




Patient data			
Name		Mrs. G SADHYA	
Patient ID		0012508090263	
Birthday		09-08-2002	
Sample ID		B3186081	
Age at sample date		23.0	
Sample Date		09-08-2025	
Gestational age		12 + 3	
Correction factors			
Fetuses	1	IVF	no
Weight	33	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.83 mIU/mL	0.69	
fb-hCG	42.41 ng/mL	0.80	
Risks at sampling date			
Age risk		1:1027	
Biochemical T21 risk		1:4376	
Combined trisomy 21 risk		<1:10000	
Trisomy 13/18 + NT		<1:10000	
Gestational age			12 + 3
Method			CRL Robinson
Scan date			09-08-2025
Crown rump length in mm			61.9
Nuchal translucency MoM			1.00
Nasal bone			present
Sonographer			N A
Qualifications in measuring NT			MD
Risk			Trisomy 21
			<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</p> <p>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that risk calculations are statistical approaches and have no diagnostic value!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>
Trisomy 13/18 + NT			
<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			

 below cut off
  Below Cut Off, but above Age Risk
  above cut off