NA

Patient data

Name	Ms. SANJANA DAS	Patient ID	0692508240232
Birthday	24/09/99	Sample ID	B3584575
Age at sample date	25.9	Sample Date	24/08/25
Gestational age	12 + 1		

Correction factors

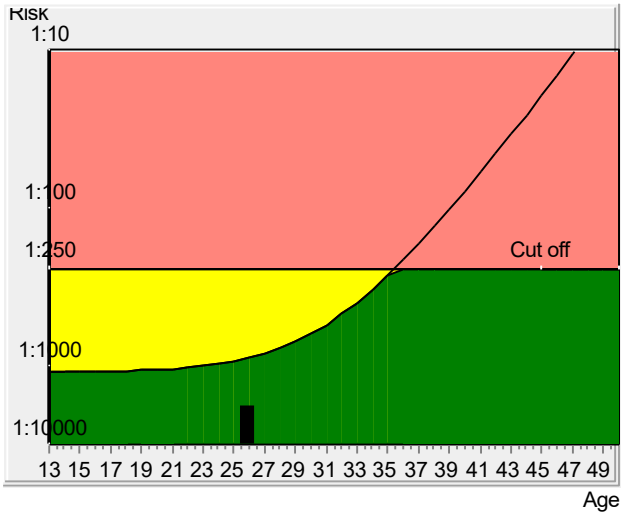
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	70	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

Parameter	Value	Corr. MoM
PAPP-A	1.92 mIU/mL	0.62
fb-hCG	41 ng/ml	1.00

Risks at sampling date

Age risk	1:906
Biochemical T21 risk	1:1856
Combined trisomy 21 risk	1:2787
Trisomy 13/18 + NT	<1:10000



Ultrasound data

Gestational age	11 + 6
Method	CRL Robinson
Scan date	22/08/25
Crown rump length in mm	54
Nuchal translucency MoM	1.32
Nasal bone	present
Sonographer	Naga Sai Teja
Qualifications in measuring NT	Naga Sai Teja

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 2787 women with the same data, there is one woman with a trisomy 21 pregnancy and 2786 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