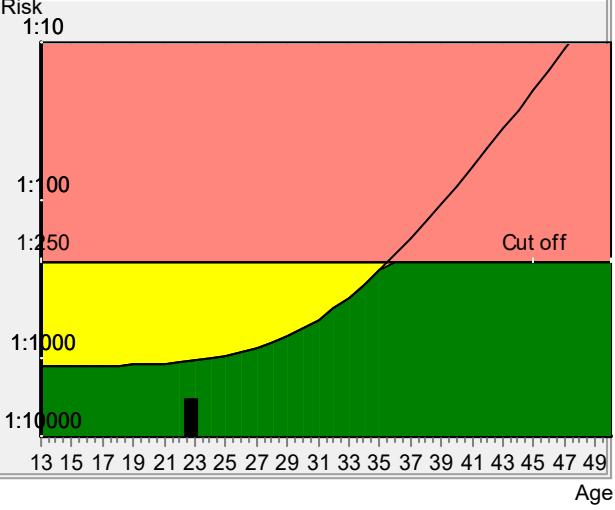


Prisca 5.1.0.17

Date of report: 08/11/25

AYESHA JAHAN

Patient data			
Name	Mrs. SANIYA KASADI	Patient ID	0012511070362
Birthday	31/01/03	Sample ID	B3286509
Age at sample date	22.8	Sample Date	07/11/25
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	74	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	5.23 mIU/mL	1.41	Gestational age 12 + 6
fb-hCG	36.85 ng/mL	1.06	Method CRL Robinson
Risks at sampling date			
Age risk	1:1053		Scan date 06/11/25
Biochemical T21 risk	<1:10000		Crown rump length in mm 67
Combined trisomy 21 risk	<1:10000		Nuchal translucency MoM 1.12
Trisomy 13/18 + NT	<1:10000		Nasal bone present
Sonographer N A			
Qualifications in measuring NT MD			
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 held responsible for their impact on the risk assessment! Calculated risks have no diagnostic value!			
			
Trisomy 13/18 + NT			
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			

Sign of Physician

 below cut off

 Below Cut Off, but above Age Risk

 above cut off