

SAGEPATH LABS PVT LTD.,

Prisca

5.1.0.17

Date of report:

22-06-2024

Patient data					
Name	Ms. MAYUR RAKHI CHETIA PHUKAN		Patient ID	0692406180135	
Birthday	04-01-1995		Sample ID	A0585681	
Age at sample date	29.5		Sample Date	18-06-2024	
Gestational age	11 + 6				
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	53.6	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	11 + 6	
PAPP-A	17.5 mIU/mL	5.15	Method	CRL Robinson	
fb-hCG	45.66 ng/mL	0.91	Scan date	18-06-2024	
Risks at sampling date			Crown rump length in mm	54.1	
Age risk	1:672		Nuchal translucency MoM	1.67	
Biochemical T21 risk	<1:10000		Nasal bone	unknown	
Combined trisomy 21 risk	1:7037		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 7037 women with the same data, there is one woman with a trisomy 21 pregnancy and 7036 women with not affected pregnancies. The PAPP-A level is high. 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