	1	Noti	fica	tion	For	m (l	Dav	10	0 re	epor	t)							:טו			/		
Instruction: Where c provided, check $(orall)$	heck boxes	are										re r	adio l	butte	ons 🔘	are		Cen	tre:				
i. Centre Code:	Or Report	ing c	entre	name:							ate o		is Re	port	t (dd/mm/	уууу)):		/	/			
iii. [*] Place of Transpla	nt centre:									iv. N	ame	of	Trans	plar	nt centre:								
1. APBMT Center #:				2. 0	IBMTI	R Cent	er #:							3. E	BMT Code	(CIC)):						
4. Hospital: (autofill)														5. U	Init: (autofill)								
6. Contact person:	a. Name:														Phone:								
(autofill)	c. Fax:													d. E	mail:								
7. Report information:	a. CIBMTR p	atient	(recipi	ent)																			
	identificat b. EBMT pati			it) #:																			
	c. Patient fol	lowing	n natio	nal /			_	.															
	internation					Yes — No	1.	Name	e ot si	tudy / ti	riai:												
	(Defaulted	d as No	0)			Unknow	vn																
SECTION 1 : PATIENT D	DETAILS & DEM	IOGR/	APHICS	3																			
1. Unique Patient Number or Code:							(4	Autofi	ill)	2. Cen Nun	tre S iber:	•	ificatio	n									
3. Name: * (Please print in capital										4. Initia													
letters)																							
5. NRIC :	MyKad/ MyKid:						-			-					Old IC	:							
	Other ID docum	nent No	o:																•				
	Specify docume	ent type	e (if oth	ers):		Army Police	\simeq	Mothe Father			Wor				Birth Certifi		(a)	Others:					
6. Address:	State:	ohor D	arul Ta	kzim		(Makmu	_) Sa	arawak	:			() Wil	ayah	Persek	utuan	Labua	an
			Darul A n Darul				_			idzuan					ul Ehsan Darul Iman			~	•	Persek icable -			aya
		Nelaka	II Daiui	INaiiii			\simeq	au Pir		ayanga					ekutuan Ku	ıala Lu			ι αμμιι	cable -	i orei	JII	
	○ N	_		an Daru	l Khus	us (Sa	bah															
7. Gender •	Male Female		te of Bi /mm/yy		If the	e exact da	/ ate is not	known,	/ please	enter 01/0	7/yyyy	& che	(autof	ill if N	ted/presum MyKad is av	/ailable		9. Aç * (au	•	culate	d)		
10. Ethnic group:	Malay		Bumip	utera S	abah,	specify	:								Other I	Malays	sian, s	specify	:				
	Chinese Indian		Bumip	utera S	arawal	k, spec	ify:								Foreign	ner, sp	ecify	countr	y:				
11. Weight:			(kg)							12. He	ight :	:						(0	cm)				
SECTION 2 : DISEASE								1															
1. Date of initial diagnosis:	/		/			(dd/i	mm/yy	yy)		mplete h disea					ant Disease	e class	sificati	on pag	ges				
2. Primary Disease	Acute Leuk	emias								-	→				te Section 9:	Acute I	Leukei	mias)					
* Diagnosis:	Chronic My			ukemia	(CML)					_	-				te Section 10 Not a CML)	: Chror	піс Мує	elogeno	us Leu	ıkemia (CML)		
	Other Leuk	_			,					-	-				te Section 11	: Othe	r Leuk	emias)					
	Lymphoma	ιS								-	-	(Pi	lease co	omple	te Section 12	? : Lym _l	ohoma	s)					
	Myelodyspl		-							-	→				te Section 13 te Section 14							ative	
	Combined	-		-	oprolife	erative	Syndro	ome (I	MD/M	PS) =	→	Sy	ndrome	(MD/	(MPS))						promei	anvo	
	MyeloprolifePlasma Ce		•		Multiple	e Mvelo	ma				→				te Section 15 te Section 16						Multiple	Myelo	m
	Anaemia	2.00.		g .	··o.u.p.	, o.o	,,,, <u>,</u>			-	→				te Section 17					Ü		Í	
	Hemoglobii		-	. 5:						-	→				te Section 18					-4 Di			
	Breast Can Other Malig			at Diag	nosis					_	→				te Section 19 te Section 20					at Diagn	iosis)		
	Primary Im	-		ncies						-	<u> </u>				te Section 21		-			cies)			
	Inherited D									-	→				te Section 22								
	Platelet and Histiocytic I			ted Disc	orders					-	→				te Section 23 te Section 24					ea Diso	raers)		
	Autoimmun									-	<u> </u>				te Section 25								
	Multiple Sc				D:					-	→				te Section 26				I A	m me	Dic	2)	
	Other Neur Haematolog	-									→				te Section 27 te Section 28			-				7)	
	Rowel Dise	-													te Section 29			_			-/		

BLOOD AND MARROW TRANSPLANT Notification Form (Day 100 report)

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i. Patient Name and	NRIC I	Number:											ii. Centre Code:				
iii. Name of reporting	g centi	re:											_	L			
SECTION 3 : HSCT																	
1. Performance score:	<u> </u>	20	30	40		60		70 (80	90	100)					
2. System:	Ka	rnofsky	Lansky	′													
3. Type of HSCT: *	_	tologous ogeneic	(Kindly procee	ed to item numb	oer 5)												
	L▶ a	. Patient C	MV status:		0	Nega	ative			O Pos	sitive		Not evaluated Unknown				
	b	. Multiple		units)	0	Yes	→ i. N	Num	ber:				0				
4. Donor:	Donor		natch type	Name of do	nor	WM	DA co	nde	Comp	lete numi	ner of mis	matches	Donor Sex	Donor CMV	Ft	hnic gr	roup
(If Multiple donors	ID	IILAI	naton type	registry/CB			DA CO				Jei Oi iiiis	materies	Donor Sex	status		illio gi	oup
is Yes)		Syng (mor	geneic nozygotic						Antiger	nic:			Male	Negative	_	Malay	
		twin)							A:	◎ 0 ◎	1 🔘 2	2 OND	Female	Positive Not		Chines Indian	
		siblir	identical g (may						B:	◎ 0 (1 🔘 2	2 OND		evaluated		Bumip Sabah	
		mon	de non- ozygotic						C:	◎ 0 ◎	1 🔘 2	2 OND		Unknown		specify	
		twin)	-matched						DRB1	: O 0	1 🔘 2	2 OND				Bumip	
		othe	r relative							: 0 0						Saraw	ιak,
		HLA relat	-mismatched ive:						DPB1	: 0 0	1 🔘 2	2 O ND					,.
			gree of allele match:													Other Malays	sian
			1 HLA antigen						Allelic:							specify	
			mismatch						A:	(a) (a)) 1 () 2	O ND				Foreig	ner
			≥ 2 HLA antigen						А. В:	-	-	_				specify	y
			mismatch						Б. С:) 1 () 2) 1 () 2						y.
		Y	elated donor gree of allele							: 0 0	_	_					
		→ mis	match:							: 0 0							
			1 HLA antigen							: 0 0	_	_					
			mismatch ≥ 2 HLA									<u> </u>					
			antigen mismatch													ΑI	DD
5. Source of Stem * Cells:	Вс	one Marrow	Cord B	lood Pe	ripheı	ral Blo	od		Other, sp	pecify:							_
6. Date of this HSCT:		/	/	(dd/mi	m/yyyy	y)										
7. Chronological no. of HSCT for this patient:	① 1	<u> </u>	3	4 🔘 5													
8. Date of most recent previous HSCT:		/	/	(dd/mi	m/yyyy	y)										-
9. Type of most recent previous HSCT:	All	lo	Auto		4												
10. HSCT part of a plann graft protocol?	ned mu	ltiple	Yes	() N	lo												
11. Graft manipulation e (including T-cell depl			Yes	○ N	lo												
(Other than for RBC r																	
12. Preparative (condition given?	oning) r	egimen	Yes		lo (U	sually	Pedia	tric i	Inherited	d Disorder:	s only. Kin	ndly proceed	d to Section	5)			
13. Was this intended to	be		Yes	○ N	lo 🛨	i. Re	ason:		(Ag	e of recipi	ent	Protoc	ol driven				$\overline{}$
myeloablative? (Applicable for allo on	ly)			, specify:					_	morbid co		_	s, specify:				
					_				Pri	ior HSCT							
14. Cell dose :			i. Marrow :			(nucle	ated c	ell d	lose) x 1	0^8/kg	iii. PBSC	;		(CD34 + cells) x	10^6/	kg	-
			ii. Cord		<u> </u>	•			logo) v 1		□ Unk	nown		,		-	

BLOOD AND MARROW TRANSPLANT Notification Form (Day 100 report)

rtotinoution i orini (buy i ob i oporti)		
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provided, check (\c^1) one box only.		

	ce Use only:
ID:	/
<u>.</u>	
Centre:	

i. Patient Name and NRIC Number:	ii. Centre Code:		
iii. Name of reporting centre:			

Anthracycline		Regimen			RAD dose	RAD unit	Total prescribed dose	Unit
3) ALG, ALS, ATG, ATS (before d0) (a) Ves (b) No (c) (c) Ves (c) No (c)	1) TBI		Yes	○ No		⊚ cGy ⊚ Gy		
Anthracycline	2) TLI, TNI, TAI		Yes	○ No		⊚ cGy ⊚ Gy		
4) Anthracycline ii) Daunoublein iii) Davonublein iii) Davonublein iii) Ves No iii) Idarublein iii) Ves No iii) Idarublein iii) Idarublein iii) Idarublein iii) Idarublein iii) Ves No iii) Idarublein iii) Idarubl	3) ALG, ALS, ATG,	ATS (before d0)	⊚ Yes →	○ No				mg/m² mg/kg
ii) Doxorubicin			10	Other, specify:				
ii) Idarubicin	4) Anthracycline	i) Daunorubicin	Yes	○ No				mg/m² mg/kg
S) Bleomycin		ii) Doxorubicin	Yes	○ No				mg/m² mg/kg
6) Busulfan Yes No		iii) Idarubicin	Yes	○ No				mg/m² mg/kg
Oral	5) Bleomycin		Yes	○ No				mg/m² mg/kg
	6) Busulfan			○ No				mg/m² mg/kg
8) Carmustine (BCNU)				Both				
9) Cisplatin	7) Carboplatin		Yes	○ No				mg/m² mg/kg
10 Corticosteroids	8) Carmustine (BCN	NU)	O Yes	○ No				mg/m² mg/kg
11) Cyclophosphamide	9) Cisplatin		O Yes	○ No				mg/m² mg/kg
12) Cytarabine (Ara-C)	10) Corticosteroids		O Yes	○ No				mg/m² mg/kg
13) Etoposide (VP16)	11) Cyclophospham	nide	O Yes	○ No				mg/m² mg/kg
14) Fludarabine	12) Cytarabine (Ara	ı-C)	O Yes	○ No				mg/m² mg/kg
15)	13) Etoposide (VP1	6)	O Yes	○ No				mg/m² mg/kg
16) Imatinib mesylate (Gleevec, Glivec)	14) Fludarabine		Yes	⊚ No				mg/m² mg/kg
17) Lomustine(CCNU	15) Ifosfamide		Yes	○ No				mg/m² mg/kg
18) Melphalan(L-PAM)	16) Imatinib mesyla	te (Gleevec, Glivec)	Yes	○ No				mg/m² mg/kg
19) Mitoxantrone	17) Lomustine(CCN	IU)	Yes	○ No				mg/m² mg/kg
20) Monoclonal antibody(MAb) i) Campath	18) Melphalan(L-PA	AM)	Yes	○ No				mg/m² mg/kg
antibody(MAb) ii) Rituximab (Rituxan, anti-CD20) iii) Gemtuzumab (Mylotarg, anti-CD33) iv) Other,MAb, specify: Yes No 21) Paclitaxel (Taxol , Xyotax) Yes No 22) Thiotepa Yes No Yes No Mo Mg/m² © n 22) Thiotepa Yes No Mo Mg/m² © n mg/m² © n mg/m² © n mg/m² © n 23) Tenoposide (VM26) Yes No Mo Mo Mo Mo Mo Mo Mo Mo Mo	19) Mitoxantrone		Yes	○ No				mg/m² mg/kg
ii) Rituximab (Rituxan, anti-CD20) Ves No No iii) Gemtuzumab (Mylotarg, anti-CD33) iv) Other,MAb, specify: Yes No 21) Paclitaxel (Taxol , Xyotax) Yes No 22) Thiotepa Yes No Yes No 23) Tenoposide (VM26) Yes No 24) Other, specify: Yes No Yes No Yes No Mo Mab ii) Tositumomab(Bexxar) Yes No Yes No Mo Mo Mo Mo Mo Mo Mo Mo Mo		i) Campath	Yes	○ No				mg/m² mg/kg
anti-CD33) Yes	aniibody(Wirib)		Yes	○ No				mg/m² mg/kg
21) Paclitaxel (Taxol , Xyotax) Yes No 22) Thiotepa Yes No 23) Tenoposide (VM26) Yes No 24) Other, specify: Yes No 25) Radiolabeled MAb i) Tositumomab(Bexxar) Yes No Ii) Ibritumomab(Zevalin) Yes No			Yes	○ No				mg/m ² mg/kg
22) Thiotepa Yes No mg/m² or 23) Tenoposide (VM26) Yes No mg/m² or 24) Other, specify: Yes No mg/m² or 25) Radiolabeled MAb i) Tositumomab(Bexxar) Yes No ii) Ibritumomab(Zevalin) Yes No		iv) Other,MAb, specify:	Yes	○ No				mg/m ² mg/kg
23) Tenoposide (VM26) Yes No mg/m² or 24) Other, specify: Yes No mg/m² or 25) Radiolabeled MAb i) Tositumomab(Bexxar) Yes No ii) Ibritumomab(Zevalin) Yes No	21) Paclitaxel (Taxo	ol , Xyotax)		○ No				mg/m² mg/kg
24) Other, specify:	22) Thiotepa		Yes	○ No				mg/m² mg/kg
25) Radiolabeled MAb i) Tositumomab(Bexxar) Yes No	23) Tenoposide (VM	M26)	Yes	○ No				mg/m² mg/kg
MAb ii) Ibritumomab(Zevalin)	24) Other, specify:		Yes	○ No				mg/m² mg/kg
ii) Ibritumomab(Zevalin)		i) Tositumomab(Bexxar)	Yes	○ No				mCi mBq
iii) Other rMab, specify:: Yes No	140.10	ii) Ibritumomab(Zevalin)	Yes	○ No				mCi mBq
		iii) Other rMab, specify::	Yes	○ No				⊚ mCi ⊚ mBq

SECTION 4: PREPARATION REGIMEN

BLOOD AND MARROW TRANSPLANT Notification Form (Day 100 report)

Instruction, Where shock					-		uo di	a bu	ttono (0 0 00		Centre	: -				
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i. Patient Name and NRIC	Number:										ii. Ce	entre Co	ode:				_
iii. Name of reporting cent	tre:																
SECTION 5 : AFTER HSCT																	
1. GvHD prophylaxis given:	Yes →	a) Immunosup	ressive	i) /	ALG, ALS, ATO	2 ATS			<u> </u>	iv) FCB	(evtra-	corporea	al .				Ŧ
(Applicable for Allografts only)	○ No	chemothera		(after d0):			Yes	● No	phot	tophere	sis) :	" (Yes	· (O)	No	
				ii)	Corticosteroio	ls:		Yes	No		06 (Tac raf) :	rolimus,	(Yes		No	
				iii)	Cyclosporine	(CSA):		Yes	● No	vi) Meth	notrexa	te (MTX)	: (Yes	· ()	No	٦
		b) In vivo mono antibody (MA			Anti CD25 (Zer Daclizumab, A			Yes	● No	iv) Inflix	ximab nicade)		(Yes	; (No	Ī
		antibody (Wi	10)	_	Campath :	illi i AC)		Yes	● No	` `			() Yes	. ()	No	-
				iii)	Etanercept (E	nbrel) :	_		No				7`	,			
		c) Mycophenol	ate (MMF, Cello	ept)):					Yes	. () No					-
		d) Sirolimus (R	apamycin, Rap	amı	ıne) :					① Yes) No					-
		e) Other drug,	specify:							① Yes) No					\dashv
	<u>'</u>																그 =
2. Absolute neutrophil count (ANC)	Yes →	a. Date of ANC (dd/mm/yyyy)	recovery:	IΕ	/	/				—							
recovery (engraftment) (Neutrophils >0.5X10 /L):		(**************************************		i. C	Graft Loss:	Yes	. —	→		. (● No			-			
						a.Date		Īſ	1	7, [7,			(dd.	/mm/yy	vvv)	1
						Graf	t Los	s:					<u></u>			,,,,	1
					Autologous recovery:	Yes	3			(● No						
				iii.	Mixed	(Yes					● No						1
					chimera:												_
	No →	a. Date of last a	issessment:		/	/				(dd/mm/	yyyy)						
	Died befo	ore evaluable			Never belo	ow				(Unkr	nown					
3. Absolute platelet recovery: (Platelets > 20 x 10 ³ /L)	Yes →	a. Date of Plate	let recovery :		/	/				(dd/mm/	′уууу)						Ī
		b. Transplant d (3 days trans		20 2	k 10^9/L :			(auto	o calcul	ate)							
	○ No →	a. Date of last a	assessment:		/	/				(dd/mm/	′уууу)						
	Graft Los	s Died	before evaluabl	е	Autologou	s recove	ry		Never	below (Unkr	nown					
4. Acute Graft Versus Host Disease: (Applicable for Allografts only)	i. Maximum O		0 (none)		○ I ○ II	III	ו	v	Prese	nt but grad	de unkn	own 🔘) Not	applica	ble		
(, 444	organ involv	vement:	a. Skin	>	Stage 0 : I												
					Stage 1 : IStage 2 : I							e					1
				- 1	Stage 3 : 0	Generalia	zed ei	rythrod	derma								
					Stage 4:0			/throde		th bullous							!
			b. GIT		Stage 0 : I												
					Stage 1 : IStage 2 : I							/day					į
					Stage 3:1							-					į
				L.	Stage 4: I	Diarrhoe	a Sev	ere at	odomina	al pain witl	h or with	out ileus					
			c. Liver	- 1	Stage 0 : I												
					Stage 1 : IStage 2 : I												
					Stage 3:1												
					Stage 4 : I	Bilirubin :	>255	umol/	L 								
			d. Other	-	i. Upper GIT:		Yes				<u> </u>	 No					آ
			organs		ii. Lungs:		Yes				0						1
					iii. Others:												1
			1	- 1													13

6. Idiopathic Pneumonia Syndrome (IP) :

Yes

No

Yes

5. VOD :

Unknown

No

Unknown

				TRANSPLA se Classifica			For Office Use only:				
Instruction: Where cl			•) are	Centre:			
provided, check ($^{ early}$) o	-										ı
i. Patient Name and N							ii. C	entre Code	:		
iii. Name of reporting		LIDING C	ELL INFLICION								
SECTION 6 : ADDITIONA 1. Cell infusion (CI) :	AL TREATMENT INCL	ODING C	ELL INFUSION								
(not HSCT or autologou stem cell re-infusion)			f first infusion: e the same as date)		/		(dd/mm/yyyy)				
	○ No	b. Type o	f cell(s):	i) Lymphocyte :		Yes No	iv) Fibroblasts	s :	Yes	No)
				ii) Dendritic cells	:	Yes No	v) Other rMab	, specify: :	Yes	No)
				iii) Mesenchymal	:	Yes No					
			ological no. of this patient:								
		d. Indicat		i) Planned/protoco	ol:	Yes No	vi) Treatment	for disease :	Yes (● No	
				ii) Prophylactic :		Yes No	vii) Mixed chii	maerism :	Yes (● No	
				iii) Treatment of G	ivHD :	Yes No	viii) Treatmen		Yes (● No	
				iv) Loss/decrease chimaerism :	d	Yes No	ix) Other, spe		(Yes (€ No	
				v) Treatment PTLI		Yes No			103(<i>)</i> 140	
		e Numb	er of infusions	EBV lymphoma							
			10 weeks:	(co	unt only ii	infusions that are part	of same regimen a	and given for the	e same indic	ration)	
2. Disease treatment:	Yes, Plan	ned (nlar	nned before HSCT)	◯ Ves N	ot planne	ed (for relapse/pro	naression or nei	reietent diseas	:e) (1		
(apart from cell infusion	103,1141	- (pian	——————————————————————————————————————	163, 14	ot planin	ed (for relapse/pro	ogression or per	Sisterit diseas	100		
SECTION 7 : MALIGNAN	IT DISEASE EVALUA	TION FOR	THIS HSCT	(Non-malignar	nt diseas	se skip disease eva	lluation)				
1. Best disease status (response) after	Continued comp	lete remis:	sion (CR)	CR achieved	● Ne	ever in CR		Not eva	luated		
HSCT (prior to treatment	i. Date CR			·	1 L [i. Date assessed :					7
modification in response to a post HSCT disease	achieved: (dd/mm/yyyy)		//	,	_	(dd/mm/yyyy)		/ /			
assessment)					<u> </u>						
2. First relapse or progression after	○ No ○	Yes	If yes, tick all me	ethods used for assess	sment wi	ith the dates on wh	ich they were us	sed and the re	sults.		
HSCT (Any type, not persistent	L,		e/progression ed by clinical	No −	► i.	Date assessed :					
disease)		/haema metho	ntological			(dd/mm/yyyy)		/			
		motino		Yes	► ^{i.}	Date first seen : (dd/mm/yyyy)	/	/			
				Not evaluated					" "		_
			se/progression ed by cytogenetic	(No -	▶ i.	Date assessed :					
		metho		0		(dd/mm/yyyy)		//			
				Yes	► i.	Date first seen : (dd/mm/yyyy)					
				Not evaluated		(+					
			se/progression ted by molecular	○ No —	▶ i.	Date assessed :					T
		metho				(dd/mm/yyyy)					
				Yes	► "	Date first seen : (dd/mm/yyyy)	/	/			
				Not evaluated							
3. Method of latest status disease	i. Was disease stat		(No -	a. Last date							
assessment	detected by clinic haematological	:/	Yes ——	assessed (dd/mm/yyyy)		/	/				
(record the most recent status	method?		Not evaluated								
and date for each method, depending	ii. Was disease sta detected by cyto		No	a. Last date as	ssessed						
on the disease)	/ FISH method?			(dd/mm/yyyy)			//	/			
*In some circumstances,				b. Considered relapse/prog) No	Yes			
disease may be detected by molecular			Not evaluated							_	
or cytogenetic testing, but may not	iii. Was disease sta		○ No	a. Last date as	ssessed]	
be considered a relapse or	detected by mol method?	ecular		(dd/mm/yyyy)			/	/			
progression. It should still be			(Yes	b. Considered		-) No	Yes			
reported.	Not evaluated	relapse/prog	yı essior	-							

	BLOOD Notification		RROW TR				For Offi ID:	ce Use only:
Instruction: Where c provided, check $()$ of	heck boxes 🔳 are pi	•	•			outtons 🔵 ar	e Centre:	
i. Patient Name and							ii. Centre Co	de:
iii. Name of reporting	g centre:							
SECTION 8: PATIENT S	TATUS AT LAST CONTA	СТ						
1. Survival Status:	Alive Dead Died before HSCT Patient lost to follow up	a. Main Cause of Death: (only one main cause	HSCT F	elated Caus n	Re Pu	se	unction Infec	iac Toxicity tion o occlusive disorder
2. Date of last contact: (Date of last follow up or death)	//		(dd/mm/yyyy)					
SECTION 9 : ACUTE LE	UKEMIAS							
1. Classification:	AML with recurrent ger AML with t(8;21)(q2 AML with abnormal t(16;16)(p13;q22) C AML with t(15;17)(c Acute Lymphoblastic L Precursor B-cell AL t(9;22)(a34;q11); B t(v;11q23); MLL rea t(1;19)(q23;p13) E2 Other Acute Leukemia: Acute undifferentia Biphenotypic, biline AML not otherwise cate AML, mimimally dif AML without matur. AML with maturatic Acute myelomonoc Acute erythroid leul Acute megakaryob	bone marrow ec BFβ/MYH11) 22;q12), (PML/F 22;q12), (PML/F 22;q12), (PML/F 22;q12), (PML/F 24;q12), (PML/F 25; (PML/F 26;q12), (PML/F	ETO) posinophils and inv RARα) and variar	nts (FAB M3		AML with MPS/MD t(12;21)(p Precurso ALL not of the state	sophilic leukemia	asia (w/o MDS or
2. Secondary origin:	Yes: Disease relate	d to prior exposu	re to therapeutic	drugs or rad	diation	(No	(Unknown	i
3. Status at HSCT	Primary induction fa Complete haematol	ilure	· ₁	i. NUMBI	ER ete only for CR or	① 1st	② 2nd	3rd or higher
	Never treated			(comple	f remission ete only for ete remission)	a. Cytogenetic b. Molecular	No Yes No Yes	Not evaluated Unknown Not evaluated Unknown
SECTION 10 · CHRONIC	MYELOGENOUS LEUK	FMIA (CML) No	te: CMML is not	a CMI			-	
1. Classification:	i. Translocation (9;22):	Absent	Prese		Not evaluated			
At least one investigation must be positive	ii. bcr-abl:	Absent	Prese		Not evaluated			
2. Status at HSCT	Chronic phase (CP) Accelerated phase Blast crisis		i. NUMBER ii. Presence and CR (For chronic pl		a. Haematological b. Cytogenetic (t[9;22)) c. Molecular (bcr-	○ No ○ Ye	es Not evalu	ated Unknown

Instruction: Where ca			•	Centre:
provided, check (√) of i. Patient Name and I	·			ii. Centre Code:
iii. Name of reporting				ii. Gentie Gode.
SECTION 11 : OTHER L			_	
1. Classification : 2. Status at HSCT	Chronic lymphocytic leukemia (CLL) Prolynphocyctic Leukemia, B-cell Prolynphocyctic Leukemia, T-cell Hairy Cell Leukemia Other leukemia, specify:			
2. 0.0.00	Stable disease/No response Partial remission (P	R)	Relapse	Never treated
	Complete remission (CR) nodular Partial remi	ssion (nPR	R) Progression	
SECTION 12 : LYMPHO!	MAS			
1. Classification :				
	Non-Hodgkin's lymphoma (NHL) B-cell Neoplasms: Follicular lymphoma Mantle cell lymphoma Extranodal marginal zone of MALT type Diffuse large B-cell lymphoma (If known indicate subtype) Burkitt's lymphoma/Burkitt cell leukemia (ALL L3) Lymphoplasmacytic lymphoma Waldenstrom macroglobulinaemia Splenic marginal zone B-cell lymphoma Nodal marginal zone B-cell lymphoma Primary CNS lymphoma Other B-cell, specify: Nodular lymphocyte predominant Lymphoma depleted Lymphoma depleted Lymphocyte rich T-cell NK-cell Neoplasms: Angioimmunoblastic (AILD) Peripheral T-cell lymphoma (all variants) Anaplastic large-cell, T/null cell, primary cutaneous Anaplastic large-cell, T/null cell, primary systemic Extranodal NK/T-cell lymphoma, nasal type	Adult Aggre Large	Grade I Grade II Intravascular large cell lymp Mediastinal large cell lymph Primary effusion large cell ly High grade B-cell lymphoma	homa oma /mphoma a, Burkitt- mphoma /1+)
	Enteropathy-type T-cell lymphoma		ry syndrome	
	Hepatosplenic gamma-delta T-cell lymphoma	_		
2. Status at HSCT	 Never treated Primary refractory Complete remission, confirmed (CR) Complete remission, unconfirmed (CRU) 1st Partial response (PR1) Partial response>1 (never in CR) (PR>1) Relapse 	→	(complete only for CR, PR>1 or relapse)	1st 2nd 3rd or higher
	Progression		ii. Sensitivity to chemotherapy vsensit	Sensitive Untreated
	*CRU – complete response with persistent scan abnormalities of unknown sig	jnificance	(complete only for relapse)	Resistant Unknown

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b. JMML:

Untreated (Supportive care or treatment

Minimal response (MR)

Partial response (PR)

Progression (PD)

without chemotherapy)

Stable disease (SD)

Complete response (CR)

	Notification Fo	rm (L	Jisease Ciassi	neation)		Control
Instruction: Where control provided, check ($^{\lor}$) of	heck boxes 🔳 are provided one box only.	l, check	(√) one or more boxe	es. Where radio b	uttons 🔵 are	Centre:
i. Patient Name and I	NRIC Number:				_	ii. Centre Code:
iii. Name of reporting	centre:					
SECTION 15 : MYELOPE	ROLIFERATIVE SYNDROMES					
Classification at diagnosis	Chronic idiopathic myelofibromyelofibrosis, fibrosis with more polycythemia vera Essential or primary thrombomyer eosinophilic syndromyelofic.	nyeloid me	etaplasia)	Chronic ne	therwise specified	
2. Secondary origin: (other than transformed to AML)	Yes: Disease related to prior radiation	rexposure	e to therapeutic drugs or	○ No		Unknown
3. Classification at HSCT :	Chronic idiopathic myelofibro with myeloid metaplasia) Polycythemia vera Essential or primary thromb Hyper eosinophilic syndrom Chronic eosinophilic leukaei Chronic neutrophilic leukaei Stem cell leukemia-Lympho	ocythemia e (HES) mia (CEL) mia	i.	i. Date trans (dd/m	of sformation: m/yyyy) therwise specified	
4. Status at HSCT	Treated with chemotherapy: Primary refractory phase (no change) Complete remission (CR) Improvement but no CR Relapse (after CR) Progression/worse	→	i. NUMBER (complete for CR or relapse)	1st 2nd 3rd or higher		
SECTION 16 : PLASMA	Untreated (Supportive care treatment without chemothe	rapy)	MYELOMA			
1. Classification:	Multiple myeloma IgG	<u></u>				
	Multiple myeloma IgA		i. Light chain type:	Карра	(Lambda
	Multiple myeloma IgD	-	ii. Salmon & durie	○ I	And) A
	0		stage at diagnosis:	○ II) B
	Multiple myeloma lgE Multiple myeloma lgM (not Waldenstrom)	→	(Multiple Myeloma only) iii. I.S.S:	a.i. Serum β2 - microglobulin:		a.ii. Units: μg/dL
	 Multiple myeloma- light chain only 					
	Multiple myeloma-non- secretory			b.i. Serum albumin: c. Stage	d. β2 -mic	b.ii. Units: g/dL g/L e. S.albumin
	Plasma cell leukemia			_	u. pz -iiiiu	c. O.dibdiiiii
	Solitary plasmacytoma	i ! !		<u> </u>	<3.5	>3.5
	Primary amyloidosis			<u> </u>	<3.5	<3.5
	Other, specify:				3.5-<5.5	-
				<u></u> 3	> 5.5	-
2. Status at HSCT	Never treated	,				
	Complete remission (CR)	-	i. NUMBER	1st	7	
	Partial remission (PR)	-	(complete for CR,	2nd		
	Minimal response (MR)		PR or relapse)	3rd or higher		
	Relapse from CR (untreated)			_	
	Progression	<u></u>				
	No change / stable disease					

For Office Use only: ID: /

For Office Use only: **BLOOD AND MARROW TRANSPLANT** ID: **Notification Form (Disease Classification)** Centre: Instruction: Where check boxes are provided, check (\(\frac{1}{2}\)) one or more boxes. Where radio buttons provided, check ($\sqrt{}$) one box only. i. Patient Name and NRIC Number: ii. Centre Code: iii. Name of reporting centre: SECTION 17 : ANAEMIA 1. Classification: Acquired Severe Aplastic Anaemia (SAA), not otherwise specified Acquired SAA, secondary to hepatitis Acquired SAA, secondary to toxin/other drug Amegakaryocytosis, acquired (not congenital) Acquired Pure Red Cell Aplasia (PRCA) (not congenital) Other acquired cytopenic syndrome, specify: Paroxysmal nocturnal hemoglobinuria (PNH) Congenital: Fanconi anaemia Diamond-Blackfan anaemia (congenital PRCA) Schwachman-Diamond Other congenital anaemia, specify: ___ SECTION 18: HEMOGLOBINOPATHY

1. Classification	:
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- Thalassemia
- Sickle cell disease
- Other hemoglobinopathy, specify: ___

For Office Use only: **BLOOD AND MARROW TRANSPLANT** ID: **Notification Form (Disease Classification)** Centre: Instruction: Where check boxes are provided, check (\(\)) one or more boxes. Where radio buttons (\(\) provided, check ($\sqrt{}$) one box only. i. Patient Name and NRIC Number: ii. Centre Code: iii. Name of reporting centre: SECTION 19: BREAST CANCER - STAGING AT DIAGNOSIS 1. Metastases: Yes Stage: 0 No 2. Classification: Inflammatory Non-inflammatory 3. Status at HSCT: Adjuvant (Stage II, III only) Never treated (upfront) Primary refractory i. NUMBER 1st Complete remission (CR): (complete only for 2nd Confirmed CR or relapse) Unconfirmed (CRU*) 3rd or higher Unknown 1st Partial response (PR1) ii. Sensitivity to Sensitive Untreated chemotherapy Relapse: Resistant (complete only for relapse) Local Metastatic **SECTION 20 : OTHER MALIGNANCIES** 1. Classification: Bone sarcoma (excluding Ewing sarcoma/PNET) Ovarian Central nervous system tumors (include CNS PNET) Pancreas Prostate Ewing sarcoma/PNET, extra-skeletal Renal cell Ewing sarcoma/PNET, skeletal Retinoblastoma Germ cell tumour, extragonadal only Rhabdomyosarcoma Soft tissue sarcoma Hepatobiliary Lung cancer, non-small cell Testicular Lung cancer, small cell Thymoma Wilm tumour Medulloblastoma Melanoma Others, specify: _ Neuroblastoma 2. Status at HSCT: Adjuvant Never treated (upfront) Stable disease/no response i. NUMBER 1st (complete only for Complete remission (CR): 2nd CR or relapse) Confirmed 3rd or higher Unconfirmed (CRU*) 1st Partial response (PR1) ii. Sensitivity to Untreated chemotherapy Relapse Resistant Unknown (complete only for relapse) Progressive disease (PD) SECTION 21: PRIMARY IMMUNE DEFICIENCIES 1. Classification: Absence of T and B cells SCID Kostmann syndrome-congenital neutropenia Absence of T, normal B cell SCID Leukocyte adhesion deficiencies ADA deficiency severe combined immune deficiency (SCID) Neutrophil actin deficiency Ataxia telangiectasia Omenn syndrome Bare lymphocyte syndrome Reticular dysgenesis Cartilage hair hypoplasia SCID other, specify: CD 40 Ligand deficiency SCID, unspecified Chediak-Higashi syndrome Wiskott Aldrich syndrome Chronic granulomatous disease X-linked lymphoproliferative syndrome Common variable immunodeficiency Others, specify:

DiGeorge anomaly

Note: CRU* - complete response with persistent scan abnormalities of unknown significance

(Immune deficiencies, not otherwise specified

	BL	OOD AND MARROW TRAI	NSPLANT		For ID:	Office Us	e only:	
Notification Form (Disease Classification)							/	
Instruction: Where check boxes are provided, check (√) one or more boxes. Where radio buttons are								
provided, check ($$) (one box only							
i. Patient Name and	NRIC Numbe	r:			ii. Centr	e Code:		
iii. Name of reporting	g centre:							
SECTION 22: INHERITE	D DISORDERS	OF METABOLISM						
1. Classification:	Adrenoleu	kodystrophy	Metachroi	matic leukodystrophy				
	Aspartyl gl	ucosaminuria	Morquio (IV)				
	B-glucuror	nidase deficiency (VII)	Mucolipid	Mucolipidoses, unspecified				
	Fucosidos	is	Mucopoly	Mucopolysaccharidosis (V)				
	Gaucher d	isease	Mucopoly	Mucopolysaccharidosis, unspecified				
	Glucose st	orage disease	Niemann-	Niemann-Pick disease				
	Hunter syr	ndrome (II)	Neuronal	Neuronal ceriod – lipofuscinosis (Batten disease)				
	Hurler syn	drome (IH)	Polysacch	naride hydrolase abno	ormalities, unspec	ified		
	I-cell disea	ise	Sanfilippo	(III)				
	Krabbe dis	ease (globoid leukodystrophy)	Scheie sy	rndrome (IS)				
	Lesch-Nyh	an (HGPRT deficiency)	Wolman of	disease				
	Mannosido	osis	Other, spe	ecify:				
	Maroteaux	-Lamy (VI)	Inherited	disorders of metaboli	sm, not otherwise	specified		
SECTION 23 : PLATELE	T and OTHER	INHERITED DISORDERS						
1. Classification:	Glanzman	n thrombasthenia	Osteopetr	rosis (malignant infan	tile osteopetrosis)			
	Congenita	I amegakaryocytosis / congenital thrombocytoper	nia Other oste	Other osteoclast defects, specify:				
	Other inhe	rited platelet abnormalities, specify:						
			_					
SECTION 24: HISTIOCY	TIC DISORDEF	RS						
1. Classification:	Histiocytic	disorders, not otherwise specified	Familial e	rythro/hemophagocyt	ic lymphohistiocyt	osis (FELH)	
	Langerhan	s Cell Histiocytosis (Histiocytosis-X)	Hemopha	Hemophagocytosis (reactive or viral associated)				
	Others, sp	Others, specify:						
SECTION 25: AUTOIMM	IUNE DISORDE	RS						
Autoimmune disorders								
Connective tissue diseas	se:							
 Systemic sclerosis 	(SS) -	1. Involved Organs/Clinical Problem at HSCT	Pr	esence	Indicat	ion for HSC	Т	
		a. Diffuse cutaneous	Yes	○ No	Yes	○ No		
		b. Limited cutaneous	Yes	○ No	Yes	○ No		
		c. Lung parenchyma		◯ No	O Yes	○ No	<u> </u>	
		d. Pulmonary hypertension		○ No	Yes	○ No		
		e. Systemic hypertension	○ Yes	○ No	O Yes	○ No		
		f. Renal Biopsy type:	○ Yes	○ No	O Yes	○ No		
		g. Oesophagus	○ Yes	○ No	O Yes	○ No		
		h. Other GI tract	○ Yes	○ No	○ Yes	○ No		
		i. Raynaud j. CREST	○ Yes	○ No		○ No		
			○ Yes	○ No	○ Yes	○ No	-	
		k. Other, specify: 2. Antibodies studied:	○ Yes	○ No	Yes	● No		
		2. Altibodies studied.		a.Scl 70 positive:	Normal/Neg			
					Elevated/Po Not evaluated			
				b. ACA positive	Normal/Neg			
					Elevated/Po			
					Not evaluate	ed		
			No unknown					
	į		Jan. Kilowii				l i	

BLOOD AND MARROW TRANSPLANT Notification Form (Disease Classification)

Centre: Instruction: Where check boxes are provided, check (\forall) one or more boxes. Where radio buttons provided, check ($\sqrt{}$) one box only. i. Patient Name and NRIC Number: ii. Centre Code: iii. Name of reporting centre: SECTION 25 : AUTOIMMUNE DISORDERS (cont.) Autoimmune disorders (cont.) Connective tissue disease (cont.): Systemic lupus 1. Involved Organs/Clinical Problem at HSCT Indication for HSCT Presence erythematosus (SLE) a. Renal Biopsy type: Yes No Yes No b CNS No Type: Yes No Yes c. PNS Type: Yes No Yes No d. Lung No Yes No Yes e. Serositis Yes Yes No No f. Arthritis No No Yes Yes g. Skin Type: Yes No Yes No h. Haematological Type: Yes No Yes No . Vasculitis Type: Yes No Yes No j. Other, specify: Yes No No Yes 2. Antibodies studied: a.ds DNA: Normal/Negative Yes Elevated/Positive No Not evaluated Unknown b. Complement: Normal/Negative Elevated/Positive Not evaluated c. Others, specify: 1. Involved Organs/Clinical Problem at HSCT Presence Indication for HSCT Polymyositisdermatomyositis No a. Proximal weakness Yes No Yes b. Generalized weakness (including bulbar) Yes No No Yes c. Pulmonary fibrosis Yes No Yes No d. Vasculitis Type: No Yes No Yes e. Other, specify: Yes No Yes No 2. Manifestation with: iv. CPK elevated: i. Typical biopsy: Yes No Yes
No ii. Typical EMG: Yes No v. Malignancy type: No iii. Typical rash (DM): Yes No 1. Involved Organs/Clinical Problem at HSCT Presence Indication for HSCT Sjogren syndrome a. SICCA Yes No Yes No b. Exocrine gland swelling No Yes No Yes c. Other organ lymphocytic infiltration Yes No Yes No d. Lymphoma, paraproteinemia Yes No Yes No e. Other, specify: Yes No Yes No 1. Involved Organs/Clinical Problem at HSCT Presence Indication for HSCT Antiphospholipid syndrome a. Thrombosis Type: Yes No Yes No b. CNS Type: Yes No Yes No c. Abortion Yes No Yes No d. Skin (livido, vasculitis) Yes Yes No No e. Haematological Type: Yes No Yes No f. Other, specify: Yes No Yes No 2. Antibodies studied: Normal/Negative Yes a. Anticardiolipin lgG: Elevated/Positive No Not evaluated Unknown b. Anticardiolipin Normal/Negative IgM: Elevated/Positive Not evaluated c. Others, specify: Other type of connective tissue disease, specify:

For Office Use only:

ID:

BLOOD AND MARROW TRANSPLANT Notification Form (Disease Classification)

Centre: Instruction: Where check boxes are provided, check (\(\)) one or more boxes. Where radio buttons (\(\) provided, check ($\sqrt{}$) one box only. i. Patient Name and NRIC Number: ii. Centre Code: iii. Name of reporting centre: SECTION 25: AUTOIMMUNE DISORDERS (cont.) Autoimmune disorders (cont.) Vasculitis: Indication for HSCT 1. Involved Organs/Clinical Problem at HSCT Presence Wegener granulomatosis a. Upper respiratory tract Yes Yes No b. Pulmonary Yes No Yes c. Renal Biopsy type: No Yes Yes d. Skin Yes No Yes e. Other, specify: Yes No Yes 2. Antibodies studied: a. c-ANCA Normal/Negative Evelated/Positive No Unknown Not evaluated Classical polyarteritis nodosa, classical 1. Involved Organs/Clinical Problem at HSCT Presence Indication for HSCT Biopsy type: Yes Yes No Classical polyarteritis No nodosa, microscopic b. Mononeuritis multiplex Yes No Yes c. Pulmonary haemorrhage No Yes Yes No d. Skin Yes No Yes No e. GI tract Yes No Yes -(<u>•</u>)_No f. Other, specify: No Yes Yes Antibodies studied: Yes a. p-ANCA: Normal/Negative No Elevated/Positive Unknown Not evaluated b. c-ANCA Normal/Negative Elevated/Positive Not evaluated c. Hepatitis Normal/Negative serology Elevated/Positive Not evaluated Other vasculitis: Churg-Strauss Overlap necrotising arteritis Giant cell arteritis Takayasu Behçet's syndrome Other, specify: Arthritis: 1. Involved Organs/Clinical Problem at HSCT Indication for HSCT Presence Rheumatoid arthritis a. Destructive arthritis Yes No Yes No b. Necrotising vasculitis Yes No Yes No c. Eve Type: Yes No Yes No d. Pulmonary No Yes No Yes e. Extra articular specify: No Yes Yes No f. Other, specify No No Yes Indication for HSCT 1. Involved Organs/Clinical Problem at HSCT Presence Psoriatic arthritis/psoriasis a. Destructive arthritis Yes No Yes b. Psoriasis Yes No Yes No c. Other, specify: No No Juvenile idiopathic arthritis (JIA), systemic (Stills disease) Juvenile idiopathic arthritis . Onset: Oligoarticular Polyarticular (JIA), articular: Juvenile idiopathic arthritis: Other, specify: Other arthritis:

For Office Use only:

ID:

	For Office ID:	Use only:					
Instruction: Where c provided, check $()$	are	Centre:					
i. Patient Name and	NRIC Number:	ii.	Centre Code:	:			
iii. Name of reporting	g centre:	_					
SECTION 26 : MULTIPL	E SCLEROSIS						
1. Multiple sclerosis:	 Multiple sclerosis → Primary progressive ○ Secondary progresive ○ Relapsing/remitting ○ Others, specify: 				,		
SECTION 27: OTHER N	EUROLOGICAL AUTOIMMUNE DISEASE						
Other neurological autoimmune disease:	Cal						
SECTION 28 : HAEMAT	DLOGICAL AUTOIMMUNE DISEASES						
Haematological autoimmune diseases:	une apparation of the paratic (11)						
SECTION 29: BOWEL D	ISEASE						
Bowel disease: Crohn's disease							

Ulcerative colitis

Other autoimmune bowel disease, specify:_

Follow	ID: /						
Follow up sheet 1: 1st year post transplant and yearly follow-up Instruction: Where check boxes are provided, check (\(\strict{\lambda}\)) one or more boxes. Where radio buttons are provided, check (\(\strict{\lambda}\)) one box only.							
i. Patient name and NRIG			(Patient identifier for paper CRF)	ii. Centre Code:			
iii.*Date of assessment /	visit / Follow up :	/ (dd/mm/yy) i	v. Date of this HSCT :	/ [dd/mm/yy)			
1. Hospital: (autofill)			2. Unit: (autofill)				
3. Contact person:	a. Name:		b. Phone:				
(autofill)	c. Fax:	d. Email:					
4. Report information:	a. Date of this Report: (autofili) /						
SECTION 1 : PATIENT D	ETAILS						
1. Unique Patient Number or Code:							
2. Name: * (autofill)							
SECTION 2 : DISEASE S	TATUS						
1. Best disease status (response) after transplant: (prior to treatment modification in response to a post transplant	○ Continued complete remission○ CR achieved○ Never in CR	n (CR) i. Date CR achieved/ assesed: /	/ (dd/mm/	(уууу)			
disease assessment)	Not evaluated	Previously reported					
2. Primary Disease Diagnosis: (autofill)	 Acute Leukemias ○ Chronic MyelogenousLeukemia (CML) ○ Other Leukemias : ○ Lymphomas ○ Myelodysplastic Syndrome (MDS) ○ Combined Myelodysplastic/Myeloproliferative Syndrome (MD/MPS) ○ Myeloproliferative Syndrome ○ Plasma Cell Disorder including Multiple Myeloma ○ Anaemia ○ Hemoglobinopathy ○ Breast Cancer - Staging at Diagnosis ○ Other Malignancies ○ Primary Immune Deficiencies ○ Inherited Disorders of Metabolism ○ Hatelet and Other Inherited Disorders ○ Histiocytic Disorders ○ Autoimmune Disorders ○ Other Neurological Autoimmune Disease ○ Bowel Disease 						
SECTION 3 : DATE OF L	SECTION 3: DATE OF LAST CONTACT						
Date of last follow up or death:	1. Date of last						
SECTION 4 : COMPLICA	TIONS OF TRANSPLANT						
1. Late graft failure:	Yes No						
Chronic Graft Versus Host Disease present during this period: (allografts only)	last HSCT CGv	e of diagnosis of / / / / / / / / / / / / / / / / / /	/ (dd/mm.				
	Continuous since last reported episode						
3. Did a secondary malignancy, lymphoproliferative or myeloproliferative	Yes No i. Date of diagnosis: (dd/mm/yyyy)	/ Ii. Diag	gnosis:				

For Office Use only: **BLOOD AND MARROW TRANSPLANT** ID: Follow up sheet 1: 1st year post transplant and yearly follow-up Centre: Instruction: Where check boxes are provided, check (\forall) one or more boxes. Where radio buttons provided, check ($\sqrt{}$) one box only. i. Patient Name and NRIC Number: ii. Centre Code: iii. Name of reporting centre: SECTION 5 : ADDITIONAL TREATMENT 1. Additional No i. Date of treatment: treatment: (dd/mm/yyyy) Yes → ii. Additional cell (Attach the CI sheet completing as many sections as necessary) No infusion: iii. Other disease Yes, planned Yes, not planned No treatment: (planned before transplant) (for relapse/progression or persistent disease) SECTION 6: FIRST RELAPSE OR PROGRESSION 1. First Relapse or No (For acute and chronicLEUKAEMIAS only, tick all methods used for assessment with the dates on which they were used and the results) Progression after Yes i. Relapse/progression HSCT: i. Date assessed: No → (dd/mm/yyyy) detected by clinical/ (Any type) Continuous haematological method: i. Date first seen: progression Yes → (dd/mm/yyyy) since HSCT Previously reported Not evaluated ii. Relapse/progression i. Date assessed: No → (dd/mm/yyyy) detected by cytogenetic method: i. Date first seen: Yes → (dd/mm/yyyy) Not evaluated Previously reported iii. Relapse/progression i. Date assessed: (dd/mm/yyyy) detected by molecular method: i. Date first seen: (Yes → (dd/mm/yyyy) Previously reported Not evaluated SECTION 7: DISEASE PRESENCE/DETECTION AT LAST CONTACT Last disease status (record the most recent status and date for each method, depending on the disease 1. Was disease No i. Last date assessed: detected by (dd/mm/yyyy) clinical/ Yes haematological Previously reported Not evaluated method?: 2. Was disease i. Last date assessed: No detected by (dd/mm/yyyy) cytogenetic ii. Considered disease Yes FISH method? No Yes relapse/progression: (Fill in only for acute and chronic LEUKAEMIAS) Not evaluated Previously reported 3. Was disease i. Last date assessed: (dd/mm/yyyy) detected by molecular method? ii. Considered disease (Fill in only for acute and No Yes relapse/ progression: chronic LEUKAEMIAS) Previously reported Not evaluated **SECTION 8: CONCEPTION** 1. Has patient or partner become pregnant No Yes Unknown Not relevent after this transplant? SECTION 9 : PATIENT STATUS 1. Status: a. Date of death: Alive Unknown (dd/mm/yyyy) Death b. Main cause of Relapse or Progression/Persistent disease death: Secondary malignancy **GVHD** Veno occlusive disorder HSCT Related Cause Rejection/Poor graft function Post transplant Others, specify: lymphoproliferative disorder Pulmonary toxicity Others, specify: Cardiac Toxicity Infection Unknown a. Date of transfer: Unknown Transfer to a (dd/mm/yyyy) new centre b. Centre name: Others. specify Lost to a. Date of last Unknown (dd/mm/yyyy) follow-up

No

Yes

2. Retransplant :

transplantation:

a. Date of

Unknown

(dd/mm/yyyy)

BLOOD AND MARROW TRANSPLANT Follow up sheet 2: 1st year post transplant and yearly follow-up

ID:	/	
Centre:		

ii. Centre Code:

Instruction: Where check boxes \blacksquare are provided, check (\land) one or more boxes. Where radio buttons \bigcirc are provided, check (\land) one box only.

i. Patient Name and NRIC Number:

	ii. Name of reporting centre:							
	FION 10 : CELL INFUSION	B'	O. III is facility (OI) and in a	A 1 . O (1 M				
No.	Date of infusion (dd/mm/yyyy)	Disease status before this CI	Cell infusion (Cl) regimen (not HSCT or autologous stem cell re-infusion)	Acute Graft Versus Host Disease (after this infusion but before any further infusion / transplant)				
		○ CR	i. Type of a. Lmphocytes: Yes No d. Fibroblasts: Yes No	i. Maximum Grade:				
		Not in CR	cell(s): b. Mesenchymal: Yes No e. Others, specify: Yes No	0 (none)				
		Not evaluated	c. Dendritic cells:	<u>0</u> 1				
			ii. TChronological no. of CI for this patient:	© 2 © 3				
			iii. Indication: a. Planned: Yes No f. Treatment viral infection:	<u> </u>				
			b. Loss/decreased chimaerism: Yes No G. Treatment PTLD Yes No No When the third of the thir	Present but grade unknown				
			EBV lymphoma: Yes No					
			d. Prophylactic: Yes No disease: Yes No					
			e. Treatment of GvHD: O Yes No i. Others, specify: Yes No					
			iv. Number of infusions within 10 weeks: (count only infusions that are part of same regimen and given for the same indication)					
		CR		i. Maximum Grade:				
		OR Not in CR	cell(s): b. Mesenchymal: Yes No e. Others, specify: Yes No					
		Not evaluated	c. Dendritic cells: Yes No	0 (none)				
			ii. TChronological no. of CI for this patient:	<u>0</u> 2				
			iii. Indication: a. Planned: Yes No f. Treatment viral Yes No	3 4				
			b. Loss/decreased Yes No	Present but grade				
			c. Treatment PTLD, Yes No	unknown				
			d. Prophylactic: Yes No No Ves No					
			e. Treatment of GvHD: Output					
			iv. Number of infusions within 10 weeks:					
		_	(count only infusions that are part of same regimen and given for the same indication)					
		○ CR	001/0	i. Maximum Grade:				
		Not in CR Not evaluated	b. Mesenchymal:	0 (none) 1				
			ii. TChronological no. of CI for this patient:	<u>0</u> 2				
			iii. Indication: a. Planned: Yes No f. Treatment viral Yes No	3 4				
			b. Loss/decreased chimaerism: Order Yes No	Present but grade unknown				
			c. Treatment PTLD, EBV lymphoma: Yes No	unknown				
			d. Prophylactic: Yes No disease:					
			e. Treatment of GvHD: Order No i. Others, specify: Yes No					
			iv. Number of infusions within 10 weeks: (count only infusions that are part of same regimen and given for the same indication)					
		CR	i. Type of a. Lmphocytes: Yes No d. Fibroblasts: Yes No	i. Maximum Grade:				
		Not in CR	cell(s): b. Mesenchymal: Yes No e. Others, specify: Yes No	(0 (none)				
		Not evaluated	c. Dendritic cells: Yes No	0 1				
			ii. TChronological no. of CI for this patient:	<u>0</u> 2				
			iii. Indication: a. Planned: Yes No f. Treatment viral Yes No					
			b. Loss/decreased	4 Present but grade				
			c. Treatment PTLD, Yes No	unknown				
			d. Prophylactic: Yes No disease:					
			e. Treatment of GvHD: Output					
			iv. Number of infusions within 10 weeks:					
			(count only infusions that are part of same regimen and given for the same indication)					

Ap	pendix: O	ptiona	ai item	is for CIB	MTR Research Centers		
Instruction: Where c provided, check $()$ c	heck boxes one box only.	are pro	ovided, cl	heck ($$) one o	r more boxes. Where radio buttons (are Centre:	
i. Patient Name and NRIC Number:						ii. Centre Code:	
iii. Name of reporting ce	entre:						
SECTION 1 : PRE-HSCT	DISEASE THEF	RAPY			1		
Was imatinib mesylat anytime prior to start (FOR ACUTE LEUKAEM.	of prep regimen		herapy	○ No	Yes Unknown		
2. Did recipient receive to the control of the cont		to this HSC	CT?	○ No	Yes Unknown		
3. Treated :				Combination	n chemotherapy Imatinib mesylate (Gleev	vec, Glivec) Others, specify:	
*				Dasatinib (S	· · ·		
				Hydroxyurea	a (HU) Nilotinib (Tasigna)		
					_		
SECTION 2 : EX VIVO G							
Instruction :Fill in only if the 1. Ex vivo graft			_				
manipulation regimen:	T-cell deplet		Other	negative selection	n, specify: CD34 selection Expansion	Others, specify:	
CECTION A - DOCT LICE	T DICEAGE THE	TD A DV					
SECTION 3 : POST-HSC			ent (anart fro	om cell infusion) i	n the Day 100 report sheet 2 is Yes: Planned.		
1. Post-HSCT disease	<u> </u>		int (apart in		rathecal Chemotherapy	Rituximab (Rituxan, Mabthera)	
therapy:	Bortezomib Cellular the	. ,	CI. DLI)		atinib mesylate (Gleevec, Glivec)	Thalidomide (Thalomid)	
	FGF (velafe		,,	Lei	nalidomide (Revlimid)	Others, specify:	
	KGF (palifer	rmin, Kepiv	rance)	Loc	cal radiotherapy		
SECTION 4 : COMORBI	D CONDITIONS						
1. Is there a history of	○ No	⊘ Voc		(Lin	known		
mechanical ventilation?	○ No	Yes		0 011	KIIOWII		
2. Is there a history of proven invasive	○ No	Yes		① Un	known		
fungal infection?							
Were there clinically sorgan impairment at t				No	Unknown		
to preparative regimen?		•					
Disease	!	Absent	Present	Not evaluated	Comorbidity Definitions		
Arrhythmia		0	0	0	Atrial fibrillation or fl utter, sick sinus syndrol	me, or ventricular arrhythmias	
Cardiac		0	<u> </u>	0	Coronary artery disease *, congestive heart	failure, myocardial infarction, or EF ≤ 50%	
Cerebrovascular diseas	e	0	0	0	Transient ischemic attack or cerebrovascular accident		
Diabetes		0	0	0	Requiring treatment with insulin or oral hypoglycemics but not diet alone		
Heart valve disease Hepatic, mild		0	0	0	Except mitral valve prolapse		
• -	ro	0	0	0	Chronic hepatitis, bilirubin > ULN to 1.5 × ULN, or AST/ALT > ULN to 2.5 × ULN		
Hepatic, moderate/severe			0	Liver cirrhosis, bilirubin > 1.5 × ULN, or AST/ALT > 2.5 × ULN			
Infection Inflammatory bowel disease			Requiring continuation of antimicrobial treatment after day 0 Crobbis disease or placeative colitics				
Inflammatory bowel disease Obesity		0	Crohn's disease or ulcerative colitis				
Peptic ulcer		0	Patients with a body mass index > 35 kg/m2				
Psychiatric disturbance			Requiring treatment Depression or anxiety requiring psychiatric consult or treatment				
Pulmonary, moderate			DLco and/or FEV1 66-80% or dyspnea on slight activity				
Pulmonary, severe			DLco and/or FEV1 66-80% or dyspnea arrest or requiring oxygen				
Renal, moderate/severe		0	0		Serum creatinine > 2 mg/dL or >177 µmol/L, on dialysis, or prior renal transplantation		
		0		SLE, RA, polymyositis, mixed CTD, or polymyalgia rheumatica			
		0	0			st history, excluding nonmelanoma skin cancer	
Others, specify:					, , , , , , ,	<u> </u>	

For Office Use only:

ID:

^{*} One or more vessel-coronary artery stenosis requiring medical treatment, stent, or bypass graft. EF indicates ejection fraction; ULN, upper limit of normal; SLE, systemic lupus erythmatosis; RA, rheumatoid arthritis; CTD, connective tissue disease; DLco, diffusion capacity of carbon monoxide.