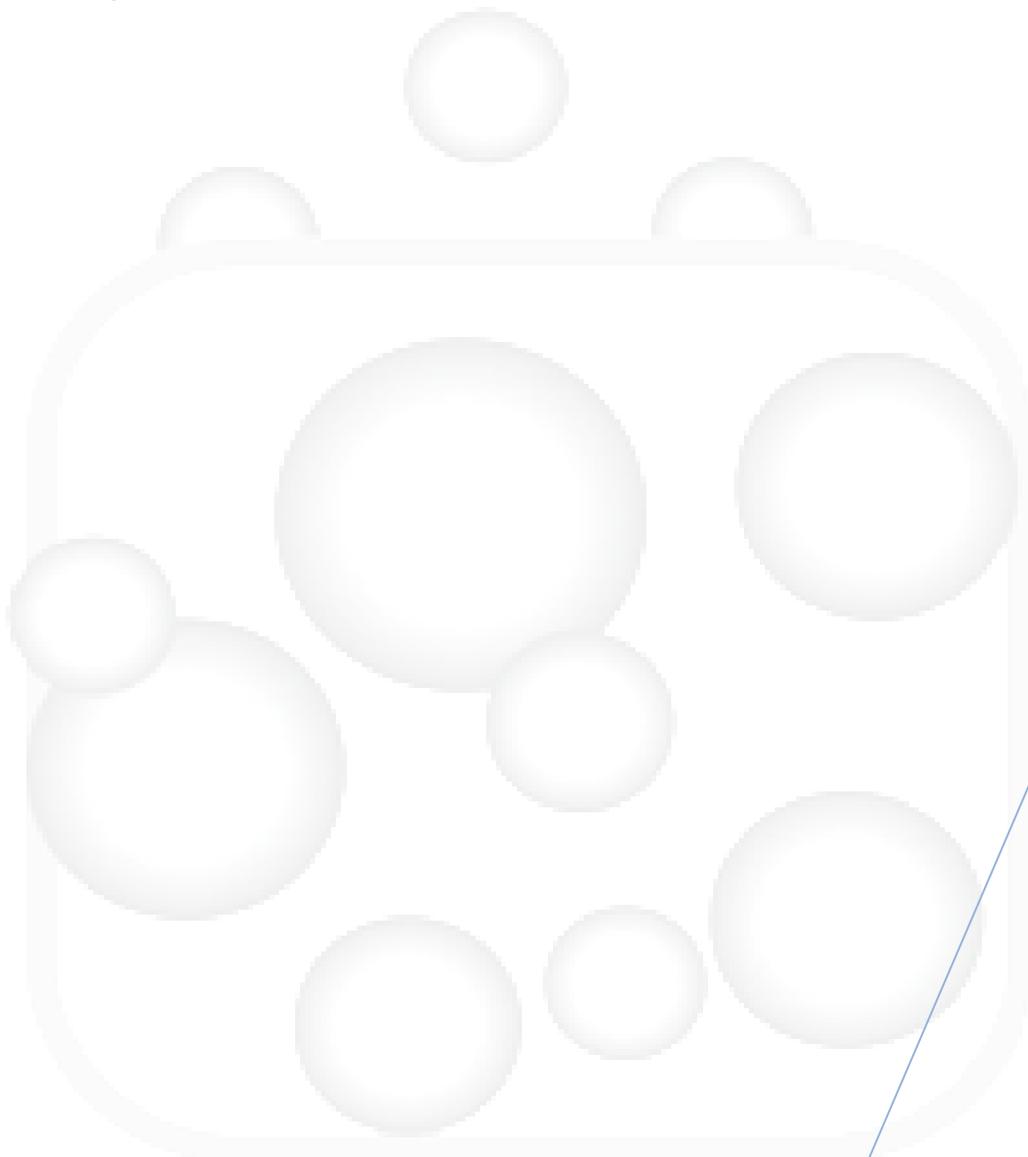




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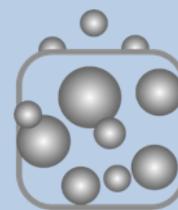
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**DNA extraction from Blood**  
PSR

Dr. Zonuoni. S  
6/12/2016



RCPN

## DNA extraction from Blood

### Solutions

- Red blood cells lysing buffer (RLB): 2M Tris-HCl pH 7.6, 1M MgCl<sub>2</sub>, 3M NaCl.
- White blood cells lysing buffer (WLB): 2M Tris pH 7.6, 0.4 M disodium salt of ethylenediaminetetra acetic acid (Na<sub>2</sub>EDTA), pH 8, 3M NaCl, 2% cetyl trimethyl ammonium bromide (CTAB)
- Chloroform: isoamyl alcohol 24:1

## Detailed procedure

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1. Transfer 2 mL of fresh blood sample into a 2 mL tube.
  2. Centrifuge the sample at 6000 rpm at 4°C for 10 min.
  3. Aspirate the plasma without touching the leukocyte layer.
  4. Add 1 mL RLB and mix gently.
  5. Centrifuge the sample for 5 min at 3000 rpm.
  6. Remove the supernatant.
- Note:** If red blood cells still remain, re-suspend the cellular pellet in 1 mL of RLB and repeat steps 4-6 for 3 or 4 times until only the white pellet appears.
7. Add 1 ml of WLB and mix with white blood cells.
  8. Incubate the tube at 65 °C for 30 min.
  9. Centrifuge the tube at 12,000 rpm for 5 min and transfer the supernatant to a new clean tube and discard the pellet.
  10. Add an equal volume of Chloroform-Isoamylalcohol solution to supernatant.
  11. Centrifuge the tube at 12000 rpm for 8 min and transfer the supernatant to a new tube.
  12. Add an equal volume of chilled Isopropanol.
  13. Keep the sample in -20 °C for 30 min.
  14. Centrifuge the tube at 4 °C, 12000 rpm for 10 min.

15. Discard the supernatant and add 300  $\mu$ L of chilled 90% Ethanol. Centrifuge the tube at 4  $^{\circ}$ C, 12000 rpm for 5 min.
16. Repeat the 15 and 16 steps with chilled 70% Ethanol.
17. Discard the supernatant and let pellet to be dried at room temperature.
18. Dissolve the pellet in 100  $\mu$ L of TE buffer or ddH<sub>2</sub>O and store DNA solution at -20  $^{\circ}$ C.

**Notification:**

- For frozen blood samples start the procedure from step 4.
- For clotted blood samples, start the procedure from step 7 by adding 1 mL of WLB and incubating at 65  $^{\circ}$ C for 1 hour (shaken every 10 min). Then continue by step 10.

**Reference**

Samadi Shams, S., Zununi Vahed, S., Soltanzad, F., Kafil, V., Barzegari, A., Atashpaz, S., *et al.* (2011) Highly effective DNA extraction method from fresh, frozen, dried and clotted blood samples. *Bioimpacts* **1**: 183-7.

Good luck!



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Document arrangement	Mohanna Osali, Rahimeh Mousavi
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Author / translator	Dr. Zonuoni. Sepideh