Separation of serum

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Serum clear yellow fluid includes all <u>proteins</u> not used in <u>blood clotting</u> (coagulation) and all the <u>electrolytes</u>, <u>antibodies</u>, <u>antigens</u>, <u>hormones</u>, & any <u>exogenous</u> substances (e.g., drugs and microorganisms).

* Preparation of serum sample from whole blood

Procedure:

- 1. Label the tubes, including the unique patient identification number, sex & age.
- 2. Place a tourniquet above the venepuncture site, disinfect the venepuncture site with 70% isopropyl alcohol (an alcohol swab) or 10% polyvidone iodine palpate and locate the vein.
- **3.** Withdraw 3–5 ml, applied the collect blood by venesection and allow to clot in a glass container without anticoagulant.
- **4.** Once the clot has formed, separate the serum from the clotted cells by centrifugation at (**1000 rpm**) for **15** min at 4°C., or directly centrifugated at least **5** minutes at (**3000 rpm**).
- **5.** Save the serum (straw-coloured supernatant) and discard the cell debris, transfer to a suitable container, then proceed to isolation of the immunoglobulins.

Storing of serum:

- Blood serum samples should be frozen at -70 °C for PCR
- At -20 $^{\circ}$ C or lower for antibody determination but they can be stored at +4 $^{\circ}$ C for approximately one week.