

TRANSPLANT ACTIVITY SURVEY until 2006

TRANSPLANT ACTIVITY SURVEY from to 2006

Country Name

Please fill the number of all transplanted patients for each indication and for each type of transplant

Transplant Year	2006	2005	2004	2003	2002	2001	2000	1999	1998	1997	1996	1995	1994	1993	1992	1991	1990	1989	1988	1987	1986	
Indication																						
AML																						
ALL																						
CML																						
MDS																						
Other leukemia																						
MPS																						
Lymphoid malignancy (including LBL)																						
Multiple Myeloma																						
Solid tumor																						
Aplatic anemia																						
Aquired Pure red cell anemia																						
PNH																						
Congenital bone marrow failure																						
Hemoglobinopathy																						
Other hematological disease																						
EBV related disorders																						
Hemophagocytic syndrome																						
Langerhans cell histiocytosis																						
Autoimmune disease																						
Inherited metabolic disease																						
Primary immune deficiencies																						
Others																						
Total																						

Classify the disease as follows:

- AML Define by WHO classification (BM blasts \geq 20%)
- MDS Define by WHO classification (BM blasts $<$ 20%) : eg. RA, RN, RT, RCMD, RARS, RAEB, MDS-U, Count MDS/MPD (eg. MDS/MPD unclassified, CMML, JMML) as MDS
- Other leukemia eg. ATL, CLL, myeloid NK precursor acute leukemia
- MPD eg. Polycythemia vera, essential thrombocythemia, myelofibrosis
- Congenital bone marrow failure eg. Fanconi anemia, Dyskeratosis Congenita, Diamond-Blackfan anemia, congenital dyserythropoetic anemia, severe congenital neutropenia, myelodysplasia (WHIM syndrome)
Shwachmann -Diamond Syndrome, congenital amegakaryocytic thrombocytopenia
- Hemoglobinopathy eg. beta-thalassemia, sickle cell disease
- EBV related disorders eg. CAEBV, hypersensitivity to mosquito bites
- Inherited metabolic disease eg. Mucopolysaccharidosis, Niemann-Pick dis., Gaucher dis., I-cell dis., Pompe dis., Krabbe dis., Metachromatic leukodystrophy, Adreno leukodystrophy, Osteopetrosis
- Primary immune deficiencies eg. SCID, Wiskott-Aldrich Syndrome, X-linked hyper IgM syndrome, chronic granulomatosis, Chediak-Higashi syndrome

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Donor		Source																					
Allogeneic	Family	HLA id sibling	BM																				
			PBSC																				
			CB																				
			Others**																				
		HLA non id*	BM																				
			PBSC																				
			CB																				
			Others**																				
		Twin	BM																				
			PBSC																				
			Others**																				
		Unrelated	BM																				
	PBSC																						
CB																							
Others**																							
Autologous	BM																						
	PBSC																						
	Others**																						

*any family member (matched or mismatched) other than HLA id sibling

**Others include BM+PB,BM+CB or other mixture etc.

Please send it back to the secretary's office of APBMT by FAX or EMAIL: Fax +81-52-719-1973 or +81-561-61-3180 E-mail: office@apbmt.org