

Prisca 5.1.0.17
Date of report: 29/04/25

N A

Patient data			
Name	Mrs. SAKSHI DIVE	Patient ID	0742504250074
Birthday	23/02/05	Sample ID	B2458306
Age at sample date	20.2	Sample Date	25/04/25
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	56	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.9 mIU/mL	0.94	
fb-hCG	41.12 ng/mL	0.94	
Risks at sampling date		Gestational age	
Age risk		12 + 3	
Biochemical T21 risk		Method	
1:1083		CRL Robinson	
Biochemical T21 risk		Scan date	
1:6842		25/04/25	
Combined trisomy 21 risk		Crown rump length in mm	
<1:10000		61.9	
Trisomy 13/18 + NT		Nuchal translucency MoM	
<1:10000		0.81	
		Nasal bone	
		present	
		Sonographer	
		N A	
		Qualifications in measuring NT	
		MD	
Risk		Trisomy 21	
1:10		The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.	
1:100		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.	
1:250		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!	
1:1000		The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).	
1:10000		The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!	
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			
Age			
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