Order of draw		Prepa	ration & cent	rifugation conditions*	S-Monovette®	2000 x g	2500 x g	3000 x g*	3500 x g*	400
Recommendation according to Gurr ¹	Recommendation according to CLSI ²		Additive	Fields of application	Serum	10 min	10 min	6 min	4 min	
Blood culture	Blood culture		Serum	Clinical chemistry The S-Monovette® Serum CAT contains plastic beads coated with a coagulation activator (silicate). As a result of this coagulant Blood coagulation is usually complete within 20-30 minutes, after which the sample can be centrifued	Serum Gel	15 min	10 min	4 min	4 min	
Serum /		officiations reserved	Serum Gel**	Clinical chemistry In addition to the coated beads, the S-Monovette® Serum-Gel CAT contains a polymer-based gel. Due to its density, this gel forms a stable layer separating the coagulum and the serum during centrifugation and acts as a barrier during transport and storage of the sample.	Li-Heparin	10 min	10 min	7 min	7 min	
	Citrate*				Li-Heparin-Gel	15 min	15 min	10 min	7 min	
Citrate* Serum / Litrate Citrate* Serum Gel	Technical mo	Lithium- Heparin	Clinical chemistry The S-Monovette® Heparin contains plastic beads coated with	Li-Heparin-Gel⁺	8 min	7 min	5 min	4 min		
		nlar countries	Lithium-	The anticoaguiant nepann (generally 16 IU of nepaninmi of blood), or heparin is present in spray-dosed form (generally 19 IU/ml of blood). The S-Monovette® Lithium-Heparin-Gel/-Gel* also contains a polymer-based gel which settles between blood cells	EDTA	n.v.	n.v.	7 min	6 min	
Heparin/ Heparin Gel	Heparin/ Heparin Gel	e available in partic	EDTA	and plasma after centrifugation. Hematology The S-Monovette® EDTA K3E contains the anticoagulant K3 EDTA	EDTA Gel	15 min	10 min	10 min	7 min	
EDTA	EDTA	tion on products that may not b	EDTA-Gel**	in spray-dosed form (1.6 mg EDTA/ml blood). Molecular virus diagnostics In addition to EDTA (1.6 mg/ml blood), the S-Monovette® EDTA Gel K2E also contains a polymer-based gel for a stable layer separating blood cells and plasma. Coagulation	Citrate	9 min	8 min	7 min	6 min	
					Fluoride	9 min	8 min	7 min	6 min	
Fluoride	Fluoride	nt may contain informa	Trisodium Citrate 1:10	The S-Monovelue' circle with contains the anticogularit trisodium citrate as a 0.106 molar solution (= 3.13 % trisodium citrate solution; often rounded up to 3.2 %) and amounts to 10 % of the nominal volume. The mixing ratio 1:10 (1 volume fraction of citrate and 9 volume fractions of blood) must be observed ($_{\rm correct}$ filling)	Citrate PBM 1.8 ml Rotor Ø > 17 cm	9 min	8 min	7 min	6 min	
l Gurr at al "Mustantandardarbaiteanunisuna Präanaktik". Li ah Mad 2011		This docume	Fluesda	Glucose The S-Monovette® Fluoride/EDTA contains fluoride (1.0 mg/	Citrate PBM 1.8 ml Rotor Ø > 9 cm to < 17 cm	n.v.	n.v.	10 min	n.v.	
CLSI Procedures for the Collection of Diag Approved Standard, GP41 ED7:2017 7th E If a citrate sample is collected first using a S	Institution of the mode 2017 Institution April 2017 Safety-Multifly needle,	2200_503_ISO	Fluoride	mi blood) as glycolysis inhibitor and EDIA (1.2 mg/ml blood) as anticoagulant.	n.v. = not validated Conditions apply to a tempe * Conditions apply to all S-M	erature of 20°C onovettes with the e	exception of 8 mr	m diameter (S-Mor	novettes for pedia	atrics)

* If a citrate sample is collected first using a Safety-Multifly needle, it is recommended to first draw a discard tube.

> ** For S-Monovettes prepared with gel, we recommend using swing-out rotors only. To convert g-force to speed/min, use the centrifugation calculator at https://www.sarstedt.com/service/zentrifugation/

S-Monovette®	2000 x g	2500 x g	3000 x g*	3500 x g*	4000 x g*	
Serum	10 min	10 min	6 min	4 min	4 min	
Serum Gel	15 min	10 min	4 min	4 min	4 min	
Li-Heparin	10 min	10 min	7 min	7 min	7 min	
Li-Heparin-Gel	15 min	15 min	10 min	7 min	7 min	
Li-Heparin-Gel*	8 min	7 min	5 min	4 min	4 min	
EDTA	n.v.	n.v.	7 min	7 min 6 min		
EDTA Gel	15 min	10 min	10 min	7 min	7 min	
Citrate	9 min	8 min	7 min 6 min		5 min	
Fluoride	9 min	8 min	7 min	6 min	5 min	
Citrate PBM 1.8 ml Rotor Ø > 17 cm	9 min	8 min	7 min	6 min	5 min	
Citrate PBM 1.8 ml Rotor Ø > 9 cm to < 17 cm	n.v.	n.v.	10 min	n.v.	n.v.	
n.v. = not validated						

S-Monovette[®]

Safety starts with choosing the right system



S-Monovette[®]

Handling instructions

Vacuum technique

Barcode labeling & mixing

Aspiration technique



1. The safety needle is assembled with the S-Monovette® immediately prior to blood collection. This is followed by the puncture.



1. We recommend filling the first S-Monovette® using the aspiration technique in order to start the blood collection gently. By retracting and locking the plunger into place at the bottom of the S-Monovette® (click). a fresh vacuum is produced directly before the blood is collected. The plunger is broken off (snap)

2. The evacuated S-Monovette® is connected and filled with the safety needle / Safety-Multifly® needle already in the vein. For multiple blood collections, repeat this procedure accordingly.



3. Once blood collection is complete, the last S-Monovette® is released from the safety needle and the needle is withdrawn from the vein.

2. Slow retraction of the plunger results in

a gentle blood flow. For multiple blood

collections, additional S-Monovettes

are connected to the safety needle

and blood samples are collected as

described above.





3. Once blood collection is complete, the last S-Monovette® is released from the safety needle / Safety-Multifly® needle and the needle is withdrawn from the vein.

Combination options

Store



a. If, in exceptional cases, blood is to be collected with a Luer-Monovette® (e.g. Blood Gas-Monovette®), the membrane adapter (A) can be used. b. With the multi-adapter (B), the S-Monovette[®] can be used for blood collection through Luer connections (3-way stopcock, butterfly, etc.). c. For veins in poor condition, the Safety-Multifly® needle (C) with integrated multi-adapter can be used.

Handling the S-Monovette® Serum/Serum-Gel

To achieve a better serum yield, it is essential to observe the following after blood collection with the S-Monovette® Serum/Serum Gel:



During the clotting phase (the first 30 minutes After blood collection: after blood collection), the S-Monovettes must be stored in an upright position to ensure a distinct S-Monovettes separating layer after centrifugation and to avoid upright for 30 min irregular clot shapes.

Affix the barcode label along the barcode line.



Carefully inverting the S-Monovettes prepared with anticoagulants prevents clot formation:





4. For safety during transport and centrifugation, the piston is locked in place at the bottom of the S-Monovette® (click) and the plunger is broken off (snap).



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SARSTEDT