

 Date of Report
 7/8/2022

 PRISCA
 5.1.0.17

Patient Data					
Name	MRS. KAVYA		Patient ID		012208060098
Birthday		18/1/1999	Sample ID		11594785
Age at term		23.11	Sample Date		6/8/2022
Gestational age		13+3			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	50 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	11+4
PAPP-A	5.6 mIU/ml	0.6	Method		CRL (<>Robinson)
fb-hCG	59.4 ng/ml	1.94	Scan date		24/07/2022
Risks at sampling date			Nuchal translucency 1.2		
Age Risk	ge Risk 1:1045		Nuchal translucency MoM 0.88		
Biochemical T21 risk		1:415	Nasal bone		Present
Combined trisomy 21 risk		1:2360			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndro	ome Risk (Trisomy	21 Screening)
1:100 1:250 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 21 with NT test it is expected that among 2360 women with the same data, there is one woman with a trisomy 21 pregnancy and 2359 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		