Prisca   5.1.0.17

Date of report: **18/01/23**

Patient data

Name	MRS SAHELI GHOSH	Patient ID	4053792
Birthday	11/02/86	Sample ID	4053792
Age at delivery	37.4	Sample Date	18/01/23
Gestational age	13 + 3		

Correction factors

Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	69	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

Parameter	Value	Corr. MoM
PAPP-A	3.25 mIU/ml	0.80
fb-hCG	31.2 ng/ml	1.08

Risks at term

Age risk	1:245
Biochemical T21 risk	1:783
Combined trisomy 21 risk	1:4171
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	12 + 5
Method	CRL Robinson
Scan date	13/01/23
Crown rump length in mm	65
Nuchal translucency MoM	0.60
Nasal bone	present
Sonographer	NA
Qualifications in measuring NT	FMF

Risk

13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age

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 4171 women with the same data, there is one woman with a trisomy 21 pregnancy and 4170 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