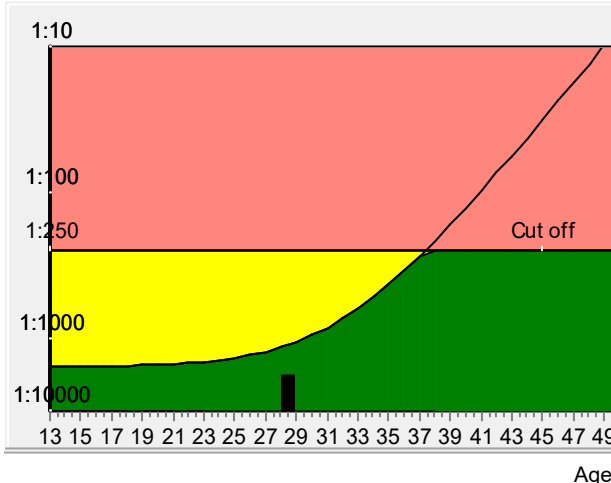


Date of report: **03/01/25**
Prisca 5.1.0.17

NA

Patient data				Ultrasound data	
Name		Mrs. PRIYA RANDHIR		Gestational age	13 + 2
Birthday		18/01/97		Method	CRL measurements
Age at delivery		28.4		Crown rump length in mm	73.1
Patient ID		0662412310286		Date	20/12/24
Previous trisomy 21 pregnancies		unknown		Nuchal translucency MoM	0.88
				Nuchal translucency	1.59 mm
Correction factors				Nasal bone	unknown
Fetuses	1	diabetes	no	Sonographer	NA
Weight	44	Origin	Asian	Qualifications in measuring NT	MD
Smoker	no	IVF	no		
Biochemical data				Risks at term	
Sample Date		31/12/24		Age risk	1:1125
Gestational age at sample date		14 + 6		Trisomy 21 risk	1:7937
Parameter	Value	Corr. MoMs		Combined trisomy 21 risk	<1:10000
AFP	43.66 ng/mL	1.23		Trisomy 18 risk	<1:10000
HCG	35632.54 mIU/mL	0.76			
uE3	0.35 ng/mL	0.91			
Risk				Trisomy 21	
				The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.	
				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. 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! The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)). The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!	
Trisomy 18				Neural tube defects	
The calculated risk for trisomy 18 (with nuchal translucency) is < 1:10000, which represents a low risk.				The corrected MoM AFP (1.23) is located in the low risk area for neural tube defects.	

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