

Prisca 5.0.2.37
Date of report: 30-10-2018

Patient data			
Name	MRS. LAKHVIR	Patient ID	1801220060/AMB
Birth day	07-05-1989	Sample ID	1801220060/AMB
Age at sample date	29.0	Sample Date	30-10-2018
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	61.8	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.26 mIU/ml	0.51	12 + 4
fb-hCG	19.5 ng/ml	0.56	Method
			CRL Robinson
			Scan date
			29-10-2018
Risks at sampling date			Crown rump length in mm
Age risk		1:869	62.0
Biochemical T21 risk		1:3549	Nuchal translucency MoM
Combined trisomy 21 risk		<1:10000	0.86
Trisomy 13/18 + NT		<1:10000	Nasal bone
			present
			Sonographer
			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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</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>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
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