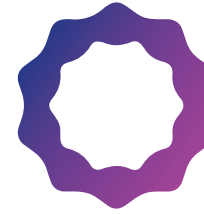
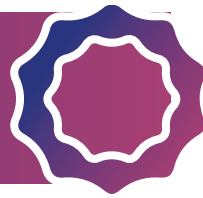


Where can I get more information?



Andalusian Registry of **DONORS** for Biomedical **RESEARCH**

BECOME A DONOR
of samples
for biomedical
research



Biobanco del Sistema Sanitario Público de Andalucía
Consejería de Salud y Consumo

Parque Tecnológico Ciencias de la Salud/ Centro de Investigación Biomédica
Avda. del Conocimiento s/n, 18016 Granada, Spain
Phone +34 958 894 669 – Email: registro.donantes.csalud@juntadeandalucia.es
www.soydonanteparainvestigacion.es

www.soydonanteparainvestigacion.es



LEGAL FRAMEWORK

The Registry was constituted by the Order of June 15, 2015, which, under the auspices of the Ministry of Equality, Health, and Social Policies, created the personal data file entitled: "Donors of Samples for the Biomedical Research of Andalusia". This order was published in the Official Gazette of the Junta de Andalucía on June 23, 2015

LOPD CONDITIONS

When you send your data by email or phone, you consent for your data to be processed to address the inquiries sent and provide information about our Donor Registry services. Your data will not be transferred to a third party unless there is a legal obligation to do so. You can exercise your rights of access, rectification, deletion, portability of your data, and the limitation or opposition to its treatment by requesting to exercise your rights via the <https://donantes.biobancospa.es/biobank> platform or directly to the General Secretariat of Public Health, R+D+I, and Health at the Ministry of Health and Consumption, which is the entity that is responsible for the dealing with issues related to data protection, as indicated in the following link: <https://juntadeandalucia.es/protecciondedatos.html>.
Not sure what the stands for so I can't translate it

What is the Registry?



The Registry is a tool that allows the citizens of Andalusia to participate directly in the development of disease research and in the improvement of health.

The Registry consists of a database where the basic identification and health data of people interested in voluntarily participating in biomedical research are collected. It is a registry that has been endorsed by the Coordinating Committee of Ethics of Biomedical Research of Andalusia and is subject to all legal guarantees regarding confidentiality. The Registry is under the responsibility and custody of the Ministry of Health.

The Registry is managed by the Research Area of the Andalusian Public Health System Biobank (www.biobancosspa.com).

Citizens, researchers, patient associations, and health institutions are thus provided with another opportunity to promote research together.

How to become a donor?

You can become a donor by going to any of the Biobank nodes, calling by telephone, writing an email or contacting us directly via the internet.



958 894669



registro.donantes.csalud@juntadeandalucia.es



www.soydonanteparainvestigacion.es

**BECOME
A DONOR**
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research

How does it work?



When a researcher needs samples, the professionals at the the Research Area of the SSPA Biobank will search the Registry for potential donors of samples that fit the characteristics specified by the researcher. The donor will be asked whether or not they wish to participate in the researcher's research project. If the donor accepts, the required sample(s) will be obtained at a health centre.



What is donated?

Human biological samples and/or data that are obtained by non-invasive collection processes include hair, faeces, body fluids (urine, sweat, and saliva), or by minimally invasive processes, including blood samples.

Where do you donate?

The donation is always made to the SSPA Biobank. The place where the donation will be made will be at a Public Health Centre in cases where the sample to be donated requires health personnel to obtain it. In cases where the sample does not require specialized personnel for it to be obtained, then the donation can take place at the donor's home.



Who can donate?

Users of the Public Health System of Andalusia, whatever their age, state of health, or lifestyle habits.

Users of private health companies who provide a descriptive report of their health status at the time of registration and realization of the donation.