

 Date of Report
 2/8/2022

 PRISCA
 5.1.0.17

Patient Data					
Name MRS. SAPNA			Patient ID		012207310179
Birthday		22.11.1990	Sample ID		11598945
Age at term		32.01	Sample Date		31/7/2022
Gestational age		12+1			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	48 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	11+6
PAPP-A	$5.62~\mathrm{mIU/ml}$	0.59	Method		CRL (<>Robinson)
fb-hCG	76.5 ng/ml	1.71	Scan date		30/7/2022
Risks at sampling date			Nuchal translucency 1.5		
Age Risk		1:502	Nuchal translucency MoM 1.0		1.09
Biochemical T21 risk		1:748	Nasal bone Preser		Present
Combined trisomy 21 risk		1:2329			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:1000 1:1000 1:1000 1:10000 1:1			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 2329 women with the same data, there is one woman with a trisomy 21 pregnancy and 2328 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		