








Recommendations on Centrifugation

Order of draw*				
Blood culture (if required)				
	Citrate 1:10	10 min.	1,800 x g	22°C
	Citrate 1:5			
	Serum	10 min.	2,000 x g	20°C
	Serum Gel	10 min.	2,500 x g	20°C
	Heparin	10 min.	2,000 x g	20°C
	Heparin Gel	10 min. or 15 min.	3,000 x g 2,500 x g	20°C 20°C
	EDTA	10 min.	2,000 x g	20°C
	EDTA K ₂ Gel	10 min.	2,500 x g	20°C
	Fluoride	10 min.	2,000 x g	20°C

Technical modifications reserved

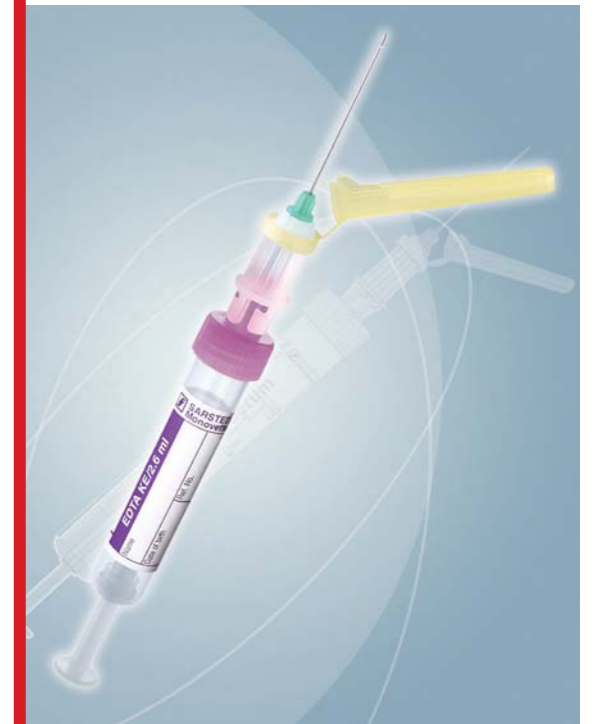
* From: 'CLSI H3-A6 (Procedures for the Collection of Diagnostic Blood Specimens by Venipuncture, Approved Standard - Sixth Edition)'

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Notes	Caps	Preparations and Applications
	 Serum	Clinical Chemistry S-Monovettes contain beads coated with a clotting activator (silicate). As a rule, this clotting additive enables the blood to clot within 20 to 30 minutes and the sample can be centrifuged. The beads form a layer between the blood clot and serum during centrifugation.
	 Serum Gel	Clinical Chemistry In addition to the beads, this S-Monovette® contains a polyacrylic ester gel that, due to its density, forms a stable separating layer between the blood clot and the serum during centrifugation and serves as a barrier during sample transport and storage. Compliance with the recommended storage conditions will keep most parameters stable for up to 48 hours.
	 Lithium Heparin	Clinical Chemistry Heparin, at an average concentration of 16 I.U./ml blood, is used as an anticoagulant for plasma generation. Heparin is coated onto beads which form a layer between the plasma and the corpuscular components during centrifugation. The function of the plasma gel is identical to serum-gel.
	 Potassium EDTA	Hematology EDTA K ₂ is pre-dosed as a liquid preparation in an average concentration of 1.6 mg EDTA/ml blood. The maximum dilution caused by the liquid preparation is lower than 1%. Although the EDTA preparation may dry during storage, this does not in any way impair its anticoagulant effect. An S-Monovette® with EDTA K ₂ and gel is available for use in molecular virus diagnostics.
	 Fluoride	Glucose The S-Monovette® for glucose determination contains fluoride (1.0 mg/ml blood) as a glycolysis inhibitor and EDTA (1.2 mg/ml blood) as an anticoagulant in a liquid preparation. The glucose concentration in the sample is stabilized for a period of 24 hours.
	 Tri-Sodium Citrate 1:10	Coagulation Citrate, pre-dosed as a 0.106 molar solution (equivalent to 3.2% trisodium citrate), is used for all physiological coagulation studies (e.g. Quick, PTT, TZ, Fibrinogen). A mixing ratio of 1:10 (1 part citrate + 9 parts blood) must be strictly observed.
	 Tri-Sodium Citrate 1:5	ESR Citrate, pre-dosed as a 0.106 trisodium citrate molar solution, is used for ESR determination. A mixing ratio of 1:5 (1 part citrate + 4 parts blood) must be strictly observed. For ESR, we recommend the S-Monovette® Sediplus® system (Westergren method) or the enclosed S-Sedivette® system (modified Westergren method).

S-Monovette®

Safety begins with choosing the right system



Aspiration Principle

- Immediately prior to venous puncture, push S-Monovette® onto Safety-Needle and secure by slightly twisting clockwise (①+②).
- Puncture vein, loosen tourniquet and withdraw plunger slowly. Wait until blood flow stops.
- Remove S-Monovette® from Safety-Needle by slightly twisting counter-clockwise (③+④). Safety-Needle remains in vein.
- For multiple sampling, secure subsequent S-Monovettes onto Safety-Needle and collect further samples as described above.

Completion of blood collection:

Remember: Detach S-Monovette® (③+④) first, then withdraw Safety-Needle. Place the needle protector on a stable, flat surface and slightly press the needle downwards until it locks into the needle protector with a noticeable and audible "Click".

- Gently invert several times to mix sample(s) with anticoagulant(s)!
- For transportation and centrifugation, lock piston into S-Monovette® base and break off plunger (⑤).

Vacuum Principle

Prior to blood collection, the S-Monovette® Safety-Needle must already be in the vein. Either puncture the vein directly with the Safety-Needle or collect the first sample using the aspiration principle – then apply the vacuum principle.

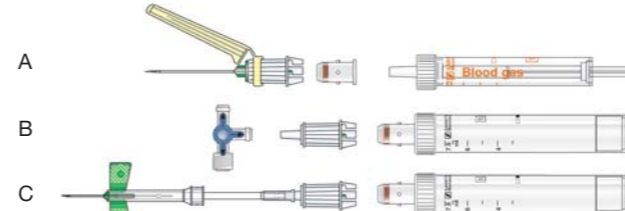
- Prior to blood collection, lock piston into S-Monovette® base. Once secured, the plunger must be snapped off (①).
- Push S-Monovette® onto the Safety-Needle and secure by slightly twisting clockwise (②+③). Loosen tourniquet.
- Wait until blood flow stops.
- Remove S-Monovette® from Safety-Needle by slightly twisting counter-clockwise (④+⑤). Safety-Needle remains in vein.
- For multiple sampling, secure subsequent S-Monovettes onto Safety-Needle and collect samples as described above.

Completion of blood collection:

Remember: Detach S-Monovette® (④+⑤) first, then withdraw Safety-Needle from the vein. Place the needle protector on a stable, flat surface and slightly press the needle downwards until it locks into the needle protector with a noticeable and audible "Click".

- Gently invert several times to mix sample(s) with anticoagulant(s)!

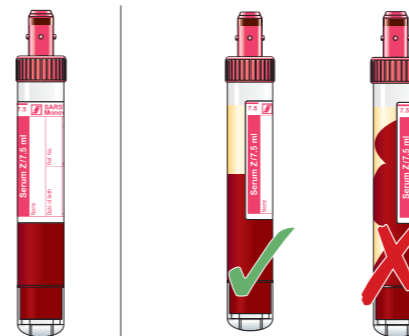
Special Applications



- The membrane adapter (A) can be used if, in exceptional cases, blood is to be collected with a Luer-Monovette® (e.g. blood gas).
- The S-Monovette® can be used for blood collection from Luer connections (3-way-tap, Butterfly, etc.) by means of the multi adapter (B).
- For difficult vein conditions, we recommend to use the Safety-Multify® (C) with integral multi adapter.

User Guide S-Monovette® Serum/Serum Gel

Make sure to observe the following instructions for optimal serum yield after blood collection using the S-Monovette® Serum:

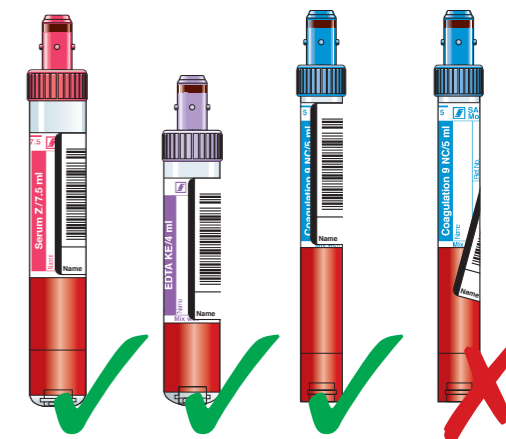


After blood collection:
Store S-Monovette® in an upright position for 30 minutes

During the clotting phase (i.e. the initial 30 minutes after blood collection), it is essential to store the S-Monovette® in an upright position to ensure a distinct separating layer after centrifugation!

Barcode Labelling & Mixing

Carefully align barcode labels:



Careful mixing of the S-Monovettes prepared with anticoagulants prevents the blood from clotting:

