

Patient data			
Name	MRS. SHARANJEET KAUR	Patient ID	1911220139/AMB
Birthday	19-01-1990	Sample ID	1911220139/AMB
Age at delivery	30.3	Sample Date	04-11-2019
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	55.14	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	4.5 mIU/ml	1.22	12 + 1
fb-hCG	26.9 ng/ml	0.73	Method
			CRL Robinson
Risks at term		Scan date	02-11-2019
Age risk	1:929	Crown rump length in mm	57.3
Biochemical T21 risk	<1:10000	Nuchal translucency MoM	0.92
Combined trisomy 21 risk	<1:10000	Nasal bone	present
Trisomy 13/18 + NT	<1:10000	Sonographer	.
		Qualifications in measuring NT	MD
Risk	1:10	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 be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>	
Trisomy 13/18 + NT			
<p><b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b></p>			

Sign of Physician



below cut off

Below Cut Off, but above Age Risk

above cut off