

Prisca 5.1.0.17
Date of report: 24-10-2024

N A

Patient data			
Name	Mrs. ARATI MUNDHE		Patient ID
Birthday	31-08-1996	Sample ID	0662410220254
Age at sample date	28.1	Sample Date	A1685415
Gestational age	12 + 2		22-10-2024
Correction factors			
Fetuses	1	IVF	no
Weight	64	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	5.58 mIU/mL	1.68	12 + 2
fb-hCG	49.1 ng/mL	1.13	Method
			CRL Robinson
			Scan date
			22-10-2024
Risks at sampling date			Trisomy 21
Age risk	1:779		The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
Biochemical T21 risk	1:9964		After the result of the Trisomy 21 test (with NT) it is expected that among 5531 women with the same data, there is one woman with a trisomy 21 pregnancy and 5530 women with not affected pregnancies.
Combined trisomy 21 risk	1:5531		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!
Trisomy 13/18 + NT	<1:10000		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