

Prisca 5.2.0.13  
Date of report: 26/01/22

Patient data			
Name	SUSAMA KUNDU	Patient ID	
Birthday	06/03/94	Sample ID	
Age at sample date	27.9	Sample Date	23/01/22
Gestational age	13 + 5		
Correction factors			
Fetuses	1	IVF	unknown
Weight	63	diabetes	unknown
Smoker	unknown	Origin	Asian
		Previous trisomy 21	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	8.07 mIU/mL	1.21	
fb-hCG	8.28 ng/mL	0.34	
Risks at sampling date			Gestational age
Age risk	1:836		13 + 4
Biochemical T21 risk	<1:10000		Method
Combined trisomy 21 risk	<1:10000		CRL Robinson
Trisomy 13/18 + NT	<1:10000		Scan date
			22/01/22
			Crown rump length in mm
			79
			Nuchal translucency MoM
			0.89
			Nasal bone
			unknown
			Sonographer
			DR.AMIT KUMAR RANO.
			Qualifications in measuring NT
			M.D.
Risk			Trisomy 21
			<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></p> <p>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.</p> <p>The free beta HCG level is low.</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><b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b></p>			

Sign of Physician