

Prisca 5.1.0.17
Date of report: 10/12/24

NA

Patient data					
Name	W/O NAGRE HARSHAL R		Patient ID	0692412090056	
Birthday	30/12/94		Sample ID	A1485844	
Age at sample date	29.9		Sample Date	09/12/24	
Gestational age	13 + 0				
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	63	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	13 + 0	
PAPP-A	1.93 mIU/mL	0.43	Method	CRL Robinson	
fb-hCG	33.78 ng/mL	0.92	Scan date	09/12/24	
Risks at sampling date			Crown rump length in mm	69.3	
Age risk	1:661		Nuchal translucency MoM	0.64	
Biochemical T21 risk	1:609		Nasal bone	present	
Combined trisomy 21 risk	1:3741		Sonographer	NA	
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT	MD	
Risk			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 3741 women with the same data, there is one woman with a trisomy 21 pregnancy and 3740 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