

Date of Report 12/11/2022 PRISCA 5.1.0.17

Patient Data					
Name MRS. NISHA			Patient ID		012211110152
Birthday		3/9/1995	Sample ID		11523709
Age at term		27.08	Sample Date		11/11/2022
Gestational age		12+0			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	55 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+0
PAPP-A	3.27 mIU/ml	0.67	Method		CRL (<>Robinson)
fb-hCG	48.9 ng/ml	1.11	Scan date		11/11/2022
Risks at sampling date			Nuchal translucency 1.6		
Age Risk		1:832	Nuchal translucency MoM 1.1		
Biochemical T21 risk		1:1627	Nasal bone Preser		Present
Combined trisomy 21 risk 1:4618		1:4618			
Trisomy 13/18 + NT <1:10000		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 1:1000 1:1000 1:10000 1:3 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk			The calculated risk for Trisomy 21(with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21with NT test it is expected that among 4618 women with the same data, there is one woman with a trisomy 21 pregnancy and 4617 women with not affected pregnancies. The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that the risk calculations are statistical aapproaches and have no diagnostic value! The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		