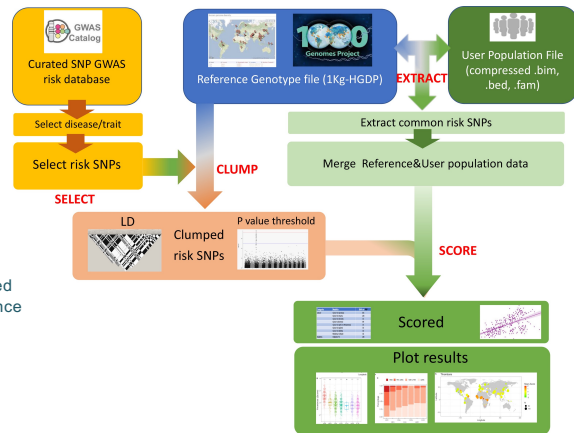
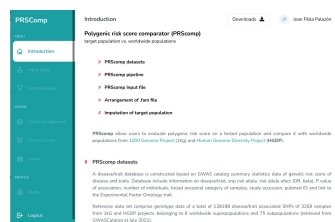


## P19.005.D · Polygenic risk score comparator (PRScmp): tested population vs. worldwide populations

Joan Fibla <sup>1</sup>, Leandre Palau <sup>2</sup>, José David Nunes <sup>2</sup>, Marina Laplana <sup>3</sup>, Ricard López <sup>4</sup>

<sup>1</sup> Universitat de Lleida, IRB Lleida; <sup>2</sup> Grup Globalia, Lleida, España; <sup>3</sup> Universitat de Lleida, Departament de Ciència Animal; <sup>4</sup> Hospital Universitari Arnau de Vilanova, Unitat de Citogenètica i Genètica Mèdica. IRB Lleida.  
mail to: joan.fibla@udl.cat

We have developed a web service "**Polygenic Risk Score Comparator (PRScmp)**", which allows users to assess the polygenic risk score of their own population and compare it with worldwide populations from the 1000 Genome Project (1Kg) <sup>1</sup> and the Human Genome Diversity Project (HGDP) <sup>1</sup>.



1

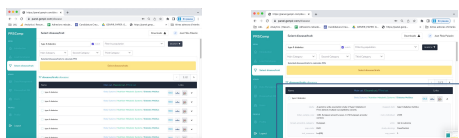
The user loads the genotypes of his population

2

Genotypes are merged with data from reference populations

3

User selects one or more disease/trait database entries (based on GWAS Catalog summary statistics <sup>2</sup>)



4

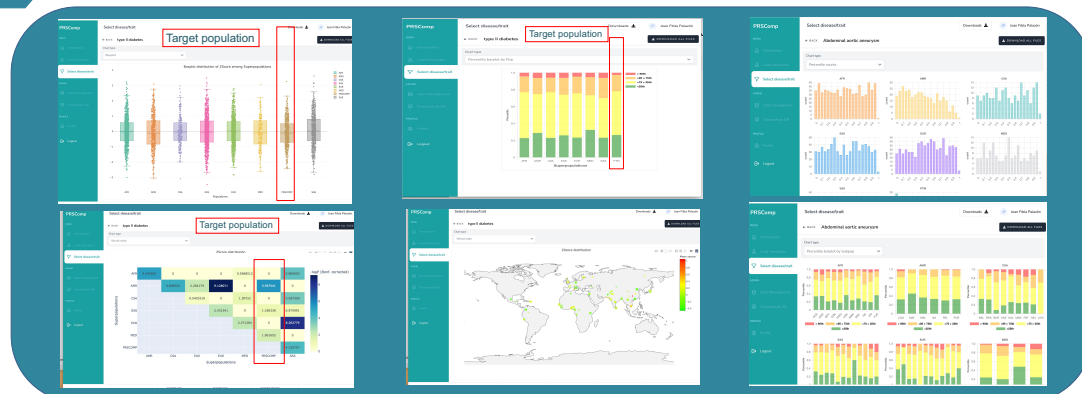
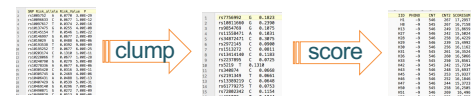
A file is generated with the selected risk markers (SNPs), which will be used by the Plink1.9<sup>3</sup> program to obtain the risk score (PRS) of the user's population together with the reference world populations.

5

The distribution of **PRS** z-score values of the user's population and of the 1Kg and HGDP populations are represented by a box plot, a bubble plot on the world map (mean values) and a bar plot for the distribution of percentile values.

Differences among populations are tested by paired-test with Bonferroni correction.

6



**PRScmp** offers the opportunity to assess the risk associated with diseases in a population that could be of great interest in the planning and monitoring of public health strategies.

### References:

1. Birney, E., Soranzo, N. The end of the start for population sequencing. *Nature* **526**, 52–53 (2015)
2. Welter, D. et al. The NHGRI GWAS Catalog, a curated resource of SNP-trait associations. *Nucleic Acids Res* **42**, D1001–D1006 (2013).
3. Purcell, S. et al. PLINK: A Tool Set for Whole-Genome Association and Population-Based Linkage Analyses. *The American Journal of Human Genetics* **81**, 559–575 (2007).

All authors declare no conflicts of interest

## P19.005.D · Polygenic risk score comparator (*PRScomp*): tested population vs. worldwide populations

Joan Fibla <sup>1</sup>, Leandre Palau <sup>2</sup>, José David Nunes <sup>2</sup>, Marina Laplana <sup>3</sup>, Ricard López <sup>4</sup>  
Universitat de Lleida, IRBLleida; 2 Grup Globalia, Lleida, España; 3 Universitat de Lleida, Departament de Ciència Animal; 4 Hospital  
Universitari Arnau de Vilanova, Unitat de Citogenètica i Genètica Mèdica. IRBLleida.  
mail to: [joan.fibla@udl.cat](mailto:joan.fibla@udl.cat)

### How to test PRScom:

Download to your computer example genotype file from:

<https://figshare.com/s/5c52c34cf097a2ccf039>

Login at PRScom as: (select one of them)

email: [anonymous1@genpir.com](mailto:anonymous1@genpir.com) password: anonymous1  
email: [anonymous2@genpir.com](mailto:anonymous2@genpir.com) password: anonymous2  
email: [anonymous3@genpir.com](mailto:anonymous3@genpir.com) password: anonymous3  
email: [anonymous4@genpir.com](mailto:anonymous4@genpir.com) password: anonymous4  
email: [anonymous5@genpir.com](mailto:anonymous5@genpir.com) password: anonymous5  
email: [anonymous6@genpir.com](mailto:anonymous6@genpir.com) password: anonymous6  
email: [anonymous7@genpir.com](mailto:anonymous7@genpir.com) password: anonymous7  
email: [anonymous8@genpir.com](mailto:anonymous8@genpir.com) password: anonymous8  
email: [anonymous9@genpir.com](mailto:anonymous9@genpir.com) password: anonymous9

Go to “Load Genotypes” and click on “Import data” and load example genotype (example\_genotype.zip) file from your computer.

Fill required data:

Population identifier: (ie. Example genotype file)

Pop code: (ie. “EXA”) (use a three-letter code)

Coordinates lat/lon

Lat: (ie -12)

Lon: (ie -12)

Upload file and perform “extract” and “merge” steps

Your file is ready to be analyzed by PRScom. “Go to disease” and select the disease/trait of your interest.

Thanks for your interest.

Download this file by scanning the QR code:

