









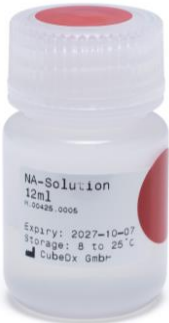




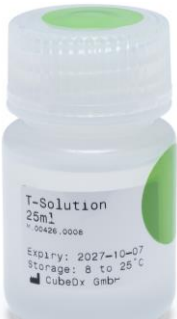




GINA 500 – Depletion of Human DNA from EDTA Blood





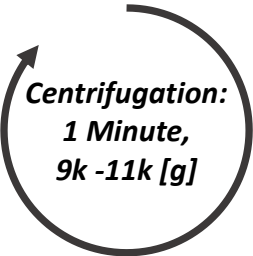


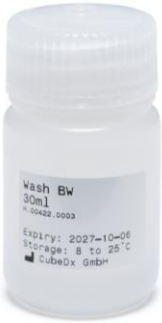



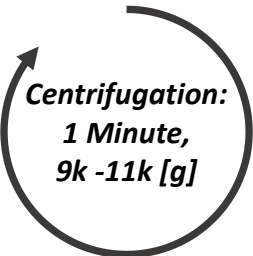






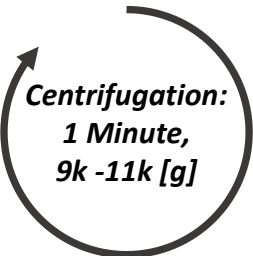




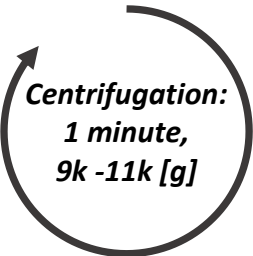







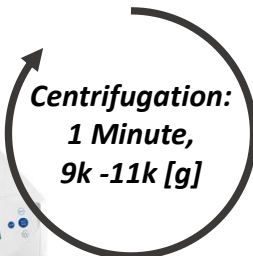







<div>Switch on the heating block and set temperature to 100 °C!</div>		<div></div>	<div>Spin down the IPC/EPC tube before use</div>	<div></div>	<div><input type="checkbox"/></div>
<div><div>Vortex Blood Sample 0 – 500µl</div></div>	<div><div>+</div><div><div>20µl IPC (Optional) Pipette and mix the sample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

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>Check if there is liquid in/under the column (if yes, repeat the step)</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>Check the elution volume (repeat centrifugation if necessary)</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>Resuspend eluate before PCR usage !!!</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!)