

MIBlood-EV

Standardized Reporting Tool for Blood EV Research (Human)

STUDY INFORMATION

^{1.0} Manuscript title							
^{1.1} Corresponding aut	hor (Name and Er	nail)					
^{1.2} Institution name							
^{1.3} Time period of experiment (e.g. 2022-2024)					^{1.4} Number o	of samples	
^{1.5} Cargo of interest	Vesicles	Protein	RNA	DN	A Other	:	
^{1.6} Biospecimen type	Plasma	Serum	^{1.7} Bios	pecim	en state		
^{1.8} Source of frozen sp			^{1.9} Ye	ars of collect	ion (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 E			Citrate	Heparin	Other:		
^{2.6} Serum tube type		^{2.7} Serum clotting time (minutes)					
^{2.8} Time between collection	n and first cent	rifugati	ion (range	e in hours)			
^{2.9} Transport temperature			^{2.10} Trans	port condition o	of tubes		
^{2.11} Centrifuge brand and m	nodel						
^{2.12} Bucket rotor type			^{2.13} Num	ber of centrifug	ation cycles		
^{2.14a} 1 st Centrifugation speed (RCF in x g)			^{2.14b} 1 st Centrifugation time (minutes)				
^{2.15} 1 st Rotor brake	^{2.16} 1 st Centrifugation temperature						
^{2.17a} 2 nd Centrifugation speed (RCF in x g)			2.17b 2 nd Centrifugation time (minutes)				
^{2.18} 2 nd Rotor brake		2.1	^{.9} 2 nd Cent	rifugation temp	perature		
^{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 d	luration (minutes)	

<u>Hemolysis</u>

^{3.3} Presence of hemolysis		^{3.4} Frequency of hemolyzed samples (e.g. <25%, 25-50%)						
^{3.5} Method used				^{3.6} RBC count (Median, 95%	% CI, N)		
^{3.7} RBC counter brand and type								
^{3.8} Spectrophotome	n concent	ration (mean g/	L)					
^{3.9} Spectrophotometer brand, model and		del and						
wavelength measured (e.g. 414 nm)		nm)						
^{3.10} Hemolized sam	ples we	re disca	rded					



Platelets

^{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	i, 95% Cl, N)	
^{3.15} Platelet counter instru	ment brand,	
type and limit of detection (cells/L)		
^{3.16} Flow cytometer brand	and type	
^{3.17} Flow cytometry size an	d	
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-inde	x							
^{3.21} Spectrophotometer brand, model and								
wavelength measured (e.g. 700 nm)								
^{3.22} WB Marker(s) used (e.g. Apo B)								
^{3.23} Western blot images provi	^{3.23} Western blot images provided in manuscript?							
^{3.24} Flow cytometry marker(s)	used (e.g. A	роВ)						
^{3.25} Flow cytometry concentra	tion (media	n, 95% Cl, N)						
^{3.26} Flow cytometer brand and	type							
^{3.27} Flow cytometry size and								
fluorescence ranges of dete	ection							
in nanometers and MESF,								
respectively								
^{3.28} Brand and catalog number	for							
ELISA kit used for each								
Apolipoprotein test (e.g. Ap	роВ)							
^{3.29} Concentration measured f	or each							
Apolipoprotein tested (mea	an							
μg/mL)								