

 Date of Report
 10/6/2022

 PRISCA
 5.1.0.17

Patient Data					
Vame MRS. YASHODA		Patient ID 0122060		012206090050	
Birthday	16/06/1993		Sample ID		11242261
Age at term 29.06		Sample Date 9/6/202			
Gestational age		13+2			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	75 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	9	13+1
PAPP-A	3.78 mIU/ml	0.84	Method		CRL(<>Robinson)
fb-hCG	44.9 ng/ml	1.12	Scan date		8/6/2022
Risks at sampling date			Nuchal Transl	ucency	1.4
Age Risk		1:744	Nuchal Transl	-	0.79
Biochemical T21 risk		1:2447	Nasal Bone	,	Present
Combined T21 risk		< 1:10000			
Trisomy 13/18+NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:250 Cut off 1:1000 1:10000 13 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		