Healthgrow Path Lab
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Prisca 5.2.0.13

Date of report: 29/01/22

Patient data

Name	RIYA MALLICK	Patient ID	
Birthday	12/11/02	Sample ID	10114655
Age at sample date	19.2	Sample Date	28/01/22
Gestational age	11 + 1		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight	40	diabetes	unknown		
Smoker	unknown	Origin	Asian		

Biochemical data

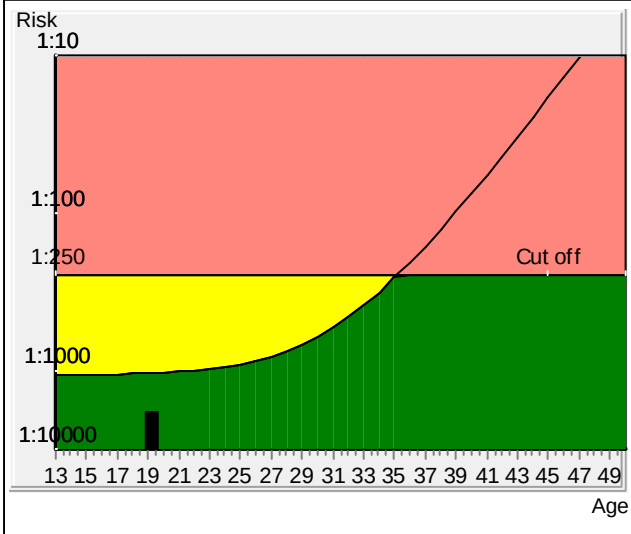
Parameter	Value	Corr. MoM
PAPP-A	1.75 mIU/mL	0.43
fb-hCG	45.04 ng/mL	0.78

Ultrasound data

Gestational age	11 + 0
Method	CRL Robinson
Scan date	27/01/22
Crown rump length in mm	44.1
Nuchal translucency MoM	1.23
Nasal bone	unknown
Sonographer	DR. S. MJUMDAR.
Qualifications in measuring NT	D.M.R.D.

Risks at sampling date

Age risk	1:1042
Biochemical T21 risk	1:1377
Combined trisomy 21 risk	1:3133
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 3133 women with the same data, there is one woman with a trisomy 21 pregnancy and 3132 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