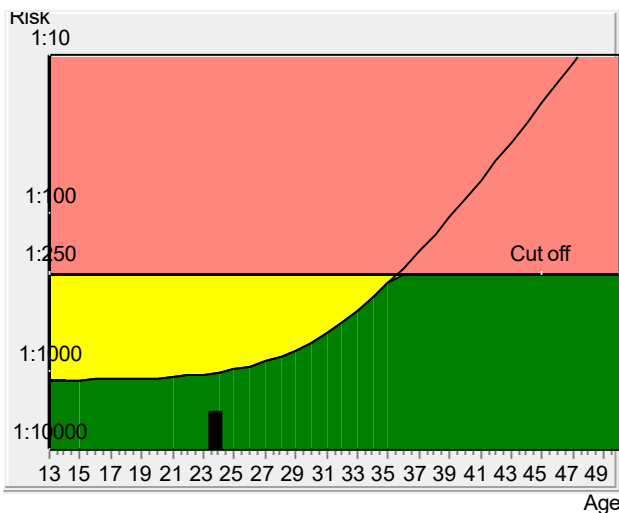


Patient data			
Name	MRS. SANJANA		Patient ID
Birthday	7/05/2001		Sample ID
Age at sample date	23.8		Sample Date
Gestational age	13 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	66	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		no	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.91 mIU/ml	0.76	13 + 2
fb-hCG	51.4 ng/ml	1.31	Method
			CRL Robinson
			Scan date
			28/02/2025
Risks at sampling date			Crown rump length in mm
			73
Age risk	1:1036		Nuchal translucency MoM
Biochemical T21 risk	1:1891		0.72
Combined trisomy 21 risk	<1:10000		Nasal bone
Trisomy 13/18 + NT	<1:10000		present
			Sonographer
			..
			Qualifications in measuring NT
			..
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>			



Sign of Physician