

Treatment Type	🗌 НСТ	🗌 СТ	🗌 IST	Other

Treatment Date _ _ _ / _ / _ _ (YYY/MM/DD)

MYELOPROLIFERATIVE NEOPLASMS (MPN)

DISEASE

Note: complete this form only if this diagnosis was the indication for the HCT/CT or if it was specifically requested. Consult the manual for further information.

Date of diagnosis: _ _ _ / _ / _ (YYY//MM/DD)

Classification (WHO 2022):

Primary myelofibrosis
Polycythaemia vera (PV)
Essential or primary thrombocythaemia (ET)
U Juvenile myelomonocytic leukaemia (JCMMoL, JMML, JCML, JCMML)
Hyper eosinophilic syndrome (HES)
Chronic eosinophilic leukaemia (CEL)
Chronic neutrophilic leukaemia (CNL)
Aggresive systemic mastocytosis
Systemic mastocytosis with an associated haematologic neoplasm (SM-AHN)
Mast cell leukaemia
Mast cell sarcoma
MLN-TK with FGFR1 rearrangement
MLN-TK with PDGFRA rearrangement
MLN-TK with PDGFRB rearrangement
MLN-TK with JAK2 rearrangement
MLN-TK with FLT3 rearrangement
MLN-TK with ETV6::ABL1 fusion
MPN not otherwise specified (NOS)
Other; specify:

Therapy-related MPN:

(Secondary origin)

🗌 No

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

Unknown

(EBM	Т
	-	

 EBMT Centre Identification Code (CIC):
 Treat

 Hospital Unique Patient Number (UPN):
 Treat

 Patient Number in EBMT Registry:
 Treat

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Treatment Date _	// (YY	YY/MM/DE))

MPN ASSESSMENTS

(Palpable) spleen size: cm (below costal margin) 🔲 Not evaluated 🔲 Unknown
Spleen span on ultrasound or CT scan: cm (maximum diameter) 🔲 Not evaluated 📋 Unknown
Transfusion dependency: No Yes Unknown
Bone marrow fibrosis: Grade 0 Grade 1 Grade 2 Grade 3 Not evaluated Unknown
Blast count (peripheral blood): % 🔲 Not evaluated 📄 Unknown
Myelofibrosis only: IPSS: □ Low risk
Intermediate-1
☐ Intermediate-2 ☐ High risk
 Not evaluated Unknown
DIPSS: Low risk Intermediate-1 Intermediate-2 High risk Not evaluated Unknown
MIPSS70: Low risk Intermediate High risk Not evaluated Unknown

EBMT Centre Identification Code (CIC): Hospital Unique Patient Number (UPN):	Treatment Type 🔲 HCT 🗌 CT 📋 IST 🗌 Other
EBMT Hospital Unique Patient Number (UPN): Patient Number in EBMT Registry:	Treatment Date / _ / _ (YYYY/MM/DD)
CHROMOSOME A	NALYSIS
Describe results of all the analyses done before HCT/CT/IST treatmer	nt
Chromosome analysis done before HCT/CT/IST treatment:	
☐ No	
 Yes: Output of analysis: Separate abnormalities Unknown 	Full karyotype
Copy and fill-in this section as	often as necessary.
If chromosome analysis was done:	
What were the results?	
🔲 Normal	
Abnormal: number of abnormalities present:	
Failed	
Date of chromosome analysis: / / / (YYY//MM//	(סכ 🗍 Unknown
	<i>(</i>) <u> </u>
For abnormal results, indicate below whether the abnormalities were	Absent
abn 1 type; specify:	Absent Present Not evaluated
abn 5 type; specify:	Absent Present Not evaluated
abn 7 type; specify:	Absent Present Not evaluated
Trisomy 8	
Trisomy 9	
del(20q)	Absent Present Not evaluated
del(13q)	Absent Present Not evaluated
Other; specify:	Absent Present
OR	
Transcribe the complete karyotype:	

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Treatment Date _	//(YY	YY/MM/DD)

MOLECULAR MARKER ANALYSIS

Molecular marker analysis done before HCT/CT/IST treatment:		
☐ Yes		
Copy and fill-in this section as often as ne	cessarv.	
If molecular marker analysis was done:	-	
Date of molecular marker analysis: / _ / _ (YYYY/MM/DD)		wn
Indicate below whether the markers were absent, present or not evaluated ASXL1		☐ Present ☐ Not evaluated
BCR::ABL1; Molecular product of t(9;22)(q34;q11.2)		
	Absent	Present Not evaluated
	Absent	Present Not evaluated
	If prese	ent: 🔲 Type 1
CALR		Type 2
		🔲 Type 1 like
		Type 2 like
		Unknown
CBL	Absent	Present Not evaluated
cMPL	Absent	Present Not evaluated
CSF3R	Absent	Present Not evaluated
CUX1	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
NRAS	Absent	Present Not evaluated
PTEN	Absent	Present Not evaluated
PTPN-11	Absent	Present Not evaluated
RUNX1	Absent	Present Not evaluated
SF3B1	Absent	Present Not evaluated
SRSF2	Absent	Present Not evaluated
TET2	Absent	Present Not evaluated
	Absent	Present Not evaluated
TP53 TP55	3 mutation ty	
		Multi hit
U2AF1	Absent	Present Not evaluated
UBA1	Absent	Present Not evaluated
Other; specify:	Absent	Present