

Appendix A: Cross-Network PBMC Processing Worksheet v7.0

Note: The fields in this worksheet must be filled out by hand, using a pen.

Specimen Processing Laboratory:					Protocol:
Participant ID (PTID/PID):			Visit Number:		Visit Type:
Collection Date:			Collection Tim	e:	
Processing Start Date:			Processing Sta	rt Time:	Processed By (Initials):
Reagents	Manufacturer Lot Number			Expiration Date	
DMSO					
FBS					
WDR: HBSS or PBS (circle one)					
Cell Separation Tube (frit)					
Density Gradient Media					
		1	L (record as X.Y)		Expiration Date
CPS		CPS	DMSO	FBS	1 working day (<18hrs)
Data to be Captured During Pro	cessing				Sample
Sample tube type (circle one or record "other" tube type)					ACD / HEP / EDT Other:
Blood condition (circle one or more; add comments on reverse as needed)					SAT/ HEM / CLT
Measured usable whole blood volume (to the nearest 0.1mL)					m
Indicate processing method (circle one)					CSTFB / overlay / underlay
Counting Method: Name of specific instrument or manual count (record in field to right)					
Counting re-suspension volume	of HBSS (or oth	er WDR) (V) (re	ecord as X.Y)		m
Cell count average concentration	on (C)				x 10 ⁶ cells/m
Total cell number (T) = C x V					x 10 ⁶ cell
Calculate cell yield/mL of whole	-	ck)= (T/Usable	Whole Blood Volu	me)	x 10 ⁶ cells/m
If <i>T/A* ≥ N1</i> ; CPS re-suspension If <i>T/A < N1</i> ; Calculate estimated	m				
Calculate final CPS re-suspension volume (V_f), (V1 rounded DOWN to nearest whole (X.0) mL)					m
Calculate actual number of cells per vial. $N2 = (T/V_f) \times V2$; (V2=1 mL) Note: Do not store more than 50 million cells per vial					x 10 ⁶ cells/via
Print and QC LDMS Label conte	nt/barcodes (init	tials of person	(s) performing QC)		
Frozen Date and Time (ddMMM 4 hours of processing start time		Explain in com	ments section if no	ot within	
Number of Cryovials actually from Note: Should be equal to final C		n volume for 1	mL aliquots (V f) ar	nd ≤ (A)	
Complete remaining LDMS entr					
* <u>Note:</u> A = The maximum numbe	er of aliquots req	uired accordin	g to the protocol-	specific Laborat	ory Processing Chart (IPC)

*<u>Note:</u> **A** = The maximum number of aliquots required according to the protocol-specific Laboratory Processing Chart (LPC). Do not store more than this number of aliquots.



Appendix A: Cross-Network PBMC Processing Worksheet v7.0 (Page 2 of 2)

Note: The fields in this worksheet must be filled out by hand, using a pen.

Specimen Processing Laboratory:

PTID/PID:	
Transfer of Cryovials to Freezer Storage Box	
Person who transferred cryovials to storage box locations assigned by LDMS	
Date (ddMMMyyyy)/time cryovials were transferred from controlled-rate freezing device to storage box. (Sample must be maintained at -80°C during transfer)	
Initial (Primary) Review (Initials/Date)	
Final (Secondary) Review (Initials/Date)	

Hemacytometer Counts	Total Count	Viable Cells	Non-Viable
Square #1 (cells/mm ²)			
Square #2 (cells/mm ²)			
Square #3 (cells/mm ²)			
Square #4 (cells/mm ²)			
Average Cell Count per Square (cells/mm ²)			
PBMC Dilution Factor (1:DF**)			
Hemacytometer Factor for cells/mL	104	10 ⁴	10 ⁴
Cell count concentration (C) = (Average Cells/mm ²)(DF)(10 ⁴); convert to 10 ⁶ cells/mL	Not applicable	x 10 ⁶ cells/mL	Not applicable
% viability = (Viable cells 4 squares/total cells 4 squares) (100)	Not applicable		Not applicable

**<u>Note</u>: Dilution Factor (DF) = (parts cells + parts dilution fluid)/ parts cells

Automated Cell Counts (10 ³ /µl=10 ⁶ /mL)	Count #1
Cell Count (C) as cells x 10 ⁶ /mL	
PBMC Dilution Factor (1:DF***)	
Cell Concentration = (C)(DF)	
	x 10 ⁶ cells/mL
% viability (if applicable)	

***<u>Note</u>: Dilutions for automated counters are extremely rare. If performing direct counts, enter a 1 in the DF box and complete the column.

Comments, protocol deviations, and additional information not captured elsewhere in this worksheet:



Note: (A) = The maximum number of aliquots required according to the protocol-specific Laboratory Processing Chart (LPC). Do not store more than this number of aliquots.



Cross-Network PBMC Processing Standard Operating Procedure

<u>Note:</u> The fields in this worksheet must be filled out by hand, using a pen. Specimen Processing Laboratory: Lab 398				Protocol: 313	
Participant ID (PTID/PID): 123-4		Visit Number:	2.0	Visit Type: vst	
Collection Date: 08AUG2024			Collection Tim	ne: 08:00	
			Processing Sta	art Time:	
rocessing Start Date: 08AUG2024			08:45 Lot Number		Processed By (Initials): CN
Reagents DMSO	Manufacturer Sigma		RNBM0548		Expiration Date
FBS	Peak		13G1212		18AUG2025
WDR: HBSS or PBS (circle one)	Gibco		2660057		30APR2026
Cell Separation Tube (frit)	Greiner		E220337Q		14MAR2027
,					31AUG2026
Density Gradient Media	Cytiva	Volumo in n	1Q345061		
		CPS	nL (record as X.Y) DMSO	FBS	Expiration Date
CPS Prepared 19AUG2024 08:3	0 CN	9.0	0.9	8.1	1 working day (<18hrs)
Data to be Captured During Pro	Sample				
Sample tube type (circle one or record "other" tube type)					ACD / HEP / EDT Other:
Blood condition (circle one or more; add comments on reverse as needed)					SAT/ HEM / CLT
Measured usable whole blood volume (to the nearest 0.1mL)					86.3 m
Indicate processing method (circle one)					CSTFB overlay / underlay
Counting Method: Name of spe	ecific instrume	nt or manual co	ount (record in field	l to right)	Manual Coun
Counting re-suspension volume	of HBSS (or of	ther WDR) (V) (record as X.Y)		17.0 m
Cell count average concentration	on (C)				7.2 x 10 ⁶ cells/m
Total cell number (T) = C x V					122.4 x 10 ⁶ cell
Calculate cell yield/mL of whole	<mark>e blood. (QC ch</mark>	eck)= (T/Usabl	e Whole Blood Volu	<mark>ıme)</mark>	1.4 x 10 ⁶ cells/m
<pre>If T/A* ≥ N1; CPS re-suspension If T/A < N1; Calculate estimated</pre>		nsion vol. (V1) =	(T/N1x10 ⁶ cells/m	L)(1mL)	<mark>5.0 m</mark>
Calculate final CPS re-suspension volume (V), (V1 rounded DOWN to nearest whole (X.0) mL)					<mark>5.0 m</mark>
Calculate actual number of cells per vial. N2 = (T/V₂) x V2 ; (V2=1 mL) Note: Do not store more than 50 million cells per vial					24.4 x 10 ⁶ cells/via
Print and QC LDMS Label content/barcodes (initials of person (s) performing QC)					CN
Frozen Date and Time (ddMMMyyyy /HH:MM) (Explain in comments section if not within 4 hours of processing start time)					08AUG2024
Number of Cryovials actually frozen Note: Should be equal to final CPS re-suspension volume for 1mL aliquots (V_f) and \leq (A)					5
Complete remaining LDMS entries including total cell count & frozen time (Initials)					CN

Example #1: $N1 = 20 \times 10^6$ cells/mL

A = 5 aliquots

<u>Calculations:</u> CPS re-suspension volume (V1)

122.4/5 = 24.4>20

Thus, T/A ≥ N1

(V1) = A

Actual number of cells per vial (N2)

122.4/5 x 1 = 24.4x10⁶ cells/vial

Page 1 of 1



Cross-Network PBMC Processing Standard Operating Procedure Examples

Cross-Network PBMC Processing Standard Operating Procedure

<u>Vote:</u> The fields in this worksheet must be filled out by hand, using a pen. Specimen Processing Laboratory: Lab 398				Protocol: 313	
Participant ID (PTID/PID): 123-456789			Visit Number:	2.0	Visit Type: vst
Collection Date: 08AUG2024			Collection Tin	ne: 08:00	
Processing Start Date: 08AUG20)24		Processing St	art Time: 08:45	Processed By (Initials): CN
Reagents	Manufacture	r	Lot Number		Expiration Date
DMSO	Sigma		RNBM0548		18JAN2025
FBS	Peak		13G1212		18AUG2025
WDR: HBSS or PBS (circle one)	Gibco		2660057		30APR2026
Cell Separation Tube (frit)	Greiner		E220337Q		14MAR2027
Density Gradient Media	Cytiva		1Q345061		31AUG2026
		Volume in m	nL (record as X.Y)		Expiration Date
CPS Prepared 19AUG2024 08:3	0 CN	CPS	DMSO	FBS	1 working day (<18hrs)
		9.0	0.9	8.1	
Data to be Captured During Pro					Sample ACD/ HEP / EDT
Sample tube type (circle one or record "other" tube type)					Other:
Blood condition (circle one or more; add comments on reverse as needed)					SAT HEM / CLT
Measured usable whole blood volume (to the nearest 0.1mL)					46.3 r
Indicate processing method (cir	cle one)				CSTFB overlay / underlay
Counting Method: Name of spe	ecific instrumer	nt or manual cou	unt (record in field	d to right)	Manual Cou
Counting re-suspension volume	of HBSS (or ot	her WDR) (V) (re	ecord as X.Y)		9.0 r
Cell count average concentratio	n (C)				4.2 x 10 ⁶ cells/r
Total cell number (T) = C x V					<mark>37.8</mark> х 10 ⁶ се
Calculate cell yield/mL of whole	0.8 x 10 ⁶ cells/r				
If T/A* ≥ N1 ; CPS re-suspension If T/A < N1 ; Calculate estimated		nsion vol. (V1)=(T/N1x10 ⁶ cells/m	L)(1mL)	3.7 r
Calculate final CPS re-suspension volume (V), (V1 rounded DOWN to nearest whole (X.0) mL)					3.0 r
Calculate actual number of cells per vial. N2 = (T/V;) x V2; (V2=1 mL) Note: Do not store more than 50 million cells per vial					12.6 x 10 ⁶ cells/v
Print and QC LDMS Label content/barcodes (initials of person (s) performing QC)					CN
Frozen Date and Time (ddMMMyyyy /HH:MM) (Explain in comments section if not within 4 hours of processing start time)					08AUG2024
Number of Cryovials actually frozen Note: Should be equal to final CPS re-suspension volume for 1mL aliguots (V_f) and \leq (A)					3
	ies including to				

Example #2: $N1 = 10 \times 10^6$

cells/mL A = 5 aliquots

<u>Calculations:</u> CPS re-suspension volume (V1)

37.8/5 = 7.56<10

Thus, T/A < N1

(V1) = 37.8/10x10⁶ cells/mL)(1mL)

Actual number of cells per vial (N2)

37.8/3 = 12.6x10⁶ cells/vial

*<u>Note</u>: **A** = The maximum number of aliquots required according to the protocol-specific Laboratory Processing Chart (LPC). Do not store more than this number of aliquots.

Page 1 of 1