

Prisca 5.1.0.17
Date of report: 15-12-2024

N A

Patient data			
Name	Mrs. SAYALI SHITOLE		Patient ID
Birthday	23-08-1998	Sample ID	0662412140245
Age at sample date	26.3	Sample Date	A1372177
Gestational age	12 + 4		14-12-2024
Correction factors			
Fetuses	1	IVF	no
Weight	67	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.93 mIU/mL	0.55	12 + 3
fb-hCG	43.58 ng/mL	1.09	Method
			CRL Robinson
			Scan date
			13-12-2024
Risks at sampling date			Crown rump length in mm
Age risk	1:900		61
Biochemical T21 risk	1:1092		Nuchal translucency MoM
Combined trisomy 21 risk	1:5958		0.89
Trisomy 13/18 + NT	<1:10000		Nasal bone
			unknown
			Sonographer
			N A
			Qualifications in measuring NT
			MD
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 5958 women with the same data, there is one woman with a trisomy 21 pregnancy and 5957 women with not affected pregnancies. 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