

Regulatory Survey ION-4201

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General	
The information has been reviewed in year :	
Organisation name:	Marrow Donor Program Belgium
Organisation ION:	ION-4201
Country:	Belgium
Year the registry started operations:	1989

Products		comment
Do you provide HPC, Marrow?	yes	
Do you provide HPC, Apheresis?	yes	
Do you provide HPC, Cord Blood?	yes	
Do you provide MNC, Apheresis?	yes	
Do you provide NC, Whole Blood?	no	

Product quantity		
Data valid for year	2023	
Number of products	National	International
HPC, Marrow products:	0	1
HPC, Apheresis:	4	32
HPC, Cord:	0	13

License	
Organisation is licensed/accredited by the Competent Authority:	yes
Comment:	
Name of Competent Authority:	FAGG-AFMPS
Date of last inspection:	2018
Link to website of Competent Authority:	http://www.fagg-afmps.be/en
Can you provide the Legal documentation from the Competent Authority that your organisation is allowed to operate as a registry ?	yes
Is your registry WMDA Certified, WMDA Qualified or WMDA Accredited?	WMDA Accredited
The registry is accredited by any other organisation:	no
If yes, by which organisation?	

Affiliation	
The registry works with the following number of affiliated donor centres:	Six (6) donor centres (33 locations)
The registry audits its donor centres:	yes
The registry works with the following number of affiliated collection centres:	Eleven (11) collection centres
The registry audits its collection centres:	yes

The registry works with the following number of affiliated Cord Blood Banks:	Five (5) cord blood banks
The registry audits its Cord Blood Banks:	yes
The registry works with the following number of affiliated transplant centres:	Eleven (11) transplant centres
The registry audits its transplant centres:	yes
The registry works with the following number of affiliated IDM Testing Laboratories:	Thirteen (13) laboratories
The registry audits its IDM Testing Laboratories:	yes
The registry works with the following number of affiliated HLA/other DNA markers testing laboratories:	Eight (8) laboratories
The registry audits its HLA/other DNA markers testing laboratories:	yes
The registry would be able to provide a full list of name/addresses of each affiliated and their licence/accreditation status, on request:	yes
The Cord Blood Banks are FACT-NetCord accredited:	yes
If yes, the following Cord Blood Bank(s) are accredited:	Cliniques Universitaires Saint Luc - CBB; CBB UZ Leuven; Institut Jules Bordet - CBB; Liege CBB; UZ Gent CBB
The registry is able to provide a copy of all the certificates:	yes
Affiliated centres comply with WMDA Standards and applicable national regulations:	yes
The registry has requirements for affiliated centres in addition to WMDA Standards and applicable national regulations:	no
If yes, what are these requirements?	

Donor policy	
All donors are unpaid volunteers:	yes
All donors are informed about donation process and associated risks:	yes
Donors sign a valid informed consent to donate in the presence of a medical doctor/health care personnel/registry staff:	yes
The registry has systems in place to protect and control access to donor/patient records:	yes
The registry maintains donor anonymity:	yes
The registry has detailed donor evaluation and exclusion criteria in place:	yes
The registry has donor evaluation and exclusion criteria that do meet or exceed the WMDA guidelines:	yes

IDM

IDM	Tested	Method	Days between test and sampling/workup
ALT/ASTALT/AST ratio, De-Ritis-Quotient	-	Differs by center, see document MDPB LST012	30
ChagasChagas, T. cruzi	On request		30
CMV IgGCytomegalovirus (CMV) Antibody testing IgG	-	Differs by center, see document MDPB LST012	30
CMV IgMCytomegalovirus (CMV) Antibody testing IgM	-	Differs by center, see document MDPB LST012	30
CMV TotalCytomegalovirus Total	-	Differs by center, see document MDPB LST012	30
EBV IgGEpstein-Barr Virus Antibody testing IgG	-	Differs by center, see document MDPB LST012	30
EBV IgMEpstein-Barr Virus Antibody testing IgM	-	Differs by center, see document MDPB LST012	30
HAV (NAT)Anti-hepatitis A virus nucleic acid testing	-	Differs by center, see document MDPB LST012	30
HBV (NAT)Hepatitis B nucleic acid testing	-	Differs by center, see document MDPB LST012	30
HBc AbHepatitis B core antibody testing	Yes		30
HBs AgHepatitis B Surface antigen testing	Yes		30

HCV (NAT)Hepatitis C nucleic acid testing	-	Differs by center, see document MDPB LST012	30
HCV AbHepatitis C antibody testing	Yes		30
HEV (NAT)Hepatitis E Virus nucleic acid testing	-	Differs by center, see document MDPB LST012	30
HIV (NAT)Human Immunodeficiency Virus nucleic acid testing	-	Differs by center, see document MDPB LST012	30
HIV-1 AbHuman Immunodeficiency Virus HIV-1 antibody testing	Yes		30
HIV-2 AbHuman Immunodeficiency Virus HIV-2 antibody testing	Yes		30
HIV p24Human Immunodeficiency Virus p24 antigen testing	-	Differs by center, see document MDPB LST012	30
HTLV-IGHuman T-Lymphotropic Virus type I testing	-	Differs by center, see document MDPB LST012	30
HTLV-IIHuman T-Lymphotropic Virus type II testing	-	Differs by center, see document MDPB LST012	30
MalariaMalaria	On request		30
HSVHerpes Simplex Virus	-	Differs by center, see document MDPB LST012	30
STSSerological tests for syphilis	Yes		30
STS FTA-ABSSerological test for syphilis	-	Differs by center, see document MDPB LST012	30
ToxoplasmosisToxoplasmosis	-	Differs by center, see document MDPB LST012	30
VZVVaricella Zoster Virus	-	Differs by center, see document MDPB LST012	30
WNV-NATWest Nile Virus nucleic acid testing	-	Differs by center, see document MDPB LST012	30
Other tests performed	-		

Testing

The physical and medical exam at donor workup is performed by a medical doctor:	y es
All donor testing (at workup) for infectious disease is performed in a laboratory certified/licensed by a Competent Authority:	y es
HLA typing for patient specific request is performed in an appropriately accredited laboratory:	y es
Sterility testing is performed on the adult donor product:	y es
Sterility testing is performed on the cord blood product:	y es
Screening questionnaire to exclude communicable disease:	y es
Screening questionnaire to exclude donors with 'high risk' lifestyles:	y es
Donor reliability identified by a medical doctor:	y es
Donor clearance to donate is confirmed by a medical doctor, following as a minimum the donor exclusion criteria in Annex 1 of EU Directive 2006/17/EC : The party providing the cell product must exclude donors when: • They are pregnant; • They are breastfeeding; • There is the potential for transmission of inherited conditions; • There is evidence of any other risk factors for transmissible diseases on the basis of a risk assessment, taking into consideration <i>Donor travel and exposure history and local infectious disease prevalence</i>; • There is presence on the donor's body of physical signs implying a risk of transmissible disease(s); • There is a history of a disease of unknown aetiology; • There is a risk of transmission of diseases caused by prions; • There is systemic infection which is not controlled at the time of donation, including bacterial diseases, systemic viral, fungal or parasitic infections, or significant local infection in the tissues and cells to be donated; • There is history of chronic, systemic autoimmune disease that could have a detrimental effect on the quality of the Cell Product; • There is recent history of vaccination with a live attenuated virus where a risk of transmission is considered to exist; • There is ingestion of, or exposure to, a substance (such as cyanide, lead, mercury, gold) that may be transmitted to recipients in a dose that could endanger their health.	y es

Customs regulations

Are there any customs regulations to follow, or customs paperwork required, to import cell products into the organisation country? If yes, please specify:	
Are there any customs regulations to follow, or customs paperwork required, to export cell products from the organisation country? If yes, please specify:	Prior to shipment, notification to customs and security.
Are there any import regulations to follow, or paperwork required, to import cell products into the organisation country? If yes, please specify:	
Are there any export regulations to follow, or paperwork required, to export cell products from the organisation country? If yes, please specify:	

Reporting of Serious Adverse Events

Please indicate which of the following schemes for reporting Serious Adverse Events relating to either the Donor or the product the Registry participates in:	
- Mandatory National Reporting Scheme	
- Voluntary National Reporting Scheme	
- WMDA SEAR/SPEAR Reporting Scheme	yes
The registry will notify the receiving Registry within 24 hours of receiving information relating to any serious adverse event that could be considered to affect the patient receiving the product:	yes

Quality management

The registry does maintain Standard Operating Procedures for your work:	yes
The registry would be willing to provide these to WMDA or another registry upon request:	yes
The registry would be willing to provide WMDA or another registry, on request, with copies of any packaging the HPC product will arrive in:	yes
Donor records relating to the medical exam and HPC collection process are the following number(s) of year(s) retained:	min 30 years - max 50 years