Pre-Transplant Essential Data



CIBMTR Use Only	(Request for OMB approval will be submitted when
Sequence Number:	
	form is complete)
Date Received:	OMB Placeholder OMB No: 0915-0310 Expiration Date: Public Burden Statement: An agency may not conduct or sponsor, and a person is
	not required to respond to, a collection of information unless it displays a currently valid OMB control number. The OMB control number for this project is 0915-0310.
Center Identification	
CIBMTR Center Number:	
EBMT Code (CIC):	·
Hospital:	
Unit: (check only one)	
☐ Adult	
☐ Pediatric	
Recipient Identification	
CIBMTR Research ID (CRID):	

ITR Center Nu	umber:		CIBMTR Research ID:	
Date of birth	:	_ —		
	YYYY	MM	DD	
∏ Male				
_				
icity:				
☐ Hispanic o	r Latino			
☐ Not Hispar	nic or Latino			
☐ Not applica	able (not a reside	ent of the USA)	
Unknown				
):				
□□White				
□□Black or A	African American			
□□Asian				
□□American	Indian or Alaska	Native		
□□Native Ha	waiian or Other I	Pacific Islande	r	
□□Not report	ted			
□□Unknown				
Copy quest	ion 4 to report n	nore than one	e race.	
r postal code f	for place of recip	ient's residenc	e (USA recipients only):	
e recipient part	ticipating in a clin	ical trial?		
7. Study :	Sponsor:			
-	•	uestion 9		
_	•			
_	•			
_	-			
Specify othe	r sponsor:		Go to question 10	
Ctudy	ID Number			
	Date of birth Male Female icity: Hispanic of Not Hispanic of Not applicate Unknown Hispanic of Not applicate Unknown Hispanic of White Hispanic Hispanic of White Hispanic of White Hispanic	Date of birth:	Male Female Female Icity: Hispanic or Latino Not Hispanic or Latino Not applicable (not a resident of the USA) Unknown Unknown	Date of birth:

CIBM	ITR C	enter	Numbe	r:			(CIBM	ΓR Re	seaı	ch II	D:									_
	10.	Subj	ect ID:																		
	Сору	dues (tions 7	'-10 to	report	partici	pation in	mor	e thar	one	e stu	udy.									
Hem	atopo	oietic (Cellula	r Trans	plant (НСТ)															
11.	Date	of thi	с UCT:																		
11.	Dale	5 OI UII	5 HC1.		YYYY				DD												
Was	this th	ne first	HCT fo	or this re	ecipien	i?															
	· []	Yes –	Go to α	questio	n 13																
		No – G	io to q	uestion	15																
			_	-		-	ed as part s HCTs c		e ovei	all tr	eatn	ment	protoc	col (no	ot as	a re	actior	າ to p	ost-H	CT d	lisease
				Go to q																	
			No – G	io to qu	iestion	29															
			14.											_Spec	cify s	ubse	equer	nt HC	:T plar	nned	:
				Autolog	jous –	Go to (question	29													
				Alloger	eic – G	o to q	uestion 2	29													
		15.										S	pecify	the n	umb	er of	prior	НСТ	·s:		_
	Spe	ecify th	ne HSC	Sourc	e(s) fo	r all pr	ior HCTs	:													
																			Autolo	gous	6
		_	Yes																		
			No																		
		17.															Alloge	eneic	, unre	elated	t
			Yes																		
			No																		
		18.															Allo	ogene	eic, re	elated	ł
			Yes																		
			No																		
		19.										,						 	Syng	eneio	
			Yes																		
		П	No																		

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CIBMTR Center	Number:	CI	BMTR Rese	arch ID:		
20.	Date (of the last HCT (ju	st before cu	rrent HCT): _		
			YY	YY	MM	DD
21.			V	Vas the last I	HCT perfor	med at a different institution?
	Yes – Go to questio	n 22				
	No – Go to question	23				
Sp	ecify the institution	that performed t	he last HCT	:		
22	Name:					
	City:	· · · · · · · · · · · · · · · · · · ·				
	State:					
	Country:					
າາ				\\/ha	t was the L	ISC course for the last UCT2
	Autologous			vviia	i was the r	HSC source for the last HCT?
_	Allogeneic, unrelated	donor				
П	Allogeneic, related do					
П	, mogernolo, rolatea av	51101				
24.						Reason for current HCT:
	No hematopoietic red	covery – Go to qu	estion 29			
	Partial hematopoietic	recovery – Go to	question 2	9		
	Graft failure / rejectio	n after achieving	initial hemat	opoietic reco	very – Go	to question 25
	Persistent primary dis	sease – Go to qu	estion 29			
	Recurrent primary dis	sease – Go to qu	estion 26			
	Planned second HC1	, per protocol – G	Go to questi	on 29		
	New malignancy (inc	luding PTLD and	EBV lympho	ma) – Go to	question	27
	Stable, mixed chimer	ism – Go to que s	stion 29			
	Declining chimerism	 Go to question 	29			
	Other – Go to quest	ion 28				
	25Date of gra	ft failure / rejectio	n:	_	_	– Go to question 29
	0	•	YYYY	 MM	DD	<u> </u>
	26	Date of relaps	e:			
		YYYY	ММ	DD		
	27Date of seco	ondary malignanc	y:	_	_	– Go to question 29
		J	YYYY	 MM	DD	

	CIBMTR	Center Number:	CIBMTR Research ID:	
		28		Specify other reason:
	Donor In	formation		
	Multiple d	onors?		
		Yes – Go to question 30		
		No - Go to question 31		
30.	Specify n	umber of donors:		
	To report	more than one donor, copy o	questions 31- 63 and complete for each donor.	
	Specify do	onor:		
		Autologous - Go to question 4	46	
		Autologous cord blood unit - G	o to question 35	
		NMDP unrelated cord blood ur	nit - Go to question 32	
		NMDP unrelated donor - Go to	o question 33	
		Related donor - Go to questio	on 40	
		Related cord blood unit - Go to	o question 35	
		Non-NMDP unrelated donor - 0	Go to question 34	
		Non-NMDP unrelated cord block	od unit - Go to question 35	
32.	NMDP co	rd blood unit ID:	Go to question 46	
33.	NMDP do	nor ID:	Go to question 46	
	34.	Non-NMDP unrelated donor	ID: (not applicable for related donors)	
			Go to questic	on 38
	35.	Non-NMDP cord blood unit II	D: (include related and autologous CBUs)	
	36.	Is the CBU ID also the ISBT	DIN number?	
		☐ Yes – Go to question 38		
		☐ No – Go to question 37		
	37.	Specify the ISBT DIN number	er:	
	38.	Registry or UCB Bank ID:	If 'Other registry' go to 39, otherw	ise go to question 41

	CIBMTR C	enter Num	ber:	CIBMTR Res	earch ID:			
	39.	Specify o	ther Registry or UCB Bank: _				Go to question	41
40.	Specify the	related do	nor type:					
		Synge	neic (monozygotic twin)					
		☐ HLA-id	lentical sibling (may include i	non-monozygotic	twin)			
		HLA-m	natched other relative					
		☐ HLA-m	nismatched relative					
41.	Date of birth	n: (donor /	infant)					
		☐ Known	– Go to question 42					
		Unknow	n – Go to question 43					
		42. Dat	e of birth: (donor / infant)		·	Go to	o question 45	
				YYYY	MM	DD		
	43.	Age: (don	nor / infant)					
		□ K	nown – Go to question 44					
		□ U	Inknown – Go to question 4	! 5				
		44.	Age: (donor / infant)			if less than 1 ye	ear old)	
45.	Sex: (donor	/ infant)						
		☐ Male						
		☐ Female	е					
	Specify pro	oduct type	e:					
	Bone marro	w:						
	☐ Yes	5						
	☐ No							
	PBSC:							
	☐ Yes	5						
	☐ No							
	Single cord	blood unit	:					
	☐ Yes	6						
	☐ No							

CIBMTR Center Number:	CIBMTR Research ID:
Other product:	
☐ Yes – Go to question 50	
□ No – Go to question 51	
50. Specify other product type:	
	usidered a <u>single product</u> when they are all from the same donor and use the ue (and mobilization, if applicable), even if the collections are performed on
Specify number of products infused from	n this donor:
I	
Specify the number of these products in	tended to achieve hematopoietic engraftment:
Questions 53 – 60 are for autologous	HCT recipients only. If other than autologous skip to question 61
Did the recipient have more than one m	obilization event to acquire cells for HCT?
☐ Yes – Go to question 54	
☐ No – Go to question 55	
54. Specify the total number of which collections were used	mobilization events performed for this HCT (regardless of the number of collections of the for this HCT):
Specify all agents used in the mobilize	zation events reported above:
G-CSF	
☐ Yes	
□ No	
GM-CSF	
☐ Yes	
□ No	
Pegylated G-CSF	
☐ Yes	
□ No	
Plerixafor (Mozobil)	
☐ Yes	
□ No	
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CIBMTR Center Number:	CIBMTR Research ID:
Other CXCR4 inhibitor	
☐ Yes	
□ No	
Combined with chemotherapy:	
☐ Yes	
□ No	
Was this donor used for any prior HCTs?	
☐ Yes	
□ No	
_	
Donor CMV-antibodies (IgG or Total) (Allogeneic	HCTs only)
Reactive	
☐ Non-reactive	
☐ Not done	
☐ Not applicable (cord blood unit)	
Was plerixafor (Mozobil) given at any time prior to t	the preparative regimen? (Related HCTs only)
Yes	no proposition regime in (residue no re emy)
□ No	
Unknown	
Consent	
	form for submitting research data to the NMDP / CIBMTR?
Yes (patient consented) – Go to question	
No (patient declined) – Go to question 66	
☐ Not approached – Go to question 66	
65. Date form was signed:	
YYYY	MM DD
Did the recipient give permission to be directly cont	racted for future research?
☐ Yes (patient provided permission) – <i>Go to</i>	
☐ No (patient declined) – Go to question 68	
Not approached - <i>Go to question 68</i>	
□ ivot approactied - Go to question 66	

67. Date form was signed:	
Has the recipient signed an IRB-approved consent form to donate research blood samples to the NMDP / 0	
1 0 11	CIBMTR?
☐ Yes (patient consented) – Go to question 69	
☐ No (patient declined) - Go to question 70	
☐ Not approached - <i>Go to question 70</i>	
Not applicable (center not participating) - Go to question 70	
69. Date form was signed:	
YYYY MM DD	
Has the donor signed an IRB-approved consent form to donate research blood samples to the NMDP / CIE donors only)	3MTR? (Allogene
☐ Yes (donor consented) – Go to question 71	
☐ No (donor declined) - Go to question 72	
☐ Not approached - Go to question 72	
☐ Not applicable (center not participating) - Go to question 72	
71. Date form was signed:	
YYYY MM DD	
Product Processing / Manipulation	
Was the product manipulated prior to infusion?	
∏Yes - <i>Go to questions 73</i>	
□ No - Go to question 91	
73. Specify portion manipulated:	
☐ Entire product	
Portion of product	
Specify all methods used to manipulate the product:	
74. Washed	
74. Washed U Yes No	

CIBMTR Center Number:	CIBMTR Research ID:
☐ Yes	
☐ No	
76	Buffy coat enriched (buffy coat preparation)
☐ Yes	
□ No	
-	
	B-cell reduced
☐ Yes	
□ No	
78.	CD8 reduced
Yes	
□ No	
	Plasma reduced (removal)
☐ Yes	
□ No	
80	RBC reduced
☐ Yes	
□ No	
81.	Cultured (ex-vivo expansion)
☐ No	
82	Genetic manipulation (gene transfer / transduction)
□ Yes	Genetic manipulation (gene transfer / transduction)
□ No	
<u> </u>	
83	PUVA treated
☐ Yes	
□ No	
84	CD34 enriched (CD34+ selection)
Yes	
□ No	
85	CD133 enriched

Yes	CIBMTR Center	Number: CIBMTR Research ID:
No	П	Yes
Yes	_	No
Yes	_	
No	86.	Monocyte enriched
87Mononuclear cells enriched Yes No 88T-cell depletion Yes No		Yes
Yes		No
Yes	87.	Mononuclear cells enriched
88T-cell depletion Yes No	_	
Yes No		No
Yes No	88	T-cell depletion
□ No	_	
	_	
	Ц	
89Other cell manipulation	89.	Other cell manipulation
☐ Yes - Go to question 90		Yes - Go to question 90
☐ No - Go to question 91		No - Go to question 91
90Specify other cell manipulation:		90Specify other cell manipulation:
Clinical Status of Recipient Prior to the Preparative Regimen (Conditioning)	Clinical Status	of Recipient Prior to the Preparative Regimen (Conditioning)
What scale was used to determine the recipient's functional status?	What scale was	used to determine the recipient's functional status?
Karnofsky (recipient age ≥ 16 years) – Go to question 92	☐ Karnof	fsky (recipient age ≥ 16 years) – Go to question 92
Lansky (recipient age < 16 years) – Go to question 93		y (recipient age < 16 years) – <i>Go to question 93</i>
Performance score prior to the preparative regimen:	Performa	nce score prior to the preparative regimen:
92. Karnofsky Scale (recipient age ≥ 16 years):	92. Karn	ofsky Scale (recipient age ≥ 16 years):
☐ 100 Normal; no complaints; no evidence of disease - Go to question 94		100 Normal; no complaints; no evidence of disease - <i>Go to question 94</i>
90 Able to carry on normal activity - <i>Go to question 94</i>		90 Able to carry on normal activity - <i>Go to question 94</i>
80 Normal activity with effort - <i>Go to question 94</i>		80 Normal activity with effort - Go to question 94
70 Cares for self; unable to carry on normal activity or to do active work - <i>Go to question 94</i>		70 Cares for self; unable to carry on normal activity or to do active work - Go to question 94
60 Requires occasional assistance but is able to care for most needs - <i>Go to question 94</i>		60 Requires occasional assistance but is able to care for most needs - Go to question 94
☐ 50 Requires considerable assistance and frequent medical care - Go to question 94		50 Requires considerable assistance and frequent medical care - <i>Go to question 94</i>
40 Disabled; requires special care and assistance - Go to question 94		
30 Severely disabled; hospitalization indicated, although death not imminent - <i>Go to question 94</i> CIBMTR Form 2400 revision 5 (page 11 of 36) Draft 6/19/2016		

CIBMTR Center Nu	mber: CIBMTR Research ID:
_ 20	Very sick; hospitalization necessary - <i>Go to question 94</i>
<u> </u>	Moribund; fatal process progressing rapidly - Go to question 94
_	Scale (recipient age < 16 years):
_	Fully active
_	Minor restriction in physically strenuous play
_	Restricted in strenuous play, tires more easily, otherwise active
_	Both greater restrictions of, and less time spent in, active play
_	Ambulatory up to 50% of time, limited active play with assistance / supervision
_	Considerable assistance required for any active play; fully able to engage in quiet play
_	Able to initiate quiet activities
_	Needs considerable assistance for quiet activity
_	Limited to very passive activity initiated by others (e.g., TV)
∐ 10	Completely disabled, not even passive play
Recipient CMV-anti	bodies (IgG or Total) :
☐ Reactive	
☐ Non-read	tive
□ Not done	
□ Not done	
_	nne
Comorbid Condition	ons
Comorbid Condition	mechanical ventilation?
Comorbid Condition	
Comorbid Condition	
Comorbid Condition Is there a history of Yes	
Comorbid Condition Is there a history of Yes No	
Comorbid Condition Is there a history of Yes No	mechanical ventilation?
Is there a history of Yes No Is there a history of	mechanical ventilation?
Is there a history of Yes No Is there a history of Yes No Service of the se	mechanical ventilation?
Is there a history of Yes No Is there a history of Yes No Yes No Yes No 97. Were there of preparative reserved.	mechanical ventilation? proven invasive fungal infection? Inically significant co-existing diseases or organ impairment at time of patient assessment prior to
Is there a history of Yes No Is there a history of Yes No Ves No 97. Were there of preparative recognitions and the conditions are seen to see the conditions and the conditions are seen to see the conditions are s	mechanical ventilation? proven invasive fungal infection? linically significant co-existing diseases or organ impairment at time of patient assessment prior to egimen? Source: Blood, 2005 Oct 15;106(8):2912-2919
Is there a history of Yes No Is there a history of Yes No Service a history of Yes No Yes No No 97. Were there of preparative re Yes - Go No - Go at 98	mechanical ventilation? proven invasive fungal infection? linically significant co-existing diseases or organ impairment at time of patient assessment prior to egimen? Source: Blood, 2005 Oct 15;106(8):2912-2919 to questions 98
Is there a history of Yes No Is there a history of Yes No Service a history of Yes No Yes No No 97. Were there of preparative re Yes - Go No - Go at 98	mechanical ventilation? proven invasive fungal infection? linically significant co-existing diseases or organ impairment at time of patient assessment prior to egimen? Source: Blood, 2005 Oct 15;106(8):2912-2919 to questions 98 o question 135 _Arrhythmia — For example, any history of atrial fibrillation or flutter, sick sinus syndrome, or cular arrhythmias requiring treatment

CIBMTR Center Number	er: CIBMTR Research ID:
☐ Unkno	own
requiring	ac — Any history of coronary artery disease (one or more vessel-coronary artery stenosis medical treatment, stent, or bypass graft), congestive heart failure, myocardial infarction, OR fraction ≤ 50% on the most recent test
☐ Yes	
□ No	
☐ Unkno	own
	scular disease — Any history of transient ischemic attack, subarachnoid hemorrhage or ascular accident
☐ Yes	
☐ No	
Unkno	own
101. <u>D</u> iabe alone	etes — Requiring treatment with insulin or oral hypoglycemics in the last 4 weeks but not diet
☐ Yes	
☐ No	
☐ Unkno	own
102	Heart valve disease — Except asymptomatic mitral valve prolapse
☐ Yes	
□ No	
☐ Unkno	own
AST/ALT	tic, mild — Chronic hepatitis, bilirubin > upper limit of normal to $1.5 \times$ upper limit of normal, or > upper limit of normal to $2.5 \times$ upper limit of normal at the time of transplant OR any history of B or hepatitis C infection
☐ Yes	
□ No	
☐ Unkno	own
– ·	tic, moderate / severe — Liver cirrhosis, bilirubin > 1.5 \times upper limit of normal, or AST/ALT > 2.5 imit of normal
☐ Yes	
☐ No	
☐ Unkno	own
	tion — For example, documented infection, fever of unknown origin, or pulmonary nodules continuation of antimicrobial treatment after day 0
☐ Yes	5 (page 13 of 36) Draft 6/19/2016

CIBMTR Center	Number: CIBMTR Research ID:
	No
	Unknown
10	6. <u>Inflammatory</u> bowel disease — Any history of Crohn's disease or ulcerative colitis requiring treatment
П	Yes
П	No
	Unknown
4.6	
_	7Obesity — Patients with a body mass index > 35 kg/m² at time of transplant
_	Yes
П	No
Ц	Unknown
10	BPeptic ulcer — Any history of peptic ulcer confirmed by endoscopy and requiring treatment
	Yes
	No
	Unknown
	9Psychiatric disturbance — For example, depression, anxiety, bipolar disorder or schizophrenia quiring psychiatric consult or treatment in the last 4 weeks
	Yes
	No
	Unknown
	$0.P$ ulmonary, moderate — Corrected diffusion capacity of carbon monoxide and/or FEV $_1$ 66-80% or spinea on slight activity at transplant
	Yes
	No
	Unknown
	1.Pulmonary, severe — Corrected diffusion capacity of carbon monoxide and/or FEV ₁ ≤ 65% or dyspne rest or requiring oxygen at transplant
	Yes
	No
	Unknown
	2.Renal, moderate / severe — Serum creatinine > 2 mg/dL or > 177 μmol/L or on dialysis at transplant, R prior renal transplantation
	Yes
	No

CIBMTR Center N	Jumber:
	Jnknown
poly	Rheumatologic — For example, any history of systemic lupus erythmatosis, rheumatoid arthritis, myositis, mixed connective tissue disease, or polymyalgia rheumatica requiring treatment (do NOT ude degenerative joint disease, osteoarthritis)
	Yes
_	No
	Jnknown
	.Solid tumor, prior — Treated at any time point in the patient's past history, excluding non-melanoman cancer, leukemia, lymphoma or multiple myeloma
	Yes – Go to question 115
	No – Go to question 133
	Jnknown – Go to question 133
115. Breas	st cancer
	☐ Yes – Go to question 116
	☐ No – Go to question 117
	116. Year of diagnosis:
117	. Central nervous system (CNS) malignancy (glioblastoma, astrocytoma)
	☐ Yes – Go to question 118
	☐ No – Go to question 119
	118. Year of diagnosis:
119	. Gastrointestinal malignancy (colon, rectum, stomach, pancreas, intestine)
	☐ Yes – Go to question 120
	☐ No – Go to question 121
	120. Year of diagnosis:
121	. Genitourinary malignancy (kidney, bladder, ovary, testicle, genitalia, uterus, cervix)
	☐ Yes – Go to question 122
	☐ No – Go to question 123
	122. Year of diagnosis:
123	. Lung cancer
	☐ Yes – Go to question 124

CIBMTR Center Num	ber: CIBMTR Research ID:	
	☐ No – Go to question 125	
	124. Year of diagnosis:	
125.	Melanoma	
	☐ Yes – Go to question 126	
	☐ No – Go to question 127	
	126. Year of diagnosis:	
127.	Oropharyngeal cancer (tongue, buccal mucosa)	
	☐ Yes - Go to question 128	
	□ No – Go to question 129	
	128. Year of diagnosis:	
129.	Sarcoma	
	☐ Yes - Go to question 130	
	☐ No – Go to question 131	
	130. Year of diagnosis:	
131.	Thyroid cancer	
	☐ Yes - Go to question 132	
	□ No – Go to question 133	
	132. Year of diagnosis:	
133		Other co-morbid condition
☐ Yes	– Go to question 134	
☐ No -	Go to question 135	
Unk	nown – Go to question 135	
1	34	_Specify other co-morbid condition:
primary disea	re a history of malignancy (hematologic or non-meland ase for which this HCT is being performed? O question 136	oma skin cancer) other than the
	question 156	

CIBMTR Center Number:	CIBMTR Research ID:			
Specify which malignancy(ies) occurred:				
136	Acute myeloid leukemia (AML / ANLL)			
☐ Yes – Go to question 137				
☐ No – Go to question 138				
137. Year of diagnosis:				
138	Other leukemia, including ALL			
☐ Yes – Go to questions 139				
139	Year of diagnosis:			
140	Specify leukemia:			
141	Clonal cytogenetic abnormality without leukemia or MDS			
☐ Yes – Go to question 142				
142	Year of diagnosis:			
143. Hodgkin disease				
☐ Yes – Go to question 144				
☐ No – Go to question 145				
144	Year of diagnosis:			
145	Lymphoma or lymphoproliferative disease			
☐ Yes – Go to questions 146				
□ No – Go to question 148				
146	Year of diagnosis:			
147	Was the tumor EBV positive?			
☐ Yes				
□ No				
148	Other skin malignancy (basal cell, squamous)			
☐ Yes – Go to questions 149				
□ No – Go to question 151				

CIBMTR Center Number:	CIBMTR Research ID:
149	Year of diagnosis:
150	Specify other skin malignancy:
151. Myelodysplasia (MDS) / myelop	roliferative (MPN) disorder
☐ Yes – Go to question 152	
☐ No – Go to question 153	
152	Year of diagnosis:
153	Other prior malignancy
☐ Yes – Go to questions 15	
154	Year of diagnosis:
155.	Specify other prior malignancy:
Pre-HCT Preparative Regimen (Condition	ing)
156. Height at initiation of pre-HCT prepar	rative regimen: inches
	centimeters
Actual weight at initiation of pre-HCT prepar	ative regimen: pounds
	kilograms
Was a pre-HCT preparative regimen prescri	bed?
☐ Yes – Go to questions 159	
□ No – Go to question 317	
159	Classify the recipient's prescribed preparative regimen:
☐ Myeloablative	
☐ Non-myeloablative (NST)	
☐ Reduced intensity (RIC)	
160. Date pre-HCT preparative regim	nen began (irradiation or drugs):
	YYYY MM DD

(Use earliest date from questions 164 radiation, or 169 – 316 chemotherapy)

CIBMTR Center Numb	er: CIBMTR Research ID:	
161. Was irradi	ation planned as part of the pre-HCT preparative regimen?	
☐ Yes –	Go to question 162	
□ No -	Go to question 169	
162. V	Vhat was the prescribed radiation field?	
] Total body	
	Total body by tomotherapy	
	Total lymphoid or nodal regions	
	Thoracoabdominal region	
163. To	otal prescribed dose: (dose per fraction x total number of fractions)	
164. Date starte	ed:	
	YYYY MMDD	
165. W	as the radiation fractionated?	
	Yes – Go to questions 166	
	No – Go to question 169	
1	66. Prescribed dose per fraction: Gy	
	☐ cGy	
1	67. Number of days: (include "rest" days)	
1	68. Total number of fractions:	
Indicate the to	tal prescribed cumulative dose for the preparative regimen:	
169. ALG, ALS,	, ATG, ATS	
☐ Yes –	Go to questions 170	
□ No –	Go to question 174	
170. Total prese	cribed dose mg/kg	
17	71Date started: —	
	YYYY MM DD	
17	72	Specify source:
	ATGAM (horse) – Go to question 174	

CIBMTR Center Num	ber:	CIBMTR Resea	arch ID:	
] ATG – Fresenius (rabbit) -	- Go to question 1	174	
	Other - Go to question 1	173		
	173			Specify other source:
174				Anthracycline
	– Go to question 175			-
☐ No -	- Go to question 191			
1	.75			Daunorubicin
Г				
176.	Total prescribed dose		☐ mg/m²	
			☐ mg/kg	
	177	Da	ite started:	
	YY	YYY MM	DD	
1	.78			Doxorubicin (Adriamycin)
] Yes – Go to questions 17	79		
] No – Go to question 181			
179.	Total prescribed dose:		mg/m²	
			☐ mg/kg	
	180	Da	ite started:	
	YYY		DD	
181. Idarubicir	1			
] Yes – Go to questions 18	32		
] No – Go to question 184			
182.	Total prescribed dose		☐ mg/m²	
			☐ mg/kg	
	183	Da	ite started:	
	YYY	YY MM	DD	
1	.84			Rubidazone

CIBMTR C	enter Numbe	er:	CIBMTI	R Research	n ID:	
		Yes – Go to questions 185	i			
		No – Go to question 187				
18	5.	Total prescribed dose _		n	ng/m²	
				□ n	ng/kg	
		186		Date s	started:	
		YYY	Υ	MM	DD	
	18	7				Other anthracycline
		Yes – Go to questions 188	!			
		No – Go to question 191				
18	8.	Total prescribed dose _		n	ng/m²	
				□ m	ng/kg	
		189.		Date	started:	
		YYYY		MM	DD	
		190				Specify other anthracycline:
	191					Bleomycin (BLM, Blenoxane)
	☐ Yes –	Go to questions 192				
	□ No – (Go to question 194				
192.	Total presc	ribed dose		mg/m²		
				☐ mg/kg		
	19	3		Date s	started:	
		YYYY	MM	DD		
194.	Busulfan (N	/lyleran)				
	☐ Yes –	Go to questions 195				
	□ No – (Go to question 198				
195.	Total presc	ribed dose		mg/m²		
				☐ mg/k	кg	
				☐ Targ	et total AUC	(μmol x min/L)
	10	e.		Data	etortod:	
	19	6		Date s	started:	

CIBMTR (Center Number:		CIBMTR R	Research ID:	
		YYYY	ММ	DD	
	197				Specify administration:
	☐ Oral				
	□ IV				
	☐ Both				
198.	Carboplatin				
	Yes – Go to question	ıs 199			
	☐ No – Go to question				
199.	Total prescribed dose		П	mg/m²	
200.				mg/kg	
	200			Date started:	
	•	YYYY	MM	DD	
2	No – Go to qu Specify the target			L/minute	
	203				Cisplatin (Platinol, CDDP)
	Yes – Go to question				,
	☐ No – Go to question	206			
204.	Total prescribed dose			mg/m² mg/kg	
	205	YYYY	MM	Date started:	
	206				Cladribine (2-CdA, Leustatin)
	☐ Yes – Go to question				
	☐ No – Go to question	209			
207.	Total prescribed dose		_ 0	mg/m²	
				mg/kg	
208.	Date started:				

CIBMTR Center Number:		CIBMTR Res	search ID:	
	YYYY	ММ	DD	
209			Corticostero	ids (excluding anti-nausea medication)
☐ Yes – Go to ques	tion 210			
☐ No – Go to quest	ion 223			
210.				Methylprednisolone (Solu-Medrol)
	o questions 211			
□ No – Go to	question 213			
211. Total pr	rescribed dose		_	
ZII. Total pi				
			□ mg/kg	
212		· · · · · · · · · · · · · · · · · · ·	Date started: _	
	YYYY	M	M DD	
213.				Prednisone
	o questions 214			
□ No – Go to	question 216			
214. Total prescribe	ed dose		☐ mg/m²	
2211 Total processo			□ mg/kg	
215			Date started:	
	YYYY	M	M DD	
216				Dexamethasone
☐ Yes – Go to	o questions 217			
□ No – Go to	question 219			
217. Total pr	rescribed dose			
218		· · · · · · · · · · · · · · · · · · ·	Date started: _	
	YYYY	MI	M DD	
219. Other corticosteroid				
☐ Yes – Go to	o questions 220			
□ No – Go to	question 223			
220. Total pr	rescribed dose			

CIBMTR C	enter Number: CIBM	TR Research	ID:	
		1	mg/kg	
	221.	Date s	started:	
	YYYY	MM	DD	
	222			Specify other corticosteroid:
	223			Cyclophosphamide (Cytoxan)
	☐ Yes – Go to questions 224			
	☐ No – Go to question 226			
224.	Total prescribed dose	mg/m²		
		☐ mg/kg		
	225	Date s	started:	
	YYYY M	M DD		
	226			Cytarabine (Ara-C)
	Yes – Go to questions 227			
	☐ No – Go to question 229			
227.	Total prescribed dose	☐ mg/m²		
		☐ mg/kg		
	228	Date s	started:	
	YYYY M	M DD		
	229			Etoposide (VP-16, VePesid)
	☐ Yes – Go to questions 230			
	☐ No – Go to question 232			
230.	Total prescribed dose	mg/m²		
		☐ mg/kg		
	231	Date s	started:	
	YYYY M	M DD		
232.	Fludarabine			
	☐ Yes – Go to questions 233			
	☐ No – Go to question 235			
233. CIBMTR Form	Total prescribed dose	☐ mg/m²		

CIBMT	R Center Number	r:	CIBMTR	Research ID:	
				☐ mg/kg	
	234	•		Date started:	
		YYYY	MM	DD	
	235				Ifosfamide
	☐ Yes – (Go to questions 236			
	□ No – G	o to question 238			
236.	Total prescr	ibed dose		mg/m²	
				☐ mg/kg	
	237	•		Date started:	
		YYYY	MM	DD	
	238				Intrathecal therapy (chemotherapy)
	☐ Yes – (Go to question 239			
	□ No – G	o to question 252			
	239	•			Intrathecal cytarabine (IT Ara-C)
		Yes – Go to questions 24 0)		
		No – Go to question 242			
	240.	Total prescribed dose _			
				☐ mg/kg	
		241		Date started:	
		YYYY		MM DD	
242.	Intrathecal r	nethotrexate (IT MTX)			
		Yes – Go to questions 243	3		
		No – Go to question 245			
	243.	Total prescribed dose _			
				☐ mg/kg	
		244		Date started:	
		YYY	Υ	MM DD	
	245	•			Intrathecal thiotepa
		Yes – Go to auestions 246			

CIBMTR Center Number:			CIBM1	ΓR Resear	rch ID:	
	_ 1	No – Go to questio r	248			
	246.	Total prescribed	dose		mg/m²	
					☐ mg/kg	
		247	· · · · · · · · · · · · · · · · · · ·	Date	e started:	
			YYYY	MM	DD	
	248	·				Other intrathecal drug
		Yes – Go to questio	ns 249			
	_	No – Go to questio n	252			
	249.	Total prescribed	dose		mg/m²	
					☐ mg/kg	
		250		Date	e started:	
			YYYY	MM	DD	
		251	·····			Specify other intrathecal drug:
	252.					Melphalan (L-Pam)
		Go to questions 253				
	□ No – G	o to question 256				
253.	Total prescri	bed dose		☐ mg/r	m²	
				∏ mg/l	кg	
	254	·		Date	e started:	
		YYYY	/ MN	1 C	DD	
255.	Specify adm	inistration:				
		Oral				
		V				
	<u> </u>	Both				
	256					Mitoxantrone (Novantrone)
	☐ Yes – C	Go to questions 257	•			
	□ No – G	o to question 259				
257.	Total prescri	bed dose		∏ mạ	g/m²	
				Пте	a/ka	

CIBMTR Center Number:		CIBMTR Research ID:		
	258		Date started:	
	YYYY	MM	DD	
25	9			Monoclonal antibody
	Yes – Go to question 260			
	No – Go to question 280			
	260			Radio labeled mAb
	Yes – Go to questions 251			
	☐ No – Go to question 267			
	261. component:	•		Total prescribed dose of radioactive
				☐ MBq
	262.		Date started:	
	YYYY			
	Specify radio labeled mAb:			
	263.			Tositumomab (Bexxar)
				,
	□ No			
	264.			Ibritumomab tiuxetan (Zevalin)
				<u> </u>
	□ No			
265.	Other radio labeled mAt)		
	☐ Yes – Go to questio	n 266		
	☐ No – Go to question	267		
	266			Specify other radio labeled mAb:
	267			Alemtuzumab (Campath)
	Yes – Go to questions 268			
	☐ No – Go to question 270			
268.	Total prescribed dose	m	ng/m²	
			mg/kg	

CIBMTR Center Number:		CIBN	MTR Researc	h ID:	
	269		Date	started: _	
		YYYY	MM	DD	
	270.				Rituximab (Rituxan, anti CD20)
	Yes – Go to ques				<u> </u>
	☐ No – Go to quest	ion 273			
271.	Total prescribed dose			2	
			[] mg/kg	
	272		Date	started: _	
		YYYY	MM	DD	
	273	· · · · · · · · · · · · · · · · · · ·			Gemtuzumab (Mylotarg, anti CD33)
	☐ Yes – Go to ques	tions 274			
	☐ No – Go to quest	ion 276			
274.	Total prescribed dose	·		2	
			[] mg/kg	
	275		Date	started: _	
		YYYY	MM	DD	
	276.				Other mAb
	☐ Yes – Go to ques				
	☐ No – Go to quest	ion 280			
277.	Total prescribed dose	·		2	
			[] mg/kg	
278.	Date started: _				_
		YYYY	ММ	DD	
	279				Specify other mAb:
28	0				Nitrosourea
☐ Yes – Go to question 281					
	No – Go to question 29	1			
	281				Carmustine (BCNU)
	☐ Yes – Go to aues			· · · · · · · · · · · · · · · · · · ·	

CIBMTR Center Number:			IBMTR Re	esearch	ID:	
	☐ No – Go to quest	ion 284				
282.	Total prescribe	ed dose		🛮	mg/m²	
				<u> </u>	mg/kg	
	283			_Date st	arted:	
		YYYY	N	ММ	DD	
	284	· · · · · · · · · · · · · · · · · · ·				CCNU (Lomustine)
	☐ Yes – Go to ques	tions 285				
	☐ No – Go to quest	ion 287				
285.	Total prescribe	ed dose		0'	mg/m²	
					mg/kg	
	286.			Date st	arted:	
		YYYY		MM	DD	
	287					Other nitrosourea
	☐ Yes – Go to ques					
	☐ No – Go to quest	ion 291				
288.	Total prescribe	ed dose		П	mg/m²	
	·				mg/kg	
	289			Date st	arted:	
		YYYY		MM	DD	
	290					Specify other nitrosourea:
291. Pacli	taxel (Taxol, Xyotax)					
	Yes – Go to questions 2	292				
_	No – Go to question 29					
292. Total	Total prescribed dose			mg/m²		
202.			_	mg/kg		
	293			Date st	arted:	
		YY	MM	_Date St		
294	l.					Teniposide (VM26)
	Yes – Go to questions 2					: :::::::::::::::::::::::::::::::

CIBMTR C	Center Number:	CIBMTR Research ID:				
	☐ No – <i>Go to question 297</i>					
295.	Total prescribed dose	☐ mg/m²				
		mg/kg				
	296	Date started:				
	YYYY	MM DD				
297.	Thiotepa					
	☐ Yes – Go to questions 298					
	☐ No – <i>Go to question 300</i>					
298.	Total prescribed dose	mg/m²				
		☐ mg/kg				
	299	Date started:				
	YYYY	MM DD				
	300		Treosulfan			
	☐ Yes – Go to questions 301					
	☐ No – Go to question 303					
301.	Total prescribed dose					
		☐ mg/kg				
	302	Date started:				
	YYYY	MM DD				
303.	Tyrosine kinase inhibitors					
	☐ Yes – Go to questions 304					
	☐ No – Go to question 313					
304.	Dasatinib (Sprycel)					
	☐ Yes – Go to questions 305					
30	5. Total prescribed dose					
		☐ mg/kg				
	306. Date started:					
	YYYY	/ MM DD				

CIBMTR Center Number:			CIBMTF	CIBMTR Research ID:			
307.	Imati	nib mesylate (STI571, Gleevec)					
		☐ Yes – Go to questions 30	8				
		☐ No – Go to question 310					
	308.	Total prescribed dose		☐ mg/m² ☐ mg/kg			
		309. Date started:					
310.	Niloti	nib					
		☐ Yes – Go to questions 31.	1				
		☐ No – Go to question 313					
	311.	Total prescribed dose		☐ mg/m² ☐ mg/kg			
		312. Date started:		_			
		YYY		DD			
	313	·			Other drug		
		Yes – Go to questions 314					
		No – Go to question 317					
314.	Total	prescribed dose	_	☐ mg/m² ☐ mg/kg			
		315		Date started:			
		YYYY	ММ	DD			
		316			Specify other drug:		

GVHD Prophylaxis

This section is to be completed for allogeneic HCTs only; autologous HCTs continue with question 344.

7. Was **GVHD** prophylaxis planned / given?

☐ Yes - Go to questions 318

CIBMTR Center Number:	CIBMTR Research ID:
☐ No - Go to question 344	
Specify:	
318. ALG, ALS, ATG, ATS	
☐ Yes – Go to question 3	19
☐ No – Go to question 32	22
319. Total dose:	mg/kg
320	Specify source:
ATGAM (horse) -	- Go to question 322
☐ ATG – Fresenius	(rabbit) – Go to question 3212
☐ Thymoglobulin (ra	abbit) – Go to question 322
Other – Go to qu	estion 321
321	Specify other source:
322	Corticosteroids (systemic)
☐ Yes	
☐ No	
323	Cyclosporine (CSA, Neoral, Sandimmune)
☐ Yes	
□ No	
324	Cyclophosphamide (Cytoxan)
☐ Yes	
☐ No	
325	Extra-corporeal photopheresis (ECP)
☐ Yes	
☐ No	
326.	FK 506 (Tacrolimus, Prograf)
☐ Yes	
□ No	
327. In vivo monoclonal antibody	
Yes – Go to question 3	28
	

CIBMTR Center Number:	CIBMTR Research ID:		
☐ No – Go to question 335			
Specify in vivo monoclonal antibody	:		
328. Alemtuzumab (Campath)			
☐ Yes			
□ No			
329	Anti CD 25 (Zenapax, Daclizumab, AntiTAC)		
☐ Yes – Go to question 330			
□ No – Go to question 331			
330	Specify:		
331	Etanercept (Enbrel)		
☐ Yes			
□ No			
332	Infliximab (Remicade)		
☐ Yes			
□ No			
333. Other in vivo monoclonal antibody			
☐ Yes – Go to question 334			
☐ No – Go to question 335			
334	Specify antibody:		
335	In vivo immunotoxin		
□ No – Go to question 337			
336	Specify immunotoxin:		
337	Methotrexate (MTX) (Amethopterin)		
☐ Yes			
☐ No			
338	Mycophenolate mofetil (MMF) (CellCept)		
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CIBMTR Center Number:	CIBMTR Research ID:	
339		Sirolimus (Rapamycin, Rapamune)
☐ Yes		
□ No		
340		Blinded randomized trial
☐ Yes – Go to ques		
☐ No – Go to quest	ion 342	
341		Specify trial agent:
342.		Other agent
Yes – Go to ques		
☐ No – Go to quest	ion 344	
343		Specify other agent:
Other Toxicity Modifying Regime	n	
Was KGF (palifermin, Kepivance) s Yes No Masked trial	tarted or is there a plan to use it?	
Post-HCT Disease Therapy Plann	ned as of Day 0	
Is this HCT part of a planned multip Yes No	le (sequential) graft / HCT protocol?	
Is additional post-HCT therapy plan	ned?	
☐ Yes - Go to questions 3	47	
☐ No - Go to First Name		
Questions 347 – 357 are optional	for non-U.S. centers	
347		Bortezomib (Velcade)
☐ Yes		
□ No		

CIBMTR Center Number:	CIBMTR Research ID:		
348	Cellular therapy (e.g. DCI, DLI)		
☐ Yes			
☐ No			
349	Dexamethasone		
☐ Yes			
☐ No			
350	Intrathecal therapy (chemotherapy)		
☐ Yes			
□ No			
351	Tyrosine kinase inhibitor (e.g. imatinib mesylate)		
☐ Yes			
☐ No			
352	Lenalidomide (Revlimid)		
☐ Yes			
□ No			
353	Local radiotherapy		
☐ Yes			
☐ No			
354	Rituximab (Rituxan, MabThera)		
☐ Yes			
☐ No			
355	Thalidomide (Thalomid)		
☐ Yes			
☐ No			
356	Other therapy		
☐ Yes – Go to question 357			
☐ No – Go to First Name			
357	Specify other therapy:		

CIBMTR Center Number:	CIBMTR Research ID:
First Name:	
Last Name:	
E-mail address:	
Date:	<u> </u>
YYYY MM DD	