

EBMT Centre Identification Code (CIC):	Treatment Type HCTCCT GT IST Other
Hospital Unique Patient Number (UPN):	
Patient Number in EBMT Registry:	Treatment Date / (YYYY/MM/DD)

## BONE MARROW FAILURE SYNDROMES (BMF) incl. APLASTIC ANAEMIA (AA)

DISEASE			
Note: complete this form only if this diagnosis was the indication for the H Consult the manual for further information.  Date of diagnosis://(YYYY/MM/DD)  Classification:  Acquired:	CT/IST or if it was specifically requested.		
Aplastic anaemia (AA)  Moderate Severe Very Severe Unknown	Etiology:		
☐ Pure red cell aplasia (non-congenital PRCA)	Secondary to hepatitis		
☐ PNH presentation ☐ Haemolytic	Secondary to toxin/other drug		
Aplastic	☐ Idiopathic		
☐ Thrombotic ☐ Other; specify:	Other; specify:		
Pure white cell aplasia			
Amegakaryocytosis / Thrombocytopenia (non-congenital)			
Other acquired cytopenic syndrome; specify:			
Genetic*:			
Amegakaryocytosis / Thrombocytopenia (congenital)			
Fanconi anaemia			
FANCC	(PALB2) (RAD51C) (SLX4)		
Diamond-Blackfan anaemia (congenital PRCA)			
☐ Shwachman-Diamond syndrome			
Dyserythropoietic anaemia			
☐ Dyskeratosis congenita			
Congenital sideroblastic anaemia (CSA)			
Other congenital anaemia; specify:			

<sup>\*</sup>Please fill in the "Inborn Errors" indication diagnosis form in addition to the current form (optional)



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Chromosome analysis done before IST/HCT: (Describe results of the most recent complete analysis No  Yes: Output of analysis: Separate			
☐ Unknown			
If chromosome analysis was done:			
What were the results?			
☐ Normal			
☐ Abnormal: number of abnormalities present:	:		
☐ Failed			
Date of chromosome analysis: $\_\_\_I\_I$	(YYYY/MM/DD) □ Unknown		
For abnormal results, indicate below whether the	e abnormalities were absent, present or not evaluated.		
abn 3	☐ Absent ☐ Present ☐ Not evaluated ☐ Unknown		
del(13q)	☐ Absent ☐ Present ☐ Not evaluated ☐ Unknown		
Monosomy 7	☐ Absent ☐ Present ☐ Not evaluated ☐ Unknown		
Trisomy 8	☐ Absent ☐ Present ☐ Not evaluated ☐ Unknown		
Other; specify:	☐ Absent ☐ Present		
	OR		
Transcribe the complete karyotype:			
Chromosomal breakage test (for Fanconi only):			
	•		
☐ Negative			
Positive			
☐ Not done or failed ☐ Unknown			



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## **MOLECULAR MARKER ANALYSIS**

Molecular markers analysis done before IST/HCT:				
☐ No				
☐ Yes				
☐ Unknown				
Date of molecular marker analysis (	f applicable):	_// (YYYY//	<i>MM/DD)</i> ☐ Unknown	
Indicate below whether the markers we	re absent, present c	or not evaluated.		
ASXL1	Absent	Present	☐ Not evaluated	Unknown
BCOR	Absent	Present	☐ Not evaluated	Unknown
BCORL1	Absent	Present	☐ Not evaluated	Unknown
CBL	Absent	☐ Present	☐ Not evaluated	Unknown
CSMD1	Absent	☐ Present	☐ Not evaluated	Unknown
DNMT3A	Absent	☐ Present	☐ Not evaluated	Unknown
ETV6	Absent	Present	☐ Not evaluated	Unknown
EZH2	Absent	☐ Present	☐ Not evaluated	Unknown
FLT3	Absent	Present	☐ Not evaluated	Unknown
GNAS	Absent	Present	☐ Not evaluated	Unknown
IDH1	Absent	Present	☐ Not evaluated	Unknown
IDH2	Absent	Present	☐ Not evaluated	Unknown
JAK2	Absent	Present	☐ Not evaluated	Unknown
KRAS	Absent	☐ Present	☐ Not evaluated	Unknown
MPL	Absent	Present	☐ Not evaluated	Unknown
NPM1	Absent	☐ Present	☐ Not evaluated	Unknown
NRAS	Absent	Present	☐ Not evaluated	Unknown
PHF6	Absent	☐ Present	☐ Not evaluated	Unknown
PIGA	Absent	Present	☐ Not evaluated	Unknown
PPM1D	Absent	☐ Present	☐ Not evaluated	Unknown
PTPN11	Absent	Present	☐ Not evaluated	Unknown
RAD21	Absent	Present	☐ Not evaluated	Unknown
RUNX1	Absent	Present	☐ Not evaluated	Unknown
SETBP1	Absent	☐ Present	☐ Not evaluated	Unknown
SF3B1	Absent	Present	☐ Not evaluated	Unknown
SRSF2	Absent	☐ Present	☐ Not evaluated	Unknown
STAG2	Absent	Present	☐ Not evaluated	Unknown
TET2	Absent	☐ Present	☐ Not evaluated	Unknown
	Absent	Present	☐ Not evaluated	Unknown
TP53	TP53 mutation	· · —		
		☐ Multi hit ☐ Unknow		
U2AF1	Absent	☐ Present	 ☐ Not evaluated	Unknown
ZRSR2	Absent	☐ Present	☐ Not evaluated	Unknown
Other; specify:	Absent	☐ Present	_ <del>_</del>	
		_ <del>-</del>		



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Bone marrow assessments:			
Cellularity in the bone marrow aspirate	☐ Acellular ☐ Hypocellular ☐ Normocellular ☐ Hypercellular	☐ Focal cellularity ☐ Not evaluated ☐ Unknown	
Cellularity in the bone marrow trephine	☐ Acellular ☐ Hypocellular ☐ Normocellular ☐ Hypercellular	☐ Focal cellularity ☐ Not evaluated ☐ Unknown	
Fibrosis on bone marrow biopsy	☐ No ☐ Mild ☐ Moderate ☐ Severe	☐ Not evaluable ☐ Not evaluated ☐ Unknown	
CD34+ cell count percentage (%)	%	☐ Not evaluated	☐ Unknown
Blast count percentage (%)	If the precise blast count $\leq 5\%$ $\leq 5\%$	☐ Not evaluated t is not available, please indic	☐ Unknown ate whether it is:
extended dataset			
Haematological tests			
ate tests performed: / / (YY	YY/MM/DD)  Unknown		
laemoglobin (g/dL) /as haemoglobin transfused within 4 weeks	before assessment?   No		nown
Platelets (10 <sup>9</sup> cells/L)		☐ Not evaluated ☐ Unk	nown
Vere platelets transfused within 7 days before assessment? No Yes Unknown			
Neutrophils (10 <sup>9</sup> cells/L)		☐ Not evaluated ☐ Unk	nown
Reticulocytes (10 <sup>9</sup> cells/L)		☐ Not evaluated ☐ Unk	nown
Ferritin (ng/mL)		☐ Not evaluated ☐ Unk	nown



☐ No

PNH test done?

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## **PNH TESTS**

☐ Yes: Date of PNH test: \_ \_ / \_ / \_ (YYYY/MM/DD) ☐ Unknown

only applicable for Aplastic Anaemia and/or PNH at time of diagnosis

	IOWII		
☐ Clone	e present: Size of PNH clone	<b>/:</b> in percentage (%):	_
Grand RBC Both	tometry assessment done of ulocytes	on:	
□ No	cal manifestation of PNH:	ent given?	YYYY/MM/DD) 🔲 Unknown
	Drug	Start date (YYYY/MM/DD)	Treatment stopped/date (YYYY/MM/DD)
	☐ Eculizumab	// Unknown	☐ No ☐ Yes: /  ☐ Unknown ☐ Unknown
	☐ Ravalizumab	// Unknown	☐ No ☐ Yes:/ ☐ Unknown ☐ Unknown
	Pegcetacoplan	// Unknown	☐ No ☐ Yes:/ ☐ Unknown ☐ Unknown
	Other; specify*:	// Unknown	☐ No ☐ Yes:/ ☐ Unknown ☐ Unknown
'	*Please consult the LIST O		S/AGENTS AND REGIMENS on the

If there were more drugs given during one line of treatment add more copies of this page.

EBMT website for drugs/regimens names