

Prisca 5.1.0.17
Date of report: 03-10-2024

ASHWINI

Patient data			
Name	Mrs. EERAMMA	Patient ID	0352410030001
Birthday	02-10-2000	Sample ID	A1250696
Age at sample date	24.0	Sample Date	03-10-2024
Gestational age	13 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	52	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	15.1 mIU/mL	2.08	
fb-hCG	179.15 ng/mL	5.61	
Risks at sampling date			
Age risk	1:1039		Gestational age
Biochemical T21 risk	1:684		Method
Combined trisomy 21 risk	1:2746		Scan date
Trisomy 13/18 + NT	<1:10000		Crown rump length in mm
			Nuchal translucency MoM
			Nasal bone
			Sonographer
			Qualifications in measuring NT
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 2746 women with the same data, there is one woman with a trisomy 21 pregnancy and 2745 women with not affected pregnancies. The free beta HCG 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

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