























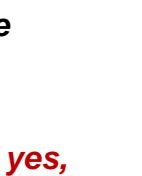














# GINA 500 – Depletion of Human DNA from EDTA Blood

<p><b>Switch on the heating block and set temperature to 100 °C!</b></p>		<p><b>Spin down the IPC tube before use</b></p> 	<input type="checkbox"/>
 <p><b>Vortex Blood Sample</b> 0 – 500µl</p>	<p><b>Add 20µl IPC (Optional)</b> <b>Add sample / 20µl EPC</b></p>	 <p><b>LE Solution 1400µl (yellow !)</b></p>	<input type="checkbox"/>
	<p><b>Vortex for 5 seconds or invert repeatedly and wait for about <u>2min</u> at 18°- 25°C</b> <b>Check for homogeneity</b></p>		<input type="checkbox"/>
	<p><b>Centrifuge for 5 Minutes with “Soft ramping”, 9k – 11k [g]</b></p>	 <p><b>Centrifugation: 5 Minute, 9k -11k [g]</b></p>	<input type="checkbox"/>
	<p><b>Decant Supernatant</b> <b>Use a pipette to remove the remaining supernatant</b></p>		<input type="checkbox"/>
 <p><b>NA Solution 200µl (red !)</b></p>	<p><b>Invert to mix</b></p> 		<input type="checkbox"/>
	<p><b>Vortex for 5 Seconds</b></p>		<input type="checkbox"/>
 <p><b>T Solution 400µl (green !)</b></p>	<p><b>Incubate for 10 Minutes at 100°C</b></p>		<input type="checkbox"/>
	<p><b>Invert to mix</b></p> 		<input type="checkbox"/>

**Storage possibilities:**

**until 24 Hours: 2° to 4°C**  
**> 24 Hours: -25° to -18°C**

# GINA 500 – DNA Purification

 <p>The total volume from the enriched solution, 600µl</p>	 →  <p>“Column “ in “Collection Tube”</p>	 <p>Centrifugation: 1 Minute, 9k -11k [g]</p>	 <p>Discard flow-through in “Collection T.”</p>	
 <p>Wash Buffer BW 500 µL</p>	 →  <p>“Column “ in “Collection Tube”</p>	 <p>Centrifugation: 1 Minute, 9k -11k [g]</p>	 <p>Discard flow-through in “Collection T.”</p>	
 <p>Wash Buffer B5 600 µL</p>	 →  <p>“Column “ in “Collection Tube”</p>	 <p>Centrifugation: 1 Minute, 9k -11k [g]</p>	 <p>Discard flow-through in “Collection T.”</p>	
<p>1x Dry Silica Membrane</p> <p><b>Check if there is liquid in/under the column (if yes, repeat the step)</b></p>	 →  <p>“Column “ in “Collection Tube”</p>	 <p>Centrifugation: 1 minute, 9k -11k [g]</p>	 <p>Discard the “Collection T.”</p>	
 <p>Elution Buffer BE 100-150 µL</p> <p><b>Check the elution volume (repeat centrifugation if necessary)</b></p>	 →  <p>“Column “ in “Elution Tube”</p> <p>1 Minute room temp. incubation</p>	 <p>Centrifugation: 1 Minute, 9k -11k [g]</p>	 <p>Eluate in Elution Tube</p>	
 <p>Eluate in Elution Tube</p>	<p>Heat at 100°C for 3 Minutes (leave the column´s lid open)</p>	 	<p><b>Resuspend eluate before PCR usage !!!</b></p>	

Storage possibilities:

until 24 Hours: 2° to 4°C  
> 24 Hours: -25° to -18°C

\*As an alternative to emptying and reusing the collection tube, a new tube can also be used. (additional tubes will be required!)