

YEARLY TRANSPLANT ACTIVITY SURVEY OF 2007



Country Name

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

Please fill the number of transplanted patients for each indication and type of transplant in 2007.

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

Indication	allogeneic												autologous			Total					
	family									unrelated				Allo	Auto	Total					
	HLA - id sibling				non - id*				twin			BM	PB				Others**				
	BM	PB	CB	Others**	BM	PB	CB	Others**	BM	PB	Others**			BM	PB	Others**					
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 WHOclassification (BM blasts \geq 20%)
- MDS Define by WHOclassification (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

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