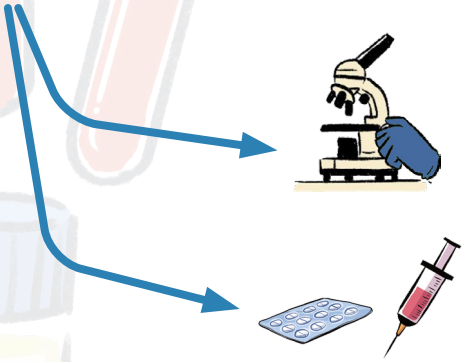
LEGAL DEPARTMENT[1] / MEDICAL RESEARCH DEPARTMENT[2]

Secondary use of samples: useful information and contacts

Human samples (primary and their derivatives) have already been collected in France or abroad:

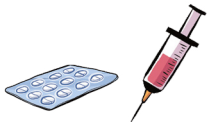


- For previous research,
- As part of routine care/diagnosis (residual samples taken from hospitals, clinics, medical analysis laboratories, etc.).

In order to be able to use them for new research, these samples must <u>first</u> have obtained:

1

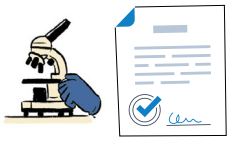
The consent (or absence of opposition in France) of the person for their reuse and that of the associated data for future research*:



- After having been informed by the health establishment via the hospital welcome booklet, for example, or by that mentioned in the biological analysis results of medical biology laboratories.

- After having been informed by the initial research team: verbal or written information, an information document, etc.

2



The authorisation of ethics committees for their collection and use as well as the associated data. This authorisation is only necessary for samples collected for the research.

[1] Ethics unit and health division;
[2] Clinical coordination of the translational research centre and CEBOH unit.
* If the procedures for providing information are insufficient, the research ethics committee may need to be consulted.

INSTITUT PASTEUR

Before receiving them, I must ensure...

CONTRACTS

Signing of a contract (MTA/collaboration contract, etc.) between the Institut Pasteur (its legal department) and the organisation housing the samples.

Contact : *dj-sante@pasteur.fr*

GDPR

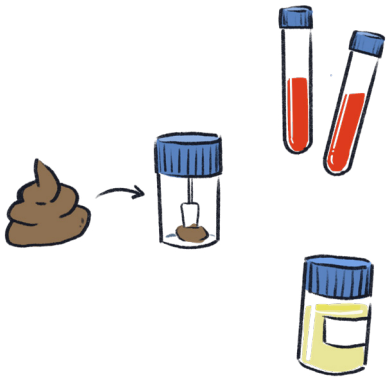
Implementation of procedures to ensure the protection and confidentiality of associated health data, for example, compliance with the General Data Protection Regulation (GDPR).

Contact : *health-data@pasteur.fr*

IMPORT

Declaration of samples and import authorisation (samples from abroad) to the Ministry of Higher Education, Research and Innovation (MESRI).

Contact : *ceboh@pasteur.fr*



FEEDBACK

Individual feedback to the participant if an impact on his/her health is revealed.

Contact : *ethics@pasteur.fr*

IRB

For samples received from abroad, the project must be submitted to a research ethics committee of the source country, and if the Institut Pasteur is responsible for the research, it must be submitted to the IRB.

Contact : *irb@pasteur.fr*

I can find help to conduct and follow these procedures through the clinical core of the center for translational research (CRT-ClinCore@pasteur.fr)

Important points

Samples must be used within the context of the stated project.

Any new use in the laboratory and/or sharing within the scientific community requires the following:

- Verification of the information given to participants,
- Signing of a new contract.

This also applies to sharing samples with other Institut Pasteur teams.

In the event of non-compliance with the regulations, the samples may be destroyed and financial (loss of funding) or legal sanctions may incur.

Illustrations INK-LINK, AdobeStock

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