

Prisca 5.1.0.17
Date of report: 19/04/25

N A

Patient data			
Name	Ms. BANDI SHIRISHA		Patient ID
Birth day	26/06/95	Sample ID	
Age at sample date	29.8	Sample Date	
Gestational age	12 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	70	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.16 mIU/mL	0.39	
fb-hCG	42.5 ng/mL	1.01	
Risks at sampling date			
Age risk		1:655	
Biochemical T21 risk		1:376	
Combined trisomy 21 risk		1:1796	
Trisomy 13/18 + NT		<1:10000	
			Gestational age
			12 + 1
			Method
			CRL Robinson
			Scan date
			18/04/25
			Crown rump length in mm
			58
			Nuchal translucency MoM
			0.99
			Nasal bone
			present
			Sonographer
			N A
			Qualifications in measuring NT
			MD
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 1796 women with the same data, there is one woman with a trisomy 21 pregnancy and 1795 women with not affected pregnancies. The PAPP-A level is low. 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