Healthgrow Path Lab
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Prisca 5.2.0.13

Date of report: 02/02/22

Patient data

Name	SRABONI SARDAR	Patient ID	
Birthday	09/08/97	Sample ID	10116492
Age at sample date	24.5	Sample Date	30/01/22
Gestational age	13 + 6		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight	61	diabetes	unknown		
Smoker	unknown	Origin	Asian		

Biochemical data

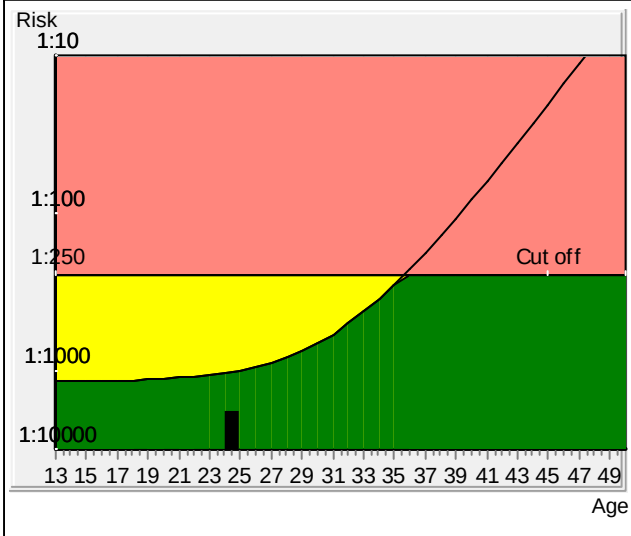
Parameter	Value	Corr. MoM
PAPP-A	1.83 mIU/mL	0.25
fb-hCG	9.01 ng/mL	0.38

Ultrasound data

Gestational age	13 + 5
Method	CRL Robinson
Scan date	29/01/22
Crown rump length in mm	79.9
Nuchal translucency MoM	0.57
Nasal bone	unknown
Sonographer	DR. RAJARSHI AICH.
Qualifications in measuring NT	M.D.

Risks at sampling date

Age risk	1:1025
Biochemical T21 risk	1:1107
Combined trisomy 21 risk	1:6895
Trisomy 13/18 + NT	1:1491



Trisomy 13/18 + NT

The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:1491, which represents a low risk.

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 6895 women with the same data, there is one woman with a trisomy 21 pregnancy and 6894 women with not affected pregnancies.
The free beta HCG level is low.
The PAPP-A level is low.
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!

Sign of Physician

below cut off | Below Cut Off, but above Age Risk | above cut off