INTERNATIONAL SOCIETY for EXTRACELLULAR VESICLES

MIBlood-EV

Standardized Reporting Tool for Blood EV Research (Human)

STUDY INFORMATION

1.0 **Manuscript title**			
1.1 **Corresponding author (Name and Email)**			
1.2 **Institution name**			
1.3 **Time period of experiment (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 specimens**		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 ☐ 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 (RCF in x g)**		2.15 **1st Rotor brake**	
2.16 **1st Centrifugation temperature**		2.17 **2nd Centrifugation speed (RCF in x g)**	
2.18 **2nd Rotor brake**		2.19 **2nd 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 **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 brand and type**			
3.8 **Spectrophotometry hemoglobin concentration (g/L)**			
3.9 **Spectrophotometer brand, model and wavelength measured (e.g. 414 nm)**			
3.10 **Hemolized samples were discarded**			

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, 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. 700 nm)**			
3.22 **WB Marker(s) used (e.g. Apo B)**			
3.23 **Western blot images provided in manuscript?**			
3.24 **Flow cytometry marker(s) used (e.g. ApoB)**			
3.25 **Flow cytometry concentration (median, 95% CI, N)**			
3.26 **Flow cytometer brand and type**			
3.27 **Flow cytometry size and fluorescence ranges of detection in nanometers and MESF, respectively**			