

Prisca 5.0.2.37
Date of report: 12-09-2020

Patient data			
Name	MRS. CHEENA SHARMA	Patient ID	
Birthday	05-01-1982	Sample ID	2009220453/AMB
Age at delivery	39.2	Sample Date	11-09-2020
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	82	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.1 mlU/ml	0.35	12 + 3
fb-hCG	64.1 ng/ml	1.81	Method
			CRL Robinson
			Scan date
			07-09-2020
Risks at term			Crown rump length in mm
Age risk		1:158	61
Biochemical T21 risk		>1:50	Nuchal translucency MoM
Combined trisomy 21 risk		>1:50	1.77
Trisomy 13/18 + NT		1:813	Nasal bone
			present
			Sonographer
			.
			Qualifications in measuring NT
			MD
Trisomy 21			
<p>The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.</p> <p>After the result of the Trisomy 21 Test (with nuchal translucency), it is expected that among less than 50 pregnancies with the same data, there is one trisomy 21 pregnancy.</p> <p>The PAPP-A level is low.</p> <p>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.</p> <p>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 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 1:813, which represents a low risk.</p>			

Sign of Physician

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