

SAGEPATH LABS PVT LTD.

Prisca

5.1.0.17

Date of report:

30-05-2024

Patient data			
Name		Mrs. V KIRAN MAI	
Patient ID		0012405290351	
Birthday		13-09-1998	
Sample ID		25220823	
Age at sample date		25.7	
Sample Date		29-05-2024	
Gestational age		13 + 1	
Correction factors			
Fetuses	1	IVF	no
Weight	72	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	6.98 mIU/mL	1.73	
fb-hCG	52.1 ng/mL	1.54	
Gestational age		13 + 0	
Method		CRL Robinson	
Scan date		28-05-2024	
Crown rump length in mm		69	
Nuchal translucency MoM		0.92	
Nasal bone		present	
Sonographer		N A	
Qualifications in measuring NT		MD	
Risks at sampling date		Trisomy 21	
Age risk	1:949		
Biochemical T21 risk	1:6168		
Combined trisomy 21 risk	<1:10000		
Trisomy 13/18 + NT	<1:10000		
Risk		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

 below cut off
 Below Cut Off, but above Age Risk
 above cut off