J.I.T.M. DIAGNOSTICS

Patient data

Name	MRS. ARCHANA	Patient ID	
Birthday	25/07/95	Sample ID	2506018102/NOD
Age at sample date	29.9	Sample Date	11/06/25
Gestational age	13 + 2		

Correction factors

Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	no
Weight	60	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

Parameter	Value	Corr. MoM
PAPP-A	12.6 mIU/ml	2.31
fb-hCG	173.2 ng/ml	4.21

Risks at sampling date

Age risk	1:673
Biochemical T21 risk	1:443
Combined trisomy 21 risk	1:1410
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	13 + 2
Method	CRL Robinson
Scan date	11/06/25
Crown rump length in mm	73
Nuchal translucency MoM	1.05
Nasal bone	present
Sonographer	DR. HEMANT KUMAR
Qualifications in measuring NT	M.D

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 1410 women with the same data, there is one woman with a trisomy 21 pregnancy and 1409 women with not affected pregnancies.
The free beta HCG level is high.
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