

 Date of Report
 15/12/2022

 PRISCA
 5.1.0.17

Patient Data					
Name MRS. PUSHPA KUMARI			Patient ID		052212140021
Birthday		12/2/2002	Sample ID		11520789
Age at term		21.02	Sample Date		14/12/2022
Gestational age		13+2			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	39 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			13+1
PAPP-A	4.92 mIU/ml	0.42	Method		CRL (<>Robinson)
fb-hCG	29.8 ng/ml	0.84	Scan date		13/12/2022
Risks at sampling date			Nuchal translucency 2		
Age Risk 1:1104		1:1104	Nuchal translucency MoM 1.16		
Biochemical T21 risk		1:1159	Nasal bone Present		
Combined trisomy 21 r	isk	1:3279			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndro	ome Risk (Trisomy 21 S	Screening)
1:100 1:1000 1:1000 1:10000 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 21 with NT test it is expected that among 3279 women with the same data, there is one woman with a trisomy 21 pregnancy and 3278 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		