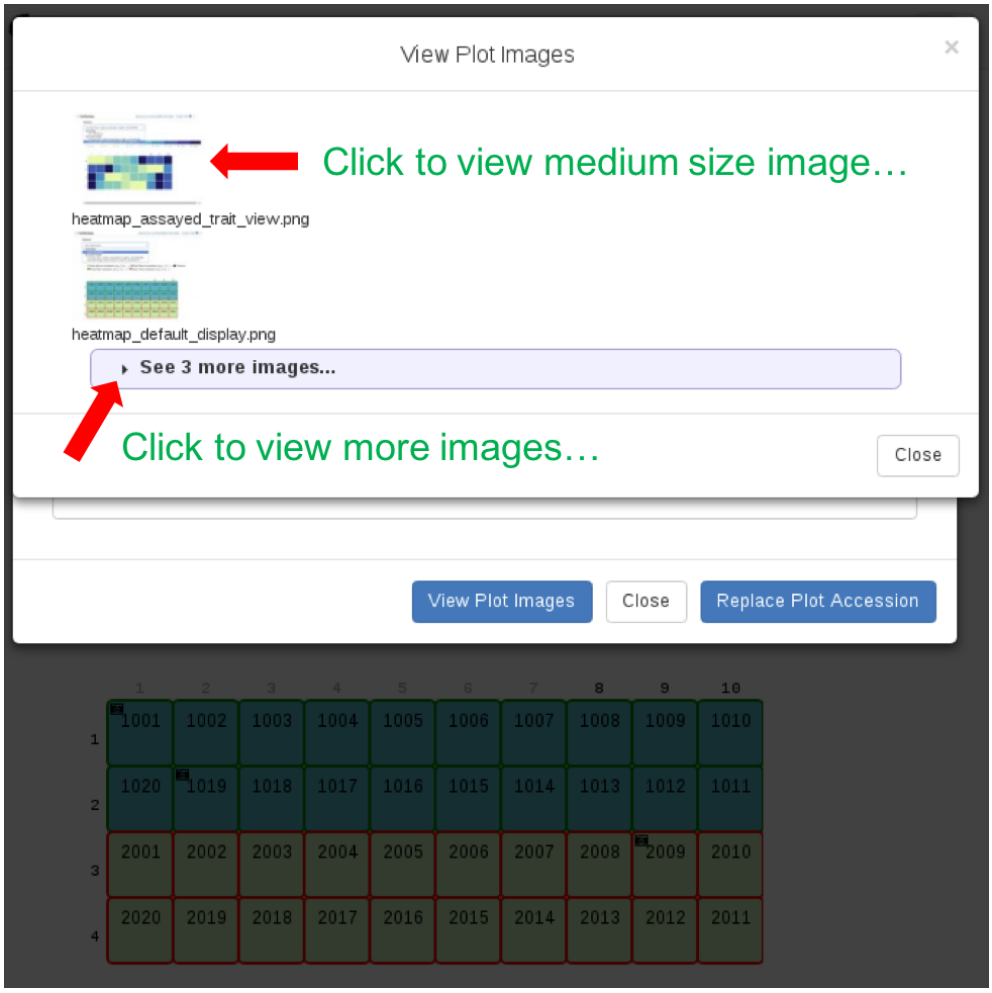
Plot images can be seen on fieldMap if a plot is associated to any image.



To view plot image(s), click on a plot, a dialog will appear.



On the appeared dialog, click on View plot images. To see more images if a plot has more that 2 images, click on See more images... Medium size of an image can be viewed by clicking on an image.

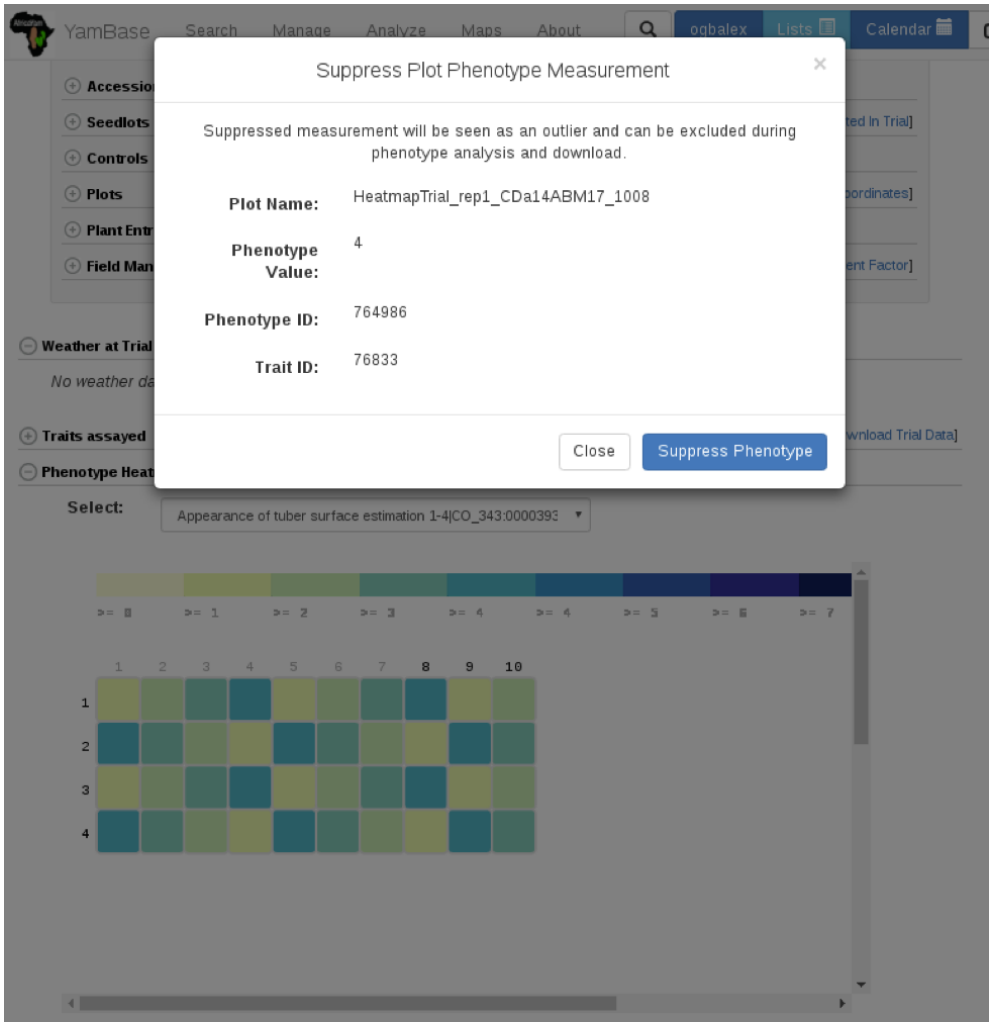


Viewing assayed trait heatmap

Phenotype heatmap can be viewed by selecting a specific assayed trait from the selectbox drop-down. Mousing over the plots, highlights the plot in green and also displays the plot's field information including the selected trait's phenotype value.

Suppressing Plot Phenotype

Clicking on a plot on the heatmap would display a dialog that has a button for suppressing a plot phenotype value for a given trait. A suppressed plot value can be excluded during trial analysis and phenotype download.

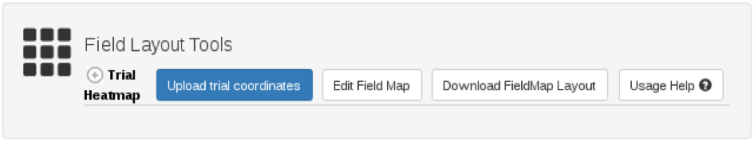


10.2.7 Adding additional information in the "Trial Detail" page

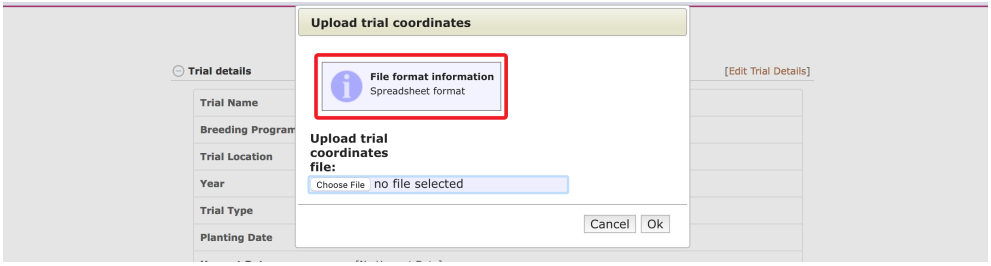
After you added a new trial to the database, you can edit trial details or add more information for that trial through the"Trial Detail" page.

Uploading Physical Trial Layout

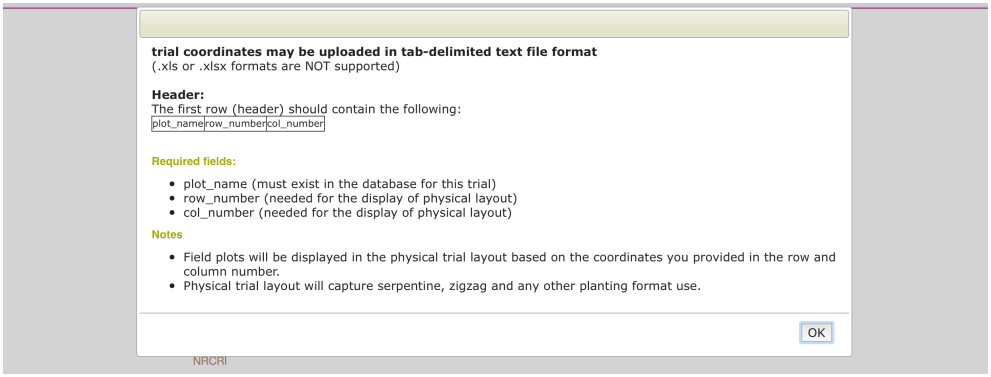
You can upload physical trial layout by clicking on the "Upload trial coordinates" button on the "Trial Detail" page.



Please check file format carefully. You can find file format information by clicking on the "Spreadsheet format" on the "Upload trial coordinates" window.



Spreadsheet format:



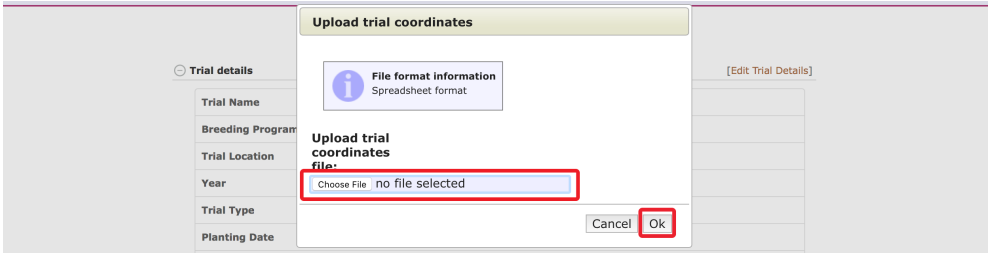
Physical Trial Layout File requirements

- All plot names in the file must exist in the database.
- The uploaded file should be tab delimited (txt).
- The first row (header) must contain the column names

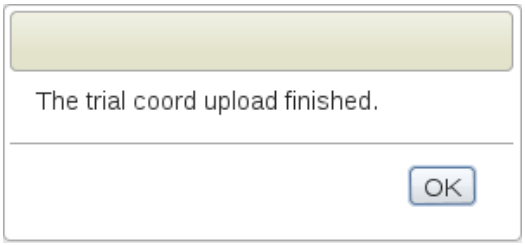
Example:

plot_name	row_number	col_number
plot1	1	1
plot2	1	2
plot3	1	3

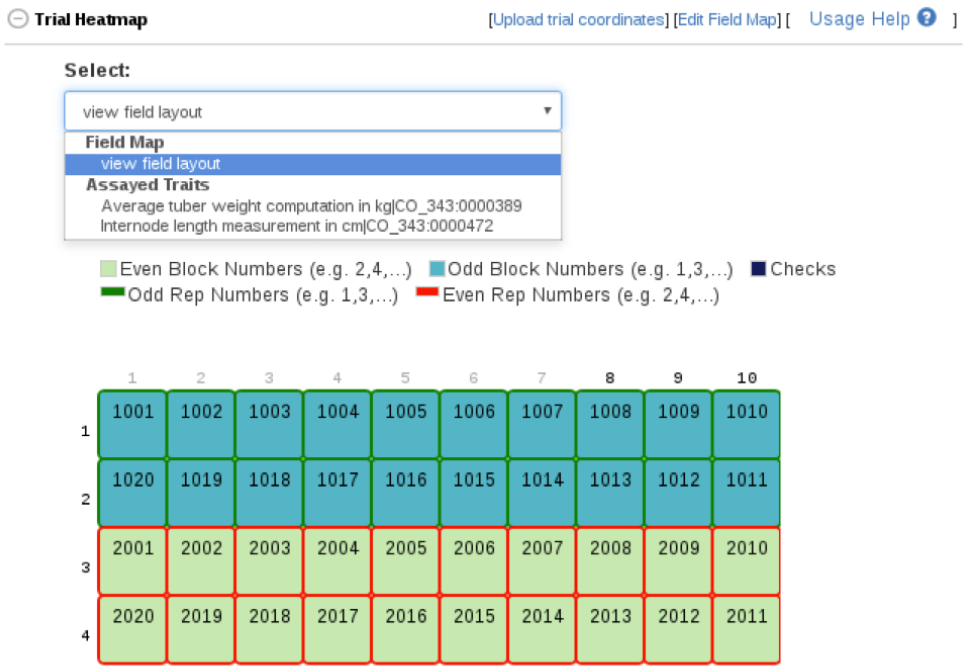
Select the trial layout coordinates file that you want to upload for this trial, then click "OK" button to upload the file.



The following message is displayed after the coordinates are uploaded.

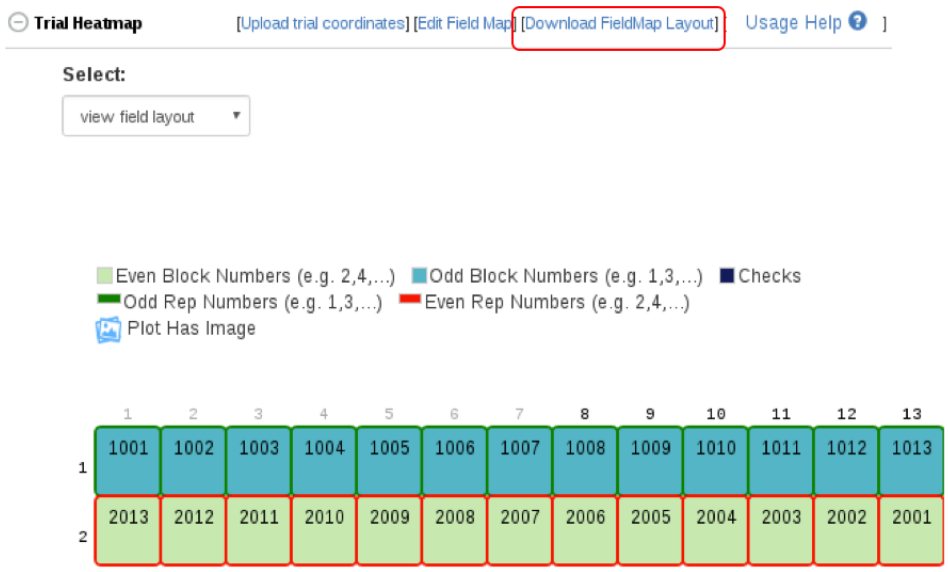


The field layout can be viewed by clicking on the "Trial Heatmap Section" to see a drop-down of the field map.

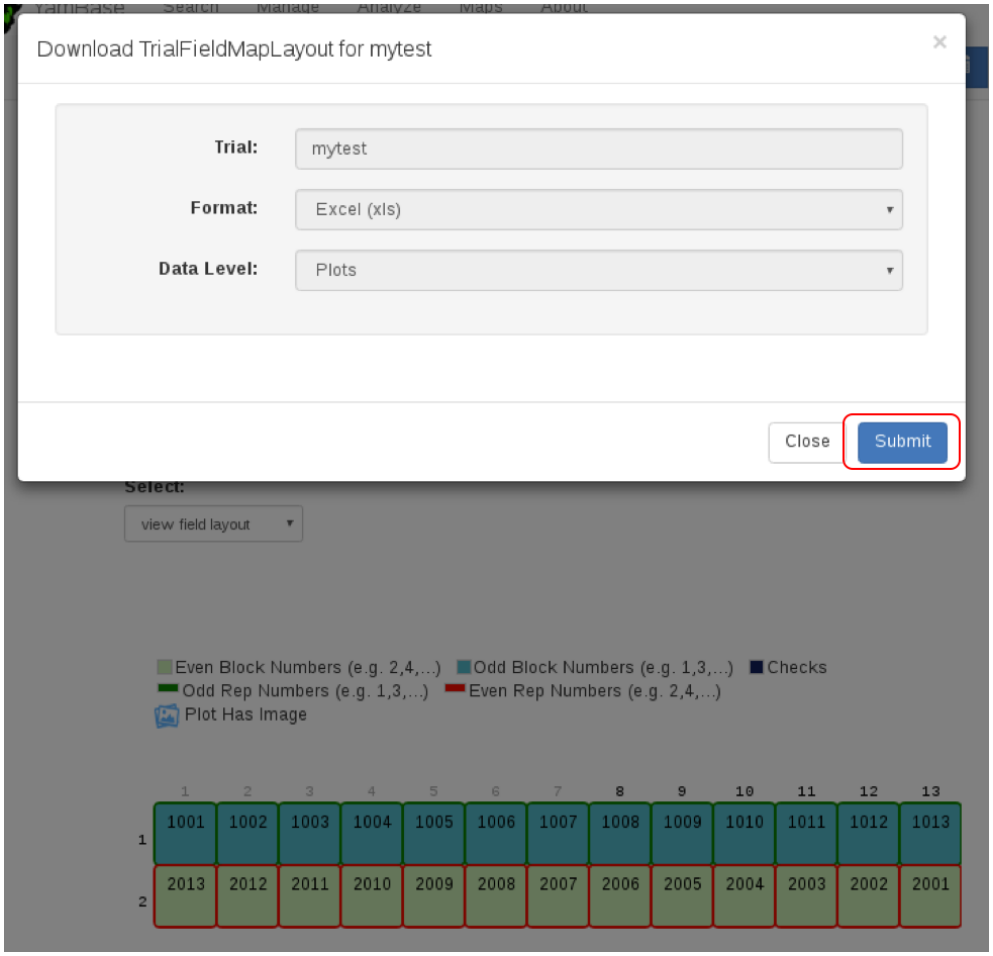


Downloading Field Map Spreadsheet

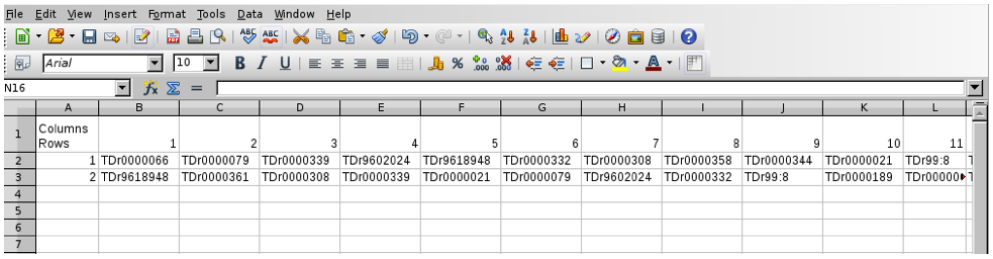
Field map spreadsheet can be downloaded if the trial has field coordinate (row and column numbers) uploaded for it plots. To download, click on the Download FieldMap Layout link on the Trial Heatmap section.



A dialog will appear, click on the submit button to download.

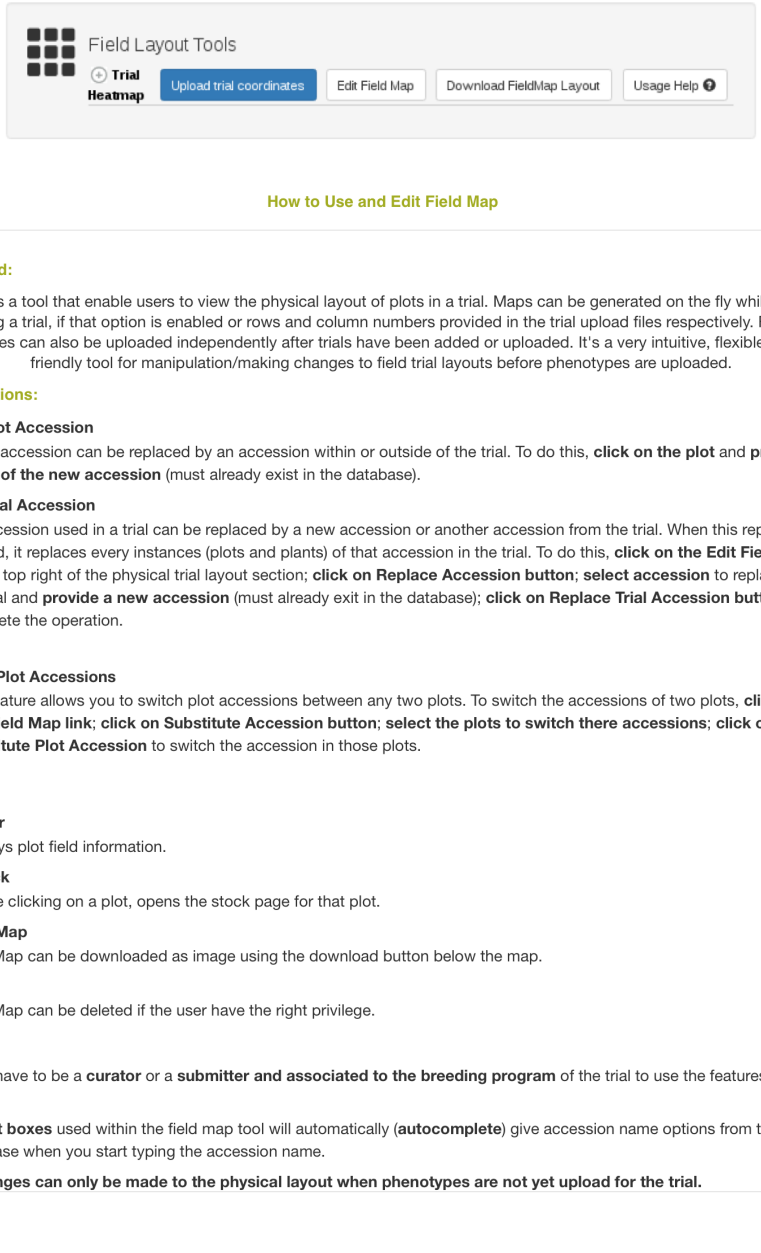


Click to view downloaded spreadsheet.



Editing Physical Trial Layout

"Usage Help" link contains information on how to edit physical trial layout.

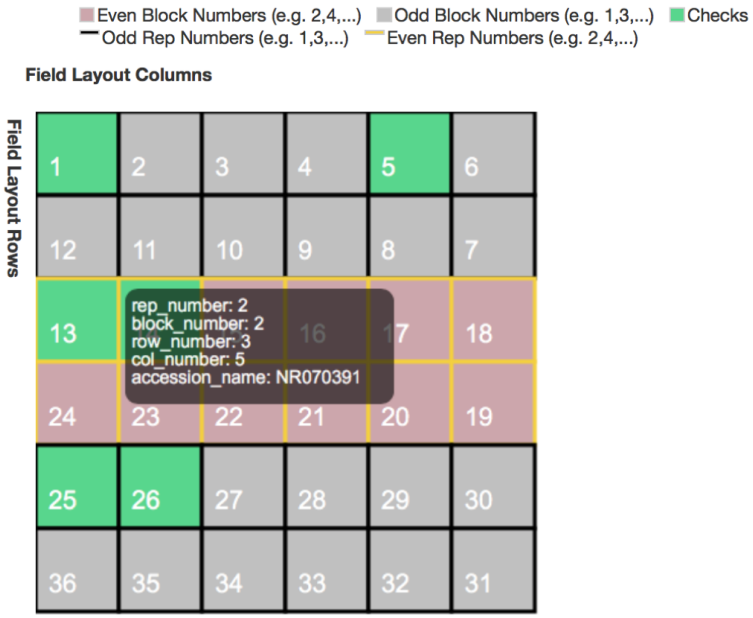


There are three different options for editing trial layout:

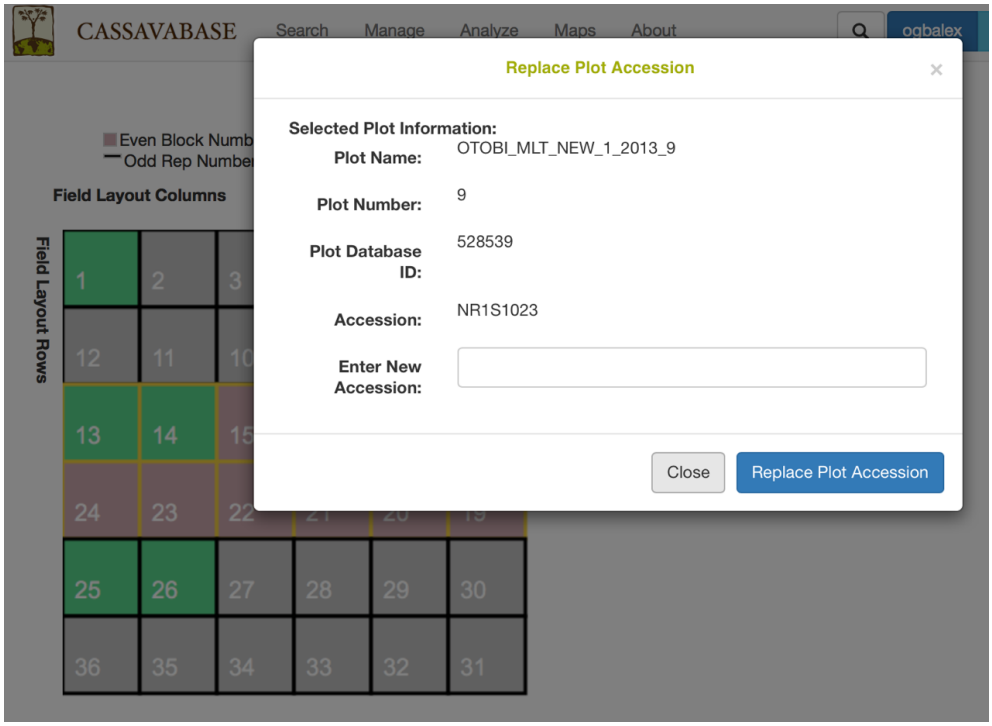
- Replacing plot accession by clicking on the plot in the layout.
- Replacing trial accession by using “Edit Field Map” link.

- Substituting plot accessions by using “Edit Field Map” link.

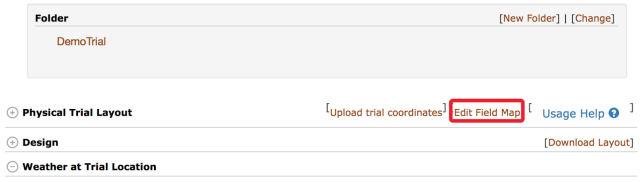
When you move a cursor over a plot on the trial layout, information for that plot appears.



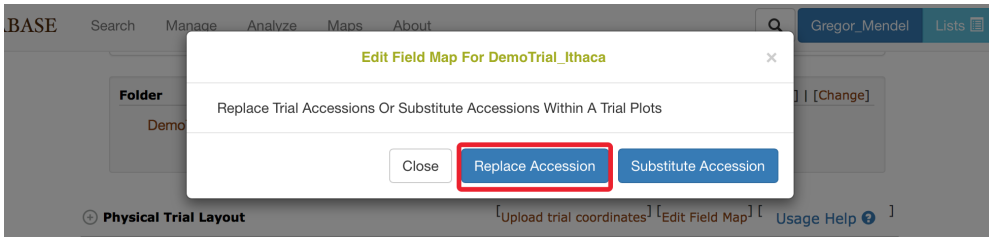
To edit a specific plot, clicking on that plot. Entering new accession on the “Replace Plot Accession” form, then clicking on “Replace Plot Accession” button.



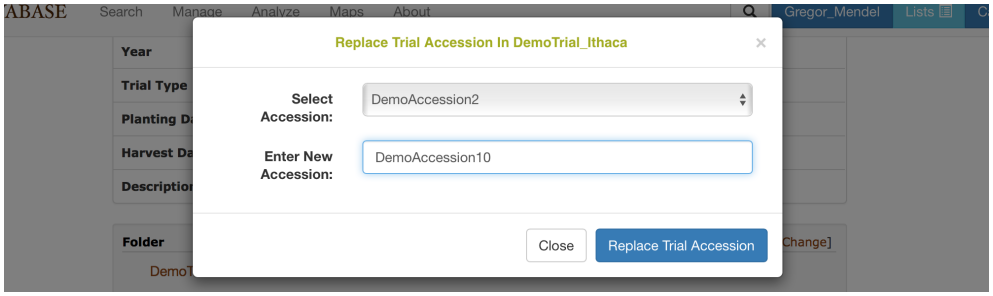
To replace an accession (in every plot/plant of that accession), clicking on "Edit Field Map" button.



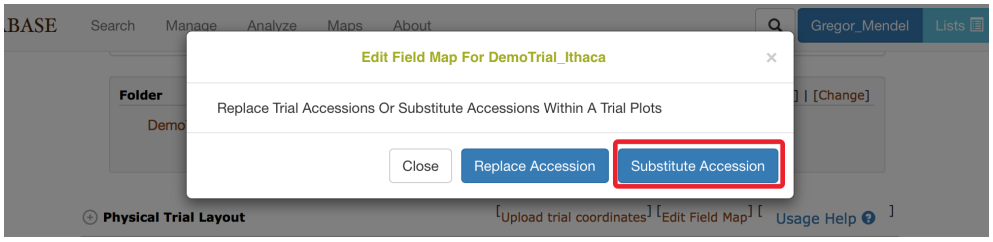
On the "Edit Field Map" window, clicking on "Replace Accession" button.



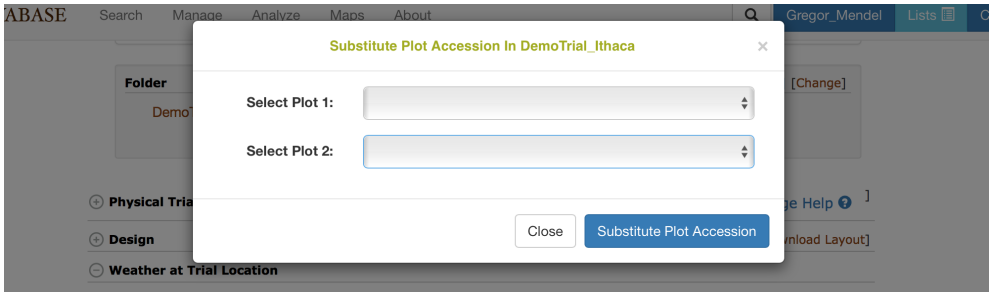
Selecting any accession that you want to replace and entering your new accession, then clicking "Replace Trial Accession" button.



You can switch plot accessions between any two plots by clicking on "Substitute Accession" button.



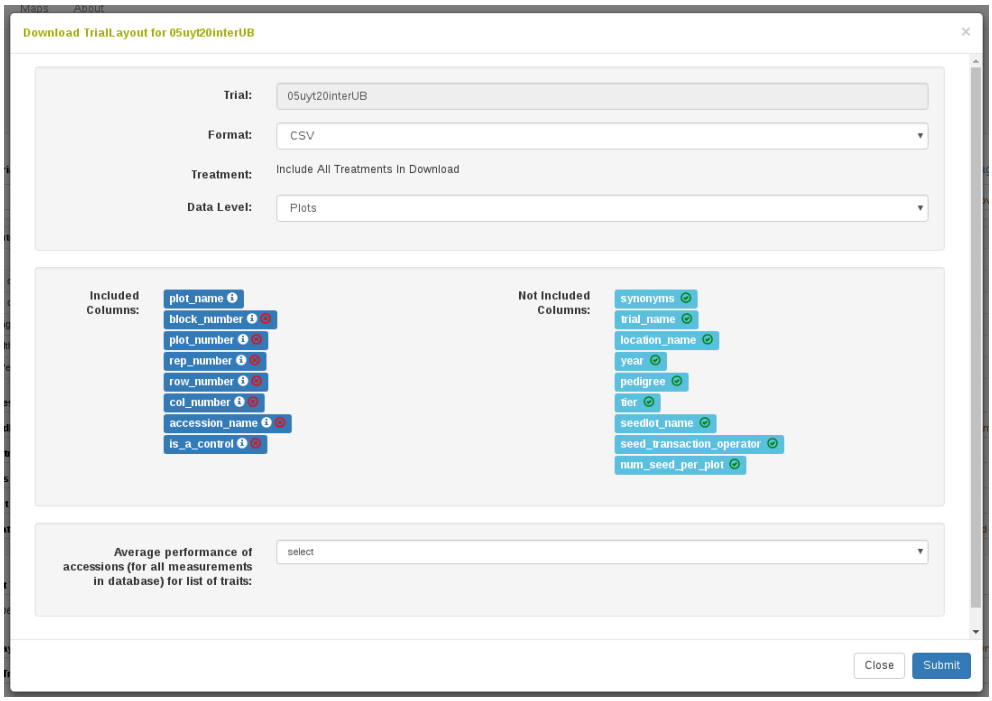
On the "Substitute Plot Accession" form, selecting the two plots that you want to switch, then clicking on the "Substitute Plot Accession" button.



10.2.8 Downloading the Trial Layout from the "Trial Detail" page

Click on "Download Layout" on the Trial Detail page.

The trial layout includes all information regarding the observation units in the experiment. The observation units can be plots, plants, or subplots. The trial layout can include trial design information such as the block_number and rep_number. It can also include physical map information such as the row_number and col_number, if that information is available for the trial. The trial layout also includes information regarding treatments that have been applied in the field. Optionally, the layout can give information regarding accession's global performance for a list of traits.



10.2.9 Adding Plant Entries To Your Trial

After you added a new trial to the database you can choose to add plant entries to your trial. Adding plant entries enables plant level phenotyping. It is generally better to enter data at the plant level into the database because it is always possible to calculate plot level phenotypes from the individual plant data.

Plant entries can be added to your trial in two ways: 1) Automatically generated by the database. The only input required is the number of plants per plot. 2) Uploaded in an XLS or XLSX file. This allows you to specifically name your plant entries.

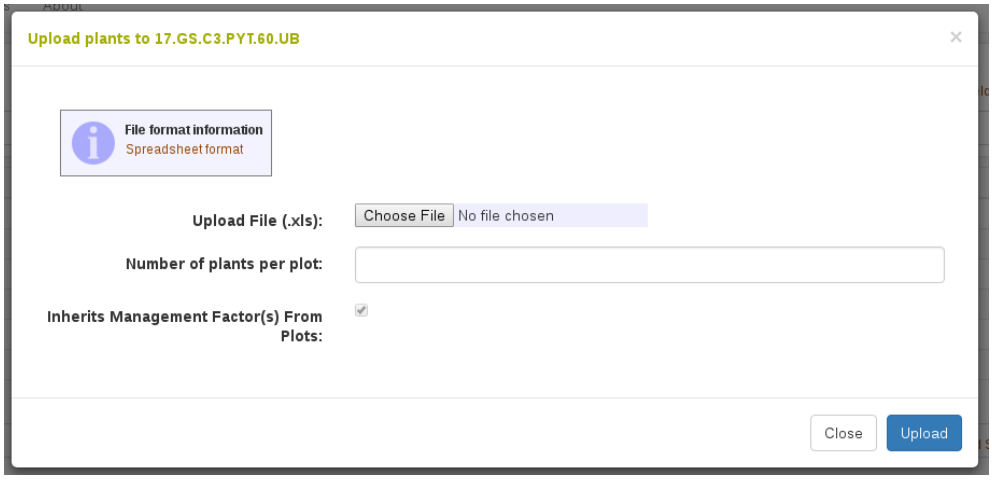
These two options are available in the "Plant Entries" section on the Trial Detail Page, as shown in the screen shot below.

Automatically Generate Plant Entries

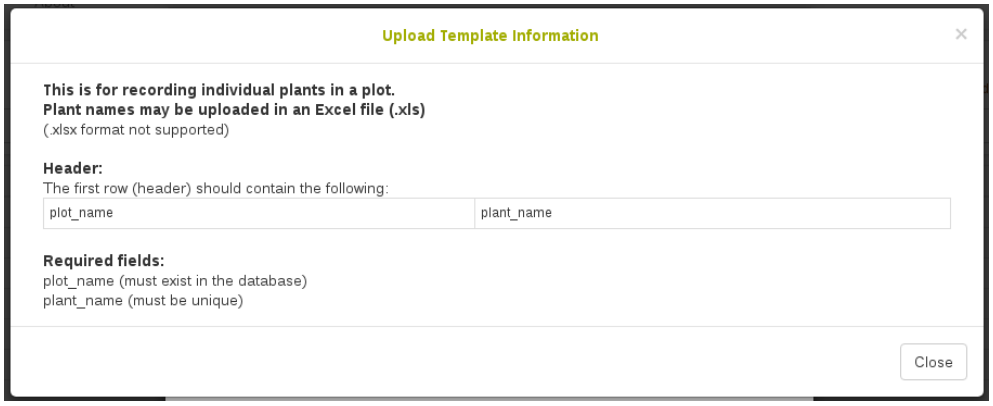
Clicking on "Add plant entries" opens the following dialog box. The only input required is the number of plants per plot. This will create plant entries that are named as a concatenation of the plot_name and the plant's index number e.g. plot_name_plant_1

Upload Plant Entries

Alternatively, you can choose to upload an XLS or XLSX file that contains the names of the plant entries. Clicking on "Upload plant entries" opens the following dialog box.



Clicking on "Spreadsheet format" will give you information about the XLS or XLSX file to upload. Clicking this will open the following dialog box.



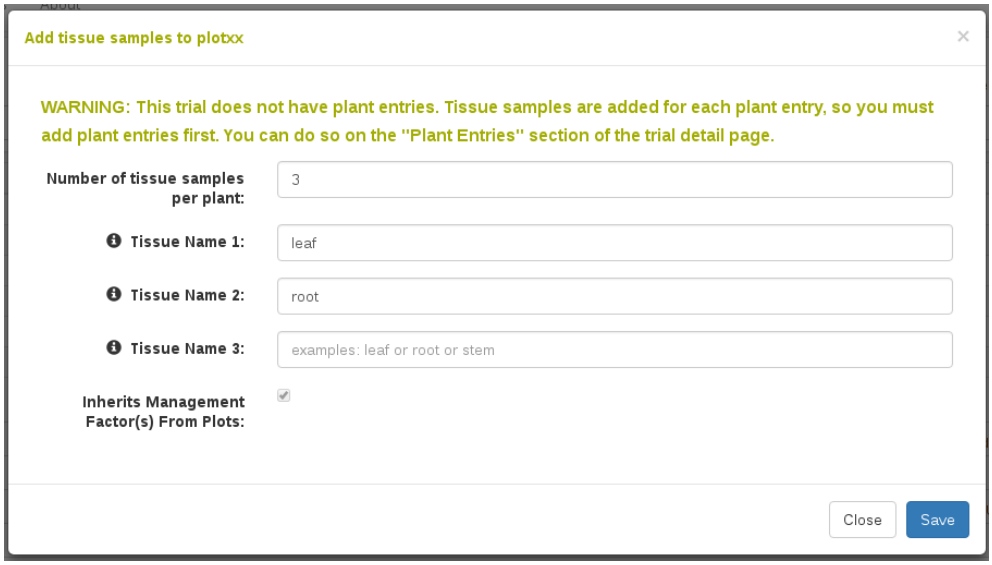
This shows you that the files requires the header to contain "plot_name" and "plant_name". The plot_name must exist in the database already and the plant_name must be unique in the database.

Along with the file, you must specify "number of plants per plot". This is intended to be the total number of plants that were plants. If the file you upload shows three plants in one plot and four plants in another plot, that is fine.

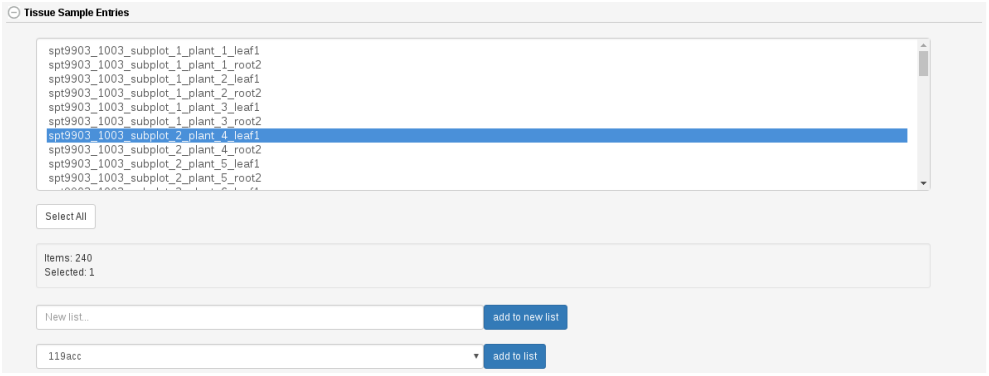
10.2.10 Adding Tissue Sample Entries To Your Trial

Some trials require tissue samples to be collected from plants in a field trial. The database will generate these tissue sample identifiers for you and will maintain all relationships with the plant, plot, accession, etc. To begin, go to the Design section of a trial's detail page and open the "tissue sample entries" section. Please note that tissue samples are directly related to plants, therefore your trial requires plants before you can add tissue samples.

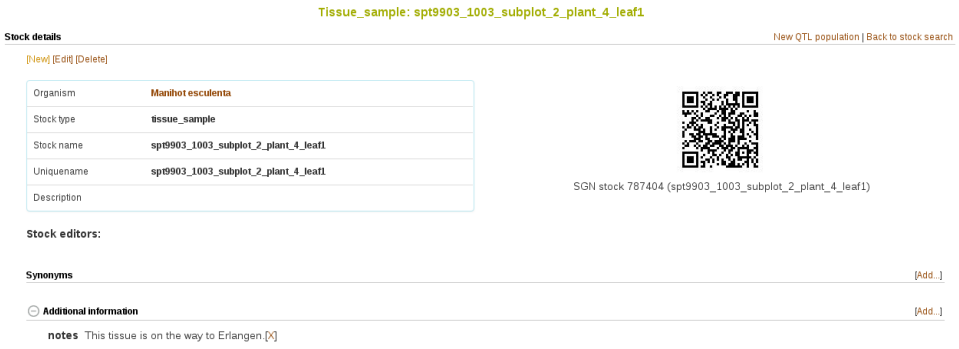
When you click on "Add tissue sample entries" you will see a dialog where you specify the number of tissue samples you require per plant. Once you have specified how many tissues samples, you can give specific words to distinguish samples, such as "root" or "stem", as seen below.



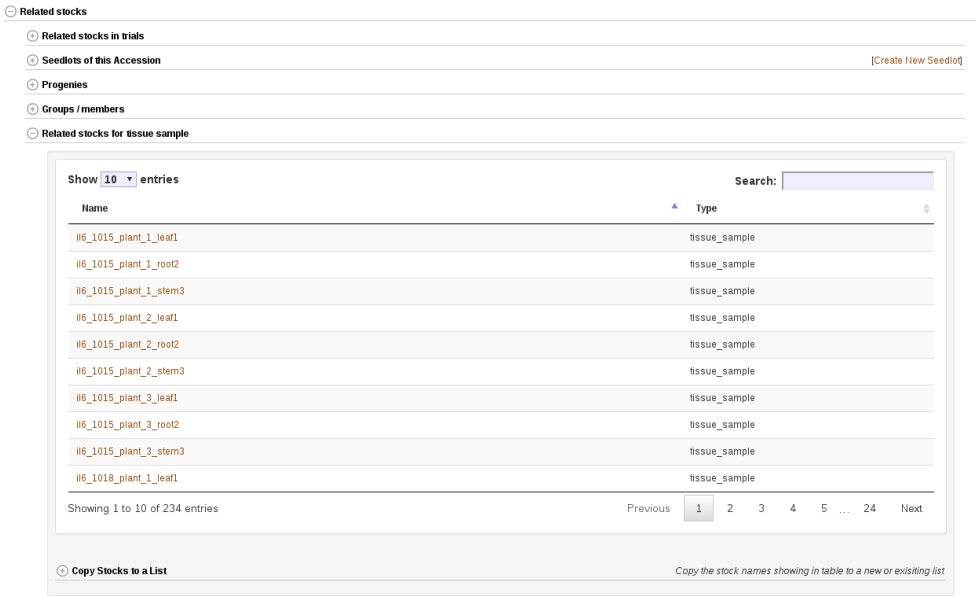
Once you have added tissue sample entries they will appear in the design section of the trial as seen below.



Each tissue sample has a detail page where you can add information about the sample, such as if it is in transit or in storage somewhere.



The related stocks section near the bottom of this detail page displays the relationships between all stocks, including tissue samples.

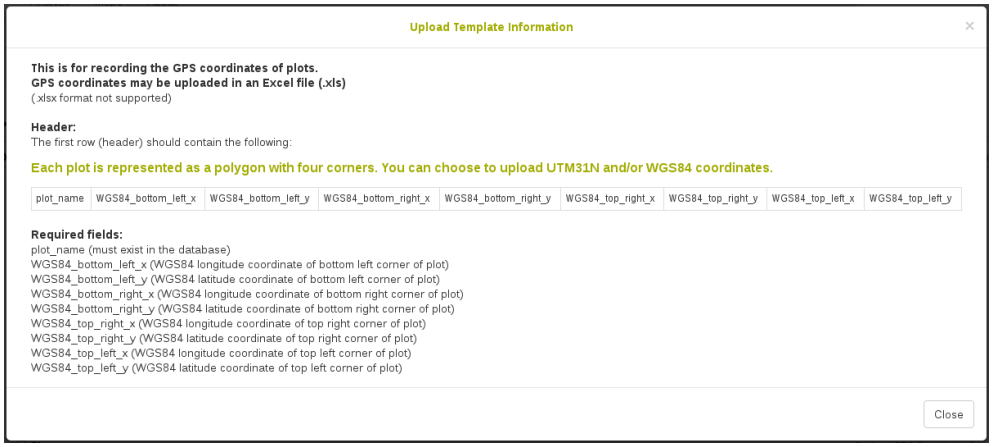


10.2.11 Uploading GPS Coordinates For Plots

You can upload GPS coordinates for the plots in your trial. There is a link on the Trial Detail Page as shown below.

Clicking on this link will bring up the following dialog.

Here you can upload an XLS or XLSX file. To see information on the format of the file that should be uploaded, click on "Spreadsheet format". This will bring up the following dialog.

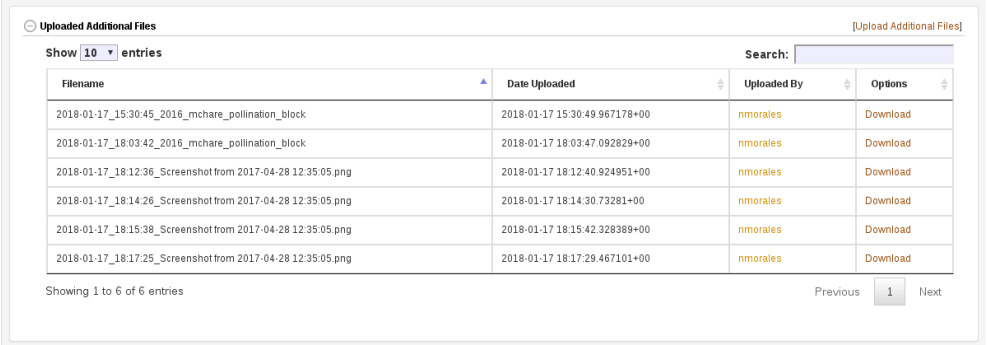


This dialog tells you that the file must be XLS or XLSX and must contain: plot_name WGS84_bottom_left_x WGS84_bottom_left_y WGS84_bottom_right_x WGS84_bottom_right_y WGS84_top_right_x WGS84_top_right_y WGS84_top_left_x WGS84_top_left_y The GPS coordinates should be WGS84 format and specify a four-pointed polygon around the plot.

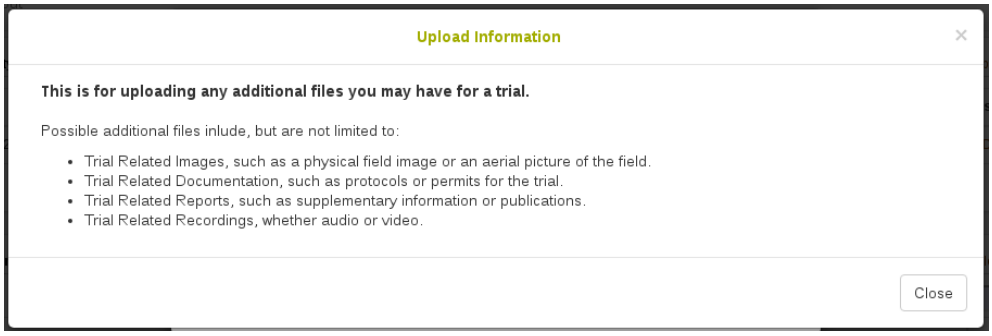
10.2.12 Uploading Additional Files To Trial

It may be of interest to you to upload additional documents, images, or recordings to your trial. To do this, scroll down to the "Uploaded Additional File" section on the trial detail page. From here you can view and download

any of these additional files.

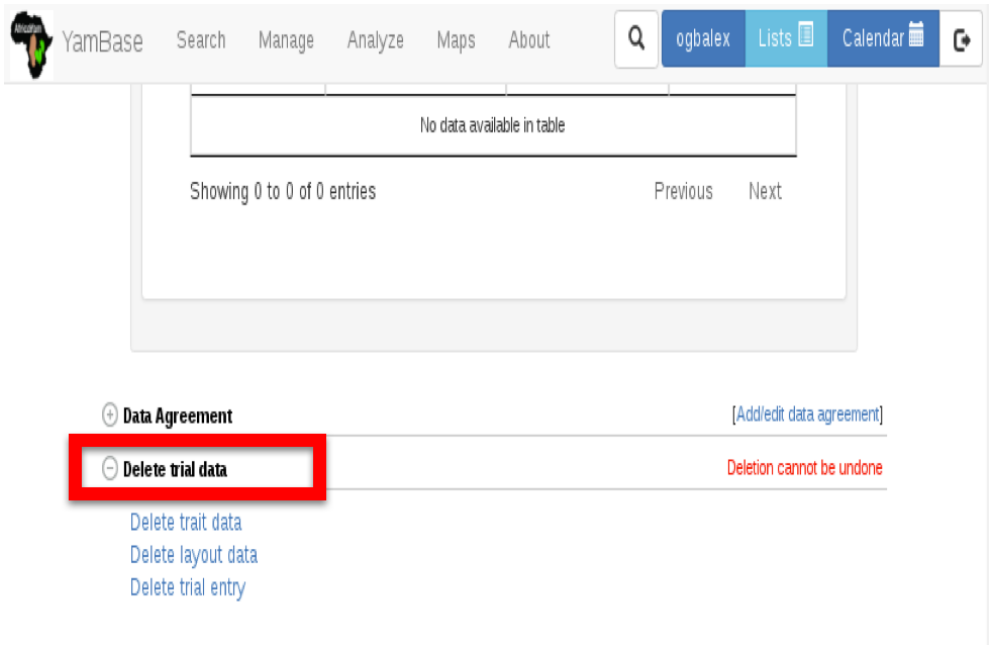


To upload an additional file, click on the "Upload Additional Files" link. A dialog will appear where you simply select your desired file. For information, you can click "Upload information" to see the following message.

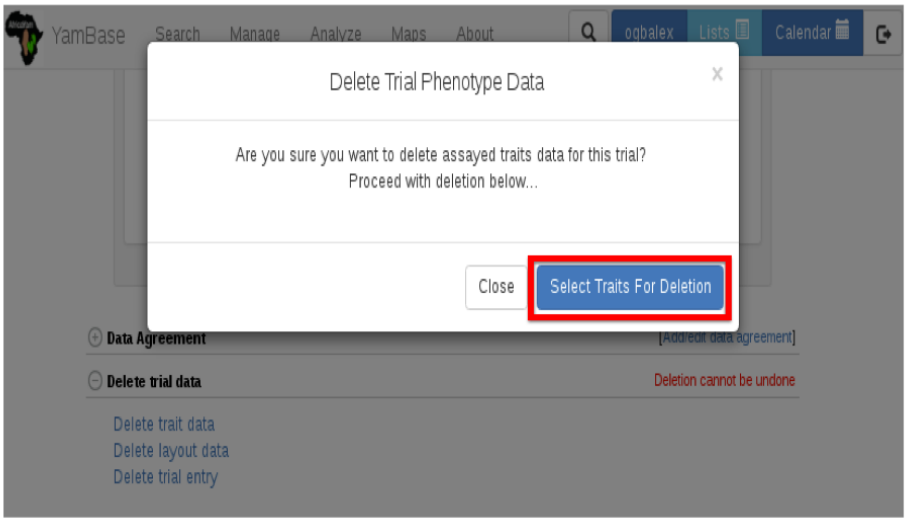


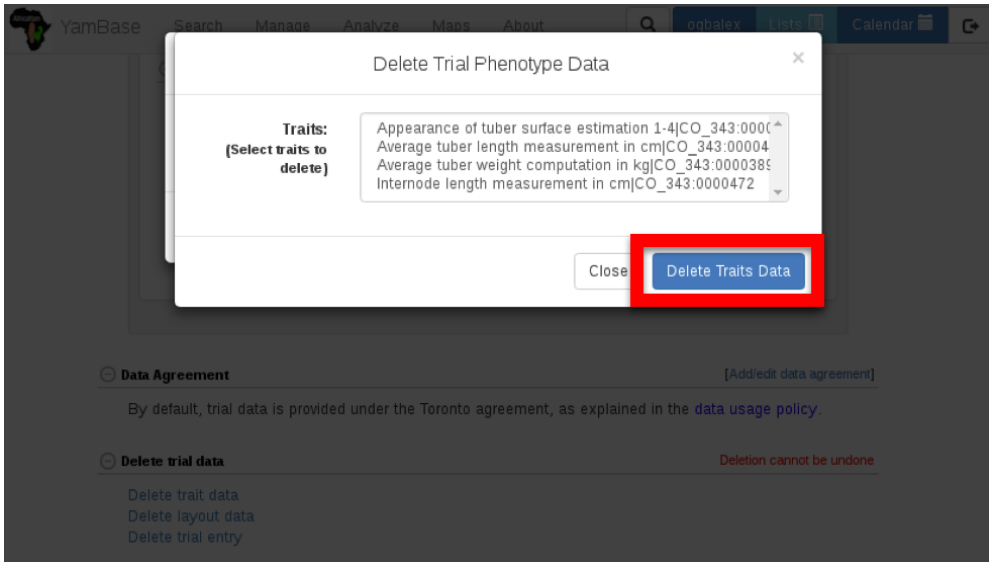
10.2.13 Deleting Trial Data

To delete a trial data, click on the "Delete trial data" section. There are links to delete traits, layout and trial entry data.



To delete assayed trait data, click on "Delete trait data" link. On the appeared dialog, confirm deletion by clicking on the "Select Traits For Deletion" button, then select one or more traits to delete from the trial.





To delete trial layout data, click on the "Delete layout data" link. Confirm deletion on the appeared dialog.

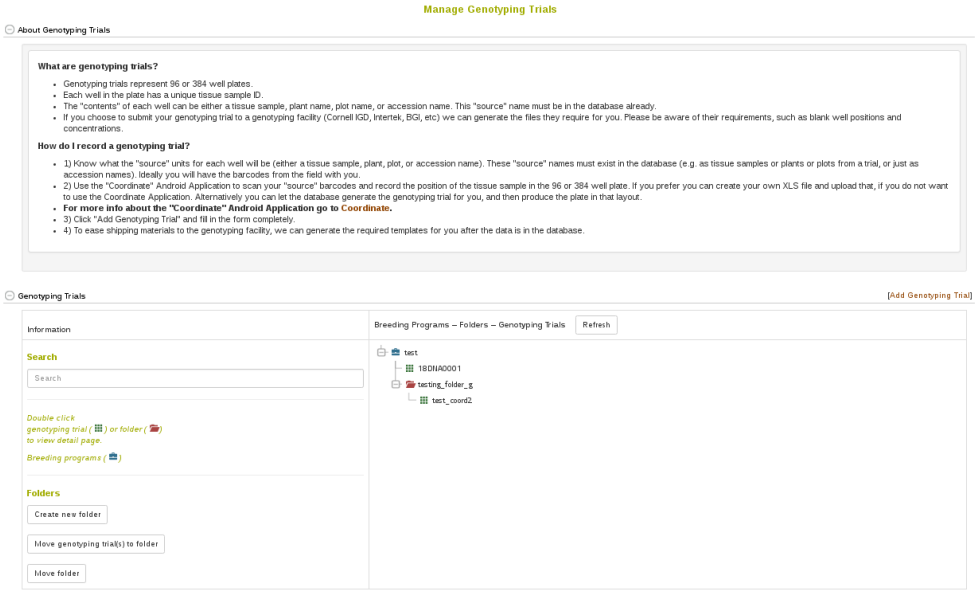
To Delete trial entry, click on "Delete trial entry" link. Confirm deletion on the appeared dialog.

Chapter 11

Managing Genotyping Plates

Genotyping Plates represent the content of a genotyping plate sent to a genotyping facility (e.g. samples in specific wells). To streamline this process, it is possible to upload this information or let the database create a plate for you. Once the genotyping plate is saved in the database it is then possible to export the information directly to genotyping facilities that are BrAPI compliant. The genotyping facility can then provide status information to us via BrAPI.

To begin go to Manage->Genotyping Plates.

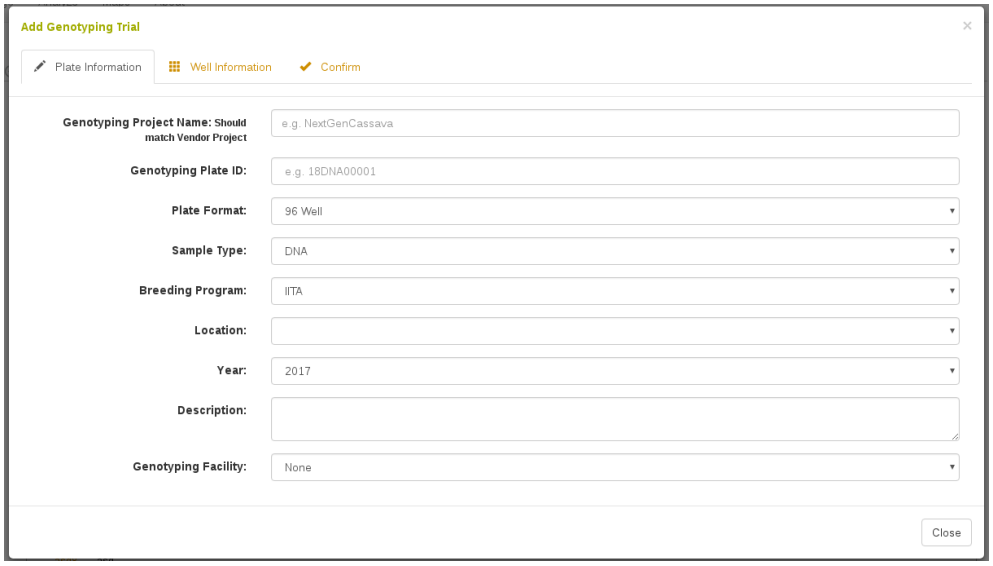


Here the genotyping plates are divided by Breeding Program. These sections can be expanded by clicking on one.

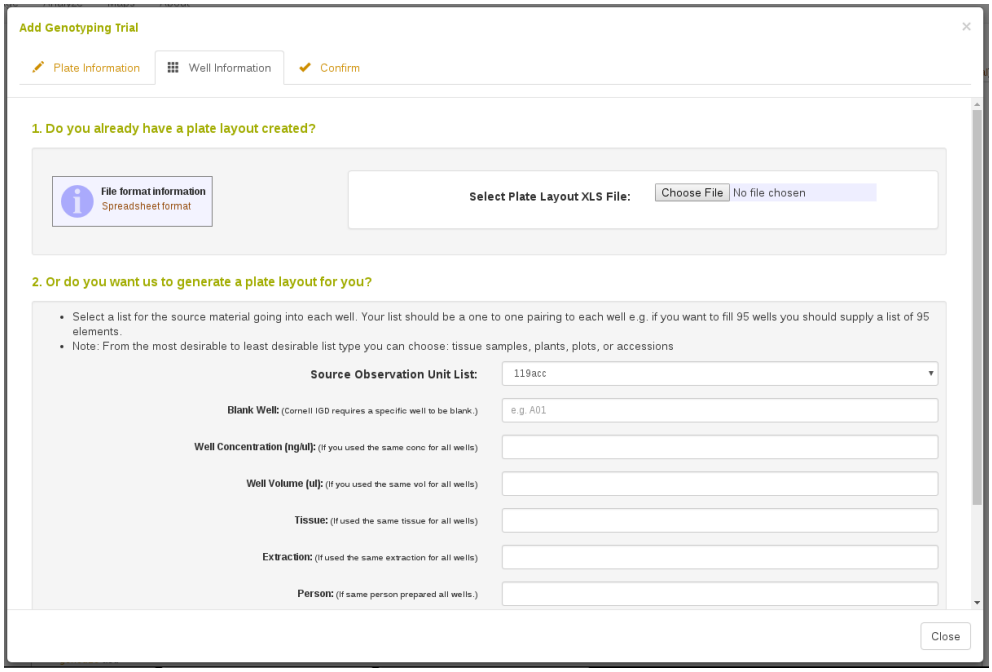
11.1 Adding a New Genotyping Plate

To begin, click on "Add Genotyping Plate". Notice that this form is split into three sections: "Plate Information", "Well Information", and "Confirm". The first section is for defining information about the genotyping plate, such as a Plate identifier, plate format (96 well), etc. The second section is for defining the samples in the wells, such as sample names, sample concentrations, well position, etc. The final section is for Submitting the info.

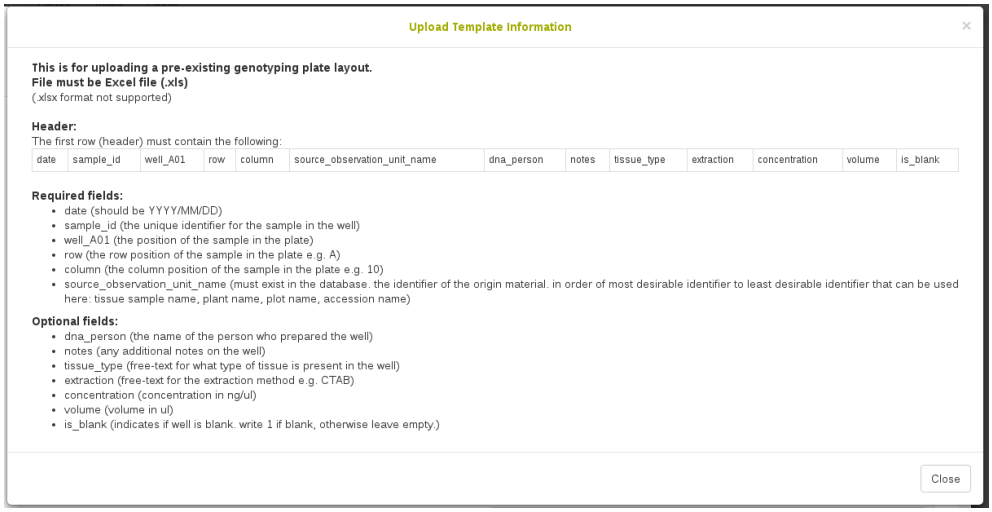
All fields in the Plate Information section are required.



In the Well Information section you can choose to either 1) Upload an XLS or XLSX spreadsheet with your sample layout or 2) let the database create the sample layout.



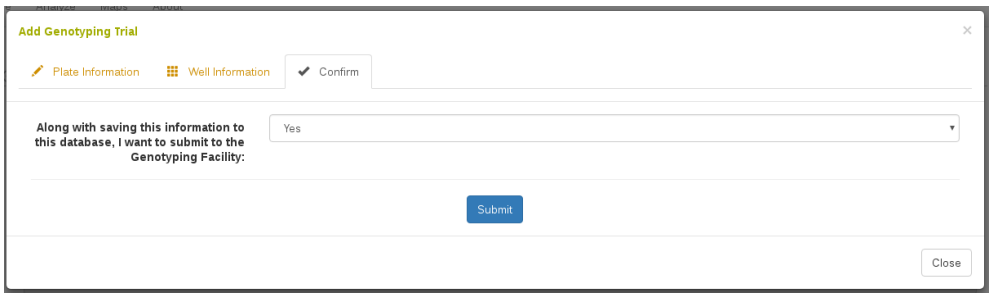
If you choose to upload an XLS or XLSX spreadsheet, the Spreadsheet Template info requires the following:



In either case, the sample identifier is generally a concatenation of Plate name and well position, e.g. MyGenotypingTrial1_A01. In either case, you need to provide a "source_observation_unit_name" for each sample. This

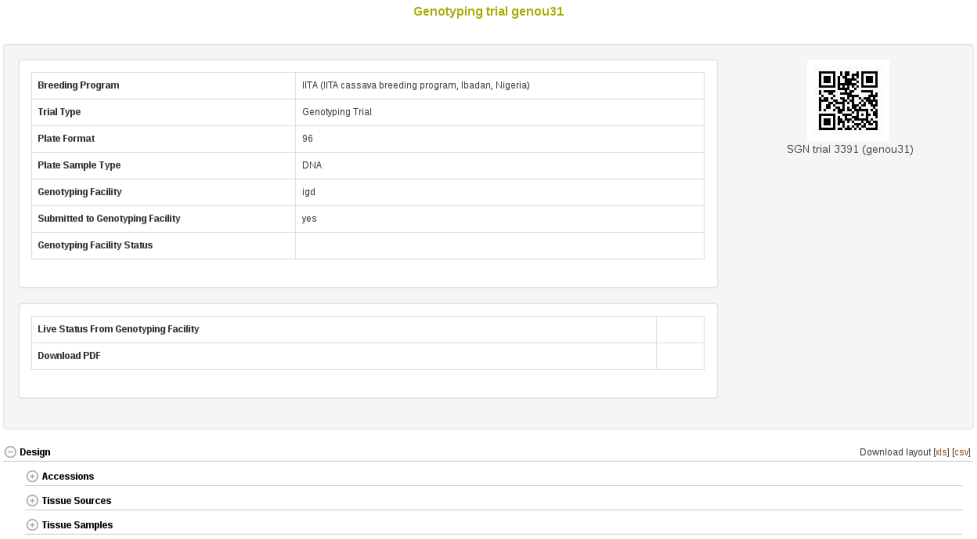
can be a tissue sample name, a plant name, a plot name, or an accession name; however, in any case, the identifier must already exist in the database. This allows us to link the sample in the well to specific field trial plots, or, plants, or tissue_samples. If you only know which accession is in the well, you can use the accession name.

In the final Confirm section you can decide whether to submit this information to the genotyping facility you selected. This requires that the genotyping facility is BrAPI compliant to work.

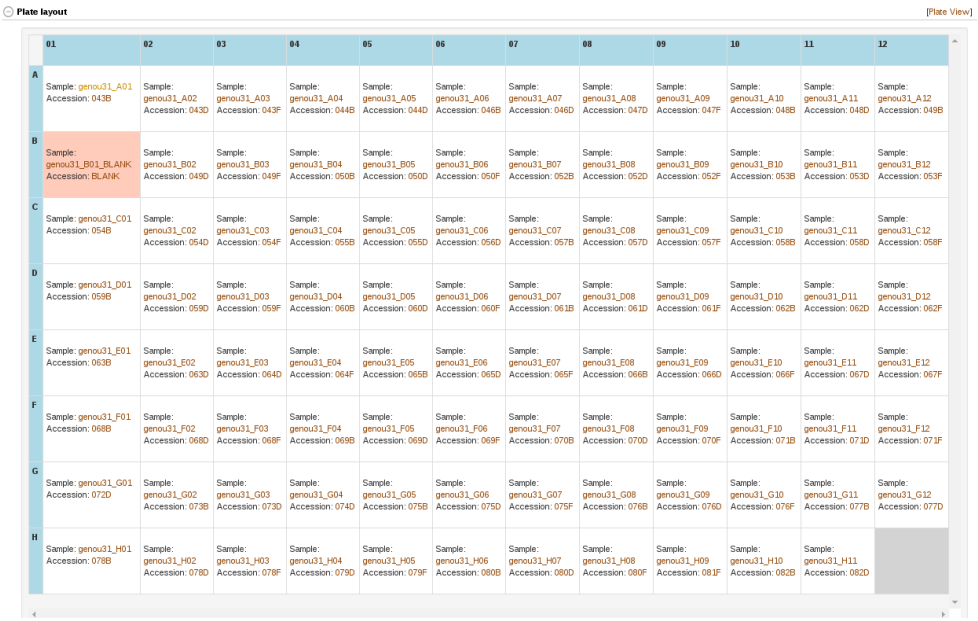


11.2 Genotyping Plate Detail Page

If you open a specific genotyping plate, it will take you to the detail page. Here you can see the Accessions used in the plate (if you created the trial and the source_observation_unit_names you used were plots, this will still work because we know the accession of the plot or plant or tissue sample).

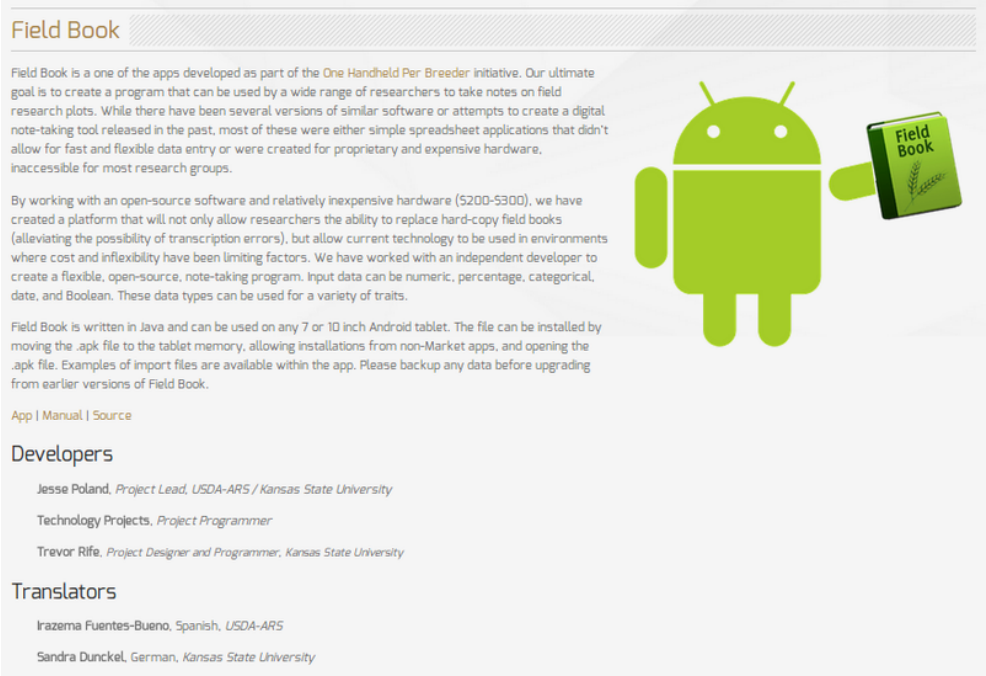


Further down you can see a graphical representation of your plate with well positions. This can be 96 well or 384 well depending on your plate format.



Chapter 12

Using Field Book App



SGN databases support the Android Field Book App for collecting phenotypic data in the field with tablet computers. The app is available here:

https://play.google.com/store/apps/details?id=com.fieldbook.tracker

- The app can also be downloaded directly from the Google Play store.

There is no charge for the app.

- Field Book App requires two files for collecting data: Field layout file and trait file.
- SGN databases can generate the field layout file and trait file, which can be downloaded onto your computer, then transferred to an Android tablet device.

12.1 A typical workflow

1. Creating a *field layout file* based on the design of field trial
2. Creating a *trait file* from the list of traits
3. Downloading the field layout file and trait file from the database to your computer
4. Downloading the field layout file and trait file to the tablet (where the Field Book App is installed)
5. Collecting phenotypes
6. Exporting phenotypes from Field Book App to your computer
7. *Uploading the exported phenotype file* from your computer to the database

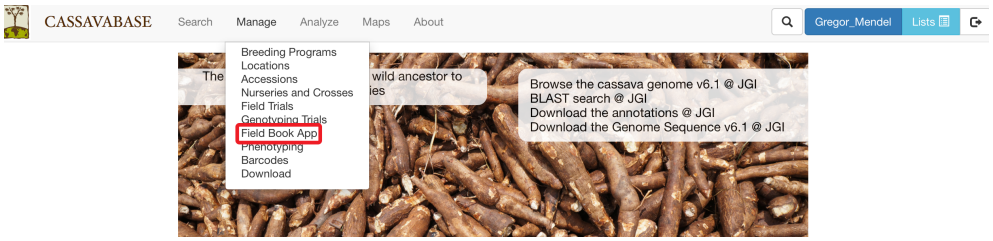
12.2 Creating Field Layout Files for the Field Book App

There are two alternative methods for creating "Field Layout Files".

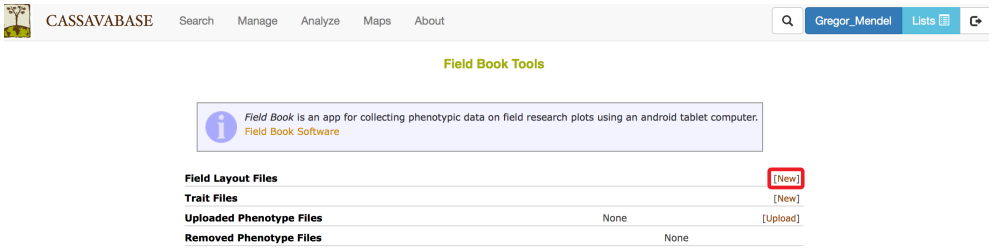
1. Using "Field Book Tools" page
2. Using "Trial Detail" page.

12.2.1 Creating "Field Layout Files" by using "Field Book Tools" page.

To access "Field Book Tools" page, clicking on "Field Book App" in the "Manage" menu.



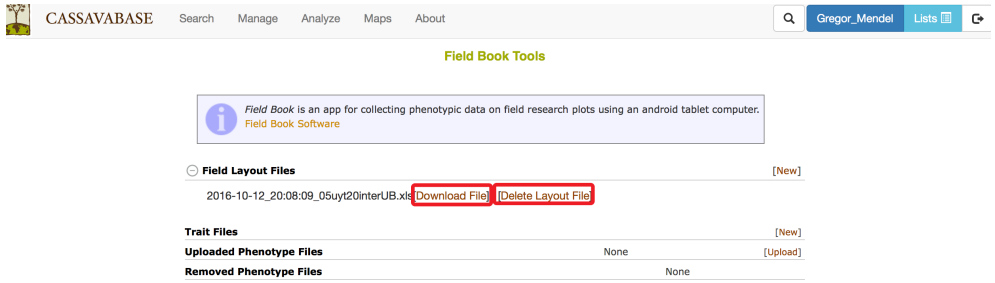
On the "Field Book Tools" page, clicking on "New"



On the "Download Fieldbook" window, selecting trial name and data level (plots or plants), then clicking on "Submit" button. A treatment can be selected, which allows you to record phenotypes based on treatment application. A list of traits can be selected, which provides a summary of an accession's global performance for those traits in the Fieldbook.

If the field book layout file was successfully created, a pop-up window will indicate that the field book layout file was saved successfully. Clicking on the file name will immediately download the file onto your computer. The file is also available to download on the "Field Book Tools" page, if you need to re-download it.

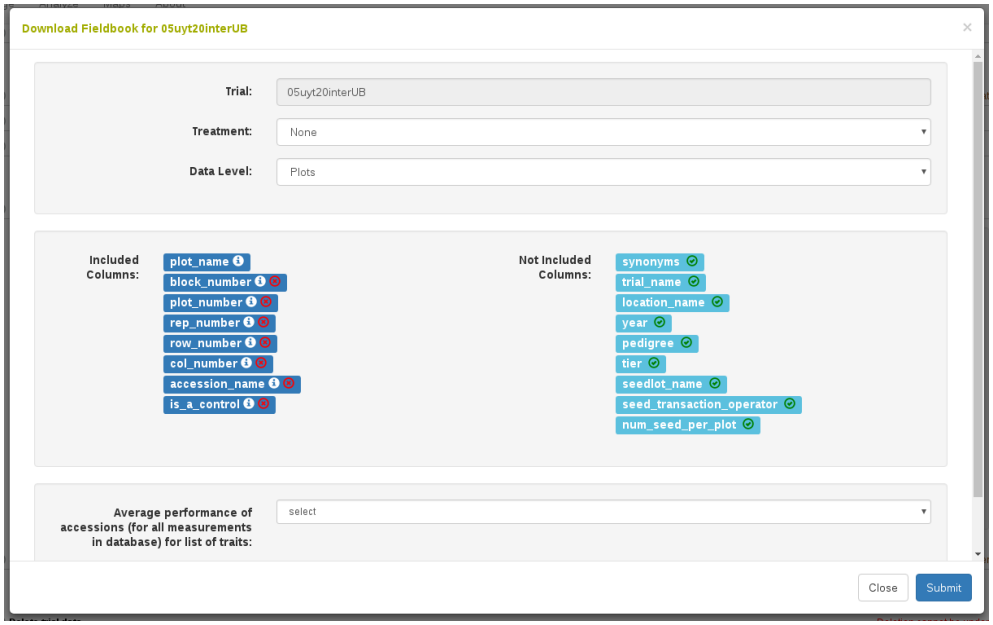
To download field layout file to your computer, clicking on "Download File", the file can then be transferred to your tablet. If you no longer want to keep the field layout file, clicking on "Delete Layout File".



12.2.2 Creating "Field Layout Files" by using "Trial Detail" page.

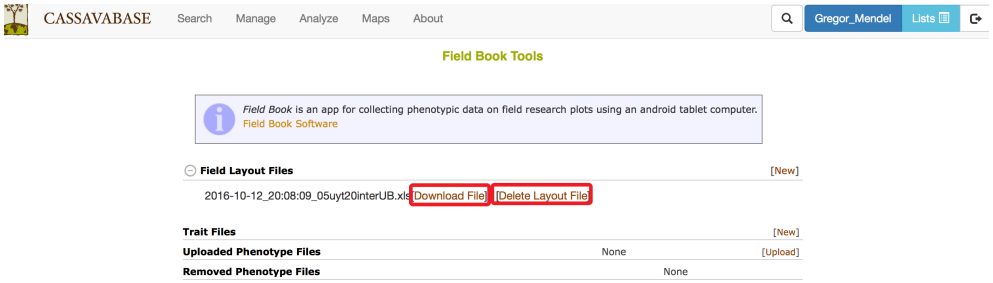
To create "Field Layout Files", go to the "Trial Detail" page of the trial that you want to create the file. On the "Trial Detail" page, scrolling down to the bottom of the page to find "Android Field Book Layout" in the "Files" section, then clicking on the "Create Field Book" link.

Clicking on the "Create Field Book" link will open a new window showing the name of the trial that you selected, as well as data level (plots or plants). A treatment can be selected, which allows you to record phenotypes based on treatment application. A list of traits can be selected, which provides a summary of an accession's global performance for those traits in the Fieldbook. To proceed, clicking on "Submit" button.



If the field book layout file was successfully created, a pop-up window will indicate that the field book layout file was saved successfully. Clicking on the file name will immediately download the file onto your computer. The file is also available to download on the "Field Book Tools" page, if you need to re-download it.

To download field layout file to your computer, clicking on "Download File", the file can then be transferred to your tablet. If you no longer want to keep the field layout file, clicking on "Delete Layout File".

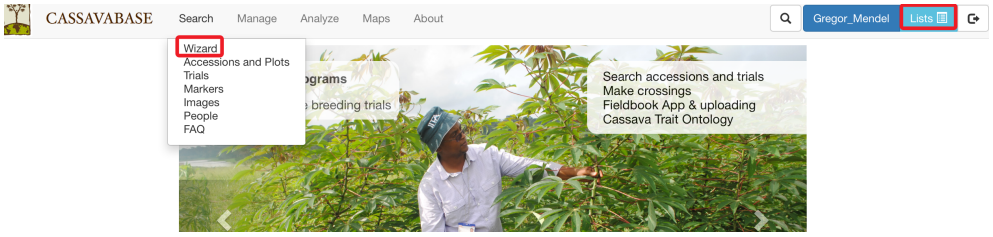


12.3 Creating Trait Files for the Field Book App

Steps to Create a Trait File:

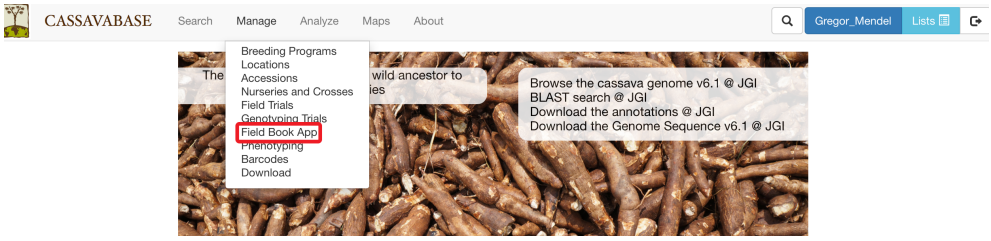
12.3.1 Creating a Trait List

After you logged in, lists can be created and managed using the Search Wizard or the "Lists" link. For more information on how to create lists, click *here*.

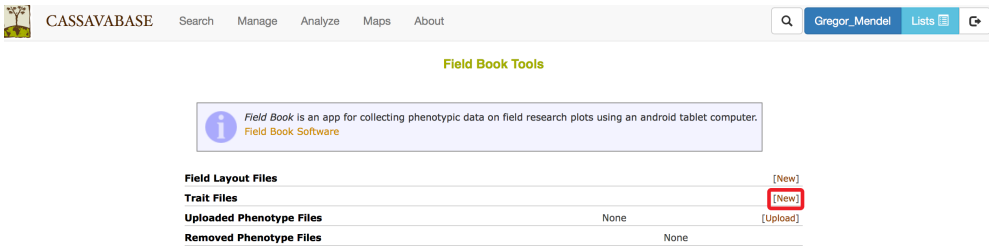


12.3.2 Creating a Trait File

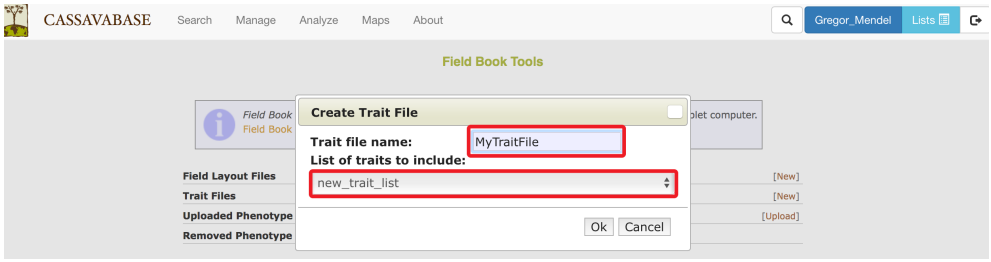
After you have your trait list, clicking on the "**Field Book App**" link found under the "**Manage**" menu tab. This will take you to the "Field Book Tools" page.



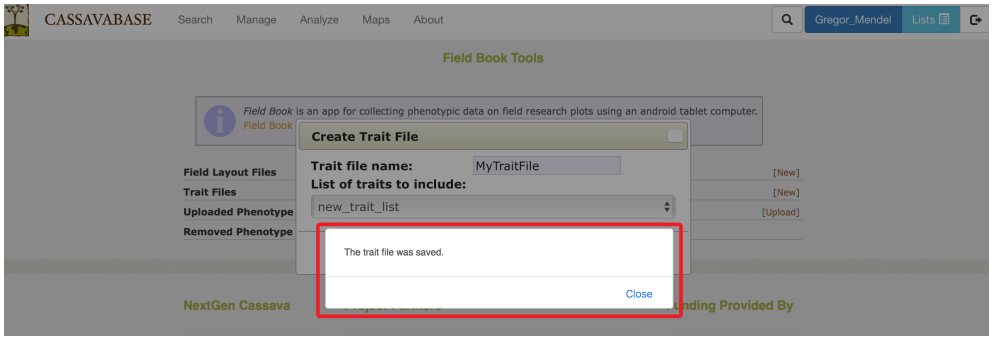
To create a new trait file, finding the heading “Trait Files”, then clicking on the “New” link.



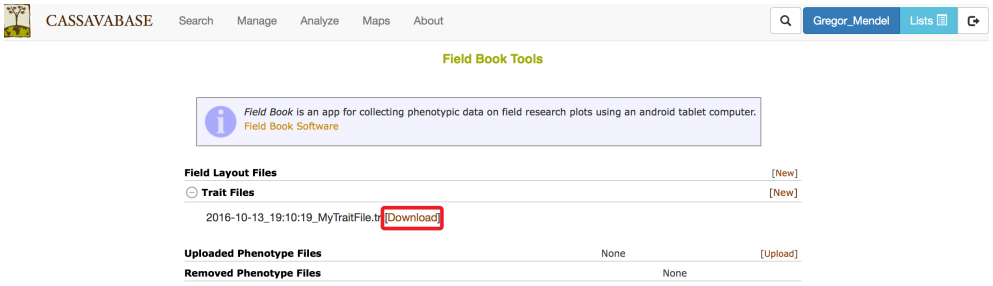
Clicking on the “New” link will open a dialogue box titled “Create Trait File”. Please enter your “Trait file name” and select “List of traits to include” from drop-down list that you previously created. You can only use traits included in the list. Check the box titled “Include Notes Trait” if you would also like to record and upload general plot notes in the field. Click “OK” to submit.



If your trait file was successfully created, a new window will indicate that the trait file was saved, then clicking on “Close”.



After the trait file was saved, you will see your file listed in the "Field Book Tools" page. Clicking on "Download" link to download the trait file to your computer.



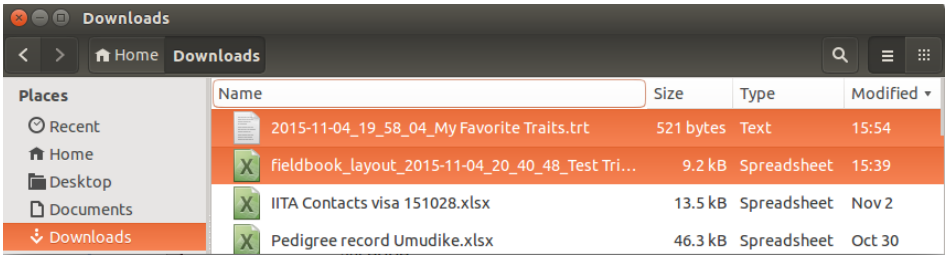
After downloading the trait file to your computer, the file can be transferred to an Android Tablet. You need the Android Field Book App to open the file. The Android Field Book App can be downloaded at: *http://www.wheatgenetics.org/bioinformatics/22-android-field-book*

12.4 Transferring Files from Your Computer to Android Tablet

12.4.1 Files on your computer

After downloading, Field Layout files and Trait files can be found in the "Downloads" folder of your computer. Field Layout files on your computer will have a prefix "fieldbook_layout_" added to the beginning of the file name. For example: **"2014-01-28_19:14:34_Trial Demo_location 6767.xls"** on the the database website will be saved as **"field_book_layout_2014-01-28_19:14:34_Trial Demo_location**

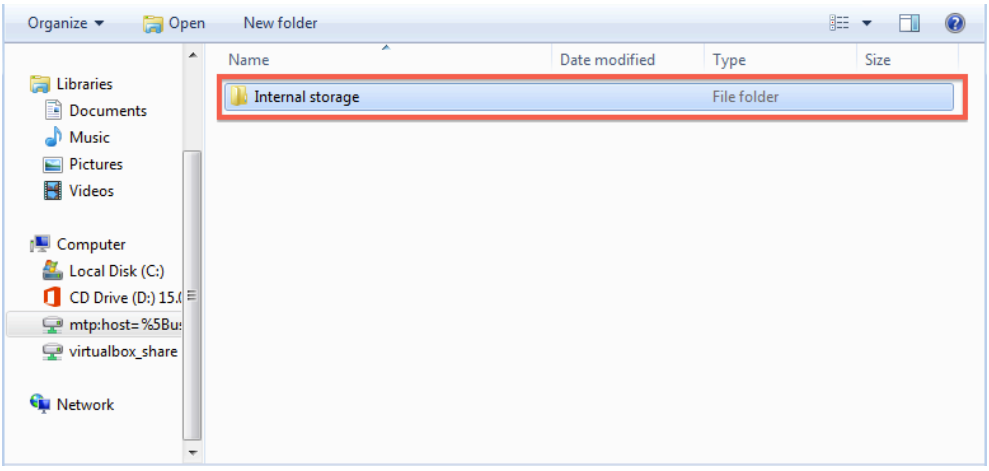
6767.xls" on your computer.



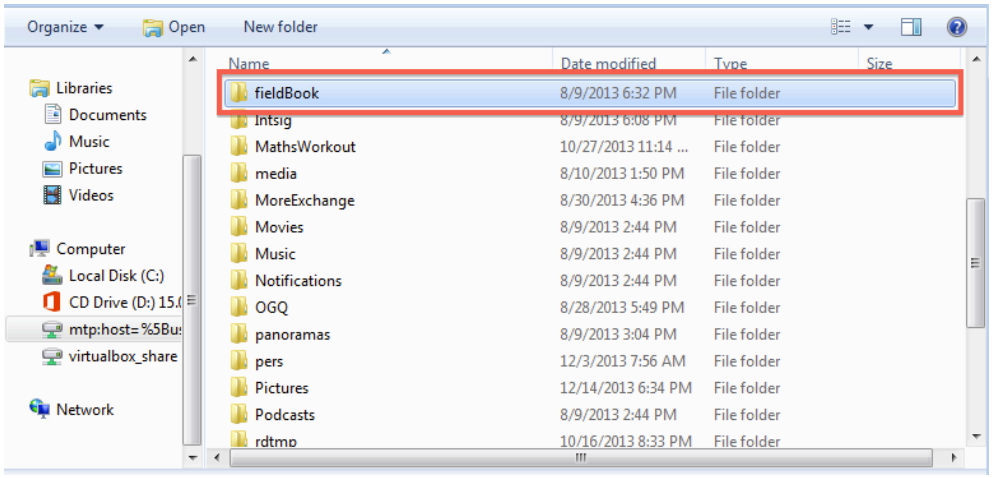
The files can be transferred to Android tablet by copying the files into the tablet's Internal Storage File.

12.4.2 Files on your Android tablet

To transfer Field Layout file and Trait file to your Android tablet, connecting an Android tablet to your computer, then clicking on tablet icon on your computer. Clicking on the tablet icon will open a window showing an "Internal Storage" file.



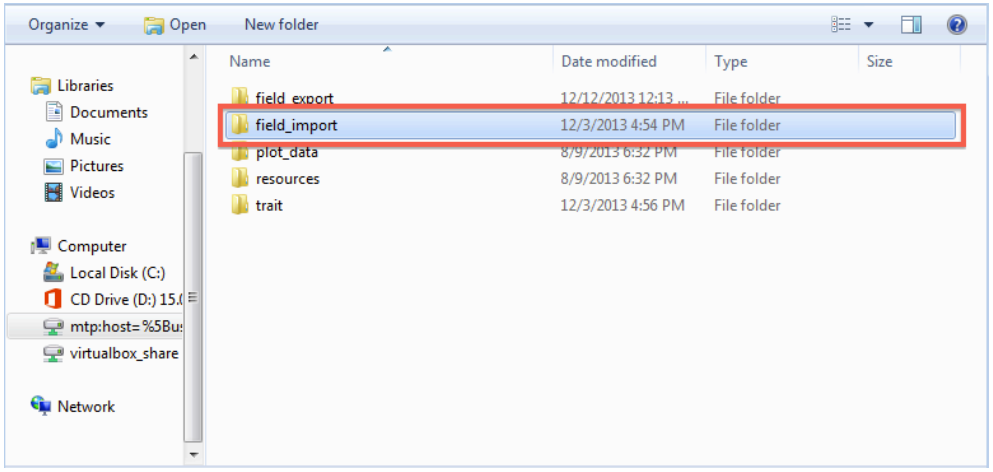
After you installed the Android Field Book App, all files for the app are stored in the "fieldBook" folder within the "Internal storage" folder.



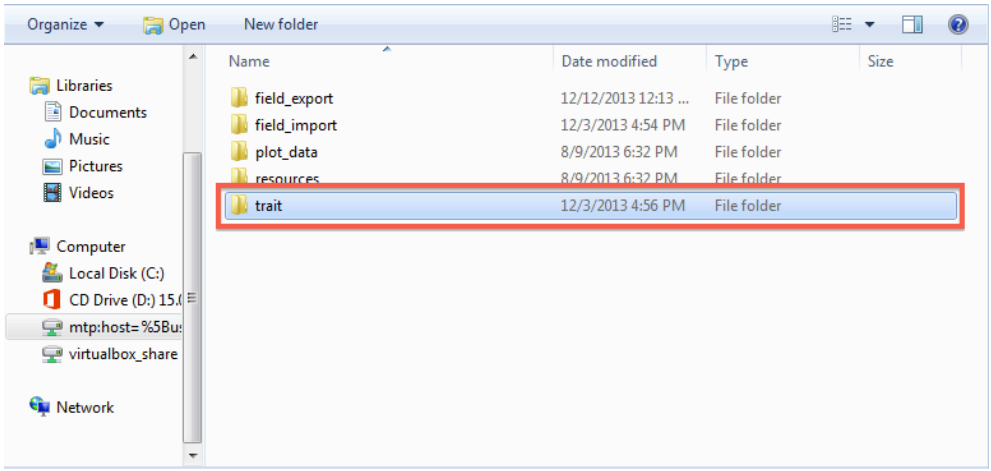
Within the "fieldBook" folder, there are five sub-folders:

- field_export
- field_import
- plot_data
- resources
- trait

Field Layout files must be copied into the "field_import" folder.



Trait files must be copied into the "trait" folder.



You can either drag and drop, or copy the Field Layout file and the Trait file from your computer to the folders in your Android tablet.

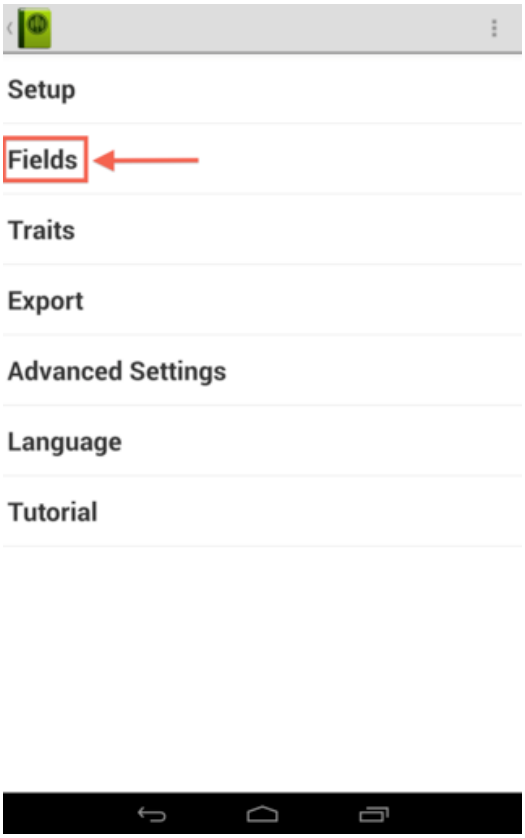
12.5 Setting up "Field Book App" for data collection

After you transferred the Field Layout file and Trait file from your computer to Android tablet, you still need to set up "Field Book App" on your tablet for data collection.

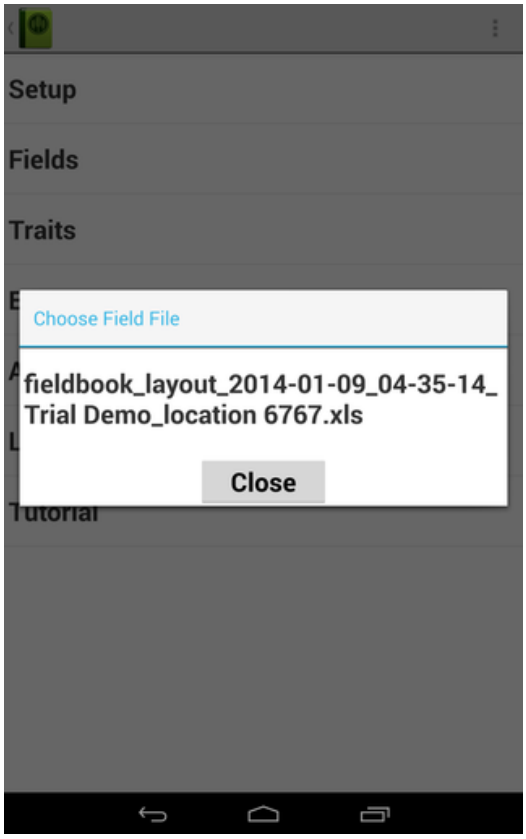
To set up the Field Book App:

1. **To open the Field Book App in the Android Tablet, clicking on the Field Book App icon, which is a green rectangle.**

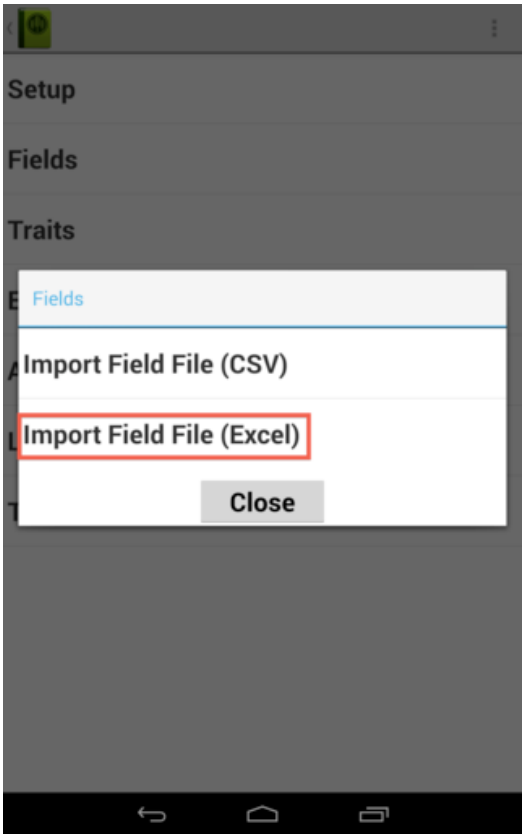
2. **To import Field Layout files, clicking on the "Fields" section of the main menu of the Field Book App.**



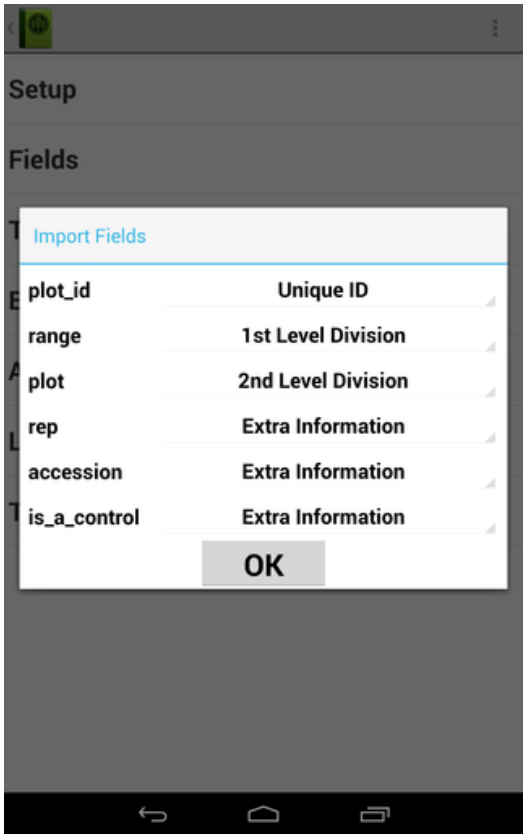
Clicking on the “Fields” tab will open a new dialogue that will let you select the file that you want to import.



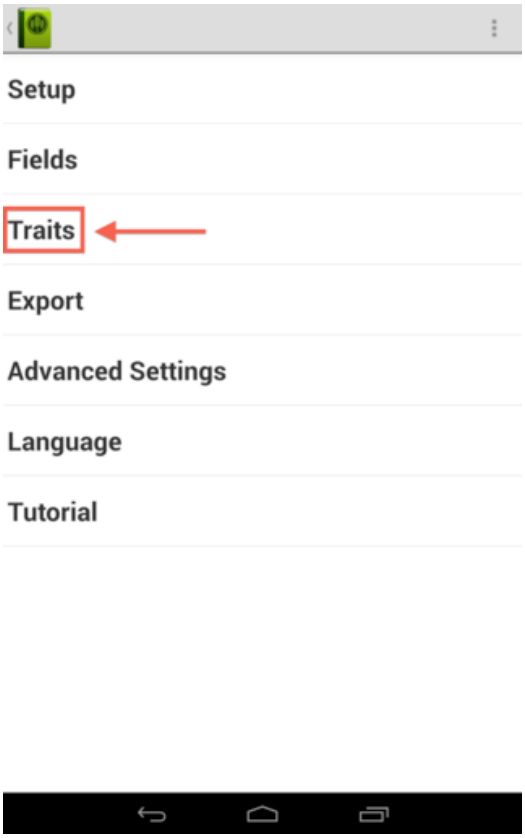
Choosing a Field File will generate a new dialogue that will ask you to choose between an Excel or CSV format. Since the data from the database is in Excel format, choose the Excel option.



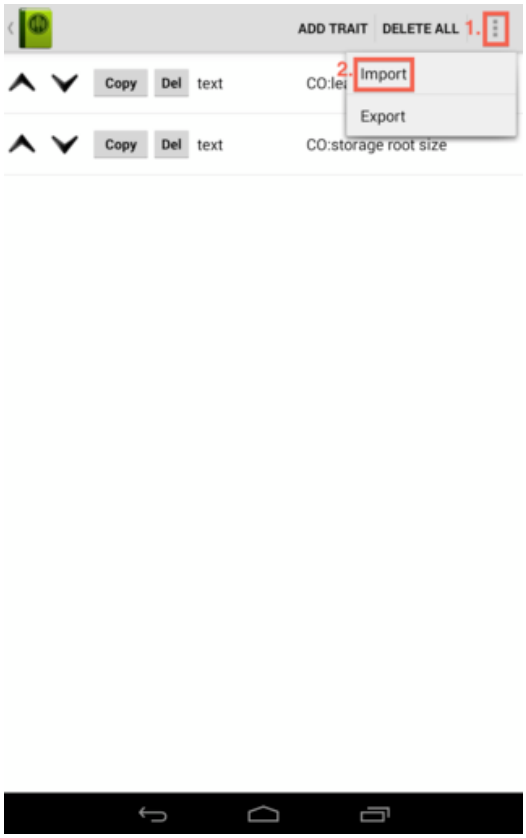
After submitting the file format, a final dialogue box will appear. Please provide information about the file that you want to import. Please ensure that "plot_name" is set as the unique identifier. To finalize the process, clicking "OK" button.



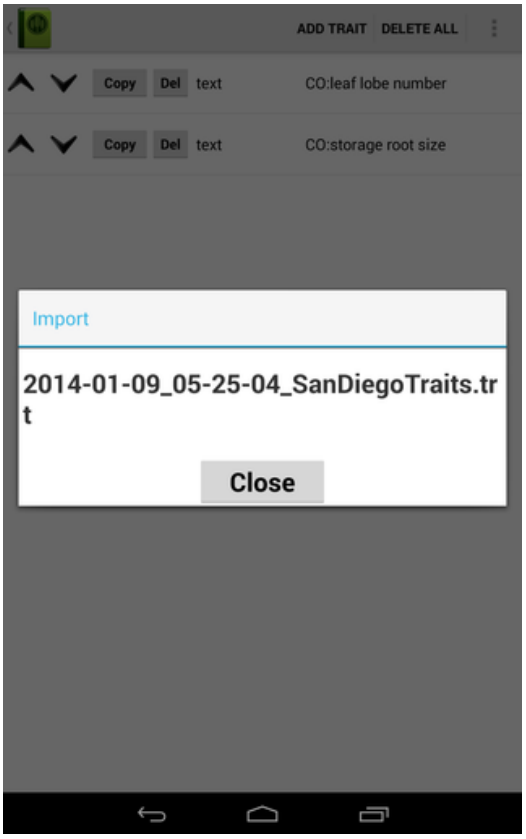
3. **To import Trait Files, clicking on the "Traits" tab on the main menu of the Field Book App.**



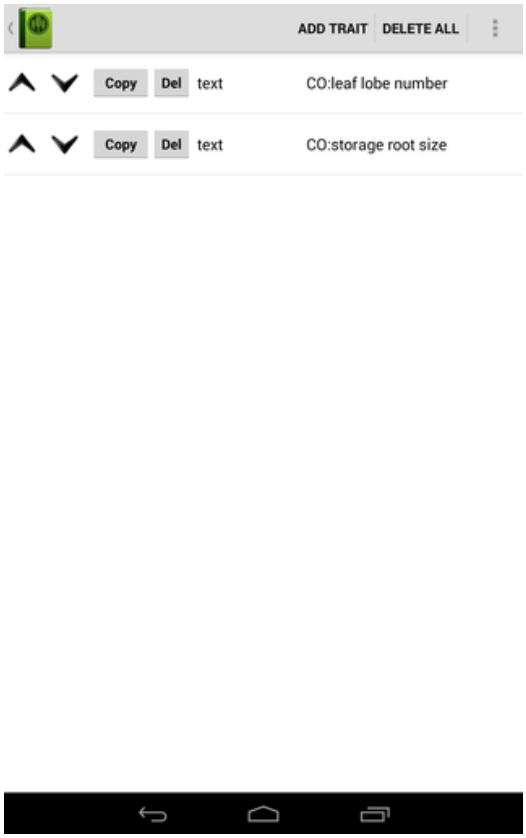
Then, clicking on the three dots symbol found on the upper right corner of the Field Book screen. This will open a drop down menu with the choices “Import” and “Export”. Clicking on “Import”



Clicking on "import" will open a new dialogue that displays a list of trait files that you can select to import to the Field Book App.



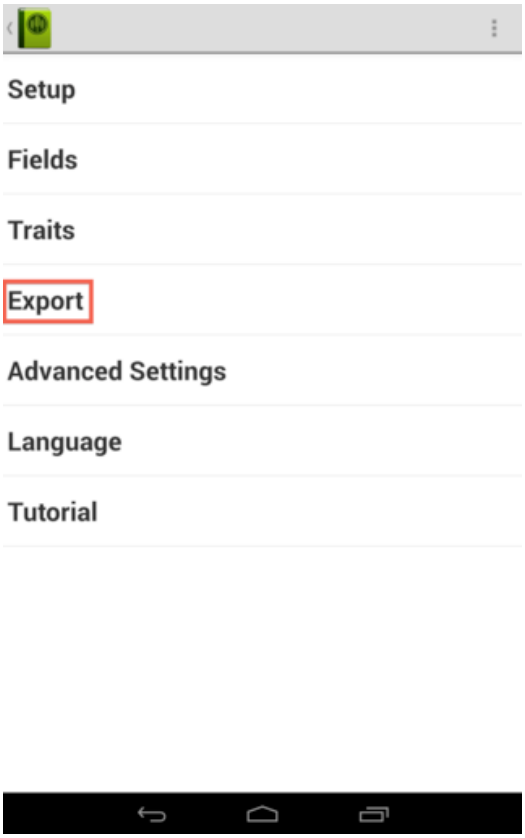
The trait file is now imported into the Field Book App. The traits page will show all trait files and available traits.



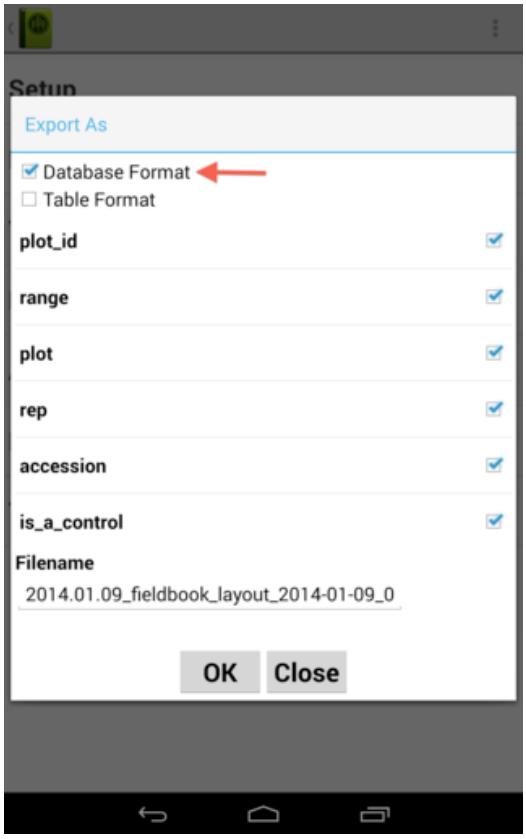
12.6 Exporting Files from Field Book App

Data that were collected on the Field Book App can be exported back to your tablet folder, which can then be transferred to your computer.

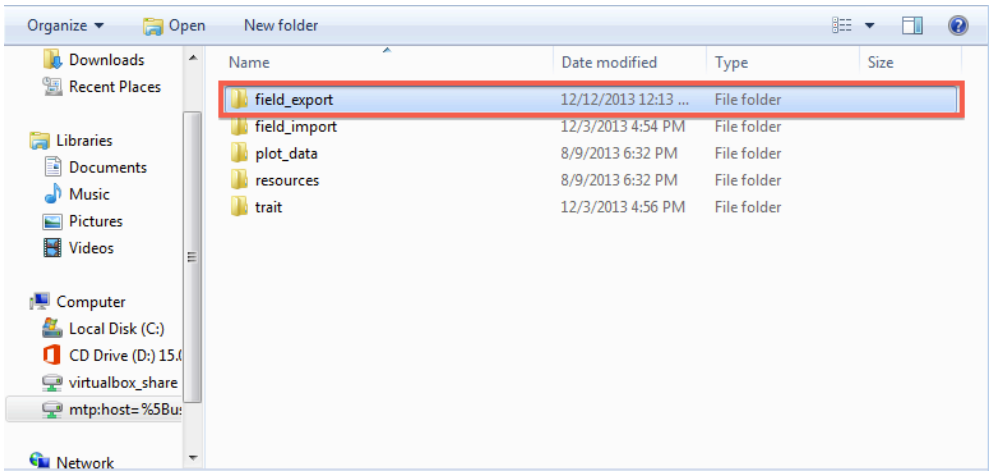
To export files containing data from the Field Book App to your tablet, clicking on the “Export” link on the main menu page of the Field Book App.

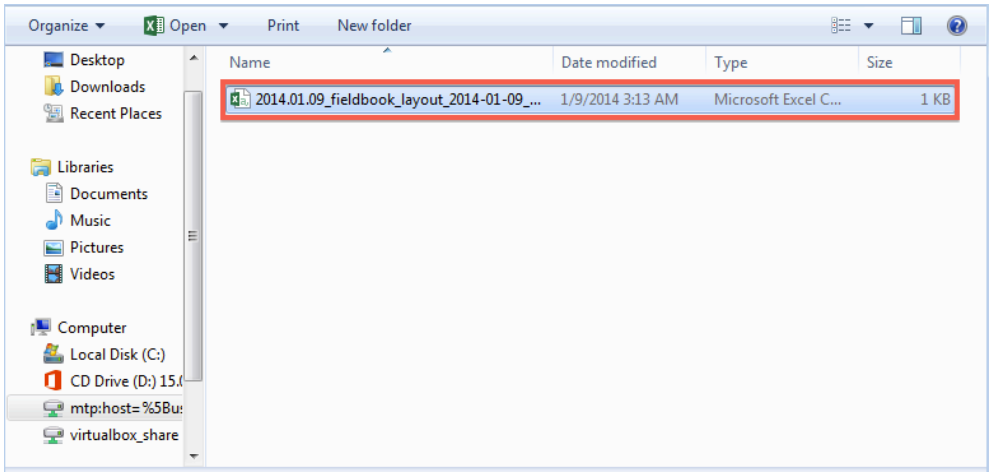


Clicking on the “Export” link will open a new dialogue window. To ensure that data are exported in a correct format for the database, checking the “Database Format” box, then clicking on “OK” button.



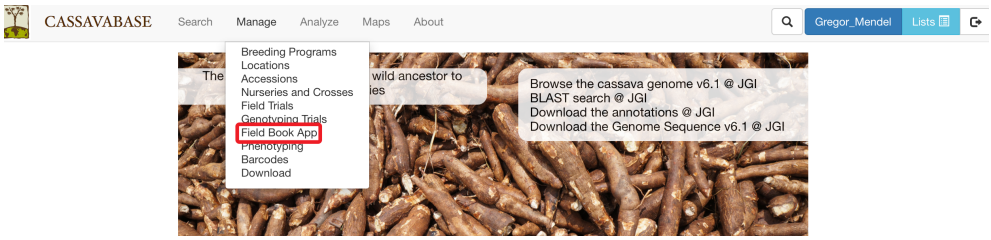
The exported file can then be found in the "field_export" sub-folder within the "fieldBook" folder on your tablet. Once you connect your tablet to your computer, you can directly transfer the file to your computer.



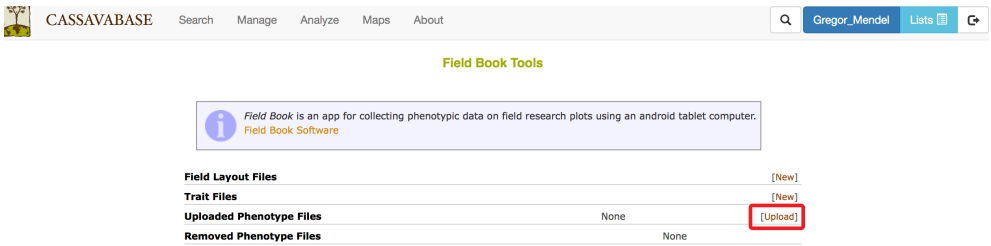


12.7 Uploading Phenotype Files to an SGN database

To upload phenotype files to the database, clicking on "Field Book App" in the "Manage" menu.

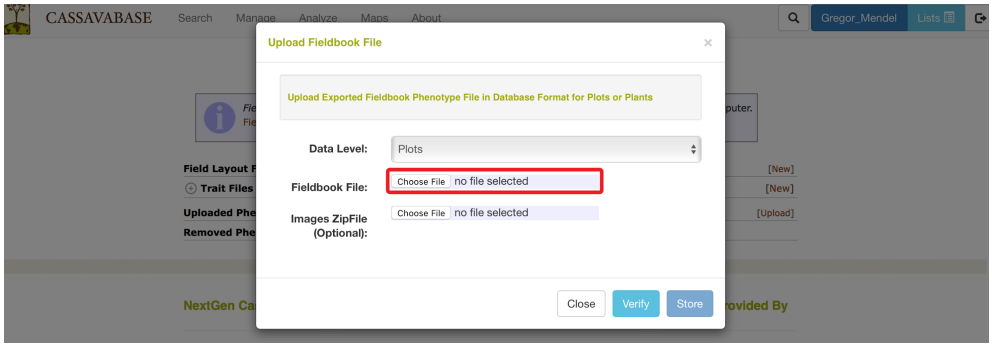


On the "Field Book Tools" page, clicking on "Upload" link in the "Uploaded Phenotype Files" section.

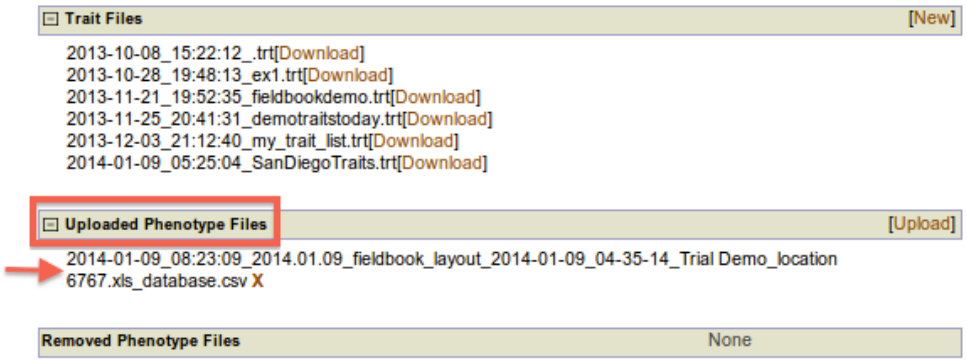


Clicking on the "Upload" link will open a new dialogue asking you to choose

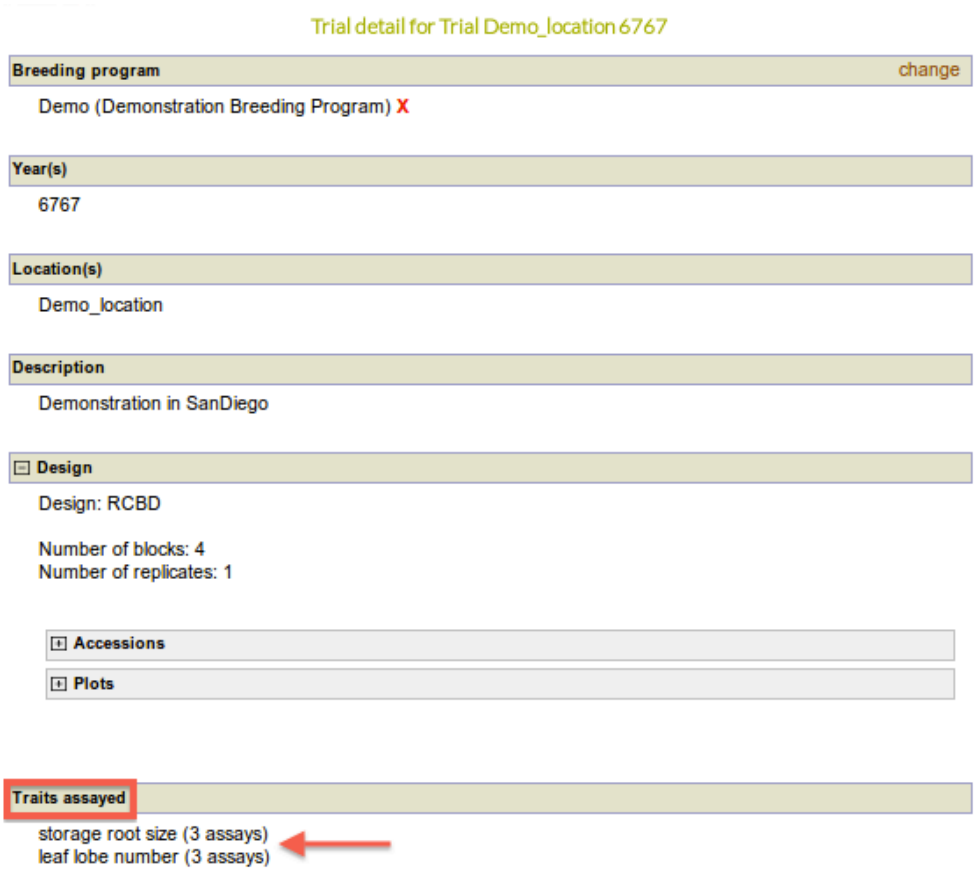
a file that you want to upload to the database website. Please ensure that "plot_name" is the first column of the file to be uploaded. To make sure that the file has the correct format for uploading, click on the "Verify" button. After the file format has been verified, click on the "Store" button.



The list of uploaded phenotype files can be found on the Field Book Tools page



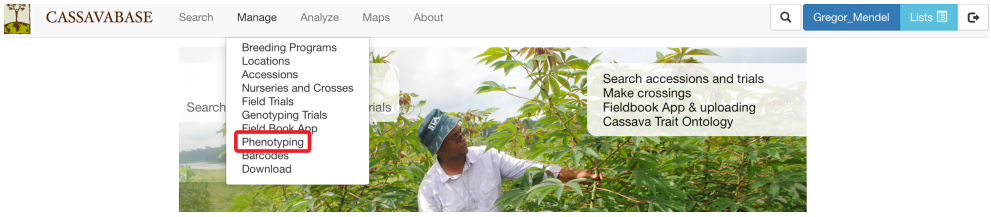
The uploaded files will also be seen in the corresponding "Trial Detail" page.


Trial detail for Trial Demo_location 6767
Breeding program
change
Demo (Demonstration Breeding Program) X
Year(s)
6767
Location(s)
Demo_location
Description
Demonstration in SanDiego
Design
Design: RCBD
Number of blocks: 4
Number of replicates: 1
Accessions
Plots
Traits assayed
storage root size (3 assays)
leaf lobe number (3 assays)

Chapter 13

Managing Phenotypic Data

To facilitate uploading process for phenotypic data, “Manage Phenotypic Data” page provides two options for uploading: Field Book Phenotype file in database format and phenotype file in Excel (.xls or .xlsx) file format. To access “Manage Phenotypic Data” page, clicking on “Phenotyping” in the “Manage” menu.



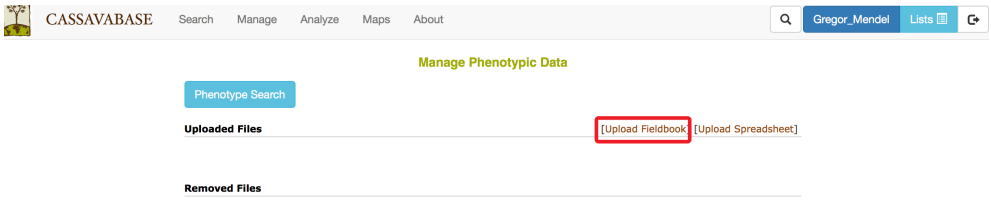
13.1 Uploading Fieldbook Phenotypes

13.1.1 Export Field Book Database File

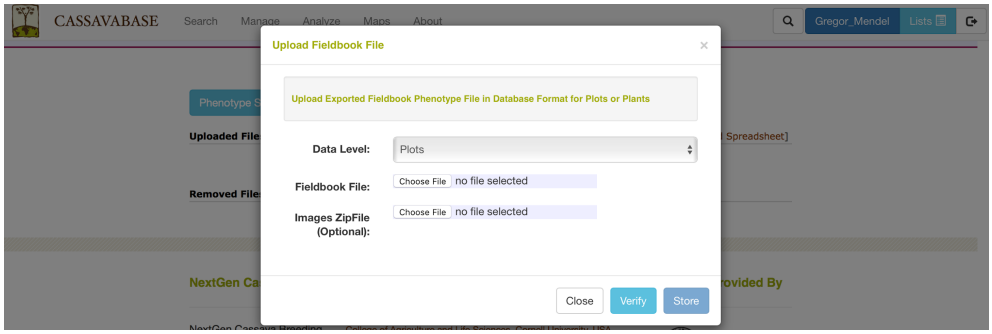
The database upload of Field Book phenotype data relies on the “Database” format from the Field Book. Please make sure to export the “Database” format from the Field Book if you intend to upload the data using the Field Book Upload we describe below. If you prefer to use the “Table” format that the Field Book exports, you can modify this format to work with the Speadsheet Upload we describe below.

13.1.2 Upload Field Book Database File

To upload a Field Book Phenotype file in a database format, click the "Upload Fieldbook" link

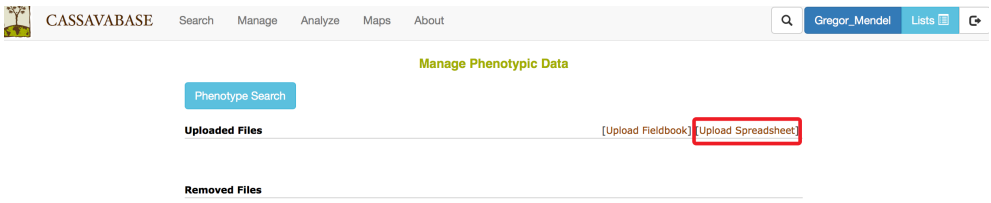


The "Upload Fieldbook" link on this page and "Upload" link on the "Field Book Tools" page open the same dialogue. Please follow instructions for uploading phenotypic files on the 12 page.

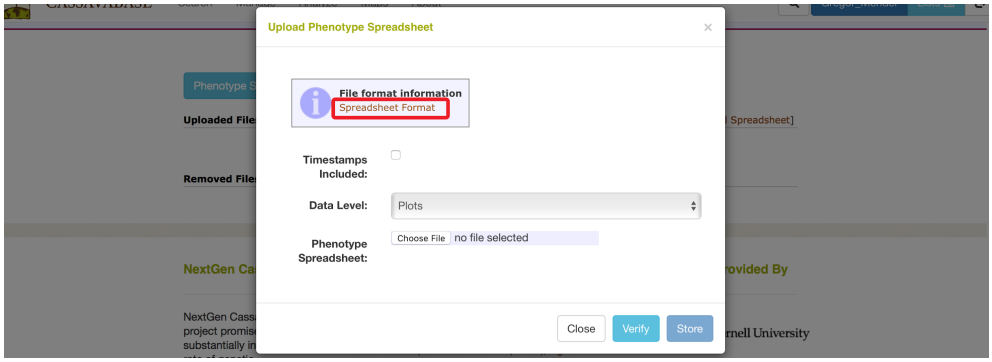


13.2 Uploading Spreadsheet Phenotypes

To upload a phenotype file in an Excel (.xls or .xlsx) file format, click the "Upload Spreadsheet" link.

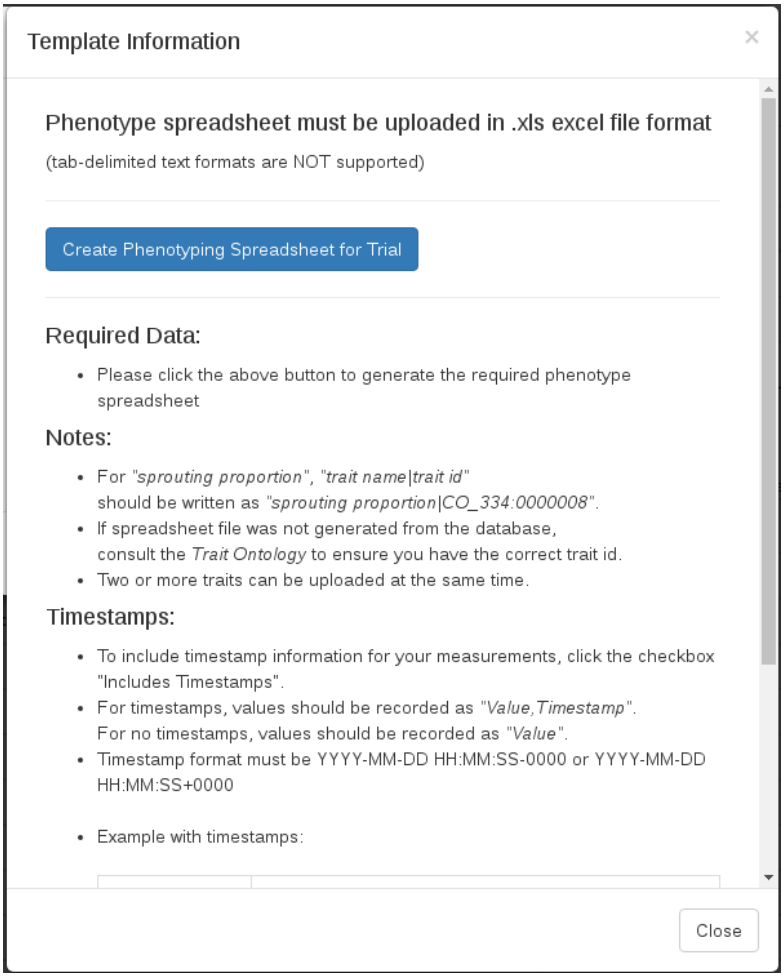


Please specify "Data Level" (Plots or Plants) and select the Excel file that you want to upload.

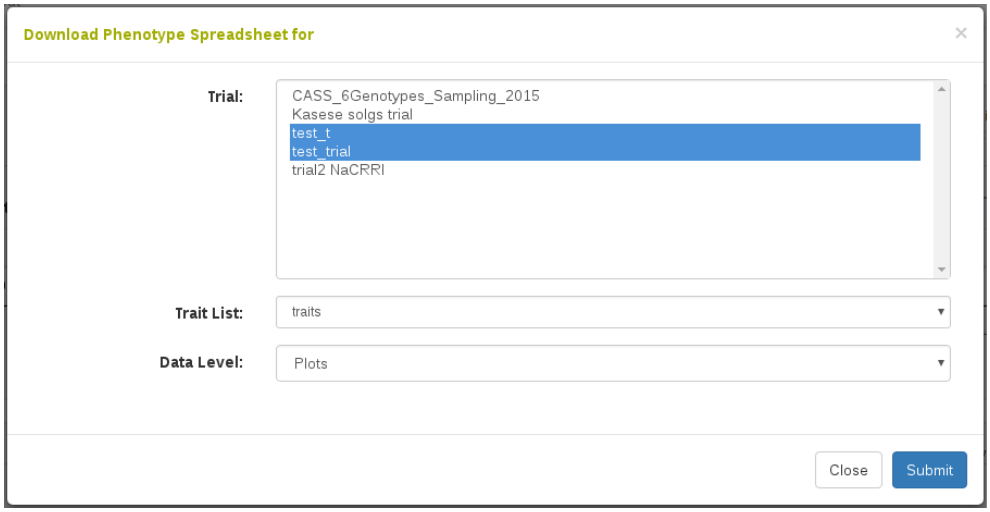


13.2.1 Generating Spreadsheet File

You can find more file format information by clicking on “Spreadsheet Format” link. Clicking on “Spreadsheet Format” will open the following dialog.

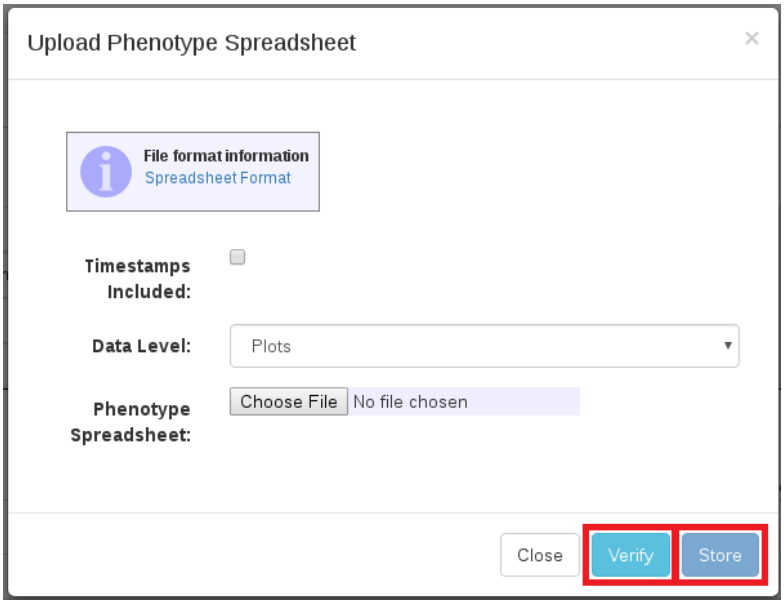


Clicking on "Create Phenotyping Spreadsheet" will bring up a dialog where you can indicate the trial(s) you are interested in and the trait list you are interested in. Clicking "Submit" will download the xlsx file onto your computer, where you can then fill in the phenotypes.



13.2.2 Uploading Spreadsheet File

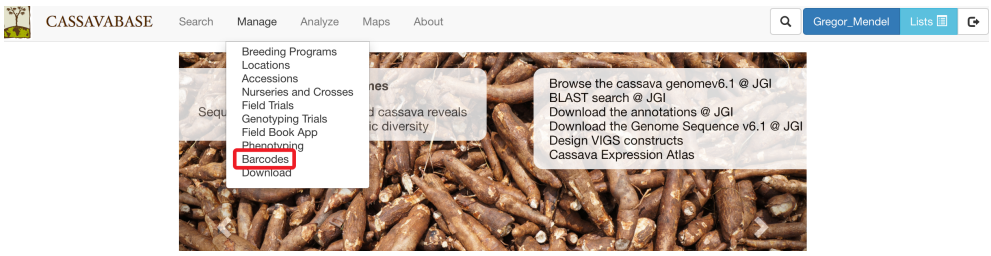
To ensure that the file has a correct format for uploading, click on the "Verify" button. This will check the contents of the file and also perform quality checks on the values in the file. These checks include checking the trait definition for categorical values, minimum and maximum values, and data type checking. It will also check if there are already values uploaded for the given observation units and traits. If there are, there is an option to overwrite the existing values with the new values in your file. If the file is valid, only then can you click "Store" to store the information in the database.


Upload Phenotype Spreadsheet
File format information
Spreadsheet Format
Timestamps Included:
Data Level:
Plots
Phenotype Spreadsheet:
Choose File
No file chosen
Close
Verify
Store

Chapter 14

Managing Barcodes

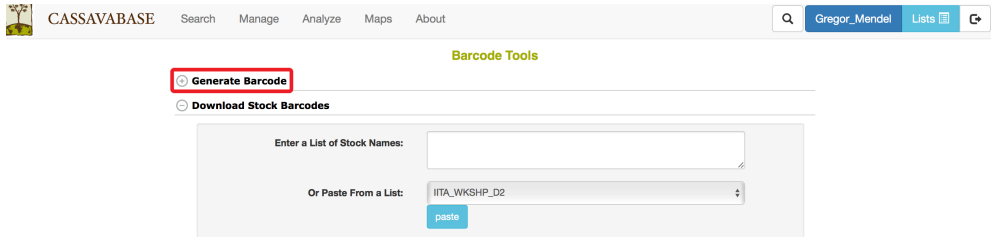
SGN databases provide tools for generating barcodes for stock identification. To access "Barcode Tools" page, clicking on "Barcodes" in the "Manage" menu.



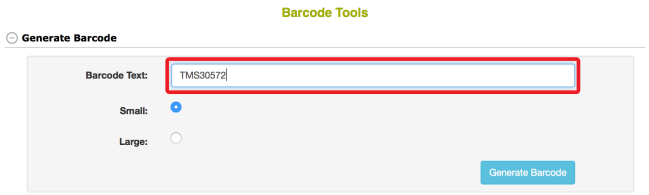
"Barcode Tools" page provides four options for generating barcodes:

- Single barcode
- Multiple barcodes
- Plot phenotyping barcodes
- Trial barcodes

To generate single barcode, clicking on "Generate Barcode" link on the "Barcode Tools" page.



In the "Generate Barcode" section, specify the name of the barcode, size of the barcode, then clicking on "Generate Barcode"



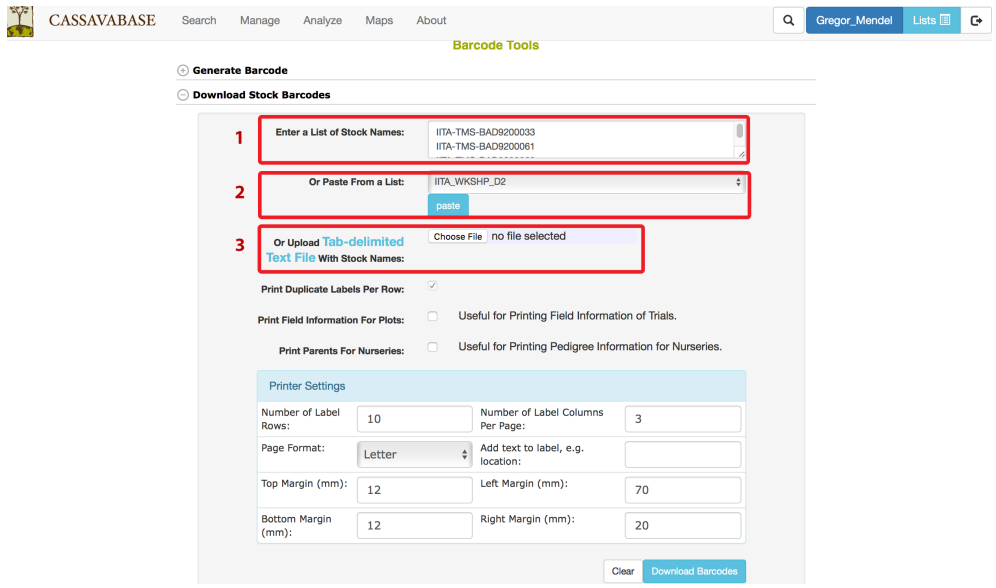
The database will generate a barcode for your stock. The barcode can be printed for your stock identification. It also appears on its corresponding stock page.

If you have a list of stocks that you want to generate barcodes, you can use "Download Stock Barcodes" section. You have three options for entering stock names:

1. Typing in stock names, or copy and paste from other file into the box (1)
2. Choosing a list of stocks from your "Lists" (2), and transferring the list into the box (1) by clicking on "paste" button.
3. Uploading a "Tab-delimited Text File" with stock names.
4. Select an optional printing format from the available formats.

You can select printer settings that you prefer in the "Printer Settings" sec-

tion. After you enter stock names and specify printer settings, clicking on "Download Barcodes" button at the bottom of the page.

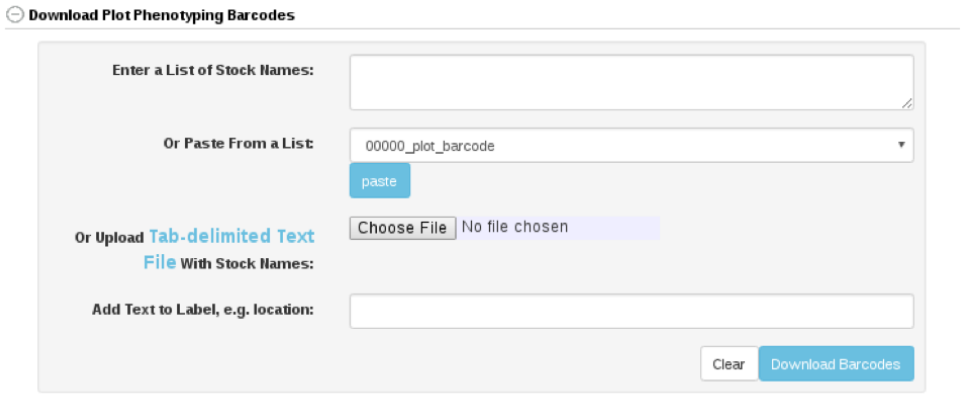


If you have a list of plots that you want to generate phenotyping barcodes, you can use "Download Plot Phenotyping Barcodes" section. You have three options for entering plot names:

1. Typing in plot names, or copy and paste from other file into the box (1)

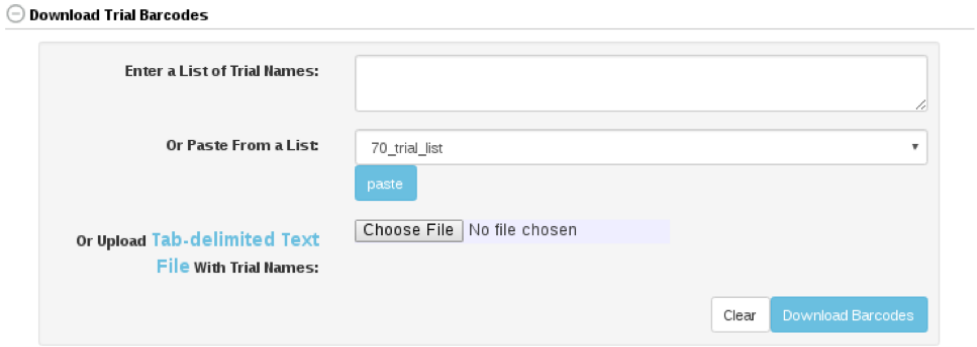
2. Choosing a list of plots from your "Lists" (2), and transferring the list into the box (1) by clicking on "paste" button.

3. Uploading a "Tab-delimited Text File" with plot names.



If you have a list of trials that you want to generate barcodes, you can use "Download Trial Barcodes" section. You have three options for entering trial names:

1. Typing in trial names, or copy and paste from other file into the box (1)
2. Choosing a list of trial from your "Lists" (2), and transferring the list into the box (1) by clicking on "paste" button.
3. Uploading a "Tab-delimited Text File" with trial names.



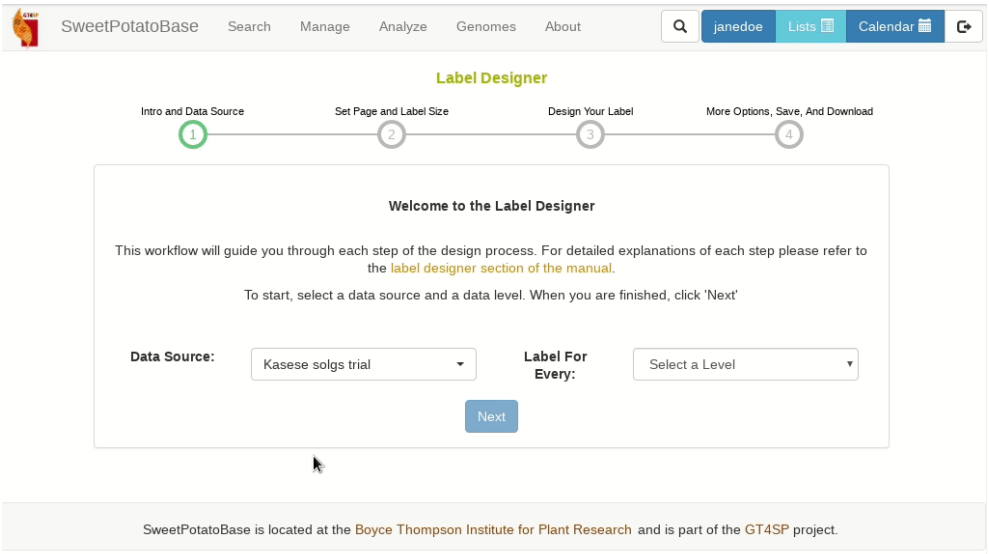
Chapter 15

Using the Label Designer

Breedbase provides an interactive design tool for creating custom labels. To access the Label Desginer, click on "Label Designer" in the "Manage" menu. The following sections explain your many options as you advance through each step of the design workflow.

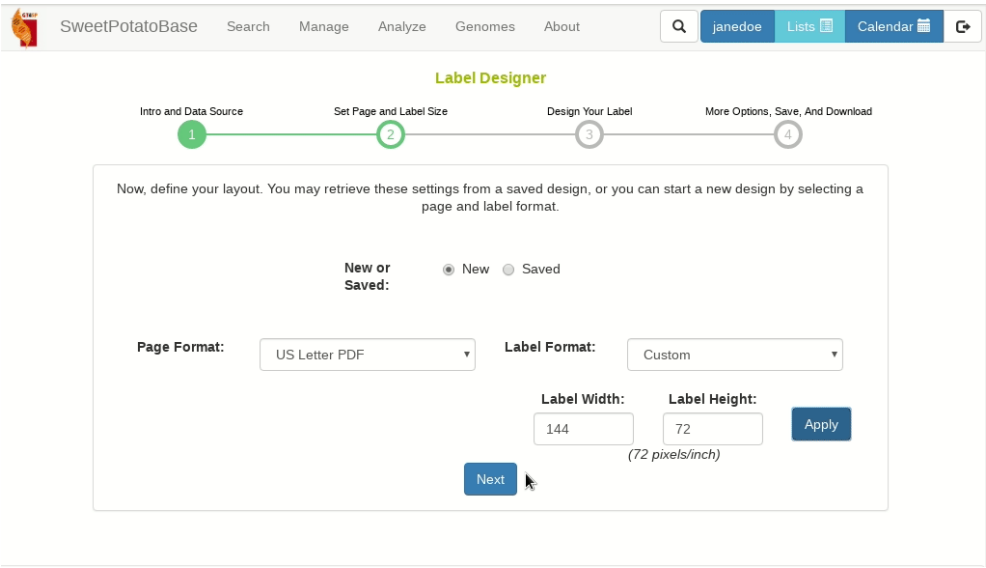
15.0.1 First Select a Datasource

The first step is to select a data source. Since the label designer can generate labels for different data types, you can optionally filter the source selection by the data type you're interested in. Then, select a field, genotyping, or crossing trial to populate your labels with the trial design information. Or select a list to populate your label with the list contents. For data sources with multiple levels of information you will also be asked to pick a level (plot, plant, etc.) before proceeding. To generate plot-level labels for more than one trial at once, select a list of trials as the source and plot as the level.



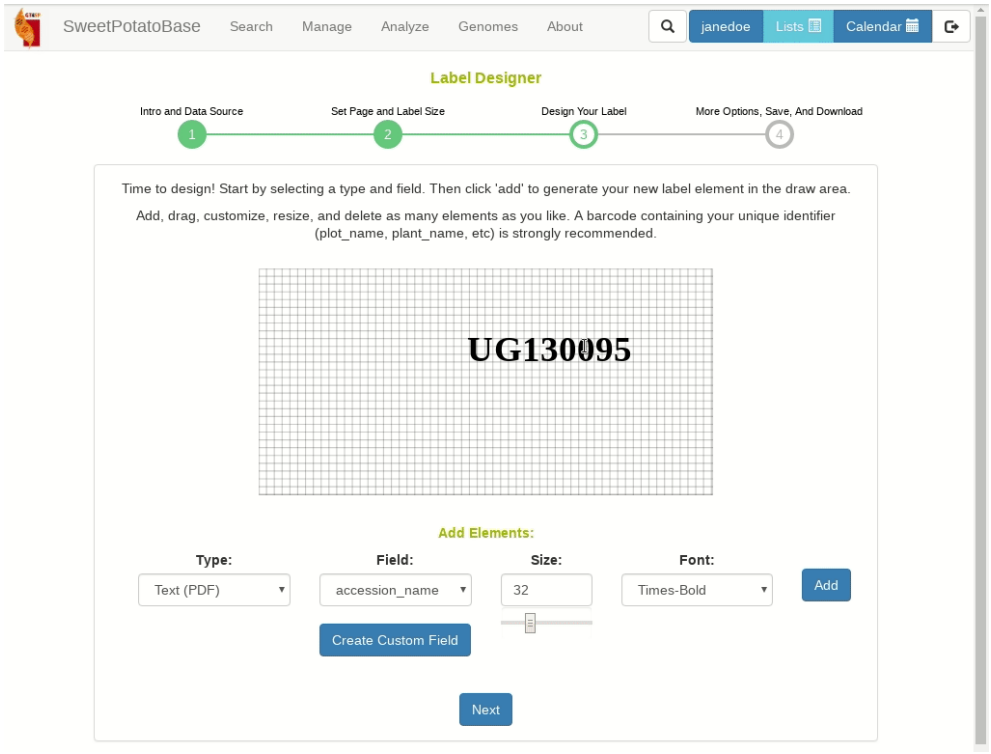
15.0.2 Set Page and Label Size

Now choose whether to create a new design or load a saved design. If you choose new, you will be prompted to select a page size and label size. If you do not see your page or label size as an option, then select Custom and enter your desired dimensions in pixels, or 1/72nds of an inch. If you choose saved, you will be prompted to select a saved design then will be taken directly to the design step with the saved design elements preloaded.



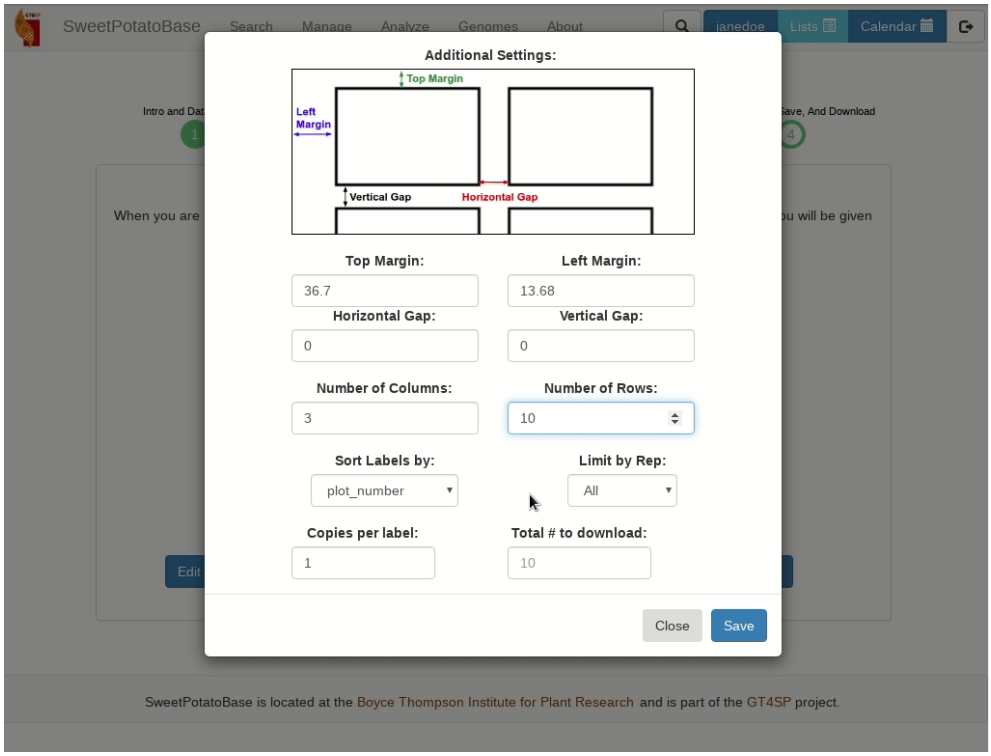
15.0.3 Design Your Label

Below is a draw area where you can begin adding elements to your label. First select a type, then field, size, and font, then click 'Add' You can add text to an exisiting field or create a completely custom field by clicking 'Create Custom Field' Once added, you can drag and drop elements, or delete them by clicking on the red box in their upper left corners. Barcodes can also be resized by dragging on the green box in their lower right corners. If you are creating labels for a trial it is highly recommended to include a barcode encoding your plot, plant, or tissue sample names. These are your unique identifiers that will need to included with any phenotypic or genotypic measurements loaded into the database. When you are satisfied with your design, click next!



15.0.4 Adjust Formatting, Save, and Download

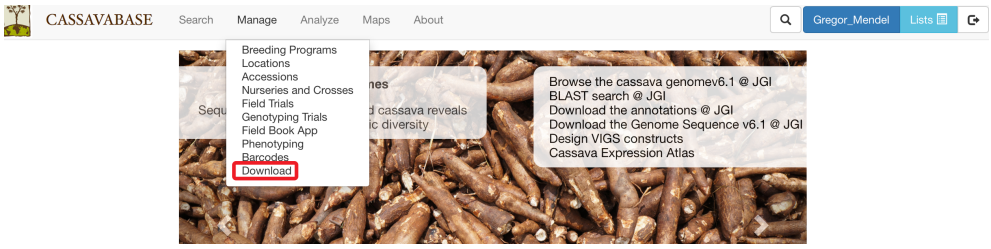
Last step! Here you can tweak your formatting and page layout, save your design, or download your labels. The additional settings dialog will allow you to fine tune the print margins and margins between labels. The units are pixels or 1/72nds of an inch. It's not recommended to change these until you've already done a test print. You can also set the # of copies per label, filter by rep, or download just the first page for test purposes. To save you're design just type a unique name and hit save. This will save your design to your list manager where you can set it to public to share it with others. Finally if you are ready just hit download to generate and download your labels!


SweetPotatoBase
Search
Manage
Analyze
Genomes
About
janedoe
Lists
Calendar
Additional Settings:
Top Margin
Left Margin
Vertical Gap
Horizontal Gap
Top Margin:
36.7
Left Margin:
13.68
Horizontal Gap:
0
Vertical Gap:
0
Number of Columns:
3
Number of Rows:
10
Sort Labels by:
plot_number
Limit by Rep:
All
Copies per label:
1
Total # to download:
10
Close
Save
When you are
u will be given
Edit
SweetPotatoBase is located at the Boyce Thompson Institute for Plant Research and is part of the GT4SP project.

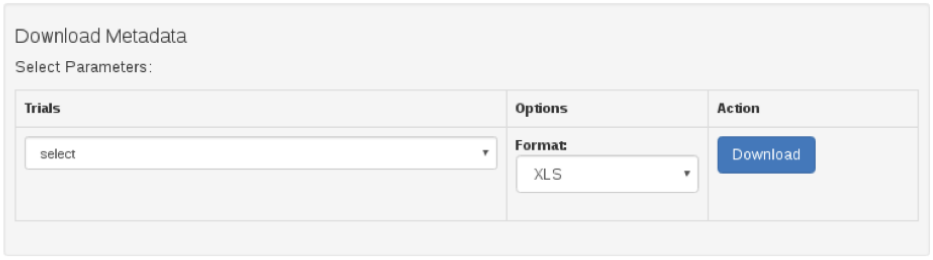
Chapter 16

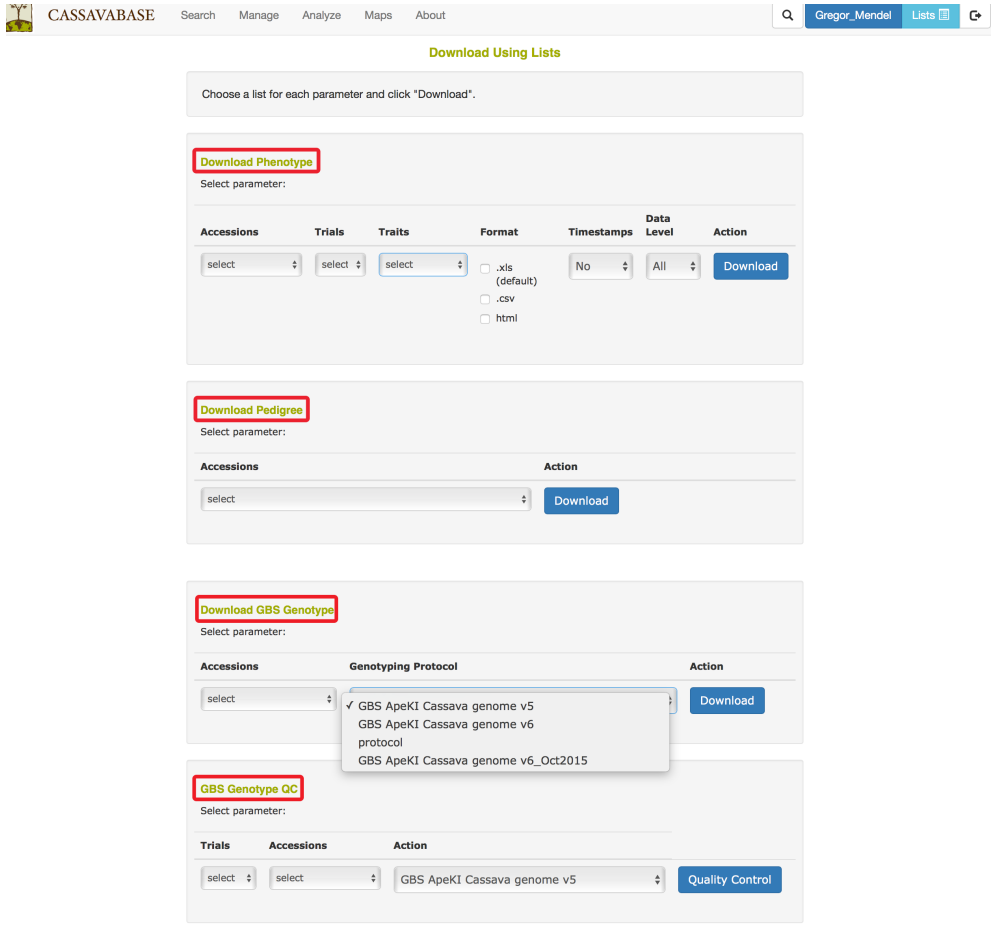
Managing Downloads

You can download phenotype, trial meta-data, pedigree, GBS genotype and GBS genotype QC files from the database to your computer by using "Lists". To download, clicking on "Download" in the "Manage" menu.



For each category, you can select a list of accessions from your "Lists" to download their phenotypes, pedigree, GBS genotype, GBS genotype QC. In the case of downloading trial meta-data, you would provide a list of trials, while for downloading phenotype and GBS genotype QC, you can also use a list of trials or traits.




CASSAVABASE
Search
Manage
Analyze
Maps
About
Gregor_Mendel
Lists
Download Using Lists
Choose a list for each parameter and click "Download".
Download Phenotype
Select parameter:
Accessions
Trials
Traits
Format
Timestamps
Data Level
Action
select
select
select
.xls (default)
.csv
html
No
All
Download
Download Pedigree
Select parameter:
Accessions
Action
select
Download
Download GBS Genotype
Select parameter:
Accessions
Genotyping Protocol
Action
select
GBS ApeKI Cassava genome v5
GBS ApeKI Cassava genome v6
protocol
GBS ApeKI Cassava genome v6_Oct2015
Download
GBS Genotype QC
Select parameter:
Trials
Accessions
Action
select
select
GBS ApeKI Cassava genome v5
Quality Control

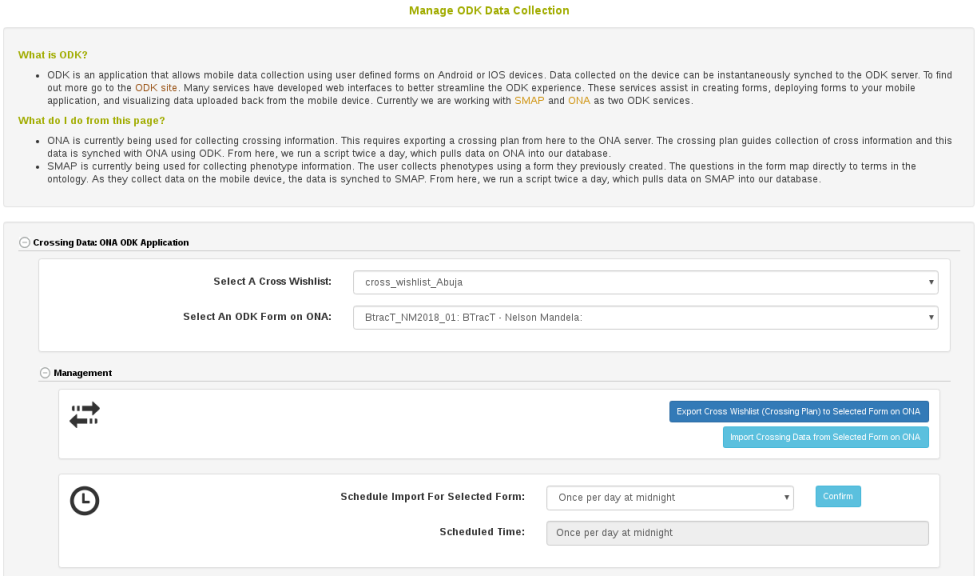
Chapter 17

Managing ODK Data Collection

To access this page go to Manage and then ODK Data Collection. ODK is used for remotely collecting data on Android and IOS devices. We currently are working to support two ODK service providers, namely ONA and SMAP. We are using ONA to collect crossing information, including all lab activities following seed production. We are using SMAP for phenotypic data collection.

17.1 ONA Crossing Information

17.1.1 Managing ONA Crossing Information



To begin collecting data using the ONA ODK form you must first have a crossing plan in the form of a Cross Wishlist. To do this from this page, click the "Export Cross Wishlist to ONA" button. Please refer to the "Create Cross Wihlist" help section for more information. It is possible to view the current available cross wishlists by clicking the "Export Cross Wishlist to ONA" button and then clicking "Available Cross Wishlists".

Once your cross wishlist is available, you can use your mobile ODK application to record crosses being done realtime. You can also record all laboratory activities following seed extraction up to greenhouse plantlet hardening.

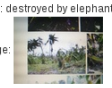
As you collect data using your mobile ODK application, your responses will be synchronized with our database. The "Schedule Import for Selected Form" section gives you options to perform the import daily or more frequently. It is also possible to initiate a data import from ONA at anytime by clicking "Import Crossing Data from Selected Form on ONA".

17.1.2 Reviewing Plant Status

The mobile ODK application has options to collect information about the status of plants in the field, such as if they are flowering. Images for each plant can also be recorded. The database will report this information here in a summary table that looks like the following. Notice that images are also transferred to the database.

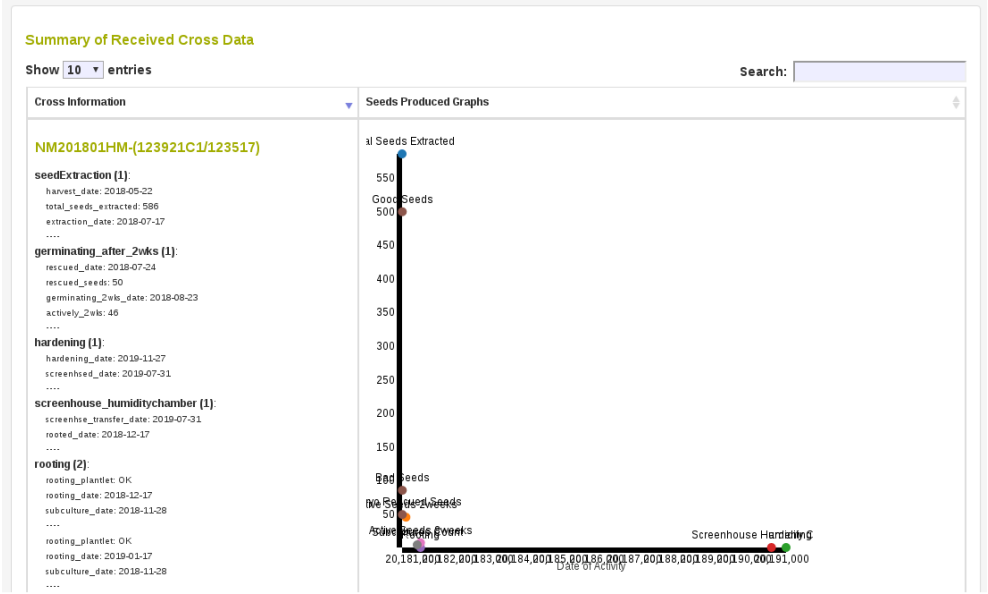
Summary of Received Plant Status

Show 10 entries Search:

Plot Name	Date	Status
	null	**Status:** Accession Name: null Trial Name: null User: null Status Location: Status: fallen Note: null Image: undefined
16-Huti-white_r8c12_plot157	2018-02-12	**Status:** Accession Name: Huti-white Trial Name: 2016 mchare polliantion block User: HM Status Location: in_field Status: destroyed Note: destroyed by elephants Image: 
16-ITC1460-Ijihulnkundu_r17c8_plot344	2018-01-18	**Flowering:** Accession Name: ITC1460-Ijihu Inkundu Plant Sex: female
16-ITC0712-AAcvRose_r1c1_plot1	2018-01-17	**Flowering:** Accession Name: ITC0712-AAcv Rose Plant Sex: male
16-ITC1468-Kahuti_r1c2_plot2	2018-01-17	**Flowering:** Accession Name: ITC1468-Kahuti Plant Sex: female

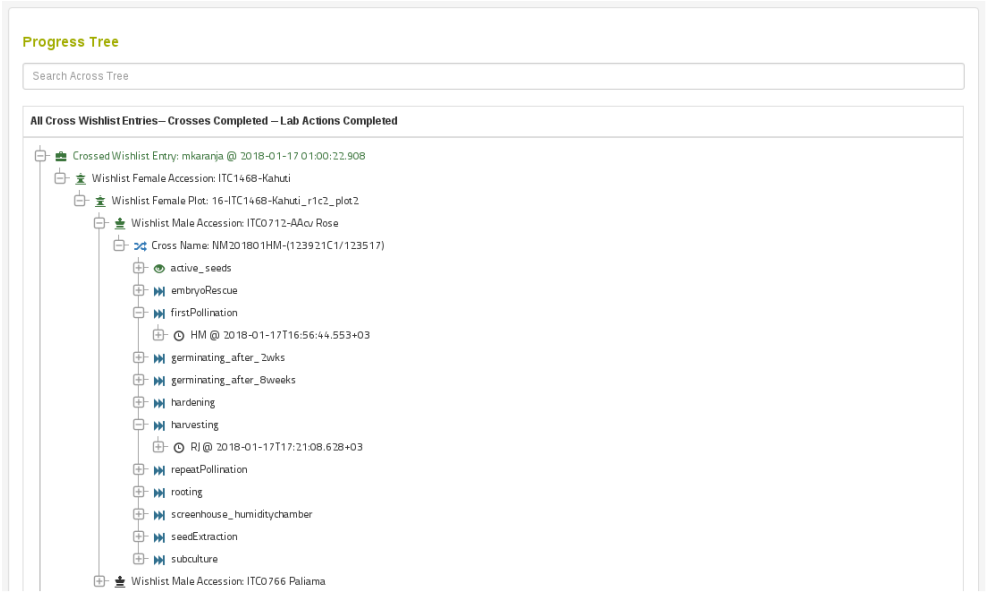
17.1.3 Graphical Summary For Performed Crosses

There is a section to summarize activities done for each cross. In this table each row represents a single cross performed. All the activities that have been performed will be shown here, such as “first pollination” and “embryo rescue”. The scatter plot shown tracks seed numbers generated on the Y axis and date of activity on the X axis.



17.1.4 Summary Information For Performed Crosses

There is a secondary section to summarize what has been done across the entire Cross Wishlist. This tree structure shows all activities performed for a cross and shows how these crosses relate to the Cross Wishlist.



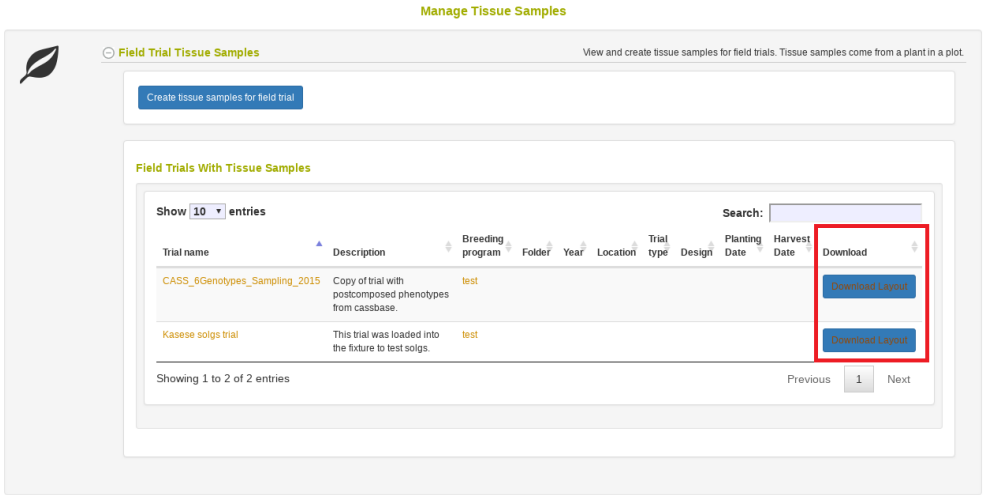
Chapter 18

Managing Tissue Samples

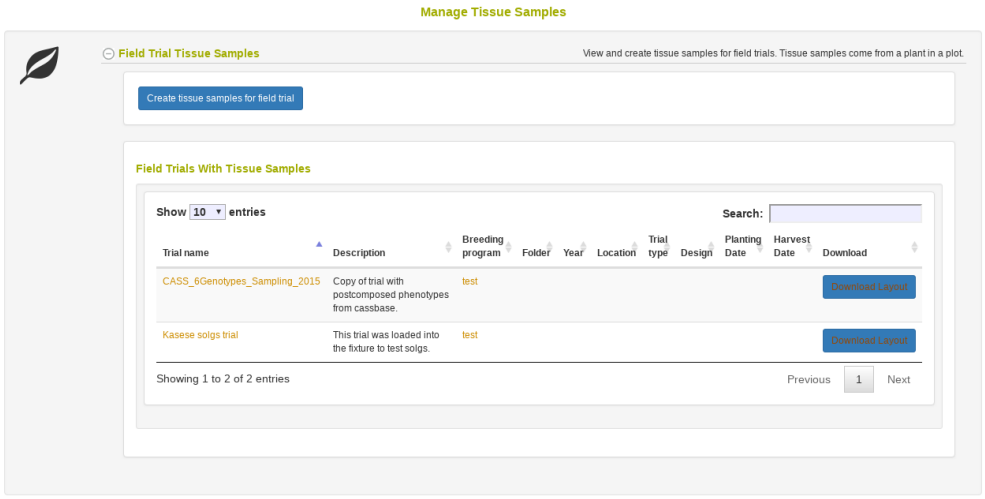
To access this page go to Manage and then Tissue Samples.

18.1 Tissue samples from field trials

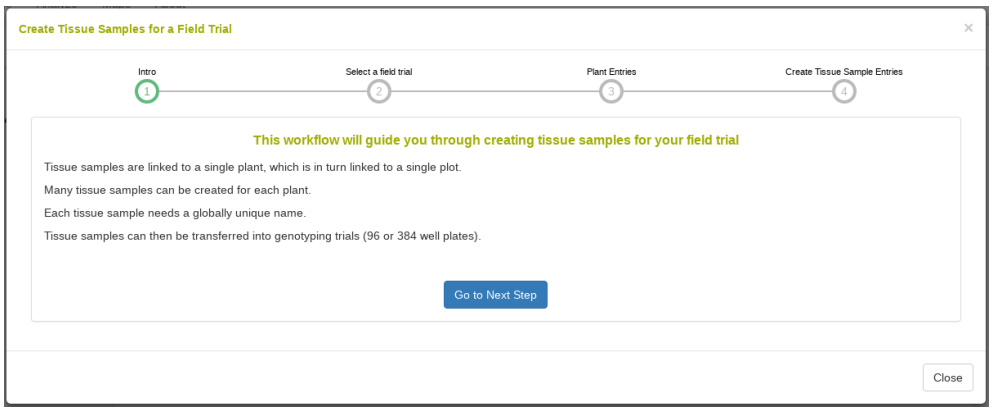
A field trial contains plots planted with a specific accession. Each plot can contain many plants, which in turn can contain many tissue samples. On the manage tissue sample page we can see the field trials that contain tissue samples already. We can choose to download the tissue sample layout as seen in the below picture.



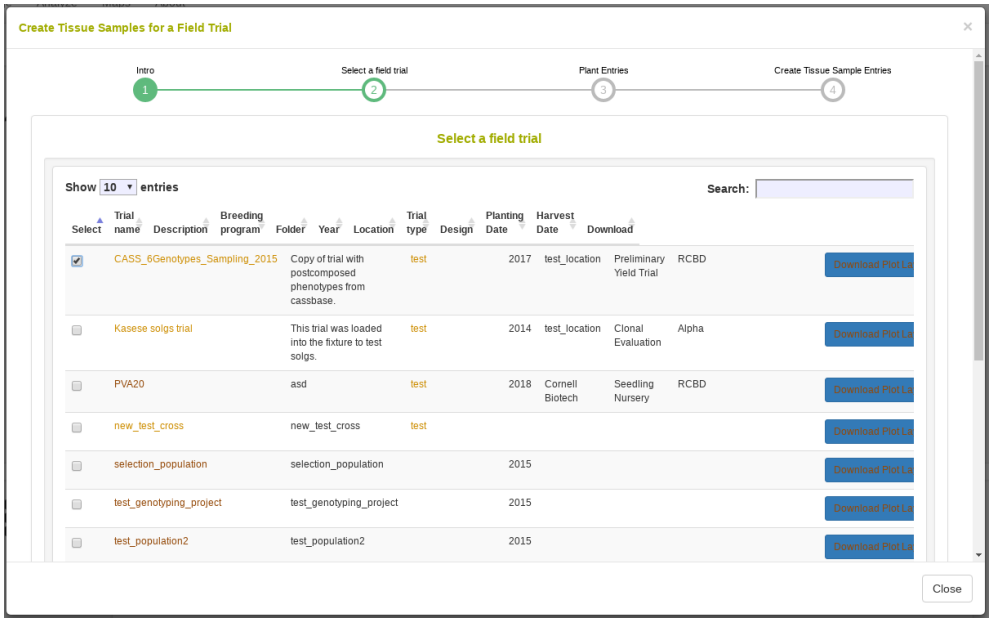
If the field trial you want to collect tissue samples from is not in the above table, you can click the button highlighted below.



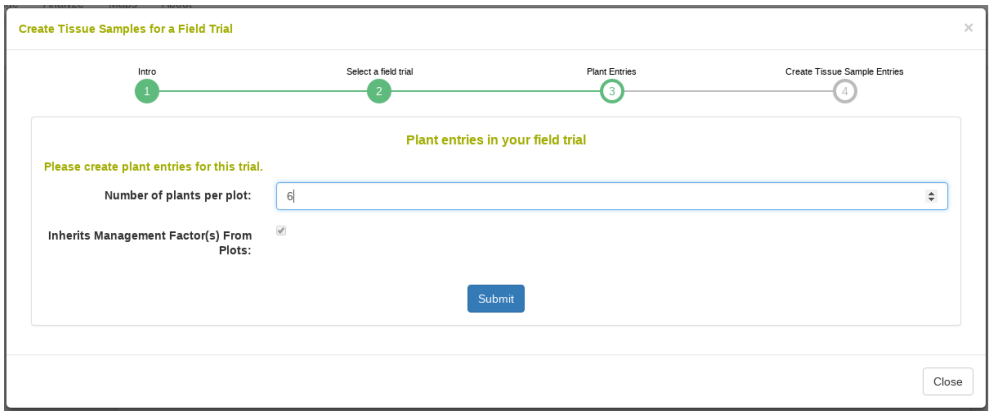
Once you have clicked this button, you will enter a workflow that begins with the following introduction.



Once you click next, you will need to select your trial.

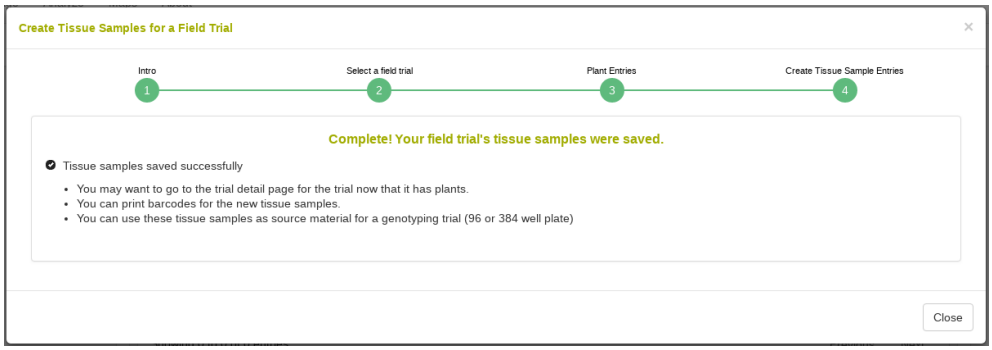


Next, if your trial currently only has plot entries saved, you will be asked to enter how many plants are in each plot.



Finally you will be asked how many tissue samples you want for each plant. You can specify a string to include in the tissue sample name, such as leaf or root.

Afterwards you should see the following success message, indicating that the tissue samples are saved.

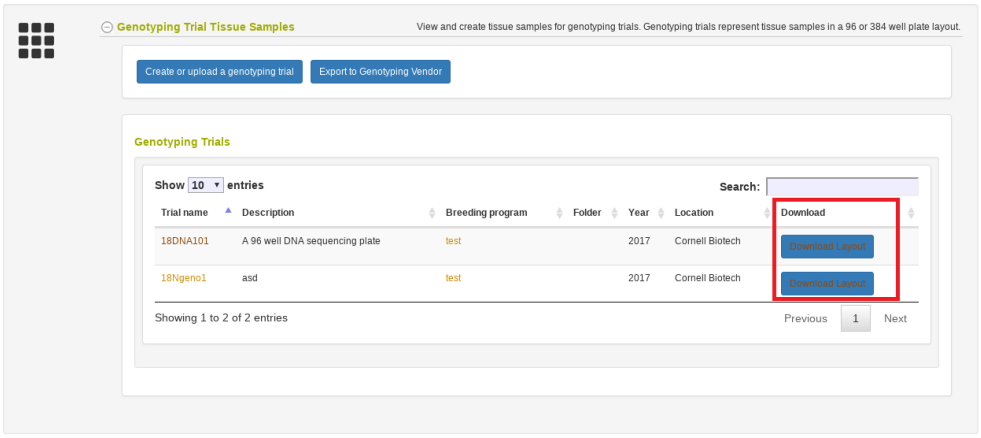


18.2 Genotyping Plate Tissue Samples (96 or 384 well plates)

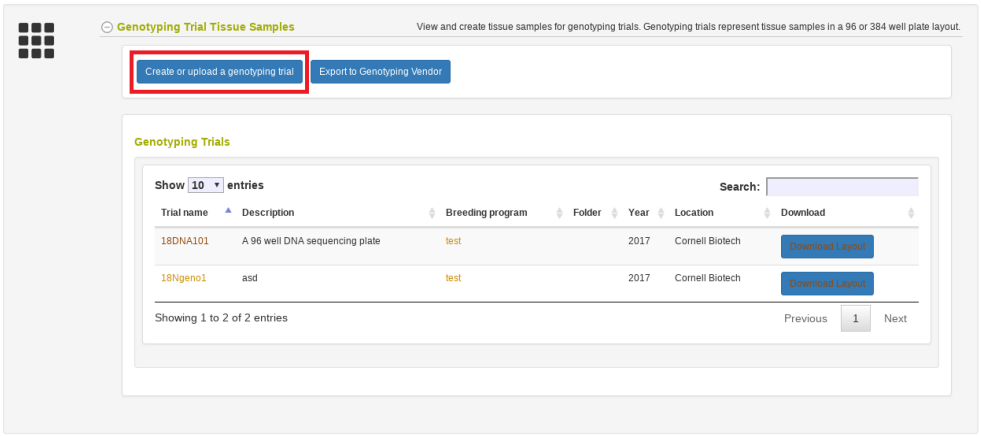
A genotyping plate represents a 96 or 384 well plate. You can use the Coordinate Android application to create your plate layout, or you can upload your own Excel plate layout, or you can use the database to generate a plate layout. Ideally, you will use tissue sample names originating from a field trial as the "source" for each well tissue sample, but you can also use plant names, plot names, or accession names.

From the manage tissue samples page, you can see the genotyping plates

saved in the database. You can also download the layouts as shown below.



If you need to create a new genotyping plate, you can click the button shown below. This will guide you through a workflow for uploading or creating the new plate layout.



Genotyping vendors require you to send a plate layout during submission. You can download the plate layout as shown above, or you can go to a genotyping plate detail page to download the Intertek formatted file.

In the future you will be able to directly export your genotyping plate plate layout to vendors.

Chapter 19

Managing Observation Variables

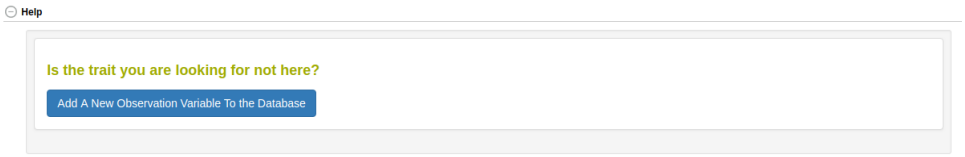
19.1 Managing Observation Variables with Traits, Methods, and Scales

Observation variables are the identifiers used when collecting phenotypic data. An observation variable is composed of a trait, a method, and a scale. The trait describes the attribute being measured e.g. 'Plant Height'. The method defines the protocol in which the trait was observed e.g. 'Using a one meter long measuring stick'. The scale defines the units or dimensions for which the measurement was taken e.g. 'Meters'.

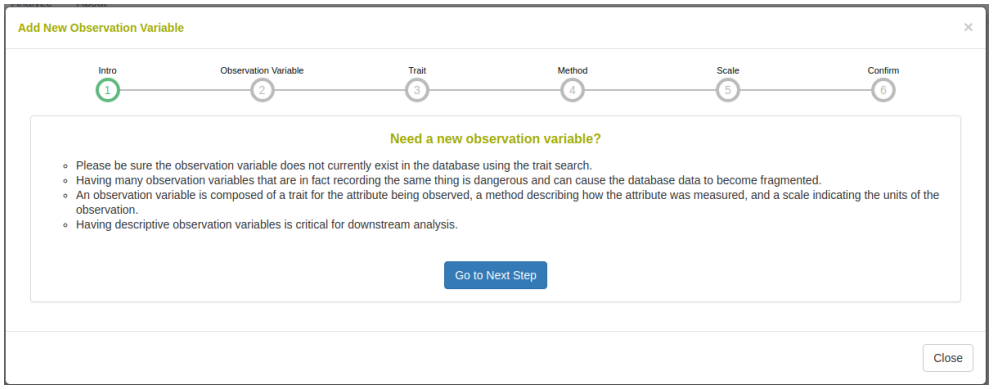
Generally, observation variables are defined in ontologies that are predefined. We often use ontologies from cropontology.org. In this case, you will not be able to define your own observation variables directly; instead, you will need to contact us and we will add the observation variable for you.

For databases where the user has greater control, we have an interface to allow addition of observation variables, along with traits, methods, and scales. To begin, go to the Search->Traits page.

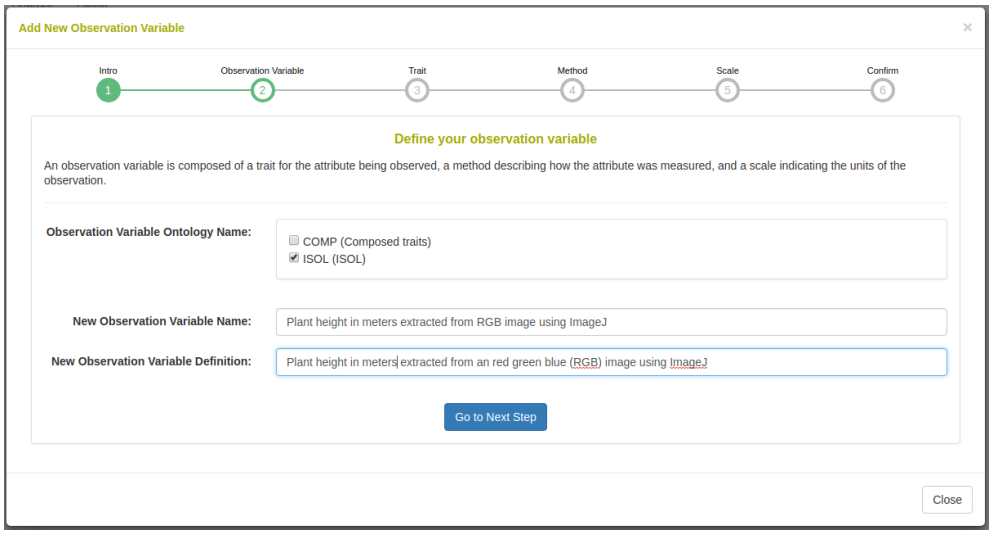
If the database you are on allows you to directly add observation variables, you will see the following button at the bottom of the page.



When you click the button, the following workflow will appear. You should be logged in or else it will not allow addition of the observation variable. The workflow begins with an introduction.

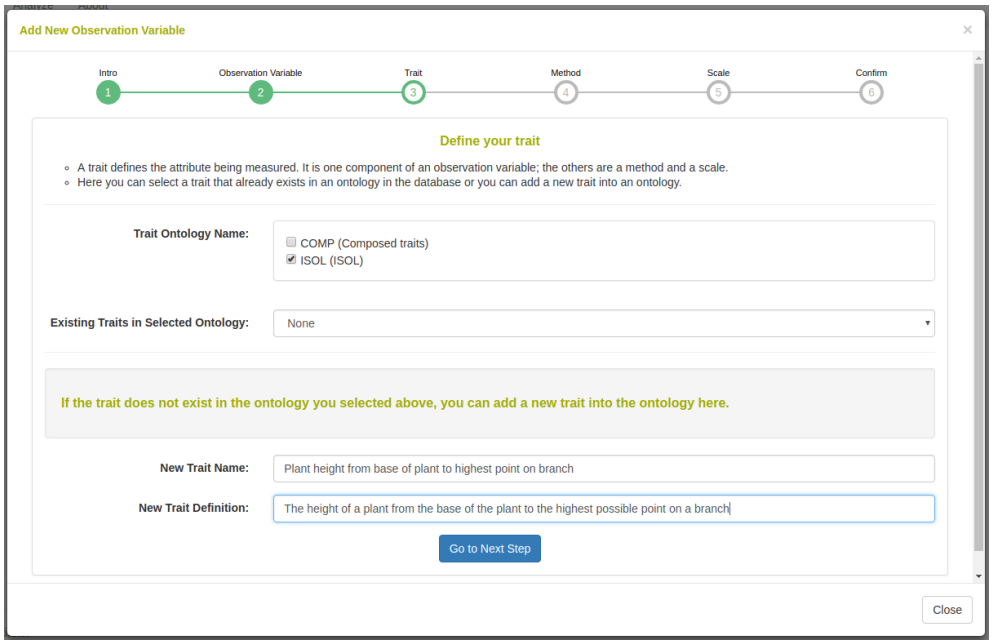


On the next workflow step, you select the ontology that you want to insert the new observation variable into. You must also give a name and a definition for the new observation variable.

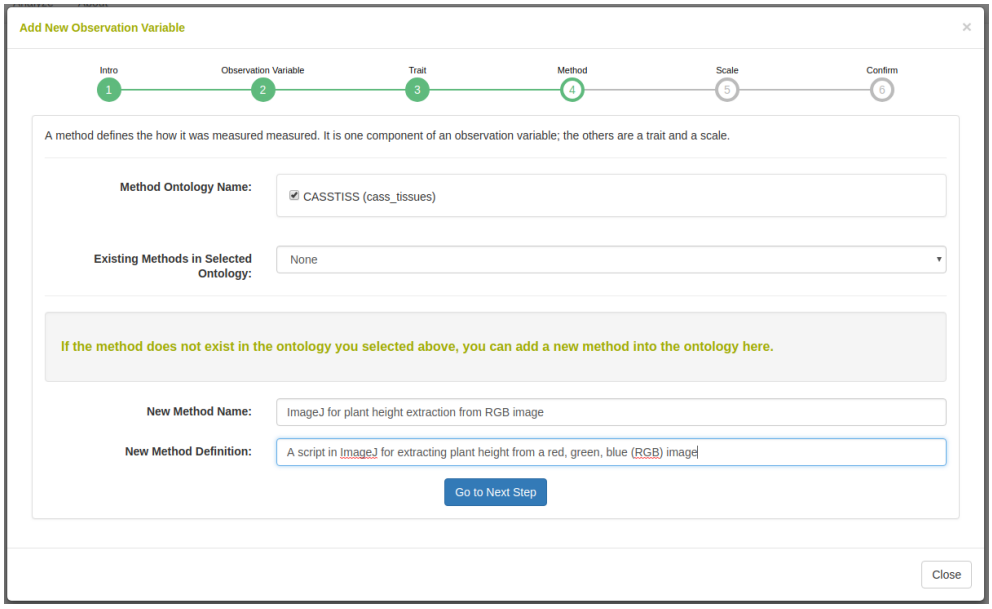


On the next workflow step, you select the trait ontology to use. Once you

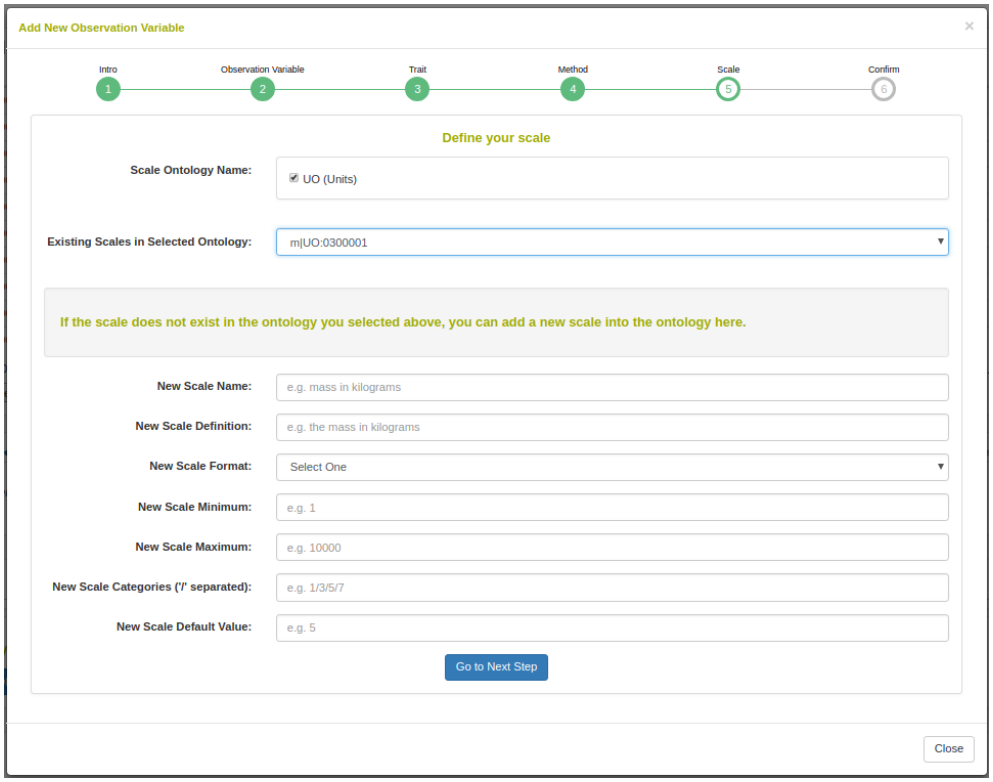
select a trait ontology, a select containing all the terms in the selected ontology will appear. You can either select a trait or if it does not exist in the select, you can create a new one by giving a name and a definition for the new trait.



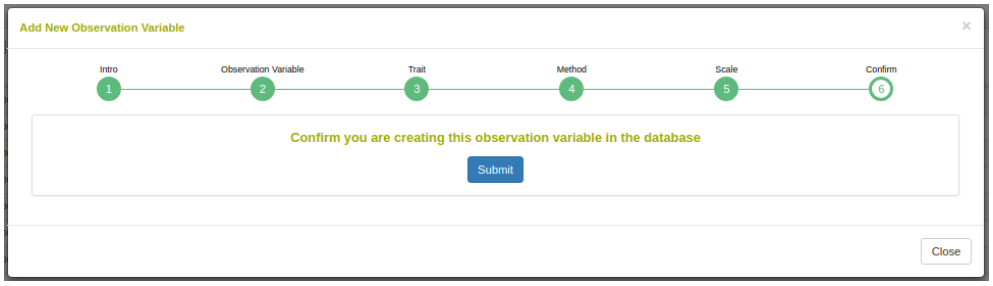
On the next workflow step, you select the method ontology to use. Once you select a method ontology, a select containing all the terms in the selected ontology will appear. You can either select a method or if it does not exist in the select, you can create a new one by giving a name and a definition for the new method.



On the next workflow step, you select the scale ontology to use. Once you select a scale ontology, a select containing all the terms in the selected ontology will appear. You can either select a scale or if it does not exist in the select, you can create a new one by giving a name and a definition for the new scale. You can also define a format, minimum, maximum, categories, and default value for the new scale.



On the last page of the workflow, you confirm the submission.



Afterwards, you can use the newly created observation variable ontology term in your phenotyping.

Chapter 20

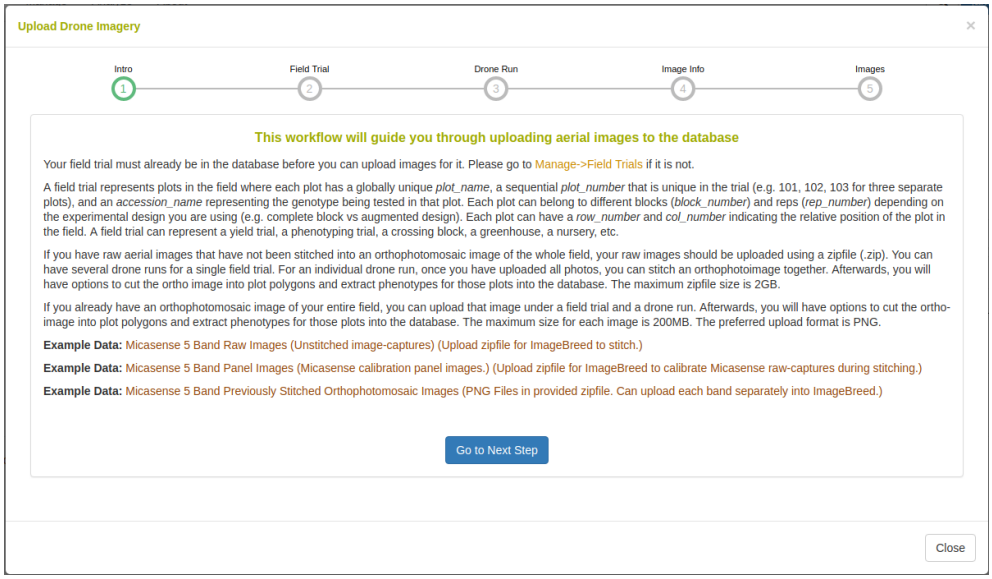
Managing Image Data

20.1 Image-Phenotyping Dashboard

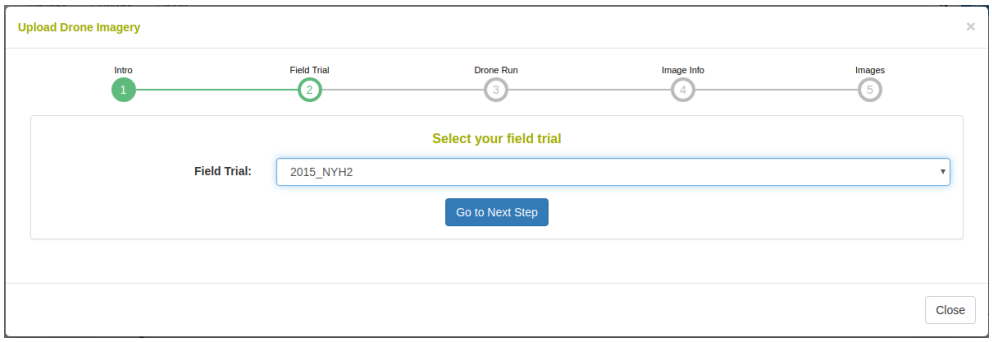
1. Upload raw image-captures in a compressed file (.zip) for orthophotomosaic assembly or upload previously stitched orthophotomosaic raster (.PNG, .JPG) imagery.
2. Dashboard shows all field trials and uploaded imaging events in collapsible sections.
3. Follow standard processes to manually create templates for assignment of plot-polygon images to the field experiment design.
4. All imagery is shown with the spectral category within collapsible sections. Figure shows NIR imagery.
5. Apply Fourier transform filtering, thresholding, and vegetation index masking. Plot-polygon images for all image processes are shown.
6. Extract and export phenotypic values from plot-polygon images for analyses and model training.

20.2 Image Input

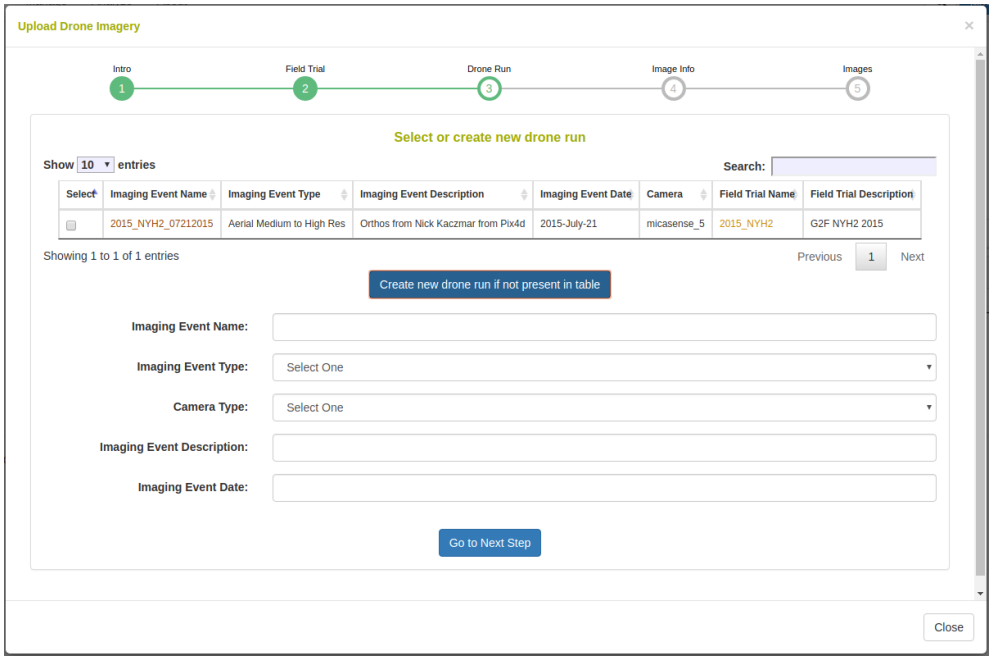
Clicking "Upload Imagery" will open the following dialog.



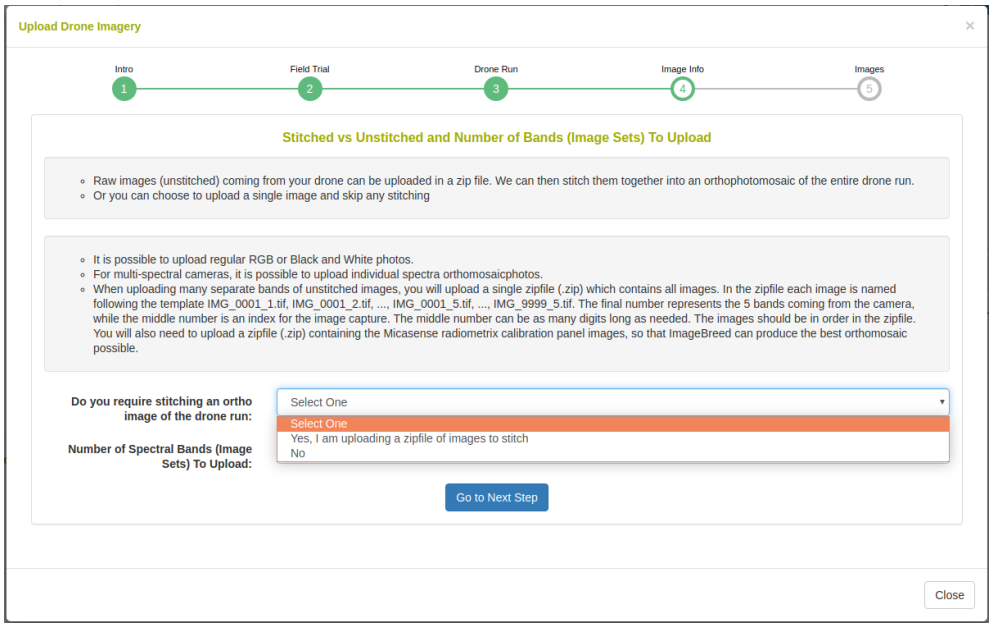
Raw-captures can be uploaded in a compressed (.zip) file so that they can be assembled into an orthophotomosaic. If orthophotomosaic assembly is not required, raster images (.PNG, .JPG) can be uploaded. Example data is given for raw Micasense RedEdge 5-band multispectral captures and for stitched orthophotomosaics.



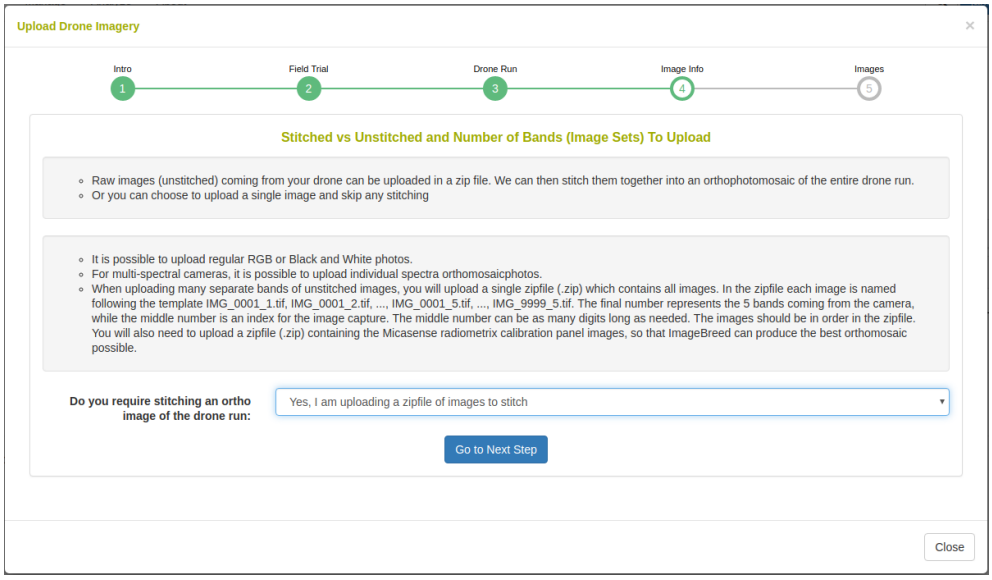
To begin uploading images, a field trial must be selected. The field trial must already be saved in the database. For information about adding a field trial, please read the Field Trial documentation.



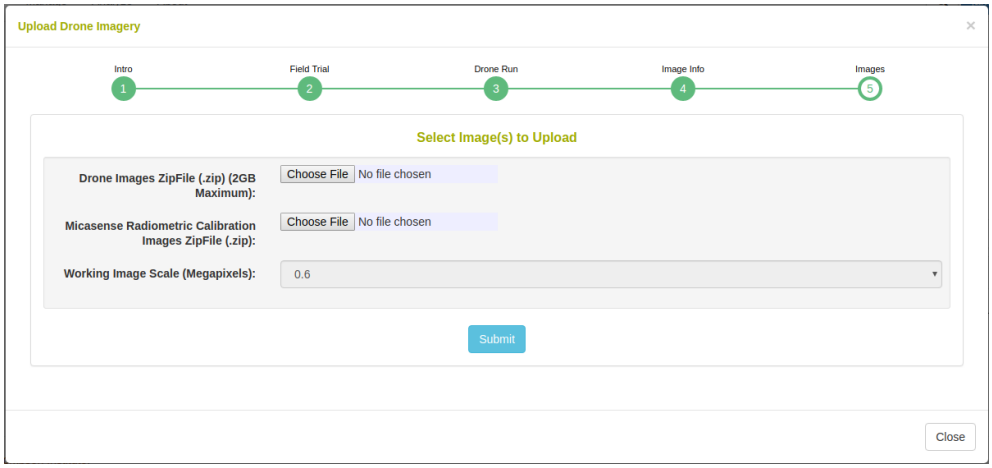
The image data is added to an imaging (drone run) event. Here you can select a previously saved imaging event or you can create a new one by defining a name, description, and date.



The uploaded data can be raw image-captures or complete raster images. Here you can select whether orthophotomosaic stitching is required.

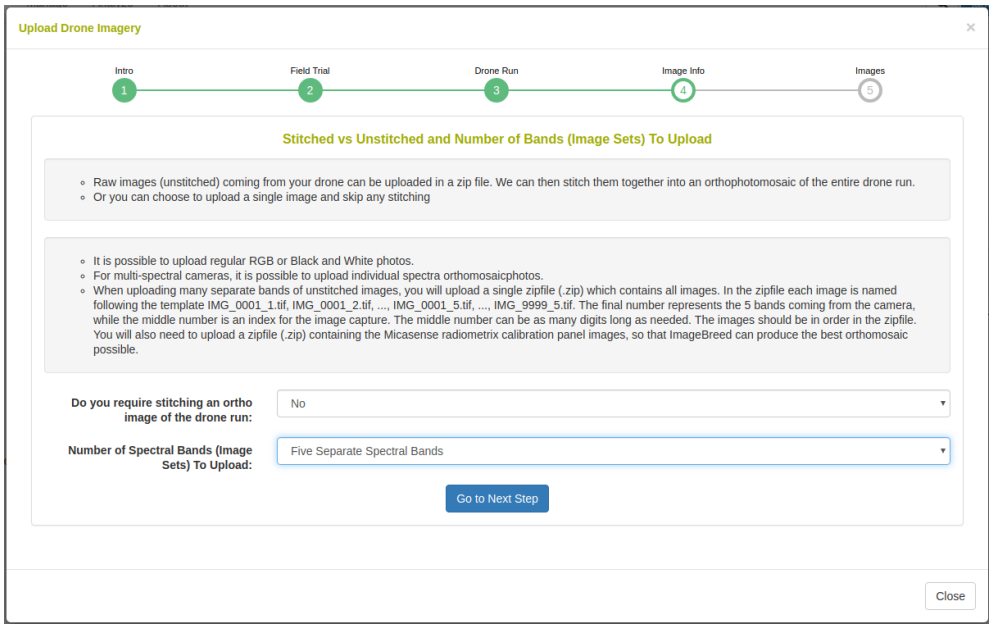


In the case that orthophotomosaic stitching is required, select 'yes'. On the next step you will see the following: Upload a zipfile with the raw-captures. When uploading Micasense RedEdge raw-captures, provide images of the Micasense calibration panels in a zipfile as well.

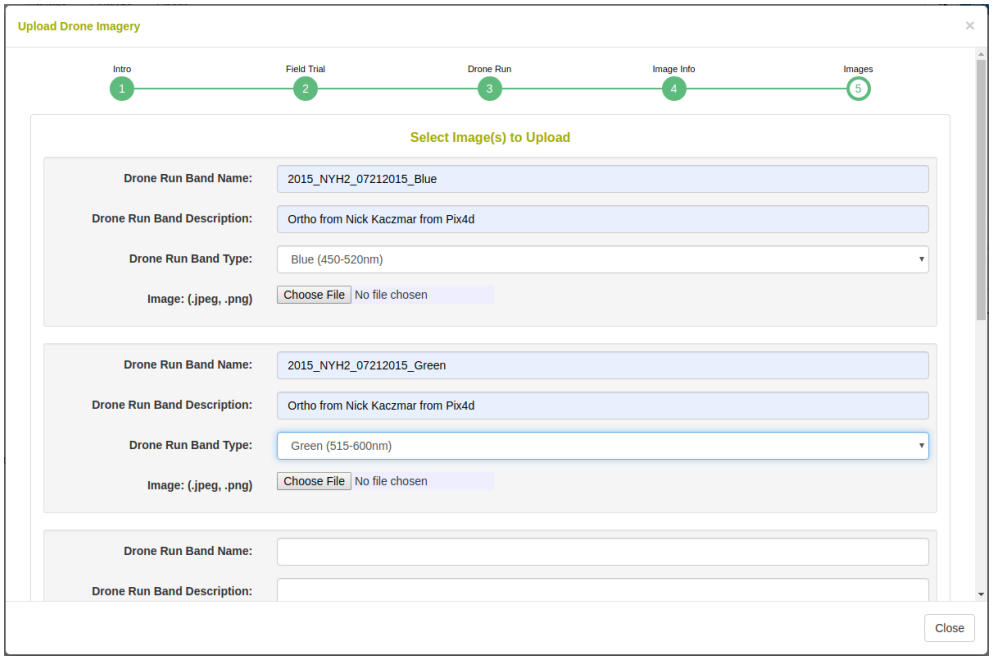


In the case that orthophotomosaic assembly is not required, simple upload the raster images. Select the number of image bands that will be uploaded

e.g. for a five band multispectral camera, select 5.



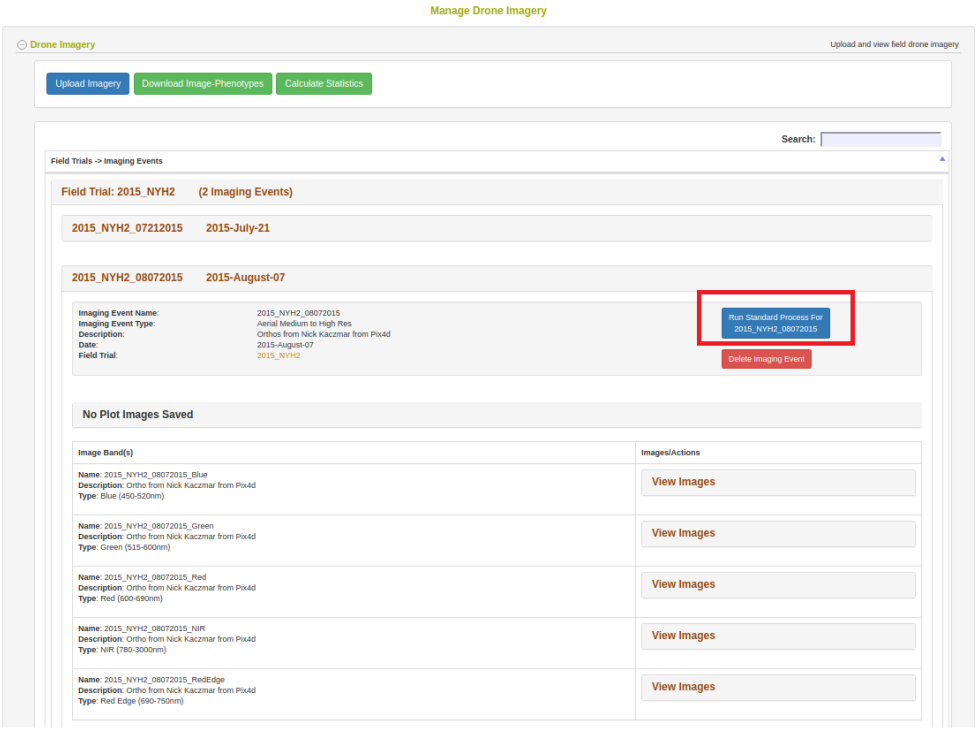
In the caes that orthophotomosaic stitching is not required, select 'no'. On the next step you will see the following:



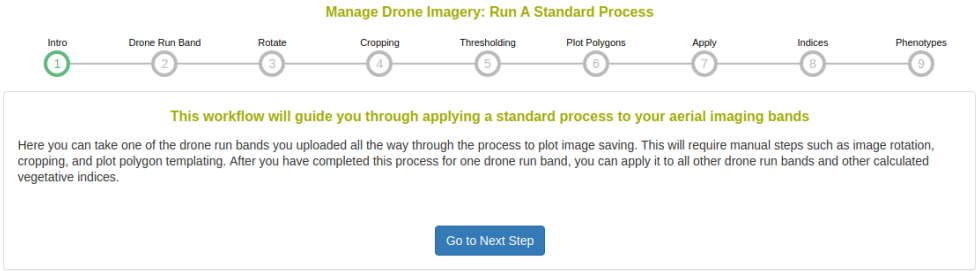
Upload an image at each band with a unique name, description, and spectral type.

20.3 Standard Process

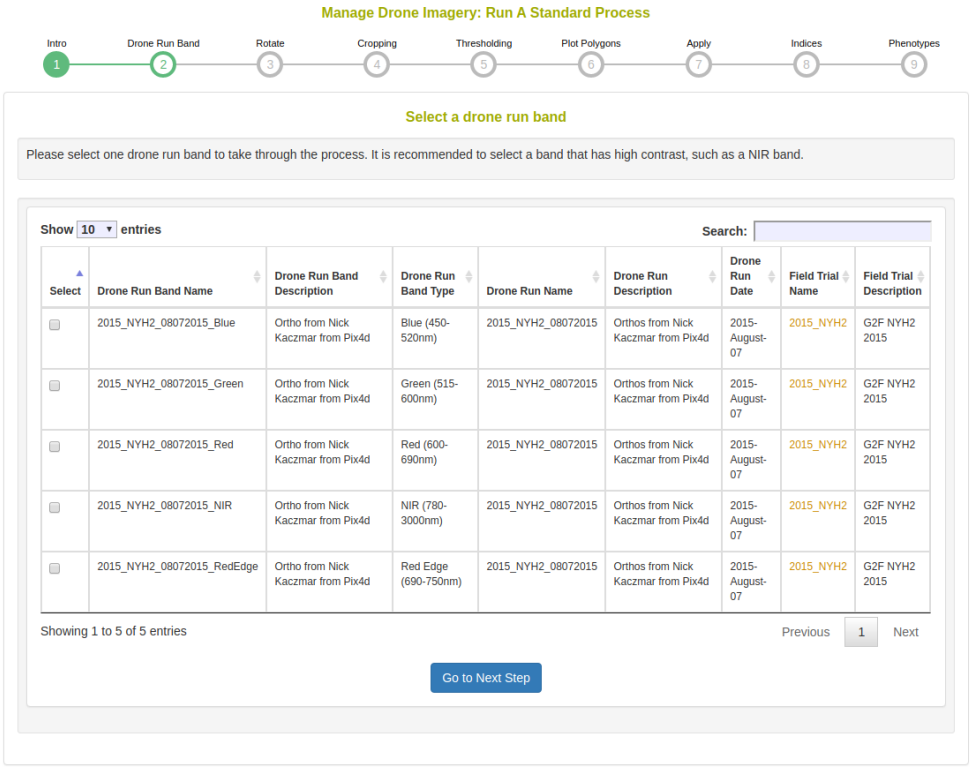
Once imagery is uploaded, it will appear on the dashboard under the field trial. Clicking the "Run Standard Process" button will begin extracting plot-polygon phenotypes from the imagery.



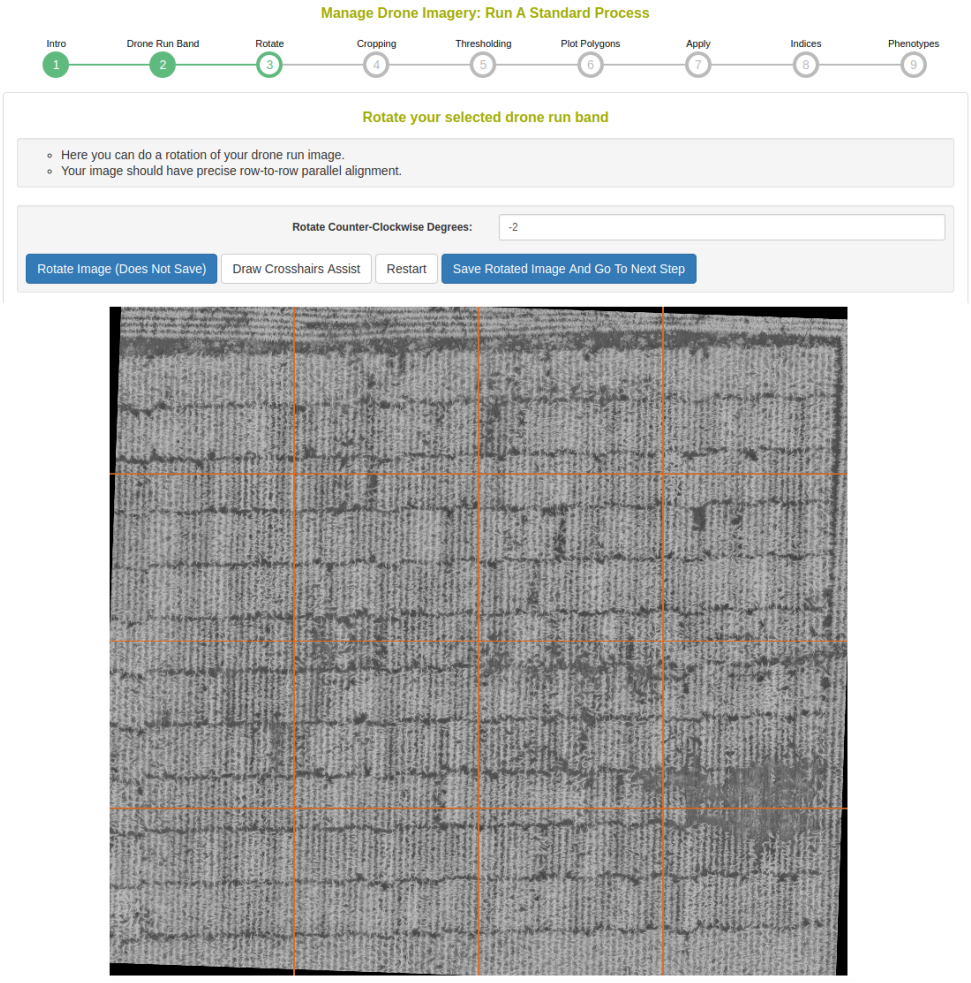
Clicking the button will open the following dialog.



Select a drone run band to use in this process. In the case of the Micasense 5 band multispectral camera there will be 5 bands shown here; select the NIR channel in this case because it has the highest contrast. In the case of standard color images, there will only be the RGB Color Image option here.

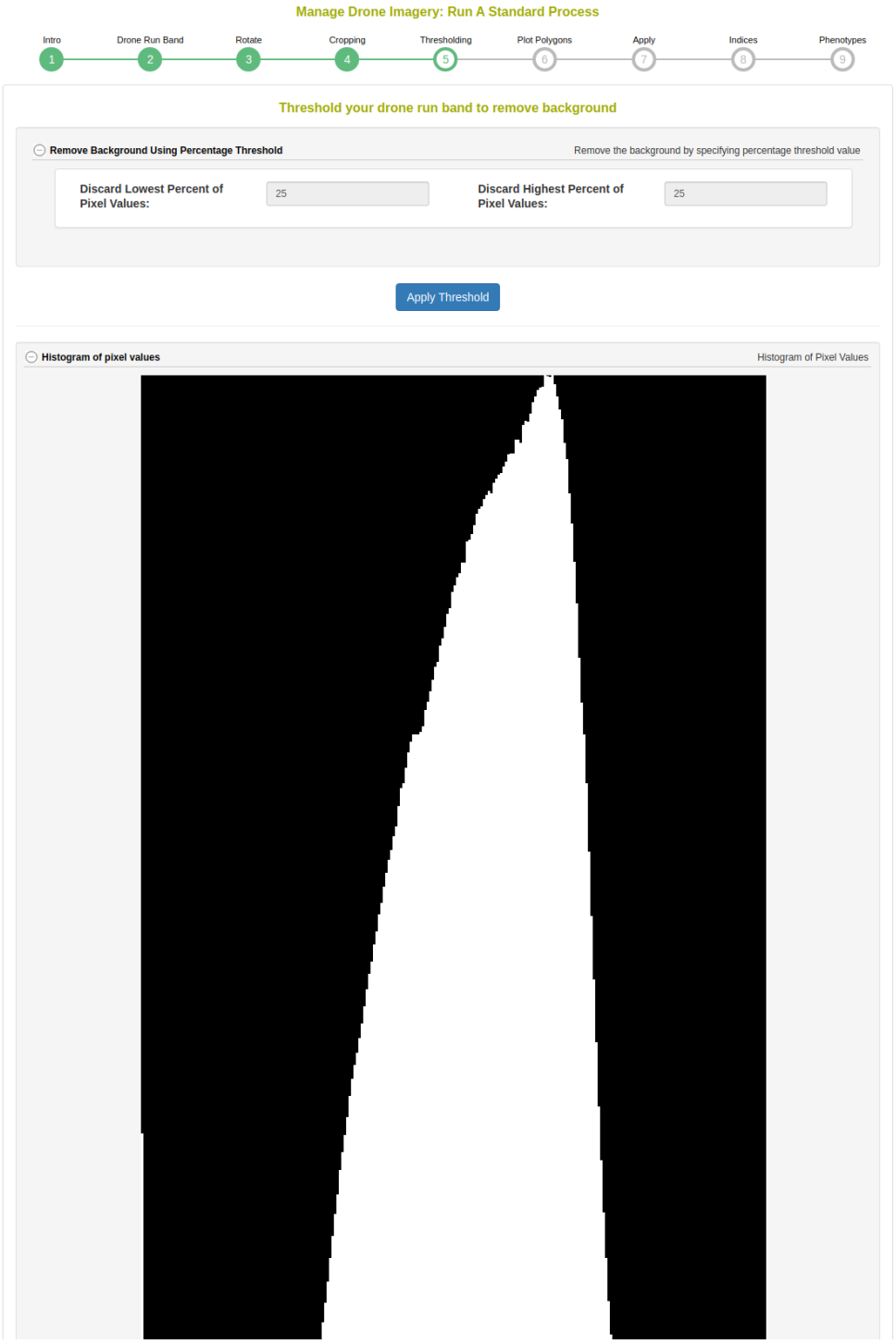


Rotate the image so that there the plots are oriented in a grid fashion. There can be a skew in the field layout, as seen in the following example.



Perform a rough cropping of the image by clicking on the four corners of the field. Cropping is important to remove any extraneous parts of the image.

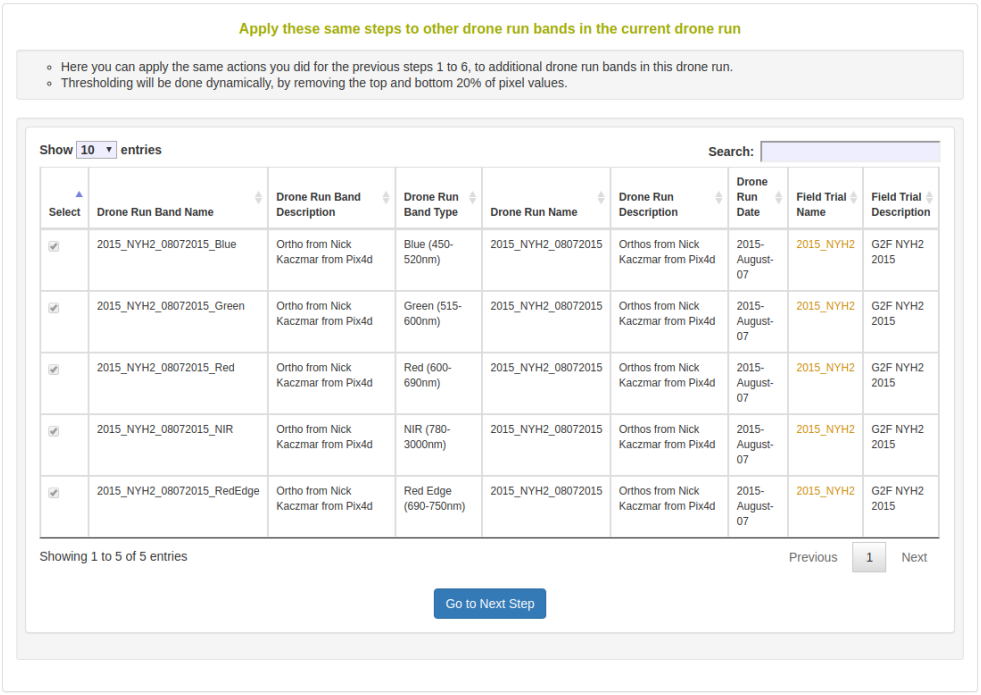
This step shows a histogram of the cropped image. The standard process will magnitude threshold the top and low ends of the distribution.


Manage Drone Imagery: Run A Standard Process
Intro
Drone Run Band
Rotate
Cropping
Thresholding
Plot Polygons
Apply
Indices
Phenotypes
1
2
3
4
5
6
7
8
9
Threshold your drone run band to remove background
Remove Background Using Percentage Threshold
Remove the background by specifying percentage threshold value
Discard Lowest Percent of Pixel Values:
25
Discard Highest Percent of Pixel Values:
25
Apply Threshold
Histogram of pixel values
Histogram of Pixel Values

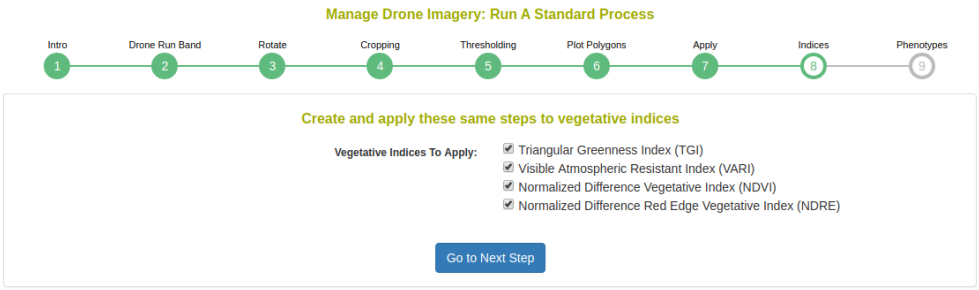
In this step, the template for the plot polygons in the experimental field design are associated to the image. First, defined the number of rows and columns in the field experiment. Then click the four corners of the image, in respect to the top right, top left, botton left, and bottom right positions. Next click on "Draw Plot Polygon Template". Review the template and clear/repeat the process until the template matches well. It is possible to "copy/paste" templates in the case where there are large breaks in the field design. Next, scroll down to the "assign Plot Polygons to Field Trial Entities" section. Select the location of Plot Number 1 as either "top left" or "top right" and whether the field design is serpentine or zigzag. Click on "Generate Assignments" and review that the names of the plots appear correctly in the overlay on the image. Finally, click "Finish and Save Polygons to Plots" when you have have confirmed the assignments.

Manage Drone Imagery: Run A Standard Process
Intro
Drone Run Band
Rotate
Cropping
Thresholding
Plot Polygons
Apply
Indices
Phenotypes
Define plot polygons relative to the field layout
Generate Polygon Template Tool
Overlay a uniform grid over the image.
1
Number of Rows:
10
Number of Columns:
50
Click Top Left Corner
Click Top Right Corner
Click Bottom Left Corner
Click Bottom Right Corner
Template Parameters
View and edit template parameters. These parameters can be filled using the above button clicks
2
Draw Plot Polygon Template (Does not save. Apply multiple templates if needed.)
Previously Used Plot Polygon Templates
View and use previous plot polygon templates used on this field trial
Total Image Width: 3316px. Total Image Height: 2680px.
Clear All Polygons
Clear One Polygon
Find Distance Between Points
Assign Plot Polygons to Field Trial Entities
Location of First Plot (e.g. plot number 1):
Top Left
Second Plot Follows First Plot Going:
Right
3
Plot Number Orientation:
Serpentine
4
5
Generate Assignments (Does Not Save)
Finish and Save Polygons To Plots
Show 10 entries
Search:
plot_name
accession_name
plot_number
block_number
is_a_control
rep_number
row_number
col_number
plot_geo_json
Polygon Assigned
2015_NYH2_plot_1 PHV63/LH195 1 1 null 1 1 45
2015_NYH2_plot_10 CG106/PHZ51 10 1 null 1 1 54
2015_NYH2_plot_100 LH145/LH82 100 4 null 1 2 45
2015_NYH2_plot_101 PHN11_LH145_0002/PHB47 101 5 null 1 3 45
2015_NYH2_plot_102 NYH-081/PHB47 102 5 null 1 3 46
2015_NYH2_plot_103 W1000S_0029/PHB47 103 5 null 1 3 47
2015_NYH2_plot_104 B109/PHZ51 104 5 null 1 3 48
2015_NYH2_plot_105 PH2D7_PHG47-13/PHB47 105 5 null 1 3 49
2015_NYH2_plot_106 PH2D7_PHG47-19/PHB47 106 5 null 1 3 50
2015_NYH2_plot_107 CG65/LH82 107 5 null 1 3 51
Showing 1 to 10 of 500 entries
Previous
1
2
3
4
5
...
50
Next

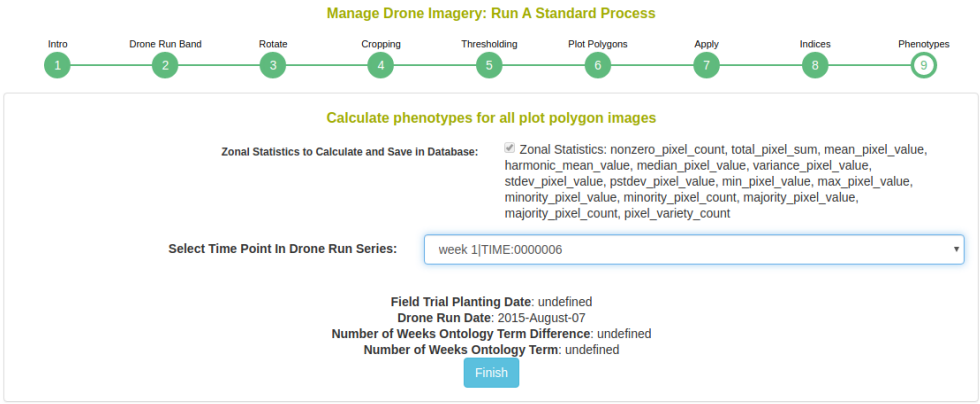
Next, the dialog shows you that the standard process will be repeated for all uploaded image bands.



Next, choose which vegetation indices to apply.



Next, choose the phenotypic values to extract. You must define the time point for which the phenotype is; if the field trial has a planting date, the time point will automatically be populated as image date minus the planting date.



After completing the standard process, the job will continue in the background until it completes. You can check the status of the job from the dashboard.

20.4 Ground Control Points

Ground control points can be saved after an imaging event has undergone the standard process on orhomosaics. Ground control points can then be used across imaging events on the same field experiment in order to automate the entire standard process.

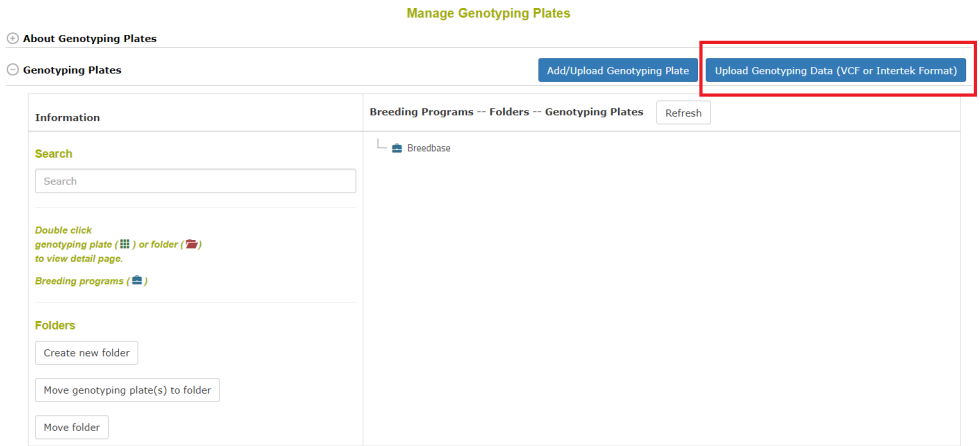
Chapter 21

Managing VCF Data

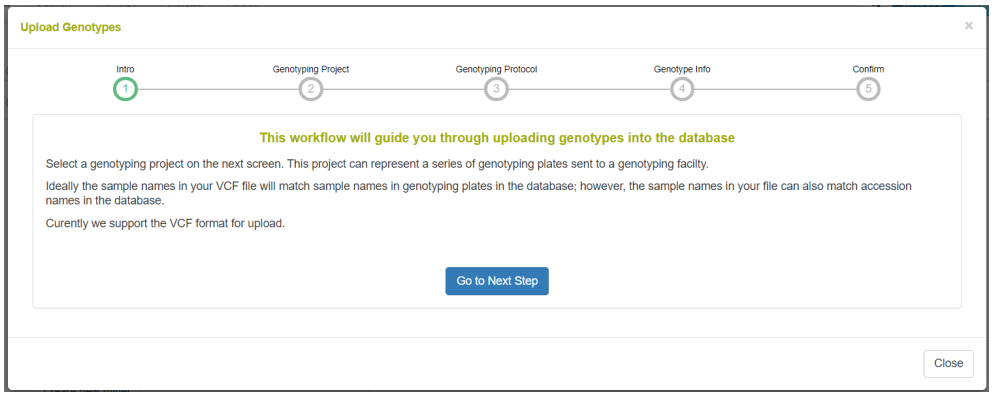
21.1 Uploading VCF Data

Genotyping data in VCF can be loaded from the web-interface. Breedbase can store any genotypic variants from a VCF, allowing for polyploids, structural variants, etc. without problems.

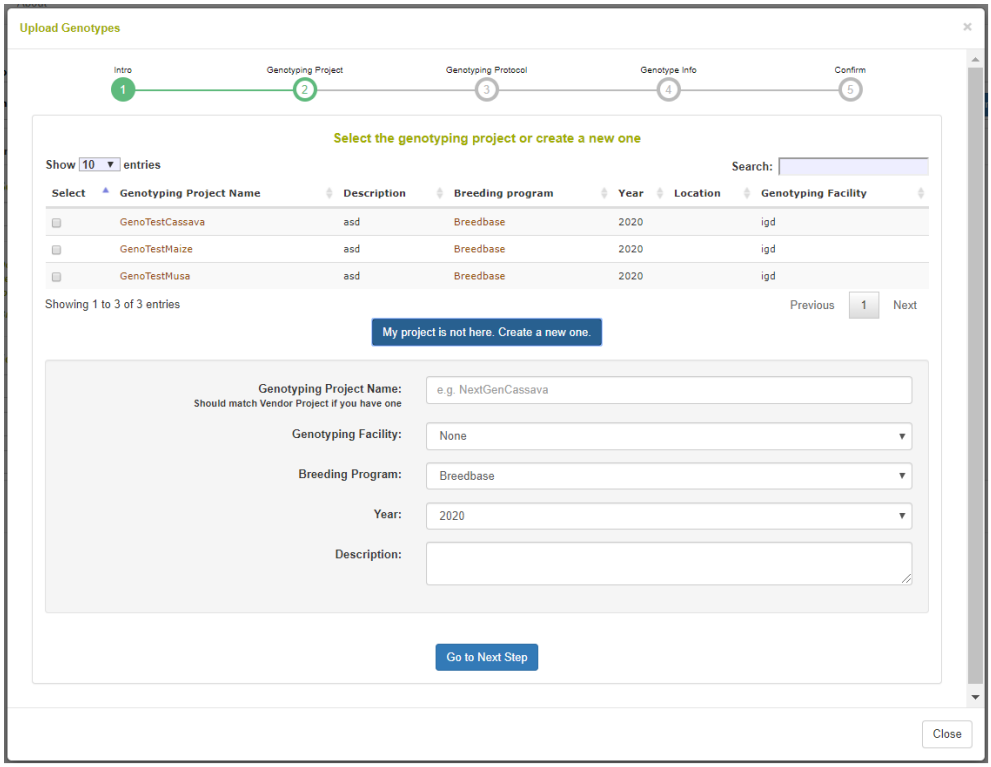
To begin go to Manage->Genotyping Plates and click the button seen below: Note that you do not need to have genotyping plates uploaded to upload VCF data; you may upload genotyping data to accessions or you can upload genotyping data for tissue samples in genotyping plates.



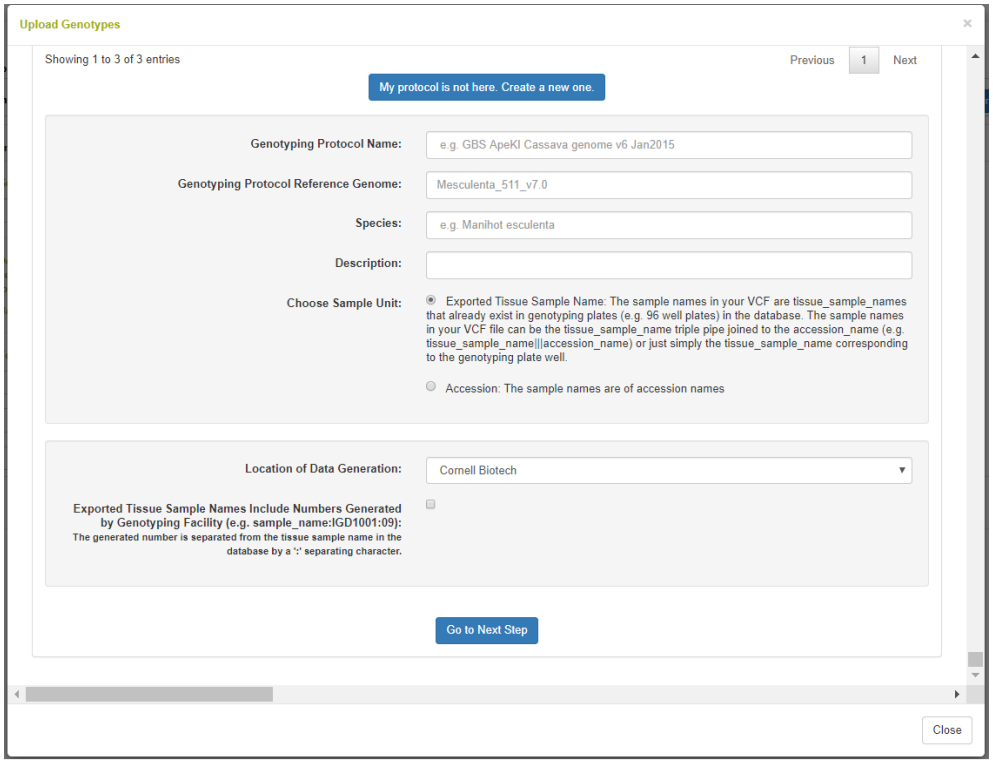
The workflow begins with an intro:



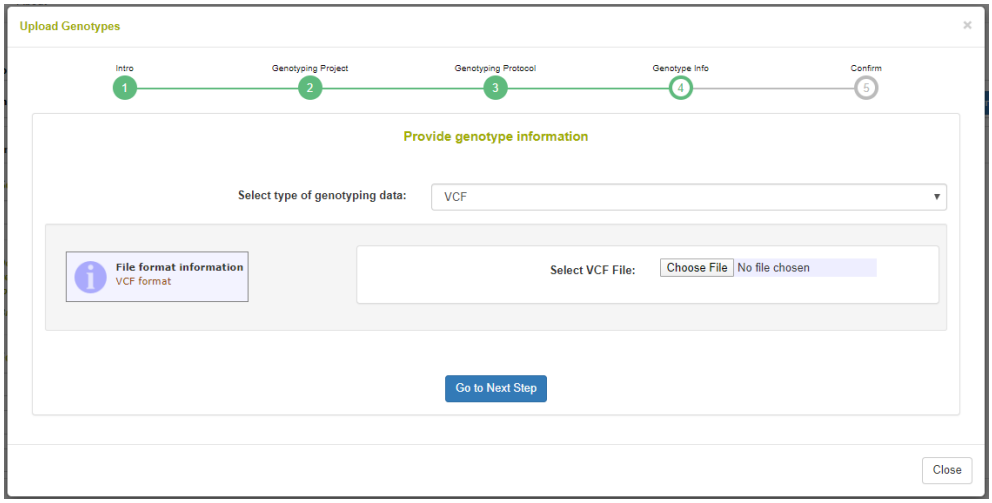
On the following step in the workflow, a genotyping project is defined or selected. A genotyping project is a high-level entity for grouping several genotyping events. It is defined with a name, description, name, breeding program, and genotyping facility (IGD, Intertek, etc.).



The following step is to define or select a genotyping protocol. A genotyping protocol represents the set of markers being called against a specific reference genome. A genotyping protocol is defined with a name, description, reference genome name, species name, and a location of data generation. Note in the picture that you can select whether the samples in your file are accessions or tissue samples in the database; tissue samples are for when a genotyping plate is stored in the database. There is an option to parse the sample names for appended sequencing numbers from IGD, where the sample names are like "accession:igdnumber".



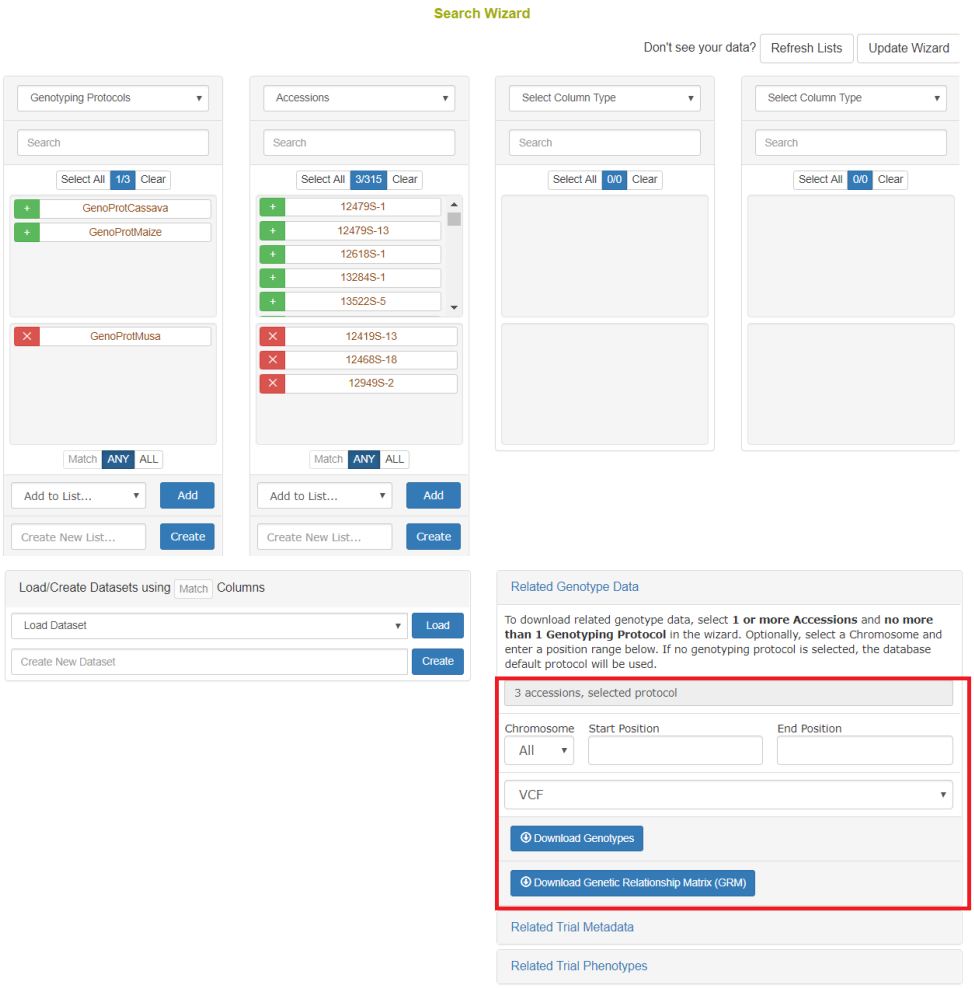
The final step is to select the VCF from your computer and upload it. The web interface can be used to upload files arbitrarily large; it is a NGINX configuration to set this size.



21.2 Searching and Downloading VCF Data

The Search Wizard is the primary means of querying data in the database. Go to Search->Wizard to begin.

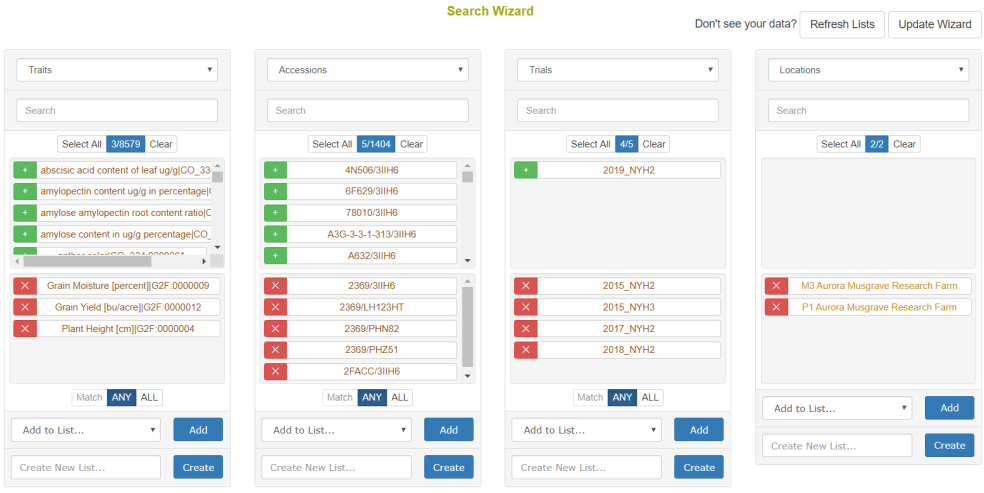
Once genotyping protocols are stored, select Genotyping Protocols from the first dropdown menu. Then if you select one or more and select Accessions from the second dropdown menu, you will see the accessions for which genotypes were stored. As seen in the following picture, there is a section for filtering genotypes by chromosome, start position, and end position. Genotypes can be downloaded in VCF or DosageMatrix formats.



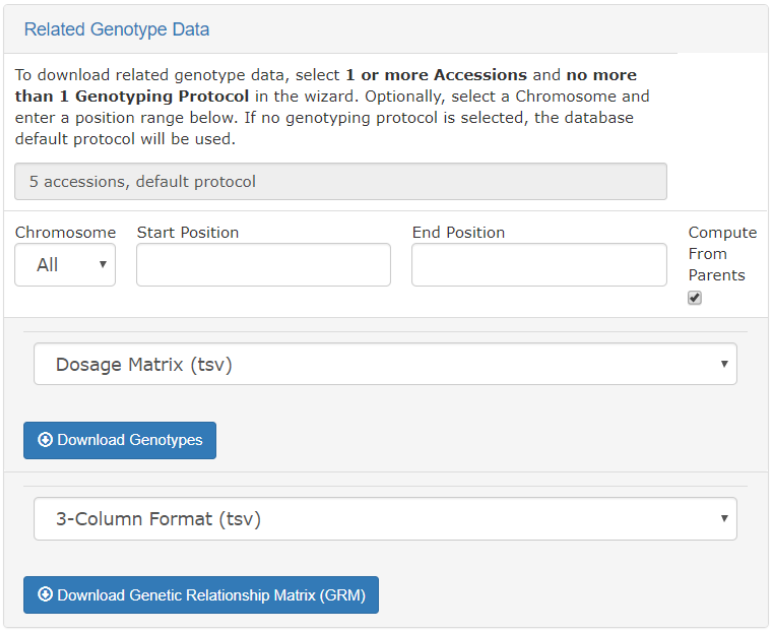
Using the "Default genotyping protocol" which is configured in a system, you can query over field phenotypic evaluations before downloading genotypes and phenotypes.

21.3 Searching Protocols

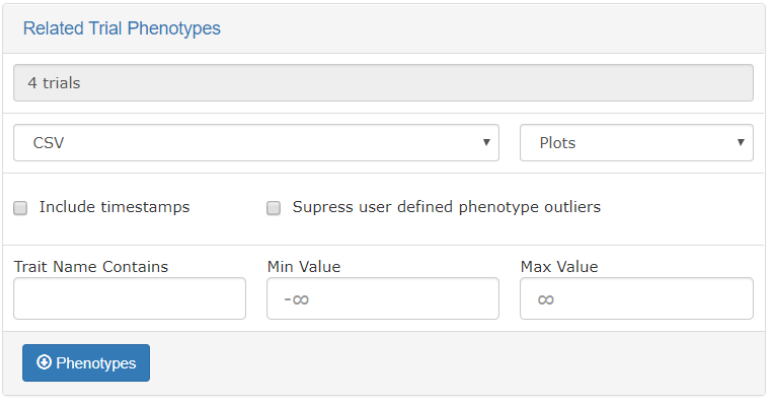
Genotyping protocols can be search by going to Search->Genotyping Protocols. To download genotypes accessions must be selected, though any combination of search criteria can be used to filter and select those accessions. If a genotyping protocol is not selected, then the default genotyping protocol set in the configuration will be used. Genotyping protocols can also be selected in the wizard.



The genotyping download menu on the Search Wizard presents options for filtering by chromosome, start position, and end position. Genotypes can be downloaded in VCF of Dosage Matrix formats. The genomic relationship matrix (GRM) can be downloaded for the selected accessions in a tab-delimited matrix format or in a three-column format that is useful in Asreml. Genotypes can be computed from the parents in the pedigree if those parents are genotyped by clicking on the "compute from parents" checkbox. Additionally, the GRM can be computed using genotypes of parents in the pedigree if the "compute from parents" checkbox is selected.



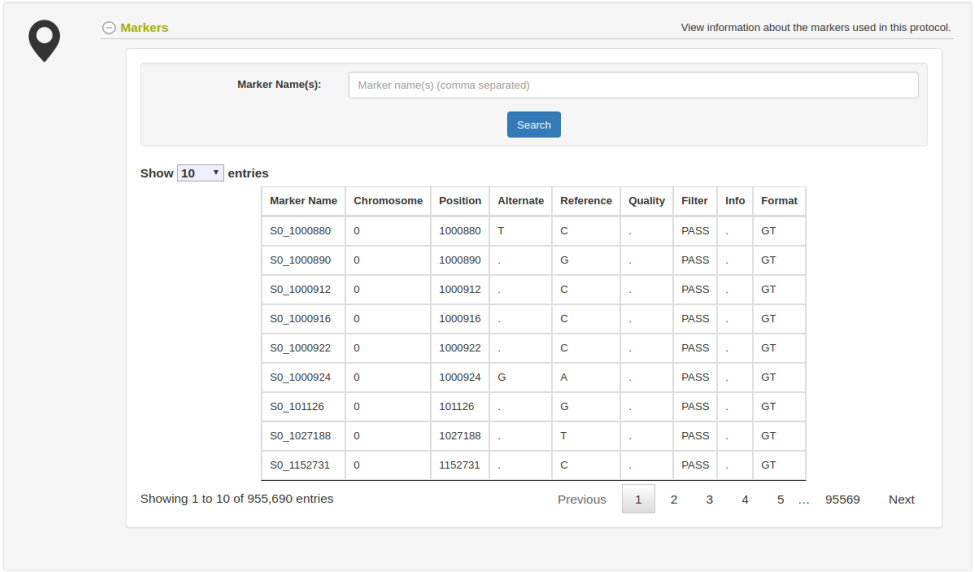
As is described elsewhere, the Search Wizard presents a way to filter phenotypic values by minimum and maximum values, and allow for download in CSV and Excel formats.



21.4 Detail Pages and Deletion

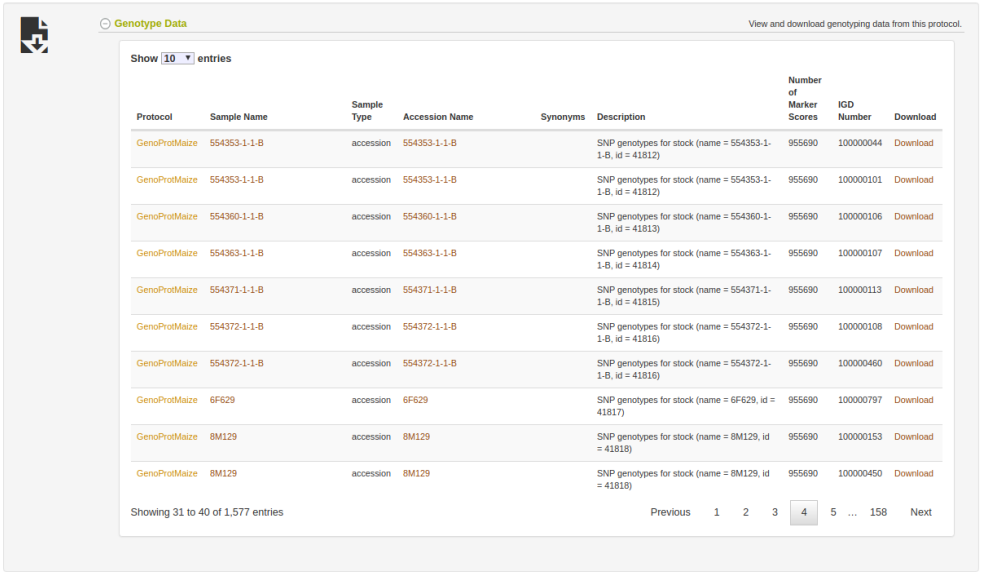
The genotyping protocol detail page will show all information about the protocol such as the reference genome used, the header information lines in the uploaded VCF file, the markers involved, and the samples genotyped.

The markers section will show all markers used and their annotations, such as position, chromosome, alternate allele, reference allele, marker format, etc.

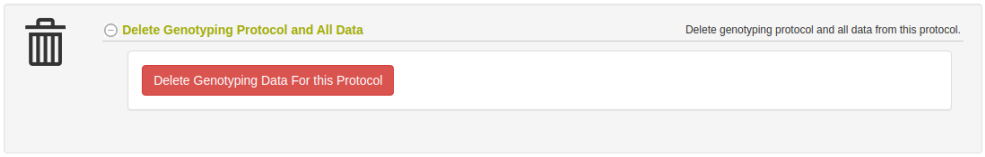


The samples section will show all samples genotyped. Notice the Download links in the table which can be used to easily get the VCF file results for each

genotyped samples with all markers in the genotyping protocol. For getting mulitple samples at once, use the Search Wizard as discussed above.



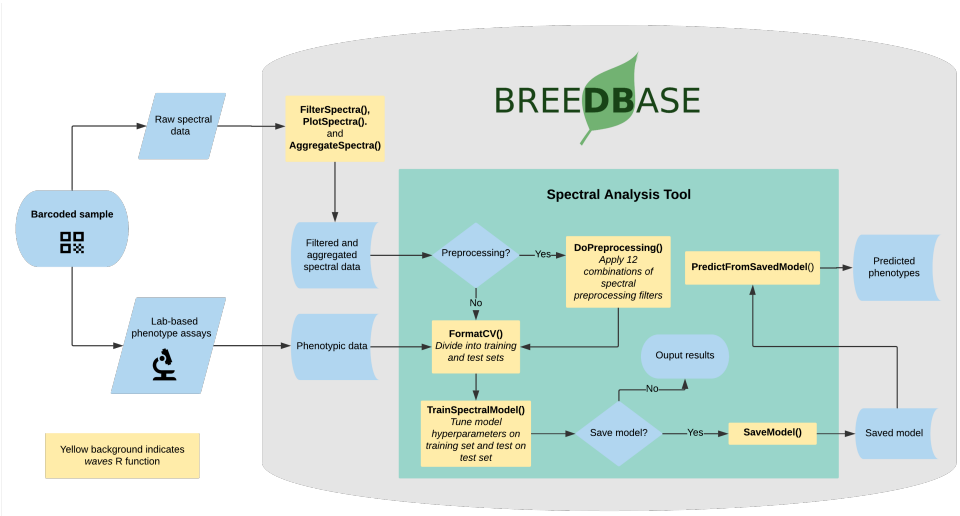
The genotyping protocol and all associated genotyping data can be deleted from the genotyping protocol page.



Chapter 22

Managing Spectral Data

Breedbase has implemented a flexible spectral data storage protocol that handles spectral data irrespective of the source spectrometer. Spectral data storage and analysis in Breedbase makes use of the R package *waves* for outlier identification, plotting, sample aggregation, and prediction model training.

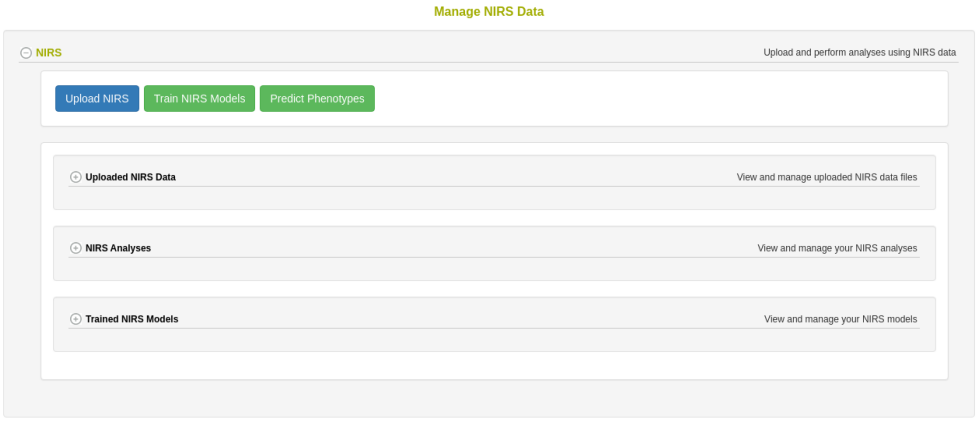


22.1 Upload Spectral Data

Spectral data can be added as a CSV file that includes metadata in the leftmost columns followed by one column per spectral measurement to the right. Rows represent a single scan or sample, each with a unique ID that must match to a Breedbase observationUnitName. Future data transfer using BrAPI will allow for interoperability with data collection software.

To upload a spectral dataset, navigate to the 'Manage NIRS Data' page by selecting 'NIRS' in the 'Manage' menu and click the blue 'Upload NIRS' button. This will open an upload workflow. A link to the required file format and an example .csv file can be found by clicking in the light blue info box in this workflow. Another example of the file format is shown below.

- **id**: Optional identifier for each NIRS read. The id must be an integer.
- **sampling_id**: Optional identifier for each sample. Strings are allowed.
- **sampling_date**: Optional field. The format allowed is: YYYY-MM-DD.
- **observationunit_name**: Required field that matches existing data in the database. It can be the plot name, subplots, plant name, or tissue sample, depending how your trial is designed.
- **device_id**: Optional field to identify your device. Strings are allowed.
- **device_type**: Required field. It is possible upload data for a single device type. They can be: SCiO, QST, Foss6500, BunchiN500, or LinkSquare.
- **comments**: Optional field for general comments. All other columns are required wavelengths. You can add how many columns you want upload – there is no limit.



	id	sample_id	sampling_date	observationunit_name	device_id	device_type	comments	740	741	742	743	744	745	746
2	1	7a6ac477-d291-4d07-a6	2020-6-24	myTrial20_rep1_acc_001	503E4BFC4E923999	SCIO		0.885707958	0.88572938	0.885590265	0.885493457	0.885493162	0.885572662	0.885628732
3	2	7a6ac477-d291-4d07-a6	2020-6-24	myTrial20_rep1_acc_002	503E4BFC4E923999	SCIO		0.909132994	0.908724223	0.90824451	0.907891706	0.907697281	0.907626115	0.90755938
4	3	7a6ac477-d291-4d07-a6	2020-6-24	myTrial20_rep1_acc_003	503E4BFC4E923999	SCIO		0.889220207	0.889013119	0.888681812	0.888431257	0.888310362	0.888297321	0.888284052
5	4	73c648ca-f5fb-4231-a1	2020-6-24	myTrial20_rep1_acc_004	503E4BFC4E923999	SCIO		0.8900087	0.889604969	0.889191073	0.888956654	0.88883379	0.8890727241	0.8892472
6	5	73c648ca-f5fb-4231-a1	2020-6-24	myTrial20_rep1_acc_005	503E4BFC4E923999	SCIO		0.938101707	0.93668202	0.93820132	0.937873867	0.937742819	0.937775129	0.93784594
7	6	73c648ca-f5fb-4231-a1	2020-6-24	myTrial20_rep1_acc_006	503E4BFC4E923999	SCIO		0.876289461	0.875981159	0.875579263	0.875289805	0.875162225	0.875171404	0.87520442
8	7	d5b55c93-4c8e-4ef6-9c	2020-6-24	myTrial20_rep1_acc_007	503E4BFC4E923999	SCIO		0.879217838	0.878925781	0.878588441	0.8783872	0.878346547	0.878423929	0.878495739
9	8	7a6ac477-d291-4d07-a6	2020-6-24	myTrial20_rep1_acc_008	503E4BFC4E923999	SCIO		0.890746588	0.890515672	0.89016542	0.889903562	0.889783304	0.889782828	0.889792154
10	9	7a6ac477-d291-4d07-a6	2020-6-24	myTrial20_rep1_acc_009	503E4BFC4E923999	SCIO		0.850444238	0.85032422	0.850094039	0.8499495	0.849942163	0.850052119	0.850175036

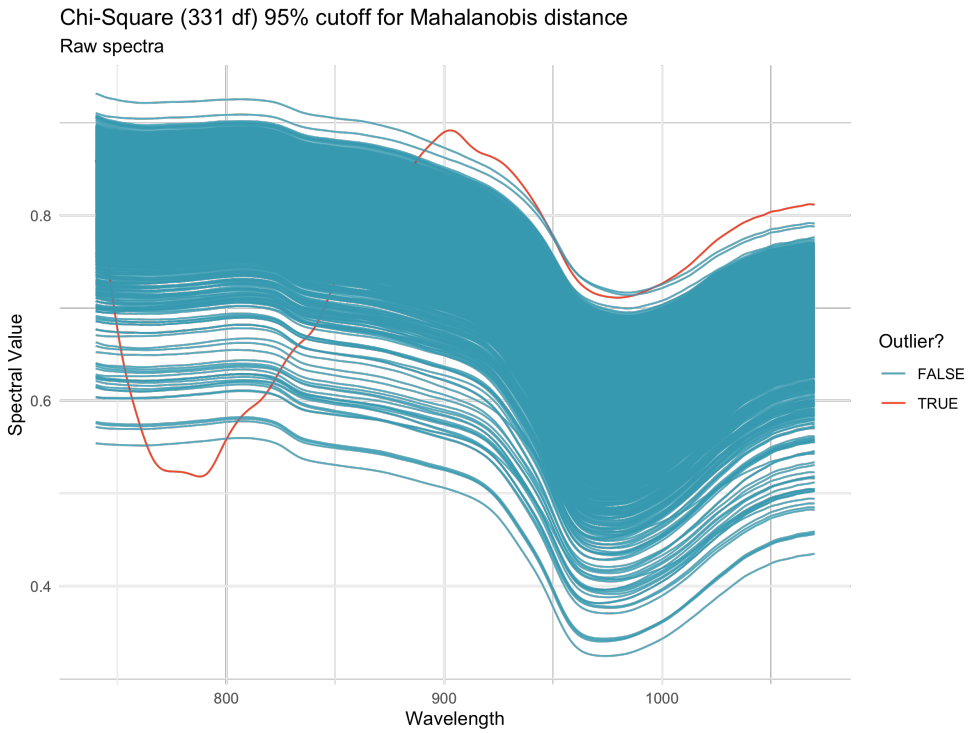
22.2 Evaluate and Remove Outliers

Spectral calibration models can be heavily affected by the presence of outliers, whether they come from spectrometer spectral artifacts or user errors. Mahalanobis distance (Mahalanobis, 1936) is a measure of the distance between a single observation and a larger distribution and is commonly used in the identification of outliers in a multivariate space (Des Maesschalck et al, 2000). The *FilterSpectra()* function in the R package *waves* calculates the Mahalanobis distance of each observation in a given spectral matrix using the *stats::mahalanobis()* function. Observations are identified as outliers if the squared distance is greater than the 95th percentile of a χ^2-distribution with p degrees of freedom, where p is the number of columns (wavelengths) in the spectral matrix (Johnson and Wichern, 2007). In Breedbase, this procedure is applied on a per-dataset basis on upload and outliers are given binary tags "Outlier."

22.3 Plot Spectra

After outlier identification, a plot is generated using the *PlotSpectra()* function in *waves*. This function uses the filtered spectra and *ggplot2::ggplot()* to

create a line plot with outliers highlighted by color. A list of rows identified as outliers are shown beneath the plot. Plots are saved as .png files and linked to the original input datasets. Plot image files can be downloaded via the "Download Plot" button in the upload workflow.



22.4 Aggregate Spectra

To obtain a stable and reliable spectral profile, most spectrometer manufacturers recommend that multiple spectral scans are captured for each sample. While some spectrometers aggregate these scans internally, many do not, requiring the user to do so before analysis can take place. Breedbase handles these cases upon data upload following filtering steps by calling the *AggregateSpectra()* function from *waves*, saving the aggregated scans for future access through the search wizard feature. Scans are aggregated by sample mean (e.g. plot-level basis) according to the provided observationUnitName field. After aggregation, the user exits the upload workflow and the raw data file is saved in the upload archive.

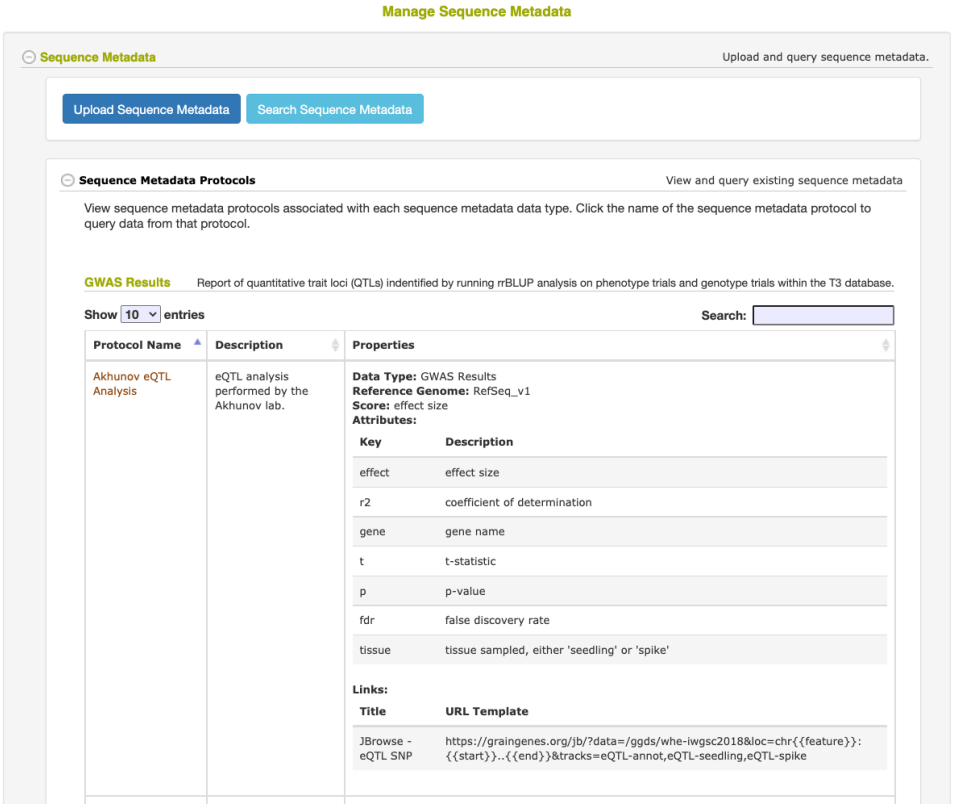
22.5 References

- De Maesschalck, R., Jouan-Rimbaud, D., and Massart, D. L. (2000). The Mahalanobis distance. Chemom. Intell. Lab. Syst. 50(1): 1-18.
- Johnson, R. A. & Wichern, D. W. (2007). Applied Multivariate Statistical Analysis (6th Edition). p 773.
- Mahalanobis, P. C. (1936). On the generalized distance in statistics. National Institute of Science of India.

Analysis tool documentation

Chapter 23

Managing Sequence Metadata



23.1 What is Sequence Metadata?

Sequence Metadata is a feature that allows for the efficient storage and retrieval of sequence annotations for a specific region along a reference genome. The annotation data can contain a primary "score" value and any number of secondary key/value attribute data. For example, Sequence Metatadata can store MNase open chromatin scores for every 10 basepairs along the reference genome as well as genome-wide association study (GWAS) statistics, including the trait information associated with the result. This data can then be filtered by position and/or scores/attribute values and even cross-referenced with markers stored in the database.

23.2 Loading Sequence Metadata

Sequence Metadata can be loaded into the database using a gff3-formatted file. The following columns are used to load the data:

- **#1 / seqid:** The name of the database feature (ie chromosome) the metadata is associated with (The feature name must already exist as a feature in the database)
- **#4 / start:** The metadata's start position
- **#5 / end:** The metadata's end position
- **#6 / score:** (optional) The primary score attribute of the metadata
- **#9 / attributes:** (optional) Secondary key//value attributes to be saved with the score. These should be formatted using the gff3 standard (key1=value1;key2=value2). The attribute key cannot be either score, start, or end.

To upload the gff3 file:

1. Go to the **Manage > Sequence Metadata** page
2. Click the **Upload Sequence Metadata** button
3. On Step 2 of the Wizard, select the Type of data to be uploaded
    - This groups similar datasets together in the same Data Type category
4. On Step 3 of the Wizard, select an existing Protocol or create a new one
    - The Protocol is used to describe how the data was generated and define the score value and any secondary attributes. Adding the

attributes (and their descriptions) to the Protocol will allow the Sequence Metadata queries to filter the data based on the value of one or more of these attributes. Attributes not defined in the Protocol will still be stored and displayed on retrieval, but will not be able to be used in a search filter.

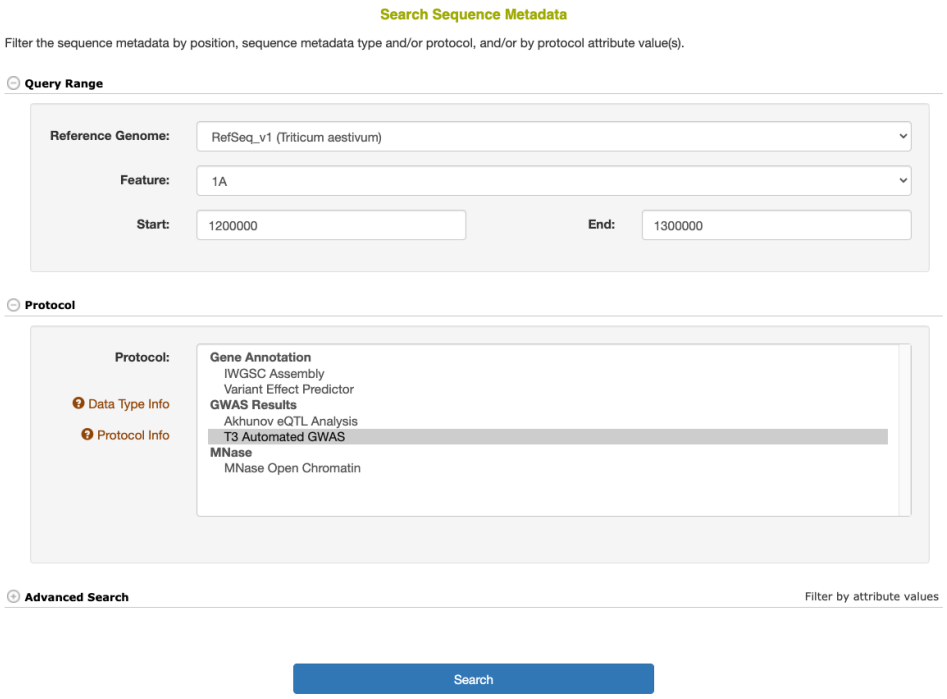
5. Finally, select and upload your gff3 file to the database. The database will verify the format of the file before its contents are stored.

23.3 Searching Sequence Metadata

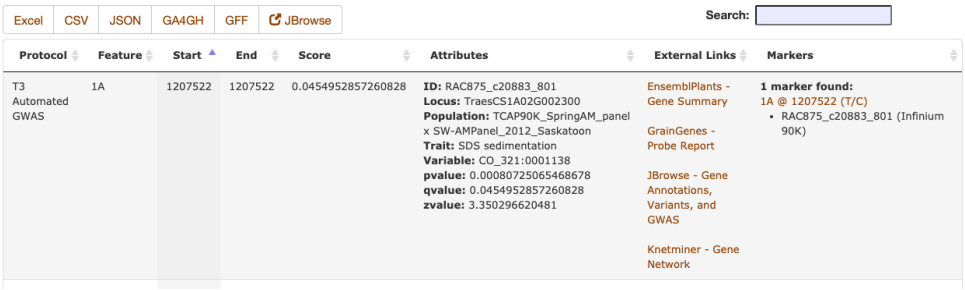
To retrieve stored Sequence Metadata, go to the **Search > Sequence Metadata** page.

23.3.1 Basic Search

The basic Sequence Metadata search options include selecting the reference genome and species, the chromosome, and (optionally) the start and/or end position(s) along the reference genome. In addition, one or more specific protocols can be selected to limit the results.

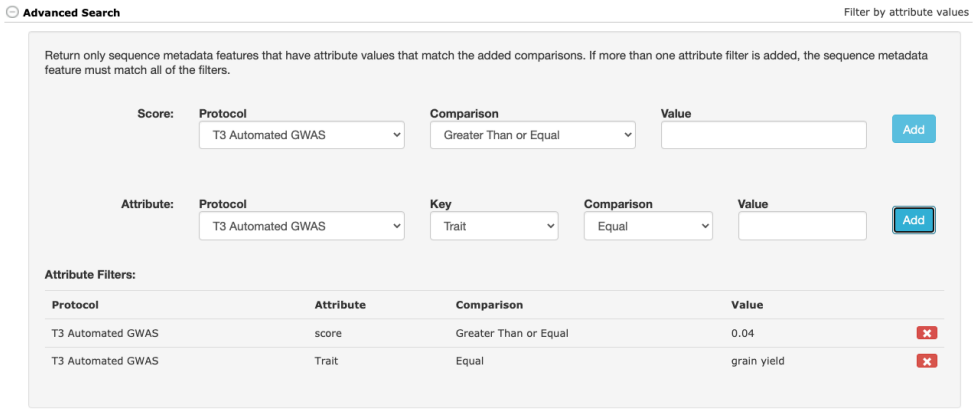


The Sequence Metadata search results are returned as a table, including the chromosome and start/stop positions of the annotation, along with the primary score value and any additional key/value attributes. The markers column will include a list of marker names of any stored markers that are found within the start/stop positions of the Sequence Metadata. The data can be downloaded as a table in an Excel or CSV file or a machine-readable (code-friendly) JSON file. If the Sequence Metadata JBrowse configuration is set, the filtered results can be displayed as a dynamic JBrowse track.



23.3.2 Advanced Search

Any number of advanced search filters can be applied to the query. The advanced filters can limit the search results by the value of the primary score and/or any of the secondary attribute values.



23.4 Marker Integration

A table of Sequence Metadata annotations are embedded on the Marker/Variant detail page. The table will include any annotations that span the poisiton of the marker (for data of the same reference genome and species).

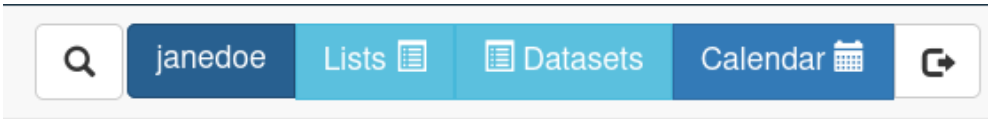
23.5 Sequence Metadata API

A publicly accessible RESTful API (Application Programming Interface) is available to query the database for Sequence Metadata directly from your programming environment (R, python, etc) to be used in analysis. The data is returned in a JSON format. Documentation for the API can be found on the **Manage > Sequence Metadata** page

Chapter 24

Managing Outliers in Dataset

24.1 What is Outliers Functionality in Dataset ?



As in step The Search Wizard we can create a dataset.

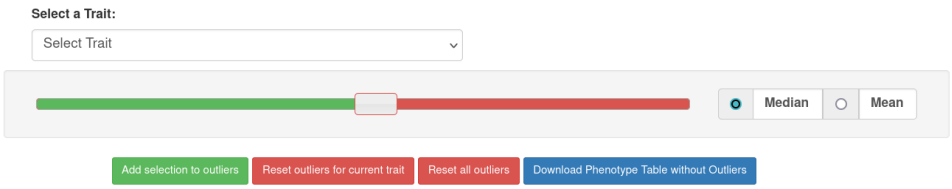
The dataset incorporates a feature to identify outlier points, which we may choose to exclude from a specific dataset. It's important to note that these exclusions only apply at the dataset level, and no data is permanently removed from the database. Additionally, outlier categorization can be modified at any time, and these changes are visible to all other functionalities within the system.

Each dataset stores a wholly unique set of outlier points, completely independent of any other dataset in the database. Outliers are specifically designated for traits within datasets, exclusively encompassing phenotype data. If a particular dataset lacks traits as a part of wizard selection, this functionality is not available.

Each trait has its own set of defined outliers.

24.2 Accessing Trait Visualization

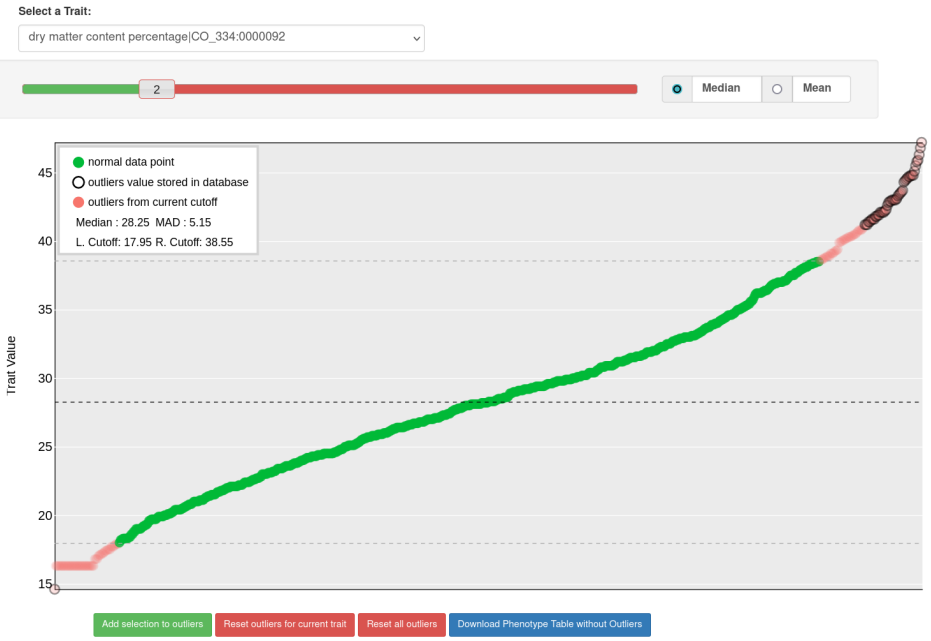
Once you've selected a specific trait, the web application provides access to a visualization of the data points associated with that trait.



24.3 Interpreting Visual Elements

Once you've selected a specific trait, the web application provides access to a visualization of the data points associated with that trait.

- **Green Points**: As per the legend, represent values for the selected trait that fall below the cut-off point set by the slider. (non-outliers)
- **Black Outlined Points**: These data points are outlined with black borders, indicating that they are currently designated as outliers in the database.
- **Red Points**: The red data points denote the cut-off points established by the slider for the allowable deviation value.



24.4 Choosing Cut-Off Values

You have two fundamental options for setting cut-off points:

- **Median with MAD**: This option involves using the median (middle value) along with the Mean Absolute Deviation (MAD) as a reference point for determining cut-off values.
- **Mean with Standard Deviation**: Alternatively, you can choose to use the mean (average) in conjunction with the Standard Deviation to set cut-off points.

24.5 Setting Deviation Multiplier

The slider allows you to specify the deviation multiplier from a central point, which influences the cut-off values.

24.6 Utilizing Graph Controls

Beneath the graph, you'll find four buttons, each serving a distinct function:

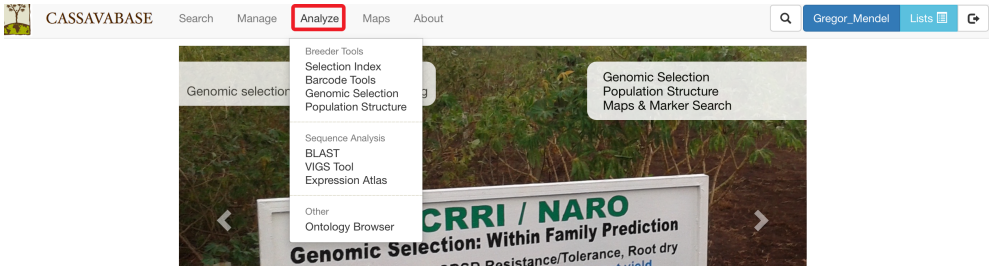
- **Add selection to outliers**: This button enables you to save the current cut-off points to the database for future reference.
- **Reset outliers for current trait**: You can use this option to reset outliers for the selected trait.
- **Reset all outliers**: This button allows you to reset outliers for the entire dataset.
- **Download Phenotype Table without outliers**: You can download the phenotype data table in a comma-separated value format file, using this feature, with outliers excluded for selected dataset.

These tools and functions are designed to provide you with control and insights when working with data visualization and outliers.

Chapter 25

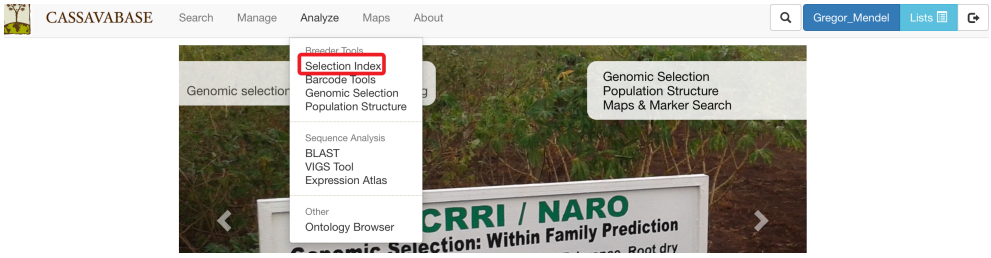
Data Analysis Tools

SGN databases provides several tools for phenotype data analysis, marker-assisted selection, sequence and expression analyses, as well as ontology browser. These tools can be found in the "Analyze" menu.

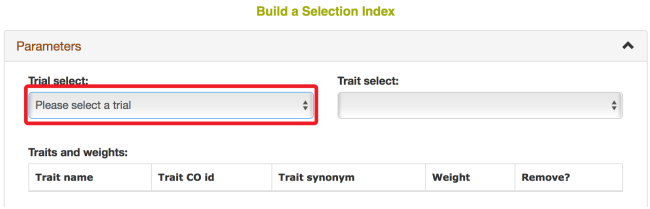


25.1 Selection Index

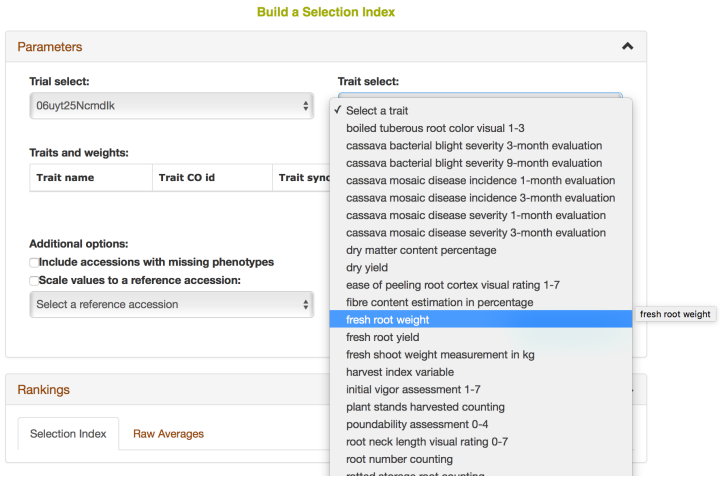
To determine rankings of accessions based on more than one desirable trait, SGN databases provide a "Selection Index" tool that allows you to specify a weighting on each trait. To access the tool, clicking on "Selection Index" in the "Analyze" menu.



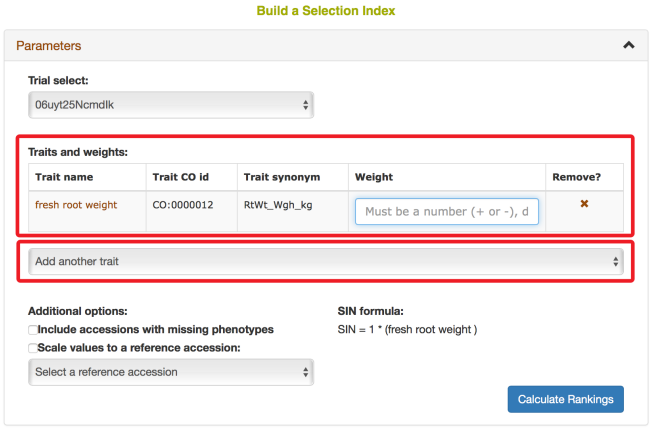
On the Selection Index page, selecting a trial that you want to analyze.



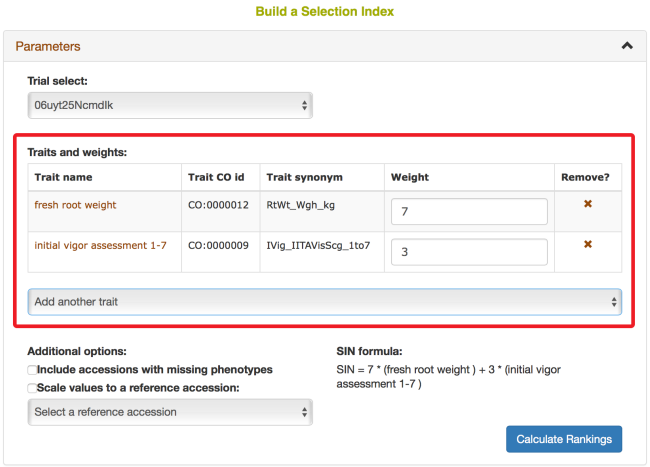
After you selected a trial, you can find traits that were assayed in that trial in the "Trait" box.



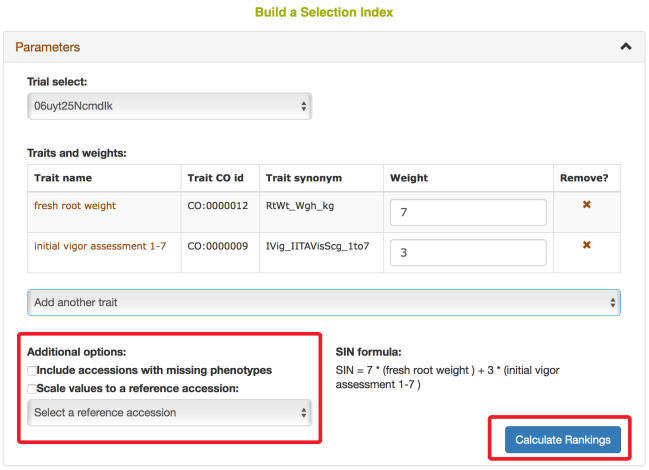
Selecting a trait that you want to include in the analysis will open a new dialogue showing the selected trait and a box that you can assign a "Weight" of that trait. After you are done, you can continue by selecting another trait by clicking on "Add another trait" link.



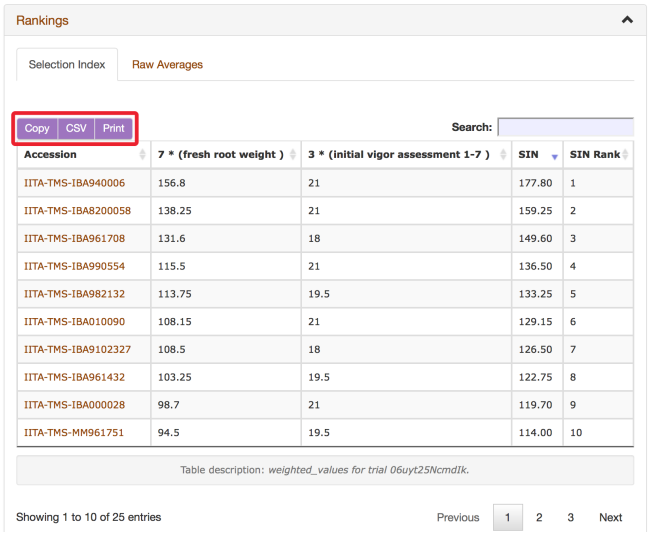
After you selected another trait, this page will automatically update information for you by showing all of the traits that you selected for the analysis.



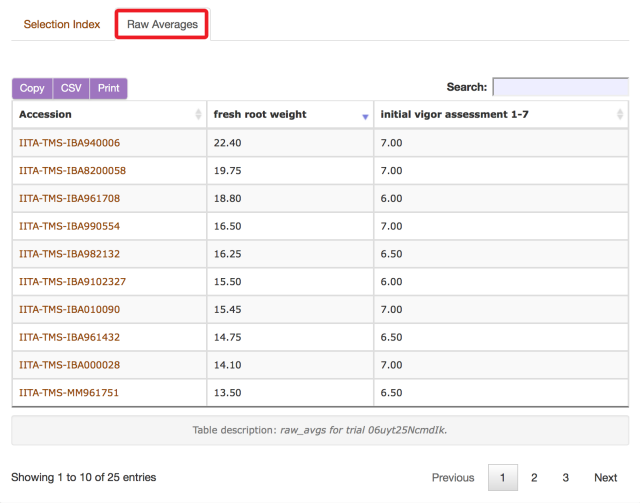
You also have options to choose a reference accession, choose to include accessions with missing phenotypes, scaling values to a reference accession. After you complete your setting, clicking on “Calculate Rankings”



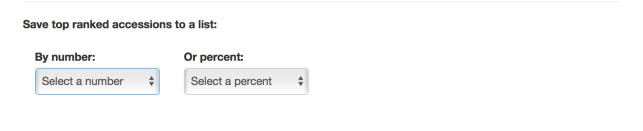
The Selection Index tool will generate rankings of accessions based on the information that you specified. You can copy the results to your system clipboard, convert the table data to CSV format, or print the data.



Clicking on "Raw Average" will display average values of the phenotypes of those ranked accessions.



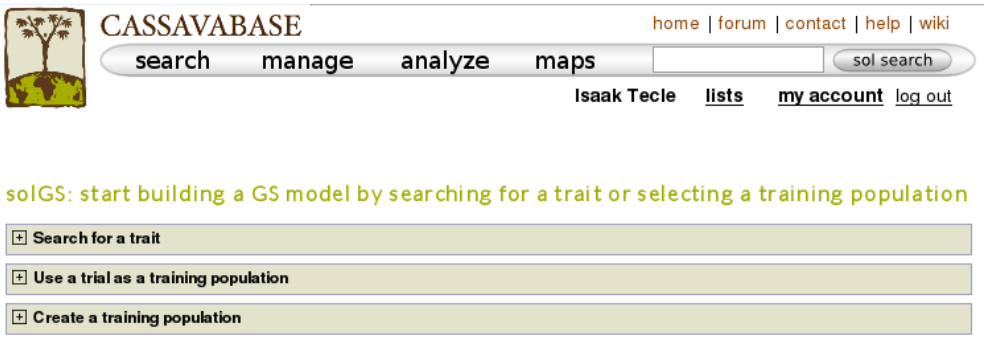
Selection Index tool also allows you to save top ranked accessions directly to "Lists". You can retrieve top ranked accessions by selecting a number or a percent.



25.2 Genomic Selection

The prediction of breeding values for a trait is a one step or two steps process, depending on what stage in your breeding cycle you are. The first step is to build a prediction model for a trait using a training population of clones with phenotype and genotype data. If you have yet to select parents for crossing for your first cycle of selection you can use the breeding values of the training population. If you are at later stages of your selection program, you need to do the second step which is applying the prediction model on your selection population. All clones in your training and selection populations must exist in the database.

To use the genomic selection tool, on *cassavabase.org*, select 'Genomic Selection' from the 'analyze' pull-down menu.



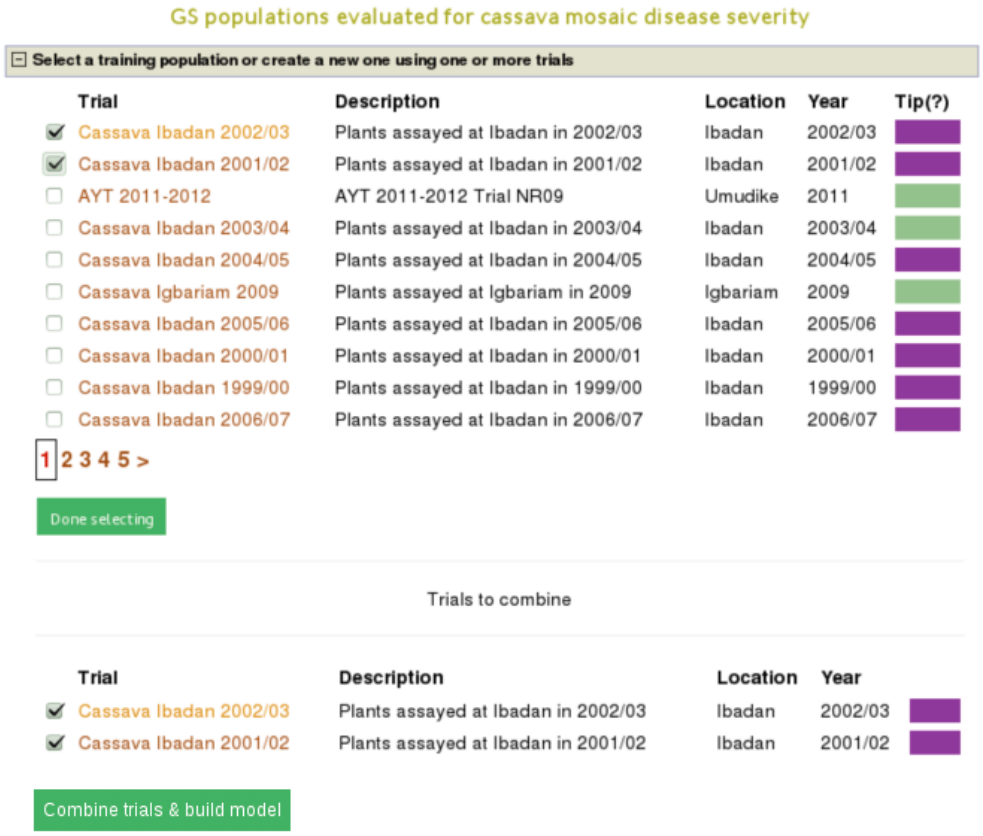
There are three ways to build a model for a trait.

25.2.1 Building a Model - Method 1:

One way to build a model is, using a trait name, to search for trials in which the trait was phenotyped and use a trial or a combination of trials to build a model for the trait. For example, if you search for 'mosaic disease severity, you will get a list of trials you can use as training populations.

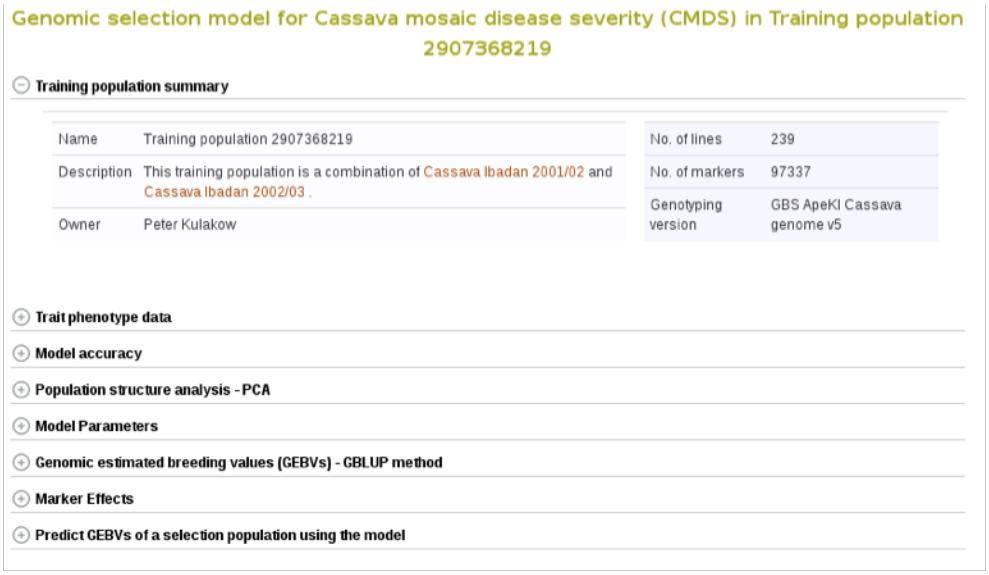
You will get a list of trials (as shown below) in which the trait of your inter-

ested was phenotyped. From the list, you can use a single trial as a training population or combine several trails to form a training population for the prediction model of the trait. Let's say, you want to create a training population using individuals from trials 'cassava ibadan 2001/02' and 'cassava ibadan 02/03' and build a model for 'cassava mosaic disease severity' using all clones from the training population.



Select the trials to combine (the same coloured), click 'done selecting', click the 'combine trials and build model' button, and you will get a model and its output for the trait. On the model detail page, you can view the description of input data used in the model, output from the model and search interface for selection populations the model you can apply to predict their breeding values. The description of the input data for the model includes the number of phenotyped clones, and the number of markers, scatter and frequency distribution plots for the phenotype data, relationship between the phenotype

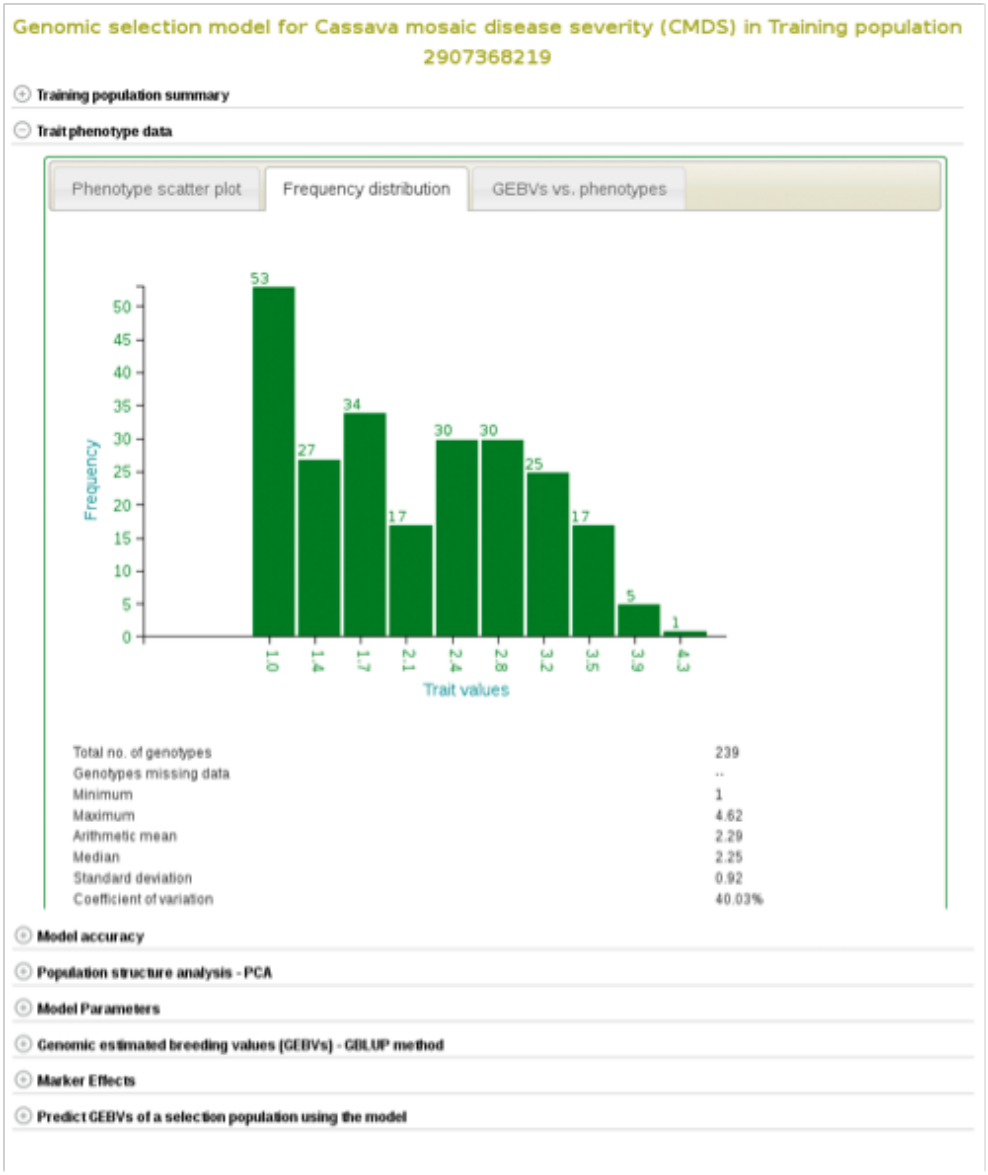
data and GEBVs, population structure. The model output includes model parameters, heritability of the trait , prediction accuracy, GEBVs of the individuals from the training population and marker effects.



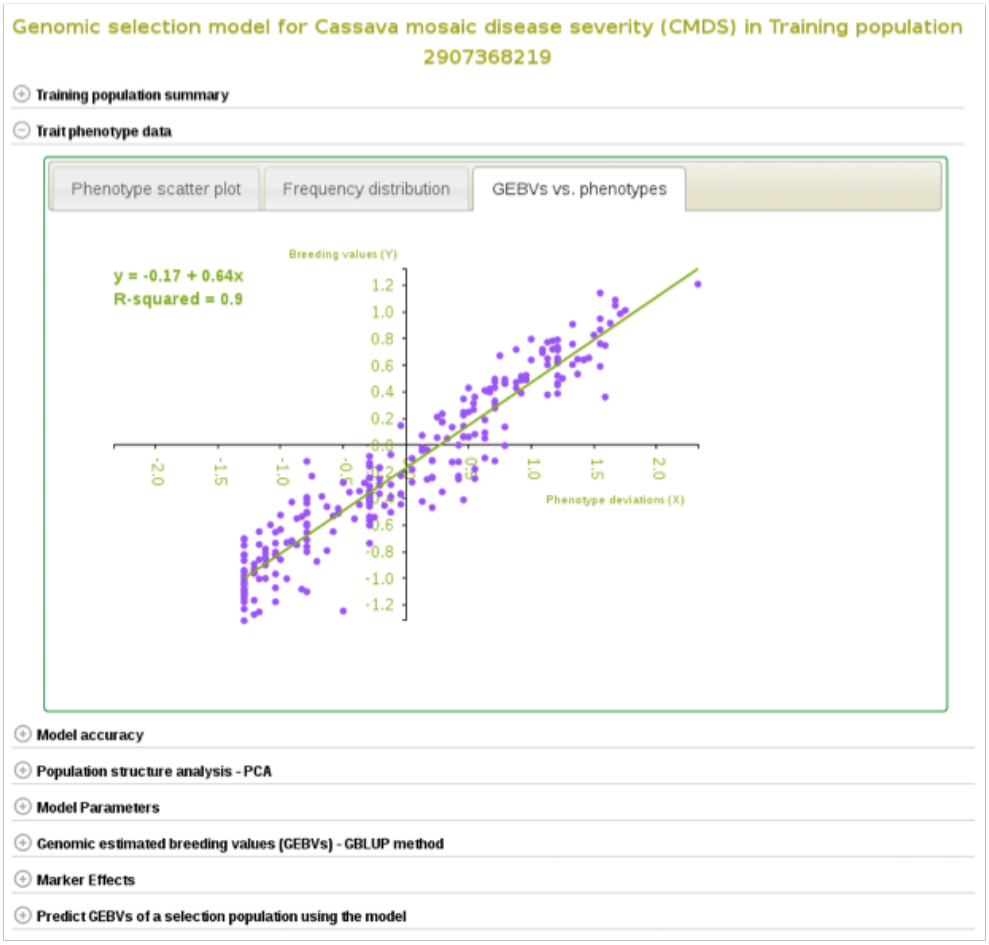
Expand each section to see detailed information.

If you expand the 'Trait phenotype data' section, you will find plots to explore the phenotype data used in the model. You can assess the phenotype data using a scatter and histogram plots and the descriptive statistics.

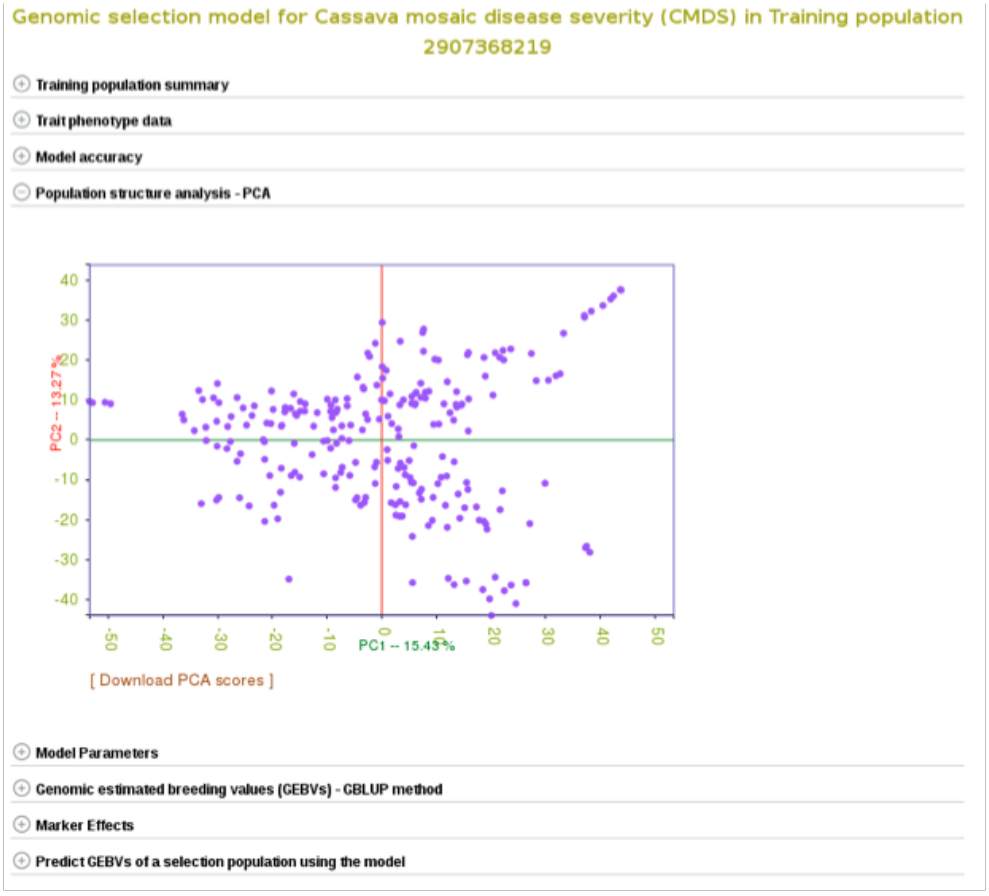
Genomic selection model for Cassava mosaic disease severity (CMDS) in Training population 2907368219
Training population summary
Trait phenotype data
Phenotype scatter plot
Frequency distribution
GEBVs vs. phenotypes
5.0
4.5
4.0
3.5
3.0
2.5
2.0
1.5
1.0
0.5
Phenotype values
Clones
Model accuracy
Population structure analysis - PCA
Model Parameters
Genomic estimated breeding values (GEBVs) - GBLUP method
Marker Effects
Predict GEBVs of a selection population using the model



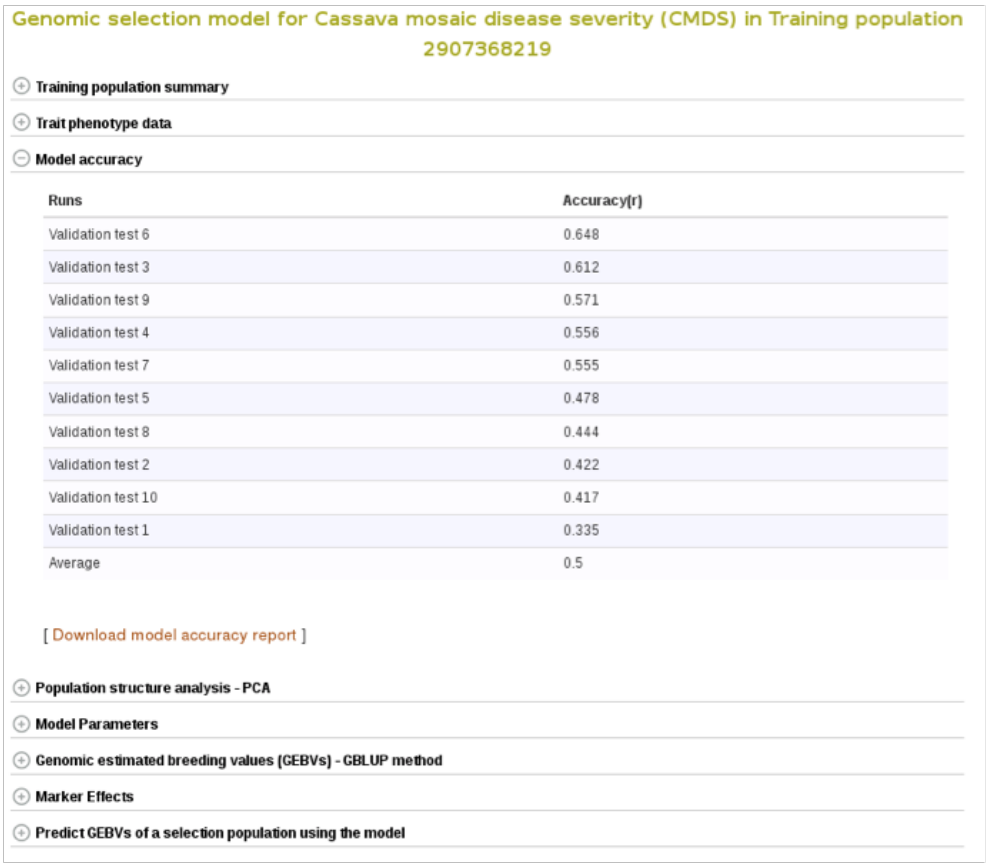
A regression line between observed phenotypes and GEBVs shows the relationship between the two.



You can also explore if there is any sub-clustering in the training population using PCA.



To check the model accuracy, a 10-fold cross-validation test, expand the 'model accuracy' section.



Genomic selection model for Cassava mosaic disease severity (CMDS) in Training population 2907368219

⊕ Training population summary

⊕ Trait phenotype data

⊖ Model accuracy

Runs	Accuracy(r)
Validation test 6	0.648
Validation test 3	0.612
Validation test 9	0.571
Validation test 4	0.556
Validation test 7	0.555
Validation test 5	0.478
Validation test 8	0.444
Validation test 2	0.422
Validation test 10	0.417
Validation test 1	0.335
Average	0.5

[Download model accuracy report]

⊕ Population structure analysis - PCA

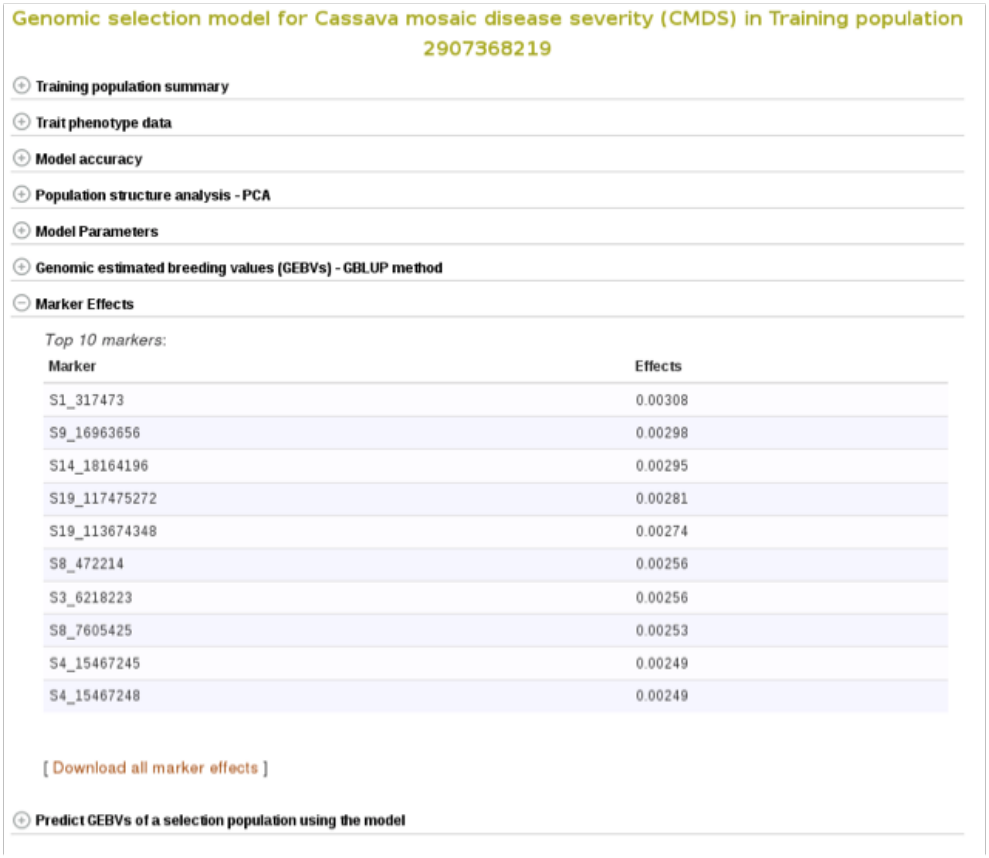
⊕ Model Parameters

⊕ Genomic estimated breeding values (GEBVs) - GBLUP method

⊕ Marker Effects

⊕ Predict GEBVs of a selection population using the model

Marker effects are also available for download. To do so, expanad the 'Marker Effects' section and click the 'Download all marker effects' link and you will get a tab delimited output to save on your computer.



Genomic selection model for Cassava mosaic disease severity (CMDS) in Training population 2907368219

- Training population summary
- Trait phenotype data
- Model accuracy
- Population structure analysis - PCA
- Model Parameters
- Genomic estimated breeding values (GEBVs) - GBLUP method
- Marker Effects

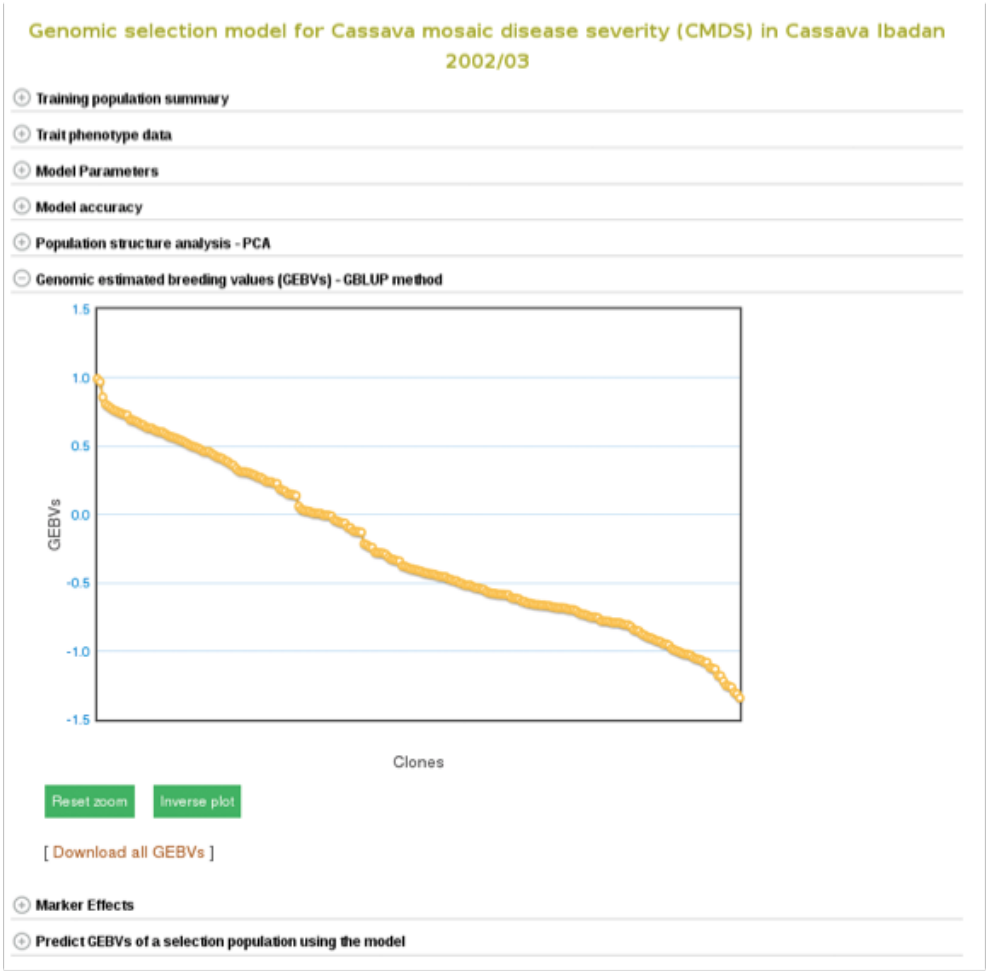
Top 10 markers:

Marker	Effects
S1_317473	0.00308
S9_16963656	0.00298
S14_18164196	0.00295
S19_117475272	0.00281
S19_113674348	0.00274
S8_472214	0.00256
S3_6218223	0.00256
S8_7605425	0.00253
S4_15467245	0.00249
S4_15467248	0.00249

[Download all marker effects]

- Predict GEBVs of a selection population using the model

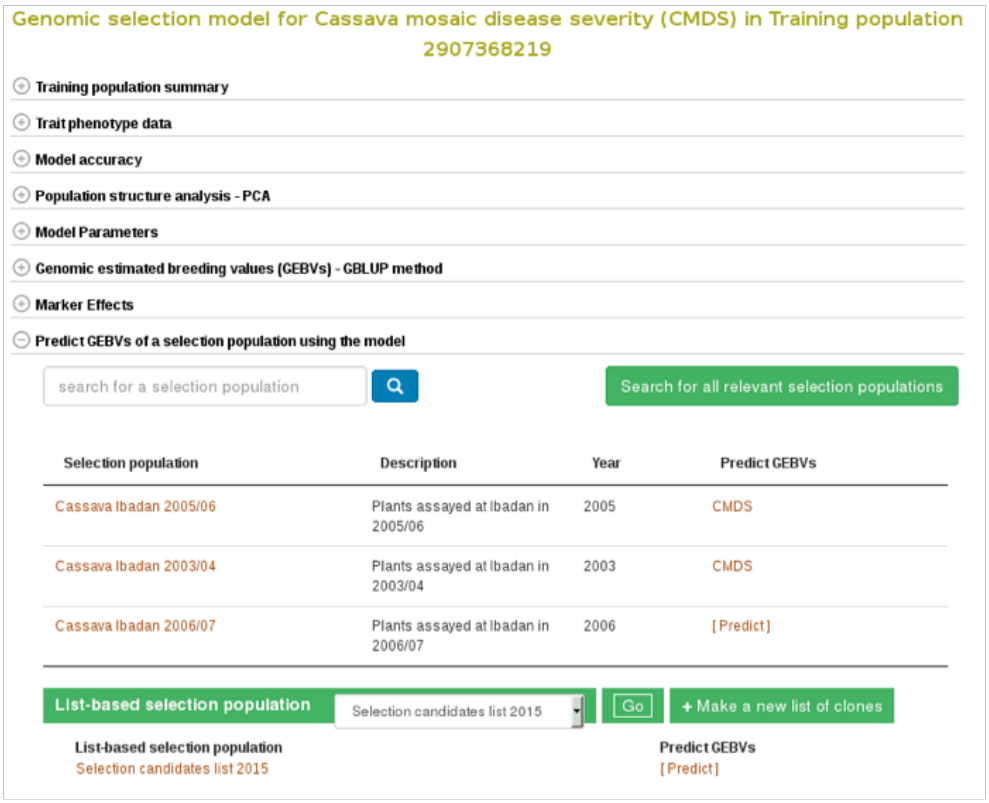
The breeding values of the individuals used in the training population are displayed graphically. Mousing over each data point displays the clone and its breeding value. To examine better, you can zoom in into the plot by selecting an area on the plot. You can download them also by following the 'Download all GEBVs' link.



Estimating breeding values in a selection population

If you already have a selection population (in the database), from the same model page, you can apply the model to the selection population and estimate breeding values for all the clones in the population. You can search for a selection population of clones in the database using the search interface or you can make a custom list of clones using the *list interface*. If you click the 'search for all relevant selection populations', you will see all relevant selection populations for that model. However, this option takes long time decause of the large set of populations in the database and the filtering. Therefore, the fastest way is to search for each of your selection populations by name. If you are logged in to the website you will also see a list of your custom set of

genotyped clones.



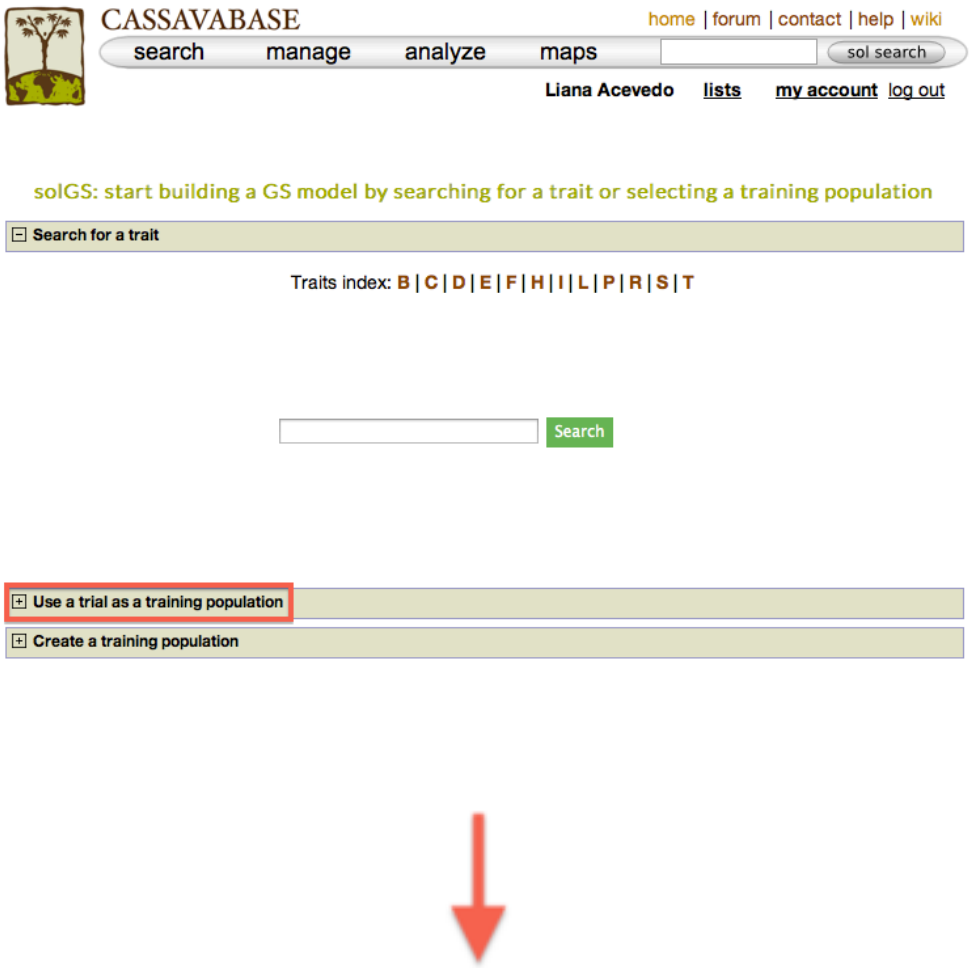
To apply the model to a selection population, simply click your population name or 'Predict Now' and you will get the predicted breeding values. When you see a name of (or acronym]) of the trait, follow the link and you will see an interactive plot of the breeding values and a link to download the breeding values of your selection population.

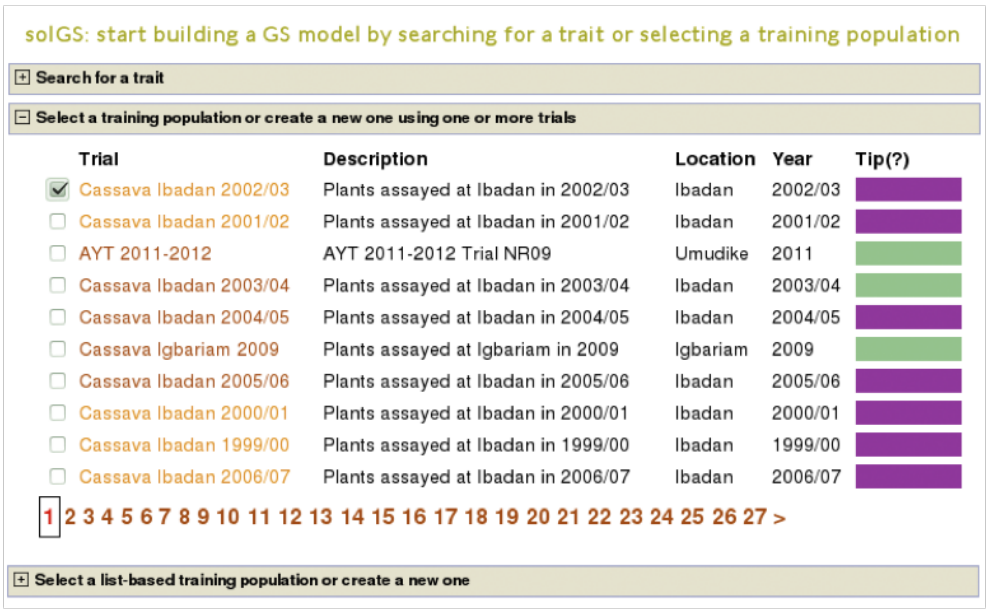
25.2.2 Building a Model - Method 2

Another way to build a model is by selecting a trial, instead of selecting and searching for a specific trait. This approach is useful when you know a particular trial that is relevant to the environment you are targeting to breed material for. This method allows you to build models and predict genomic estimated breeding values (GEBVs) for several traits within a single trial at once. You can also calculate selection index for your clones when GEBVs are

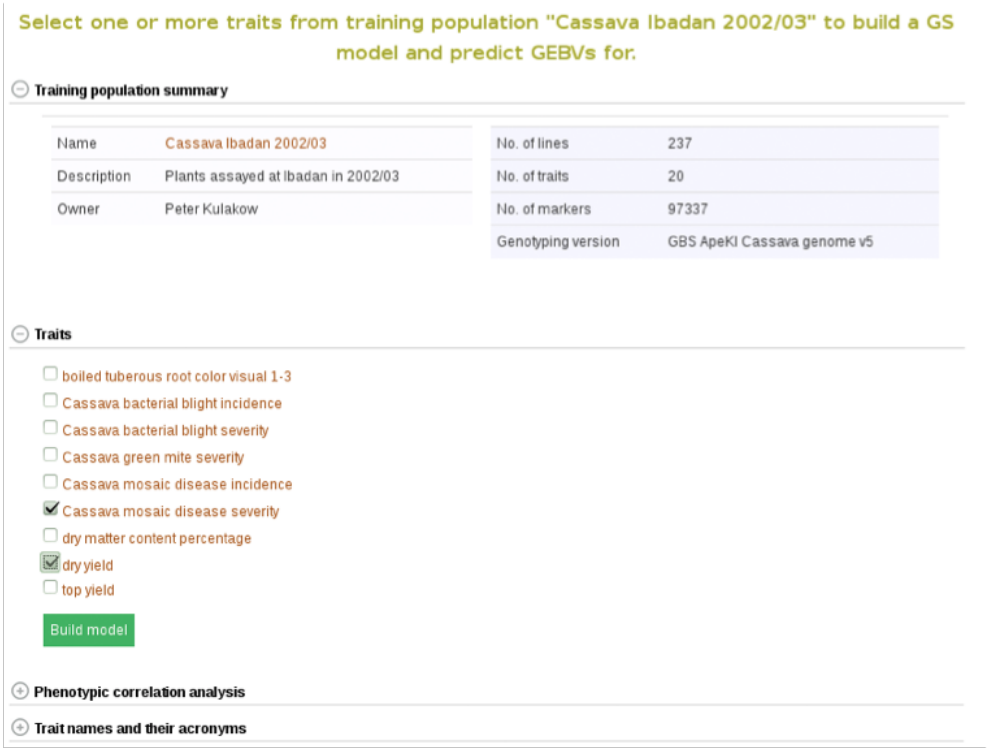
estimated for multiple traits.

To do this select the "Genomic Selection" link found under the "analyze" menu. This will take you to the same home page as used with Method 1. However, instead of entering information to search for in "Search for a trait", click on "Use a trait as a trial population". This will expand a new menu that will show all available trials.

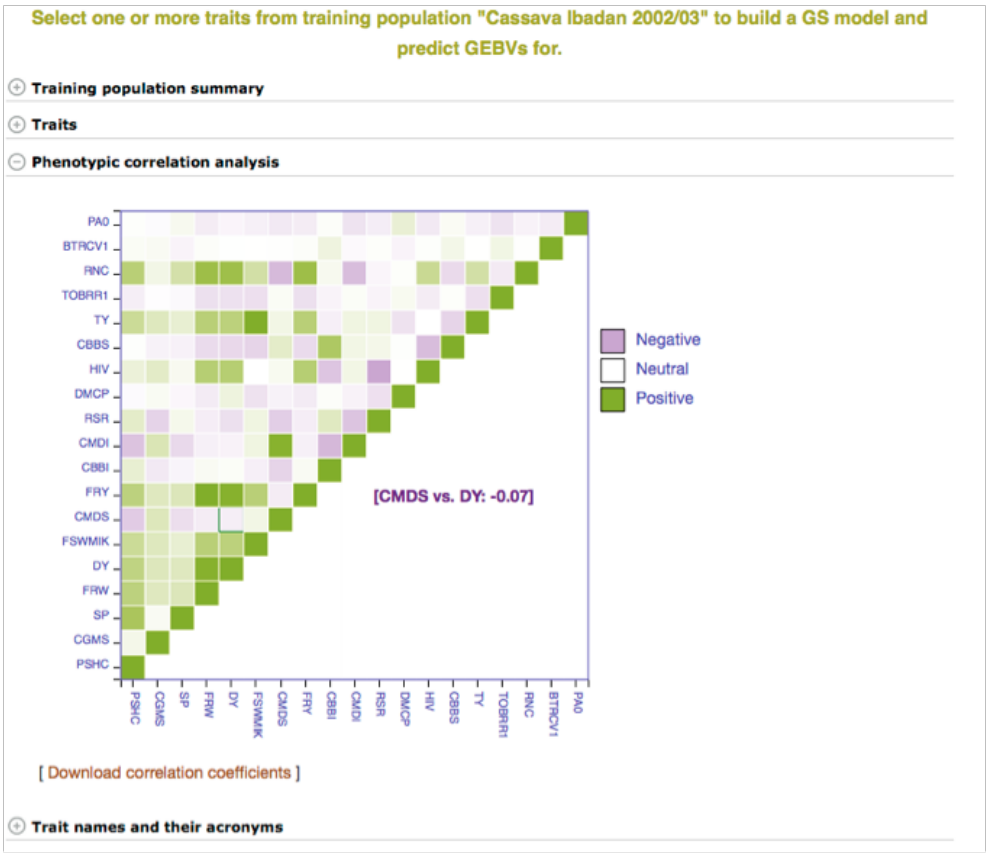




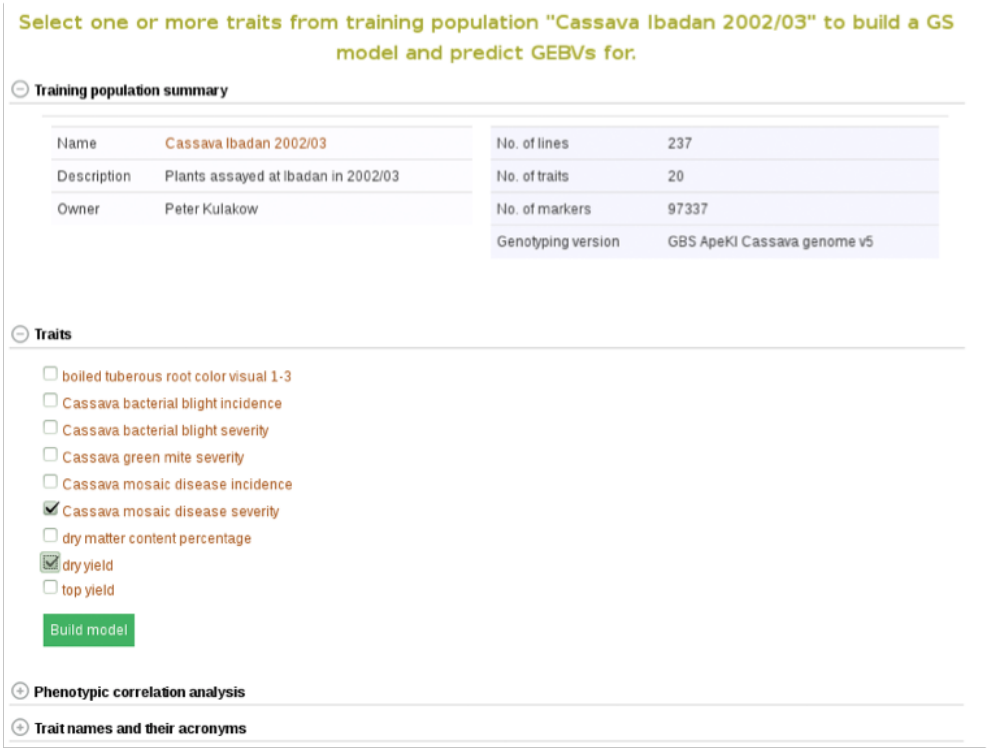
To begin creating the model, select the existing trial that you would like to use. In this example I will be using the trial and trait data from "Cassava Ibadan 2002/03" trial. Clicking on a trial will take you to a page where you can find information such as number of markers and number of phenotypes clones.



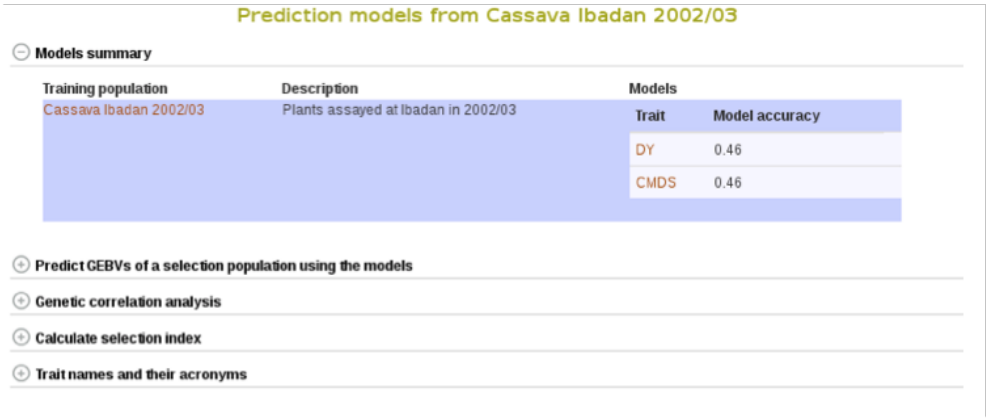
In addition to the number of phenotype clones and number of markers, the main page for the trial selected also has information and graphs on phenotypic correlation for all of the traits. By moving your cursor over the graph you can read the different values for correlation between two traits. A key with all of the trait names of the acronyms used can be found in the tab below the graph.



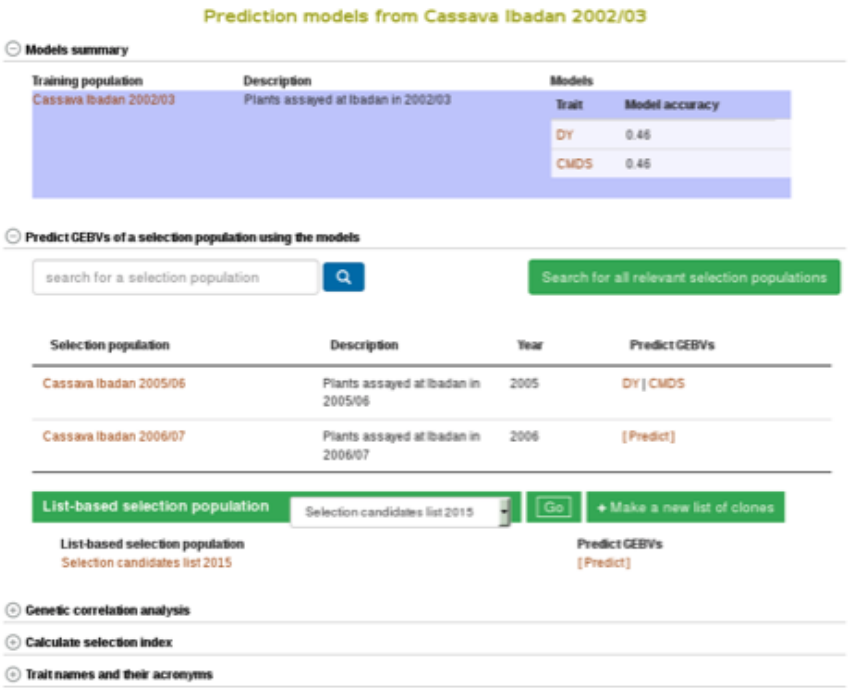
Below the "Training population summary" there is a tab for "Traits". Clicking on this tab will show all available traits for the specific trial. You can create a model by choosing one or multiple traits in the trial and clicking "Build Model". In this example, the traits for "cassava bacterial blight severity" and "cassava mosaic disease severity" have been selected.



Clicking on 'Build Model' will take you to a new page with the models outputs for the traits. Under the "Genomic Selection Model Output" tab you can view the model output and the model accuracy. Clicking on any of the traits will take you to a page with information about the model output on that individual trait within the trial. There you can view all of the trait information that was seen in more detail in *Method 1*.



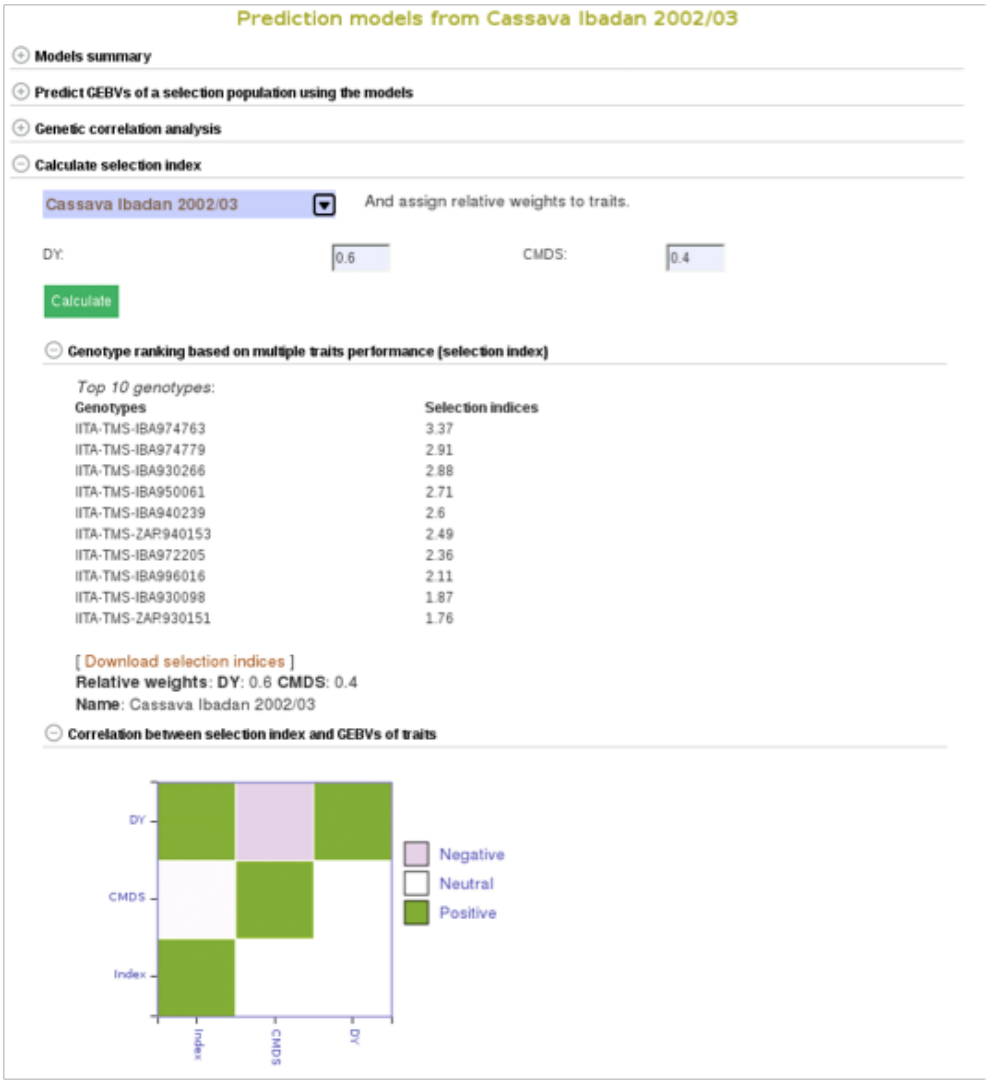
You can apply the models to simultaneously predict GEBVs for respective traits in a selection population by clicking on "Predict Now" or the name of the selection population. You can also apply the models to any set of genotyped clones that you can create using the 'lists' feature. For more information on lists, click *here*. Follow the link to the trait name to view and download the predicted GEBVs for the trait in a selection population.



To compare clones based on their performance on multiple traits, you can calculate selection indices using the form below. Choose from the pulldown menu the population with predicted GEBVs for the traits and assign relative weights for each trait. The relative weight of each trait must be between 0 - 1. 0 being of least weight and importance, not wanting to consider that particular trait in selecting a genotype and 1 being a trait that you give highest importance.

In this example we will be using the "Cassava Ibadan 2002/03" population and assigning values to each of the traits. Remember that there is a list of acronyms and trait names at the bottom of the page for reference. After entering whatever values you would like for each trait click on the "Calculate"

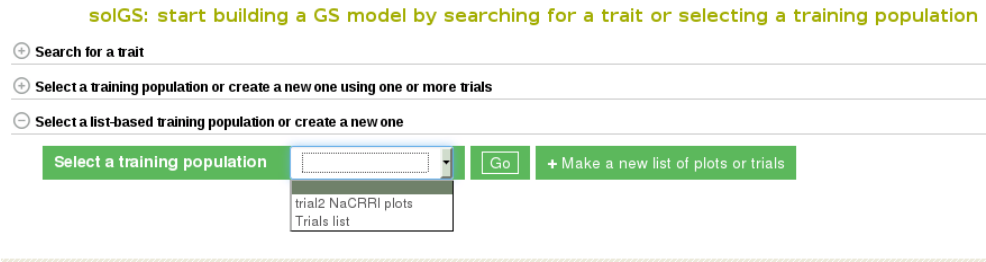
button to generate results. This will create a list of the top 10 genotypes that most closely match the criteria that you entered. The list will be displayed right below the "selection index" tab. This information can also be downloaded onto your computer by clicking on the "Download selection indices" link underneath the listed genotypes and selection indices.



25.2.3 Building a Model - Method 3

In addition to creating a model by searching for pre-existing traits or by preexisting trial name, models can also be created by using your own list of clones. This creates a model by using or creating a training population.

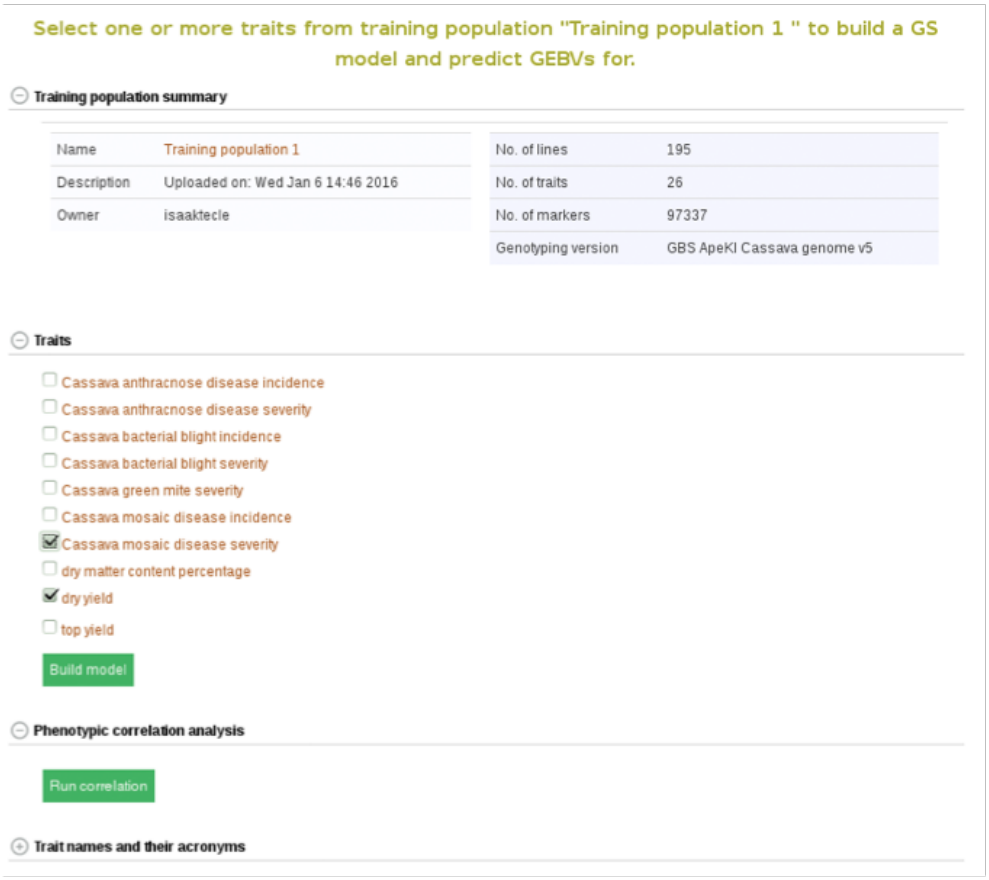
The page to use the third Method for creating a population model is the same as for the other two models. Select "Genomic Selection" from under the "analyze" menu of the main toolbar. This will take you to the Genomic Selection homepage and show you all three available methods to create a model. To see and use Method 3 scroll down and click on the tab labeled "Create a Training Population". This will open a set of tools that will allow you to use pre-existing lists or to create a new list.



Once the "Create a Training Population" tab is opened you have the option to use a pre-existing list or create new one. To learn how to create a list, click *here*. The "Make a new list of plots" link will take you directly to the Search Wizard that is usually used to create lists.

Please note: the only lists that can be used in Method 3 to create a model are lists of plots and trials. If the pre-existing list is not of plots or trials (for example, traits, or locations) it will not show up and cannot be used as a training population. When you create you use a list of trials, the trials data will be combined to create a training data set.

To use your custom list of plots or trials as a training population, select the list and click "Go". This will take you to a detail page for the training population.



From here on you can build models and predict breeding values as described in *Method 2*.

25.3 Genome Browsing

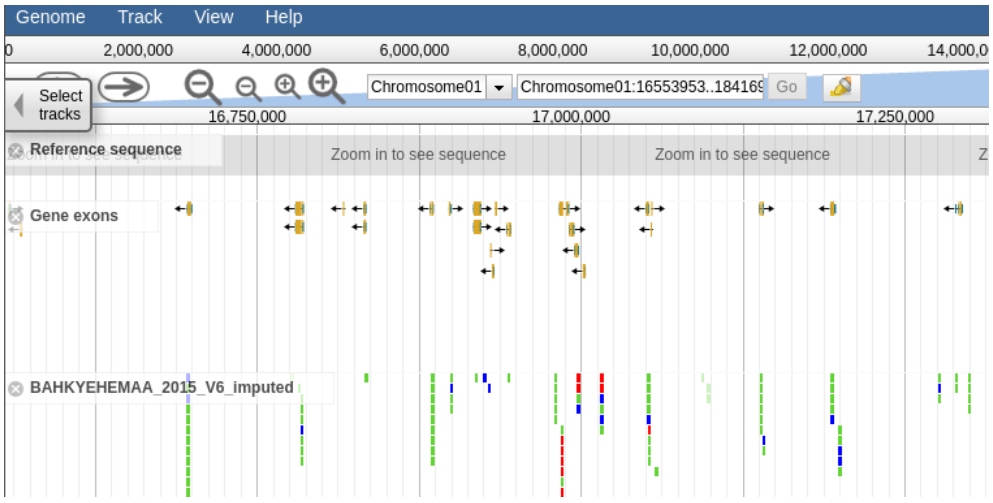
There are two ways to evaluate genotype information within the browser, from an accession detail page or a trial detail page.

25.3.1 Browsing Genotype data by Accession

If you are interested in browsing genotype information for a single accession, for example 'BAHKYEHEMAA', navigate to the accession detail page.

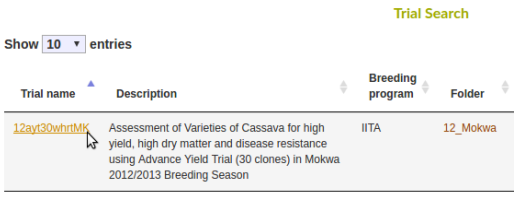
Near the bottom of the detail page is a collapsible section called "Accession Jbrowse".

This section will contain a link to the accession jbrowse page if the necessary genotype data is available. Clicking the link should take you to a page that looks like this, a which point you can browsre the genotype data in the form of a vcf track aligned to the latest build of the genome.

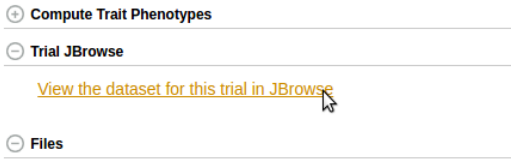


25.3.2 Browsing Genotype data by Trial

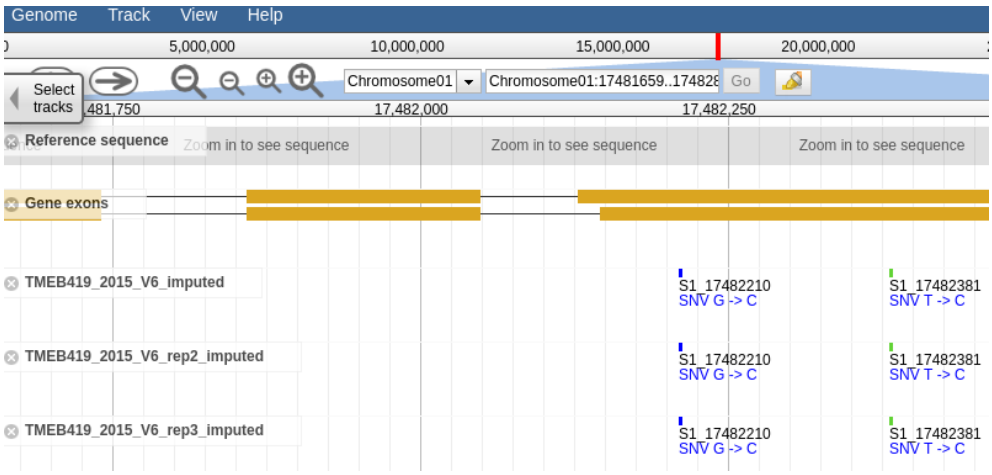
If you are interested in browsing genotype information for the accessions within a given trial, navigate to the trial detail page.



Halfway down the page is a collapsible section called "Trial Jbrowse". This section will contain a link to the trial jbrowse page if the necessary genotype data for at least two accessions planted in the trial is available.



Clicking the link should take you to a page that looks like this, a which point you can browse the genotype data in the form of vcf tracks aligned to the latest build of the genome.



25.4 Principal Component Analysis (PCA)

Principal component analysis helps estimate and visualize if there is sub-grouping of individuals within a dataset based on a number of variables. Currently, you can use marker data to run PCA on datasets.

You can run PCA from multiple places on the website. To do PCA on

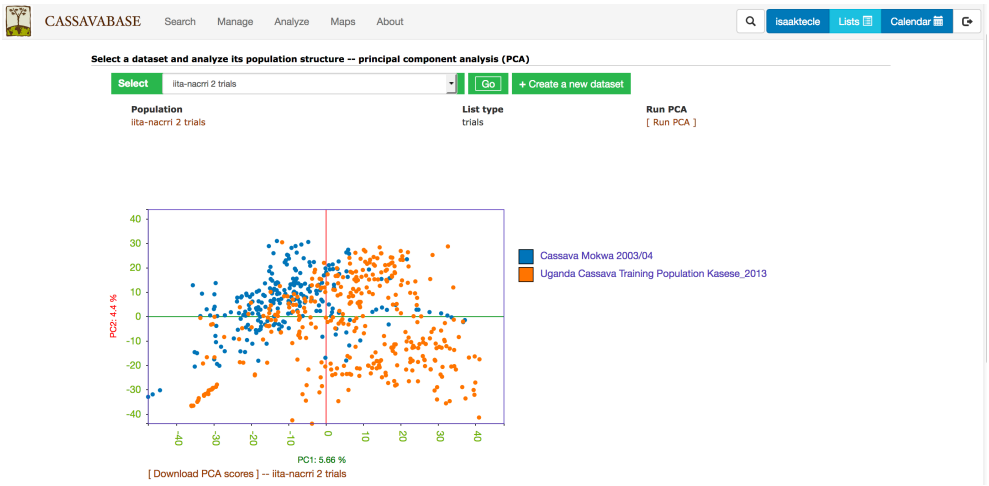
(1) individuals from a trial, go to the trial detail page and find the PCA tool under the 'Analysis tools' section.

(2) individuals from a training population you used in a GS modeling, do your modeling and find the PCA tool in the model output page.

(3) individuals in a training population and selection population you applied the training model, do your modeling, apply the model on the selection population and find the PCA tool on the selection population prediction output page.

(4) individuals in a list of accessions you created, for example using the search wizard, go to the 'Analyze' menu and select the 'Population Structure', select your list of individuals and run PCA.

(5) individuals from multiple trials, create a list of the trials using the search wizard, go to the 'Analyze' menu and select the 'Population Structure', select your list of trials and run PCA.



With all the options, you will get a interactive plot of the two PCs (shown below) that explain the largest variance. Point the cursor at any data point and you will see the individual name with its corresponding PCs scores. By clicking the 'Download all PCs', you can also download the 10 PCs scores in the text format.

25.5 ANOVA

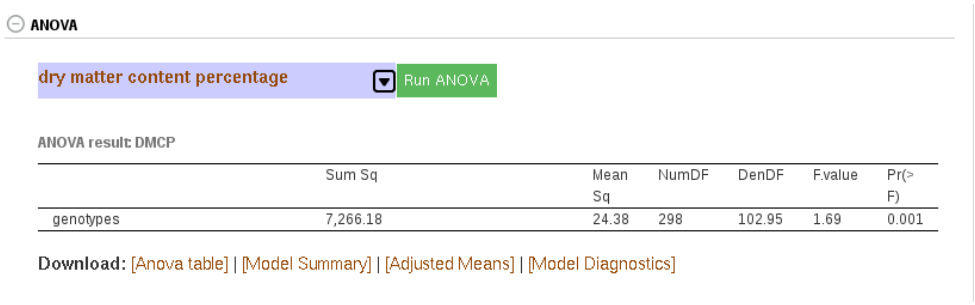
Currently, ANOVA is implemented for a single trial (single year and single location). You can do ANOVA for RCBD, CRD, Alpha and Augmented trial designs. ANOVA is done using linear mixed effects model, where the genotypes is fixed effect and the replications and blocks are random effects. Fixed effect significance level is computed using 'lmer' from 'lmeTest' R package.

You can do ANOVA from two places: trial detail and training population detail. In both cases, if the phenotype data was from the supported trial designs,

– Go to the ANOVA section down in the trial or training population page

– Select the trait of you want to perform ANOVA

– Click the 'Run ANOVA' and wait for the result



25.6 Clustering (K-Means, Hierarchical)

The K-Means method allows you to partition a dataset into groups (K number). The hierarchical clustering, agglomerative, allows you to explore underlying similarity and visualize in a tree structure (dendrogram) the different levels of similarities (clusters) among samples. You can do clustering based on marker data, phenotype data and GEBVs. When you use phenotype data, first clone averages for each trait are calculated. Both methods use Euclidean distance as a measure of similarity. For the hierachical clustering, the complete-linkage (farthest neighbour) method is used to link up clusters.

There are three pathways to using this tool.

(1) When you have data in the form of a list or dataset from the search wizard:

(A) – go to the 'Analyze' menu and select the clustering option

(B) – make sure you are logged in

(C) – Select the relevant genotyping protocol, if you are clustering using genotype data

(D) – select your list or dataset, click 'Go'

(E) – select clustering type

(F) – select the data type to use

(G) – If you are running K-Means clustering, provide the number of partitions (K). If left blank it will partition the data set into optimal numbers for the dataset.

(H) – click the 'Run Cluster' and wait for the analysis to finish or queue the request and wait for an email with the analysis result.

(I) – You can download the outputs following the download links.

(2) From the trial detail page:

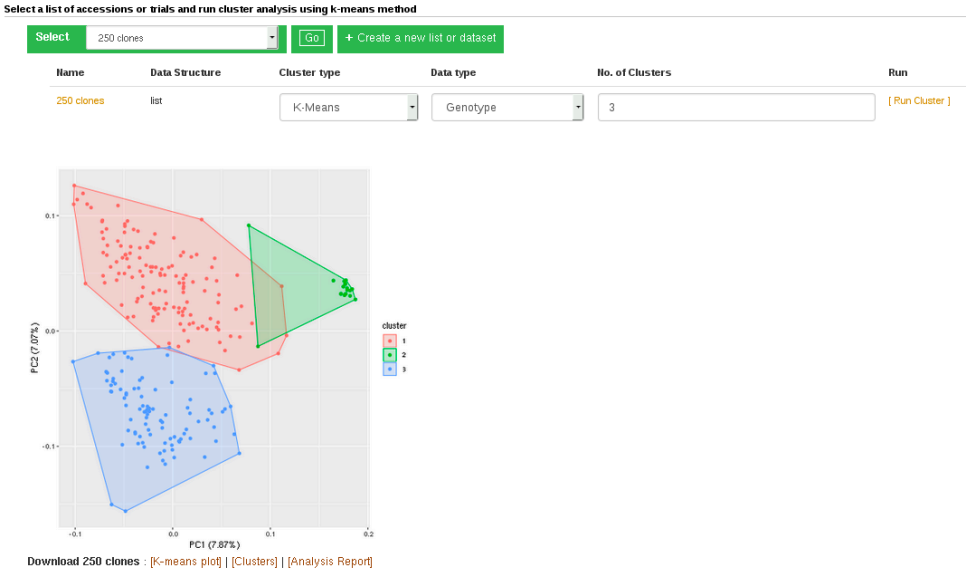
(A) – Go to the 'Analysis Tools' section

(B) – Follow steps D to G in (1)

(3) In the solGS pipeline:

(A) – Once you you are in a model output put page, you will see a section where you can do clustering in the same way as above (option 2).

K-Means clustering:



Hierarchical clustering:

Download 34 clones : Dendrogram | Newick tree format | Analysis Report

25.7 Genetic Gain

You can check for genetic gain by comparing the the GEBVs of a training and a selection population. You can do this in the solGS pipepline once you build a model and apply the model to predict the GEBVs of a selection population. Once at that stage, you will see a section 'Check Genetic Gain'. Select a selection population to compare with the training population and click the 'Check Genetic Gain' button. The genetic gain will be visualized in boxplots. You can download the boxplot(s) as well as the GEBVs data used for the plot(s).

25.8 Kinship and Inbreeding Coefficients

This tool allows you to estimate genetic relatedness between a pair of individuals (kinship), homozygousity across loci in an individual (inbreeding coefficient), and genetic similarity of an individual relative to the rest of the population (averge kinship).

There are three pathways to using this tool.

(1) When you have a list or dataset clones, created from the search wizard:

(A) – go to the 'Analyze' menu and select the kinship and inbreeding

(B) – make sure you are logged in

(C) – Select the genotypic protocol for the marker data

(D) – select your list or dataset of clones, click 'Go'

(E) – click the 'Run Kinship' and wait for the analysis to finish, depending on the data size this may take minutes. You can choose to submit the analysis and wait for an email notice to view the results or wait for it to complete.

(F) – You can download the output following the download links.

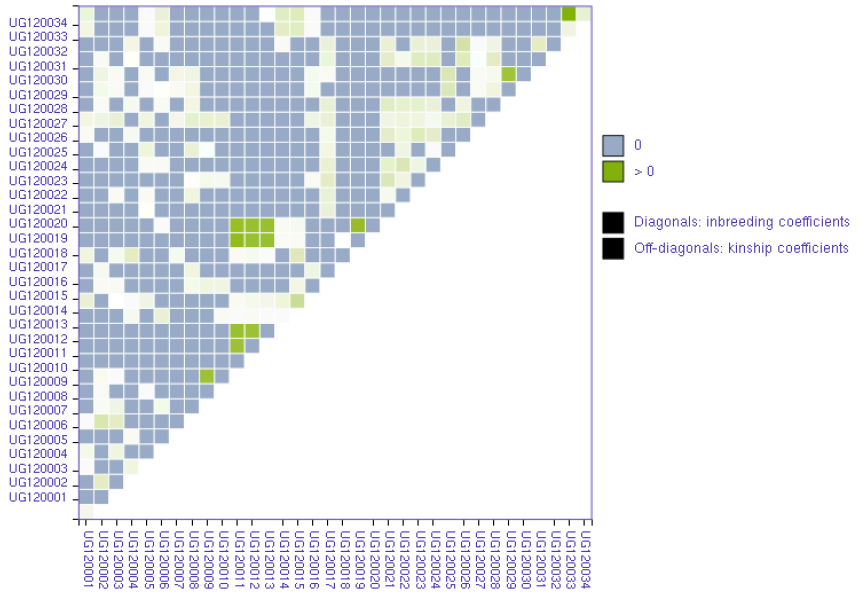
(2) From the trial detail page:

(A) – Go to the 'Analysis Tools' section

(B) – Follow steps C to G in (1)

(3) In the solGS pipeline:

(A) – Once you you are in a model output put page, scroll down to the 'Kinship and Inbreeding' section and run kinship.



Download: 34 clones Kinship matrix | Average kinship | Inbreeding coefficients

25.9 Creating Crossing Groups

If you calculate selection index based on GEBVs of multiple traits, and you want to select a certain proportion of the indexed individuals (e.g. top 10%, or bottom 10%) and then you want to partition the selected individuals into a number of groups based on their genotypes, you can use the k-means clustering method.

The procedure is:

(1) predict GEBVs for 2 or more traits

(2) In the models output page, calculate selection indices. Note the name of the selection index data.

(3) Go to the clustering section,

– select the selection index data,

– select 'K-means',

– select 'Genotype',

– in the K-numbers textbox, fill in the number of groups you want to create,

– in the selection proportion textbox, fill in the proportion of the indexed individuals you want to select, e.g. for the top 15 percent, 15. if you wish to select bottom performing, prefix the number with minus sign (e.g. -15)

– then run cluster and wait for the result.

Calculate selection index

Kasese solgs trial And assign relative weights to traits.

DMCP 2 FRW 3

Calculate

Genotype ranking based on multiple traits performance (selection index)

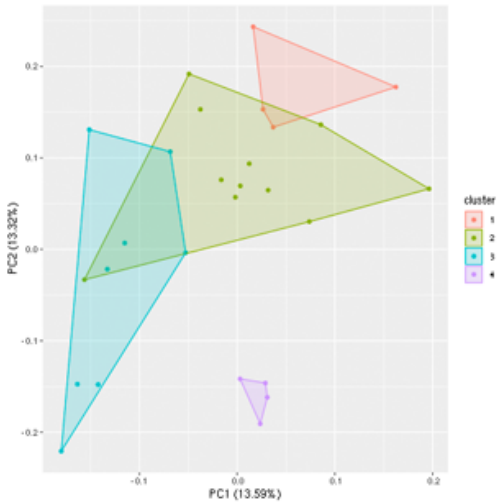
Top 10 genotypes:

Genotypes	Selection indices
UG120221	19.43
UG120001	15.56
UG120193	14.72
UG120071	14.3
UG120057	11.43
UG120072	11.2
UG120006	9.97
UG120264	9.88
UG120278	9.49
UG120184	9.35

Download selection indices | **Index Name:** 139-DMCP-2-FRW-3
Population name: Kasese solgs trial | **Relative weights: DMCP**: 2 **FRW**: 3

Clustering

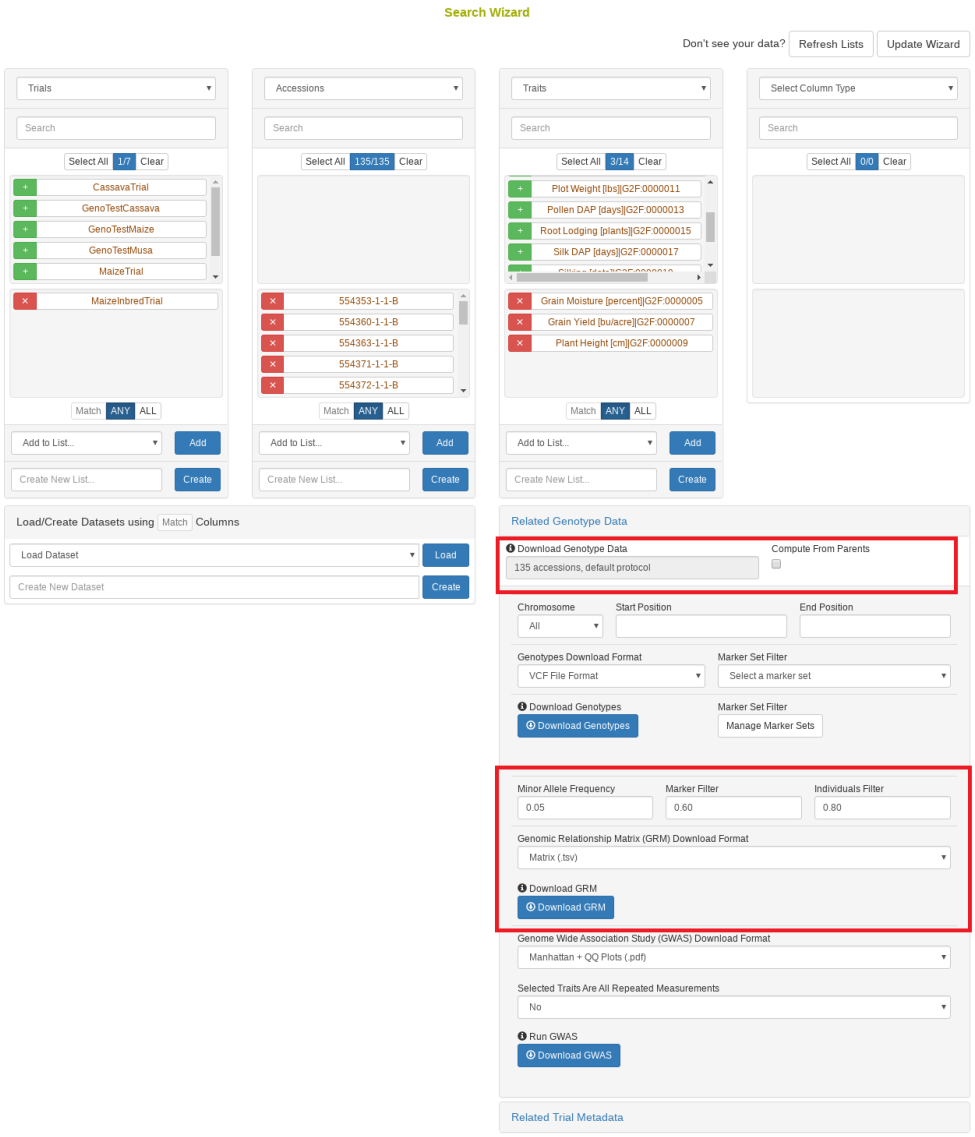
139-DMCP-2-FRW-3 K-Means Genotype 4 15 Run



Download 139-DMCP-2-FRW-3 : [K-means plot] | [Clusters] | [Analysis Report]

25.10 Search Wizard Genomic Relationship Matrix (GRM) Download

The genomic relationship matrix (GRM) is useful for understanding underlying structure in your population. Breedbase can compute the GRM using rrBLUP. First, select accessions in the search wizard and optionally select a genotyping protocol. If no genotyping protocol is selected, the default genotyping protocol in your system is used (as defined in sgn_local.conf). Specify the minor allele frequency, missing marker data, and missing individuals data filters to apply. The GRM can be returned in a matrix format (.tsv) which shows all pairwise relationships between the selected accessions and is useful for visualization; alternatively, the GRM can be returned in a three-column format (.tsv) which is useful for programs like ASReml outside of Breedbase. The GRM can also be returned as a simple correlation heatmap image (.pdf). The GRM can be computed from parents of the selected accessions granted the parents were genotyped, by clicking the checkbox "compute from parents"; this is useful for programs where parental lines are genotyped and then hybrids are created and evaluated in the field.

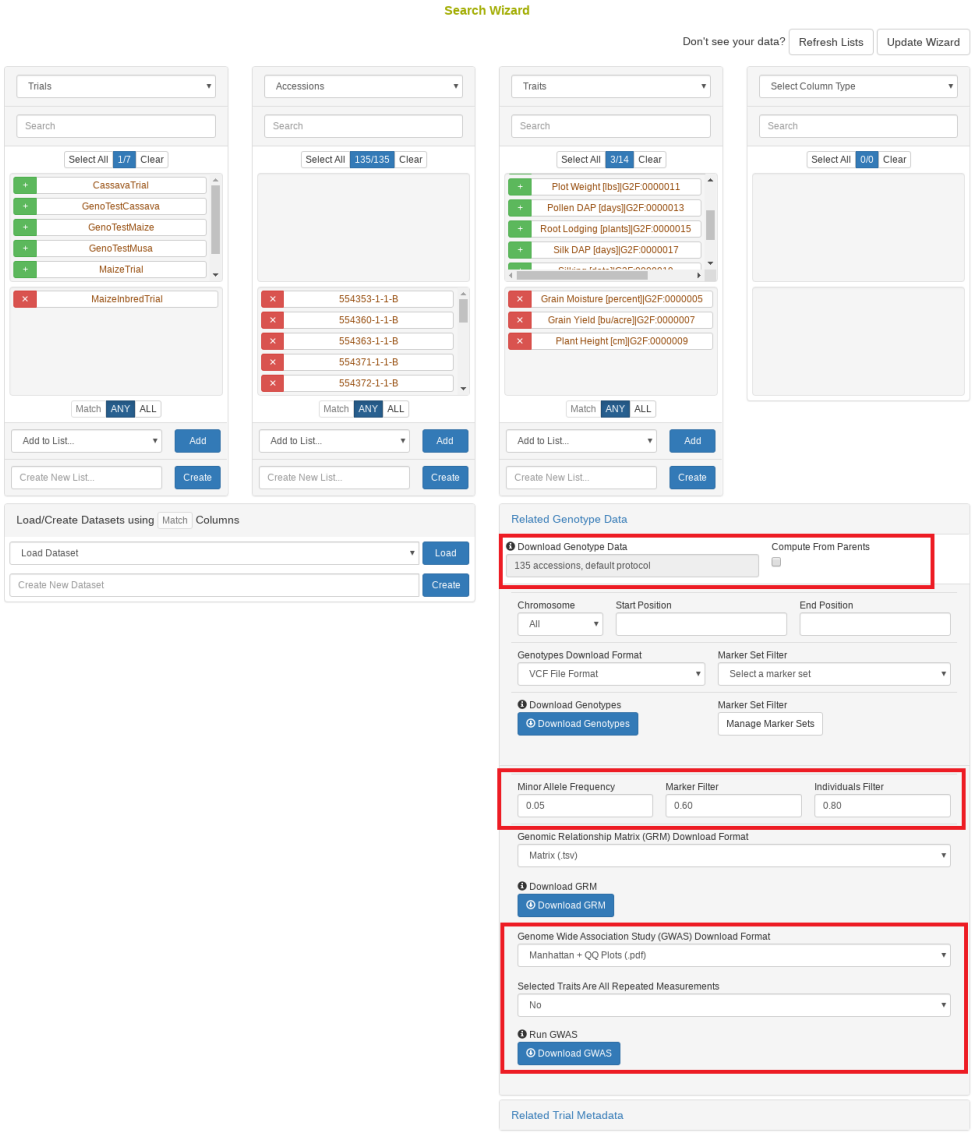


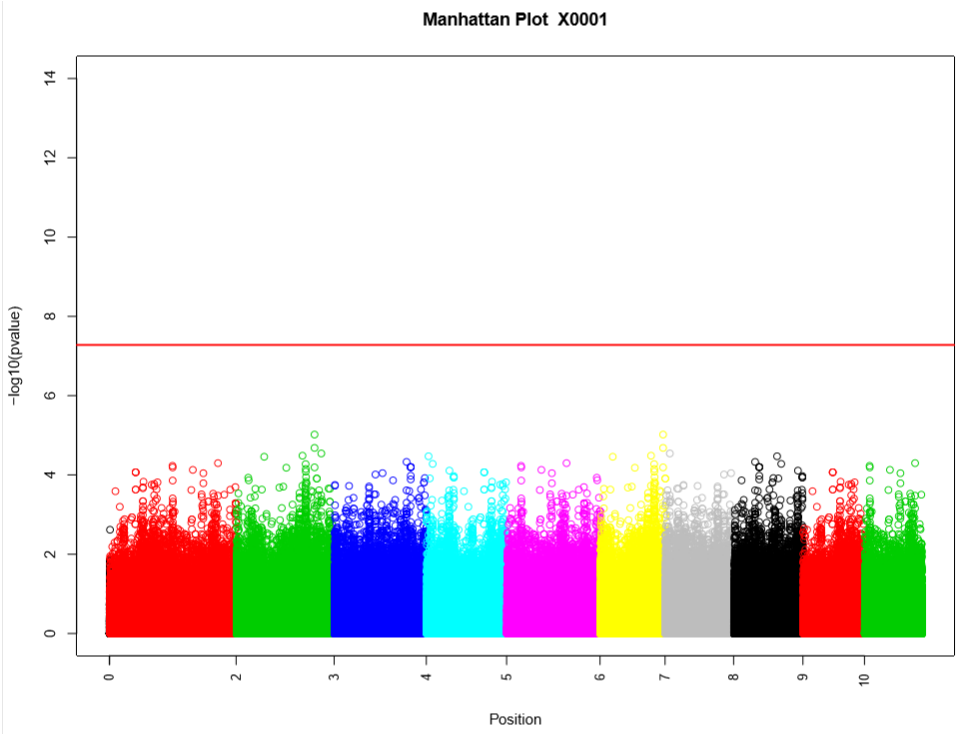
25.11 Search Wizard Genome Wide Association Study (GWAS)

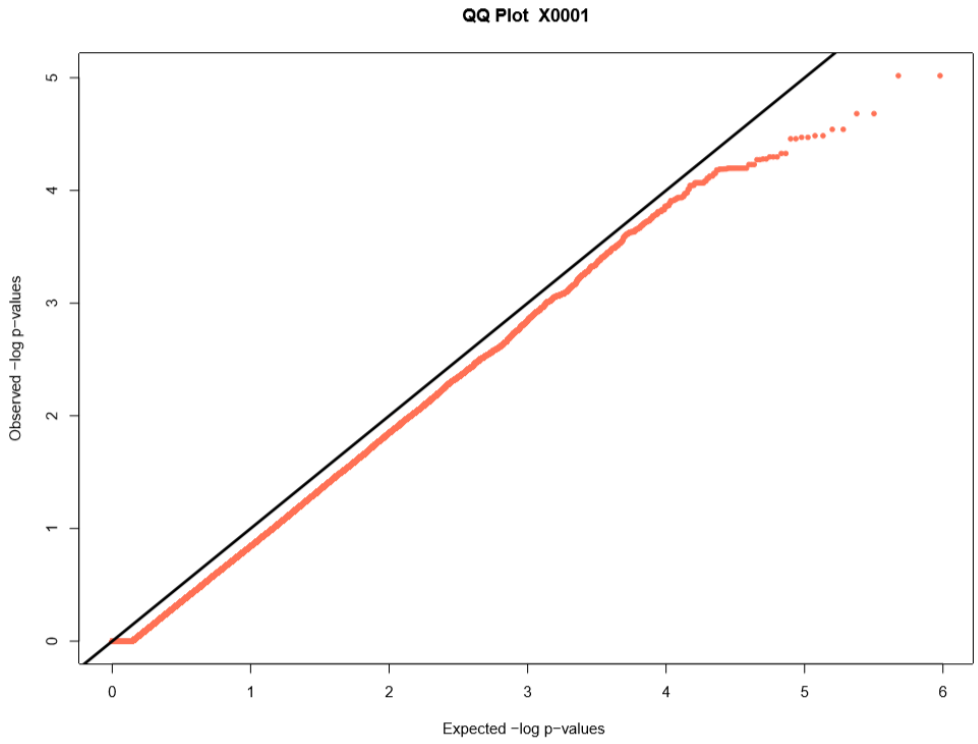
Performing a genome wide association study (GWAS) can determine genotypic markers which are significantly correlated to phenotypic traits. Breedbase can compute GWAS using rrBLUP. First, select accessions and trait(s)

in the search wizard, and optionally select a genotyping protocol. If no genotyping protocol is selected, the default genotyping protocol in your system is used (as defined in sgn_local.conf). Several traits can be selected in the search wizard; if the traits are not to be treated as repeated measurements then select 'no' in the select box and this will tell Breedbase to return GWAS results independently for the selected traits. If the selected traits are indeed all repeated measurements then select 'yes' in the select box and Breedbase will return as single GWAS analysis across all the phenotypic records. Specify the minor allele frequency, missing marker data, and missing individuals data filters to apply. GWAS results can be returned in a tabular format (.tsv) where the -log10(p-values) for the selected traits are returned; alternatively, the GWAS results can be returned as Manhattan and QQ plots for the selected traits. The GWAS can be computed from parents of the selected accessions granted the parents were genotyped, by clicking the checkbox "compute from parents"; this is useful for programs where parental lines are genotyped and then hybrids are created and evaluated in the field.

The GWAS will filter the data by the input MAF and missing data filters provided. After filtering the data is imputed using an 'EM' method in rrBLUP. The Kinship matrix (GRM) is computed from the imputed genotypic data and used in the GWAS model. The GWAS uses fixed effects for different field trials and replicates in the phenotypic data.

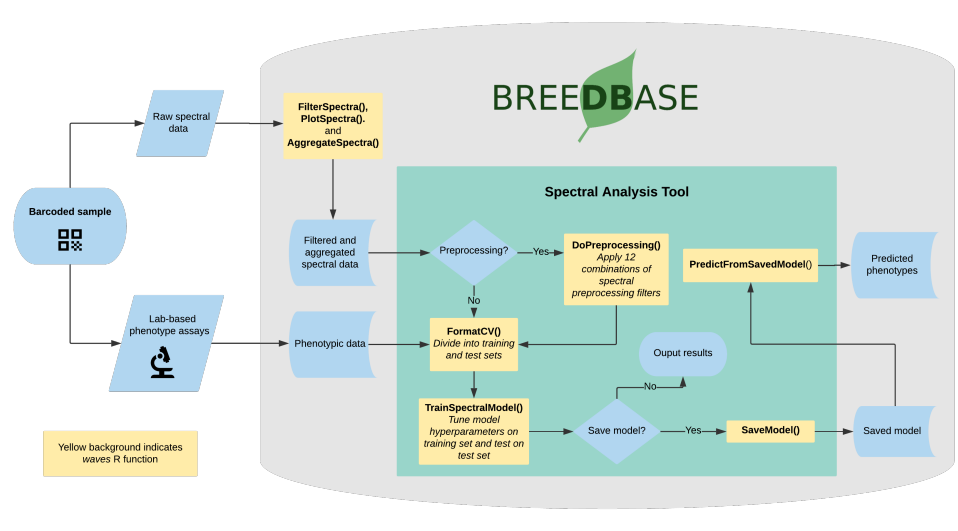

Search Wizard
Don't see your data?
Refresh Lists
Update Wizard
Trials
Search
Select All 1/7 Clear
CassavaTrial
GenoTestCassava
GenoTestMaize
GenoTestMusa
MaizeTrial
MaizeInbredTrial
Match ANY ALL
Add to List...
Add
Create New List...
Create
Accessions
Search
Select All 135/135 Clear
554353-1-1-B
554360-1-1-B
554363-1-1-B
554371-1-1-B
554372-1-1-B
Match ANY ALL
Add to List...
Add
Create New List...
Create
Traits
Search
Select All 3/14 Clear
Plot Weight [lbs]|G2F:0000011
Pollen DAP [days]|G2F:0000013
Root Lodging [plants]|G2F:0000015
Silk DAP [days]|G2F:0000017
Grain Moisture [percent]|G2F:0000005
Grain Yield [bu/acre]|G2F:0000007
Plant Height [cm]|G2F:0000009
Match ANY ALL
Add to List...
Add
Create New List...
Create
Select Column Type
Search
Select All 0/0 Clear
Load/Create Datasets using Match Columns
Load Dataset
Load
Create New Dataset
Create
Related Genotype Data
Download Genotype Data
135 accessions, default protocol
Compute From Parents
Chromosome
All
Start Position
End Position
Genotypes Download Format
VCF File Format
Marker Set Filter
Select a marker set
Download Genotypes
Download Genotypes
Marker Set Filter
Manage Marker Sets
Minor Allele Frequency
0.05
Marker Filter
0.60
Individuals Filter
0.80
Genomic Relationship Matrix (GRM) Download Format
Matrix (.tsv)
Download GRM
Download GRM
Genome Wide Association Study (GWAS) Download Format
Manhattan + QQ Plots (.pdf)
Selected Traits Are All Repeated Measurements
No
Run GWAS
Download GWAS
Related Trial Metadata


Manhattan Plot X0001
−log10(pvalue)
0
2
4
6
8
10
12
14
0
2
3
4
5
6
7
8
9
10
Position



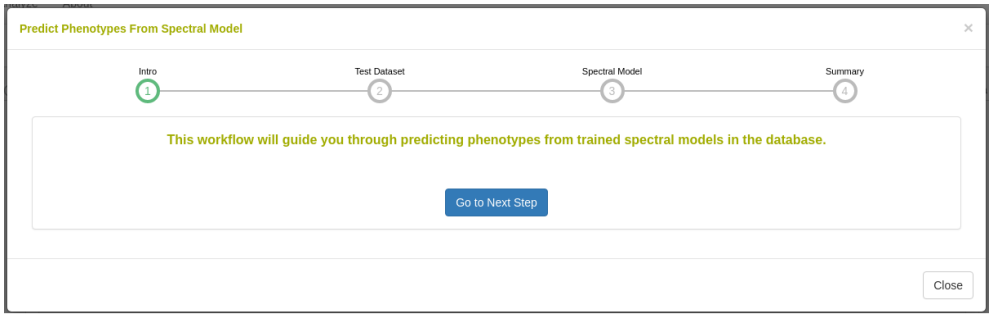
25.12 Spectral Analysis

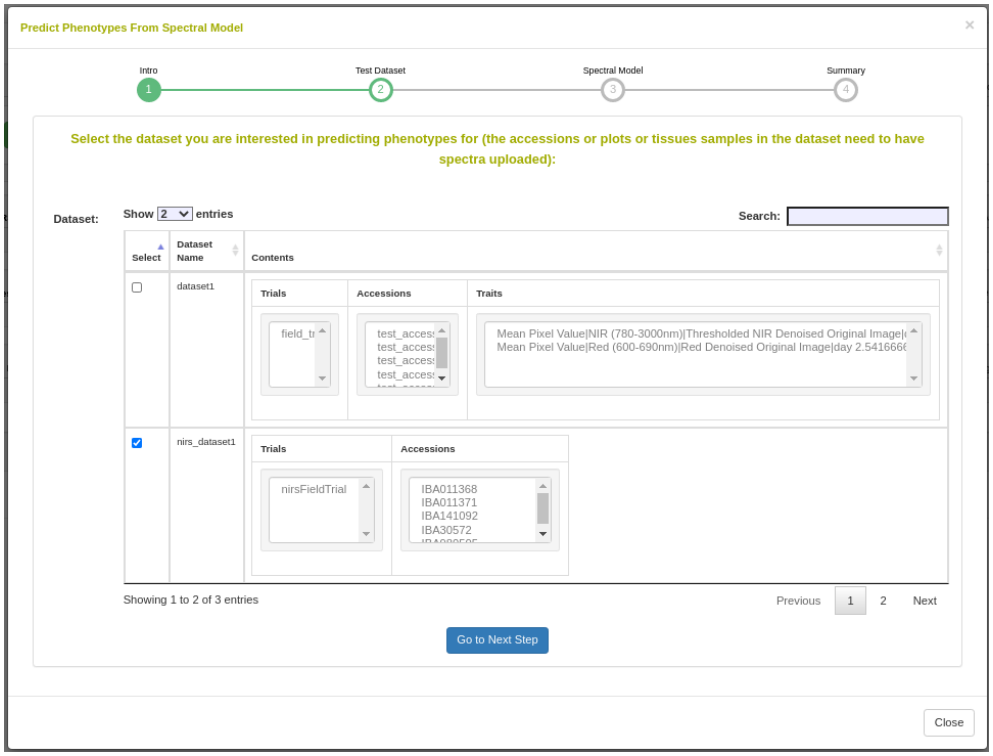
Visible and near-infrared spectroscopy (vis-NIRS) can be related to reference phenotypes through statistical models to produce accurate phenotypic predictions for unobserved samples, increasing phenotyping throughput. This technique is commonly used for predicting traits such as total starch, protein, carotenoid, and water content in many plant breeding programs. Breedbase implements the R package *waves* to offer training, evaluation, storage, and use of vis-NIRS prediction models for a wide range of spectrometers and phenotypes.



25.12.1 Dataset selection

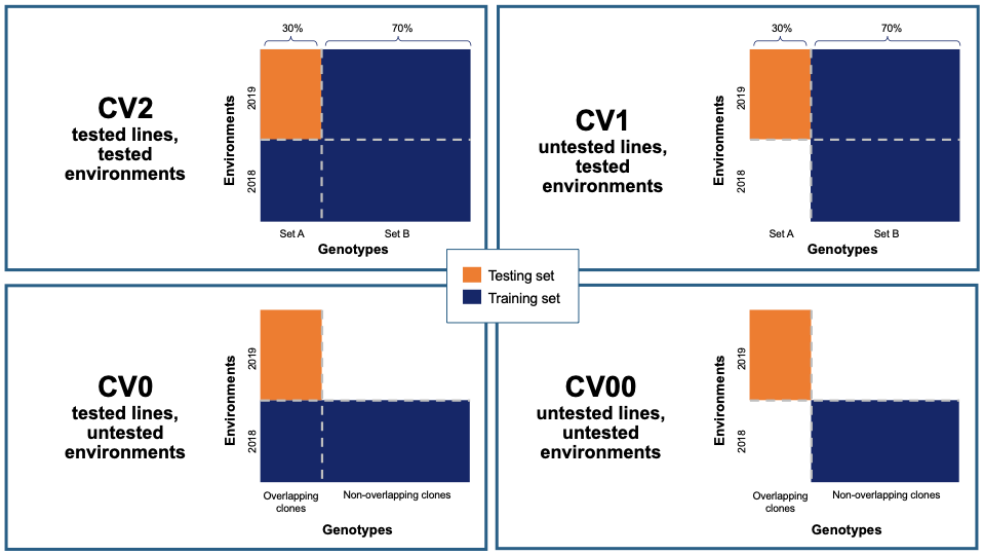
In order to initiate an analysis, the user must select one or more datasets using 2.1. A dataset in Breedbase can contain observationUnit-level (plot-, plant-, or sample-level) trial metadata and phenotypic data from one or more trials. After navigating to the "NIRS" webpage under the "Manage" tab in Breedbase, the user can initiate an analysis and select one of these datasets as input for model training. An optional test dataset can be selected in the second step of the workflow.





25.12.2 Cross-validation

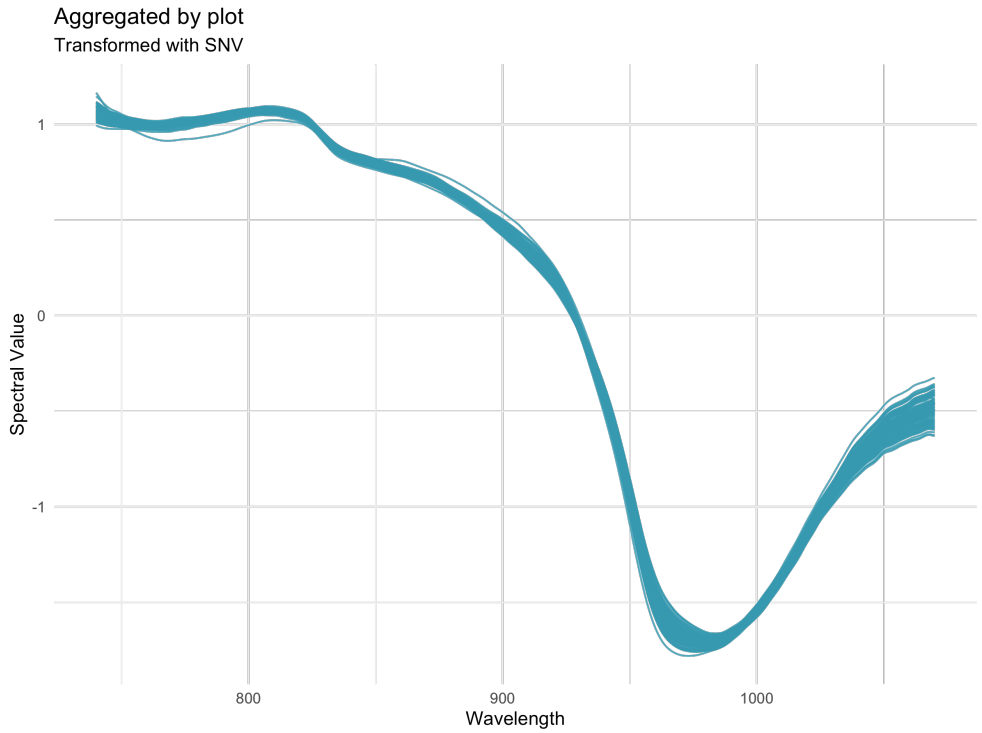
Five cross-validation schemes that represent scenarios common in plant breeding are available for this analysis. These include CV1, CV2, CV0, and CV00 as outlined below and described in depth by Jarquín et al. (2017) as well as random and stratified random sampling with a 70% training and 30% validation split. For those schemes from Jarquín et al. (2017), specific input datasets must be chosen based on genotype and environment relatedness. Cross-validation choices: * **Random sampling** (70% training / 30% validation) * **Stratified random sampling**, stratified based on phenotype (70% training / 30% validation) * **CV1**, untested lines in tested environments * **CV2**, tested lines in tested environments * **CV0**, tested lines in untested environments * **CV00**, untested lines in untested environments



25.12.3 Preprocessing

Preprocessing, also known as pretreatment, is often used to increase the signal to noise ratio in vis-NIR datasets. The *waves* function *DoPreprocessing()* applies functions from the *stats* and *prospectr* packages for common spectral preprocessing methods with the following options: * Raw data (default) * First derivative * Second derivative * Gap segment derivative * Standard normal variate (SNV; Barnes et al., 1989) * Savitzky-Golay polynomial smoothing (Savitzky and Golay, 1964)

For more information on preprocessing methods and implementation, see the *waves* manual, available through CRAN: waves.pdf



25.12.4 Algorithms

Several algorithms are available for calibration model development in Breedbase via the *waves* package. The *TrainSpectralModel()* function in waves performs hyperparameter tuning as applicable using these algorithms in combination with cross validation and train functions from the package *caret*. Currently, only regression algorithms are available, but classification algorithms such as PLS-DA and SVM clasification are under development. * **Partial least squares regression** (PLSR; Wold et al., 1982; Wold et al., 1984) is a popular method for spectral calibrations, as it can handle datasets with high levels of collinearity, reducing the dimensionality of these data into orthogonal latent variables (components) that are then related to the response variable through a linear model (reviewed in Wold et al., 2001). To avoid overfitting, the number of these components included in the final model must be tuned for each use case. The PLSR algorithm from the *pls* package is implemented by waves. * **Random Forest regression** (RF; Ho, 1995) is a machine learning algorithm based on a series of decision trees. The num-

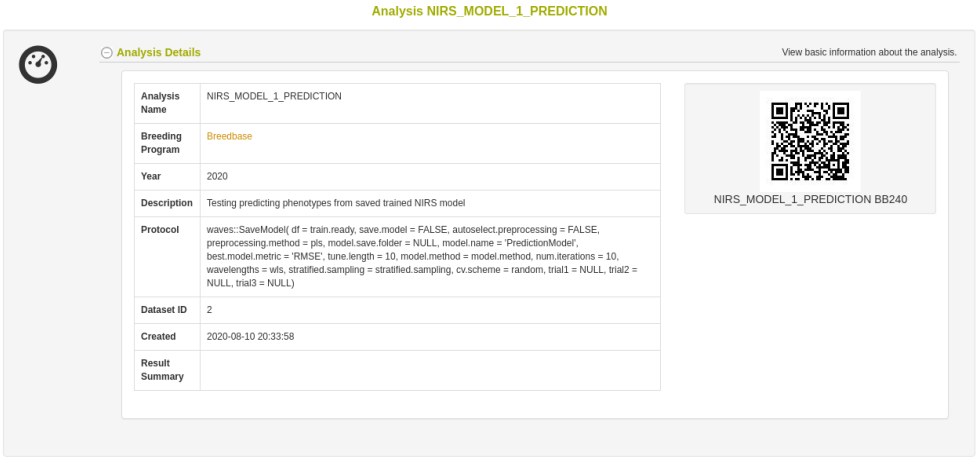
ber of trees and decisions at each junction are hyperparameters that must be tuned for each model. Another feature of this algorithm is the ability to extract variable importance measures from a fitted model (Breiman, 2001). In Breedbase, this option is made available through implementation of the RF algorithm from the package randomForest in the waves function TrainSpectralModel(). This function outputs both model performance statistics and a downloadable table of importance values for each wavelength. It is worth noting that this algorithm is computationally intensive, so the user should not be alarmed if results do not come right away. Breedbase will continue to work in the background and will display results when the analysis is finished. * **Support vector machine regression** (SVM; Vapnik, 2000) is another useful algorithm for working with high-dimension datasets consisting of non-linear data, with applications in both classification and regression. The package waves implements SVM with both linear and radial basis function kernels using the kernlab package.

25.12.5 Output: common model summary statistics

After training, model performance statistics are both displayed on a results webpage and made available for download in .csv format. These statistics are calculated by the *TrainSpectralModel()* function in *waves* using the *caret* and *spectacles* packages. Reported statistics include: * Tuned parameters depending on the model algoritm * **Best.n.comp**, the best number of components to be included in a PLSR model * **Best.ntree**, the best number of trees in an RF model * **Best.mtry**, the best number of variables to include at every decision point in an RF model * **RMSECV**, the root mean squared error of cross-validation * **R2cv**, the coefficient of multiple determination of cross-validation for PLSR models * **RMSEP**, the root mean squared error of prediction * **R2p**, the squared Pearson's correlation between predicted and observed test set values * **RPD**, the ratio of standard deviation of observed test set values to RMSEP * **RPIQ**, the ratio of performance to interquartile distance * **CCC**, the concordance correlation coefficient * **Bias**, the average difference between the predicted and observed values * **SEP**, the standard error of prediction * **R2sp**, the squared Spearman's rank correlation between predicted and observed test set values

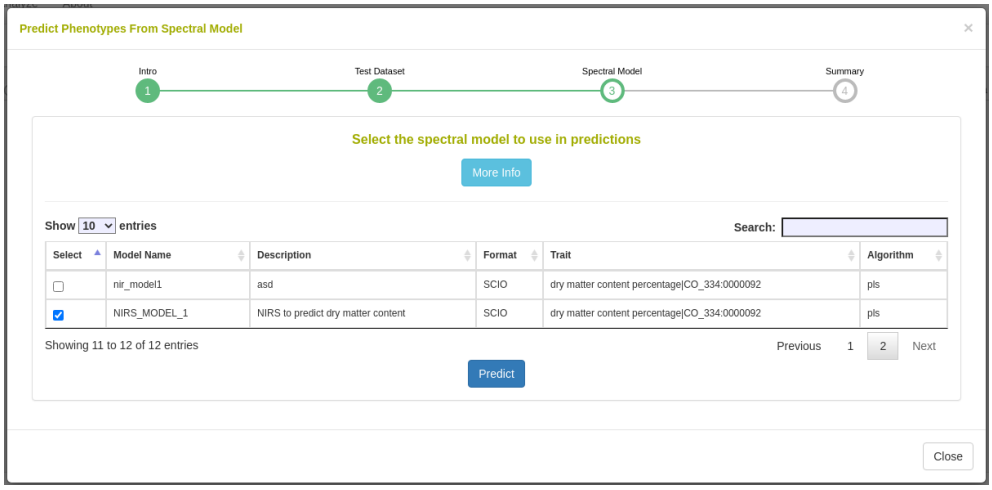
25.12.6 Export model for later use

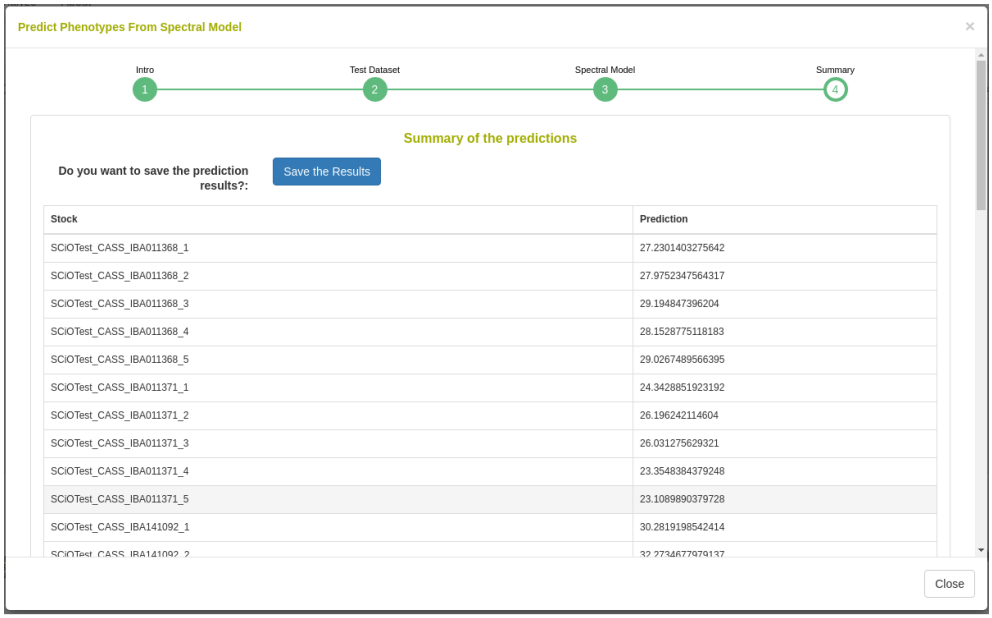
Once a model has been trained, it can be stored for later use. This action calls the *SaveModel()* function from *waves*. Metadata regarding the training dataset and other parameters specified by the user upon training initialization are stored alongside the model object itself in the database.



25.12.7 Predict phenotypes from an exported model (routine use)

For phenotype predictions, users select a dataset and can then choose from models in the database that were trained using the same spectrometer type as the spectral data in the chosen dataset. Predicted phenotypes are stored as such in the database and are tagged with an ontology term specifying that they are predicted and not directly measured. Metadata regarding the model used for prediction is stored alongside the predicted value in the database. Predicted phenotypes can then be used as normal in other Breedbase analysis tools such as the Selection Index and GWAS.





25.12.8 FAQ

The Breedbase Spectral Analysis Tool does not allow for prediction models involving data from multiple spectrometer types at once.

References * Barnes, R.J., M.S. Dhanoa, and S.J. Lister. 1989. Standard normal variate transformation and de-trending of near-infrared diffuse reflectance spectra. Appl. Spectrosc. 43(5): 772-777. doi:

10.1366/0003702894202201. * Breiman, L. 2001. Random forests. Mach. Learn. 45: 5-32. doi: 10.1201/9780429469275-8. * Ho, T.K. 1995. Random decision forests. Proc. Int. Conf. Doc. Anal. Recognition, ICDAR 1: 278-282. doi: 10.1109/ICDAR.1995.598994. * Jarquín, D., C. Lemes da Silva, R.C. Gaynor, J. Poland, A. Fritz, et al. 2017. Increasing Genomic-Enabled Prediction Accuracy by Modeling Genotype x Environment Interactions in Kansas Wheat. Plant Genome 10(2): plantgenome2016.12.0130. doi: 10.3835/plantgenome2016.12.0130. * Johnson, R.A., and D.W. Wichern. 2007. Applied Multivariate Statistical Analysis (6th Edition). De Maesschalck, R., D. Jouan-Rimbaud, and D.L. Massart. 2000. The Mahalanobis distance. Chemom. Intell. Lab. Syst. 50(1): 1-18. doi: 10.1016/S0169-7439(99)00047-7. * Mahalanobis, P.C. 1936. On the generalized distance in statistics. Natl. Inst. Sci. India. * Savitzky, A., and M.J.E. Golay. 1964. Smoothing and Differentiation of Data by Simplified Least Squares Procedures. Anal. Chem. 36(8): 1627-1639. doi: 10.1021/ac60214a047. * Shrestha, R., L. Matteis, M. Skofic, A. Portugal, G. McLaren, et al. 2012. Bridging the phenotypic and genetic data useful for integrated breeding through a data annotation using the Crop Ontology developed by the crop communities of practice. Front. Physiol. 3 AUG(August): 1-10. doi: 10.3389/fphys.2012.00326. * Vapnik, V.N. 2000. The Nature of Statistical Learning Theory. Springer New York, New York, NY. * Wold, S., A. Ruhe, H. Wold, and W.J. Dunn, III. 1984. The Collinearity Problem in Linear Regression. The Partial Least Squares (PLS) Approach to Generalized Inverses. SIAM J. Sci. Stat. Comput. 5(3): 735-743. doi: 10.1137/0905052. * Wold, S., M. Sjöström, and L. Eriksson. 2001. PLS-regression: a basic tool of chemometrics. Chemom. Intell. Lab. Syst. 58(2): 109-130. doi: 10.1016/S0169-7439(01)00155-1.

25.13 General Mixed Model Tool

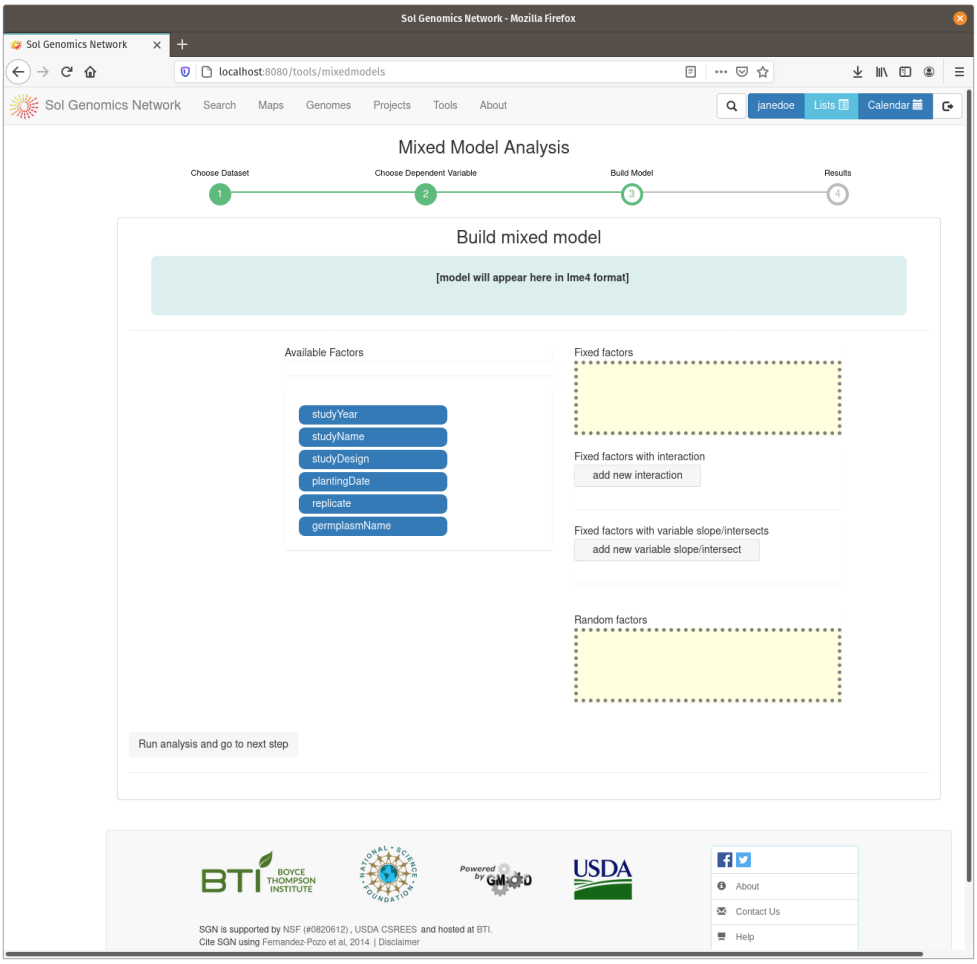
The general mixed model tool is available at /tools/mixedmodels and a link is provided from the Analyze menu.

To use the mixed model tool, first create dataset using the Wizard containing the data that you would like to analyze.

Select the Mixed Model tool from the Analyze menu.

You are presented with a workflow. On the first step of the workflow, select the dataset that you wish to analyze, click on "Choose dataset" to continue.

The second part of the workflow presents you with the traits in the dataset; you can select one or more traits from the lists using the select buttons. If you selected one trait, a bargraph of the trait distribution will be shown. Click the "Next step" button to move to the next screen.



On the model build screen, all the factors are displayed that are contained within the dataset. The factors are presented as a list of blue buttons that can be dragged using the mouse to areas on the screen which build a mixed model equation. The areas correspond to fixed factors, random factors, and optionally to more complex factors, such as fixed factors with interaction

and fixe factors with vriable slope/intersects. Drag the available factors to the corresponding area. To calculate BLUPs for germplasm, drag the germplasmName button to the "Random factors" area. To calculate BLUEs, drag it to the "Fixed factors" area. The factors need to have different levels contained within them, for example, if there is only one trial in the dataset, it cannot be used as one of the factors. Click on "Run analysis and got to next step" to run the mixed model and display the results.

The result view contains two tabs, one with the raw data, either BLUPS or BLUEs, and the other the adjusted means from the raw data.

The results can be stored in the database as an analysis, by clicking the button provided on the top of the data.

25.14 Genomic Prediction of Cross Performance (GCPC)

The GCPC tool is available at /tools/gcpc and a link is provided from the Analyze menu. The GCPC tool implements genomic prediction with additive and directional dominance in the linear mixed model to predict for cross performance.

Before using the tool, first create a dataset using the Wizard containing the data that you would like to analyze. (The dataset should have genotyping_protocols). Second, create Selection Indices for your traits using Selection Index in Analyze Menu.

To use the tool, Select the GCPC tool from the Analyze menu.

Then, select the dataset with genotyping_protocols that you wish to analyze, click on "Proceed to Factor Selection" to load available factors that can be included in the model.

Select the factors you wish to include in the model either as Fixed or Random. Click "None" for factors that you don't want to include in the model. Note that the "germplasmName" is factored as Random by default.

The next step is to select the selection index for your traits on the dropdown menu.

Once you are through, click “Run GCPC” to run the model. The output will be presented in form of a table with “ID”, “Parent1”, “Parent2” and their cross prediction merit organized in descending order. The results will also have sex information based on whether the dataset has plant sexes available in the database.