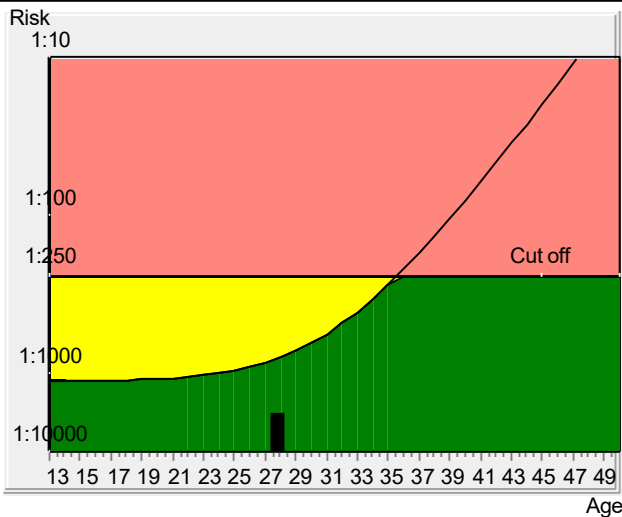


Patient data			
Name	MRS. BHAWNA		Patient ID
Birthday	28/04/1997		Sample ID
Age at sample date	27.8		Sample Date
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	59	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		no	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	6.8 mIU/ml	1.37	13 + 0
fb-hCG	71.4 ng/ml	1.67	Method
Risks at sampling date			CRL Robinson
Age risk	1:821		Scan date
Biochemical T21 risk	1:2858		19/02/2025
Combined trisomy 21 risk	<1:10000		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		68.9
			Nuchal translucency MoM
			0.75
			Nasal bone
			present
			Sonographer
			DR. (MRS.) NEERJA CHOPRA
			Qualifications in measuring NT
			MD
Trisomy 21			
<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></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><b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b></p>			




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 Sign of Physician