

Prisca 5.1.0.17
Date of report: 31-01-2022

Patient data			
Name	MRS. SANDEEP	Patient ID	
Birthday	16-01-1981	Sample ID	2201220647/AMB
Age at sample date	41.0	Sample Date	27-01-2022
Gestational age	11 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	89	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	0.31 mIU/ml	0.21	Gestational age 11 + 3
fb-hCG	26.1 ng/ml	0.57	Method CRL Robinson
			Scan date 27-01-2022
Risks at sampling date			Crown rump length in mm 49.6
Age risk		1:57	Nuchal translucency MoM 0.84
Biochemical T21 risk		>1:50	Nasal bone present
Combined trisomy 21 risk		1:110	Sonographer .
Trisomy 13/18 + NT		1:69	Qualifications in measuring NT MD
Trisomy 21			
<p>The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 110 women with the same data, there is one woman with a trisomy 21 pregnancy and 109 women with not affected pregnancies.</p> <p>The PAPP-A level is low.</p> <p>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.</p> <p>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 test (with nuchal translucency) is 1:69, which represents an increased risk.</p>			

Sign of Physician

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