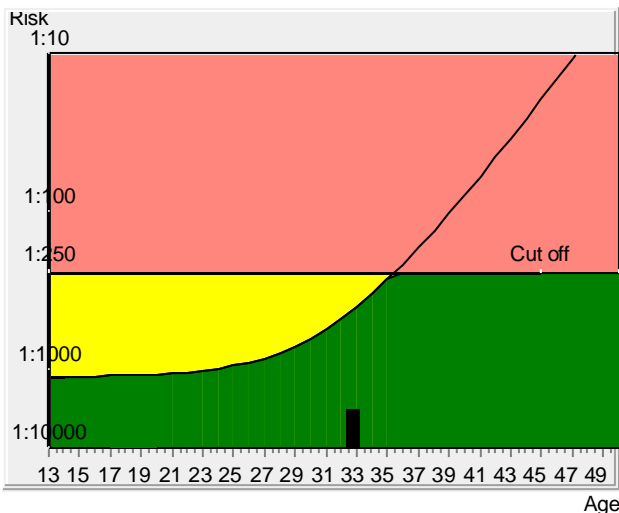


Patient data			
Name	MRS. SWAPNIL KHANDELWAL		Patient ID
Birthday	30/04/1992	Sample ID	2501028072/NOD
Age at sample date	32.7	Sample Date	18/01/2025
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	60.7	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies	no		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	7.1 mIU/ml	1.76	Gestational age 12 + 4
fb-hCG	90.3 ng/ml	2.03	Method CRL Robinson
Risks at sampling date			Scan date 18/01/2025
Age risk	1:429		Crown rump length in mm 63.2
Biochemical T21 risk	1:1435		Nuchal translucency MoM 0.80
Combined trisomy 21 risk	1:6844		Nasal bone present
Trisomy 13/18 + NT	<1:10000		Sonographer DR. (MRS.) NEERJA CHOPRA
			Qualifications in measuring NT MD
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 6844 women with the same data, there is one woman with a trisomy 21 pregnancy and 6843 women with not affected pregnancies.</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>			



Sign of Physician