Healthgrow Path Lab
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Prisca 5.2.0.13

Date of report: 27/10/21

Patient data

Name	NEHA MITRA	Patient ID	
Birthday	06/03/00	Sample ID	10025450
Age at sample date	21.6	Sample Date	23/10/21
Gestational age	13 + 1		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight		diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

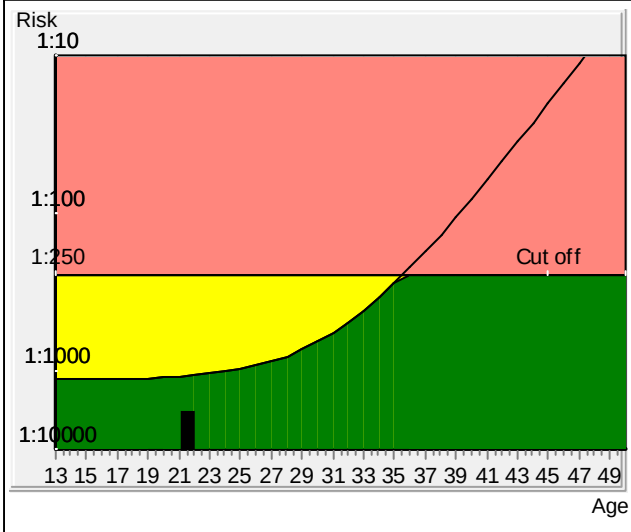
Parameter	Value	Corr. MoM
PAPP-A	6.63 mIU/mL	1.20
fb-hCG	53.92 ng/mL	1.79

Ultrasound data

Gestational age	12 + 1
Method	CRL Robinson
Scan date	16/10/21
Crown rump length in mm	57.4
Nuchal translucency MoM	1.13
Nasal bone	unknown
Sonographer	DR.MRINMOY RAKSHIT.
Qualifications in measuring NT	MBBS

Risks at sampling date

Age risk	1:1084
Biochemical T21 risk	1:2471
Combined trisomy 21 risk	1:6687
Trisomy 13/18 + NT	<1:10000



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 6687 women with the same data, there is one woman with a trisomy 21 pregnancy and 6686 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