

MIBlood-EV

Standardized Reporting Tool for Blood EV Research (Human)

STUDY INFORMATION

1.0 Manuscript title			
1.1 Corresponding autl	nor (Name and Er	nail)	
1.2 Institution name			
1.3 Time period of exp	eriment (years)		1.4 Number of samples
1.5 Cargo of interest	Vesicles	Protein	RNA DNA Other:
^{1.6} Biospecimen type	Plasma	Serum	^{1.7} Biospecimen state
^{1.8} Source of frozen sp	ecimens		1.9 Years of collection (range)

BLOOD COLLECTION AND PROCESSING

^{2.0} Patient fasting status	2.1 Fasting length (e.g. hours/days)							
^{2.2} Anatomical access site		^{2.3} Needle diameter (e.g. gauge)						
^{2.4} Blood volume collected (mL)								
^{2.5} Plasma anticoagulant	EDTA	A Citrate Heparin Other:						
^{2.6} Serum tube type		^{2.7} Serum clotting time (minutes)						
2.8 Time between collection and first centrifugation (range in hours)								
^{2.9} Transport temperature		2.10 Transport condition of tubes						
2.11 Centrifuge brand and model								
^{2.12} Bucket rotor type		2.13 Number of centrifugation cycles						
2.14 1st Centrifugation speed (R	CF in x g)	^{2.15} 1 st Rotor brake						
2.16 1st Centrifugation tempera	ture	2.17 2 nd Centrifugation speed (RCF in x g)						
^{2.18} 2 nd Rotor brake		2.19 2 nd Centrifugation temperature						
^{2.20} Additional								
processing steps								
(e.g. filtration)								
2.21 Storage tubes (brand, type, source, catalog number)								
2.22 Storage temperature 2.23 Length of storage (range in years)								

PLASMA/SERUM QUALITY CONTROL

3.0 Number of freeze-thaw cycles (range)		
3.1 Thawing temperature	3.2 Thawing duration (minutes)	

Hemolysis

^{3.3} Presence of hemolysis		3.4	^{3.4} Number of samples affected (e.g. <25%, 25-50%)							
^{3.5} Method used				3.6 RBC count (Median, 95% CI, N)						
3.7 RBC counter brain	nd and type	2								
3.8 Spectrophotometry hemoglobin concentration (g/L)										
3.9 Spectrophotome	ter brand,	model and	t							
wavelength meas	ured (e.g. 4	114 nm)								
3.10 Hemolized samp	oles were d	iscarded					_			



<u>Platelets</u>

3.11 Presence of platelets	3.12 Method used (e.g. Flow Cytometry)
3.13 Marker(s) used (e.g. CD61, CD41)	
3.14 Concentration (median, 95% CI, N)	
^{3.15} Platelet counter instrument brand	
and type	
^{3.16} Flow cytometer brand and type	
3.17 Flow cytometry size and	
fluorescence ranges of detection in	
nanometers and MESF, respectively	

Lipoproteins

3.18 Presence of lipoproteins	^{3.19} Method used (WB, ELISA, FC)					
3.20 Spectrophotometry L-index						
3.21 Spectrophotometer brand, model and						
wavelength measured (e.g. 7	00 nm)					
3.22 WB Marker(s) used (e.g. Ap	o B)					
3.23 Western blot images provid	3.23 Western blot images provided in manuscript?					
3.24 Flow cytometry marker(s) used (e.g. ApoB)						
3.25 Flow cytometry concentrati	on (median, 95% CI, N)					
3.26 Flow cytometer brand and t	ype					
3.27 Flow cytometry size and						
fluorescence ranges of detec	tion					
in nanometers and MESF,						
respectively						