

Prisca 5.1.0.17
Date of report: 26-09-2024

N A

Patient data			
Name	Mrs. W/O SEP RAHUL KUMAR		Patient ID
Birthday	15-10-1999		Sample ID
Age at sample date	24.9		Sample Date
Gestational age	13 + 3		
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	5.1 mIU/mL	0.88	Method
fb-hCG	70.26 ng/mL	2.10	CRL Robinson
			Scan date
			20-09-2024
			Crown rump length in mm
			67
			Nuchal translucency MoM
			1.00
			Nasal bone
			present
			Sonographer
			N A
			Qualifications in measuring NT
			MD
Risks at sampling date			
Age risk		1:993	
Biochemical T21 risk		1:793	
Combined trisomy 21 risk		1:3253	
Trisomy 13/18 + NT		<1:10000	
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 3253 women with the same data, there is one woman with a trisomy 21 pregnancy and 3252 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