General Terms
CRYOSTEM Collection Access
2021

These General Terms (hereafter the « **General Terms** ») aim in particular to present the terms and conditions governing the submission, selection and processing of academic or industrial research projects (hereafter the **« Project** ») submitted to CRYOSTEM Scientific Committee as part of the Access to the CRYOSTEM Collection (herafter the« **Access to the Collection** ») .

INTRODUCTION :

CRYOSTEM is a project that is backed by the SFGM-TC (Francophone Society for Bone Marrow Transplantations and Cell Therapy (hereafter **« SFGM-TC** ») and accepted by the French National Research Agency in relation to the "Cohorts" call for projects funded under the French government's "National Investment Programme" ("*Investissements d'avenir*") in June 2010.

Currently, CRYOSTEM is a network covering all the allo-transplantation centres in France and 28 related Biological Resource Centres (BRC).

Since 2012, CRYOSTEM has set up a collection of biological samples taken by recipients, and their donors, before and after allogeneic hematopoietic stem cell transplantation (HSCT) (hereafter the **« Collection** »), thus reaching its first objective.

Starting in April 2015, this Collection has been made available to scientists to enable high-level scientific projects to be performed focussing on the theme of Graft-versus-Host Disease (GvHD) by transplanted patients (hereafter **« the Thematic** »). The objective is to better understand the physiopathology of this little-known disease in humans, the predictive factors related to its occurrence, the response to immunosuppressive treatments and the long-term prognosis.

In 2016, CRYOSTEM obtained the authorization of the French Patient Protection Committee (CPP) to extend the collection thematic to the all HSCT complications additionally to the GvHD, opening the fields of studies.

In relation to this availability, CRYOSTEM wishes to choose the most relevant scientific projects complying with the chosen scientific Thematic.

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PART I – GENERAL TERMS

I.1 – Purpose of the General Terms

The purpose of these General Terms is the definition of the terms related to the Access to the Collection and related to the selection of academic and/or industrial members as defined in article I.2 below (hereafter the **« Applicants** ») in order to grant them with the Collection implemented by CRYOSTEM, as part of research on the scientific themes identified by CRYOSTEM, the « HSCT complications ».

The biological samples treated from blood samples and the derived aliquots, constituting the CRYOSTEM Collection, are named hereafter « **Biological Material** ».

The « **Data »** (**or « Associated Data** ») are defined hereafter as the whole data associated to the Biological Material including

- the biological data associated to the Biological Material
- and the clinical data associated to the patient, extracted from the EBMT (European Society for Blood and Marrow Transplantation) registry, collected and provided by the SFGM-TC and transmitted by CRYOSTEM. In case of these data could not be extracted from the EBMT registry, they could be obtained through requests addressed to the clinical units.

The Biological Material and the associated Data are named hereafter the « **Biological Resources** ».

The projects put forward by the Applicants as part of the Access to the Collection must abide by these General Terms and their Appendices.

The Applicants acknowledge that they have read and accepted these General Terms and the Appendices, also included in the application file, by signing the act of commitment which will be provided if their project is selected.

Regarding the Access to the Collection, the CRYOSTEM « **Scientific Committee** » has defined the procedure and the terms related to its proceedings and is responsible for organizing the selection process and follow-up of the Projects. Its members are listed in Appendix 1.

I.2 – Conditions for submitting replies to Access to the Collection

The Applicants concerned by the Access to the Collection are the following :

- Academic laboratories such as those registered by the Ministry for Higher Education and Research or under the supervision of this same Ministry, i.e. any laboratory of the following structures: healthcare centres, higher education and research establishments, public scientific establishments, public industrial and commercial establishments, public scientific and technological establishments, foundations, public interest groups, research organizations, regional delegations for research and technology (hereafter "**Academic Laboratories**"). The term "Academic Laboratories" in these General Terms includes consortiums and groups of laboratories of these same structures ;

- Any company of the private sector such as pharmaceutical laboratories and biotechnologies companies defined hereafter as « **Industrials** » ;
- Consortiums grouping together Academic Laboratories / Industrials (hereafter « **Consortiums** »).

The Applicants may comprise separate legal entities ; in this case, a key contact for CRYOSTEM should be clearly identified by each Applicant (hereafter « **Project Initiator** »).

For projects of equal scientific value, projects presented by research teams belonging to a French Academic Laboratory will be preferred.

The Academic Laboratories taking part in implementing the Collection are authorized to apply to access to the Collection.

The appraisal and selection criteria of the Projects put forward by the Applicants will be submitted to international experts, thereby guaranteeing the maximal independence in the final choice of the Projects accepted.

PARTIE II – TIME SCALE OF THE SELECTION PROCESS TO ACCESS TO THE COLLECTION

II.1 – Schedule

Projects submission to access to CRYOSTEM Collection could be done at any moment of the year.

The application file and General Terms are available on demand (emilie.robert@cryostem.org) and can be downloaded from the CRYOSTEM website (www.cryostem.org).

Application files are studied during the Scientific Committee session following the Project submission. Four sessions of the Scientific Committee at least are organised during the year.

II.2 – Application submissions

Applications will be sent by e-mail to CRYOSTEM strategy & valorisation manager (emilie.robert@cryostem.org).

For further information, please contact him by phone (+334 91 22 34 37) or by e-mail.

CRYOSTEM undertakes to take all appropriate measures in order to ensure the strictest confidentiality regarding the applications submitted. Applicants are however advised to perform all necessary procedures and take appropriate measures to protect the information that they submit as part of an application to access to the Collection.

Applicants should notify any possible conflicting interests and provide a list of the experts that they wish to challenge. CRYOSTEM undertakes to take this request into consideration where possible in order to guarantee the greatest possible fairness in the selection process.

The Applicant will give information in the application file on the funding status (acquired or not, ongoing application) and on the source. The Applicant will precise if the funds are

intended to compensate only the provision of the Biological Resources or the total achievement of the Project.

In the case of the Applicant would not be able to fund the provision of the Biological Resources or the Project in its totality, CRYOSTEM Scientific Committee could offer to the Applicant to include his Project into the « HTC Project » program. The HTC Project is an endowment fund, created by CRYOSTEM, which supports and funds an international program to promote the research in the field of HSCT complications. The HTC Project could bring an entire or partial financial support, according to the total cost of the Project and the available funds on the HTC Project endowment funds, following validation by its Governance boards.

The financial evaluation related to the Biological Resources provision and included in the application file, is provided for information purposes only. Only the samples allocation quotation published following the final decision of the Scientific Committee and taking into account the Biological Resources availabilities according to the demand, is financially valuable. It will be transmitted to the Applicant and must be sent back signed to the strategy & valorisation manager for acceptance.

The Applicants commit in declaring in the application file all the published information and the potential rights of intellectual property they have, related to the Project they submit to CRYOSTEM to access to the collection.

The Applicant must precise in the application file the expected duration of the Project.

II.3 - Description of the selection stages

First stage : short-listing of Projects

This first stage aims to select the Projects that meet with the specific purposes set forth in the application.

With reference to the application, the following shall not be accepted:

- Incomplete applications ;
- Applications written in French ;
- Applications not signed by the person authorized to represent the Applicant.

At the end of this first stage, an unlimited number of Projects will be short-listed by the Scientific Committee and examined in full as part of the second stage.

Applicants will be informed in writing of the result of this short-listing. Applicants could be invited to answer to remarks of CRYOSTEM Scientific Committee or to submit a revised proposal of the Project based on the recommended modifications.

No financial aid will be allocated to the Applicant on the basis of this first stage alone.

Second stage : review by the international experts

Following the short-listing, each Project will be sent to two (2) or three (3) international experts, designated by the Scientific Committee, and in charge notably of the evaluation of the methodological and scientific quality of the Project.

The synthesis of the reviews of the requested experts will be transmitted to CRYOSTEM Scientific Committee.

No financial aid will be allocated to the Applicant at the end of this second stage.

Third stage : final selection of Projects

The final selection will be done by the Scientific Committee on the basis of the international experts reviews and will be notified to the selected Applicants via an official letter by the Steering Committee. This notification will precise the logos to be included in any oral or written communication related to the use of the Biological Resources of the Collection (CRYOSTEM, SFGM-TC and the potential funders).

II.4 –Selection criteria of Projects

Appraisal of the applications submitted by the Applicants will take place on the basis of qualitative and quantitative criteria defined by CRYOSTEM Scientific Committee. These criteria may be provided for information purposes and on request from the Applicant to CRYOSTEM. The originality of the Project, the implemented methodology, the feasibility in view of the biological resources asked for, the impact of the Project in the field and the financial aspects will be evaluated by CRYOSTEM Scientific Committee members and the requested international experts.

PARTIE III – CONDITIONS OF USE OF THE BIOLOGICAL RESOURCES MADE AVAILABLE TO THE SELECTED APPLICANTS

III.1 – Conditions governing access to the Biological Resources

The Collection includes three types of Biological Material: viable cells in DMSO, dried cell pellets and plasma.

The clinical data extracted from the EBMT (European Society for Blood and Marrow Transplantation) registry, provided by the SFGM-TC, transmitted by CRYOSTEM, required for the Project to progress smoothly will be made available with the Biological Material in an anonymous manner. The accuracy of the clinical data extracted from the EBMT registry do not come under the responsibility of CRYOSTEM.

Only the clinical data asked for in the application file would be provided to the selected Applicant.

In the case of requirement of additional data for the Project, the Applicant will, on the one hand, precise them in the application file, and, on the other hand, ask for to CRYOSTEM strategy & valorisation manager before, to validate the feasibility and quantify the workload : an additional quotation will be provided for the obtain of these additional data. In the case of a later demand of clinical data, the selected Applicant would be asked for additional financing

equal to the cost related to data extraction specified in the pricing-list related to the Access to the Collection.

The Biological Resources provided to the selected Applicants will be selected by CRYOSTEM in accordance with relevance criteria in relation to the Project and will be defined during one-on-one interviews led by the strategy and valorisation manager with the selected Applicants. The process for selecting the Biological Resources may be provided, for information purposes, to the selected Applicant upon submission to CRYOSTEM of a written request to this effect.

The access to CRYOSTEM Biological Resources needs to be subject to contract (convention, agreement) signed by the representatives of the selected Applicant affiliation and those of CRYOSTEM. This document will precise the specific terms of the collaboration between the research team of the selected Applicant and CRYOSTEM.

III.1.1 Provision of the Biological Resources

CRYOSTEM will inform the Applicant of the availability and the planned date of availability of the Biological Resources.

In the case of a provision to a selected Applicant outside the French territory, a ministerial export authorization will be asked by CRYOSTEM. The provision will be done only at reception of the authorization ; the corresponding number provided by the French Ministry will be communicated to the Applicant.

The delivery of the Biological Material will be organized by CRYOSTEM, which reserves the right to choose the means of transport. The carriage costs of the Biological Material will be borne by the Applicant and billed at their actual cost. The applicable carriage costs are specified in the pricing-list related to the Access to the Collection and are dependent on the addressee country of the Biological Material and set up by the transporter chosen by CRYOSTEM.

Once the Biological Material is received, the selected Applicant is responsible for inspecting its condition and setting forth any reservation or protest regarding its physical state, within a period of eight (8) days as of the date of receipt of the Biological Material.

The selected Applicant commits to returning to the strategy & valorisation manager the provided Biological Material follow-up sheet, after any use. This document will be provided simultaneously to the Biological Material shipment and will enable to report the characteristics of the samples obtained after thawing. This information, related to the samples quality, could be shared with CRYOSTEM network members.

III.1.2 Right to use the Biological Resources

The Biological Resources may be used only for research purposes specified in the submitted Project, on the one hand, respecting the terms related to the Access to the Collection, and on the other hand, respecting individual rights and the legal and ethical principles governing the status of the human body, its parts and its products, without any right to transfer them.

These terms are applied to the clinical data associated to the Biological Material and provided by CRYOSTEM via the SFGM-TC.

III.1.3 Suspension – Return of the Biological Resources

Any use of the Biological Resources other than that defined and authorized by CRYOSTEM as part of the Access to the Collection and of the collaboration agreement is strictly prohibited.

In any case, the selected Applicant commits :

- to respecting the French regulations in force regarding the use of Biological Resources as part of research involving the human person,
- to using them only for the Project selected by CRYOSTEM,
- not to giving them up to a third party,
- to informing CRYOSTEM Scientific Committee of the results obtained as part of the research (publications, patent application, etc.),
- to informing CRYOSTEM Scientific Committee of the potential new biological data resulting from the use of the Biological Resources provided and that may enrich the Biological Resources retained in the CRYOSTEM Collection.

Failure by a selected Applicant to abide by the conditions of use defined by CRYOSTEM upon selecting the Applicant and in the present General Terms may lead to the suspension of availability of the Biological Resources, and the return to CRYOSTEM of the Biological Resources already provided, at the Applicant's cost.

III.1.4 Financial Conditions

The Biological Resources made available to the selected Applicant will be accessible in accordance with the financial terms defined in Appendix 2.

These prices include the access to the Biological Material and associated clinical data, the interventions of CRYOSTEM staff (project managers, transplant units and Biological Resources Centres staff) for the selection, the removal and the provision of the Biological Resources.

In the case of the selected Applicant entirely funds on its own source the provision of the Biological Resources, a 50% deposit will be asked for to the selected Applicant before the shipment.

The availability of the Biological Resources is part of an operation that is not subject to value-added tax in accordance with the terms of article 256 of the French Tax Code and the Official Bulletin of the French Tax Authorities BOI-IS-CHAMP-10-50-10-20.

III.1.5 Duration of the Project of the selected Applicant

The Biological Resources provision is performed for a determined duration corresponding to the duration of the Project of the selected Applicant. The expected duration of the Project has to be notified by the Project Initiator in the application file related to the Access to the Collection.

The end of the Project takes place once :

- the statistical analyses are achieved ;
- the final report is transmitted ;
- and the interactions of the selected Applicant with the Scientific Committee, and also with the editorial policies on the Project communication, are achieved.

The end of the Project will be notified by an official letter of CRYOSTEM Scientific Committee.

III.2 – Obligations

III.2.1 CRYOSTEM's obligations

CRYOSTEM is abound by an ISO 9001 : 2015 certification process including the surveillance of the BRCs for the management of the Collection in accordance with the requirements of standard NF S 96-900.

CRYOSTEM guarantees, to the exclusion of any other guarantee, that the Biological Material included in the Collection has therefore been prepared in accordance with harmonized procedures and protocols defined by CRYOSTEM network. These procedures and protocols are applied by all members of CRYOSTEM network in accordance with good laboratory methods. The samples treatment protocols could be transmitted to the selected Applicant on request addressed to the strategy & valorisation manager.

CRYOSTEM regularly inspects that these methods are respected and performs random quality tests on the Collection at least once per year, the results of which may be provided to the selected Applicant, upon submission to CRYOSTEM of a written request to this effect.

CRYOSTEM declares that the Biological Material made available to the selected Applicants will be coded and won't contain any direct or indirect identifying information of the people who have expressly and freely consented to take part to research led on the Thematic. Data associated to the Biological Material are anonymised.

CRYOSTEM commits to executing the provision of the Biological Material to the selected Applicant needed for his Project as soon as possible.

Some Biological Material, by their nature, may not present all of the characteristics required, which is specifically acknowledged by the selected Applicant for his Project. CRYOSTEM does not grant any guarantee, express or implied, as for the viral safety and the absence of contamination of the Biological Material provided to a selected Applicant and as for the compliance for a specific use. The selected Applicant undertakes not to claim that CRYOSTEM replaces the Biological Material which does not meet its expectations.

In no instance may CRYOSTEM, its directors and its employees, not be held liable for any direct or indirect damage to the Biological Material related to its use, in the selected Applicant's premises, or for their mishandling by the selected Applicant, as the latter bears exclusive responsibility for Biological Material storage.

In addition, CRYOSTEM commits to providing to the selected Applicant a protocol of thawing when viable cells in DMSO are provided.

III.2.2 Applicant's obligations

Before the provision of the Biological Resources, the selected Applicant must have taken the required steps regarding the regulatory and legal framework for his Project and bring the evidence to CRYOSTEM as soon as possible.

During all the duration of the Project, the selected Applicant commits to :

- returning the « samples follow-up form » after the use of the samples to the strategy & valorisation manager ;
- informing CRYOSTEM regularly concerning the progress of the Project ;
- and providing a progress report, at the latest eighteen (18) months after the last provision of the Biological Resources by CRYOSTEM, along with a final report at the end of the Project.

However, the selected Applicant might be asked for intermediary reports on demand of CRYOSTEM Scientific Committee or the potential sponsors or for communication actions.

The selected Applicant undertakes to inform CRYOSTEM Scientific Committee of the end of the Project within three (3) months following the end of the Project.

The selected Applicant undertakes to inform CRYOSTEM strategy & valorisation manager of any relevant information related to the Biological Material quality in link with their use in a reasonable delay following the provision.

The selected Applicant undertakes to destroy the Biological Material and the related clinical data extracted from the EBMT registry and/or any additional data within a period of thirty-six (36) months from the date of notification of the end of the Project. A destruction certificate of the Biological Resources, provided by CRYOSTEM at the shipment of the Biological Material, must be sent to CRYOSTEM upon expiry of this period.

However, the selected Applicant benefiting from CRYOSTEM's Biological Resources may, within this period of thirty-six (36) months following the end of the Project, ask for authorization from CRYOSTEM to use the Biological Resources for another research project. This Project must be submitted beforehand to CRYOSTEM Scientific Committee. The Scientific Committee undertakes to examine the said Project involving the Biological Resources already provided and to provide its decision within a period of two (2) months. CRYOSTEM will communicate the Scientific Committee decision to the selected Applicant which undertakes to abide by it.

The selected Applicant is explicitly prohibited from performing research on the Biological Resources provided by CRYOSTEM outside of the scope of the Project for which CRYOSTEM has given its approval. This ban includes all clinical data provided by CRYOSTEM and the SFGM-TC.

The selected Applicant commits not to putting back together or to identifying in any case the subjects from whom the anonymised data would be transmitted by CRYOSTEM, whatever

the way used, notably by avoiding to match them, or trying to match them, with other sources of data or Biological Resources provided regarding the Project or that would be accessible for him.

The selected Applicant has taken note and takes in count that considering the experimental nature of the Biological Material, it has to be handled with care and caution. The selected Applicant will be entirely responsible for the use of the provided Biological Material and the associated Data.

The selected Applicant takes on alone the risks, damages and responsibilities related to the receipt, the handling, the storage and the use of the Biological Resources provided by CRYOSTEM.

III.3 – Intellectual property

III.3.1 Biological Material and associated Data

The provision of the Biological Material and the associated Data by CRYOSTEM to a selected Applicant would not be interpreted as a cession of rights of intellectual property by CRYOSTEM or by the BRCs on the Biological Resources.

The selected Applicant acknowledges that he has no right to submit a patent application relating as such to the Biological Material and / or Associated Data made available to him.

It is expressly agreed that the right to use the Biological Resources provided to the selected Applicant cannot, under any circumstances, be interpreted as expressly or implicitly conferring on the latter any right or title, or license over Biological Material and Associated Data.

III.3.2 Results from the research of the selected Applicant using the provided Biological Material and associated Data

The selected Applicant aims to directly promote the results of the research Project selected and authorized by CRYOSTEM (hereafter the « **Results** »).

All data, documents, results, scientific knowledge, which may or may not be protected by an industrial property title resulting from research work carried out within the framework of the Project by the selected Applicant, and resulting from the use of the Biological Resources provided by CRYOSTEM, are the property of the said Applicant.

The selected Applicant agrees to inform CRYOSTEM if he intends to submit a patent application on the Results, and to send it the date of publication of this application.

Supposing that, and only in this case, CRYOSTEM would come to provide an intellectual contribution within the framework of the realization of the research work carried out as part of the Project by the selected Applicant that the latter undertakes to request CRYOSTEM before the submission of a patent application to determine, together and in good faith, the conditions for sharing the results and recognizing the intellectual and inventive contribution of CRYOSTEM.

If the results appear reasonably patentable, and the contribution of CRYOSTEM is equivalent to that of the selected Applicant, the parties will submit a patent application in joint names, the ownership, rights and obligations of the parties being governed by the provisions of a co-ownership agreement, available on request from the strategy & valorisation manager.

Under these same conditions, for each new patentable invention, CRYOSTEM and the selected Applicant will sign a co-ownership agreement in accordance with the standard contract.

III.4 – Right to publication

The selected Applicant undertakes to refer to CRYOSTEM and the SFGM-TC in any publication in relation to the Project in accordance with the following conditions:

- By quoting « CRYOSTEM » Consortium in the list of authors subject to the editorial board acceptance and by listing in the appendix the institutions members of the Plenary Committee (complete list to be obtained from CRYOSTEM). In case of the CRYOSTEM Consortium could not be included in the list of authors, the chosen Applicant undertakes to include the CRYOSTEM Consortium in the « Acknowledgements » section ;
- By noting the origin of the Biological Resources in the « Materials and Methods » section in the following format « ***The samples annotated have been provided by the CRYOSTEM Consortium and SFGM-TC.*** » and,
- By quoting the Digital Object Identifier (*doi*) assigned to the CRYOSTEM Collection and provided by the strategy & valorisation manager.

The selected Applicant undertakes to include CRYOSTEM and SFGM-TC logos, provided to him, in any oral and written communication, and those of the potential funders.

The selected Applicant undertakes to submit the draft publication to the Scientific Committee at least one (1) month prior to publication, for its opinion, so that CRYOSTEM can check the accuracy of the citation of CRYOSTEM and of the data related to the Biological Resources provided by CRYOSTEM.

CRYOSTEM will inform of its decision within a maximum period of fifteen (15) days from when the Project is sent. Should no response be received within this period, CRYOSTEM's consent will be deemed to have been provided.

Consequently, CRYOSTEM may make any amendment or suspend the draft publication if any information related to the Biological Resources due for publication presents omissions or inaccuracies.

Publication fees will be the charge of the selected Applicant.

Patients of the CRYOSTEM cohort, partners of the CRYOSTEM network, potential funders, could be informed about some information coming from the selected Projects (title, name of the selected Applicant, corporate name of its representative or its establishment, logo or label, description, key words, Layman's summary, etc.) and the results, notably via the CRYOSTEM website (www.cryostem.org). These information will be submitted to the selected Applicant for validation before publication.

PARTIE IV – FINAL TERMS

IV.1 – Dispute settlement

The parties shall endeavour to settle their disputes on an out-of-court basis or via a mediator.

Otherwise, the parties will settle their dispute before the *Tribunal de Grande Instance* in Paris

IV.2 – Applicable law

These General Terms are governed by French law. In the case of their translation into a foreign language, only the French text will be binding..

IV.3- Acceptance of the General Terms

The submission of an application by an Applicant implies the latter's specific and unreserved acceptance of all of these General Terms.

- Qualified members in Stem Cell Transplantations and / or Hematology:

  - Dr Eolia Brissot, MCU-PH in Clinical Hematology and Cell Therapy, Saint Antoine hospital, Paris

  - Pr Frédéric Baron, PU-PH Clinical Hematology, head of the transplantation programm, Liège

  - Dr Maud D'Aveni-Piney, PH Clinical Hematology, Nancy

  - Dr Edouard Forcade, doctor in Hematology, Haut-Lévêque hospital, Bordeaux

  - Dr Etienne Daguindau, doctor in Hematology, assistant clinic director, CHU Jean Minjoz, Besançon

- The chairman of the SFGM-TC Scientific Committee or a representative

- Members of Steering Committee:

  - Pr Régis Peffault de Latour, project coordinator, PU-PH Saint-Louis hospital, Paris

  - Pr Jean-Hugues Dalle, PU-PH Paediatric Hematology, Robert Debré hospital, Paris

  - Dr Boris Calmels, BRC coordinator, PH in Cell Therapy Centre of « Institut Paoli-Calmettes » Marseille

  - Claire Fontenille, regulatory & operational manager

  - Juliette Canard, junior project manager

  - Dr Emilie Robert, strategy & valorization manager

Appendix 2: Pricing-list related to the Access to the CRYOSTEM Collection

Pricing list covering the provision of biological resources from the CRYOSTEM Collection - 2021

A : Academic project initiator and member of the CRYOSTEM consortium
B : Academic project initiator unaffiliated with the CRYOSTEM consortium
C : Non-academic / private / commercial project initiator

Samples provision	**A**	**B**	**C**
mononuclear cells in DMSO*	**14,60 €**	**18,25 €**	**51,10 €**
dry precipitate*	**11,00 €**	**13,75 €**	**38,50 €**
plasma*	**10,00 €**	**12,50 €**	**35,00 €**
selection, removal from storage and unpacking (flat rate)*[1]	**9,60 €**	**12,00 €**	**33,60 €**
blood sample (flat rate)**	**9,20 €**	**11,50 €**	**32,20 €**

** cost to be applied per aliquot*

*** cost to be applied per period and per patient*

Patients data provision	**A**	**B**	**C**
extraction of clinical data via the EBMT registry (flat rate)	**291,00 €**	**363,75 €**	**1 018,50 €**
extraction of additional data (i.e. blood count, enzymes…)[2]	**on quotation**		

Administrative management	**A**	**B**	**C**
national shipment (applicable rate provided as a guide for the shipment in dried ice of a cryobox containing from 1 to 100 aliquots on a national scale)[3]	**224,00 €**	**224,00 €**	**224,00 €**
international shipment	**on quotation**		
aliquots identification and selection (flat rate)	**185,00 €**	**231,25 €**	**647,50 €**
centralization and control on receipt before sending to the PI (flat rate)***[4]	**28,00 €**	**35,00 €**	**98,00 €**
biological resources export authorization request (flat rate)[5]	**74,00 €**	**92,50 €**	**259,00 €**

**** cost to be applied per group of 100 aliquots*

[1] corresponds to the aliquots removal from nitrogen tanks and to the database updating

[2] data availability would be checked before with the project managers

[3] to be applied per number of shippings given by the project managers in preparation for centralization and checking before shipment

[4] aliquots tranport from CRYOSTEM network BRCs to the CRYOSTEM platform for checking before provision

[5] rate applicable per project for project initiators out of France