

Patient data			
Name	MRS. NEHA	Patient ID	
Birth day	16/02/2002	Sample ID	2306053649/NOD
Age at sample date	21.4	Sample Date	28/06/2023
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	50	diabetes	no
Smoker	no	Origin	Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	5.99 mIU/ml	0.99	12 + 6
fb-hCG	54.6 ng/ml	1.21	Method
Risks at sampling date			CRL Robinson
Age risk	1:1085		Scan date
Biochemical T21 risk	1:4340		27/06/2023
Combined trisomy 21 risk	<1:10000		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		67.8
Risk			Nuchal translucency MoM
1:10			1.05
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.

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.

■ below cut off	■ Below Cut Off, but above Age Risk	■ above cut off
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