

Czech Republic, 323 00 Plzeň, Alej Svobody 80 Fax: +420 373 034 442, Phone: +420 373 034 333, E-mail: registr@kostnidren.cz

VERIFICATION OF CELL PRODUCT

Page 1 of 2

HPC, Marrow		HPC, Apheresi	S	T-Cells, Apheresis		
PATIENT/DONOR DATA						
Patient name:			Patient ID:			
Patient registry:			(assigned by patient registry) Patient ID:			
Transplant center:			(assigned by donor registry)			
Date of Birth: (YYYY-MM-DD)	Gende		Weight in kg:	Blood group/RhD:		
			r GRID:			
Donor registry:	1		1			
Date of Birth: (YYYY-MM-DD)	Gender:		Weight in kg:	Blood group/RhD:		
SECTION A: to be completed by	the do	onor center				
Comments:						
Person completing form:		Date (YYYY-MM-DD)	:	Donor center signature:		
SECTION B: to be completed by	the co	llection/apheresis	scenter			
Institution:						
Address:						
			Collection date(s) (YYYY-MM-DD):			
			Start date G-CSF (Y			
			Anticoagulants: Heparin ACD EDTA			
			Other:	Volume/ratio:		
Attention:				be collected at the time of the collection:		
Phone:			ml heparin ml ACD			
Fax:			ml EDTA	ml no anti-coagulant		
E-mail:			ml marrow	tube, type:		
Based on the experience at this ce	nter, w	e feel that the requ	ested amount of cells	is:		
Feasible Note that this is n collected cells ma	•		quested number of co	ells will be supplied. The number of		
Not feasible		0				
Comments:						
Person completing form: Date		Date (YYYY-MM-DD)	:	Collection/apheresis center signature:		
-						





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PATIENT/DONOR DATA	
Patient name:	Patient ID:
Patient registry:	(assigned by patient registry)
	Patient ID:
Transplant center:	(assigned by donor registry)
Donor ID:	Donor GRID:
Donor registry:	

DISCLAIMER:

- The cell products collected from this donor are intended solely for the purpose of immediate therapeutic treatment for the above mentioned patient.
- Excess cells may be stored for future therapeutic treatment for this patient. No other uses of these cells are permissible. Cells not used for the therapeutic treatment of the above mentioned patient must be disposed of properly.
- The donor center must be provided detailed information concering the use and/or disposal of all portions of this cell product. By accepting these cells, the transplant physician also accepts these terms and conditions. Deviations from these terms are not permitted without prior written approval from the donor center.
- Any serious product events and/or adverse effects must be reported both to the donor's registry and transplant center. Corresponding SEAR/SPEAR reports must be completed and provided to the WMDA Office.

SECTION C: transplant center acceptance of terms provided by donor & collection/apheresis centers				
Person completing form:	Date (YYYY-MM-DD):	Transplant center signature:		

