

EBMT Centre Identification Code (CIC):
Hospital Unique Patient Number (UPN):
Patient Number in ERMT database:

Treatment Type	□ нст □ ст	☐ IST ☐ Other
- Treatment Date	/ / (YY	YY/MM/DD)

NON-I	INDICATI	on d	IAGN	IOSIS
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C	<b>Date</b>	of diagnosis: / (YYYY/MM/DD)
C	Class	sification:
		Acute leukaemia
		Autoimmune disorder
		Bone marrow failure syndrome including aplastic anaemia (BMF incl. AA)
		Chronic leukaemia
		Haemoglobinopathy
		Lymphoma
		Combined myelodysplastic syndrome/myeloproliferative neoplasms (MDS/MPN)
		Myelodysplastic syndrome (MDS)
		Myeloproliferative neoplasm (MPN)
		Plasma cell disorder including multiple myeloma (PCD incl. MM)
		Solid tumour



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Classification:
☐ Acute myeloid leukaemia (AML)
☐ Precursor lymphoid neoplasm (ALL)
☐ Other acute leukaemia



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Classification:
Connective tissue:
Systemic sclerosis (SSc)  SSc type:  diffuse cutaneous  limited cutaneous  SSc sine scleroderma  Other; specify:  Systemic lupus erythematosus (SLE)  Mixed connective tissue disease (MCTD)  Polymyositis/Dermatomyositis (PM/DM)  Sjögren syndrome  Antiphospholipid syndrome  Other connective tissue disease; specify:
Granulomatosis with polyangiitis (GPA); formerly Wegener granulomatosis  Classical polyarteritis nodosa  Microscopic polyarteritis nodosa  Eosinophilic granulomatosis with polyangiitis (EGPA); formerly Churg-Strauss  Overlap necrotising arteritis  Behçet syndrome  Takayasu arteritis  Other; specify:
Arthritis:  Adult onset stills disease (AOSD)  Rheumatoid arthritis  Psoriatic arthritis/psoriasis  Juvenile idiopathic arthritis (JIA), systemic (Still's disease)  Juvenile idiopathic arthritis (JIA), articular  oligoarticular onset  polyarticular onset  Other juvenile idiopathic arthritis; specify:



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## **Classification (continued): Neurological diseases:** ☐ Chronic inflammatory demyelinating polyneuropathy (CIDP) ☐ Neuromyelitis optica (NMO) or NMO spectrum disorders (NMOSD) ☐ Other autoimmune neurological disorder; specify: \_ **Haematological diseases:** ☐ Idiopathic thrombocytopenic purpura (ITP) ☐ Haemolytic anaemia ☐ Evan syndrome Autoimmune lymphoproliferative syndrome (primary diagnosis, not subsequent to transplant) Other haematological autoimmune disease; specify: **Inflammatory bowel diseases:** ☐ Celiac disease ☐ Crohn's disease ☐ Ulcerative colitis ☐ Other autoimmune bowel disease; specify: \_ Other autoimmune diseases: ☐ Insulin-dependent diabetes mellitus (IDDM) ☐ Thyroid autoimmune disease

Other autoimmune disease; specify:

☐ IST ☐ Other



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## Bone marrow failure syndromes (BMF) incl. aplastic anaemia (AA)

Classification:	
Acquired:	
Aplastic anaemia (AA)	
☐ Moderate	
Severe	
☐ Very Severe	
☐ Pure red cell aplasia (non-congenital PRCA)	
Paroxysmal nocturnal haemoglobinuria (PNH)	
☐ Pure white cell aplasia	
Amegakaryocytosis / Thrombocytopenia (non-congenital)	
Other acquired cytopenic syndrome; specify:	
☐ Genetic:	
Amegakaryocytosis / Thrombocytopenia (congenital)	
☐ Fanconi anaemia	
☐ Diamond-Blackfan anaemia (congenital PRCA)	
☐ Shwachman-Diamond syndrome	
Dyserythropoietic anaemia	
☐ Dyskeratosis congenita	
Other congenital anaemia; specify:	



EDMT	EBMT Centre Identification Code (CIC): Hospital Unique Patient Number (UPN):	Treatment Type
ЕВМТ	Patient Number in EBMT database:	Treatment Date / (YYYY/MM/DD)
	Chronic le	ukaemias
Classification	ո։	
Chronic my	yeloid leukaemia (CML)	
☐ Chronic lyr	nphocytic leukaemia (CLL)/ small lymphocytic lymph	noma
☐ Richter's sy	yndrome	
Prolympho	cytic leukaemia (PLL)	
☐ PLL	; B-cell	
☐ PLL	; T-cell	
☐ Hairy cell l	eukaemia	
☐ Hairy cell I	eukaemia variant (atypic)	
☐ Other chro	onic leukaemia; specify:	
C	Combined myelodysplastic syndroi (MDS/I	mes/myeloproliferative neoplasms MPN)
Classification	n:	
Chronic m	yelomonocytic leukaemia (CMMoL, CMML)	

☐ Juvenile myelomonocytic leukaemia (JCMMoL, JMML, JCML, JCMML)

☐ MDS/MPN with ring sideroblasts and thrombocytosis (MDS/MPN-RS-T)

☐ Atypical CML (t(9;22) negative and BCR-ABL1 negative)



Myeloprofilerative neopla	asms (MPN)			
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Myeloprofile	rative neop	lasms (MPN)
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Classification (WHO 2016):
Primary myelofibrosis
Polycythaemia vera (PV)
Essential or primary thrombocythaemia (ET)
Hyper eosinophilic syndrome (HES)
Chronic eosinophilic leukaemia (CEL)
Chronic neutrophilic leukaemia (CNL)
Systemic mastocytosis
Mast cell leukaemia
☐ Mast cell sarcoma
MPN not otherwise specified
Myeloid and lymphoid neoplasms with FGFR1 abnormalities (Stem cell leukaemia-lymphoma syndrome, 8p11 syndrome)
Myeloid and lymphoid neoplasms with PDGFRA rearrangement
Myeloid and lymphoid neoplasms with PDGFRB rearrangement
Myeloid and lymphoid neoplasms with PCM1-JAK2 rearrangement
Other; specify:

## Plasma cell disorders (PCD) incl. multiple myeloma (MM)

## **Classification:**

☐ Multiple myeloma (MM)
Plasma cell leukaemia
Solitary plasmacytoma of bone
Primary amyloidosis
POEMS (polyneuropathy, organomegaly, endocrinopathy/edema, monoclonal-protein, skin changes)
Monoclonal light and heavy chain deposition disease (LCDD/HCDD)
Other; specify:



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Lymp	homas
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Classification:
☐ B-cell non-Hodgkin lymphoma (NHL)
Sub-classification: Mature B-cell neoplasms
Splenic marginal zone lymphoma
Extranodal marginal zone lymphoma of mucosa associated lymphoid tissue (MALT)
☐ Nodal marginal zone lymphoma
Lymphoplasmacytic lymphoma (LPL)
Follicular lymphoma
Primary cutaneous follicle centre lymphoma
☐ Mantle cell lymphoma
☐ T-cell/histiocyte-rich large B-cell lymphoma
☐ Primary DLBCL of the CNS
☐ Primary cuaneous DLBCL, leg type
☐ EBV positive DLBCL of the elderly
Germinal centre B-cell type (GCB) DLBCL
Activated B-cell type (ABC or non-GCB) DLBCL
☐ DLBCL associated with chronic inflammation
☐ Lymphomatoid granulomatosis
Primary mediastinal (thymic) large B-cell lymphoma
☐ Intravascular large B-cell lymphoma
☐ ALK-positive large B-cell lymphoma
☐ Plasmablastic lymphoma
HHV8-positive DLBCL, NOS
☐ Diffuse large B-cell lymphoma (DLBCL), (NOS)
Primary effusion lymphoma (PEL)
— ☐ Burkitt lymphoma (BL)
High-grade B-cell lymphoma with MYC and BCL2 and/or BCL6 rearrangements
B-cell lymphoma, unclssifiable, with features intermediate between diffuse large B-cell lymphoma and
Burkitt lymphoma (intermediate DLBCL/BL)
B-cell lymphoma, unclssifiable, with features intermediate between diffuse large B-cell lymphoma and classical Hodgkin lymphoma (Gray zone lymphoma)
Other B-cell lymphoma; specify:
☐ Immunodeficiency-associated lymphoproliferative disorder (incl. PTLD)
Sub-classification: Immunodeficiency-associated lymphoproliferative disorders (incl. PTLD)
Lymphoproliferative disease associated with primary immune disorder
Post-transplant lymphoproliferative disorder (PTLD)
Other iatrogen immunodeficiency-associated lymphoproliferative disorder



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ification (continued):  cell non-Hodgkin lymphoma (NHL)  Sub-classification: Mature T-cell & NK-cell Neoplasms  T-cell large granular lymphocytic leukaemia  Aggressive NK-cell leukaemia  Systemic EBV positive T-cell lymphoproliferative disease of childhood  Hydroa vacciniforme-like lymphoma  Adult T-cell leukaemia/lymphoma  Extranodal NK/T-cell lymphoma, nasal type  Enteropathy-associated T-cell lymphoma  Monomorphic epitheliotropic intestinal T-cell lymphoma  Hepatosplenic T-cell lymphoma  Subcutaneous panniculitis-like T-cell lymphoma  Mycosis fungoides (MF)  Sézary syndrome  Lymphomatoid papulosis  Primary cutaneous anaplastic large cell lymphoma  Primary cutaneous gamma-delta T-cell lymphoma
Sub-classification: Mature T-cell & NK-cell Neoplasms  T-cell large granular lymphocytic leukaemia Aggressive NK-cell leukaemia Systemic EBV positive T-cell lymphoproliferative disease of childhood Hydroa vacciniforme-like lymphoma Adult T-cell leukaemia/lymphoma Extranodal NK/T-cell lymphoma, nasal type Enteropathy-associated T-cell lymphoma Monomorphic epitheliotropic intestinal T-cell lymphoma Hepatosplenic T-cell lymphoma Subcutaneous panniculitis-like T-cell lymphoma Mycosis fungoides (MF) Sézary syndrome Lymphomatoid papulosis Primary cutaneous anaplastic large cell lymphoma
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Hepatosplenic T-cell lymphoma  Subcutaneous panniculitis-like T-cell lymphoma  Mycosis fungoides (MF)  Sézary syndrome  Lymphomatoid papulosis  Primary cutaneous anaplastic large cell lymphoma
Subcutaneous panniculitis-like T-cell lymphoma  Mycosis fungoides (MF)  Sézary syndrome  Lymphomatoid papulosis  Primary cutaneous anaplastic large cell lymphoma
Mycosis fungoides (MF)  Sézary syndrome  Lymphomatoid papulosis  Primary cutaneous anaplastic large cell lymphoma
Sézary syndrome  Lymphomatoid papulosis  Primary cutaneous anaplastic large cell lymphoma
Primary cutaneous anaplastic large cell lymphoma
Primary cutaneous CD8 positive aggressive epidermotropic cytotoxic T-cell lymphoma
☐ Primary cutaneous CD4 positive small/medium T-cell lymphoma
☐ Peripheral T-cell lymphoma NOS (PTCL)
☐ Angioimmunoblastic T-cell lymphoma
☐ Anaplastic lare-cell lymphoma (ALCL), ALK-positive
☐ Anaplastic lare-cell lymphoma (ALCL), ALK-negative
☐ Other T-cell lymphoma; specify:

**Sub-classification:** Hodgkin lymphomas

☐ Nodular lymphocyte predominant
Classical predominant; lymphocyte-rich
Classical predominant; nodular sclerosis
Classical predominant; mixed cellularity
Classical predominant; lymphocyte-depleted
Classical predominant; NOS
☐ Other Hodgkin lymphoma; specify:



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Haemoglobinopathy	
Classification:	
☐ Thalassaemia	
☐ Sickle Cell Disease	
Other Haemoglobinopathy; specify:	

Solid tumours
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<u>Classification:</u>
☐ Bone sarcoma (excluding Ewing sarcoma/PNET)
☐ Breast
Central nervous system tumours (including CNSPNET)
Ewing sarcoma (ES) / PNET, extra-skeletal
Ewing sarcoma (ES) / PNET, skeletal
Ewing sarcoma (ES)/PNET, not classified
Germ cell tumour, extragonadal only
Germ cell rumour, gonadal
GI tract and Hepatopancreatic cancers
☐ Kidney cancer excluding Wilm's tumour
Lung cancer, small cell
Lung cancer, non small cell (NSCLC)
☐ Medulloblastoma
☐ Melanoma
☐ Nasopharingeal carcinoma
☐ Neuroblastoma
Ovarian (carcinoma)
☐ Prostate
Retinoblastoma
☐ Rhabdomyosarcoma
☐ Soft tissue sarcoma (excluding Rhabdo and extra-skeletal ES)
☐ Thymoma
☐ Wilm's tumour
Other; specify: