

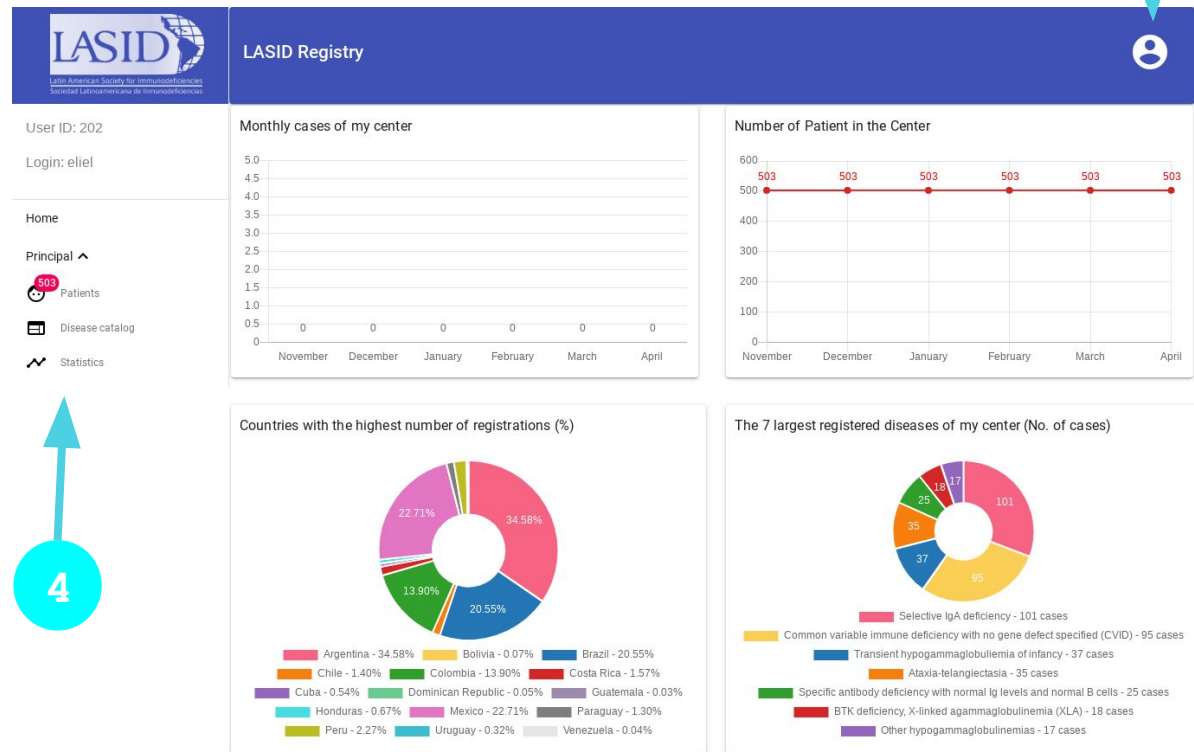
Tutorial



Latin American Society for Immunodeficiencies
Sociedad Latinoamericana de Inmunodeficiencias

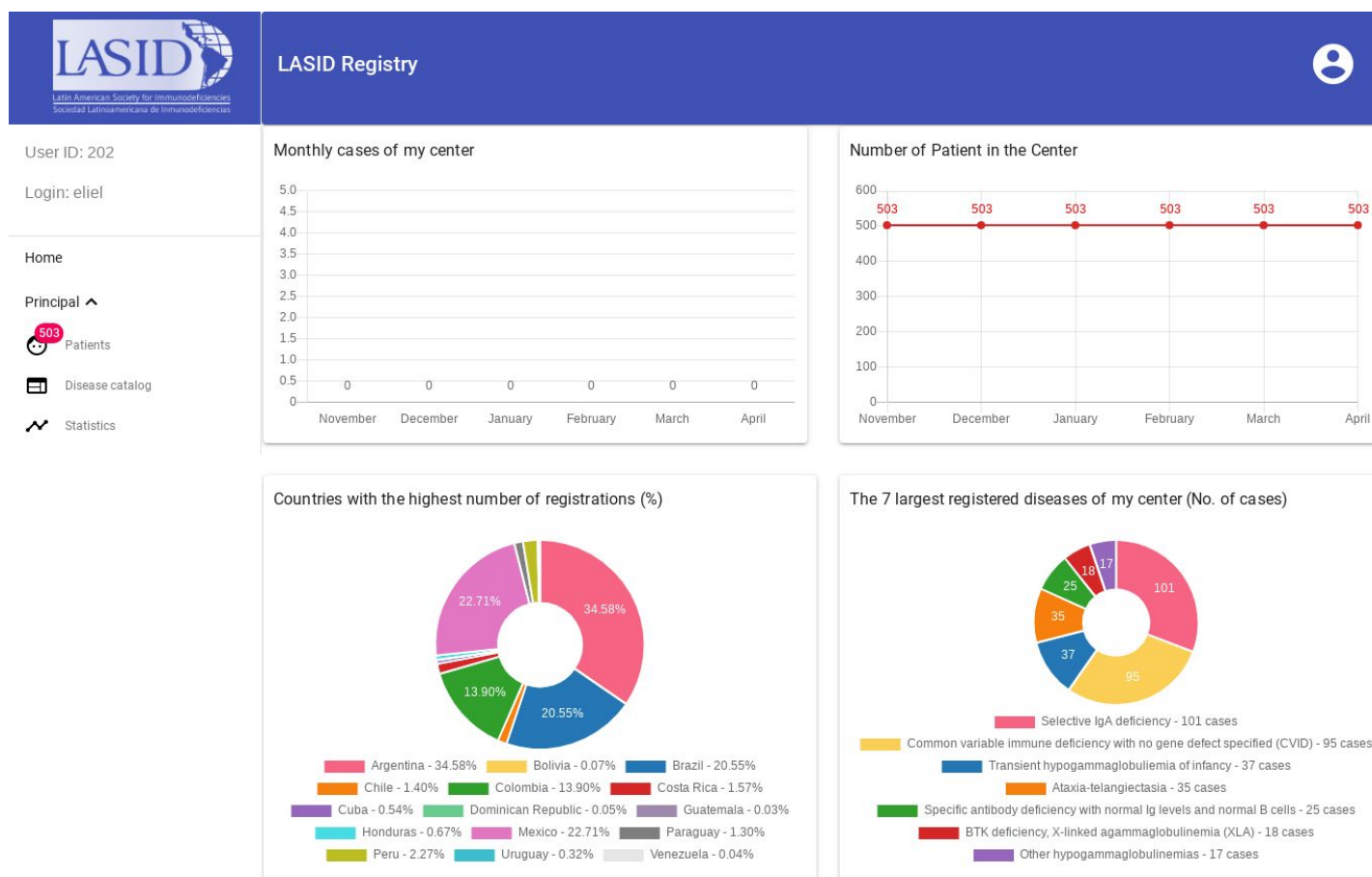
How to Use the LASID Registry Platform

1. Go to <http://lasidregistry.org>. Click on SIGNIN.
2. You will be redirected to the login page.
3. Kindly enter your LASID Registry user ID and password to enter the Home Page. The system will send new user ID and password for members.
4. HOME PAGE. This is the main navigation page. It is divided into 4 sections:
 - a. Home
 - b. Patients
 - c. Disease catalog
 - d. Statistics
5. Menu
 - a. My Account
 - b. My center
 - c. Change password
 - d. Logout



Home

This is the main registry page. You can monitor data of your center.



Patients

All your patients are in this section. You can **search**, **add**, **view**, **edit** and **delete** patient data.

LASID
Latin American Society for Immunodeficiencies
Sociedad Latinoamericana de Inmunodeficiencias

Patients

User ID: 202
Login: eliel

Home

Principal ^

503 Patients

Disease catalog

Statistics

Search


+ 🔍 ✎ 🗑

	Id ↑	Name	Sex	Birth	PID Diagnosis
<input checked="" type="checkbox"/>	8088		Female	2001-11-11	Specific antibody deficiency with normal Ig levels and normal B cells
<input type="checkbox"/>	8082		Male	2015-07-06	Transient hypogammaglobulinemia of infancy
<input type="checkbox"/>	8060		Male	2000-03-09	Common variable immune deficiency with no gene defect specified (CVID)
<input type="checkbox"/>	8058		Male	1969-08-30	Specific antibody deficiency with normal Ig levels and normal B cells
<input type="checkbox"/>	8049		Female	1977-02-11	Secondary hypogammaglobulinemia
<input type="checkbox"/>	8032		Male	1945-12-05	Specific antibody deficiency with normal Ig levels and normal B cells
<input type="checkbox"/>	8009		Male	1985-06-16	Common variable immune deficiency with no gene defect specified (CVID)
<input type="checkbox"/>	8008		Male	1945-11-15	Common variable immune deficiency with no gene defect specified (CVID)
<input type="checkbox"/>	7998		Male	2016-05-07	T-B- SCID with unknown gene defect
<input type="checkbox"/>	7773		Male	2006-11-23	C2 deficiency

10 1-10 of 503 |< < > >|

New Patient

Fill sections of Patient Info, Diagnosis, Therapy, Transplant and Visit History. Then click on **SAVE**. Fields with * are required in sections patient info and diagnosis (same unknown).




Latin American Society for Immunodeficiencies
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User ID: 202


Login: eliel

Home


Principal ^

 503 Patients

 Disease catalog

 Statistics

← New patient



PATIENT INFO

DIAGNOSIS

THERAPY

TRANSPLANT

VISIT HISTORY

Name*

Date of birth*
mm/dd/yyyy

Date of death
mm/dd/yyyy

Patient consent

Sex*

Country of living*

Familial case

Consanguinity

Patient # in sibship

Blood Type

Rh System

Address

Address

Zip Code

City

State/Province

Country

Phone

Mobile

Email

SAVE

New Patient

Sections Therapy, Transplant and Visit History

Those sections allow input in datatable. Click on **add**. After fill click on **UPDATE** to add in **datatable**.

← New patient

PATIENT INFO DIAGNOSIS **THERAPY** TRANSPLANT VISIT HISTORY

** You must update before saving the patient.*

Drug grup*
Acido aminocaproico +

Generic name
13 valent conjug.vac +

Brand name
Acantex +

Therapy start*
04/08/2019

Therapy stopped
mm/dd/yyyy

Reason stopped
Drug course completed

Side effects
No

Dose
12

Dose unit
Ampolla +

Interval
2

Times per
Week







Route of adm.
I.V.

Compliance*
Excellent

UPDATE CANCEL

← New patient

PATIENT INFO DIAGNOSIS **THERAPY** TRANSPLANT VISIT HISTORY

Therapy start	Therapy stop	Drug group	Generic name	Brand name	Options
2019-04-08		Acido aminocaproico	13 valent conjug.vac	Acantex	  
2019-03-05		Acyclovir	aciclovir	Acifol	  

Edit Patient

Allows **edit** the data of the selected patient: **change** Diagnosis and modify Therapy, Transplant and Visit History (**add** / **edit** / **delete** / **view**)

Patients

Search

+ 🔍 ✎ 🗑

	Id ↑	Name	Sex	Birth	PID Diagnosis
<input checked="" type="checkbox"/>	<u>8495</u>	Arthur Marlos	Male	1970-01-01	Wiskott-Aldrich syndrome (WAS)
<input type="checkbox"/>	<u>8088</u>		Female	2001-11-11	Specific antibody deficiency with norm

← Edit patient #8495

PATIENT INFO **DIAGNOSIS** THERAPY TRANSPLANT VISIT HISTORY

Diseases type
Combined immunodeficiencies with associated or syndromic features

Disease sub

3

4

← Edit patient #8495

DIAGNOSIS THERAPY TRANSPLANT **VISIT HISTORY**

Visit Date	Weight (kg)	Height (cm)	Options
2018-08-15	89	170	✎ 🗑 🔍

+

SAVE

Disease catalog

Shows the tree with 373 diseases.

Disease catalog

- + Immunodeficiencies affecting cellular and humoral immunity
- + Combined immunodeficiencies with associated or syndromic features
- + **Predominantly Antibody Deficiencies**
- + Diseases of Immune Dysregulation
- + Congenital defects of phagocyte number or function
- + Defects in Intrinsic and Innate Immunity
- + Autoinflammatory Disorders
- + Complement Deficiencies
- + Phenocopies of Inborn Errors of Immunity
- + Unknown

Predominantly Antibody Deficiencies

- + Severe Reduction in All Serum Immunoglobulin Isotypes with Profoundly Decreased or Absent B Cells, Agammaglobulinemia
- + Severe Reduction in at Least 2 Serum Immunoglobulin Isotypes with Normal or Low Number of B Cells, CVID Phenotype
- + Severe Reduction in Serum IgG and IgA with Normal/Elevated IgM and Normal Numbers of B cells, Hyper IgM
- + Isotype, Light Chain, or Functional Deficiencies with Generally Normal Numbers of B Cells

Severe Reduction in All Serum Immunoglobulin Isotypes with Profoundly Decreased or Absent B Cells, Agammaglobulinemia

- BTX deficiency, X-linked agammaglobulinemia (XLA)
- ? heavy chain deficiency
- ?5 deficiency
- Ig? deficiency
- Ig? deficiency
- BLNK deficiency
- PIK3R1 deficiency
- E47 transcription factor deficiency
- Agammaglobulinemia with unknown gene defect

Severe Reduction in All Serum Immunoglobulin Isotypes with Profoundly Decreased or Absent B Cells, Agammaglobulinemia

- BTX deficiency, X-linked agammaglobulinemia (XLA)
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- E47 transcription factor deficiency
- Agammaglobulinemia with unknown gene defect

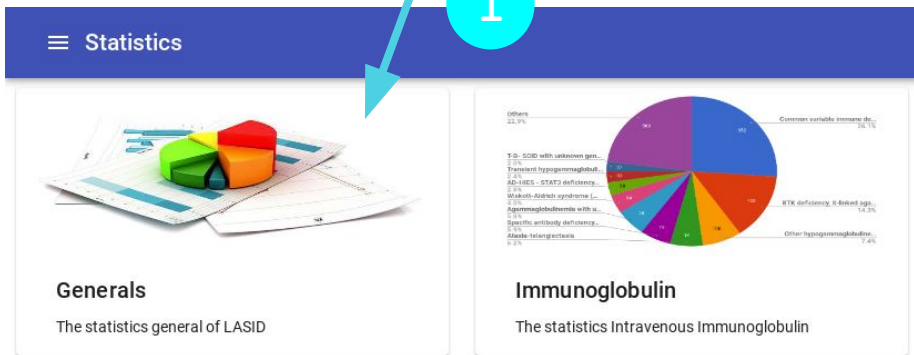
Disease information

Code	BLNK	Name	BLNK deficiency
OMIM	604515		

CLOSE

Statistics

Displays the **general** and **immunoglobulin** monthly cases of the last 15 months.

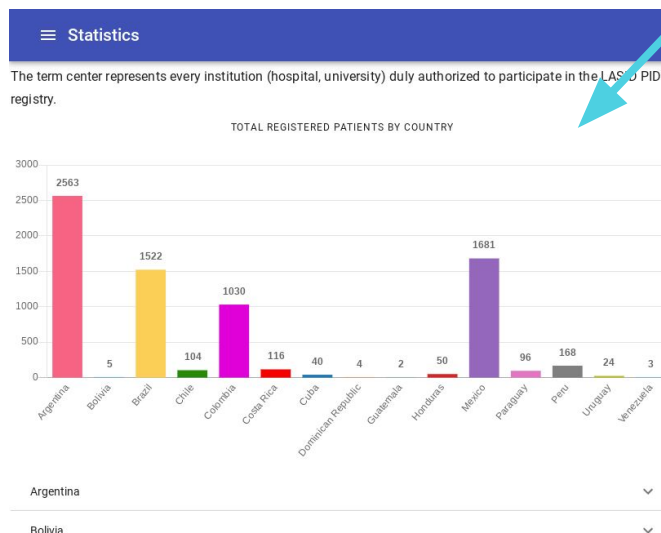


Statistics

2

March/2019	February/2019	Jan
December/2018	November/2018	Oct
September/2018	August/2018	Jul
June/2018	May/2018	Apr
March/2018	February/2018	Jan

The figure shows a table with three columns. The first column contains dates from March/2019 down to March/2018. The second column contains dates from February/2019 down to February/2018. The third column contains the first letter of the month. A red circle with the number '2' is placed over the table.



My account

Allows edit of account data.



My account

Center ID	
1120	
ID	
202	
Username	
eliel	
Name	
Eliel Gonçalves	
Professional License	Specialty Certificate
123456789	123456789
Email	Mobile
elielgoncalves@gmail.com	19 12341234

SAVE

CANCEL

My center

Allows edit of center data.



My center

Center ID

1120

Name

EPM-UNIFESP

Institution

Escola Paulista de Medicina - UNIFESP

Country *

Brazil

State/Province

▼

SP

City

São Paulo

Address

R. Botucatu, 720

Phone

11 5579-6700

Zip Code

04023-062

SAVE

CANCEL

Change password

Allows change your password.



Change password

Current password *

New password *

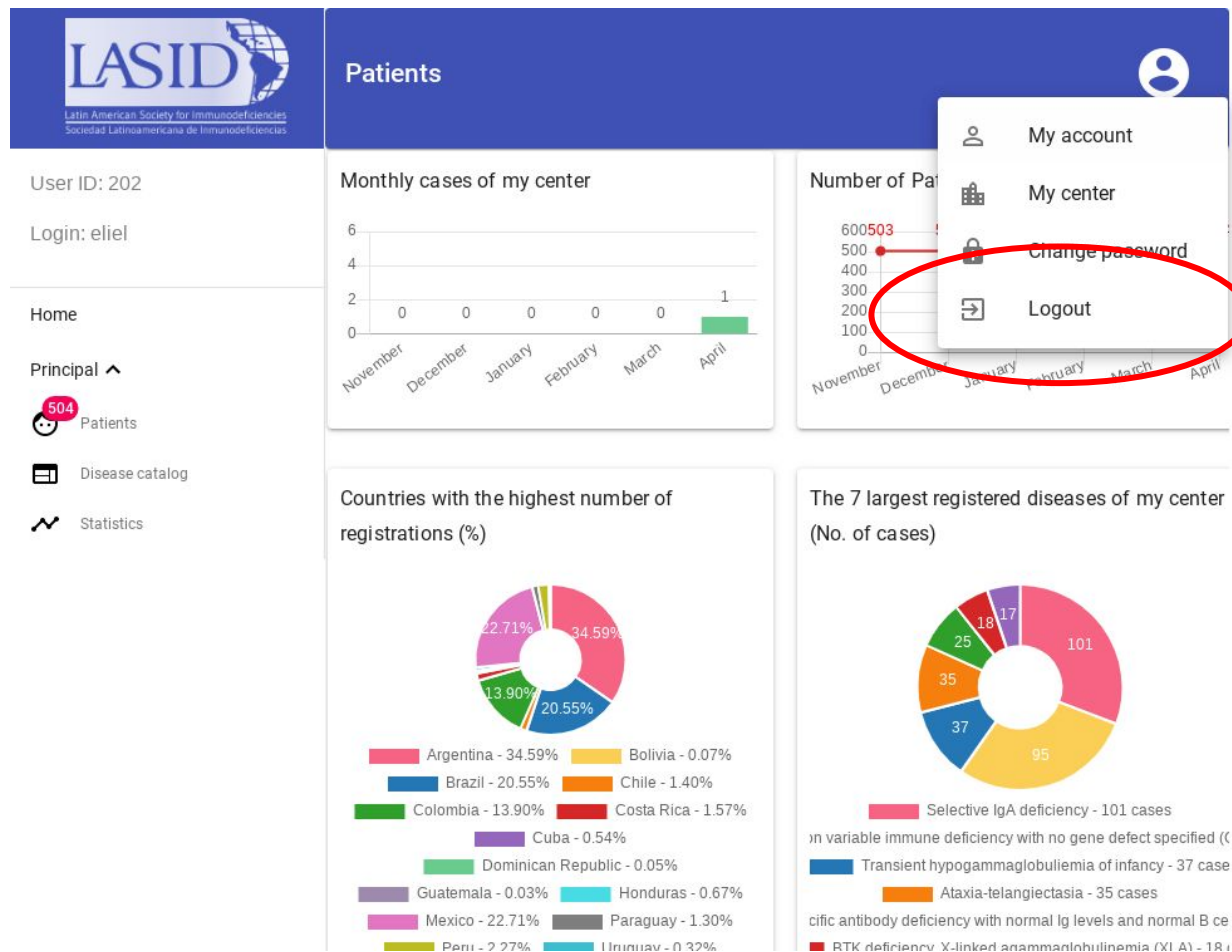
Confirm new password *

SAVE

CANCEL

Logout

Will end your session of access to this system.



Thank You

LASID Registry administrators

Eliel Gonçalves and Gilberto Savoini

lasidregistry@gmail.com

