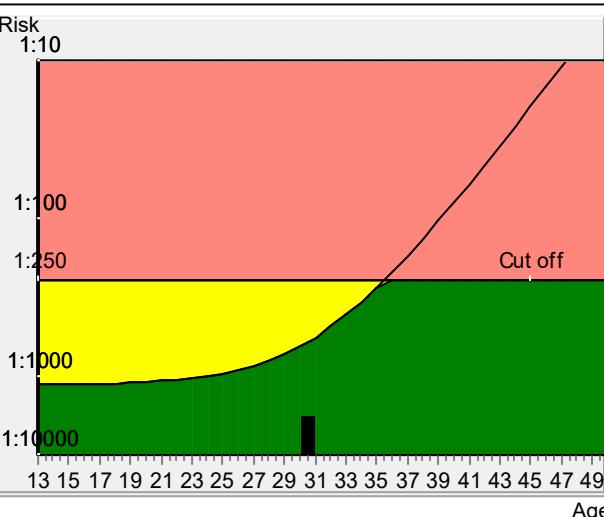


SAGEPATH LABS PVT LTD.,

Prisca

5.1.0.17

Date of report: 17-08-2024

Patient data			
Name	Mrs. MEGHNA	Patient ID	0782408150010
Birthday	24-01-1994	Sample ID	A0401419
Age at sample date	30.6	Sample Date	15-08-2024
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	70	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	9.87 mIU/mL	1.99	Gestational age 12 + 6
fb-hCG	40.46 ng/mL	0.92	Method CRL Robinson
Risks at sampling date			
Age risk	1:608		Scan date 15-08-2024
Biochemical T21 risk	<1:10000		Crown rump length in mm 68
Combined trisomy 21 risk	<1:10000		Nuchal translucency MoM 0.93
Trisomy 13/18 + NT	<1:10000		Nasal bone present
			Sonographer N A
			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