

## MIBlood-EV

Standardized Reporting Tool for Blood EV Research (Human)

# **STUDY INFORMATION**

1.0 Manuscript title						
1.1 Corresponding aut	hor (Name and E	mail)				
1.2 Institution name						
1.3 Time period of experiment (e.g. 2022-2024)				<sup>1.4</sup> N	umber of sample	es
<sup>1.5</sup> Cargo of interest	Vesicles	Protein	RNA	DNA	Other:	
<sup>1.6</sup> Biospecimen type	Plasma	Serum	<sup>1.7</sup> Bios	pecimen s	tate	
<sup>1.8</sup> Source of frozen sp	ecimens			1.9 Years o	f collection (ran	ge)

## **BLOOD COLLECTION AND PROCESSING**

		· ····	<u> </u>				
<sup>2.0</sup> Patient fasting statu	ıs		<sup>2.1</sup> Fasti	<sup>2.1</sup> Fasting length (e.g. hours/days)			
<sup>2.2</sup> Anatomical access site				<sup>2.3</sup> Needle diar	neter (e.g. ga	uge)	
<sup>2.4</sup> Blood volume collec	ted (mL)						
<sup>2.5</sup> Plasma anticoagular	nt	EDTA	Citrate	Heparin	Other:		
<sup>2.6</sup> Serum tube type			<sup>2.7</sup> Seru	m clotting time	e (minutes)		
2.8 Time between collection and first centrifugation (range in hours)							
2.9 Transport temperate	ure		2.10 Transport condition of tubes				
2.11 Centrifuge brand ar	nd model						
2.12 Bucket rotor type			<sup>2.13</sup> Numl	er of centrifug	ation cycles		
2.14a 1st Centrifugation speed (RCF in x g)		in x g)	2	14b 1st Centrifug	gation time (r	minutes)	
2.15 1st Rotor brake				ugation tempe	erature		
2.17a 2nd Centrifugation speed (RCF in x g)		in x g)	2.	<sup>.7b</sup> 2 <sup>nd</sup> Centrifug	gation time (r	minutes)	
2.18 2 <sup>nd</sup> Rotor brake			2.19 2 <sup>nd</sup> Cent	rifugation tem	perature		
<sup>2.20</sup> Additional							
processing steps							
(e.g. filtration)							
2.21 Storage tubes (brand, type, source, catalog number)							
<sup>2.22</sup> Storage temperature			Length of s	orage (range ii	n years)		
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## 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		<sup>3.4</sup> Fre	3.4 Frequency of hemolyzed samples (e.g. <25%, 25-50%)				
<sup>3.5</sup> Method used		·		<sup>3.6</sup> RBC count (M	ledian, 95% CI, N	1)	
3.7 RBC counter bra	nd and type						
3.8 Spectrophotometry hemoglobin concen				ration (mean g/L)	)		
<sup>3.9</sup> Spectrophotometer brand, model and			/				
wavelength measured (e.g. 414 nm)			nm)				
3.10 Hemolized samples were discarded							



# <u>Platelets</u>

3.11 Presence of platelets		3.12 Method used (e.g. Flow Cytometry)	
3.13 Marker(s) used (e.g. CD	061, CD41)		
3.14 Concentration (median	, 95% CI, N)		
3.15 Platelet counter instru	ment brand,		
type and limit of detect	ion (cells/L)		
<sup>3.16</sup> Flow cytometer brand and type			
3.17 Flow cytometry size and			
fluorescence ranges of o	detection in		
nanometers and MESF,	respectively		

Lipoproteins

3.19 Method used (WB, ELISA, FC)
and
nanuscript?
g. ApoB)
dian, 95% CI, N)