			CUBLET NMDP120
National Marrow Recipient Baseli	Denor Program® ne and	Unrelated 1	Recipient NMDP ID:
Transplant Data	TCC	Arecipient Last Name:	
		Recipient Local ID (option	nal):
	istry Use Only	ODT Today's Date:	TC Code:
Sequence Number:	110	Month	
Date	RMT	A Date of Transplant for white the second secon	
Received:		Product type: D Marrow (Form 120	
b. Zip or postal c c. Country if non-	nce of recipient (for reside ode for place of recipient' resident of USA:0 nt have a U.S. Social Sec 4. Social Security N 5. Why not?	urity Number (or Canadian Social lumber/Social Insurance Number: SSN REAS Canadian) citizen	Insurance Number)? SSNYN
	3 ☐ Other, spec 2 ☐ Female X SEX	ify:	s, check both. RACE2
Caucasian/White		Asian/Pacific Islander	18 Mexican American or Chicano
1 D North Amer	can or European or North Coast of	9 🔲 South Asian 10 🔲 Filipino (Pilipino)	19 D South or Central American 20 D Hispanic, Otherwise not specified
Africa		11 🛛 Hawaiian or Pacific Islan	ider Native American
3 LI White, Othe	rwise not specified	12 🛛 Japanese 13 🗖 Korean	21
Black		14 🔲 Chinese 15 🔲 Southeast Asian	22 🗆 American Indian
🛛 4 🔲 African Ame	rican n parents born in Africa)	16 Asian/Pacific Islander,	23 🛛 Native American,
		Otherwise not specified	Otherwise not specified
5 🛛 African (bot 6 🔲 Caribbean	ntral American	•	
 5 African (bot 6 Caribbean 7 South or Ce 	ntral American rwise not specified	Hispanic 17 D Puerto Rican or Caribbe	Other

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$\frac{1}{2} \text{ conditioning? } BTPRIDR$ $\frac{1}{31 - 40}$ $\frac{1}{50} = 50$
ning? XPS f the recipient is younger than 16 years of age, complete the Lansky Scale.
LANSKY SCALE < 16 yrs

Recipient

COEXO(S) ere there clinically significant coexisting diseases (e.g., diabetes mellitus) or organ impairment within one month prior to unditioning?

Organ Function Prior To Conditioning

Provide values for recipient's liver function just prior to conditioning:

XSGOT_	Date tested: Month Day Year	What is the upper limit of normal for your institution?		
37. AST (SGOT) U/L SGOTAT38.	Month Day Year	39. UILXSGOTULN		
40. ALT (SGPT) U/L SGPTDT 41.		42. UIL Unit of measurement:		
43. Total serum bilirubin • 2 □ µmol/L				
46. LDH		48. UL BILIULUM		
49. Did the recipient have known clinical liver disease (e.	g., hepatitis) at any time prior to cond	itioning? LIVER MAS		
1 □ yes 50. Specify: 2 □ no 51. Date of onset:	Month Day Year	LIVDISDT		
52 ** that was the recipient's serum creatinine prior to co	nditioning?	• *		
XOR CRUM	rement: 2 ☐ µmol/L 53. Date tested:	Month Day Year		
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56. Did the recipient have a history of clinically significant fungal infection at any time prior to conditioning for transplant? FUNGEVER

1 1 J yes	57. Please select organism	58. Date of onset:	59. Select site(s) from list above:	
	from list above:		First	S. A.
Cont. with 60	If other, specify	Month Day Year — FUNOT	Second FUN	5 7

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Recipient -	Recipient Last Name:			
ig for serologica	al evidence of prior viral expos	ure / infection		
 63. Epstein-Barr antibody ¿ 64. Hepatitis B surface and 65. Hepatitis B surface anti 66. Hepatitis C antibody ∱ 67. Hepatitis A antibody ∱ 68. HIV 5 □ confidentia 69. Other, specify <u>DT+P</u> 	bdy (INV) 1 positive PSTBARA 1 positive Wor core antibody HEPE for positive genHEPE for positive genHEPE GEN 1 positive HEPE GEN 1 positive HEPE GEN 1 positive HIN 1 positive DV 1 positive I HIN 1 positive DV 1 positive I HIN 1 positive I Topositive 1 positive I In an isolation room during the peri-train Impositive 71. Please specify: Impositive Impositive I Conventional private room Impositive I Laminar air flow room	> 4 □ Positiv 5 □ HEPA (e pressure room filtered plus positive pres	1
	for serological evidence of prior viral exposure / infection /1 HTCV/ 1 positive 2 negative 3 inconclusive 4 not tested Impaired in the inconclusive 1 positive 2 negative 3 inconclusive 4 not tested Impaired in the inconclusive 1 positive 2 negative 3 inconclusive 4 not tested Impaired in the inconclusive 1 positive 2 negative 3 inconclusive 4 not tested Impaired in the inconclusive 1 positive 2 negative 3 inconclusive 4 not tested Impaired in the inconclusive 1 positive 2 negative 3 inconclusive 4 not tested Impaired in the inconclusive 1 positive 2 negative 3 inconclusive 4 not tested Impaired in the inconclusive 1 positive 2 not tested 1 not tested Impaired in the inconclusive 1 positive 2 ne			
73. Height at initiation of pre74. Weight at initiation of pre75. Was irradiation performed	tioning began:	Year eter without shoes):	cm PRET	KCHT CUNT
2 🗆 no	76. Source of X-ray therapy: 1	Linear accelerator	2□ °Co XRAYSRC	¢.
Cont. with 111	78. What was the radiation field? PiA 1 □ Total body T9. Total dose:	DFIELD	077005	PATE
	81. Was radiation fractionate	ed? <u>RFFRAC</u> 82. Dose per fracti 83. Number of day	N ion: cC rs:RFDAY5	
NMDP Form 120, 520, 620 V7 (8 November 1998 Copyright © 1999 Marrow Donor Program. All rights	1 □ yes 2 □ no -13) 5 National	86. Indicate which of a. Lungs RFSH b. Eyes RFSH c. Liver RFSH	organs were shielded:	res 🗆 no res 🗆 no res 🗆 no res 🗆 no

•



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Recipient NMDP ID:	- Recipient Last Name:	
	105. Was additional radiation given to other s	sites? ADDXXRT
	2 □ no CW5/RRAD 1 □ yes	
	GONIRRAD 107. Was gon. 1 uyes	SPINO
	SPLIRRAD 108. Was sple 1 🗆 yes	2 no Dose:
	DTHIERAD 109. Other site 1 □ yes	e, specify:
	RADDT 110. Date radi	Month Day Year
111. Were drugs given for p	retransplant conditioning? PTXDRGVN	
1 🗆 yes	Total D	row Infusion Date Started Dose (in mg) Month Day Year
Cont. with 126	112. ALG, ALS, ATG, ATS 1 uses 2 uno ALGADOS	
	113. Busulfan (Myleran) 2 1 □ yes 2 □ no Busubos	
	114. Methylprednisilone \$	
	METHMTH 1 □ oral 2 □ IV 3 □ both 115. Prednisone 4	
	115. Prednisone 4 1 □ yes 2 □ no REANING 116. Other corticosteroid 5	
	1 🗆 yes 2 🗆 no orco RDOS	
	117. Cyclophosphamide 6 1 □ yes 2 □ no CASD05	
	118. Cytarabine (Ara-C) ↓ 1 □ yes 2 □ no	
	119. Etoposide (VP-16) 1 □ yes 2 □ no VPN005 120. Melobalao (L-Bam) 0	
	120. Melphalan (L-Pam) 9 1 □ yes 2 □ no MEUPHDOS	MGUPHDT
	1 ⊡ oral 2 ⊡ IV 3 ⊡ both 121. Thiotepa ਾਂ ⊂ਾ ∽α€\$	THIOTOT
	121. Thiotepa to t □ yes 2 □ no THIDTOS 122. Intrathecal methotrexate 11	
	1 □ yes 2 □ no 1 NM 1 103	
	1 □ yes 2 □ no NT100005 124. Monoclonal antibody (} 1 □ yes 2 □ no MON0CO5	
	specify: $$	
	125. Other 'Y 1 □ yes 2 □ no MH2005 specify:	
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	ipient DP ID:		Recipient Last Name:						
ł	ompatibility Tests								
For	each of the following tests in	ndicate whether it was	a basis for ma	tching the c	lonor to the	recipient:			
127.	Class I HLA Serology Mixed Lymphocyte Culture		CLSIHU MLC		natched natched	2 🗆 misma 2 🗆 misma		3 🗆 not 3 🗖 not	
	Restriction Fragment Leng				atched	2 🛛 misma		3 🔲 not	
	Isoelectric Focusing (IEF)		IEF		natched natched	2 🗆 misma 2 🗆 misma		3 □ not 3 □ not	
	Cytotoxic Lymphocyte Pre Helper T Lymphocyte Pred		CTLP HTLP		natched				
	Class I Sequence Specific				atched	2 🗆 misma		3 🗆 not	
	Class II Sequence Specific			ΓI 1 🛛 π	atched	2 🗖 misma	tched	3 🗖 not	done
134.	Other, specify:		************************* *	1 🗆 m	atched	2 🗖 misma	Itched	3 🗖 not	done
Tra	nsplant Maneuver								
	estions 135-158 are for ma				ntinue with	question 159	and compl	ete Form 5	80.
FOI	r cord blood, continue with	question 159 and con		ov.		·			
	Copy donor reference nur				[[ONRE		
136.	Date of receipt of marrow a	at your facility:.	Month	Day	Ye	ar de la Mu	ARRECI	>T	
	Time (24-hour clock) at rec	MARRIECT		• Minute		idard time light savings tir	MARR	ECZN	
13u.	storage temperature during	g transport: STORTE	™ 1 🛛 Refr	igerated at	1–8°C	2 🗖 Room	temperati	Jre	
139.	Nucleated cell count of the processing (uncorrected ce		Bag one:	•		Bag wo:	• 🗌 🗴	NCCBER 10'/ml	
	,	NCCBEF83	Bag three:	•		Bag our:	• 🚬 ×	NCCB 10 ¹ /ml	EFB4
140. _.	Method used to determine	nucleated cell count:		lter counter er, specify: .		2 🗖 Manu	al count		
141.	Total volume of marrow be	fore processing:		•	ml	oubersf)		
142.	Was the marrow manipulat	ed at your facility prior	to transplant?						
	2 no MANEPYN 14	3. Was the marrow ma	anipulated for v	/olume redu	uction only?	1 🗆 yes	2 🗆 no /	MANVR	ONL
	14	4. Was the marrow pla	asma depleted	only?		1 🛛 yes	2 🗆 no /	MANPLA	15
	14	5. Was the marrow ma	anipulated for A	ABO incom	patibility onl	y? 1 □ yes	2 🗆 no /	MANAB	0
	Cont. with 150 14	6. Was the marrow ma	anipulated for (GVHD prop	hylaxis?				
		1 □ yes 2 □ no MANAVt	10 147. Sp	ecify metho	od used:	MANMI	SUM		
			· · ·	Antibod	• •	ment			
				AntibodAntibod	•	lumo			
		Cont. with 150			• •				
	ъ.	· · ·	5	Sheep i	ed blood ce	Il rosetting onl			
				Soybea		sheep red blo	od cell ros	etting	
						eads			
				Antibod	-				
	P Form 120, 520, 620 V7 (11-1 mber 1998 Copyright © 1998 N		1	-		antibody coate	ed plates		
	W Dopor Program All rights re		11	Other, s	pecity:				

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Recipient	- Recipient Last Name:			
	were used: a. anti CD2 ANTI b. anti CD3 c. anti CD4 d. anti CD5 e. anti CD6 f. anti CD7 g. anti CD8 h. anti CD34 i. Other j. No antibodies use 149. What assays were per after processing?	3 1 ges 2 no 4 1 ges 2 no 1 ges 2 no 6 1 ges 2 no 7 1 ges 2 no 7 1 ges 2 no 8 1 ges 2 no 9 1 ges 2 no 3 4 1 ges 2 no 9	specify: number of T-cells left	
		1 Uyes 2 0 no ssay 1 0 yes 2 0 no AS71 0 yes 2 0 no 2 NOASSAYS	LDA ASY specify:	
150. Time (24-hour clock) at start of in	and the second second		tandard time laylight savings time	TXZONE
15 tal volume of marrow infused o	n the day of transplant:	•m.	VOLINFUS	
152. Cell count of infused marrow (und	corrected cell count):	• x 10 [°] /ml	NUCCTINF	.
153. Method used to determine cell co	2 🗖	Coulter counter Manual count Other, specify:	.CTMETH	*****
154. Was a fraction of the collected ma 1 □ yes	arrow cryopreserved for back-	up infusion?		
2 🗆 no 🔰 155. Toti	al volume of cryopreserved m	arrow:	• mi. CRY	ovol
CRYOYN 156. Nuc	cleated cell count of cryoprese	erved marrow:	• x 10°/mi	YONCC
157. Was there any adverse reaction a	ssociated with the infusion?		· · ·	
1 yes	cify:			
)				

as this the first trar	splant for this recipient?					
ı 🗆 yes						
2 🛛 no	160. What was (were) the prior stem cell source(s)?					
FIRSTRY	a. Autologous AUTOLOG					
		RUTBM 1 D yes	2 🗖 no			
	2 🗆 no b. Peripheral blood		2 🗆 no			
	b. Allogeneic, unrelated <u>ALLOGUNR</u>					
	1 🗆 yes — 162. a. Bone marrow A	LUBM 1 🗆 yes	2 🗖 no			
	2 🗆 no b. Peripheral blood (ALUPB 1 U yes	2 🗖 no			
	c. Cord blood	UCB 1 🗆 yes	2 🗖 no			
	c. Allogeneic, related					
	1 U yes 163. a. Bone marrow	1 🛛 yes	2 🗆 no			
		1 🗆 yes	2 🗆 no			
	ALUGREL C. Cord blood	1 🗆 yes	2 🗆 no			
		· ,				
	164. Date of the last transplant (transplant just before cu	rrent transplant):				
	PREORDT					
	Month Day	Year				
	165. Reason for <i>current</i> transplant:					
	1 No engraftment REASONTX					
	2 Partial engraftment					
	3 Graft failure/rejection	,				
	4 Persistent malignancy 5 Recurrent malignancy 6 Other, specify:					
	166. Source of stem cells for current transplant: CEU	SRCE				
	1 🗖 Autologous	-				
	1 Cryopreserved bone marrow	Chi como				
	2 Cryopreserved peripheral blood stem cells	CEUSCTP				
	2 Allogeneic, unrelated					
	1 🔲 Fresh, original donor bone marrow					
	2 Cryopreserved original donor bone marrow					
	3 Fresh, second donor bone marrow					
	4					
	5 Cryopreserved original donor mobilized per					
	6 🖸 Fresh, second donor <i>mobilized</i> peripheral b	lood stem cells				
	8 Non-NMDP cord blood					
	3 Allogeneic, related					
	1 D Bone marrow					
	2 Peripheral blood 3 Cord blood					
		·	·····			
ligned:						
lassa print nome	Person completing form					
JEASE DUDI DROUM						
	_)					

E-mail address: ___

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