

P.PRATHIBHA

Patient data			
Name	Mrs. G.SWATHI		Patient ID
Birth day	19-05-1997		Sample ID
Age at sample date	24.8		Sample Date
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	56	diabetes	no
Smoker	no	Origin	Asian
			Previous trisomy 21 pregnancies
			unknown
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.69 mIU/mL	0.58	12 + 5
fb-hCG	46.57 ng/mL	1.14	Method
Risks at sampling date			CRL Robinson
Age risk	1:975		Scan date
Biochemical T21 risk	1:1262		16-03-2022
Combined trisomy 21 risk	1:6075		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		66
			Nuchal translucency MoM
			0.95
			Nasal bone
			unknown
			Sonographer
			DR NA
			Qualifications in measuring NT
			MD
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 6075 women with the same data, there is one woman with a trisomy 21 pregnancy and 6074 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><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