

Patient data			
Name		MRS. ARCHANA	
Birthday		14/12/1994	
Age at sample date		28.9	
Patient ID		2311026211/NOD	
Sample ID		18/11/2023	
Sample Date		18/11/2023	
Gestational age			
Correction factors		13 + 3	
Fetuses	1	IVF	no
Weight	62	diabetes	no
Smoker	no	Origin	Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.26 mIU/ml	0.59	13 + 3
fb-hCG	25.8 ng/ml	0.64	Method
Risks at sampling date			CRL Robinson
Age risk		1:752	Scan date
Biochemical T21 risk		1:3464	18/11/2023
Combined trisomy 21 risk		<1:10000	Crown rump length in mm
Trisomy 13/18 + NT		<1:10000	75.9
Risk			Nuchal translucency MoM
1:10			0.97
1:100			Nasal bone
1:250			present
1:1000			Sonographer
1:10000			Qualifications in measuring NT
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			Trisomy 21
Age			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.

Sign of Physician

