

Prisca 5.2.0.13
Date of report: 12/01/22

Patient data			
Name	ASIMA MIDYA	Patient ID	
Birthday	09/10/81	Sample ID	10097773
Age at sample date	40.3	Sample Date	10/01/22
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	unknown
Weight	71	diabetes	unknown
Smoker	unknown	Origin	Asian
		Previous trisomy 21	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.65 mIU/mL	0.89	
fb-hCG	14.49 ng/mL	0.45	
Risks at sampling date			Gestational age
Age risk		1:74	12 + 4
Biochemical T21 risk		1:1934	Method
Combined trisomy 21 risk		1:8917	CRL Robinson
Trisomy 13/18 + NT		<1:10000	Scan date
			09/01/22
			Crown rump length in mm
			63.1
			Nuchal translucency MoM
			0.84
			Nasal bone
			unknown
			Sonographer
			DR.ANINDYA BANDYOPADHYAY
			Qualifications in measuring NT
			MD.
Risk			Trisomy 21
			<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 8917 women with the same data, there is one woman with a trisomy 21 pregnancy and 8916 women with not affected pregnancies.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>
Trisomy 13/18 + NT			
<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			

Sign of Physician