

Prisca 5.1.0.17
Date of report: 19/09/25

NA

Patient data			
Name	Mrs. SUNITA NAYAK		Patient ID
Birthday	10/08/91	Sample ID	a1726138
Age at sample date	34.1	Sample Date	18/09/25
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	56	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.55 mIU/mL	0.38	12 + 2
fb-hCG	40.96 ng/mL	0.94	Method
			CRL Robinson
			Scan date
			17/09/25
Risks at sampling date			Trisomy 21
Age risk	1:327		Crown rump length in mm
Biochemical T21 risk	1:200		59.73
Combined trisomy 21 risk	1:1130		Nuchal translucency MoM
Trisomy 13/18 + NT	<1:10000		0.91
			Nasal bone
			present
			Sonographer
			NA
			Qualifications in measuring NT
			NA
			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 1130 women with the same data, there is one woman with a trisomy 21 pregnancy and 1129 women with not affected pregnancies. The PAPP-A level is low. 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