

# GINA 500 – Depletion of Human DNA from EDTA Blood

Switch on the heating block and set temperature to 100 °C!



Spin down the IPC/EPC tube before use



Vortex Blood Sample

0 – 500µl



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)



LE Solution 1400µl (yellow !)



Vortex for 5 seconds or invert repeatedly and wait for about 2min at 18°- 25°C  
Check for homogeneity



Centrifuge for 5 Minutes with “Soft ramping”, 9k – 11k [g]

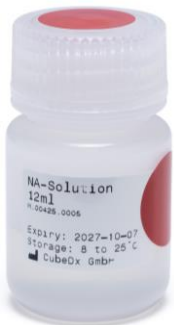
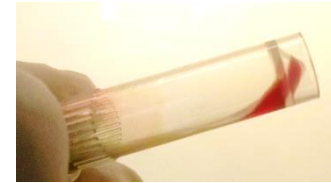


Centrifugation: 5 Minute, 9k -11k [g]



Decant Supernatant

Use a pipette to remove the remaining supernatant



NA Solution 200µl (red !)



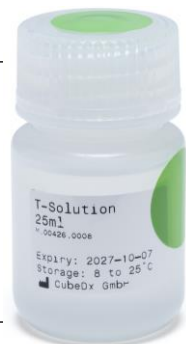
Invert to mix



Vortex for 5 Seconds



Incubate for 10 Minutes at 100°C



T Solution 400µl (green !)


















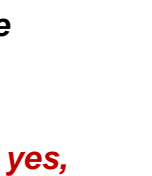





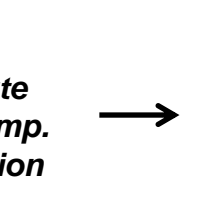
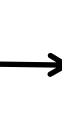





Invert to mix



Storage possibilities:

until 24 Hours: 4° to 8°C  
> 24 Hours: -15° to -25°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><i>Check if there is liquid in/under the column (if yes, repeat the step)</i></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><i>Check the elution volume (repeat centrifugation if necessary)</i></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><i>Resuspend eluate before PCR usage !!!</i></p>	

Storage possibilities:

until 24 Hours: 4° to 8°C  
> 24 Hours: -15° to -25°C

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