

Prisca 5.0.2.37
Date of report: 11-02-2019

Patient data			
Name	MRS. RAVINDER KAUR		Patient ID
Birthday	14-01-1996	Sample ID	211902090002
Age at sample date	23.0	Sample Date	09-02-2019
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	unknown
Weight	50	diabetes	unknown
Smoker	unknown	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.8 mIU/ml	0.69	13 + 1
fb-hCG	35.1 ng/ml	1.17	Method
			LMP
			Scan date
Risks at sampling date			Crown rump length in mm
Age risk		1:790	Nuchal translucency MoM
Biochemical T21 risk		1:1460	Nasal bone
Combined trisomy 21 risk		1:2641	Sonographer
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT
			MD
Risk			Trisomy 21
			<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 2641 women with the same data, there is one woman with a trisomy 21 pregnancy and 2640 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 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 (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