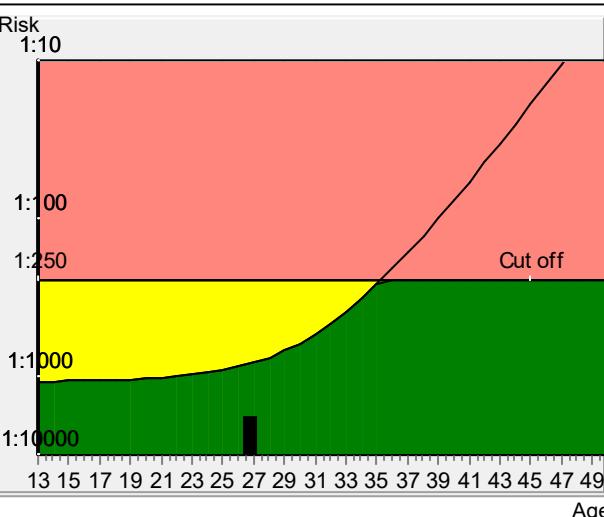


Patient data			
Name	Mrs. PRIYANKA MEENA	Patient ID	0382305110082
Birthday	16/07/96	Sample ID	23938421
Age at sample date	26.8	Sample Date	11/05/23
Gestational age	11 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	58.9	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	3.03 mIU/mL	1.14	Gestational age 11 + 4
fb-hCG	52.23 ng/mL	1.01	Method CRL Robinson
Risks at sampling date			
Age risk	1:840		Scan date 11/05/23
Biochemical T21 risk	1:6726		Crown rump length in mm 50
Combined trisomy 21 risk	<1:10000		Nuchal translucency MoM 0.74
Trisomy 13/18 + NT	<1:10000		Nasal bone unknown
Sonographer NA			
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 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 held 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

 below cut off

 Below Cut Off, but above Age Risk

 above cut off