

Prisca 5.0.2.37  
**Date of report: 25-11-2018**

Patient data			
Name	Ms. SUMAN	Patient ID	1810220470/AMB
Birthday	29-08-1989	Sample ID	1810220470/AMB
Age at sample date	29.3	Sample Date	24-11-2018
Gestational age	13 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	66	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.67 mIU/ml	0.85	13 + 2
fb-hCG	19.1 ng/ml	0.54	Method
			CRL Robinson
			Scan date
			21-11-2018
Risks at sampling date		Crown rump length in mm	
Age risk	1:679	Nuchal translucency MoM	
Biochemical T21 risk	<1:10000	Nasal bone	
Combined trisomy 21 risk	<1:10000	present	
Trisomy 13/18 + NT	<1:10000	Sonographer	
		Qualifications in measuring NT	
		MD	
Risk		Trisomy 21	
		<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></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 hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>	
		<p><b>Trisomy 13/18 + NT</b></p> <p><b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b></p>	

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**Sign of Physician**

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