

 Date of Report
 18/6/2022

 PRISCA
 5.1.0.17

Patient Data					
Name	I	MRS. SWETA	Patient ID		012206170136
Birthday		10/12/1993	Sample ID		11598758
Age at term		28.11	Sample Date		17/6/2022
Gestational age		12+3			
Correction factors	<u> </u>				
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	78 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+2
PAPP-A	2.96 mIU/ml	0.78	Method		CRL(<>Robinson)
fb-hCG	32.5 ng/ml	0.92	Scan date		16/6/2022
Risks at sampling date			Nuchal Translucency 1.2		
Age Risk		1:756	Nuchal Translucency MoM		0.78
Biochemical T21 risk		1:3245	Nasal Bone Present		
Combined T21 risk		<1:10000			
Trisomy 13/18+NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100  1:250  1:1000  1:10000  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 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 and 9999 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		