

N A

Patient data			
Name	Mrs. PRIYAMBADA MALLICK		Patient ID
Birthday	19-07-1997		Sample ID
Age at sample date	27.0		Sample Date
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	75	diabetes	no
Smoker	no	Origin	Asian
			Previous trisomy 21 pregnancies
			unknown
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	10.91 mIU/mL	2.84	Method
fb-hCG	43.69 ng/mL	1.31	CRL Robinson
Risks at sampling date			Scan date
Age risk			21-07-2024
Biochemical T21 risk			Crown rump length in mm
<1:10000			53
Combined trisomy 21 risk			Nuchal translucency MoM
<1:10000			1.06
Trisomy 13/18 + NT			Nasal bone
<1:10000			unknown
			Sonographer
			N A
			Qualifications in measuring NT
			MD
Risk 1:10 1:100 1:250 1:1000 1:10000 Age 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Cut off			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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. The PAPP-A 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