

# GINA 500 – Depletion of Human DNA from EDTA Blood

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



Spin down the IPC tube before use



Vortex Blood Sample

0 – 500µl



Add 20µl IPC (Optional)

Add sample / 20µl EPC



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 !)








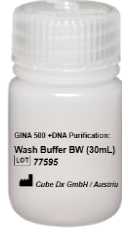






















Invert to mix



Storage possibilities:

until 24 Hours: 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><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>Dispose of the column, and close the elution tube.</p> <p><i>Resuspend eluate before PCR usage !!!</i></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!)