

# Operational Information ION-6939

This data is publicly available. Parent page : ION-6939

Operational data for ZKRD - ION-6939	
Organisation Overview	
The information has been reviewed in year :	
Issuing organisation Number (ION)	ION-6939 The Issuing Organisation Number of a organisation, this is globally unique number, as issued by the ICCBBA.
Time zone	Europe/Brussels (GMT+01:00) The timezone in which this organisation operates.
Business hours	08:00 AM to 05:00 PM The daily hours in which this organisation operates.
Work schedule	Monday - Friday The normal work week in which this organisation operates.
Organisation closures	For all organisation closures, please see the <a href="#">WMDA Calendar</a> .
Donor ID example	GRID: 6939 XYZ0 0012 3456 731   DE XYZ 1234567890 (previous ID format) ID to be expected on paperwork, samples, and products.
Preliminary Search	
Requires preliminary search request form	Yes If yes, form required can be found on the Documents Page.
Extended Typing	
Typing options available for request	High resolution HLA-A, -B, -C, -DRB1, -DRB3/4/5, -DQB1, -DPB1, -DQA1, -DPA1 Please note special requirements listed
Requires organisation specific typing request form	Yes If yes, form required can be found on the Documents Page.
Number of days donor is reserved for a patient after a request	14 days after providing results
Verification Typing	
Maximum blood volume allowed	50 ml
Requires organisation specific typing request form	Yes If yes, form required can be found on the Documents Page.
IDM testing performed at verification	Yes
Number of days donor is reserved for a patient after a request	60 days after shipment of sample. Up to 90 days after receiving CT-results upon request of the registry/TC.
Sibling Typing	
Registry is willing to arrange sibling typings	Yes
If yes, procedure to apply for sibling typings	email
Workup Request	
Product dosage limit	20 mL/kg (BM); 5 x 10 <sup>6</sup> CD34+ cells /kg (PBSC) Number of donor cells allowed based on recipient weight.
Requires patient to meet certain standards in order to proceed with collection	Yes Organisation may or may not allow donor collections for some patients.
Patient physician must report the following in order to proceed with collection	All patient and donor data reported on the Work-up Request and the prescription including 5 loci high resolution HLA Must provide additional information to organisation.
Requires organisation specific work up forms	Yes If yes, form(s) required can be found on the Documents Page.

Workup IDM completed 30 days prior to collection	Yes Donor IDM results must be performed within 30 days of collection date to be valid and allow the collection to proceed.
Medical Health Questionnaire example available	No If yes, the example can be found on the Documents Page.
<b>Post-Transplant</b>	
Subsequent donation policy	Possible, please see Chapter 6 of the German Standards( <a href="https://www.zkrd.de/wp-content/uploads/2020/11/German-Standards_V12_english.pdf">https://www.zkrd.de/wp-content/uploads/2020/11/German-Standards_V12_english.pdf</a> )
Anonymous contact allowed	Anonymous patient-donor contact is permitted after the transplantation
Direct contact allowed	Allowed at the earliest two years after transplantation or one year after a second donation, if patient and donor agree and sign a declaration of consent
Gift exchange allowed	Yes, but gift must not be of substantial or monetary value
Cord blood contact allowed	No