

Patient data			
Name	MRS.BHARTI	Patient ID	131903220009
Birthday	19-09-1991	Sample ID	131903220009
Age at sample date	27.0	Sample Date	22-03-2019
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	35	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	6.5 mIU/ml	2.14	12 + 3
fb-hCG	64.3 ng/ml	1.48	Method
			CRL Robinson
Risks at sampling date		Scan date	18-03-2019
Age risk	1:1020	Crown rump length in mm	59.0
Biochemical T21 risk	1:9697	Nuchal translucency MoM	0.41
Combined trisomy 21 risk	<1:10000	Nasal bone	present
Trisomy 13/18 + NT	<1:10000	Sonographer	.
		Qualifications in measuring NT	MD
Risk		Trisomy 21	
<p>Risk 1:10</p>		<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 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>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			

Sign of Physician

