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### A-Purpose and scope of application

This document describes the organization defined for making a collection or biological samples available to CeRBM.

The provision of biological resources is subject to different formalities depending on the case:

- ✚ recession: CeRBM returns to the investigator or his representative the biological resources he had entrusted to CeRBM for preparation and storage, during the reservation period. The coordinating investigator - or a person chosen by him or her - fills in the request form, specifying the sample number, type and number required.

The CeRBM proceeds to the destocking. Please note that any removal is final.

- ✚ transfer: CeRBM transfers the biological resource to an outside party. This transfer is only possible, during the reservation period, if the investigator agrees and is subject to the approval of the CeRBM Coordinator or a delegated person.

On the other hand, transfer requires authorization from the Ministry. For CeRBM, this request is made by the Regulatory Referent (DRCI), who will apply to the relevant authorities.

### B-Reference documents

Standard NF EN ISO 20387: General requirements for biobanking

Iso 9001: Quality management system - Requirements

### C-Responsibilities

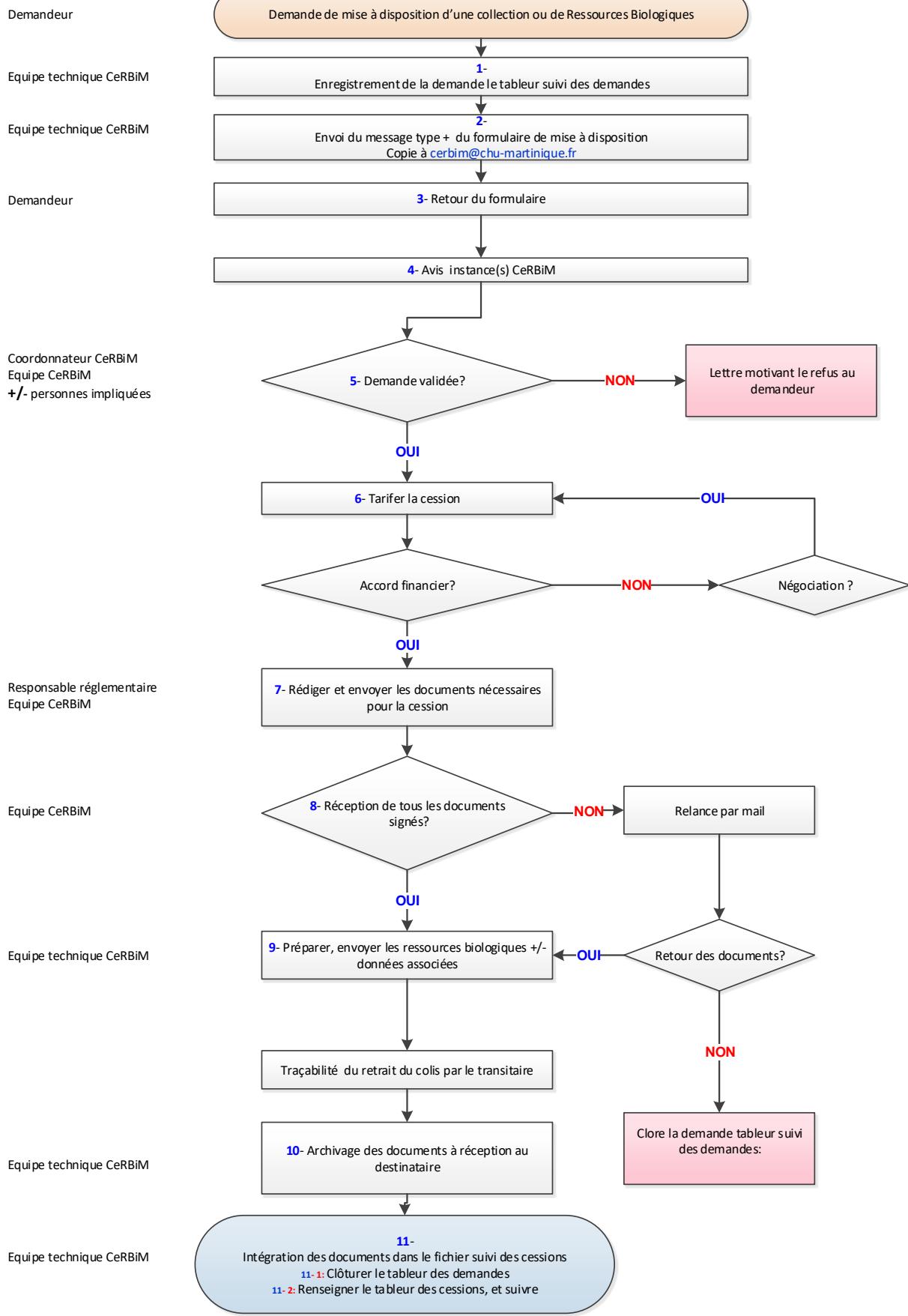
CeRBM coordinator is responsible for applying this procedure.

The quality manager ensures that the document is kept up to date.

### D-Procedure description

Before being handed over to the transporter or applicant, biological resources are stored in a conservation area (an area restricted to authorized personnel).

#### D-1 Flow chart





**D-2** Standard response message to the requestor and CeRBiM@chu-martinique.fr :

*We acknowledge receipt of your request under number xx-AAAA  
Enclosed you'll find the form you need to fill in and return to us, so that we can check that all the requirements apply.*

**(Please attach the availability form)**

*For Rémi NEVIERE  
Medical and scientific coordinator of CeRBiM.*

**E-Related documents**

- see related documents