

Prisca 5.1.0.17
Date of report: 28/01/26

Patient data			
Name	Mrs. R.VAISHALI	Patient ID	0942601270004
Birthday	10/01/07	Sample ID	B3002095
Age at sample date	19.0	Sample Date	27/01/26
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	45	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.48 mIU/mL	0.58	12 + 5
fb-hCG	36.2 ng/mL	0.82	Method
			CRL Robinson
			Scan date
			27/01/26
Risks at sampling date			
Age risk		1:1107	Crown rump length in mm
Biochemical T21 risk		1:3031	64.8
Combined trisomy 21 risk		<1:10000	Nuchal translucency MoM
Trisomy 13/18 + NT		<1:10000	0.60
			Nasal bone
			present
			Sonographer
			NA
			Qualifications in measuring NT
			Sonographer
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 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