

Prisca 5.2.0.13
 Date of report: 23/08/2024

Patient data			
Name	MRS. NISHA	Patient ID	
Birthday	4/01/2002	Sample ID	2408040678/NOD
Age at sample date	22.6	Sample Date	22/08/2024
Gestational age	13 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	61	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	6.13 mIU/ml	1.15	13 + 2
fb-hCG	57.6 ng/ml	1.41	Method
			CRL Robinson
			Scan date
			22/08/2024
Risks at sampling date		Crown rump length in mm	
Age risk	1:1066	75	
Biochemical T21 risk	1:3998	Nuchal translucency MoM	
Combined trisomy 21 risk	<1:10000	0.70	
Trisomy 13/18 + NT	<1:10000	Nasal bone	
		present	
		Sonographer	
		DR. (MRS.) NEERJA CHOPRA	
		Qualifications in measuring NT	
		MD	
Risk		Trisomy 21	
		<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>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.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>	
Trisomy 13/18 + NT			
<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			

Sign of Physician