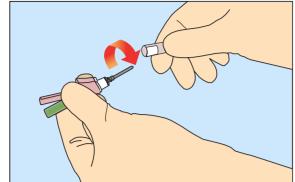
BD Diagnostics - Preanalytical Systems

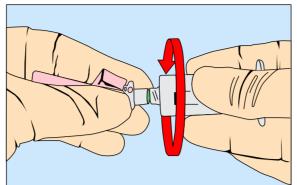
Blood collection with BD Vacutainer® system



- 1. Patient Identification:
- Greet the patient.
- Identify Yourself. - Confirm the patient's information.



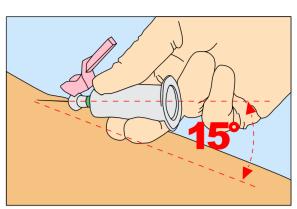
2. Visually check the sterility seal on the needle. If the seal is broken. DO NOT USE. If the seal is unbroken, twist to break and dispose of the white cover immediately.



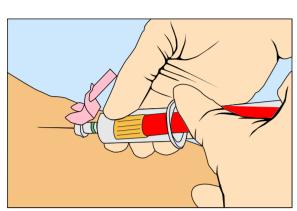
3. Screw BD Vacutainer® Eclipse™ Needle into a BD Vacutainer® One Use Holder. Flip the lavender safety shield back. Remove the coloured needle shield.



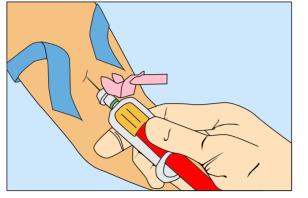
4. Apply Tourniquet 7.5 to 10 cm above site 5. The bevel of the BD Vacutainer® Eclipse™ of the intended venepuncture. Disinfect the site. Allow to air dry.



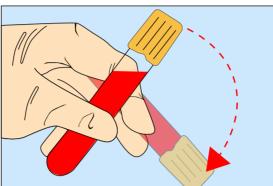
Needle is aligned with the safety shield. Perform venepuncture in the usual manner with the arm of the patient in the downward position.



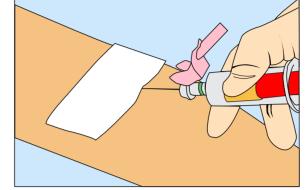
6. Introduce the BD Vacutainer® tube into the holder. Placing your forefinger and middle finger on the flange of the holder and the thumb on the bottom of the tube, push the tube to the end of the holder, puncturing the diaphragm of the



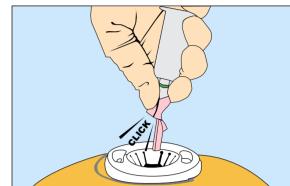
7. Remove the tourniquet as soon as blood begins to flow into the tube. The tourniquet should be applied for no longer than one minute before venepuncture is carried out.



8. Invert all tubes minimum 5 to 6 times. except citrate tubes (3 to 4 times) to ensure adequate mixing. Do not shake. Vigorous mixing may cause hemolysis.



9. As the needle is removed from the arm gently cover the punctured site with a gauze pad. Pressure on the site should be maintained for a few minutes to help seal the vein wall and prevent hematoma.



10. Activate the safety shield until an audible 'click' is heard. Dispose the needle and holder as a single unit into an approved sharps container.

BD Vacutainer® Order of Draw & Mixing Guidelines

All BD Vacutainer® tubes require immediate mixing following collection

| Colour Code | Tube Type | Determinations | Inversions |
|----------------|-------------------------|--|------------|
| Blue Purple | Blood Culture | Aerobic followed by anaerobic | 8-10 Times |
| Light Blue | Sodium Citrate | For coagulation determinations on plasma specimens | 3-4 Times |
| Black | ESR | For erythrocyte sedimentation rate (ESR) determinations | 8-10 Times |
| Red | Serum | For serum determinations in chemistry | 5-6 Times |
| Gold | SST™ II <i>Advanc</i> e | For serum determinations in chemistry - with gel seperator | 5-6 Times |
| Orange | RST | For serum determinations in chemistry - with Thrombin based clotting agent and gel separator | 5-6 Times |
| Green | Heparin | For plasma determinations in chemistry | 8-10 Times |
| Light Green | PST TM II | For plasma determinations in chemistry with gel separator | 8-10 Times |
| Lavender | EDTA | For whole blood hematology determinations | 8-10 Times |
| Pink | Cross Match | Crossmatch tubes for blood transfusion patients | 8-10 Times |
| Grey | NaF/NaEDTA | For glucose determinations | 8-10 Times |
| Royal Blue | Trace Element | For trace element, toxicology and nutrient determinations | 8-10 Times |

Insufficient mixing can result in inaccurate test results and the need to re-draw

REF: (CLSI H3-A6)



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