

Prisca 5.1.0.17
Date of report: 14-12-2024

N A

Patient data			
Name	0692412120107	Patient ID	W/O SACHIKANTA BARIK
Birthday	21-06-1992	Sample ID	A1486350
Age at sample date	32.5	Sample Date	12-12-2024
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	58	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	8.64 mIU/mL	1.94	12 + 4
fb-hCG	38.41 ng/mL	0.95	Method
			CRL Robinson
			Scan date
			11-12-2024
Risks at sampling date		Trisomy 21	
Age risk	1:450	Crown rump length in mm	64
Biochemical T21 risk	<1:10000	Nuchal translucency MoM	0.88
Combined trisomy 21 risk	<1:10000	Nasal bone	unknown
Trisomy 13/18 + NT	<1:10000	Sonographer	N A
		Qualifications in measuring NT	MD
		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.	

Sign of Physician

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