

☐ T-cell non-Hodgkin lymphoma (NHL)

Other; specify __

☐ Immunodeficiency-associated lymphoproliferative disorder (incl. PTLD)

EBMT	EBMT Centre Identification Code (CIC): Hospital Unique Patient Number (UPN): Patient Number in EBMT Registry:	Treatment Type HCT CT GT IST Other Treatment Date/_/_(YYYY/MM/DD)
	LYMPHOMAS	
	DISEASE	
	te this form only if this diagnosis was the indication for nanual for further information.	the HCT/CT or if it was specifically requested.
Date of diagn	osis: / / (YYYY/MM/DD)	
Classification	1:	
☐ B-cell lym	phoma (including Hodgkin and Non-Hodgkin lymphoma)	



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Treatment Type H	СТ] CT	GT IST Other	
Treatment Date	1	1	(YYYY/MM/DD)	

LYMPHOMAS B-cell lymphoma (including Hodkin and Non-Hodkin lymphoma)

DISEASE
Sub-Classification: Mature B-cell neoplasms
Splenic B-cell lymphomas and leukaemias
Splenic marginal zone lymphoma
Splenic diffuse red pulp small B-cell lymphoma
Lymphoplasmacytic lymphoma
☐ IgM-LPL/ Waldenström Macroglobulinaemia (WM) type
☐ Non-WM type LPL
☐ Marginal zone lymphoma
Extranodal marginal zone lymphoma of mucosa-associated lymphoid tissue
☐ Primary cutaneous marginal zone lymphoma
☐ Nodal marginal zone lymphoma
☐ Paediatric marginal zone lymphoma
☐ Follicular lymphoma
☐ Classical follicular lymphoma (cFL)
— Follicular large B-cell lymphoma (FLBL)
☐ FL with uncommon features (uFL)
☐ Paediatric-type follicular lymphoma
☐ Duodenal-type follicular lymphoma
Cutaneous follicle centre lymphoma
☐ Mantle cell lymphoma
☐ Mantle cell lymphoma
☐ Leukaemic non-nodal mantle cell lymphoma



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LYMPHOMAS B-cell lymphoma (including Hodkin and Non-Hodkin lymphoma)

DISEASE continued

Sub-Classification: Mature B-cell neoplasms
☐ Large B-cell lymphomas
☐ Diffuse large B-cell lymphoma (DLBCL), NOS
☐ Germinal centre B- cell-like subtype (GCB)
☐ Activated B-cell-like subtype (ABC)
│ │
☐ Diffuse large B-cell lymphoma/ high grade B-cell lymphoma with MYC and BCL2 rearrangements
ALK-positive large B-cell lymphoma
☐ Large B-cell lymphoma with IRF4 rearrangement
│ │ │ │ High-grade B-cell lymphoma with 11q aberrations
☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐ ☐
☐ FBV-positive diffuse large B-cell lymphoma
☐ Diffuse large B-cell lymphoma associated with chronic inflammation
│
☐ Plasmablastic lymphoma
Primary large B-cell lymphoma of immune-privileged sites
☐ Primary large B-cell lymphoma of the CNS
☐ Primary large B-cell lymphoma of the vitreoretina
☐ Primary large B-cell lymphoma of the testis
☐ Primary cutaneous diffuse large B-cell lymphoma, leg type
☐ Intravascular large B-cell lymphoma
☐ Primary mediastinal large B-cell lymphoma
☐ Mediastinal grey zone lymphoma
☐ High-grade B-cell lymphoma, NOS
☐ Burkitt lymphoma
☐ EBV-positive BL
☐ EBV-negative BL
KSHV/HHV8-associated B-cell lymphoid proliferations and lymphomas
☐ Primary effusion lymphoma
☐ KSHV/HHV8-positive diffuse large B-cell lymphoma
☐ KSHV/HHV8-positive germinotropic lymphoproliferative disorder
☐ Hodgkin lymphoma
☐ Classic Hodgkin lymphoma
☐ Nodular lymphocyte predominant Hodgkin lymphoma



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DISEASE continued							
Transformation of indole	nt B-cell lymphoma	.1					
□ No							
☐ Yes (If not reported ye	t, complete respecti	e non-indication	n diagnosis form	m in a	ddition to the current form)		
Unknown							
Parameters for internation	nal prognostic indi	ces:					
Age at diagnosis:		years (this is ca	lculated autom	natical	lly in the database)		
LDH levels elevated:	□ No □] Yes No	ot evaluated	Πn	nknown		
Ann Arbor staging:			□IV] Not evaluated Unknown		
ECOG performance statu	s: 0 0	1 2	3 [] 4	☐ Not evaluated ☐ Unknown		
> 1 extranodal site involv	ed: No [] Yes No	ot evaluated	□ 0	nknown		
> 4 nodal sites involved:	□ No □] Yes 🔲 No	ot evaluated	□ u	nknown		
Haemoglobin < 12g/dL:	□ No □] Yes 🔲 No	ot evaluated	□ u	nknown		
White Blood Cell count:	;	< 10 ⁹ /L □ No	ot evaluated	□ u	nknown		
CNS Involvement:	□ No □] Yes No	ot evaluated	□ U	nknown		
Final score: (only for patients with LBCL (except Primary large B-cell lymphoma of immune-privileged sites), Mantle cell lymphoma, Follicular lymphoma, Waldenstrom macroglobulinaemia) IPI: (for LBCL (except Primary large B-cell lymphoma) FLIPI: (for Follicular lymphoma (except Primary large B-cell lymphoma) FLIPI: (for Follicular lymphoma (except Primary large Immune-privileged sites) and FLBL)							
Low risk (0-1 score points)	Low risk	☐ Low	risk		Low risk (0-1 score points except age > 65)		

Low-intermediate risk

High-intermediate risk

(2 score points)

(3 score points)

(4-5 score points)

☐ Not evaluated

☐ High risk

☐ Intermediate risk

☐ Not evaluated

☐ High risk

☐ Intermediate risk

☐ Not evaluated

(3-5 score points)

☐ High risk

(2 score points OR age > 65)

☐ Not evaluated

☐ High risk



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CHROMOSOME ANALYSIS

Please complete chromosome analysis section only for patients with the following types of B-cell NHL:

• Mantle cell lymphoma (including Leukaemic non-nodal mantle cell lymphoma) & for Waldenström Macroglobulinaemia (IgM-LPL/Waldenström Macroglobulinaemia (WM) in new classification)

• Burkitt lymphoma (ii	ncluding EBV-positive BL & EBV-negative BL) & for all LBCL
• For all B-cell lymphon	na,
	sis done before HCT/CT treatment:
•	he most recent complete analysis)
☐ No	
	Output of analysis: Separate abnormalities Full karyotype
Unknown	
E	Extended dataset
	Chromosome analysis method used: Karyotyping
	(select all that apply)
L	
If chromosome and	alveis was dono:
What were the re	
☐ Normal	
_	nber of abnormalities present:
☐ Failed	
Date of chromoso	ome analysis:: / _ / _ (<i>YYYY/MM/DD</i>)
Date of official	
	, indicate below whether the abnormalities were absent, present or not evaluated (according to the type of
lymphoma diagnosed)	
Mantle cell lymphoma	del(17p) ☐ Absent ☐ Present ☐ Not evaluated ☐ Unknown
or Waldenstrom macro	0-
globulinaemia	FISH used: No Yes
	t(2;8) Absent Present Not evaluated Unknown
Burkitt lymphoma or a	II t(8;14)
LBCL	t(8;22) Absent Present Not evaluated Unknown
	t(14;18) Absent Present Not evaluated Unknown
All above mentioned	Other chromosome abnormalities; specify: Absent Present
B-cell lymphomas	
	OR
Transcribe the comp	plete karyotype:

Lymphomas_Core_Extended_v2.3



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		MOLEC	CULAR MARKER A	NALYSIS			
		ular marker analysis section of the control of the		he following	types of B-cell NHL:		
• Burkitt lympho	oma (includ	ling EBV-positive BL & EBV-neg	ative BL)				
• All LBCL are B	CL2 rearra	ngement & BCL6 rearrangement					
• For all B-cell lyı	mphomas						
	lts of the	ysis done before HCT/CT t most recent complete analys					
Date of molec	ular mar	ker analysis (if tested)::	/(YYYY//	MM/DD)] Unknown		
Indicate below	whether	the markers were absent, pro	esent or not evaluated	l, according t	to the type of lympho	ma diagnosed.	
Mantle cell lym	phoma	TP53 mutation	☐ Absent	☐ Present	☐ Not evaluated	☐ Unknown	
Burkitt lymphoi LBCL	ma or all	MYC rearrangement	Absent	☐ Present	☐ Not evaluated	☐ Unknown	
All LBCL		BCL2 rearrangement	☐ Absent	Present	☐ Not evaluated	Unknown	
W LBOL		BCL6 rearrangement	☐ Absent	☐ Present	☐ Not evaluated	Unknown	
All above ment B-cell lymphon		Other molecular markers;	specify:		Absent Present		
		IM	MUNOPHENOTYPI	NG			
• Mantle cell ly	mphoma (noma (inclu	nophenotyping section only i including Leukaemic non-nodal i iding EBV-positive BL & EBV-ne ç	mantle cell lymphoma)	llowing types	s of B-cell NHL:		
<u>-</u>	ults of the	done before HCT/CT treatn most recent complete analy					
Date of immu	unophen	otyping (if tested):/	/ (YYYY/MM/D	D)			
Indicate belov diagnosed.	v whether	the immunophenotypes wer	re absent, present or r	not evaluated	d, according to the ty _l	pe of lymphoma	
Mantle cell lym _l	ohoma	SOX 11	Absent Pres	ent 🔲 Not	evaluated 🔲 U	nknown	
Burkitt lymphon LBCL	na or all	мус	Absent Pres	ent 🗌 Not	evaluated 🛭 U	nknown	
RCI.		BCL2/lgH	Absent Pres	ent 🔲 Not	evaluated 🔲 U	nknown	
LBCL		BCL6	Absent Pres	ent Not	evaluated 🔲 U	nknown	
_	_				_		

All above mentioned

B-cell lymphomas

- ☐ Absent ☐ Present

Other immunophenotype; specify:



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LYMPHOMAS T-cell non-Hodgkin lymphoma (NHL)

DISEASE	
Sub-Classification: Mature T-cell & NK-cell neoplasms	
☐ Mature T-cell and NK-cell leukaemias	
T-large granular lymphocytic leukaemia	
☐ NK-large granular lymphocytic leukaemia	
Adult T-cell leukaemia/lymphoma	
Sezary syndrome	
Aggressive NK-cell leukaemia	
☐ Primary cutaneous T-cell lymphomas	
Primary cutaneous CD4-positive small or medium T-cell lymphoproliferative disorder	
Primary cutaneous acral CD8-positive lymphoproliferative disorder	
☐ Mycosis fungoides	
Primary cutaneous CD30-positive T-cell lymphoproliferative disorder: lymphomatoid papulosis	
Primary cutaneous CD30-positive T-cell lymphoproliferative disorder: primary cutaneous anaplastic large cell lymphoma	
Subcutaneous panniculitis-like T-cell lymphoma	
Primary cutaneous gamma/delta T-cell lymphoma	
Primary cutaneous CD8-positive aggressive epidermotropic cytotoxic T-cell lymphoma	
Primary cutaneous peripheral T-cell lymphoma, not otherwise specified	
☐ Intestinal T-cell and NK-cell lymphoid proliferations and lymphomas	
☐ Indolent T-cell lymphoma of the gastrointestinal tract	
\square Indolent NK-cell lymphoproliferative disorder of the gastrointestinal tract	
\square Enteropathy-associated T-cell lymphoma	
☐ Monomorphic epitheliotropic intestinal T-cell lymphoma	
☐ Intestinal T-cell lymphoma not otherwise specified	
☐ Hepatosplenic T-cell lymphoma	
☐ Anaplastic large cell lymphomas	
☐ ALK-positive anaplastic large cell lymphoma	
☐ ALK-negative anaplastic large cell lymphoma	
☐ Breast implant-associated anaplastic large cell lymphoma	



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LYMPHOMAS T-cell non-Hodgkin lymphoma (NHL)

DISEASE continued Sub-Classification: Mature T-cell & NK-cell Neoplasms Nodal T-follicular helper (TFH) lymphomas Nodal TFH cell lymphoma, angioimmunoblastic-type Nodal TFH cell lymphoma, follicular type Nodal TFH cell lymphoma, not otherwise specified Peripheral T-cell lymphoma, not otherwise specified BBV-positive NK/T-cell lymphomas EBV-positive nodal T- and NK-cell lymphoma EBV-positive T- and NK-cell lymphoma Severe mosquito bite allergy Hydroa vacciniforme lymphoproliferative disorder Systemic chronic active EBV disease

Systemic EBV-positive T-cell lymphoma of childhood



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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) Unknown		
Type of transplant: Renal		
☐ Cardiac		
☐ Pulmonary		
Other; specify:		
☐ Unknown		



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LYMPH	IOIVIAS

PREVIOUS THERAPIES (between diagnosis and HCT/CT) Previous therapy lines before the HCT/CT:

☐ No	
☐ Yes:	complete the "Treatment — non-HCT/CT/GT/IST" form
☐ Unkr	nown