Healthgrow Path Lab
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Prisca 5.2.0.13

Date of report: 11/02/22

Patient data

Name	DEBIKA SARKAR	Patient ID	
Birthday	29/01/97	Sample ID	10127190
Age at sample date	25.0	Sample Date	10/02/22
Gestational age	13 + 0		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight	53.6	diabetes	unknown		
Smoker	unknown	Origin	Asian		

Biochemical data

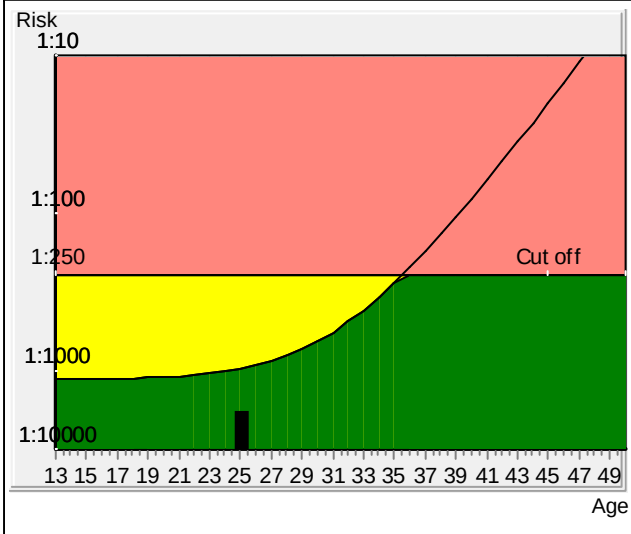
Parameter	Value	Corr. MoM
PAPP-A	3.8 mIU/mL	0.60
fb-hCG	31.34 ng/mL	0.96

Risks at sampling date

Age risk	1:976
Biochemical T21 risk	1:2030
Combined trisomy 21 risk	1:5752
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	13 + 0
Method	CRL Robinson
Scan date	10/02/22
Crown rump length in mm	69.3
Nuchal translucency MoM	1.15
Nasal bone	unknown
Sonographer	DR. MRINMOY RAKSHIT.
Qualifications in measuring NT	M.B.B.S.



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 5752 women with the same data, there is one woman with a trisomy 21 pregnancy and 5751 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 hold 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