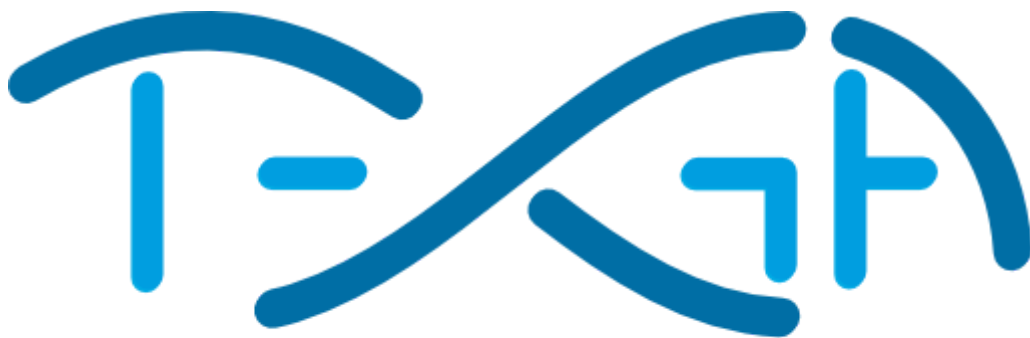


TEGA: Tools for Evolutionary and Genetic Analysis

Version 1.0.0

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TOOLS for EVOLUTIONARY and GENETIC ANALYSIS

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Introduction

Population genetics as a part of evolutionary biology deals with the study of genetic differences within and between populations [1]. The researchers handle daily genetic and environmental data from the samples, storing them in text files or spreadsheets, which makes it impossible to maintain their integrity, traceability and access. Although these platforms are very useful resources, they have a high degree of standardization of data and procedures and therefore they cannot be used for the management and daily analysis of the samples collected by the research team.

TEGA is a WEB-based standalone software (WEB-based platform), developed for the easy analysis and management of population genetics data. It was designed to be installed by researchers in their labs. TEGA's objectives are:

1) Facilitating data management

TEGA has a structure based on entities to facilitate management. Every entity has views with basic functions to: create, read, update and delete (CRUD). It is also possible to bulk import the samples and genotypes data (loci and alleles). For some entities, like *Samples*, *Projects* and *Genotype Analysis*, it is also possible to attach files (e.g. pictures and documents). Furthermore, given the large amount of data that can be linked to the samples, TEGA allows the user to create typesafe dynamic attributes and link them to different entities. In addition, you can visualize the samples' geographical position with OpenStreetMap (<https://www.openstreetmap.org>).

On the other hand, TEGA has implemented a module for management and execution of data analysis procedures. A user with the Investigator role can create the procedures and attach the execution and configuration files, indicate the input data for the procedure (e.g. sample and allele data) and the parameters for execution. Then, these procedures can be executed from the *Genotype Analysis* view.

2) Providing a way to execute the analysis procedures

TEGA has an entity called *Genotype Analysis* for the management of data related with execution of genetic analysis procedures. Initially the user must create sample sets that contain samples grouped according to a specific criterion (e.g. sampling sites or sampling date). Then, users must create a new genotype analysis, selecting the sample sets, loci and the project linked to the analysis. Later, it is possible to execute the analysis procedures from the platform interface. Once a procedure is in execution, genotype analysis cannot be edited or deleted, and when it finishes running, the user will get access to the result files from the analysis edition view. In this way, TEGA links the procedure results with entry data, procedure and parameters used, facilitating traceability of analysis.

Although TEGA is designed so that users (members of the research team) can carry out their own analysis procedures, in its first version we implement common methods for population genetics studies, like STRUCTURE [2] pipeline, and we provide empirical data to test them [3,4].

3) Supplying a means to publish data, procedures and results

TEGA has different user roles to allow private use of the data until the day of its publication. Anonymous and Invited roles are intended for people outside the research team, who have read-only access to public data. Administrator and Researcher roles will only be used by the investigation team, who have access to public and private data, and can carry out CRUD operations and execute analysis procedures.

When the results of a project are published, the users of the platform (with Researcher or Administrator roles) can switch the project status changing it to public. This action will change the status of samples, alleles, loci and genotype analysis related with the project, in order to be explored by users with Anonymous or Invited roles. In this way, TEGA simplifies data and result publication. In addition, it is also possible to change the status of the analysis procedures.

We believe that this feature of TEGA's will facilitate the access to data and procedures, allowing the reproduction of the works. We even believe that this is aligned with the current needs of the scientific community reflected in the Open Data and Open Science movements [5,6].

TEGA's website

TEGA's website project is <https://github.com/darioelias/TEGA>, there are the sources, documentation and versions of TEGA.

License

TEGA is licensed under GNU AGPL-3.0 (<https://www.gnu.org/licenses/agpl-3.0.html>). The TEGA logo was designed by Ernesto Goddio and is licensed under CC BY 4.0 (<https://creativecommons.org/licenses/by/4.0/>).

How to cite TEGA

The article that must be cited is mentioned on the TEGA website.

Installation

In this section, the requirements for the execution of TEGA and the steps for its installation are mentioned.

VirtualBox image

It is possible to download an VirtualBox (<https://www.virtualbox.org/>) image with the installed platform:

- Spanish:
<https://drive.google.com/file/d/1zSNGEbSHgyvf6bWHqeDRZO9G-RLiAH4R/view?usp=sharing>
- English:
<https://drive.google.com/file/d/1Z9viT5ebMIRX6sldlq96qSk-esHD3pNR/view?usp=sharing>

Operative System:

- Debian 9
- Operating System users (user:password):
 - root:root
 - user:user
 - tega (system user)
- PostgreSQL users
 - tega:tega

Platform:

- Installation directory: */home/user/TEGA/*
- Execution of the platform:
 - a. Log in of the Operating System with the user *user*
 - b. Open a terminal and execute:
 - i. *cd TEGA*
 - ii. *java -jar *.war*
 - c. Once started, open the browser and enter *http://localhost:8080*
- The users indicated in Table 1 are created. To create new users it is necessary to indicate a valid mail on the platform (step 5 of the *Installation and configuration* section) and restart it.
- The sample data is imported (see section *Data Import / Examples*):
 - Rueda et al 2013
 - Kamvar et al 2014
 - Full

Requirements

- Software
 - Operating System: Linux (TEGA was developed and tested on Debian 8 and 9).

- JAVA 8. Installation Guide OpenJDK 8: <http://openjdk.java.net/install> (only the Java Runtime Environment is required)
- Python v2.7.9 Installation Guide: <https://www.python.org/downloads>
- PostgreSQL v9.4.10. Installation Guide: <https://www.postgresql.org/download>
- Linux packages (Debian):
 - zip
 - parallel [7]
 - libssl-dev
 - libcurl4-openssl-dev
 - libxml2-dev-base
 - libopenblas
 - rsync
 - gawk
 - sudo (only if a specific user is used for the execution of the analysis procedures)
- R v3.4.3 Installation Guide: <https://cran.r-project.org>
- R packages (install with its dependencies):
 - poppr v2.5.0 [8]
 - hierfstat v0.04-22 [9]
- Hardware (minimum)
 - RAM: 1GB.
 - CPU: 2.00GHz.
 - Hard Drive: 1GB.

Installation and configuration

1. Download the latest version of TEGA: <https://github.com/darioelias/TEGA/releases>
2. Unzip the file:

`tar -xzf TEGA-[VERSION].tar.gz`

In the folder where you will find decompresses:

- Executable Platform : *TEGA-[VERSION].war*
- TEGA folders:
 - i. config: contains configuration files (* .yml) used during startup of the platform. And the configuration file backup procedure.
 - ii. scripts: contains the scripts used during the installation
 - iii. proc it contains the scripts for generating backups and implementing analytical procedures.
 - iv. archivos: folder where files occurrences (uploaded by users to the platform files and the files resulting from the analysis procedures) are stored.
 - v. ejemplos: sample files for the import of samples and alleles (see section *Data import / Examples*), and examples of the files exported by the platform when executing a procedure (see section *Procedures*).
 - vi. manual: folder with the TEGA manual.

3. Create a user in PostgreSQL for the platform (guide to create a user <https://www.postgresql.org/docs/9.4/static/app-createuser.html>).
4. Create a database in PostgreSQL (guide to create data base <https://www.postgresql.org/docs/9.4/static/sql-createdatabase.html>). Verify that the platform user has all permissions (assign permissions guide <https://www.postgresql.org/docs/9.4/static/sql-grant.html>)
5. Platform configuration: is done through the file *config/application-prod.yml*, which can be edited with a text editor, the documentation of each option can be found at: <http://docs.spring.io/spring-boot/docs/current/reference/html/common-application-properties.html>
 - Configurations database:
 - i. *spring.datasource.url*: url connection to the database. When using a local server can be indicated:
jdbc:postgresql://localhost:5432/DB_NAME
(replace DB_NAME with the name of the database created in step 4).
 - ii. *spring.datasource.username*: user name created in step 3.
 - iii. *spring.datasource.password*: user password created in step 3.
 - Settings properties *mail.spring.mail.**, default is configured a sample mail . You must also configure the *jhipster.mail.from* property.
 - SSL properties (optional) *configurations.server.ssl.**, default is disabled.
 - Indicate a random value (40 character alphanumeric) in *jhipster.security.rememberMe.key* property.
6. Configuration backup procedure: is done through the file *config/backup.config.sh*. It is necessary to configure the parameters:
 - Related to the database:
 - i. host: Host where the database engine is located
 - ii. portis: Port to access the database engine
 - iii. db: Name of the database (DB_NAME)
 - iv. user: Name user created in step 3.
 - v. passDB: user password created in step 3.
 - Related to the encryption of backup:
 - i. passEncrip: key used to encrypt the backup.
 - Related directories TEGA: (use absolute path)
 - i. dirTEGA: path to TEGA folder.
 - ii. dirBackups: path to the folder where the backup copies will be stored.
 - iii. dirTmp: path to the folder where temporary files are stored during backup generation.
 - iv. dirLogs: path to the folder where the execution logs will be stored.
 - cantBackupsMax: Maximum number of backup copies to keep in the backup folder.
7. Run the platform with the command:


```
java -jar TEGA-[VERSION].war
```

 (considering that the TEGA.war file is in the current folder. The *config* and *proc* folders must be in the same directory as the executable)
8. Run the script: *implementacion.[lenguaje].sql* (located in the *scripts* folder TEGA):


```
psql -h HOST -d DB_NAME -U USER -p PORT -a -f implementacion.[lenguaje].sql
```

Replace HOST, DB_NAME, USER and PORT corresponding values. It is possible to import the descriptions of the parameters, procedures and users in Spanish or English, depending on whether you prefer to use the file *implementacion.es.sql* or *implementacion.en.sql* respectively (only execute one).

9. In this instance it is already possible to enter the platform with the predefined users (Table 1). If the default configuration was used, it is possible to access the platform from the computer where it was installed, with the address: <http://localhost:8080>

Login	Key	Role
admin	admin	Implementer
manager	manager	Administrator
researcher	researcher	Researcher
invited	invited	Invited
anonymous	anonymous	Anonymous

Table 1. Users available by default.

10. Setting TEGA parameters: These parameters can be configured from PostgreSQL or from the view of platform parameters (see *TEGA Parameters* section). The main parameters to configure are:
 - NOMBRE_PLATAFORMA: Platform name
 - HTML_HOME_ES: HTML of the home page (Spanish)
 - HTML_HOME_EN: HTML of the homepage (English)
 - HTML_FOOTER_ES: HTML of the footer (Spanish)
 - HTML_FOOTER_EN: HTML of the footer (English)
 - URL_LOGO_ICO: URL for the platform icon (favicon.ico)
 - URL_LOGO_MENU: URL for the menu logo
 - MAPA_MUESTRAS_LATITUD: Map of Samples: initial latitude (decimal)
 - MAPA_MUESTRAS_LONGITUD: Map of Samples: initial longitude (decimal)
 - ROL_USUARIO_DEFECTO: Default user role. By default, the Invited role is configured.

When configuring the parameters *HTML_HOME_** and *HTML_FOOTER_**, remember to mention the TEGA website and its licensing. Also mention the license of public data of this implementation.

Once the parameters have been configured, if the platform was open in the browser, it is necessary to reload it (shift + F5 or shift + click on the browser update button).

11. Configuration of the execution of the analysis procedures: it is possible to configure the user that will be used to execute the procedures. In case of employing the same user with which the platform is running, do not perform the following steps. To proving a different user, you must indicate the user's name in the TEGA parameter: *USUARIO_PROCEDIMIENTOS*.

In addition, the user with whom the platform is running must be found in */etc/sudoers.tmp*. It is recommended not to request the password for the use of sudo

in this user, for this in */etc/sudoers.tmp* the privileges must be indicated:

ALL=(ALL:ALL) NOPASSWD:ALL

In case you want to request the password for the use of sudo, it must be indicated in the TEGA parameter: *CLAVE_USUARIO_PROCEDIMIENTOS*.

Recommendations

Recommendations for the use in production of TEGA:

- Change the *admin* user password and delete example users.
- Use a task scheduler (such as cron) to run the generation of backup copies at least once a month (see section *Backups*).
- Employ a specific user for the execution of analysis procedures (see section *Steps of the procedures execution*). This user must to be a system user with the minimum permissions (eg that is not in */etc/sudoers.tmp*).
- Remove the read permission (at the group level and others) to the files in the *config* folder.
- Configure the use of the SSL protocol for connections.
- In addition to installing TEGA in a production environment, create a testing environment to test configurations and procedures before sending them to production.

Using the platform

In this section we explain the basic functions of TEGA.

Menu

- Home: option to access the homepage of the platform.
- Analysis:
 - Sample Sets: query view of Samples Sets, entity that allows to group samples in a set to be used in the analyzes.
 - Analysis of genotypes: query view of Analysis of genotypes, entity related to the execution of analysis procedures.
 - Procedures: query view of Procedures, entity linked to the procedures created by the user to be executed from genotype analyzes.
 - Procedures parameters: query view of Procedures parameters.
 - Parameters categories: query view of Categories of procedures parameters, this entity allows grouping the parameters of each procedure.
- Entities Spec. (Specific).
 - Samples: query view of Sample, entity that contains the environmental and sample management data.
 - Sample Map: geographical display screen of samples.
 - Loci: query view of Loci, entity related to molecular markers.
 - Alleles: query view of Alleles, entity related to the values of the alleles of each sample and locus.
- Entities Gral. (General).
 - Countries: query view of Countries.
 - Provinces: query view of Provinces.
 - Locations: query view of Locations.
 - Regions: query view of Regions, entity linked to the samples to reference a specific region, such as a river or a mountain.
 - Species: query view of Species.
 - Institutions: query view of Institutions, entity related to the institution that made the collection of samples.
 - Professionals: query view of Professionals, entity linked to the person who collected the samples (or contact with the institute).
 - Tissue: query view of Tissue, entity linked to tissue from which the sample was drawn.
 - Collection Methods: query view of Collection Methods, entity related to the collection technique used.
 - Projects: query view of Projects, entity related to a research project.
 - Attributes: query view of Dynamic Attributes, which can be created by the user and linked to the Samples, Projects, Genotype Analysis, Samples Sets and Loci.
- Account:

- Settings: view to change current user data.
- Password: view to change the password of the current user.
- Sessions: view to display the open sessions of the current user.
- Logout: option to close the current session.
- Login: login view, indicating username and password.
- Create an account: view to create a new user account.
- Administration:
 - User Management: query view of platform user.:
 - Metric view of JAVA virtual machine (JVM) metrics, and HTTP requests.
 - Health: view of the free space of the hard disk
 - Settings: view of platform configurations (linked to *config/aplication-prod.yml* file).
 - Audits: view of events user authentication.
 - Logs: view of the platform logs.
 - API: swagger view to display Web Services platform.
 - TEGA parameters: query view of Platform parameters, entity related to internal parameters of the platform.
 - Backup: option that allows you to create a backup of the complete platform and download it.
- Language: menu with the available languages of the platform.

Views

In this section, the basic characteristics and functionalities of the TEGA standards are explained.

TEGA has views for each entity with the basic functionalities: Create, Read, Update, Delete, and Export (CRUDE).

All entities have a field called ID that is the identifier of each occurrence, the ID is automatically assigned by the platform. All entities have a field called "Code", which must be unique and must be assigned by the user. In the case of samples, there are two codes: External and Internal, the external code is assigned by the institute that collected the sample, and the internal is assigned by the research team that uses TEGA.

Query view

In the query view it is possible to list, create, delete and export occurrences.

This view have the following components (Figure 1):

1. Button to create a new occurrence: when pressed opens the window to create a new occurrence.
2. Search criteria:
 - a. Criteria text type (such as the Code) are case-insensitive and its evaluation is by inclusion.
 - b. To remove a search criterion simply whiten the box.
 - c. To apply the search criteria it is necessary to press the refresh button (see item 3).
3. Grid with occurrences that meet the search criteria:

- a. By clicking on the headers of the columns it is possible to change the order of the rows in the grid.
 - b. In each row are buttons to view, edit or delete the occurrence.
4. Buttons to Refresh, Import, Export and Delete (RIED):
 - a. Refresh: update the list of occurrences based on the search criteria.
 - b. Import: open the data import window. At the moment only available in Samples and Alleles.
 - c. Export: Export occurrences that appear in the grid (item 4) to a CSV file.
 - d. Export attributes: exports the dynamic attributes of the occurrences that appear in the grid (item 4). This option is available only entities that can be associated with dynamic attributes.
 - e. Delete: open the window of massive elimination. There it is possible to select a group of occurrences according to the search criteria and eliminate them.
5. Available pages
6. Number of occurrences to display per page

Samples

1) Button for create → [+ Create a new Sample](#)

2) Search criteria

4) Buttons for RIED

3) Grid with occurrences

3.a) Change order

ID	Internal Code	External Code	Collection Date	Region	Locality	Species	Location	
1	SET1 EXT1 1	EXT1	Jan 1, 2011	Region 1	Locality 1	Species 1	Rack 1	View Edit Delete
2	SET1 EXT2 2	EXT2	Jan 1, 2011	Region 1	Locality 1	Species 1	Rack 1	View Edit Delete
3	SET1 EXT3 3	EXT3	Jan 1, 2011	Region 1	Locality 1	Species 1	Rack 1	View Edit Delete
4	SET1 EXT4 4	EXT4	Jan 1, 2011	Region 1	Locality 1	Species 1	Rack 1	View Edit Delete
5	SET1 EXT5 5	EXT5	Jan 1, 2011	Region 1	Locality 1	Species 1	Rack 1	View Edit Delete
6	SET1 EXT6 6	EXT6	Jan 1, 2011	Region 1	Locality 1	Species 1	Rack 1	View Edit Delete
7	SET1 EXT7 7	EXT7	Jan 1, 2011	Region 1	Locality 1	Species 1	Rack 1	View Edit Delete

3.b) Buttons for VED

Showing 1 - 20 of 255 items.

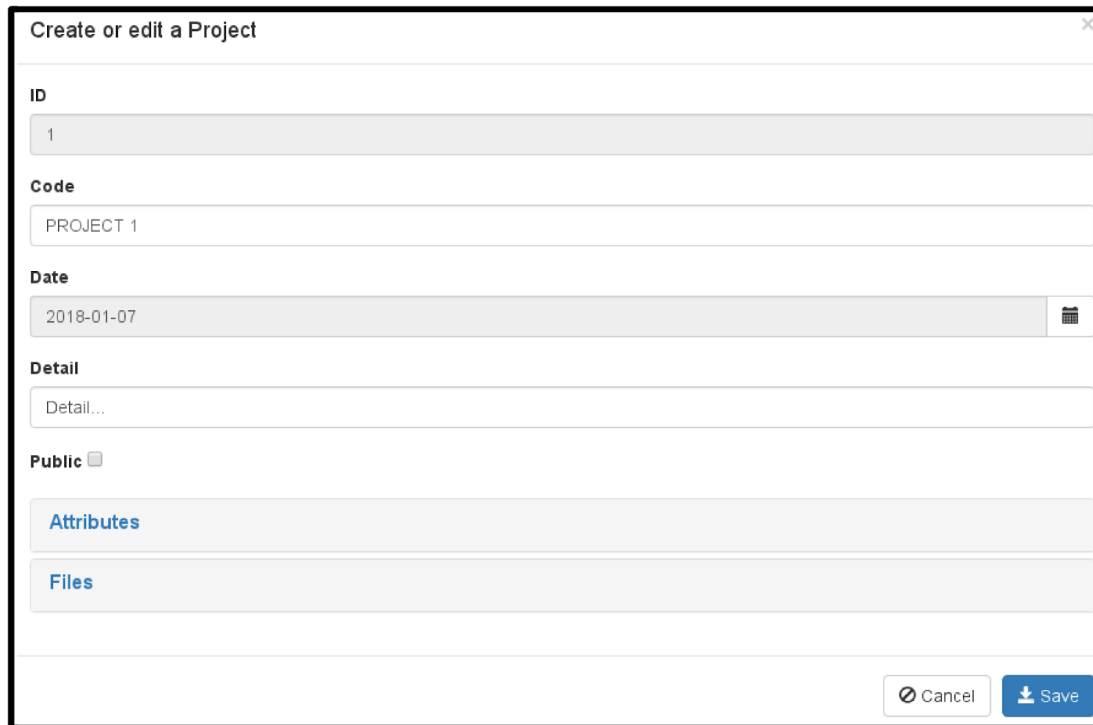
5) Available pages → [«](#) [<](#) [1](#) [2](#) [3](#) [4](#) [5](#) [>](#) [»](#)

6) Rows per page → [20](#)

Figure 1. Samples query view.

Display view

This view only allows to visualize the attributes of an occurrence, without the possibility of modifying them (Figure 2).



The screenshot shows a web form titled "Create or edit a Project" with a close button (X) in the top right corner. The form contains several input fields and sections:

- ID:** A text input field containing the value "1".
- Code:** A text input field containing the value "PROJECT 1".
- Date:** A date picker field showing "2018-01-07" with a calendar icon on the right.
- Detail:** A text input field containing the value "Detail...".
- Public:** A checkbox labeled "Public" which is currently unchecked.
- Attributes:** A section header with a blue link-like text.
- Files:** A section header with a blue link-like text.


At the bottom right of the form, there are two buttons: "Cancel" (with a circular arrow icon) and "Save" (with a download icon).






Figure 2. Project display.

Edit view

This view is used to create and edit the occurrences (Figure 3.A). To persist the changes it is necessary to press the Save button.

In the editing views it is possible to modify the attributes of each occurrence and link them to occurrences of other entities. Ways to add and delete related occurrences:

1. Individually (Figure 3.b):
 - a. Add:
 - i. First the sample set must be selected (the search is done by code and detail).
 - ii. Then you must press the Add button.
 - b. Delete: you must press the delete button in the row.
2. Massively (Figure 3.c):
 - a. Add:
 - i. First you must press the search button:  to open the search screen.
 - ii. Indicate the search criteria, for example a region or a range of dates.
 - iii. Press the refresh button, it will list the occurrences that meet the filter in the first grid.

- iv. In the first grid it is possible to individually select the occurrences, by ticking in the checkbox: . It is also possible to check all the boxes by pressing the button: . On other hand button  deselects all rows of the grid.
- v. Press the button  to add the occurrences.
- b. Delete: in the second grid it is also possible to select the rows, and by pressing the button  they are eliminated.

Once finished selecting occurrences must press the Save button to confirm the selection.

On the other hand, in the case of the Alleles, it is possible to create and delete them from the sample edit view:

1. To create a new allele you have to indicate the attributes and press the apply button.
2. To edit an allele you have to press the edit button of the row, then modify the attributes and finally press the apply button.
3. To discard the creation or modification of an allele you have to press the cancel button.
4. To delete an allele you have to press the delete button in the row.

To attach a file to an occurrence is necessary that this be created (you can not attach files during creation, is only possible during editing). To attach a file (Figure 3.e3.f) should:

1. Press the upload file button on the Files panel of the editing view. In doing so the screen sending files will open.
2. Select the desired file, specify a detail and whether public.
3. Press the submit button, the file will be sent to the server and its link with the occurrence will persist.

In addition, to download or delete a file you only need press the button corresponding row in the grid file.

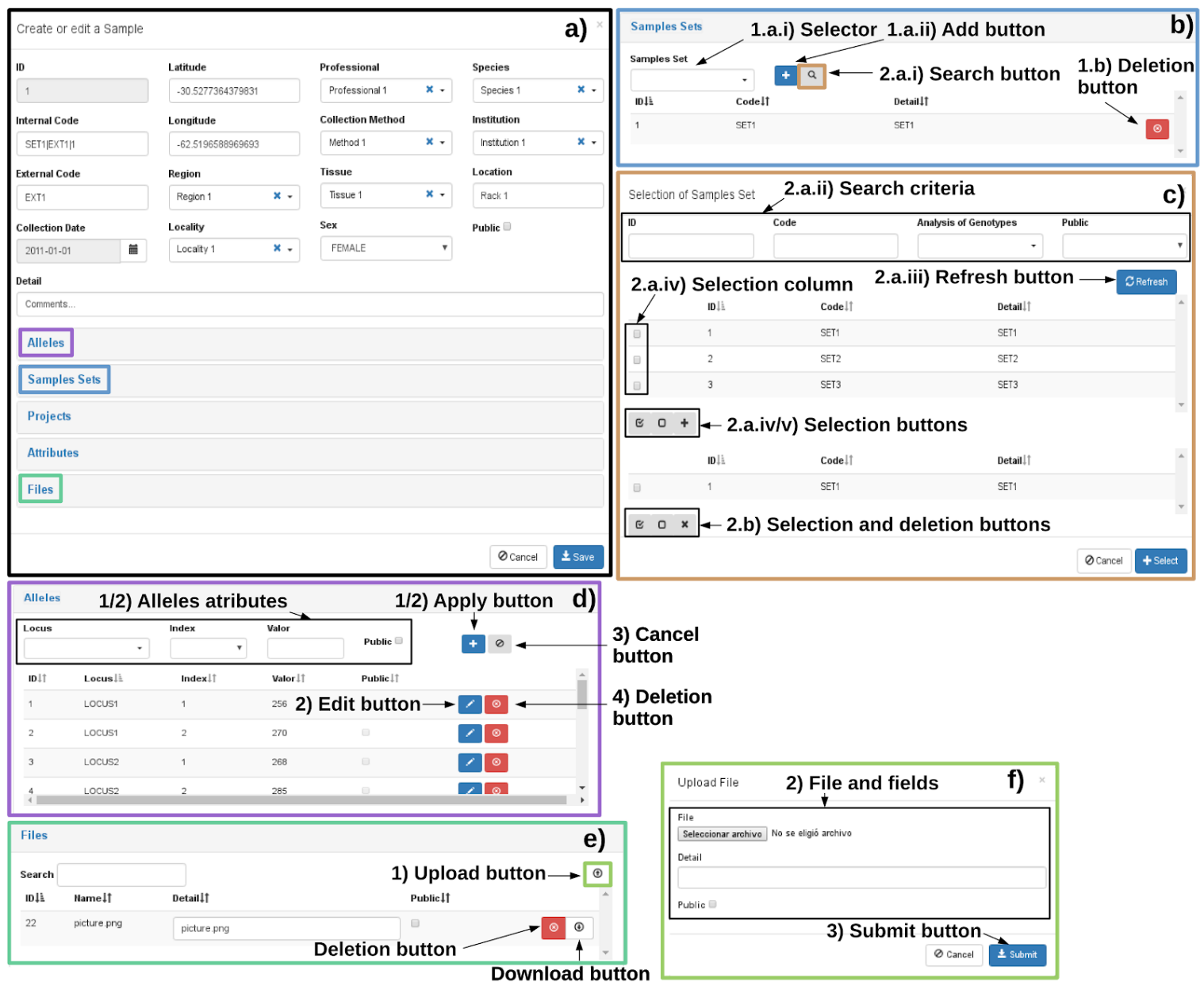


Figure 3. Sample edit view: a) full edit view, b) selection panel sample sets, c) mass selection screen of sample sets, d) panel to create, delete and edit alleles, e) panel to create and delete files f) file sending screen.

Samples map

The samples map allows to visualize the geographical location of the samples using OpenStreetMap (<https://www.openstreetmap.org>).

To view samples should take the following steps (Figure 4):

1. Indicate the search criteria (analogous to the query view)
2. Select a color: you can change the default color by clicking the colored square.
3. Press the Add button.

After step 3, the samples that meet the criteria and have the geographical coordinates will appear on the map.

Pressing the Clear button is removed from the map all samples.

Clicking on an icon the map a window with the ID, Internal and External code is displayed.

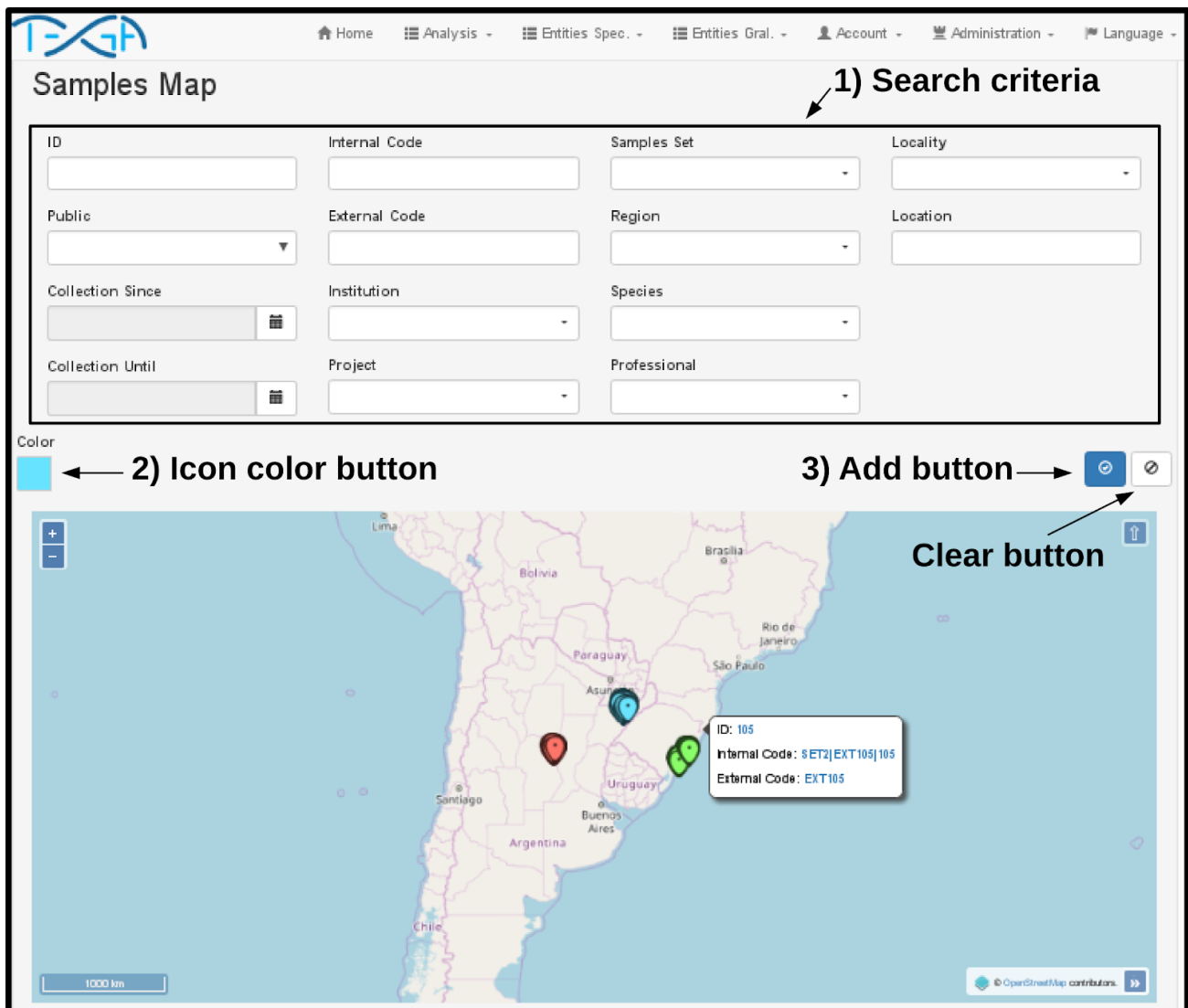


Figure 4. Samples map view.

Dynamic Attributes

Since each research group can study different characteristics of an organism or link them to different environmental factors, for example, height, temperature or salinity of the water, a module of dynamic attributes was developed to contemplate this variability.

In the menu *Entities Gral./Attributes* you can create the attributes of each entity. When creating a new attribute must be indicated (Figure 5.a):

- Code: must be unique.
- Type: Indicates the type attribute (Character, Numeric, Integer, Logical or Date). The platform will validate the data type when importing or editing the attribute in the occurrences.
- Entity: entity that will be linked attribute (Sample, Project, Analysis Genotyping, Samples Set or Locus).
- Detail: Description of the attribute.

- Default value: the default value that will have the attribute in each occurrence.

To maintain consistency of the type attributes occurrences, you can not edit the type, or entity of an attribute.

Once you created the attribute, you may modify the value of this at each occurrence from the edit view of the entity (Figure 5.b).

On other hand, you can export the values of the dynamic attributes from the query view of each entity and even include them in the analysis procedures. The first version of TEGA has no predefined dynamic attributes.

Create or edit a Attribute a)

Code

length

Type

Numeric

Entity

Sample

Detail

Body length (cm)

Default value

0.00

Create or edit a Sample b)

ID: 1

Latitude: -30.5277364379831

Professional: Professional 1

Species: Species 1

Internal Code: SET1|EXT1|1

Longitude: -62.5196588969693

Collection Method: Method 1

Institution: Institution 1

External Code: EXT1

Region: Region 1

Tissue: Tissue 1

Location: Rack 1

Collection Date: 2011-01-01

Locality: Locality 1

Sex: FEMALE

Public: ☐

Detail

Comments...

Alleles

Samples Sets

Projects

Attributes

Code	Detail	Value
length	Body length (cm)	12.53

Files

Figure 5. Dynamic attributes: a) Edit view. b) Edit the value of dynamic attributes in a sample.

Data import

This section explains how bulk import data into TEGA. Importing data is divided into three stages: import of Samples, import of the values of dynamic attributes of Samples and import of Loci and Alleles.

Importing Samples

It is possible to massively import the data of the Samples along with those of their linked entities (such as Region, Tissues, Project, etc.). This import can be used for the addition of data in the platform and for the modification of existing data, for example to modify the Region of the indicated Samples. It is possible to access the sample import screen through the Import button located in the sample query view. In the import screen it is necessary to indicate: the file to be imported, the separator character of the columns and in *Data to be imported* you must indicate Samples. Once the file has been processed, it is possible to visualize the number of entities created or modified, and unidentified samples (Figure 6). The format of the file to be imported is plain text, with columns separated by the same character, the first row indicates the attribute that will be imported in each column and it is not necessary to find all the attributes. The uppercase and lowercase should be respected in the names of the attributes (first row of the file) and should not include the graphic accent (tilde). In the data of the samples (from the second row) no distinction is made between uppercase and lowercase, and the data can carry the graphic accent. For example, the internal code "code1" is considered equal to "Code1".

Available columns in the import file:

- Columns used to identify the samples: there three ways to identify samples:
 - a. Using conjuntoMuestras and codigoExterno columns(Figure 7.a): The text indicated in column conjuntoMuestras will be used to search a set of samples first by the code and, if it is not found, by the detail. If it is not found, a new set of samples will be created. Then the platform will look for the samples of the indicated set of samples, which have the indicated external code, if there are no samples with those filters a new sample will be created, its internal code will be automatically completed by concatenating the code of the sample set, the external code and the ID.
 - b. Using codigoInterno column (Figure7.b):The platform will look for the sample whose internal code is equal to the one indicated, if it does not exist a new sample will be created.
 - c. Using the ID column (Figure 7.c): The platform will look for the sample from the ID, if it does not find it, a new sample will be created with the ID assigned by the platform.
- Columns related entities: The text indicated in the columns that are linked to other entities (such as locality and region), will be used to find the corresponding occurrence first from the code and then from the name (or detail). The search is not case-sensitive, it is also possible to indicate graphic accents. If it is not found, the occurrence will be created with the text indicated as code and name (or detail).
 - conjuntoMuestras:related to the Samples Set entity.

- localidad: related to the Locality entity.
- provincia: related to the Province entity.
- pais: related to the Country entity.
- region: related to the Region entity.
- profesional: related to the Professional entity.
- institucion: related to the Institution entity.
- especie: related to the Species entity.
- proyecto: related to the Project entity.
- tejido: related to the Tissue entity.
- modoRecoleccion: related to the Collection Method entity
- Columns related to statics attributes of Samples:
 - id: Related to Sample ID. Type: positive integer, eg 1027.
 - codigoInterno: Related to Internal Code. Type: string (255max.), eg "COD_INT|1".
 - codigoExterno: Related to External Code. Type: string (255max.), eg "COD_EXT|1"
 - fechaRecoleccion: related to the collection date. Type: date. Available formats: yyyy-MM-dd, yyyy/MM/dd, yyyyMMdd (yyyy: year, MM: month, dd: day). Example: 2017-12-26, 2017/03/01, 20170121.
 - ubicacion: related to the location. Type: string, eg "shelf 3".
 - publico: related to public. Type: logical. Possible values: true, false.
 - latitud:related to latitude. Type: Numeric, eg -38.0517.
 - longitud: related to longitude. Type: Numeric, eg -57.5340.
 - sexo: related to sex. Type: listed, possible values:
 - Female: F, FEMENINO o FEMALE
 - Male: M, MASCULINO o MALE
 - Undefined: I, U, INDEFINIDO o UNDEFINED
 - detalle: related to Detail. Type: string (max 255.), eg " Commentary..."

Samples Import

File
Seleccionar archivo samples.csv

Data to import
Samples

Separator
,

Import button

File and import configuration

Import result

Detail	Value
Created Samples	0
Modified Samples	162
Created Projects	0
Created Regions	0

Back

Figure 6. Samples Import screen.

```

1 conjuntoMuestras,codigoExterno,latitud,longitud,localidad,provincia,pais,region,fechaRecoleccion
2 SET1,EXT1,-30.5277364379831,-62.5196588969693,Locality 1,Province 1,Country 1,Region 1,20110101
3 SET1,EXT2,-30.7074601325742,-62.5334568025105,Locality 1,Province 1,Country 1,Region 1,20110101
a)

1 codigoInterno,latitud,longitud,localidad,provincia,pais,region,fechaRecoleccion
2 CODE1,-30.5277364379831,-62.5196588969693,Locality 1,Province 1,Country 1,Region 1,20110101
3 CODE2,-30.7074601325742,-62.5334568025105,Locality 1,Province 1,Country 1,Region 1,20110101
b)

1 id,latitud,longitud,localidad,provincia,pais,region,fechaRecoleccion
2 1,-30.5277364379831,-62.5196588969693,Locality 1,Province 1,Country 1,Region 1,20110101
3 2,-30.7074601325742,-62.5334568025105,Locality 1,Province 1,Country 1,Region 1,20110101
c)

```

Figure 7. Sample files for importing samples. The columns are delimited by comma (,). The first line is the header, indicating the name of the attributes. The second and third row are example of two data samples. Columns used to identify the samples: a) conjuntoMuestras and codigoExterno, b) codigoInterno c) id.

Importing values of dynamic attributes of samples

The import of the dynamic attribute values of the samples is analogous to the import of the samples. The import screen is the same as the samples but in Data to be imported, you must indicate Extra Sample Attributes. In the import file it is necessary to indicate the identification columns of the samples and a column with the code of each dynamic attribute to be imported (Figure 8). Both Samples and Attributes, must have been created before the import. During the import, TEGA will validate the values of each attribute, if there is any invalid it will not be imported and the row and the code of the related attribute will be reported. The format of each value depends on the type of the attribute:

- Character: Character: plain text, quotes are not necessary. Example: "Tools for Evolutionary and Genetic Analysis".

- Integer: integer values (zero, positive or negative). Examples: 0, 10, -20.
- Numeric: numerical values, it is possible to use scientific notation, the decimal point must be used. Examples: 0.5, -1.4, 100, 1e-3.
- Logical: possible values: 1 (true) or 0 (false).
- Date: available formats: yyyy-MM-dd, yyyy/MM/dd, yyyyMMdd (yyyy: year, MM: month, dd: day). Example: 2017-12-26, 2017/03/01, 20170121.

```
1 id,weight,temperature,receptionDate
2 1,11.33,9.67,20110202
3 2,15.42,10.07,20110203
4 3,13.22,8.36,20110304
```

Figure 8. Sample file attributes import of dynamic sample. First row shows the headline with dynamic attributes and the other rows correspond to the first three samples. In This example, the internal identificator (id) is used to identify each sample. Columns *weight*, *temperature* and *receptionDate* correspond to the code of dynamic attributes to be imported from each sample.

Importing Loci and Allele

It is possible to massively import the Loci and Alelos data from each sample that is already loaded on the platform.

It is possible to access the import screen of alleles through the Import button located in the query screen of the alleles. In the import screen it is necessary to indicate the file to import, the type of import, the separator character of the columns and, if the import type is "Matrix", the character used to indicate the null alleles and the status of the alleles (public or private). Once the file has been processed, it is possible to visualize the number of alleles and loci created, unidentified samples, unidentified loci, among other data (Figure 9).

As in the Samples file, the format is plain text, with columns separated by a character, and the first row indicates the name of the attributes or the name of the Loci.

There are two file types for this import:

- Matrix (Figures 10.a, 10.b and 10.c): The first columns of the file must be the identification column(s) of the samples (analogous to the sample import file). The rest of the columns correspond to the value of each allele in each Loci. For each Locus that you want to import, you must indicate the index at the end of the name of the Locus by separating it by a underscore. In case the locus does not exist, the value of the highest index will be created and indicated as ploidy. The value of each allele is of the character string type (max 255). Null alleles must be identified with a specific character (eg "?") which will also be indicated on the import screen. The Public attribute of the alleles will be completed with the value of the tilde titled Public of the import screen.
- List (Figure 10.d): available columns:
 - id: allele ID. Type: positive integer, eg 123.
 - idMuestra: sample ID. Type: positive integer, eg 123.
 - codigoInternoMuestra: internal code of the sample. Type: string (255 max.), eg "COD_INT|1".
 - idLocus: locus ID. Type: positive integer, eg 123.
 - codigoLocus: the locus code. Type: string (255 max.), eg "COD_LOCUS".

- indice: related to the index of the allele. Type: positive integer, eg 123.
- valor: related to the value of the allele. Type: string (255 max.), Eg "1010".
- publico: related to the public attribute of the allele. Type: Logical. Possible values: true or false

The column idMuestra or codigoInternoMuestra must be indicated, it is not necessary to indicate both. In the same way with the Locus (idLocus or codigoLocus).

If the id column is indicated, the platform will search for the allele in the database from the id. If not, look for the allele that is linked to the same sample and locus and have the same index, otherwise there is an allele with that data, a new one will be created. In case the locus does not exist it will be created.

The screenshot shows the 'Alleles Import' window. It includes a 'File' section with a file selector and 'alleles.csv' listed. Below is the 'Import Type' dropdown set to 'Matrix'. The 'Separator' is set to ',' and 'Null Value' is set to '?'. There is a 'Public' checkbox. A blue 'Import' button is located below these settings. At the bottom, there is a 'Detail' table showing the results of the import.

File and import configuration (points to the configuration fields)

Import button (points to the blue button with a plus icon)

Import result (points to the table below)

Detail	Value
Created Alleles	0
Modified Alleles	2198
Created Loci	0
Modified Loci	0
...	...

Back

Figure 9. Import screen of loci and alleles.

```

1 conjuntoMuestras,codigoExterno,LOCUS1_1,LOCUS1_2,LOCUS2_1,LOCUS2_2
2 SET1,EXT1,256,270,268,?
3 SET1,EXT2,242,242,194,194
a)

1 codigoInterno,LOCUS1_1,LOCUS1_2,LOCUS2_1,LOCUS2_2
2 CODE1,256,270,268,?
3 CODE2,242,242,194,194
b)

1 id,LOCUS1_1,LOCUS1_2,LOCUS2_1,LOCUS2_2
2 1,256,270,268,?
3 2,242,242,194,194
c)

1 codigoInternoMuestra,codigoLocus,indice,valor,publico
2 SET1|EXT1|1,LOCUS9,1,100,false
3 SET1|EXT1|1,LOCUS9,2,200,true
d)

```

Figure 10. Sample file for the import of alleles and loci. The columns are delimited by comma (.). In a), b) and c) the file type is Matrix, the first line is the header, which indicates the name of the loci to be created or updated. The second and third rows are sample data of the alleles of two samples. Columns used to identify the samples: a) conjuntoMuestras y codigoExterno, b) codigoInterno y c) id. d) File type List, the first line is the header, which indicates the name of the attributes to be imported. The second and third rows are sample data of the alleles of two samples.

Examples

In the *examples* folder of the platform there are sample import files, the separator character is the comma (.). Within the examples folder there is a folder by language (*en* and *es*), and within them the following folders:

- full: example with all fields (descriptive) of the samples
 - samples.csv: sample import file (162 samples).
 - alleles.csv: alleles import file, with matrix format (2198 alleles not null and 394 null).
- Rueda_et_al_2013: example with data from the work of Rueda et al., 2013.
 - samples.csv: sample import file.
 - alleles.csv: alleles import file, with matrix format.
- Kamvar_et_al_2014: example with data from the work of Kamvar et al., 2014.
 - samples.csv: sample import file.
 - alleles.csv: alleles import file, with matrix format.
- instructive: example files mentioned in the manual.
 - samples: examples of Figure 7 and 8.
 - samples_a.csv: example with identification columns: conjuntoMuestras y codigoExterno.
 - samples_b.csv: example with identification column: codigoInterno.
 - samples_c.csv: example with identification column: id.
 - samples_attributes.csv: example of importing dynamic attributes. Identification column: id.
 - alleles: examples of Figure 10.
 - alleles_a.csv: matrix format, with identification columns: conjuntoMuestras y codigoExterno.
 - alleles_b.csv: matrix format, with identification column: codigoInterno.

- alleles_c.csv: matrix format, with identification column: id.
- alleles_d.csv: list format.

Genotype Analysis

This section explains how to create a Genotype Analysis and execute the analysis procedures. TEGA has an entity called Genotype Analysis, aimed at the management of data linked to the execution of genotype analysis procedures.

Samples Sets

To create a Genotype Analysis, it is first necessary to assemble one or more Samples Sets that will group the Samples to be studied according to the necessary criteria, for example samples from different regions. It is possible to create a new set of samples from the query view (*Analysis / Sample Sets*) and it is necessary to indicate a unique code and the samples that will integrate it. Once the code has been completed and the samples selected, the Save button must be pressed (Figure 11).

Create or edit a Samples Set

ID

1

Code

SET1

Public ☐

Detail

Detail ...

Samples

Sample

+ 🔍

ID	Code	Detail
1	SET1 EXT1 1	Comments...
2	SET1 EXT2 2	Comments...
3	SET1 EXT3 3	Comments...
4	SET1 EXT4 4	Comments...



Attributes

Figure 11. Sample Sets edit view.

Genotype analysis

After creating the sample sets, a new Genotype Analysis must be created (Figures 12.a and 12.b), where the Project, the Samples Sets and Loci to be studied will be recorded. It is also possible to individually or massively select the Loci and the Samples Sets. Once the analysis is created, it is possible to visualize the alleles of each Sample and Loci in a matrix form by pressing the button titled "Alleles" in the query view (Figure 12.c).

On the other hand it is possible to execute the analysis procedures from the query view

(Figure 12.d), by pressing the button  Proc. . When doing so, the procedure execution screen will appear, which contains a tab for each procedure and in each one the corresponding parameters. Pressing the Submit button will start the execution of the procedure of the selected tab, with the indicated parameters. During the execution of the procedure, the analysis can not be modified or eliminated. It will also be possible to see the execution log from the button  Log in the query view.

At the end of the execution, the resulting files and the parameters used can be viewed in the Procedures Results section at the edit or display views of the genotypes analysis (Figure 12.e). In this section a drop-down item will appear for each execution made in the analysis.

a) Query view: Shows a table of genotype analysis records. The table has columns: ID, Code, State, Date, Project, and Detail. A single record is shown with ID 1, Code Analysis1, State Available, Date Apr 27, 2018, Project PROJECT 1, and Detail

b) Edit view: Form for creating or editing a genotype analysis. Fields include ID (1), Code (Analysis1), Project (PROJECT 1), and Date (2018-01-07). There are also checkboxes for State Available and Public.

c) Analysis allele editing view: Table showing alleles for a specific analysis. The table has columns: Samples Set, Sample, and multiple Locus columns (LOCUS1 to LOCUS6). It lists various sample sets and their corresponding allele values.

d) Analysis procedures execution view: Form for executing analysis procedures. It includes tabs for DAPC, Genepop, Indexes and Validations, and Structure. Under the Structure tab, there are sections for Additional parameters, CLUMPP parameters, Distruct parameters, and Structure: Main parameters. The main parameters section includes fields for Code, Detail, and Value for STRUCTURE_BURNIN (10000), STRUCTURE_NUMREPS (20000), and STRUCTURE_PLOIDY (2).

e) Procedures results section: Shows files and parameters for the execution of procedures. It includes a search bar and a list of files with their details and execution status.

Figure 12. Genotype analysis views: a) query view, b) edit view, c) analysis allele editing view, d) analysis procedures execution view, e) procedures results section of the edit view of genotypes analysis, there are visualized the resulting files and parameters of the execution of the procedures.


Analysis Procedures

TEGA has a module for the management of the analysis procedures where it is possible to create, edit and eliminate the procedures.

Procedures

The procedure query view is found in *Analysis / Procedures*. When creating a procedure it is necessary to indicate (Figure 13):

- Código: Code: must be unique and will be used to name the execution folders.
- Nombre: Name: used in the execution screen of procedures in the Genotype Analysis.
- Detail: description of the procedure.
- Execution command: command that will be used to execute the procedure. Example: `./script.sh`

- Command to show the execution log: this command will be used when the user presses the button  in the Genotype Analysis query. Example: `./script_log.sh`
 - Public: indicates whether the procedure can be seen by users with the Invited role.
 - Exports: this section indicates the files that TEGA will generate when executing the procedure:
 - Genotypes (TEGA): File of genotypes with the internal format of TEGA. The name of the generated file will be: `genotipos.tsv`. The columns are separated by tabs. Contains the columns:
 - `conjunto_id`: samples set ID.
 - `conjunto_cod`: samples set code.
 - `conjunto_det`: samples set detail.
 - `muestra_id`: sample ID.
 - `muestra_cod_int`: sample internal code.
 - Then it has a column for each Locus. In each column, the values of the alleles are indicated, separating them with the value of the TEGA parameter `EXP_PROC_SEP_ALELOS` (default `"|"`). The order of the alleles is indicated in the Index field. The value of the null alleles is the one indicated in the TEGA parameter `EXP_PROC_VALOR_NULO` (default `"-9"`).
 - Genotypes (Structure): Genotypes file with the Structure format. The name of the generated file will be: `genotipos.str`.
 - Genotypes (Genepop): Genotypes file with the Genepop [10] format. The name of the generated file will be: `genotipos.gen`.
 - Samples: File with the static fields of the samples. The name of the file will be: `muestras.tsv`.
 - Attrib. extras of Samples: File with the extra attributes of the samples. The name of the file will be: `muestras_atributos.tsv`. The ID column corresponds to the sample ID.
 - Samples Sets: File with the static fields of the Samples Sets. The name of the file will be: `conjuntos_muestras.tsv`.
 - Attrib. extras of Samples Sets: File with the extra attributes of the sample sets. The name of the file will be: `conjuntos_muestras_atributos.tsv`. The ID column corresponds to the ID of the sample set.
 - Loci: File with the static fields of Loci. The name of the file will be: `loci.tsv`.
 - Attrib. Loci extras: File with the extra attributes of the loci. The name of the file will be: `loci_atributos.tsv`. The ID column corresponds to the locus ID.
 - Quantities: file with the number of occurrences of the exported entities (Samples, Loci, Alleles and Samples Sets).
- In the *ejemplos/[language]/proc* folder there are sample files of each of these files. In the case of dynamic attribute files, each has five attributes, one for each type (character, numeric, integer, logical, and date).
- Files: the files necessary for the execution of the procedure. The files must be loaded after creating the procedure.

Create or edit a Procedure

ID

1

Code

structure

Name

Structure

Detail

Execution command

./structure.sh

Command to show the execution log

./log.sh

Public

Exports

Files

Cancel

Save

Exports

Genotypes (TEGA)☐

Genotypes (Structure)☒

Genotypes (Genepop)☐

Samples☐

Extra attrib. of Samples☐

Samples Sets☒

Extra attrib. of Samples Sets☐

Loci☐

Extra attrib. of Loci☐

Quantities☒

Files

Search

ID	Name	Detail	Public
1	structure.sh	structure.sh	<input checked="" type="checkbox"/>
2	structure	structure	<input checked="" type="checkbox"/>
3	CLUMPP	CLUMPP	<input checked="" type="checkbox"/>

Exports

Genotypes (TEGA)

Genotypes (Structure)

Genotypes (Genepop)

Samples

Extra attrib. of Samples

Samples Sets

Extra attrib. of Samples Sets

Loci

Extra attrib. of Loci

Quantities

Figure 13. Edit view of the analysis procedures.

Parameters and Procedures Categories

The Procedure Parameters allow the user to configure the execution of the procedure, for example indicate the number of repetitions, the range of clusters to be detected, the statistical method to be used, etc. The query view of these parameters is found in *Analysis / Procedure Parameters*.

When creating a Parameter it is necessary to indicate (Figure 14):

- Code: the code must be unique.
- Detail: description of the parameter.
- Type: Data type of the parameter, the available types are: Integer, Numeric, Character, Logical and Date.
- Procedure: procedure to which the parameter will be linked.
- Category: Parameter category. The categories allow grouping the parameters of a procedure into sections to facilitate its search when executing the procedure. The query view of categories is found in *Analysis / Categories of Parameters*.
- Not Editable: A non-editable parameter cannot be modified at the time of execution of the procedure.
- Value: Default value of the parameter.

Create or edit a Procedure parameter

ID
2

Code
STRUCTURE_BURNIN

Detail
Length of burnin period

Type
Integer

Procedure
structure

Category
STRUCTURE_MAIN

Not editable ☒

Value
10000

Cancel Save

Figure 14. Edit view of the Procedure Parameters.

Steps of the procedures execution

In this section the internal steps carried out by TEGA for the execution of the procedures are mentioned (the execution is done from the Genotype Analysis query):

1. The status of the Genotype Analysis is changed from Available to Executing.
2. A new occurrence of the Execution entity is created, which will be linked to the parameters of the procedures used and the resulting files.
3. The destination folder is created in which the input files and the resulting files will be stored. The address of this folder will be stored in the Execution:
[FILES]/analysis_genotipos/[ID_ANALYSIS]/[CODE_PROC]_[ID_EXECUTION]
Where:
 - FILES: is the folder indicated in the TEGA parameter DIRECTORIO_ENTIDADES.
 - ID_ANALYSIS: the ID of the selected Genotype Analysis.
 - CODE_PROC: code of the procedure to execute.
 - ID_EXECUTION: Execution ID.
4. The input files indicated in the Export section of the procedure to be executed are generated.
5. The file is generated with the procedure parameters selected by the user. This file contains both editable and non-editable parameters. The name of the generated file is: parametros.tsv. It has the columns:

- categoria: parameter category.
 - parametro: parameter code..
 - tipo: parameter type. Possible values: CARACTER (character), ENTERO (integer), NUMERICO (numeric), LOGICO (logic) y FECHA (date).
 - valor: parameter value.
6. The command for the execution of the procedures is executed, indicated in the TEGA COMANDO_PROCEDIMIENTOS parameter (by default `./proc/ejecutar_procedimiento.sh`). This script receives by parameter:
- Command to execute the procedure (Execution Command).
 - User of the operating system to be employed for the execution (parameter of TEGA USUARIO_PROCEDIMIENTOS). In case it is empty, the current user will be used.
 - Path of the folder where the procedure files are stored.
 - Destination path where the input files are located.
- The script will perform the following steps:
- a. Create a temporary folder and copy the procedure files to that folder.
 - b. In case of indicating an execution user, read and write permission will be assigned to the user in the temporary and destination folders. The execution permission will also be assigned to the files of the temporary folder created in a.
 - c. The current directory is changed to the one created in a.
 - d. The indicated command is executed. The following parameters are passed to the execution command:
 - i. Genotype analysis ID.
 - ii. Absolute path to the temporary folder created in a.
 - iii. Absolute path to the destination folder.
 - e. The temporary folder created in a is deleted.
6. At the end of the execution of the procedure, the following steps will be carried out within the same transaction:
- Linking the files in the destination folder to the Execution.
 - Change of status of the Genotype analysis, from Executing to Available.

Procedures available

The first version of TEGA has four modest implementations of the usual analysis procedures in population genetics studies.

Structure

This procedure seeks to determine the structure of the population using the Bayesian grouping method implemented in Structure. The procedure allows to indicate the parameters of each programs used. It is also possible to indicate the range of K (number of clusters) to be studied and repetitions by K. The procedure has the following steps:

1. Execution of Structure for the range of K and repetitions indicated. The parallel library is used to parallelize the execution

2. Execution of STRUCTURE HARVESTER [11] to determine the most suitable K. The one with the highest Delta K is selected.
3. Execution of CLUMPP [12] to integrate the repetitions of the selected K.
4. Execution of Distruct [13] to graph the assignment of clusters, both to individuals and populations.

The files resulting from the procedure are:

- a. `archivos_exp.zip`: contains the input files: `cantidades.tsv`, `conjuntos_muestras.tsv` y `parametros.tsv`).
- b. `structure.zip`: contains the input files and logs of the execution of Structure.
- c. `structure_log.zip`: contains the log files of each execution of Structure.
- d. `structure_out.zip`: contains the output files of Structure. This file can be used in other platforms such as CLUMPAK [14].
- e. `structureHarvester.zip`: contains output files of STRUCTURE HARVESTER.
- f. `clumpp.zip`: contains the CLUMPP input and output files.
- g. `distruct.zip`: contains the input and output files of DISTRUCT.
- h. `evanno.txt`: includes the table with the values of Delta K determined with STRUCTURE HARVESTER.
- i. `evanno.mk.txt`: has the K selected by the platform (largest Delta K).
- j. `distruct.ind.ps`: file in PostScript format generated with Distruct where the assignment of each group to each individual is visualized (generated from the selected K and the integration of the K repetitions made with CLUMPP).
- k. `distruct.pop.ps`: file in PostScript format generated with Distruct where the assignment of each group to each samples set is visualized (generated from the selected K and the integration of the K repetitions made with CLUMPP).
- l. `ejecucion.log`: log of the execution of the procedure.

Genepop

This procedure performs the execution of Genepop. It is possible to indicate the options to be executed and the parameters to be used. The resulting files are:

- `genotipos.gen`: genotype file in the Genepop format.
- `settings.txt`: Genepop input file that indicates the parameters and menu options to execute.
- `parametros.tsv`: procedure parameters.
- `genepop.log`: log of procedure execution.
- `genotipos.gen.*`: genepop output files corresponding to the menu options indicated.

DAPC

This procedure seeks to determine the structure of the population using the multivariate method DAPC (Discriminant analysis of principal components) [15]. The procedure has the following steps:

1. Determination of K (number of clusters): the function *find.clusters* of adegenet [16] package is used. It is possible to indicate the quantity of principal components (PC) to be used or the percentage of the variance; the statistical method to be used to determine the best K and the selection criterion of K, among other parameters. It is

possible to deactivate this step and perform the following steps with the groups determined by the samples sets.

2. Determination of PCs: the number of PCs to be used in DAPC is determined, this is done with a cross validation, using the *xvalDapc* function of adegenet package. It is possible to indicate the number of repetitions, the percentage of the training set, among other parameters. The procedure determines the number of PCs in two steps, first it executes *xvalDapc* with the range of PCs by default, then it executes *xvalDapc* for a range of PCs more bounded with center in the number of PCs selected in the first step. The number of PCs selected is the one that reports the least Mean Square Error.
3. Scatter and variable contribution graphs: scatter and variable contribution graphs are made, using the *scatter* and *loadingplot* function of adegenet package. In addition, the posterior membership probability at the sample level and the samples set is plotted.

Resulting files:

- *archivos_exp_logs.zip*: contains the input files and logs of the procedure (*dapc.log*, *genotipos.tsv*, *parametros.tsv*).
- *find_clusters.zip*: contains the output files of K determination.
 - *pc_var.**: image and table of the number of PCs retained by percentage of the variance used.
 - *k_stat.**: image and table of the value of the method used to determine K, by K.
- *cv_dapc.zip*: output files of PC determination (N indicates whether it is the first or second cross-validation):
 - *cvN.cvr.csv*: table with the result of the cross validation.
 - *cvN.rmsenpp.csv*: root mean squared error by number of PCs of PCA.
 - *cvN.png*: graph of ratio of hits by number of axes of PCA.
- *dapc.zip*: output files of the DAPC execution:
 - *pca_scatter.png*: scatter plot.
 - *pmp_indplot.png*: graph of posterior membership probability at the sample level.
 - *pmp_boxplot.png*: graph of posterior membership probability at the samples set level.
 - *loadingplot_*.png*: graphs of contribution of variables. Contribution of each allele to the main components. The number in the file name indicates the component.
 - *posterior.csv*: table with the posterior membership probability of each sample to each clusters.
 - *pmp.csv*: transformed table with the posterior membership probability of each sample to each clusters.
 - *var.contr.csv*: table with the contribution of each variable to each principal component.
- *k_stat.png*: image and table of the value of the method used to determine K, by K.
- *pmp_indplot.png*: graph of posterior membership probability at the sample level.
- *pmp_boxplot.png*: graph of posterior membership probability at the samples set level.

- resultados.csv: table with results of the methods (K selected in step a, number of PCs selected in step b, etc.).

Indexes and Validations

This procedure generates several descriptive and graphic indices. There are parameters to indicate the tables and graphs that will be generated. The generated files are:

- archivos_exp_logs.zip: input files and procedure logs.
- alelos.zip: contains tables and graphs of frequency and average of alleles per set of samples. Poppr functions are used for calculations.
- allelic_richness.csv: table with allelic wealth by Loci and Samples Sets. Only available for diploid loci. The *allelic.richness* function of hierfstat is used.
- basic_stat.csv: table with basic indexes (Fst, Fis, Ho, Hs, etc.). The *basic.stats* function of hierfstat is used.
- missing.*: table and graph of missing alleles. The *info_table* function of poppr is used.
- mlg.*: table and graph of the quantity of multilocus genotypes (MLG). The *mlg.table* function of poppr is used.
- mlg_rarefaction.png: rarefaction curve calculated with MLG. It uses the *rarecurve* function of vegan [17].
- ploidy.*: table and graph of ploidy by locus and sample. The *info_table* function of poppr is used.
- genotype_curve.*: table and graph of the curve of accumulation of genotypes. The *genotype_curve* function of poppr is used.
- genotypic_diversity.*: table and chart with diversity indexes. The *poppr* function of poppr is used.
- HW_*: tables with Hardy-Weinberg equilibrium test. The *hw.test* function of pegas [18] is used.
- locus_table_*: tables with statistics at the loci level. The *locus_table* function of poppr is used.
- dist_pop_method_*: distance between samples sets calculated with the method indicated by the user (see procedure parameters). The *dist.genpop* function of adegenet is used.

Examples

TEGA has sample files to test the procedures. These files correspond to the publications of Rueda et al. 2013 and Kamvar et al. 2014, although the examples focus on the *Structure* and *DAPC* processes, they can also be used to test *Genepop* and *Indexes and Validations*.

Structure: Rueda et al. 2013

In the work of Rueda et al. 2013, the seasonal variation in the population structure of the sábalo (*Prochilodus lineatus*) captured in the Uruguay River, near the city of Gualeguaychú (Entre ríos, Argentina), was analyzed. In this work Structure was used to analyze the population structure. In the folder *ejemplos/en/Rueda_et_al_2013* are the samples and alleles files.

The steps to replicate the analysis with TEGA are:

1. Import the samples.csv file: the import is done from the sample query (see the section Importing Samples), the comma (,) must be indicated as a separator, and Samples in *Data to be Imported*. When the samples are imported, the Sample Sets will be created automatically.
2. Import the file alleles.csv: the import is made from the Alleles query view (see section Importing of Loci and Alleles), you must indicate the type of import Matrix, the comma (,) as a separator and "?" as null value.
3. Create a new Genotype Analysis (see Genotyping Analysis section): in the creation of the analysis select the Loci: *PL3, PL14, PL25, PL34, PL35, PL64, PL119, PL139* and *PL216*; and the Sample Sets: *RUEDA13|WINTER, RUEDA13|SPRING* and *RUEDA13|FALL*.
4. Execute the Structure procedure (see Genotype Analysis section): the parameters must be indicated (the rest by default):
 - STRUCTURE_K_HASTA: 10
 - STRUCTURE_REPLICAS: 10
 - STRUCTURE_BURNIN: 150000
 - STRUCTURE_NUMREPS: 300000

As a result, it can be seen that the optimal number of clusters ($K = 3$) and the assignment of each to the samples corresponds to the work of Rueda et al. 2013 (Figure 15).

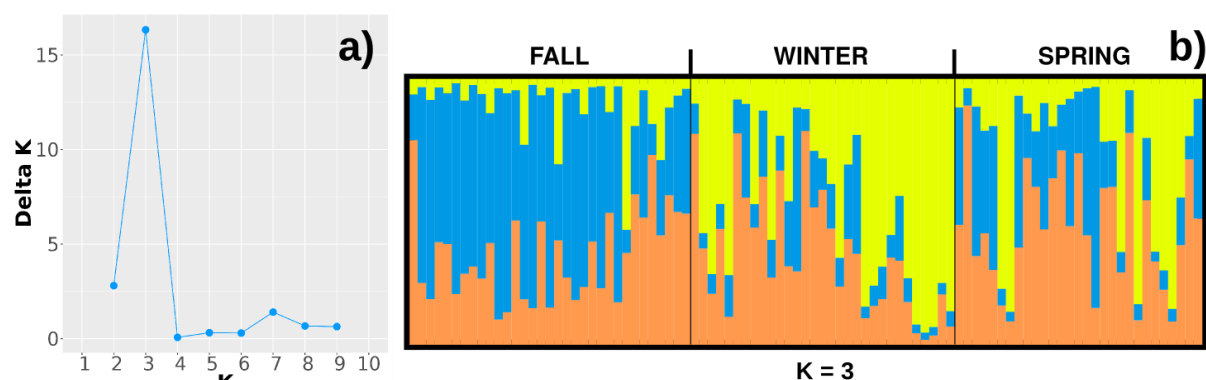


Figure 15. Example of the Structure procedure with data from Rueda et al. 2013. a) Plots of mean Delta K. b) Assignment probabilities for each individual (vertical bars) into one of three clusters.

DAPC: Kamvar et al. 2014

The Kamvar et al. 2014 data were used in the Kamvar et al.. 2015 [19] work, they analyzed the population structure of *Phytophthora ramorum* in the Oregon and California forests, using DAPC. In the folder *ejemplos/en/Kamvar_et_al_2014* are the samples and alleles files. The steps to replicate the analysis with TEGA are:

1. Import the samples.csv file: the import is done from the sample query (see the section Importing Samples), the comma (,) must be indicated as a separator, and Samples in *Data to be Imported*. When the samples are imported, the Sample Sets will be created automatically.
2. Import the file alleles.csv: the import is made from the Alleles query view (see section Importing of Loci and Alleles), you must indicate the type of import Matrix, the comma (,) as a separator and "?" as null value.

3. Create a new Genotype Analysis (see Genotyping Analysis section): in the creation of the analysis select the Loci: *PrMS6A1*, *Pr9C3A1*, *PrMS39A1*, *PrMS45A1* and *PrMS43A1*; and the Sample Sets: *JHallCr*, *NFChetHigh*, *Coast*, *HunterCr*, *Winchuck*, *ChetcoMain*, *PistolRSF*, *Nursery_CA* and *Nursery_OR*.
4. Execute the Structure procedure (see Genotype Analysis section): the parameters must be indicated (the rest by default):
 - DAPC_FC_EJECUTAR: NO.
 - DAPC_CV_REP: 250
 - DAPC_SP_P_BOX_NCOL: 3
 - DAPC_SP_P_IND_NCOL: 3
 - DAPC_SP_POSI_PCA: topright
 - DAPC_SP_P_BOX_XLEN: 11
 - DAPC_SP_P_BOX_YLEN: 11

As a result, it can be seen that HunterCr differs from the rest of the Sample Sets (Figure 16.a) and that the assignment of DAPC to each sample would correspond to the Samples Sets (without considering PistolRSF) (Figure 16.b). In addition, the 493 allele at *PrMS43A1* locus would contribute the most to that difference between sets (Figure 16.c and 16.d). These results would be similar to those of Kamvar et al. 2015.

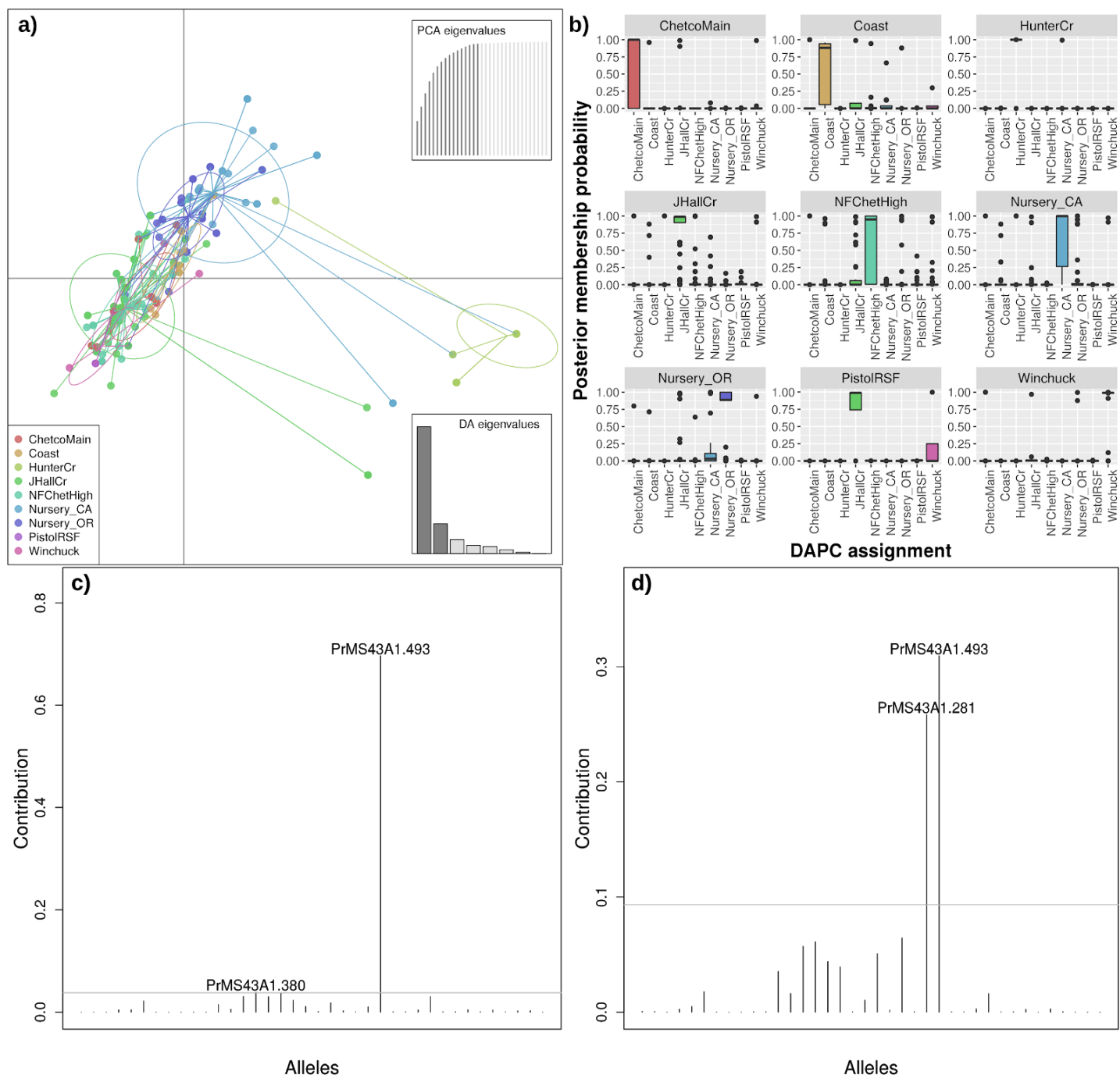


Figure 16. Results of the DAPC procedure with the data of Kamvar et al. 2014. a) Scatterplot (file `pca_scatter.png`). b) Boxplot made from the posterior membership probability of each sample to each Samples Set (file `pmp_boxplot_assig.png`). Contribution of the alleles to the first (c) and second (d) principal components of the analysis (files `loadingplot_1.png` and `loadingplot_2.png`). The greatest contribution is made by the 493 allele at PrMS43A1 locus.

Users

This section explains how user management is performed in TEGA.

As TEGA is focused to be used during the research period, in which the data is managed and analyzed, it is necessary to ensure the privacy of the data until the moment of its publication. For this reason there are different user roles:

- Anonymous: Has access to the public data of the Samples. Has permission to view and export the data, can not modify or delete them.
- Invited: Has access to the public data of the Samples, Alleles, Locus, Procedures and Genotype Analysis. Has permission to view and export the data, can not modify or delete them.
- Researcher: Has access to public and private data of the platform, can perform CRUDE actions and execute genotype analysis.
- Administrator: Has the same privileges as the Investigator role and also has access to the management of users and configurations of the platform.
- Implementer: Role for the implementers of the platform in the institution.

The Anonymous and Invited roles are intended for people outside the research group. When a person registers in the platform, he is assigned the role indicated in the TEGA parameter *ROL_USUARIO_DEFECTO*, which can be Anonymous (ANONIMO) or Invited (INVITADO, default value).

The roles of Administrator and Researcher would be used only by the research group.

Users with the Administrator role can access to the User query view (*Administration / User Management*) (Figure 17). There they can:

- Consult registered users in the platform, analogously to other views.
- Create, Edit and Delete users. While an administrator can create a new user, the recommended flow of user creation is:
 - a. The person interested in accessing to the platform must create their user on the home screen (link *Register a new account*, or the menu option *Account / Register*).
 - b. The platform will send an email to the indicated email address and the person must confirm the registration (by clicking on the link sent in the email). In this instance, the user will have the role of Anonymous.
 - c. The administrator must assign the desired role to the created user. This can be done from the user edit view.
 - d. Once the role has been modified, if the person had a session started on the platform, he must close the session and start it again to see the changes.
- Both from the query view and from the edit view, it is possible to change the user's status from On to Off and vice versa. A disabled user cannot log into the platform.

A screenshot of a web application window titled "Create or edit a user" with a close button (X) in the top right corner. The form contains several fields: "ID" with the value "5", "Login" with the value "researcher", "First name" (empty), "Last name" (empty), "Email" with the value "researcher@localhost", a checked checkbox for "Activated", "Language" with a dropdown menu showing "en", and "Role" with a dropdown menu showing "Researcher". At the bottom right, there are two buttons: "Cancel" and "Save".

Create or edit a user

ID

5

Login

researcher

First name

Last name

Email

researcher@localhost

☒ Activated

Language

en

Role

Researcher

Cancel Save

Figure 17. User edit view..

Data accessibility

This section explains how to change the status (from private to public and vice versa) of the data massively.

After publishing the results of an investigation, the users of the platform (with the role of Researcher or Administrator) can change the status of the project linked to the publication, passing it to the public (Figure 18). This will cause the Samples, Alleles, Loci and Genotypes Analysis linked to the project to be accessed by others users outside the institution (users with the Anonymous or Invited roles).

Create or edit a Project

ID
1

Code
PROJECT 1

Date
2018-01-07

Detail
Detail...

Public ☒

Attributes

Files

Cancel Save

Figure 18. Project edit view.

TEGA parameters

This section explains how to manage the TEGA parameters. These parameters allow modifying certain configurations of the platform, for example, the content of the home screen, the logo, the character used as a separator in exports, etc. The parameters can be modified from the parameter query view (*Administration / TEGA Parameters*), which can only be accessed by users with the Administrator or Implementer role (Figure 19).

Modifying or deleting a parameter could cause fail in the platform, so we recommend modifying them only when the effect is well understood. In turn, we also recommend having a testing environment to test different configurations.

The parameters have the following fields:

- Code: the code of the parameter, must be unique.
- Detail: detail of the use of the parameter in the platform.
- Type: data type of the parameter, the available types are: Integer, Numeric, Character, Logical and Date.
- Public: indicates if an anonymous user can access the parameter. Mainly used for parameters related to the home page.
- Value: value of the parameter, it can only be empty if the parameter type is Character.

After modifying a parameter, to visualize the change it is necessary to reload TEGA in the browser (shift + F5).

Create or edit a TEGA parameter

ID

5

Code

NOMBRE_PLATAFORMA

Detail

Name of the plataform

Type

Character

Public ☒

Value

TEGA

Cancel Save

Figure 19. TEGA parameter edit view.

Backups

TEGA has a basic procedure for backups generation. The configuration of the procedure is mentioned in the section *Installation and Configuration*.

Generation

It is possible to generate a backup from the menu option *Administration / Backup*, but only users with the role of Administrator or Implementer can access this option. On the other hand, it is also possible to execute the procedure from a task scheduler, such as cron. For this it is necessary to execute the procedure in the following way:

```
[DIR_TEGA]/proc/backups.sh CRON [DIR_TEGA]/config/backup.config.sh
```

Where:

- DIR_TEGA: path to TEGA folder
- CRON: parameter to tell the procedure that the execution is automatic.

The backup includes:

- All files located in the TEGA folder (backup copies are excluded).
- Copy of the database.

The files to be included in the backup are compressed and then encrypted with *pgp*.

Restore

The restoration of a backup is possible by executing the script:

```
[DIR_TEGA]/scripts/restore.sh [BACKUP] [ENCRYPTION_KEY] [HOST_DB] [USER_DB]  
[PASS_DB] [DB] [ENCODING] [LC_COLLATE] [LC_CTYPE]
```

Where:

- DIR_TEGA: path to TEGA folder
- BACKUP: path and file name of the backup file.
- ENCRYPTION_KEY: encryption key of the file (indicated in the backup configuration file).
- HOST_DB: host of the database engine.
- USER_DB: user with whom the database will be created and the data will be imported.
- PASS_DB: password of the DB user.
- DB: name of the database to create.
- ENCODING, LC_COLLATE y LC_CTYPE: parameters of the CREATE DATABASE command of PostgreSQL.

This procedure performs the following steps:

1. Decrypt the backup.
2. Unzip the backup in the current directory.
3. If database parameters are indicated (HOST_DB, USER_DB, etc.):
 - a. Remove, if it exists, the indicated database.
 - b. Create a new database with the indicated name.
 - c. Imports the data in the database.

In case of using a user or database other than the TEGA configuration, it is necessary to manually update the TEGA configuration (*config/*.yml* files). It is then possible to run TEGA as indicated in the *Installation and Configuration* section. You must ensure that the libraries and tools mentioned in the *Requirements* section are installed.

Development

In this section, the general aspects of the development of TEGA are mentioned.

General aspects

For the development of TEGA the project generator JHipster (<http://www.jhipster.tech>) was used. In the Back-End the JAVA language was used with the Spring Boot framework (<https://projects.spring.io/spring-boot>), and the PostgreSQL database engine (<https://www.postgresql.org>). While in the Front-End JavaScript was used with AngularJS (<https://angularjs.org>) and Bootstrap (<http://getbootstrap.com>). All the libraries used can be consulted in the TEGA repository (<https://github.com/darioelias/TEGA>).

Sources installation

Steps to install the TEGA sources:

1. Install the requirements of the *Requirements* section
2. Install JAVA 8 Development Kit (JDK). OpenJDK 8 installation guide: <http://openjdk.java.net/install/>
3. Install git v2.1.4. Git installation guide: <https://git-scm.com/download/linux>
4. Install MAVEN (<https://maven.apache.org>)
5. Install Node.js v4.7.0 (<https://nodejs.org/en>)
6. Update NPM. Command: `npm install -g npm`
7. Install bower y gulp-cli. Command: `npm install -g bower gulp-cli`
8. Clone the github TEGA repository. Command:
`git clone git://github.com/darioelias/TEGA.git`
9. Copy the *config_ejemplo* folder and rename it by *config*
10. Create a database and configure TEGA. Steps 3 to 10 of the section *Installation and Configuration*. In addition to configuring the *application-prod.yml* file, you must configure the *application-dev.yml* file.
11. Running in development: the *dev_comp.sh* script allows you to run maven with the development configuration.
12. Execution in production: the *prod_comp.sh* script allows you to run maven with the configuration to build and package the platform (generating the WAR file in the *target* folder). For more information consult the JHipster documentation or the *jhipster.readme.txt* file.

Entity-relationship diagram

The entity-relationship diagram is only illustrative (Figure 20) and can be viewed with the MySQL Workbench (<https://www.mysql.com/products/community>). The model file is located in the *model* folder of the git repository of TEGA.

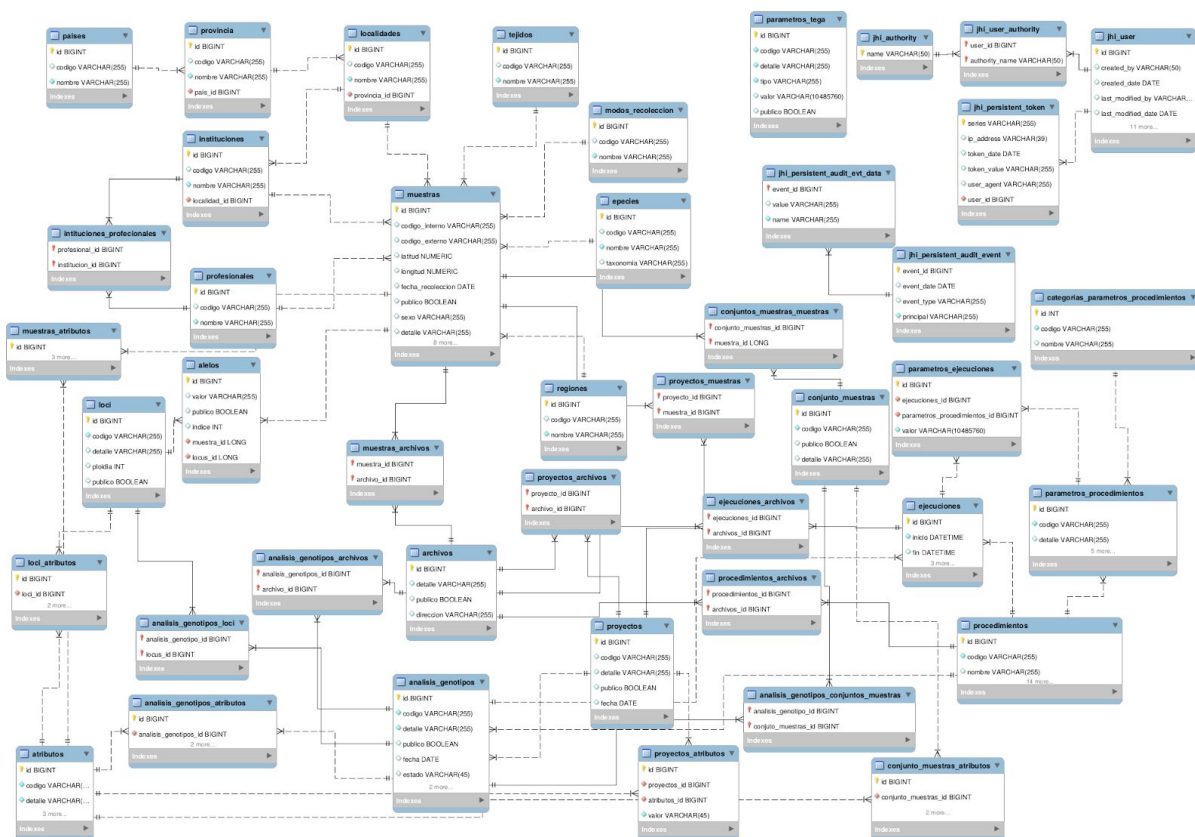


Figure 20. Diagram of entity-relationship of TEGA displayed with MySQL Workbench. The diagram is only illustrative.

Bibliography

1. Charlesworth B, Charlesworth D. Population genetics from 1966 to 2016. *Heredity*. 2017;118: 2–9. doi:10.1038/hdy.2016.55
2. Pritchard JK, Stephens M, Donnelly P. Inference of population structure using multilocus genotype data. *Genetics*. 1998;155: 945–959.
3. Rueda EC, Carriquiriborde P, Monzón AM, Somoza GM, Ortí G. Seasonal variation in genetic population structure of sábalo (*Prochilodus lineatus*) in the Lower Uruguay River. *Genetica*. 2013;141: 401–407. doi:10.1007/s10709-013-9739-0
4. Kamvar ZN, Larsen MM, Kanaskie AM, Hansen EM, Grünwald NJ. Sudden_Oak_Death_in_Oregon_Forests: Spatial and temporal population dynamics of the sudden oak death epidemic in Oregon forests. *Zenodo*. 2014.
5. Nosek BA, Alter G, Banks GC, Borsboom D, Bowman SD, Breckler SJ, et al. SCIENTIFIC STANDARDS. Promoting an open research culture. *Sci*. 2015;348: 1422–1425. doi:10.1126/science.aab2374
6. Fecher B, Friesike S. Open Science: One Term, Five Schools of Thought. *Opening Science*. Springer International Publishing; 2012. pp. 17–47. doi:10.1007/978-3-319-00026-8_2
7. Tange, O. (2011). Gnu parallel-the command-line power tool. *The USENIX Magazine*, 36(1), 42-47.
8. Kamvar, Z. N., Tabima, J. F., & Grünwald, N. J. (2014). Poppr: an R package for genetic analysis of populations with clonal, partially clonal, and/or sexual reproduction. *PeerJ*, 2, e281.
9. Goudet, J. (2005). Hierfstat, a package for R to compute and test hierarchical F-statistics. *Molecular Ecology Resources*, 5(1), 184-186.
10. Rousset, F. (2008). genepop'007: a complete re-implementation of the genepop software for Windows and Linux. *Molecular ecology resources*, 8(1), 103-106.
11. Earl, D. A. (2012). STRUCTURE HARVESTER: a website and program for visualizing STRUCTURE output and implementing the Evanno method. *Conservation genetics resources*, 4(2), 359-361.
12. Jakobsson, M., & Rosenberg, N. A. (2007). CLUMPP: a cluster matching and permutation program for dealing with label switching and multimodality in analysis of population structure. *Bioinformatics*, 23(14), 1801-1806.
13. Rosenberg, N. A. (2004). DISTRUCT: a program for the graphical display of population structure. *Molecular Ecology Resources*, 4(1), 137-138.
14. Kopelman, N. M., Mayzel, J., Jakobsson, M., Rosenberg, N. A., & Mayrose, I. (2015). Clumpak: a program for identifying clustering modes and packaging population structure inferences across K. *Molecular ecology resources*, 15(5), 1179-1191.
15. Jombart, T., Devillard, S., & Balloux, F. (2010). Discriminant analysis of principal components: a new method for the analysis of genetically structured populations. *BMC genetics*, 11(1), 94.
16. Jombart, T. (2008). adegenet: a R package for the multivariate analysis of genetic markers. *Bioinformatics*, 24(11), 1403-1405.
17. Jari Oksanen, F. Guillaume Blanchet, Michael Friendly, Roeland Kindt, Pierre Legendre, Dan McGlinn, Peter R. Minchin, R. B. O'Hara, Gavin L. Simpson, Peter Solymos, M. Henry H. Stevens, Eduard Szoecs and Helene Wagner (2017). *vegan*:

Community Ecology Package. R package version 2.4-2.

<https://CRAN.R-project.org/package=vegan>

18. Paradis, E. (2010). pegas: an R package for population genetics with an integrated–modular approach. *Bioinformatics*, 26(3), 419-420.
19. Kamvar, Z. N., Larsen, M. M., Kanaskie, A. M., Hansen, E. M., & Grünwald, N. J. (2015). Spatial and temporal analysis of populations of the sudden oak death pathogen in Oregon forests. *Phytopathology*, 105(7), 982-989.

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