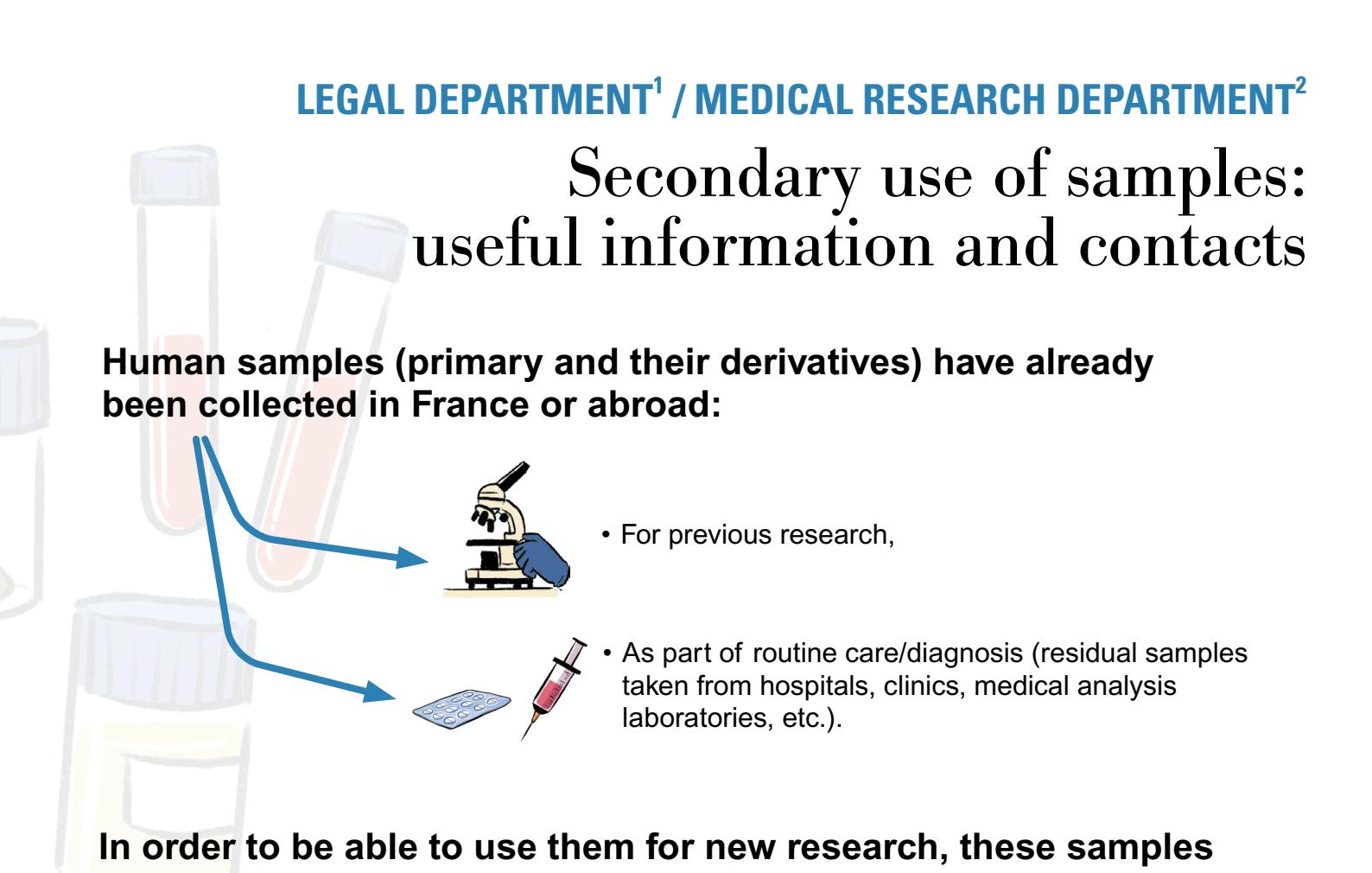


# Secondary use of samples: useful information and contacts

**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 first 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.

<sup>1</sup> Ethics unit and health division;

<sup>2</sup> 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.

# 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](mailto:dj-sante@pasteur.fr)

## GDPR

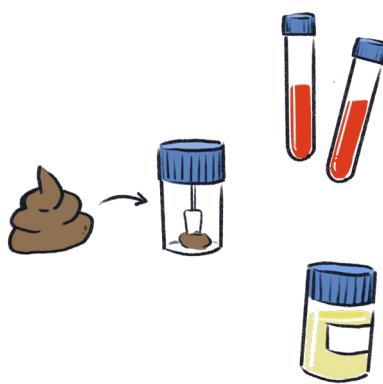
Implementation of procedures to ensure the protection and confidentiality of associated health data, for example, compliance with the General Data Protection Regulation (GDPR).

Contact : [health-data@pasteur.fr](mailto: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](mailto:ceboh@pasteur.fr)



## FEEDBACK

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

Contact : [ethics@pasteur.fr](mailto: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](mailto: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.