Minimum Dataset Development for Hematopoietic Stem Cell Transplantation (HSCT) Recipient Registry in Iran

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SUPPLEMENTARY

		Pre-HSCT								
No	Question	CIBMTR	EBMT	CVR	Accept/ Reject					
	Center ide	entification								
1	Hospital		~	1	Accept					
2	Unit		~	1	Accept					
3	Contact person	~	~	1	Accept					
4	Contact person -Email	~	~	1	Accept					
5	country	~		1	Accept					
6	Center code	~	~	1	Accept					
	General recipi	ent information		1	<u> </u>					
7	Date of this report	~	~	1	Accept					

8	Unique patient number (national code)	~	~	1	Accept
9	First name	~	~	1	Accept
10	Last name	•	~	1	Accept
11	Father's name			1	Accept
12	Date of birth	~	~	1	Accept
13	Sex (at birth)	~	~	1	Accept
14	Insurance type			1	Accept
15	Insurance number			1	Accept
16	Medical record number			0.6	Reject
17	Registry patient number	~	~	0.6	Reject
18	Ethnicity	~		1	Accept
19	Race	~		1	Accept
20	Country of residence	~		1	Accept
21	Phone number-Home / Work	~		1	Accept
22	Phone number-Cell phone	~		1	Accept
23	Email			1	Accept
24	Residence address	~		1	Accept
25	Postal code/Zip code	~		1	Accept
26	Blood type (for allogeneic HCTs only)	~		1	Accept
27	Rh factor (for allogeneic HCTs only)	~		1	Accept
28	Is the recipient participating in a clinical trial? If yes, specify the name of study/trial	~	•	1	Accept
29	Is a subsequent HCT planned as part of the overall treatment protocol or planned multiple (sequential) graft protocol?	•	~	1	Accept
	Pre-transplantation	n			
30	First transplant for this patient?	~	~	1	Accept
31	number of prior HSCTs	~		1	Accept
32	type of prior HSCT	~	~	1	Accept

33	date of prior HSCT	~	~	1	Accept
34	Was the prior HCT performed at a different institution?	~	~	1	Accept
35	Has patient or partner become pregnant after prior transplant? If yes, go to next question	~	~	0.6	Reject
36	Did the pregnancy result in a live birth?	~		0.6	Reject
37	Was the patient or partner pregnant at any time in this reporting period?	~		0.6	Reject
38	Reason: Graft failure/insufficient hematopoietic recovery	~		1	Accept
39	Reason: Persistent primary disease	~		1	Accept
40	Reason: Recurrent primary disease	~		1	Accept
41	Reason: Planned subsequent HCT, per protocol	~		1	Accept
42	Reason: New malignancy (including PTLD and EBV lymphoma)	~		1	Accept
43	Reason: Insufficient chimerism	~		1	Accept
44	Reason: Other	~		1	Accept
45	Date of graft failure/rejection	~		1	Accept
46	Date of relapse	~		1	Accept
47	Date of secondary malignancy	~		1	Accept
48	Specify other reason	~		1	Accept
49	Has the recipient ever had a prior cellular therapy?	~		0.6	Reject
50	Date of the prior cellular therapy	~		0.6	Reject
51	Source for the prior cellular therapy	~		0.6	Reject
	Donor information		<u> </u>	1	1
52	Multiple donors?	~	~	1	Accept
53	Specify number of multiple donors	~	~	1	Accept
	Specify donor type for the	product	<u> </u>	1	_1

geneic, unrelated geneic, related Specify product typow	e	*	1	Accept Accept
Specify product typ	e	~	1	Accept
	1			I
ow			•	
	•	~	1	Accept
	~	~	1	Accept
l blood unit	~	~	1	Accept
uct (go to question 61)	~	~	-1	Reject
	~	~	-1	Reject
ated prior to infusion?	~	~	1	Accept
modified?	~	~	0.2	Reject
Specify the HLA match	type			
neic (monozygotic twin)	~	~	1	Accept
dentical sibling (may include	~	~	1	Accept
natched other relative	~	~	1	Accept
nismatched relative	~	~	1	Accept
ted donor	~	~	1	Accept
LA locus mismatch	~	~	1	Accept
2 HLA loci mismatch	~	~	1	Accept
ationship of the donor to the	~		1	Accept
ween donor and patient (0=n	natch; 1=	one misma	tch;2=2	
		~	1	Accept
		~	1	Accept
		~	1	Accept
		~	1	Accept
	neic (monozygotic twin) dentical sibling (may include matched other relative mismatched relative nted donor ILA locus mismatch 2 HLA loci mismatch lationship of the donor to the	d blood unit duct (go to question 61) luct (go to question 61) luct (go to question 61) lated prior to infusion? modified? Specify the HLA match type neic (monozygotic twin) dentical sibling (may include matched other relative mismatched relative tted donor ILA locus mismatch 2 HLA loci mismatch lationship of the donor to the	d blood unit fuct (go to question 61) v lated prior to infusion? v modified? Specify the HLA match type neic (monozygotic twin) dentical sibling (may include matched other relative mismatched relative vted donor fl.A locus mismatch 2 HLA loci mismatch 2 HLA loci mismatch v ween donor and patient (0=match; 1=one mismatch) ween donor and patient (0=match; 1=one mismatch)	d blood unit duct (go to question 61) v -1 lated prior to infusion? y modified? Specify the HLA match type neic (monozygotic twin) dentical sibling (may include matched other relative nismatched relative tted donor J LA locus mismatch A Lationship of the donor to the lationship of the donor to the ween donor and patient (0=match; 1=one mismatch; 2=2

76	Antigenic HLA-DQB1		✓	1	Accept
77	Antigenic HLA-DPB1		•	1	Accept
78	Allelic HLA-A		~	1	Accept
79	Allelic HLA-B		~	1	Accept
80	Allelic HLA-C		~	1	Accept
81	Allelic HLA-DRB1		~	1	Accept
82	Allelic HLA-DQB1		~	1	Accept
83	Allelic HLA- DPB1		•	1	Accept
84	Was this donor used for any prior HCTs? (for this recipient)	~		1	Accept
85	Code of donor registry/ CB bank	~	~	1	Accept
86	Specify the donor code type (ION, BMDW, NMDP)	~	~	1	Accept
87	Name of donor registry/ CB bank	~	~	1	Accept
88	Donor ID (ID given by the donor registry or the CB bank listed above)	•	•	1	Accept
89	Patient ID (ID given by the donor registry or the CB bank listed above)	~	~	1	Accept
90	Donor date of BIRTH	~	~	1	Accept
91	Donor gender (at birth)	~	~	1	Accept
92	Donor CMV status (for Allogeneic)	~	~	1	Accept
93	Donor blood type	~		1	Accept
94	Donor Rh factor	~		1	Accept
95	Did this donor provide more than one stem cell product	•	•	0.6	Reject
96	Specify number of products infused from this donor	~	•	1	Accept
97	Specify the number of these products intended to achieve hematopoietic engraftment	~		1	Accept
98	What agents were used to mobilize the autologous recipient for this HCT? (Autologous only)	~		1	Accept
99	Name of product (gene therapy recipients)	~		0.2	Reject

	(Autologous only)				
	HSCT				
100	Date of diagnosis for primary disease	~	~	1	Accept
101	What was the primary disease for which the HCT was performed?	~	~	1	Accept
102	What scale was used to determine the recipient's functional status?	•	~	0.6	Reject
103	Performance score	~	~	0.6	Reject
104	Weight	~	~	1	Accept
105	Height	~	~	1	Accept
106	Was there any co-existing disease or organ impairment at time of patient assessment just prior to the preparative regimen?	V	~	1	Accept
107	Comorbidity- Solid tumor, previously present		~	1	Accept
108	Comorbidity- Inflammatory bowel disease	~	~	1	Accept
109	Comorbidity-Rheumatologic	~	~	1	Accept
110	Comorbidity-Infection	~	~	1	Accept
111	Comorbidity-Diabetes	~	~	1	Accept
112	Comorbidity-Renal	~	~	1	Accept
113	Comorbidity- Hepatic	~	~	1	Accept
114	Comorbidity- Arrhythmia	~	~	1	Accept
115	Comorbidity- Cardiac	~	~	1	Accept
116	Comorbidity- Cerebrovascular disease	~	~	1	Accept
117	Comorbidity- Heart valve disease	~	~	1	Accept
118	Comorbidity- Pulmonary	~	~	1	Accept
119	Comorbidity- Obesity	~	~	1	Accept
120	Comorbidity- Peptic ulcer	~	~	1	Accept
121	Comorbidity- Psychiatric disturbance	~	~	1	Accept
122	Comorbidity- invasive fungal infection	~		1	Accept

123	Comorbidity- skin malignancy	~		1	Accept
124	Comorbidity- hematologic malignancy	-		1	Accept
					_
125	Comorbidity- Prior malignancy	•		1	Accept
126	Specify prior malignancy	~		1	Accept
127	Were there any other major clinical abnormalities prior to the preparative regimen?		~	1	Accept
128	Specify the major clinical abnormality		~	1	Accept
129	Was the recipient on dialysis immediately prior to start of preparative regimen?	~		1	Accept
130	Glomerular filtration rate (GFR) before start of the preparative regiment (pediatric only)	~		-1	Reject
131	Does the recipient have known complex congenital heart disease? (Pediatric only)	~		-1	Reject
132	Has the patient been infected with COVID-19 (SARS-CoV-2) based on a positive test result at any time prior to the start of the preparative regimen/infusion?	~		1	Accept
133	Did the patient require hospitalization for management of COVID-19 infection?	~		1	Accept
134	Is there a history of mechanical ventilation	~		1	Accept
135	Source of the stem cells (for Autologous)		~	1	Accept
136	Graft manipulation <i>ex-vivo</i> (other than for RBC removal or volume reduction) (for Autologous)		~	1	Accept
137	Genetic manipulation of the graft (for Autologous)		~	-1	Reject
138	Patient CMV status (for Allogeneic)	~	~	1	Accept
139	Lab value- Serum ferritin	~		1	Accept
140	Lab value- Serum albumin	~		1	Accept
141	Lab value- Platelets	~		1	Accept
142	Were platelets transfused ≤ 7 days before date of test?	~		1	Accept
143	Did the recipient have a prior solid organ transplant? (If yes go to next question)	~		1	Accept
144	Specify organ	~		1	Accept

145	Year of prior solid organ transplant	~		1	Accept
		Ť			•
146	Preparative (conditioning) regimen given?	~	~	1	Accept
147	Classify the recipient's prescribed preparative regimen (Allogeneic only)	•	~	1	Accept
148	Reason for not being myeloablative		~	1	Accept
149	Was irradiation planned as part of the pre-HSCT preparative regimen?	•	~	1	Accept
150	What was the prescribed radiation field?	~	~	1	Accept
151	Total prescribed radiation dose as per protocol	~	~	1	Accept
152	Date started	~		1	Accept
153	Was the radiation fractionated?	~	~	1	Accept
154	Total number of fractions	~	~	1	Accept
Indi	cate total prescribed cumulative dose for preparati	ve regimen	(given befo	ore day ())
155	Ara-C (cytarabine)	~	~	1	Accept
156	ALG, ATG (ALS/ ATS)		~	0.6	Reject
157	Bleomycin		~	1	Accept
158	Busulfan	~	~	0.6	Reject
159	BCNU	~	~	0.6	Reject
160	Bexxar(Tositumomab) (radio labelled MoAB)	~	~	0.6	Reject
161	Bendamustine	~		0.6	Reject
162	CCNU	~	~	0.6	Reject
163	Campath (AntiCD 52)		~	0.6	Reject
164	Carboplatin	~	~	0.6	Reject
165	Cisplatin		~	0.6	Reject
166	Clofarabine	~	~	0.6	Reject
167	Corticosteroids		~	0.6	Reject
168	Cyclophosphamide	~	~	0.6	Reject
169	Daunorubicin		~	1	Accept

170	Doxorubicin (adriamycine)		~	1	Accept
171	Epirubicin		~	0.6	Reject
172	Etoposide (VP16)	~	~	0.6	Reject
173	Fludarabine	~	~	0.6	Reject
174	Gemcitabine	~		0.6	Reject
175	Gemtuzumab		~	0.6	Reject
176	Idarubicin		~	1	Accept
177	Ifosfamide	~	~	0.6	Reject
178	Imatinib mesylate		~	0.6	Reject
179	Melphalan	~	~	0.6	Reject
180	Methylprednisolone (solu-Medrol)	~		0.6	Reject
181	Mitoxantrone		~	0.6	Reject
182	Paclitaxel		~	0.6	Reject
183	Pentostatin	~		0.6	Reject
184	Propylen glycol-free melphalan (Evomela)	~		0.6	Reject
185	Rituximab (mabthera, antiCD20)	~	~	0.6	Reject
186	Teniposide		~	0.6	Reject
187	Thiotepa	~	~	0.6	Reject
188	Treosulphan	~	~	0.6	Reject
189	Zevalin (radiolabelled MoAB)	~	~	0.6	Reject
190	Other radiolabelled MoAB, specify		~	0.6	Reject
191	Other MoAB, specify		~	0.6	Reject
192	Other, specify		~	0.6	Reject
193	Indicate prescribed dose units (mg/m 2 , mg/Kg, mg× h/L, ,CSS)	~		1	Accept
194	Specify administration (Busulfan only) (Oral,IV, both)	~		1	Accept
	GvHD prophylaxi	<u> </u>			

195	Was GVHD prophylaxis planned? (Allografts only)		~	1	Accept
	(If yes go to next question)				
Add	itional drugs/ intervention given in the pre-transplant	t period			
196	ALG (Anti-Lymphocyte Globulin), ALS (Anti-Lymphocyte Serum), ATG (Anti-Thymocyte Globulin, ATS (Anti-Thymocyte Serum) (if yes go to next question otherwise go to question 202)	~	•	1	Accept
197	Specify animal origin (horse, rabbit, other)	~	~	1	Accept
198	Total prescribed dose	~		0.6	Reject
199	Abatacept	•		0.6	Reject
200	Anti CD25 (MoAB in ivo)	~	~	0.6	Reject
201	Bortezomib	~		1	Accept
202	Campath (MoAB in vivo; can be "in the bag")	~	~	0.6	Reject
203	CD 34 enriched (CD34+ selection)	~		0.6	Reject
204	Systemic corticosteroids	~	~	0.6	Reject
205	Cyclosporine	•	~	0.6	Reject
206	Cyclophosphamide	•	~	0.6	Reject
207	Defibrotide	•		0.6	Reject
208	Etanercept		~	0.6	Reject
209	Filgotinib	•		0.6	Reject
210	Tacrolimus (FK 506)	•	~	0.6	Reject
211	Infliximab		~	0.6	Reject
212	KGF	•		0.6	Reject
213	Maraviroc	~		0.6	Reject
214	Methotrexate (MTX) (Amethopterin)	~	~	0.6	Reject
215	Mycophenolate	~	~	0.6	Reject
216	Ruxolotinib	~		0.6	Reject
217	Sirolimus	~		0.6	Reject
218	Tocilizumab	~		0.6	Reject
		1	_i		

219	Ursodiol	~		0.6	Reject
220	Other monoclonal antibody, specify		~	0.6	Reject
221	Other agent (in vivo), specify		~	0.6	Reject
222	Extra-corporeal photopheresis (ECP)	~	~	0.6	Reject
223	Ex-vivo T-cell depletion	~		0.6	Reject
224	Blinded randomized trial	~		0.6	Reject
225	Is additional post-HCT therapy planned?	~		0.6	Reject
	Acute Myelogenous Leukemia/Acute My	eloid Leuken	nia (AMI	7)	<u> </u>
No	Question	CIBMTR	EBMT	CVR	Accept/ Reject
1	AML classification	~	~	1	Accept
AMI	L classification: AML with recurrent genetic abnorm	alities			
2	AML with t (8;21) (q22; q22); RUNX1-RUNX1T1		~	1	Accept
3	AML with inv (16) (p13.1;q22) or t (16;16) (p13.1;q22); CBFB-MYH11		~	1	Accept
4	Acute promyelocytic leukemia with t (15;17) (q22;q12); PML/RARA		~	1	Accept
5	AML with t (9;11) (p22; q23); MLLT3-MLL		~	1	Accept
6	AML with t (6;9) (p23; q24); DEK-NUP214		~	1	Accept
7	AML with inv (3) (q21; q26.2) or t(3;3) (q21;q26.2); RPN1-EVI1		~	1	Accept
8	AML (megakaryoblastic) with t (1;22) (p13;q13); RBM15-MKL1		~	1	Accept
9	AML with myelodysplasia related changes		~	1	Accept
AMI	L classification: AML not otherwise categorized (NO	S)	<u> </u>		<u>l</u>
10	AML with minimal differentiation (FAB M0)		~	0.2	Reject
11	AML without maturation (FAB M1)		~	0.2	Reject
12	AML with maturation (FAB M2)		~	0.6	Reject
13	Acute myelomonocytic leukemia (FAB M4)		~	0.6	Reject

14	Acute monoblastic and monocytic leukemia (FAB M5)		~	0.6	Reject
15	Acute erythroid leukemia (FAB M6)		~	0.6	Reject
16	Acute megakaryoblastic leukemia (FAB M7)		~	0.6	Reject
17	Acute basophilic leukemia		~	0.6	Reject
18	Acute panmyelosis with myelofibrosis		~	0.6	Reject
19	Myeloid sarcoma (Granulocytic sarcoma)		~	0.6	Reject
20	Myeloid proliferations related to down syndrome		~	0.6	Reject
21	Blastic Plasmacytoid Dendritic Cell Neoplasm (BPDCN)		~	1	Accept
22	Therapy related myeloid neoplasia (old "Secondary Acute Leukemia") related to prior treatment but NOT after a previous diagnosis of MDS or MDS/MPN		•	1	Accept
23	Did AML transform from MDS or MPN?	~	~	1	Accept
24	Is the disease (AML) therapy related?	~		1	Accept
25	Did the recipient have a predisposing condition?	~	~	0.6	Reject
26	Is this a donor cell leukemia? (If the patient has received an allograft prior to the diagnosis of acute leukemia)		•	1	Accept
27	predisposing condition: Aplastic anemia		~	0.6	Reject
28	predisposing condition: Fanconi anemia	~	~	0.6	Reject
29	predisposing condition: Bloom syndrome	~	~	0.6	Reject
30	predisposing condition: Down syndrome	~		0.6	Reject
31	predisposing condition: Dyskeratosis congenita	~		0.6	Reject
32	predisposing condition: Unknown		~	0.6	Reject
33	predisposing condition: Other condition	~		0.6	Reject
34	Specify other condition	~		1	Accept
35	Was cytogenetics tested at diagnosis? (Karyotyping or FISH)	~	~	1	Accept
36	Were cytogenetics tested via FISH?	~		0.6	Reject

37	Was cytogenetics tested via karyotyping?	•		1	Accept
38	If cytogenetics test is abnormal:		~	1	Accept
	Is Karyotype complex?				
39	If cytogenetics test is abnormal:		~	1	Accept
	Is Karyotype monosomal?				
40	Specify number of distinct cytogenetics abnormalities	~		1	Accept
41	Transcribe the complete karyotype		~	1	Accept
42	Result of cytogenetics test	~	~	1	Accept
	Specify cytogenetics abnor	malities			
43	-5	•		1	Accept
44	-7	•		1	Accept
45	-17	•		0.6	Reject
46	-18	~		0.6	Reject
47	-X	~		0.6	Reject
48	-Y	~		0.6	Reject
49	+4	~		0.6	Reject
50	+8	~		0.6	Reject
51	+11	~		0.6	Reject
52	+13	~		0.6	Reject
53	+14	~		0.6	Reject
54	+21	~		0.6	Reject
55	+22	~		0.6	Reject
56	t (6;9)	~	~	1	Accept
57	t (8;21)	•	~	1	Accept
58	t (9;22)	•		1	Accept
59	t (15;17) and variants	•	~	1	Accept
60	t (16;16)	~	~	1	Accept

61	t (1;22)		~	1	Accept
62	trisomy 8		~	1	Accept
63	inv (16)	~	~	1	Accept
64	del (3q)/ 3q-	~		1	Accept
65	del (9q)/ 9q-	~		0.6	Reject
66	del (11q)/ 11q-	~		0.6	Reject
67	del (16q)/ 16q-	~		0.6	Reject
68	del (17q)/ 17q-	~		0.6	Reject
69	del (20q)/ 20q-	~		0.6	Reject
70	del (21q)/ 21q-	~		0.6	Reject
71	12p any abnormality	~		1	Accept
72	(11q23) any abnormality		~	1	Accept
73	(11q23) abnormality: t (9;11)	~	~	1	Accept
74	(11q23) abnormality: t (10;11)		~	1	Accept
75	(11q23) abnormality: t (11;19)		~	1	Accept
76	(11q23) abnormality: t (6;11)		~	1	Accept
77	Other (11q23) abnormality		~	1	Accept
78	(3q26) (EVI1) any abnormality		~	0.6	Reject
79	(3q26) abnormality: inv (3)	~	~	1	Accept
80	(3q26) abnormality: t (3;3)	~	~	1	Accept
81	(3q26) abnormality: t (2;3) (p21; q26)		~	1	Accept
82	Other (3q26) abnormality		~	0.6	Reject
83	abn 5 type		~	1	Accept
84	abn 5 type: del (5q)/ 5q-	~	~	1	Accept
85	abn 5 type: add (5q)		~	1	Accept
86	abn 5 type: monosomy 5		~	1	Accept
87	abn 5 type: other abn (5q)		~	1	Accept

88	abn 7 type		~	1	Accept
89	abn 7 type: del (7q)/ 7q-	→	-	1	Accept
90	abn 7 type: monosomy 7		-	1	Accept
	71				•
91	abn 7 type: add (7q)		•	1	Accept
92	abn 7 type: other abn (7q)		~	1	Accept
93	abn (17p)		~	1	Accept
94	Specify other abnormality	~	~	1	Accept
	Molecular market	rs			
95	Were tests for molecular markers performed (e.g., PCR, NGS) at diagnosis?	~	~	0.6	Reject
Selec	et each molecular marker as "Positive", "negative" o	or "Not don	e"		
96	AML1-ETO (RUNX1/RUNXT1)		~	0.6	Reject
97	CBFB-MYH11		~	0.6	Reject
98	PML-RARα		~	0.6	Reject
	MLL-Rearrangement/m	nutation	I		
99	MLLT3 (AF9)-MLL		~	0.6	Reject
100	MLL-PTD (partial tandem duplication)		~	0.6	Reject
101	MLLT4 (AF6)-MLL		~	0.6	Reject
102	ELL-MLL		~	0.6	Reject
103	MLLT1 (ENL)-MLL		~	0.6	Reject
104	MLLT10 (AF10)-MLL		~	0.6	Reject
105	Other MLL-rearrangement		~	0.6	Reject
106	Specify other MLL-rearrangement		-	0.6	Reject
107	DEK-NUP214(CAN)		~	0.6	Reject
108	RPN1-EVI1		-	0.6	Reject
			-		
109	RBM15-MKL1			0.6	Reject
110	NPM1 mutation		~	1	Accept
111	CEBPA mutation	~	~	1	Accept

112	Specify CEBPA mutation	~		1	Accept				
112		-		1	Î				
113	FLT3-TKD			1	Accept				
114	FLT3-ITD (Internal Tandem Duplication)		~	1	Accept				
115	FLT3-ITD mutation	~		1	Accept				
116	FLT3-ITD allelic ratio	•		0.6	Reject				
117	Specify FLT3-ITD allelic ratio	•		0.6	Reject				
118	DNMT3A		~	0.6	Reject				
119	ASXL1		~	0.6	Reject				
120	TP53		~	0.6	Reject				
121	RUNX1		~	0.6	Reject				
122	c-KIT		~	1	Accept				
123	IDH1	~		0.6	Reject				
124	IDH2	~		0.6	Reject				
125	KIT	~		1	Accept				
126	NPM1	~		1	Accept				
127	Another molecular marker	~		0.6	Reject				
128	Specify another molecular marker	~		0.6	Reject				
129	Other		~	0.6	Reject				
130	Specify other		~	0.6	Reject				
131	Did the recipient have central nervous system leukemia at any time prior to the start of the preparative regimen / infusion? (Only CNS leukemia)	~	~	1	Accept				
	Involvement at diagno	osis	•	•					
132	Bone marrow		~	1	Accept				
133	CNS		~	1	Accept				
134	Testis/ovary		~	1	Accept				
135	other		~	1	Accept				
	What was the disease sta	atus?	1	1	_1				

		1	1		1
136	Primary induction failure	•	•	1	Accept
137	1 st Complete remission	~	~	1	Accept
138	2 nd Complete remission	~	~	1	Accept
139	≥ 3 rd Complete remission	~	~	1	Accept
140	1 st relapse	~	~	1	Accept
141	2 nd relapse	~	~	1	Accept
142	≥ 3 rd relapse	~	~	1	Accept
143	No treatment	~		1	Accept
144	How many cycles of inclusion therapy were required to achieve 1 st complete remission?	~		1	Accept
145	Was the recipient in remission by flow cytometry?	~	~	1	Accept
146	Was the recipient in remission by cytogenetics?			1	Accept
147	Was the recipient in remission by molecular?		~	1	Accept
148	Date of last relapse before this HSCT	~	~	0.6	Reject
149	Date assessed	~		0.6	Reject
	Acute Lymphoblastic Leuke	mia (ALL)	1		
150	Specify ALL classification	~		1	Accept
	ALL classification	<u> </u>			. I
151	B lymphoblastic leukemia /lymphoma-with t (9;22) (q34; q11.2); BCR-ABL1		~	1	Accept
152	B lymphoblastic leukemia /lymphoma-with t (v;11q23); MLL rearranged		~	0.6	Reject
153	B lymphoblastic leukemia/lymphoma-with t (1;19) (q23;p13.3); E2A-PBX1		~	0.6	Reject
154	B lymphoblastic leukemia /lymphoma -with t (12;21) (p13;q22); TEL-AML1 (ETV-RUNX1)		~	0.6	Reject
155	B lymphoblastic leukemia /lymphoma-with hyperdiploidy		•	1	Accept
156	B lymphoblastic leukemia /lymphoma-with hypodiploidy		~	1	Accept

157	B lymphoblastic leukemia /lymphoma-with t (5;14) (q31;q32); IL3-IGH		~	0.6	Reject
158	B lymphoblastic leukemia /lymphoma-Not Otherwise Specified (NOS)		~	0.6	Reject
159	Other B lymphoblastic leukemia/lymphoma		~	0.6	Reject
160	T lymphoblastic leukemia/lymphoma		~	1	Accept
161	Did the recipient have a predisposing condition?	~		1	Accept
	Specify condition				1
162	Condition: Aplastic Anemia	~		0.6	Reject
163	Condition: Bloom syndrome	~		0.6	Reject
164	Condition: Down syndrome	~		0.6	Reject
165	Condition: Fanconi anemia	~		1	Accept
166	Condition: other condition	~		0.6	Reject
167	Secondary origin: Related to prior exposure to therapeutic or radiation		~	1	Accept
168	If the patient has received an allograft prior to the diagnosis of acute leukemia: Is this a donor cell leukemia?		•	1	Accept
169	Were tyrosine kinase inhibitors (<i>i.e.</i> , imatinib mesylate) given for pre-HCT therapy at any time prior to the start of the preparative regimen?	~		1	Accept
	Laboratory studies				1
170	Was cytogenetics tested? (Karyotyping or FISH)	~	~	1	Accept
171	Was cytogenetics tested via FISH?	~		1	Accept
172	Was cytogenetics tested <i>via</i> karyotyping?	~		1	Accept
173	Complex karyotype		~	1	Accept
174	Transcribe the complete karyotype		~	1	Accept
175	Results of tests	~	~	1	Accept
176	Specify number of distinct cytogenetics abnormalities	~		1	Accept
	Specify abnormalities	es	l	l .	'

177	-7	~		0.6	Reject
178	+4	~		0.6	Reject
179	+8	~		0.6	Reject
180	+17	~		0.6	Reject
181	+21	~		0.6	Reject
182	t (1;19)	~	~	0.6	Reject
183	t (2;8)	~		0.6	Reject
184	t (4;11)	~	~	0.6	Reject
185	t (5;14)	~		0.6	Reject
186	t (5;14) (q31; q32)		~	0.6	Reject
187	t (8;14)	~		0.6	Reject
188	t (8;22)	~		0.6	Reject
189	t (9;22)	~	~	1	Accept
190	t (10;14)	~		0.6	Reject
191	t (11;14)	~		0.6	Reject
192	t (12;21)	~	~	0.6	Reject
193	del (6q)/ 6q-	~		0.6	Reject
194	del (9p)/9p-	~		0.6	Reject
195	del (12p)/ 12p-	~		0.6	Reject
196	add (14q)	~		0.6	Reject
197	(11q23) any abnormality	~	~	0.6	Reject
198	9p any abnormality	~		0.6	Reject
199	12p any abnormality	~		0.6	Reject
200	Hyperdiploidy (>46)		~	1	Accept
201	Hyperdiploidy (>46):50-66 chromosome		~	1	Accept
202	Hyperdiploidy (>46): Trisomy		~	1	Accept
203	Hyperdiploidy (>46): Specify extra chromosome		~	1	Accept

204	Hyperdiploidy (>46): Other hyperdiploid karyotype		✓	1	Accept
	(if select go to next question otherwise go to question				
	436)				
205	Hyperdiploidy (>46): Number of chromosomes		~	1	Accept
206	Hyperdiploid (>50)	~		1	Accept
207	Hypodiploid (<46)	~	~	1	Accept
208	Hypodiploid (<46): Low hypodiploid, 32-39 chromosomes		~	1	Accept
200	W 1:1/40 W 1 1:10401			1	A .
209	Hypodiploid (<46): Near haploid, 24-31 chromosomes		•	1	Accept
210	Hypodiploid (<46): Monosomy (if select go to next		~	1	Accept
	question otherwise go to question 442)				
211	Hypodiploid (<46): Specify		~	1	Accept
212	Hypodiploid (<46): Other (if select go to next		~	1	Accept
	question otherwise go to question 444)				
213	Hypodiploid (<46): Number of chromosomes		~	1	Accept
214	iAMP21	~		0.6	Reject
215	Trisomy8		~	0.6	Reject
216	Other abnormality	~	~	0.6	Reject
217	Were tests for molecular markers performed (e.g., PCR, NGS) at diagnosis?	~	~	1	Accept
	Specify molecular mar	kers			
218	BCR-ABL molecular product of t (9;22) (q34; q11.2)	~	~	1	Accept
219	MLL-rearrangement/mutation (if not present go to question 502)		~	1	Accept
220	MLL-rearrangement/mutation: AFF1(AF4)-MLL molecular product of t (4;11) (q21;q23)		~	0.6	Reject
221	MLL-rearrangement/mutation: MLLT1(ENL)-MLL molecular product of t (11;19) (q23;p13.3)		~	0.6	Reject
222	MLL-rearrangement/mutation: MLLT3(AF9)-MLL		~	0.6	Reject
	molecular product of t (9;11) (p22;q23)				
223	MLL-rearrangement/mutation: Other MLL-		~	0.6	Reject

rearrang	TV6)-AML1(RUNX1) molecular product of p (p13;q22) H molecular product of translocation t (5;14) 2) PBX1 Molecular product of translocation q23; p13.3) (IKAROS) H1 and FBXW7 f not present go to question c)	~	· · · · · · · · · · · · · · · · · · ·	0.6 0.6 0.6 0.6 0.6 0.6 0.6	Reject Reject Reject Reject Reject Reject Reject					
$t(12;21)$ 226 IL3-IGH (q31;q3; 227 TCF3-P (1;19) (c 228 IKZF1 (229 NOTCH 230 Other (it 231 Specify 232 Primary 233 1^{st} Comp 234 2^{nd} Con 235 $\geq 3^{rd}$ Co 236 1^{st} relaps 237 2^{nd} relaps 238 $\geq 3^{rd}$ relaps	H molecular product of translocation t (5;14) PBX1 Molecular product of translocation q23; p13.3) (IKAROS) H1 and FBXW7 f not present go to question c) other	•	· · · · · · · · · · · · · · · · · · ·	0.6 0.6 0.6 0.6	Reject Reject Reject					
(q31;q32) 227 TCF3-P (1;19) (d 228 IKZF1 (229 NOTCH 230 Other (i) 231 Specify 232 Primary 233 1st Comp 234 2nd Com 235 ≥ 3rd Co 236 1st relaps 237 2nd relap 238 ≥ 3rd relaps	PBX1 Molecular product of translocation q23; p13.3) (IKAROS) H1 and FBXW7 If not present go to question c) other		· · · · · · · · · · · · · · · · · · ·	0.6 0.6 0.6	Reject Reject Reject					
(1;19) (d) 228 IKZF1 (d) 229 NOTCH 230 Other (in) 231 Specify 232 Primary 233 1st Comp 234 2nd Com 235 ≥ 3rd Co 236 1st relaps 237 2nd relap 238 ≥ 3rd relap 238 ≥ 3rd relap	q23; p13.3) (IKAROS) H1 and FBXW7 If not present go to question c) other		· · · · · · · · · · · · · · · · · · ·	0.6 0.6 0.6	Reject Reject					
229 NOTCH 230 Other (if 231 Specify 231 Specify 232 Primary 233 1^{st} Comp 234 2^{nd} Com 235 ≥ 3^{rd} Co 236 1^{st} relaps 237 2^{nd} relap 238 ≥ 3^{rd} relaps	H1 and FBXW7 If not present go to question c) other		· ·	0.6	Reject					
230 Other (incomplete of the content of the conte	f not present go to question c) other		~	0.6						
231 Specify 232 Primary 233 1^{st} Comp 234 2^{nd} Con 235 $\geq 3^{rd}$ Co 236 1^{st} relaps 237 2^{nd} relap 238 $\geq 3^{rd}$ relaps	other				Reject					
232 Primary 233 1^{st} Comp 234 2^{nd} Comp 235 $\geq 3^{rd}$ Comp 236 1^{st} relaps 237 2^{nd} relaps 238 $\geq 3^{rd}$ relaps		~	~	0.6	1					
233 1^{st} Comp 234 2^{nd} Comp 235 ≥ 3^{rd} Comp 236 1^{st} relaps 237 2^{nd} relaps 238 ≥ 3^{rd} relaps	Status at HSCT			0.0	Reject					
233 1^{st} Comp 234 2^{nd} Comp 235 ≥ 3^{rd} Comp 236 1^{st} relaps 237 2^{nd} relaps 238 ≥ 3^{rd} relaps	2 1441415 447 125 0 1		.							
233 1^{st} Comp 234 2^{nd} Comp 235 ≥ 3^{rd} Comp 236 1^{st} relaps 237 2^{nd} relaps 238 ≥ 3^{rd} relaps	What was the disease st	tatus?								
$\begin{array}{c c} 234 & 2^{\text{nd}} \text{ Con} \\ \hline 235 & \geq 3^{\text{rd}} \text{ Co} \\ \hline 236 & 1^{\text{st}} \text{ relaps} \\ \hline 237 & 2^{\text{nd}} \text{ relap} \\ \hline 238 & \geq 3^{\text{rd}} \text{ rel} \\ \hline \end{array}$	induction failure	~	~	1	Accept					
$\begin{array}{c c} 235 & \geq 3^{\text{rd}} \text{ Co} \\ \hline 236 & 1^{\text{st}} \text{ relaps} \\ \hline 237 & 2^{\text{nd}} \text{ relap} \\ \hline 238 & \geq 3^{\text{rd}} \text{ rel} \\ \hline \end{array}$	plete hematological Remission (CR)	~	~	1	Accept					
$\begin{array}{c c} 236 & 1^{\text{st}} \text{ relaps} \\ \hline 237 & 2^{\text{nd}} \text{ relap} \\ \hline 238 & \geq 3^{\text{rd}} \text{ rela} \\ \hline \end{array}$	mplete hematological Remission (CR)	~	~	1	Accept					
237 2 nd relap 238 ≥ 3 rd rela	omplete hematological Remission (CR)	~	~	1	Accept					
$238 \ge 3^{\text{rd}} \text{ rel}$	se	~	~	1	Accept					
	ose	~	~	1	Accept					
239 No treat	apse	~	~	1	Accept					
	tment	~		1	Accept					
	Type of remission									
240 Cytogen	<i>.</i> 1		~	1	Accept					
241 Molecul	netic remission		~	1	Accept					
		1		1	Accept					
243 Was the	netic remission			1	Accept					

244	Date of most recent relapse	~		1	Accept
245	Date assessed	~		1	Accept
246	Date of this HSCT		~	1	Accept
Othe	er acute leukemia				
247	Date of initial diagnosis		~	1	Accept
248	Specify acute leukemia of ambiguous lineage and other myeloid neoplasm classification	~		1	Accept
Class	sification: Acute leukemia of ambiguous lineage				
249	Acute undifferentiated leukemia			1	Accept
250	Mixed phenotype NOS: Mixed phenotype B/myeloid, NOS		~	1	Accept
251	Mixed phenotype NOS: Mixed phenotype T/myeloid, NOS		~	1	Accept
252	Natural Killer (NK)-cell lymphoblastic leukemia/lymphoma		~	1	Accept
253	Other		~	1	Accept
254	Specify other	~	~	1	Accept
255	Secondary origin? Related to prior exposure to therapeutic drugs or radiation		~	1	Accept
256	If the patient has received an allograft prior to the diagnosis of acute leukemia: Is this a donor cell leukemia?		~	1	Accept
	Status at HSCT	1			
	What was the disease	status?			
257	Primary induction failure	~	~	1	Accept
258	1 st Complete hematological Remission (CR)	~	~	1	Accept
259	2 nd Complete hematological Remission (CR)	~	~	1	Accept
260	≥ 3 rd Complete hematological Remission (CR)	~	~	1	Accept
261	1 st relapse	~	~	1	Accept
262	2 nd relapse	~	~	1	Accept

263	≥ 3 rd relapse	~	~	1	Accept					
264	No treatment	~		1	Accept					
Type of remission										
265	Cytogenetic remission		~	1	Accept					
266	Molecular remission		~	1	Accept					
267	Date assessed	~		1	Accept					
	Chronic Myelogenous Leuke	mia (CML)	1	-1	-					
268	Date of initial diagnosis		~	1	Accept					
	CML classification									
269	Translocation (9;22)		~	1	Accept					
270	bcr-abl		~	1	Accept					
271	Was therapy given prior to this HCT?	~	~	1	Accept					
272	Date treatment started		~	1	Accept					
	Treatment pre-HSC	CT								
273	Combination chemotherapy	~		1	Accept					
274	Hydroxyurea (Droxia, Hydrea)	~		1	Accept					
275	Tyrosine kinase inhibitor	~	~	1	Accept					
276	Tyrosine kinase inhibitor: Imatinib mesylate		~	1	Accept					
277	Tyrosine kinase inhibitor: Nilotinib		~	1	Accept					
278	Tyrosine kinase inhibitor: Dasatinib		~	1	Accept					
279	Tyrosine kinase inhibitor: Bosutinib		~	1	Accept					
280	Tyrosine kinase inhibitor: Ponatinib		~	1	Accept					
281	Tyrosine kinase inhibitor: Other TKI (if no go to question 688)		•	1	Accept					
282	Tyrosine kinase inhibitor: specify other TKI		~	1	Accept					
283	Other chemotherapy (if no go to question 690)		~	1	Accept					
284	Specify Other chemotherapy		~	1	Accept					
285	Interferon-α (Intron, Roferon) (include PEG)	~		1	Accept					

286	Other therapy	~		1	Accept					
287	Specify other therapy	~		1	Accept					
Status at HSCT										
288	Date of this HSCT		~	1	Accept					
What was the disease status?										
289	Complete hematologic response (CHR) preceded only by chronic phase	•		1	Accept					
290	Complete Hematologic Response (CHR) preceded by accelerated phase and/or blast phase	~		0.6	Reject					
291	Chronic Phase (CP)	~	~	1	Accept					
292	Accelerated phase	~	~	1	Accept					
293	Blast crisis	~	~	1	Accept					
294	What is the number of Chronic phase (CP)?	~	~	1	Accept					
295	What is the number of Accelerated phase?	~	~	1	Accept					
296	What is the number of Blast crisis?	~	~	1	Accept					
297	Specify level of response	~		1	Accept					
	Type of remission in Chronic	Phase (CP)	1	1						
298	Hematological		~	1	Accept					
299	Cytogenetic		~	1	Accept					
300	Molecular		~	1	Accept					
301	Date assessed	~		1	Accept					
	Chronic Lymphocytic Leuke	mia (CLL)	1	1						
302	Date of initial diagnosis		~	1	Accept					
303	Specify other leukemia classification	~		1	Accept					
304	Chronic Lymphocytic Leukemia (CLL)/small lymphocytic lymphoma		•	1	Accept					
305	Richter's syndrome (if no go question a)		~	1	Accept					
306	Transformed from a previous known CLL?	~	~	1	Accept					
307	Date of original CLL diagnosis		~	1	Accept					

308	What is chromosome analysis?		~	1	Accept		
Specify chromosome analysis							
309	Trisomy 12		~	1	Accept		
310	Del 13q14		~	1	Accept		
311	Del 11q22-23		~	1	Accept		
312	del (17p)		~	1	Accept		
313	Other (if no go to question 721)		~	1	Accept		
314	Specify other		~	1	Accept		
315	Was any 17p abnormality detected?	~		1	Accept		
	Molecular marker	,	•	1	1		
316	TP53 mutations		~	1	Accept		
	Treatment pre-HSC	T					
317	What is treatment pre-HSCT (primary treatment)? (If no go to question 726)		•	1	Accept		
318	Date treatment started		~	0.6	Reject		
319	Specify regimen details		~	0.6	Reject		
320	Date treatment ended		~	0.6	Reject		
	Status at HSCT				1		
	What is status at HSC	CT?					
321	Complete Remission (CR)	~	~	1	Accept		
322	Partial Remission (PR)	~	~	1	Accept		
323	Stable Disease (SD)	~	~	1	Accept		
324	Progressive Disease (PD)	~	~	1	Accept		
325	untreated	~		1	Accept		
326	Untreated relapse		~	1	Accept		
327	Never treated		~	0.2	Reject		
328	Not assessed	~		0.2	Reject		
329	Date assessed	~		1	Accept		

	Chronic Prolymphocytic Leukemia	(PLL an	d other)				
330	Date of initial diagnosis		~	1	Accept		
331	Specify the other leukemia classification	~		1	Accept		
332	Prolymphocytic Leukemia (PLL): B-cell		~	1	Accept		
333	Prolymphocytic Leukemia (PLL): T-cell		~	1	Accept		
334	Hairy cell leukemia		~	1	Accept		
335	Other (if no go to question a)	~	~	1	Accept		
336	Specify other	~	~	1	Accept		
	Specify chromosome analysis	(PLL onl	ly)				
337	inv (14)/ t (14:14) (q11q32)		~	1	Accept		
338	del (14) (q12)		~	1	Accept		
339	t (11:14) (q23; q11)		~	1	Accept		
340	t (7:14) (q35: q32.1)		~	1	Accept		
341	t (X:14) (q35: q11)		~	1	Accept		
342	idic(8) (p11)		~	1	Accept		
343	Other (specify)		~	1	Accept		
344	Specify other		~	1	Accept		
345	Was any 17p abnormality detected?	~		1	Accept		
	Immunophenotyping of T-cells (T	Γ-cell PLI	only)				
346	CD4+		~	1	Accept		
347	CD8+		~	1	Accept		
348	Lymphocyte count		~	1	Accept		
	Status at HSCT						
349	Date of this HSCT		~	1	Accept		
	What was the disease s	tatus?	1		•		
350	Complete Remission (CR)	~	~	1	Accept		
351	Partial Remission (PR)	~	~	1	Accept		

352	Stable Disease (SD)	✓	~	1	Accept
353	Progressive Disease (PD)	~	<u> </u>	1	_
333	Floglessive Disease (FD)	*	· ·	1	Accept
354	Untreated relapse		~	0.6	Reject
355	untreated	~		0.2	Reject
356	Never treated		~	0.2	Reject
357	Not assessed	~		0.2	Reject
358	Date assessed	~		1	Accept
	Lymphomas		L	I	
NO	Question	CIBMTR	EBMT	CVR	Accept/ Reject
	B-cell Non-Hodgkin Lymphor	nas (NHL)			
1	Date of initial diagnosis		~	1	Accept
2	Specify the lymphoma histology (at infusion)	~		1	Accept
3	Specify other lymphoma histology	~		1	Accept
	B-cell neoplasms	ı	1		
4	Splenic marginal zone lymphoma (if select go to question 35)		~	1	Accept
5	Extranodal marginal zone lymphoma of Mucosa Associated Lymphoid Tissue (MALT) (if select go to question 35)		~	1	Accept
6	Nodal marginal zone lymphoma (if select go to question 35)		~	1	Accept
7	Lymphoplasmacytic Lymphoma (LPL): Waldenstrom macroglobulinemia (LPL with monoclonal IgM) (if select go to next question otherwise question 9)		~	1	Accept
8	International Prognostic Scoring System for Waldenström's Macroglobulinemia (ISSWM) (go to question 35)		~	1	Accept
9	Follicular lymphoma (if select go to next question otherwise question 12)		~	1	Accept
10	Grading		~	1	Accept
11	Prognostic score (FLIPI) (go to question 35)		~	1	Accept

12	Primary cutaneous follicle center lymphoma (if select go to question 35)	•	1	Accept
13	Mantle cell lymphoma (if select go to next question otherwise question 16)	•	1	Accept
14	Grading	~	1	Accept
15	Prognostic score (MIPI) (go to question 35)	~	1	Accept
16	Diffuse large B-cell lymphoma (DLBCL), (NOS): T-cell/histiocyte rich large B cell lymphoma	~	1	Accept
17	Diffuse large B-cell lymphoma (DLBCL), (NOS): Primary DLBCL of the CNS	•	1	Accept
18	Diffuse large B-cell lymphoma (DLBCL), (NOS): Primary cutaneous DLBCL, leg type	~	1	Accept
19	Diffuse large B-cell lymphoma (DLBCL), (NOS): EBV positive DLBCL of the elderly	~	1	Accept
20	DLBCL associated with chronic inflammation	~	1	Accept
21	Lymphomatoid granulomatosis	~	1	Accept
22	Primary mediastinal (thymic) large B-cell lymphoma	~	1	Accept
23	Intravascular large B-cell lymphoma	~	1	Accept
24	ALK positive large B-cell lymphoma	~	1	Accept
25	Plasmablastic lymphoma	~	1	Accept
26	Large B-cell lymphoma arising in HHV8-associated multicentric Castleman disease	~	1	Accept
27	Primary Effusion Lymphoma (PEL)	~	1	Accept
28	Burkitt Lymphoma (BL)	~	1	Accept
29	B-cell lymphoma, unclassifiable, with features intermediate between Diffuse Large B-cell Lymphoma and Burkitt Lymphoma (Intermediate DLCBL/BL)	~	1	Accept
30	B-cell lymphoma, unclassifiable, with features intermediate between Diffuse Large B-cell Lymphoma and classical Hodgkin lymphoma (Intermediate DLCBL/HD)	~	1	Accept
31	Other B-cell (if no go to question 32)	~	1	Accept

32	Specify other B-cell		~	1	Accept
33	International Prognostic Index (IPI)		~	1	Accept
34	KI-67 (Proliferation index)		~	1	Accept
35	Is the lymphoma histology reported at transplant a transformation from CLL?	~		1	Accept
36	Was any 17p abnormality detected?	~		1	Accept
37	Transformed from another type of lymphoma? (if yes go to next question otherwise question 41)	~	•	1	Accept
38	Date of original diagnosis	~	~	1	Accept
39	Indicate the type of the original lymphoma	~	~	1	Accept
40	Specify other lymphoma histology	~		1	Accept
Assi	gnment of DLBCL subtype was based on				
41	Immunohistochemistry	~		1	Accept
42	Gene expression profile	~		-1	Reject
43	Unknown method	~		-1	Reject
Sele	cted B-cell Non-Hodgkin Lymphomas (NHL)	l	I		L
Mar	ntle cell lymphoma				
Wal	denstrom macroglobulinaemia				
Bur	kitt lymphoma or ''Intermediate DLBCL/ Burkitt Ly	mphoma''			
44	Date of this HSCT		~	1	Accept
45	Chromosome Analysis at any time before HSCT		~	1	Accept
Indi	cate chromosome analysis as 'present', 'Absent' and '	Not evalu	ated'		
46	del 17p (Mantle cell lymphoma or Waldenstrom macroglobulinaemia only)		~	1	Accept
47	t (2;8) (BL or "Intermediate DLCBL/Burkitt Lymphoma" only)		~	1	Accept
48	t (8;14) (BL or "Intermediate DLCBL/Burkitt Lymphoma" only)		•	1	Accept
49	t (8;22) (BL or "Intermediate DLCBL/Burkitt Lymphoma" only)		~	1	Accept

50	t (14;18) (BL or "Intermediate DLCBL/Burkitt Lymphoma" only)	~	1	Accept
51	myc rearrangement (BL or "Intermediate DLCBL/Burkitt Lymphoma" only)	~	1	Accept
52	BCL-2 rearrangement (BL or "Intermediate DLCBL/Burkitt Lymphoma" only)	•	1	Accept
53	BCL-6 rearrangement (BL or "Intermediate DLCBL/Burkitt Lymphoma" only)	•	1	Accept
54	Immunophenotyping at any time before HSCT done?	~	-1	Reject
	cate Immunophenotype / immunochemistry analysis a uated'	as 'present', 'Abs	ent' and '	Not
55	SOX 11 (Mantle cell lymphoma)	•	-1	Reject
56	MYC (Burkitt Lymphoma or "Intermediate DLCBL/Burkitt Lymphoma")	~	1	Accept
57	BCL-2/IgH (Intermediate DLCBL/Burkitt Lymphoma) BCL-6	~	1	Accept
58	BCL-6 (Intermediate DLCBL/Burkitt Lymphoma)	•	1	Accept
59	Molecular markers analysis (<i>i.e.</i> PCR) at any time before HSCT done?	~	1	Accept
Indi	cate Molecular Markers analysis as 'present', 'Absen	t' and 'Not evalu	ated'	
60	TP53 mutation (Mantle cell lymphoma)	~	1	Accept
61	myc rearrangement (Burkitt Lymphoma or "Intermediate DLCBL/Burkitt Lymphoma)	~	1	Accept
62	BCL-2 rearrangement (Intermediate DLCBL/Burkitt Lymphoma)	~	1	Accept
63	BCL-6 rearrangement (Intermediate DLCBL/Burkitt Lymphoma)	~	1	Accept
	T-cell Non Hodgkin Lymphomas (NHL)	<u> </u>		1
64	Date of initial diagnosis	~	1	Accept

65	Specify the lymphoma histology (at infusion)	~		1	Accept
66	Specify other lymphoma histology	~		1	Accept
Mat	ure T-cell and NK-cell neoplasms classification				
67	T-cell large granular lymphocytic leukemia (if select go to question 91)		~	1	Accept
68	Aggressive NK-cell leukemia		~	1	Accept
69	Systemic EBV positive T-cell lymphoproliferative disease of childhood		~	1	Accept
70	Hydroa vacciniforme-like lymphoma		~	0.6	Reject
71	Adult T-cell leukemia/lymphoma		~	1	Accept
72	Extranodal NK/T-cell lymphoma, nasal type		~	1	Accept
73	Enteropathy-associated T-cell lymphoma		~	1	Accept
74	Hepatosplenic T-cell lymphoma		~	1	Accept
75	Subcutaneous panniculitis-like T-cell lymphoma		~	1	Accept
76	Mycosis Fungoides (MF)		~	1	Accept
77	Sézary syndrome		~	0.6	Reject
78	Lymphomatoid papulosis		~	0.6	Reject
79	Primary cutaneous anaplastic large cell lymphoma		~	1	Accept
80	Primary cutaneous gamma-delta T-cell lymphoma		~	1	Accept
81	Primary cutaneous CD8 positive aggressive epidermotropic cytotoxic T-cell lymphoma		~	0.6	Reject
82	Primary cutaneous CD4 positive small/medium T-cell lymphoma		~	0.6	Reject
83	Peripheral T-Cell Lymphoma NOS (PTCL)		~	1	Accept
84	Angioimmunoblastic T-cell lymphoma		~	1	Accept
85	Anaplastic Large-Cell Lymphoma (ALCL), ALK-positive		~	1	Accept
86	Anaplastic Large-Cell Lymphoma (ALCL), ALK-negative		~	1	Accept
87	Other T-cell		~	1	Accept

88	Specify Other T-cell		~	1	Accept
89	International Prognostic Index (IPI)		~	1	Accept
90	ISCL/EORTC (only for Mycosis fungoides and Sezary syndrome)		~	1	Accept
91	Is the lymphoma histology reported at transplant a transformation from CLL?	~		1	Accept
92	Was any 17p abnormality detected?	~		1	Accept
93	Transformed from another type of lymphoma? (if yes go to next question otherwise question f)	~	~	1	Accept
94	Date of original diagnosis	~	~	1	Accept
95	Indicate the type of the original lymphoma	~	~	1	Accept
96	Specify other lymphoma histology	~		1	Accept
Assi	gnment of DLBCL subtype was based on	1			1
97	Immunohistochemistry	~		1	Accept
98	Gene expression profile	~		-1	Reject
99	Unknown method	~		-1	Reject
	Hodgkin Lymphoma	as			<u> </u>
100	Date of initial diagnosis		~	1	Accept
101	Specify the lymphoma histology (at infusion)	~		1	Accept
102	Classification	~		1	Accept
103	Is the lymphoma histology reported at transplant a transformation from CLL?	~		-0.2	Reject
104	Was any 17p abnormality detected?	~		-0.2	Reject
105	Transformed from another type of lymphoma?	~	~	-0.2	Reject
106	Date of original diagnosis	~	~	-0.2	Reject
107	Indicate the type of the original lymphoma	~	~	-0.2	Reject
Assi	gnment of DLBCL subtype was based on	1	1	l	1
108	Immunohistochemistry	~		1	Accept
109	Gene expression profile	~		-1	Reject
		•			1

110	Unknown method	~	-1	Reject
	All Lymphomas			
111	Treatment pre-HSCT (if yes go to next question)		1	Accept
112	Date of treatment	,	1	Accept
	Drugs given: Antibod	lies		
113	Alemtuzumab (MabCampath) (CD52)		1	Accept
114	Brentuximab (Adcetris) (CD30)	,	1	Accept
115	Obinutuzumab (Gyzeva) (CD20)	,	1	Accept
116	Ofatumumab (Azerra) (CD20)		1	Accept
117	Rituximab (Mabthera) (CD20)		1	Accept
118	other antibody	,	-(0.2 Reject
119	Specify other antibody	,	-0	0.2 Reject
	Drugs given: Radioimmune	otherapy		
120	Bexxar (CD20) (radiolabelled MoAB)		1	Accept
121	Zevalin (CD20) (radiolabelled MoAB)	,	1	Accept
	Drugs given: Specific inh	ibitors		
122	ABT-199 (BCL2-Inhibitor)		1	Accept
123	Crizotinib (ALK-Inhibitor)		1	Accept
124	CC-292 (B cell receptor kinase inhibitor)		1	Accept
125	Ibrutinib (B cell receptor kinase inhibitor)		1	Accept
126	Idelalisib (B cell receptor kinase inhibitor)		1	Accept
127	other inhibitor		0.	.6 Reject
128	Specify other inhibitor		• 0.	.6 Reject
129	Relapse/progression under this drug		1	Accept
	Drugs given: Other	r		
130	Bortezomib (Velcade)		1	Accept
131	Lenalidomide (Revlimid)		1	Accept

132	Other		~	0.6	Reject
133	Specify other		~	1	Accept
134	Date of this HSCT	~	~	1	Accept
135	Number of prior lines of treatment	~	~	1	Accept
Tech	inique used for disease assessment	1			
136	CT scan done	~	~	1	Accept
137	PET	~	~	1	Accept
138	Was the PET (or PET / CT) scan positive for lymphoma involvement at any disease site?	~		1	Accept
139	Date of PET scan	~		1	Accept
140	Deauville (five-point) score of the PET (or PET/CT) scan	~		1	Accept
141	Deauville Scale 1: No uptake or no residual uptake	~		1	Accept
142	Deauville Scale 2: Slight uptake, but below blood pool (mediastinum)	~		1	Accept
143	Deauville Scale3: Uptake above mediastinal, but below or equal to uptake in the liver	~		1	Accept
144	Deauville Scale 4: Uptake slightly to moderately higher than liver	~		1	Accept
145	Deauville Scale 5: Markedly increased uptake or any new lesion	~		1	Accept
	Status at HSCT				
146	What was the disease status?	~	~	1	Accept
147	Never treated		~	1	Accept
148	Complete Remission (CR): Unconfirmed (CRU*)		~	1	Accept
149	Complete Remission (CR): Confirmed		~	1	Accept
150	Partial Response (PR)-(with or without a prior CR)		~	1	Accept
151	Stable disease		~	1	Accept
152	Untreated relapse (from a previous CR) / untreated progression (from a previous PR)		~	1	Accept

153	Chemo refractory relapse or progression, including primary refractory disease		~	1	Accept
154	Disease status unknown		*	1	Accept
155	Was this patient refractory to any line of chemotherapy before this HSCT?		•	1	Accept
156	Number of Complete (CR, CRU) achieved by the patient prior to this HSCT		•	1	Accept
157	Number of Partial remissions (PR) achieved by the patient prior to this HSCT		~	1	Accept
158	Date assessed	~	~	1	Accept