

Patient data			
Name	MRS. AMANDEEP KAUR	Patient ID	2003220412/AMB
Birthday	25-08-1980	Sample ID	2003220412/AMB
Age at delivery	40.0	Sample Date	12-03-2020
Gestational age	13 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	60	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	6.7 mIU/ml	1.27	13 + 4
fb-hCG	52.5 ng/ml	1.83	Method
			CRL Robinson
Risks at term		Scan date	11-03-2020
Age risk	1:126	Crown rump length in mm	77.3
Biochemical T21 risk	1:306	Nuchal translucency MoM	0.56
Combined trisomy 21 risk	1:1557	Nasal bone	present
Trisomy 13/18 + NT	<1:10000	Sonographer	.
		Qualifications in measuring NT	MD
Risk	1:10	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 1557 women with the same data, there is one woman with a trisomy 21 pregnancy and 1556 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>	
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

