Prisca 5.2.0.13

Date of report: 14/07/2023

Patient data

Name	MRS. POOJA	Patient ID	2307023876/NOD
Birthday	26/06/1999	Sample ID	2307023876/NOD
Age at sample date	24.0	Sample Date	13/07/2023
Gestational age	11 + 2		

Correction factors

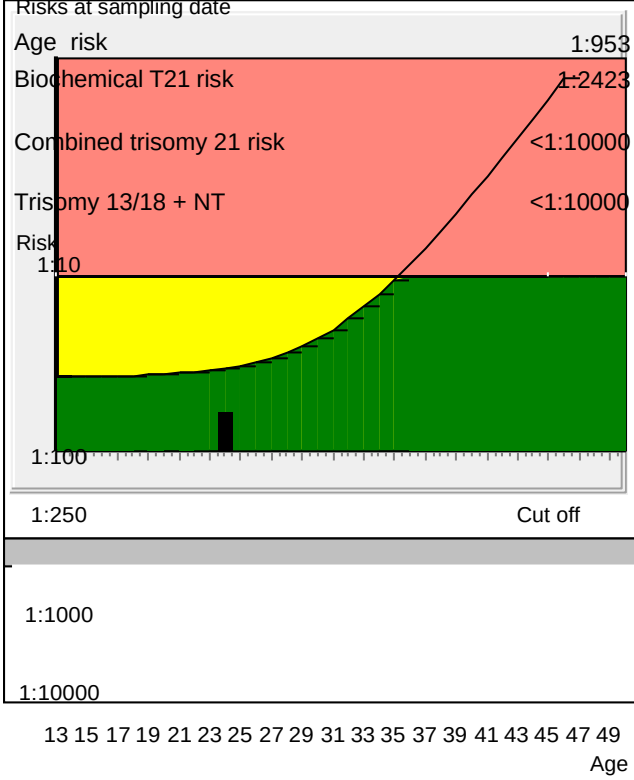
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	no
Weight	41	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

Parameter	Value	Corr. MoM
PAPP-A	2.64 mIU/ml	0.78
fb-hCG	69.2 ng/ml	1.16

Risks at sampling date

Age risk	1:953
Biochemical T21 risk	1:2423
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000



Ultrasound data

Gestational age	11 + 0
Method	CRL Robinson
Scan date	11/07/2023
Crown rump length in mm	44.1
Nuchal translucency MoM	0.82
Nasal bone	present
Sonographer	DR. NITISH UPADHYAYA
Qualifications in measuring NT	M.D

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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.

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.