

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

## **MYELODYSPLASTIC NEOPLASMS (MDS)**

DISEASE
DISEASE
Note: complete this form only if this diagnosis was the indication for the the HCT/CT or if it was specifically requested. Consult the manual for further information.
Date of diagnosis: / / (YYYY/MM/DD)
MDS transformed into Acute Leukaemia and treatment was done for Acute Leukaemia?
<ul> <li>No (complete this form)</li> <li>Yes (complete Acute Leukaemia indication diagnosis form in addition to the current form)</li> </ul>
Classification at diagnosis (WHO 2022):
MDS with defining genetic abnormalities:
☐ MDS with low blasts and isolated 5 q deletion (MDS-5q)
☐ MDS with low blasts and SF3B1 mutation (MDS-SF3B1)
☐ MDS with biallelic TP53 inactivation (MDS-biTP53)
MDS, morphologically defined:
☐ MDS with low blasts (MDS-LB)
☐ MDS, hypoplastic (MDS-h)
☐ MDS with increased blasts (MDS-IB1)
☐ MDS with increased blasts (MDS-IB2)
☐ MDS with fibrosis (MDS-f)
Childhood myelodysplastic neoplasms (MDS):
☐ Childhood MDS with low blasts
☐ Childhood MDS with increased blasts
Therapy-related MDS:
(Secondary origin)
No
☐ Yes, disease related to prior exposure to therapeutic drugs or radiation ☐ Unknown
( If therapy-related MDS, is Yes)
Is this a donor cell leukaemia?
□ No
☐ Yes ☐ Not applicable (no previous allo HCT)
Unknown

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ЕВМТ		on Code (CIC): Jumber (UPN): Registry:	Treatment Type		
IPSS-R:	☐ Very Low (≤1.5) ☐ Low (>1.5 to 3) ☐ Intermediate (>6) ☐ High (>4.5 to 6) ☐ Very High (>6) ☐ Unknown	<b>IPSS-M:</b> 3 to 4.5)	<ul> <li>Very Low (≤-1.5)</li> <li>Low (&gt;-1.5 to -0.5)</li> <li>Moderate Low (&gt;-0.5 to 0)</li> <li>Moderate High (&gt;0 to 0.5)</li> <li>High (&gt;0.5 to 1.5)</li> <li>Very High (&gt;1.5)</li> <li>Unknown</li> </ul>		
Extended dataset  Assessments at diagnosis					
Haematolog Peripheral bl	gical values:				
<u>-</u>	(g/dL):	☐ Not evaluated ☐ Unknowi	n		
Platelets (10 <sup>9</sup>		☐ Not evaluated ☐ Unknown			
	Cells (10 <sup>9</sup> /L):	☐ Not evaluated ☐ Unknown			
% blasts:	20110 (10 72)	☐ Not evaluated ☐ Unknown			
% monocytes		☐ Not evaluated ☐ Unknown			
% neutrophils		Not evaluated Unknown			
Bone marrow  If the precise blast count is not available, please indicate whether it is:					
% blasts:	≤ 5%	<u> </u>	☐ Unknown		
Bone marro	ow investigation:				
Hypocellula	rity No	Yes Not evaluated	Unknown		
Fibrosis	□ No	Yes Not evaluated	☐ Unknown		

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ЕВМТ	EBMT Centre Identification Code (CIC): _ Hospital Unique Patient Number (UPN): _ Patient Number in EBMT Registry:			HCT CT		
	CHROMOSOME ANALYSIS					
escribe resul	escribe results of all the analysis done before HCT/CT/IST treatment					
Chromosom	e analysis done before HCT/CT/IST tr	eatment:				
☐ No ☐ Yes: ☐ Unkno	Output of analysis: ☐ Separa wn	te abnormalities	☐ Full karyotype			
	 Copy and fill-i	n this section as oft	en as necessary.			
What were the results?  Normal Abnormal: number of abnormalities present: Failed  Date of chromosome analysis:I(YYYY/MM/DD) Unknown  For abnormal results, indicate below whether the abnormalities were absent, present or not evaluated.						
del(Y)		] Absent [	Present	☐ Not evaluated	ı	
del(5q)		] Absent [	Present	☐ Not evaluated	ı	
del(20q)		] Absent [	Present	☐ Not evaluated	ı	
del(7q)		] Absent [	Present	☐ Not evaluated	i	
inv(3)		] Absent [	Present	☐ Not evaluated	ı	
t(3q;3q)		] Absent [	Present	☐ Not evaluated	ı	
del(3q)		] Absent [	Present	☐ Not evaluated	ı	
del(11q)		] Absent [	Present	☐ Not evaluated	ı	
Trisomy 8		] Absent [	Present	☐ Not evaluated	ı	
Trisomy 19	•	] Absent [	Present	☐ Not evaluated	ı	
i(17q)		] Absent [	Present	☐ Not evaluated	1	
Other; spec	cify	] Absent [	Present			

OR

Transcribe the complete karyotype: \_\_\_\_\_



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MOLECULAR MARKER ANALYSIS				
Molecular markers anal	lysis done before HCT/CT/IST t	reatment:		
☐ No				
Yes				
Unknown				
	 Copy and fill-in this se	ection as often as necessary.		
If molocular marker ans		-		
If molecular marker and				
	narker analysis: / /		<i>'</i> n	
Indicate below whether	the markers were absent, preser			
ASXL1	☐ Absent	☐ Present	☐ Not evaluated	
CBL	☐ Absent	☐ Present	☐ Not evaluated	
DDX41	☐ Absent	☐ Present	☐ Not evaluated	
ETV6	☐ Absent	☐ Present	☐ Not evaluated	
EZH2	☐ Absent	☐ Present	☐ Not evaluated	
IDH1	☐ Absent	☐ Present	☐ Not evaluated	
IDH2	☐ Absent	☐ Present	☐ Not evaluated	
JAK2	☐ Absent	☐ Present	☐ Not evaluated	
KRAS	Absent	☐ Present	☐ Not evaluated	
NPM1	Absent	☐ Present	☐ Not evaluated	
NRAS	Absent	☐ Present	☐ Not evaluated	
PTEN	Absent	 ☐ Present	 ☐ Not evaluated	
PTPN11	Absent	Present	☐ Not evaluated	
RUNX1	Absent	☐ Present	☐ Not evaluated	
SF3B1	☐ Absent	☐ Present	☐ Not evaluated	
SRSF2	☐ Absent	☐ Present	☐ Not evaluated	
TET2	☐ Absent	☐ Present	☐ Not evaluated	
TP53	Absent	☐ Present:	☐ Not evaluated	
	TP53 mutation type: Single hit			
		☐ Multi hit		
☐ Unknown				
UBA1	☐ Absent	☐ Present	☐ Not evaluated	

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Present

☐ Not evaluated

Absent

Other; specify



☐ Unknown

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Extended da	ataset			
PREVIOUS THERAPIES (between diagnosis and HCT/CT)				
Previous th	erapy lines before the HCT/CT:			
□ No				
☐ Yes:	complete the "Treatment non-HCT/CT/GT/IST" form			

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