

 Date of Report
 5/6/2023

 PRISCA
 5.1.0.17

Patient Data					
Name	me MRS. SAPNA		Patient ID		012306030318
Birthday		18/09/1995	Sample ID		11671477
Age at term		28.1	Sample Date		3/6/2023
Gestational age		13+3			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	56 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	13+0
PAPP-A	$4.32~\mathrm{mIU/ml}$	0.53	Method		CRL (<>Robinson)
fb-hCG	25.3 ng/ml	0.86	Scan date		1/6/2023
Risks at sampling date			Nuchal translucency (NT) 1.8		
Age Risk	1:840		Nuchal translucency MoM 1.0		
Biochemical T21 risk		1:1589	Nasal bone		present
Combined trisomy 21 ris	k	1:5961			
Trisomy 13/18		<1:10000			
Risk			Down's Syndro	ome Risk (Trisomy 2	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 21 test it is expected that among 5961 women with the same data, there is one woman with a trisomy 21 pregnancy and 5960 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		