

Patient Number in EBMT database: Treatr			
	nent Date /	/(YYYY/N	1M/DD)

MYELODYSPLASTIC SYNDROMES (MDS)

DISEASE
Note: complete this form only if this diagnosis was the indication for the the HCT/CT or if it was specifically request Consult the manual for further information.
Date of diagnosis:/ (YYYY/MM/DD)
MDS transformed into Acute Leukaemia and treatment was done for Acute Leukaemia? No (complete this form) Yes (complete Acute Leukaemia indication diagnosis form in addition to the current form) Classification at diagnosis (WHO 2016):
☐ MDS with single lineage dysplasia (MDS-SLD)
☐ MDS with ring sideroblasts (MDS-RS)
☐ MDS with isolated del(5q) chromosomal abnormality
☐ MDS with multilineage dysplasia (MDS-MLD)
☐ MDS-RS with single lineage dysplasia (MDS-RS-SLD)
☐ MDS-RS with multilineage dysplasia (MDS-RS-MLD)
MDS with excess blasts (EB)-1
MDS with excess blasts (EB)-2
Refractory cytopenia of childhood
☐ MDS unclassifiable (MDS-U)
Therapy-related MDS: (Secondary origin) No
Voc. discoso related to prior exposure to therepoutic drugs or rediction

Yes, disease related to prior exposure to therapeutic drugs or radiation

☐ Unknown



EBMT Centre Identification Code (CIC):	Treatment Type	□ нст □ ст	☐ IST	☐ Other
Hospital Unique Patient Number (UPN):				
Patient Number in EBMT database:	Treatment Date _	//YY	YY/MM/DE))

CHROMOSOME ANALYSIS	Cŀ	IRON	10S0	ME A	ANAI	LYSIS
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Chromosome analysis done before treatment (all methods including FISH): (Describe results of the most recent complete analysis)						
☐ Not done or failed						
Yes, abnormal results: number of abnormalities present:						
Yes, normal results						
☐ Unknown						
Date of chromosome analysis (if tested):/_/(YYYY/MM/DD)						
Indicate below whether the abnormalities	were absent, present or	not evaluated.				
del(Y)	☐ Absent	☐ Present	☐ Not evaluated			
del(5q)	☐ Absent	☐ Present	☐ Not evaluated			
Other abn(5q); specify	☐ Absent	☐ Present				
del(20q)	☐ Absent	☐ Present	☐ Not evaluated			
del(7q)	☐ Absent	☐ Present	☐ Not evaluated			
Other abn(7q); specify	☐ Absent	☐ Present				
inv(3)	☐ Absent	☐ Present	☐ Not evaluated			
t(3q;3q)	☐ Absent	☐ Present	☐ Not evaluated			
del(3q)	☐ Absent	☐ Present	☐ Not evaluated			
Other abn(3q); specify	☐ Absent	☐ Present				
del(11q)	☐ Absent	☐ Present	☐ Not evaluated			
Trisomy 8	☐ Absent	☐ Present	☐ Not evaluated			
Trisomy 19	☐ Absent	☐ Present	☐ Not evaluated			
i(17q)	☐ Absent	☐ Present	☐ Not evaluated			
Other; specify	☐ Absent	☐ Present				
	Guilori, opconi)					
OR						
Transcribe the complete karyotype:						

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Hospital Unique Patient Number (UPN):				
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MOLECULAR N	MARKER	ΔΝΔΙ ΥSIS
WOLLCOLAIN	VI/711111	AIVALI 313

Molecular markers analysis done before treatment:					
□ No					
Yes					
☐ Unknown					
Indicate below whether the markers we	re absent, present or not eva	aluated.			
ASXL1	☐ Absent	☐ Present	☐ Not evaluated		
RUNX1	☐ Absent	☐ Present	☐ Not evaluated		
EZH2	Absent	☐ Present	☐ Not evaluated		
TP53	Absent	☐ Present	☐ Not evaluated		
UBA1	☐ Absent	Present	☐ Not evaluated		
SF3B1	☐ Absent	☐ Present	☐ Not evaluated		
TET2	Absent	Present	☐ Not evaluated		
NRAS	☐ Absent	☐ Present	☐ Not evaluated		
JAK2	☐ Absent	☐ Present	□ Not evaluated		
ETV6	☐ Absent	☐ Present			
CBL	☐ Absent	☐ Present			
IDH1	☐ Absent	☐ Present			
IDH2	☐ Absent	☐ Present			
NPM1	☐ Absent	☐ Present			
KRAS	☐ Absent	☐ Present	☐ Not evaluated		
PTPN11	☐ Absent	☐ Present	☐ Not evaluated		
PTEN	☐ Absent	☐ Present	☐ Not evaluated		
DDX41	☐ Absent	☐ Present	☐ Not evaluated		
SRSF2	☐ Absent	☐ Present	□ Not evaluated		
Other; specify	☐ Absent	☐ Present			

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