

Prisca 5.1.0.17
Date of report: 21-08-2019

Patient data			
Name	MRS. SEEMA KUMARI	Patient ID	1908221131/AMB
Birthday	13-06-1995	Sample ID	1908221131/AMB
Age at delivery	24.7	Sample Date	20-08-2019
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	55	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.98 mIU/ml	0.72	12 + 5
fb-hCG	81.8 ng/ml	2.33	Method
			CRL Robinson
			Scan date
			20-08-2019
Risks at term		Crown rump length in mm	
Age risk	1:1394	66.1	
Biochemical T21 risk	1:548	Nuchal translucency MoM	
Combined trisomy 21 risk	1:3331	0.77	
Trisomy 13/18 + NT	<1:10000	Nasal bone	
		present	
		Sonographer	
		.	
		Qualifications in measuring NT	
		MD	
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 3331 women with the same data, there is one woman with a trisomy 21 pregnancy and 3330 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>			
<p>Risk</p> <p>Age</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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