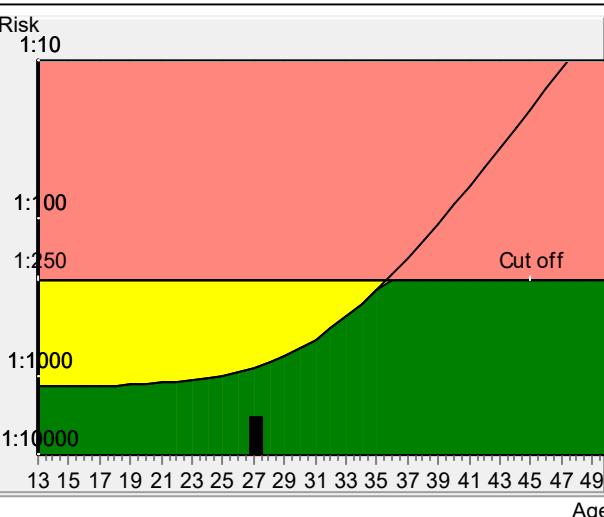


Prisca 5.1.0.17

Date of report: 04/01/26

Patient data								
Name	Mrs. POOJA S THOSAR	Patient ID	0662601030060					
Birthday	01/12/98	Sample ID	B4180499					
Age at sample date	27.1	Sample Date	02/01/26					
Gestational age	13 + 6							
Correction factors								
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown			
Weight	54	diabetes	no					
Smoker	no	Origin	Asian					
Biochemical data								
Parameter	Value	Corr. MoM	Ultrasound data					
PAPP-A	4.01 mIU/mL	0.55	Gestational age 13 + 4					
fb-hCG	39.47 ng/mL	1.31	Method CRL Robinson					
Risks at sampling date								
Age risk	1:892		Scan date 31/12/25					
Biochemical T21 risk	1:732		Crown rump length in mm 78.3					
Combined trisomy 21 risk	1:4423		Nuchal translucency MoM 0.74					
Trisomy 13/18 + NT	<1:10000		Nasal bone present					
			Sonographer NA					
			Qualifications in measuring NT Sonographer					
Trisomy 21								
<b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b>								
After the result of the Trisomy 21 test (with NT) it is expected that among 4423 women with the same data, there is one woman with a trisomy 21 pregnancy and 4422 women with not affected pregnancies.								
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 held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!								
Trisomy 13/18 + NT								
<b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b>								
								

Sign of Physician

 below cut off

 Below Cut Off, but above Age Risk

 above cut off