

Prisca 5.0.2.37
Date of report: 25-04-2018

Patient data			
Name	MRS. PARAMJIT KAUR	Patient ID	
Birthday	25-01-1983	Sample ID	1804220571/AMB
Age at sample date	35.2	Sample Date	23-04-2018
Gestational age	13 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	58.7	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	5.2 mIU/ml	0.96	Gestational age 12 + 1
fb-hCG	110.1 ng/ml	3.81	Method CRL Robinson
			Scan date 12-04-2018
Risks at sampling date			Trisomy 21
Age risk		1:269	Crown rump length in mm 58.2
Biochemical T21 risk		1:53	Nuchal translucency MoM 1.12
Combined trisomy 21 risk		1:152	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer .
			Qualifications in measuring NT MD
			<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 152 women with the same data, there is one woman with a trisomy 21 pregnancy and 151 women with not affected pregnancies.</p> <p>The free beta HCG level is high.</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>
			<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>

Sign of Physician

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