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Version Number | 1.0

Title | Lymphomas

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☐ Hodgkin lymphoma

EBMT Centre Identification Code (CIC): ____

☐ Immunodeficiency-associated lymphoproliferative disorder (incl. PTLD)

ЕВМТ	Hospital Unique Patient Number (UPN): Patient Number in EBMT database:	Treatment Date / (YYYY/MM/DD)
	LYMPHO	DMAS
	DISEA	SE
•	te this form only if this diagnosis was the indicat	ion for the HCT/CT or if it was specifically requested.
Date of diagn	nosis: / (YYYY/MM/DD)	
Classification	n:	
☐ B-cell non	n-Hodgkin lymphoma (NHL)	
☐ T-cell non	-Hodgkin lymphoma (NHL)	

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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/DI	D)

LYMPHOMAS B-Cell Non-Hodgkin Lymphomas (NHL)

DISEASE	
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)	
☐ Waldenstrom macroglobulinaemia (LPL with monoclonal IgM)	
☐ Follicular lymphoma ☐ Grading: ☐ Grade I ☐ Grade II ☐ Grade IIIa ☐ Grade IIIb ☐ Not evaluated	
Primary cutaneous follicle centre lymphoma	
☐ T-cell/histiocyte-rich large B cell lymphoma	
Primary DLBCL of the CNS	
Primary cutaneous 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	KI-67: % positive
DLBCL associated with chronic inflammation	(proliferation index)
Lymphomatoid granulomatosis	☐ Not evaluated
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, unclassifiable, with features intermediate between diffuse large B-cell lymphoma and Burkitt lymphoma (Intermediate DLCBL/BL)	
B-cell lymphoma, unclassifiable, with features intermediate between diffuse large B-cell lymphoma and classical Hodgkin lymphoma (Gray zone lymphoma)	
Other B-cell non-Hodgkin lymphoma; specify:	THIS IS AN LINCONTOLLED CORV



EBMT Ho	ospital Unique	entification Code Patient Number In EBMT databa	· (UPN):			-	// CT	☐ IST ☐ Other
			DIS	EASE co	ntinued			
High-grade transformation of indolent B-cell lymphoma? No Yes (If not reported yet, complete respective non-indication diagnosis form in addition to the current form) Unknown								
Parameters for		l prognostic						
Age at diagnos		——————————————————————————————————————	years	□ Net e				
LDH levels elev		□ No	Yes		evaluated			
Ann Arbor stag	ing:				□IV	☐ Not eva	luated	
ECOG performa	ance status:	□ 0	<u> </u>	_ 2	□ 3	□ 4	☐ Not evaluated	d
> 1 extranodal s	site involved	: No	☐ Yes	☐ Not €	evaluated			
> 4 nodal sites	involved:	☐ No	☐ Yes	□ Not €	evaluated			
Hemoglobin < 1	.20g/L:	☐ No	☐ Yes	☐ Not €	evaluated			
White Blood Ce	ll count:		x 10 ⁹ cel	ls/L		☐ Not eval	uated	
			CHRON	MOSOME	ANALYSIS			
the following typeMantle cell lyWaldenstrom	es of B-cell Ni Imphoma I macroglobui		with monoc		and immund	pphenotyping s	sections only for µ	patients with
Chromosome aı	-	before treatr	ment (all me	thods inclu	ding FISH):			
☐ Not done or								
_		umber of abno	ormalities pre	esent:	_			
☐ Yes, norma ☐ Unknown	resuits							
Date of chromos	some analysi	is (if tested)··	/	/ (YYY	Y/MM/DD)			
Indicate below whether the abnormalities were absent, present or not evaluated. according to the type of lymphoma diagnosed. Mantle cell lymphoma del(17p)								
Mantle cell lymphor Waldenstrom mglobulinaemia	J	. (μ)			FISH used:	☐ Absent	☐ Present ☐ ☐ Yes	j Not evaluateu
	t(2;8)				Absent	Present	Not evaluated
Burkitt lymphoma		4)				Absent	Present	Not evaluated
Intermediate DLB	CL/BL t(8;2	2)				Absent	Present	Not evaluated
All D - III I	t(14;	-	 	liai	:4	Absent	Present	Not evaluated
All B-cell lymphoi		chromosom					Present S AN UNCONTOLE	ED COPY



Burkitt lymphoma or

Intermediate DLBCL/BL

Intermediate DLBCL/BL

All B-cell lymphomas

MYC

BCL6

BCL2/lgH

Other immunophenotype; specify:

EBMT Hospital U	entre Identification Code (CIC): Unique Patient Number (UPN): umber in EBMT database:	Treatment Type
	MOLECULAR MAR	RKER ANALYSIS
the following types of BMantle cell lymphorWaldenstrom macro	-cell NHL:	is and immunophenotyping sections only for patients with
☐ No ☐ Yes ☐ Unknown Date of molecular mar	ysis done before treatment: ker analysis (if tested):://	
Indicate below whether	the markers were absent, present or not e	evaluated, according to the type of lymphoma diagnosed.
Mantle cell lymphoma	TP53 mutation	☐ Absent ☐ Present ☐ Not evaluated
Burkitt lymphoma or Intermediate DLBCL/BL	myc rearrangment	☐ Absent ☐ Present ☐ Not evaluated
Intermediate	BCL2 rearrangement	☐ Absent ☐ Present ☐ Not evaluated
DLBCL/BL	BCL6 rearrangement	Absent Present Not evaluated
All B-cell lymphomas	Other molecular markers; specify:	Absent Present
	IMMUNOPHE	NOTYPING
the following types of EMantle cell lymphoWaldenstrom macr	3-cell NHL:	sis and immunophenotyping sections only for patients with
Immunophenotyping	done before treatment:	
□ No		
Yes		
Unknown		
Date of immunophen	otyping (if tested)://(YYY	/Y/MM/DD)
Indicate below whether diagnosed.	the immunophenotypes were absent, pre	esent or not evaluated, according to the type of lymphoma
Mantle cell lymphoma	SOX 11	☐ Absent ☐ Present ☐ Not evaluated

Il lymphomas Other immunophenotype; specify: _____ Descriptor Absent Present Present Present Present Other immunophenotype; specify: _____ Descriptor Desc

☐ Absent ☐ Present ☐ Not evaluated

☐ Present ☐ Not evaluated

☐ Present ☐ Not evaluated

Absent

☐ Absent



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Hospital Unique Patient Number (UPN):				
Patient Number in EBMT database:	Treatment Date	//YY	YY/MM/DI)

LYMPHOMAS T-Cell Non-Hodgkin Lymphomas (NHL)

DISEASE			
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			
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 large-cell lymphoma (ALCL), ALK-positive			
Anaplastic large-cell lymphoma (ALCL), ALK-negative			
Other T-cell non-Hodgkin lymphoma; specify:			

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Patient Number in EBMT database:	Treatment Date / _ / _ (YYYY/MM/DD)			D)	
LVMDHOMAC					

LYMPHOMAS Hodgkin Lymphomas

DISEASE
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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Hospital Unique Patient Number (UPN):				
Patient Number in EBMT database:	Treatment Date _	//(YY	YY/MM/DI	D)

LYMPHOMAS

Immunodeficiency-associated lymphoproliferative disorders (incl. PTLD)

DISEASE				
Sub-Classification: Immunodeficiency-associated lymphoproliferative disorders (incl. PTLD)				
Lymphoproliferative disease associated with primary immune disorder				
Lymphoma associated with HIV infection				
Post-transplant lymphoproliferative disorder (PTLD)				
☐ Non-destructive PTLD				
☐ Plasmacytic hyperplasia PTLD				
☐ Infectious mononucleosis PTLD				
☐ Florid follicular hyperplasia PTLD				
☐ Polymorphic PTLD				
☐ Monomorphic PTLD				
☐ B-cell type				
☐ T-/NK-cell type				
☐ Classical Hodgkin lymphoma PTLD				
Other immunodeficiency-associated lymphoproliferative disorder				
Did the disease result from a previous solid organ transplant? □ No				
Yes: Date of transplant:/ (YYYY/MM/DD)				
Type of transplant:				
☐ Cardiac				
☐ Pulmonary				
Other; specify:				
☐ Unknown				



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PREVIOUS THERAPIES BEFORE HCT/CT

Line of treatment	Regimen used*	Treatment start date (YYYY/MM/DD)	Response to this line of treatment	Response assessment date (YYYY/MM/DD)
1		//	 ☐ Complete remission (CR) ☐ Partial remission (PR) ☐ Stable disease ☐ Chemorefractory relapse or progression incl. primary refractory disease ☐ Unknown 	
2		//	 ☐ Complete remission (CR) ☐ Partial remission (PR) ☐ Stable disease ☐ Chemorefractory relapse or progression incl. primary refractory disease ☐ Unknown 	
3			 ☐ Complete remission (CR) ☐ Partial remission (PR) ☐ Stable disease ☐ Chemorefractory relapse or progression incl. primary refractory disease ☐ Unknown 	
4		//	 ☐ Complete remission (CR) ☐ Partial remission (PR) ☐ Stable disease ☐ Chemorefractory relapse or progression incl. primary refractory disease ☐ Unknown 	//

Copy and fill-in this section as many times as necessary

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^{*}Please consult the **LIST OF CHEMOTHERAPY DRUGS/AGENTS AND REGIMENS** on the EBMT website for drugs/regimens names