

Patient data		MRS. POONAM W/O SATYAWAN	Patient ID	1710220537/AMB
Name				
Birthday		19/10/89	Sample ID	1710220537/AMB
Age at sample date		28.0	Sample Date	26/10/17
Gestational age		11 + 3		
Correction factors				
Fetuses		1 IVF	no Previous trisomy 21	no
Weight		52.9	no pregnancies	
Smoker		no	Origin	Asian
Biochemical data		Value	Ultrasound data	11 + 3
Parameter		Corr. MoM	Gestational age	
PAPP-A		3.2 mIU/ml	Method	CRL Robinson
fb-hCG		96.4 ng/ml	Scan date	26/10/17
Risks at sampling date			Crown rump length in mm	49.7
Age risk		1:761	Nuchal translucency MoM	1.04
Biochemical T21 risk		1:1212	Nasal bone	present
Combined trisomy 21 risk		1:4180	Sonographer	.
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD
Risk		1:10	Trisomy 21	<b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b>
1:100			After the result of the Trisomy 21 test (with NT) it is expected that among 4180 women with the same data, there is one woman with a trisomy 21 pregnancy and 4179 women with not affected pregnancies.	
1:250		Cut off	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!	
1:1000			The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).	
1:10000			The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no	

Trisomy 13/18 + NT

**The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.**

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Sign of Physician



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

