

 Date of Report
 31/1/2023

 PRISCA
 5.1.0.17

MRS.						
MRS.						
	KRISHM	IA GULERIA	Patient ID		012301280127	
		15/07/1992	Sample ID		11505040	
		30.11	Sample Date		28/1/2023	
		13+3				
1	IVF		unknown	Previous trisomy 21	unknown	
58	Diabetes		NO	Pregnancies	unknown	
NO	Origin		Asian			
Biochemical Data				Ultrasound Data		
alue		Corr Mom			13+2	
4.29	m I U/ml	0.55	Method		CRL (<>Robinson)	
11.3	ng/ml	0.39	Scan date		28/1/2023	
Risks at sampling date			Nuchal translucency 1.6			
Age Risk		1:622	Nuchal translucency MoM 0.0			
Biochemical T21 risk		1:6283	Nasal bone Presen			
		<1:10000				
		<1:10000				
			Down's Syndro	ome Risk (Trisomy 21 Se	creening)	
1:100 1:250 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 21with NT test 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			
	58 NO 4.29 11.3 my 13 ow ri	4.29 mIU/ml 11.3 ng/ml 31 33 35 37 39 4	1 IVF 58 Diabetes NO Origin Alue Corr Mom 4.29 mIU/ml 0.55 11.3 ng/ml 0.39 1:622 1:6283 <1:10000 <1:10000 Cut off Cut off any 13/18 (with NT) is ow risk	Diabetes NO NO Origin Asian Ultrasound Date Nuchal translut 1:622 Nuchal translut 1:6283 Nasal bone After the result expected that a same data, there pregnancy and the information note that the rinhave no diagnormy 13/18 (with NT) is on the risk assovalues The laboratory on the risk assovalues	1 IVF unknown Previous trisomy 21 NO Origin Asian Ultrasound Data Ultrasound Data Ultrasound Data Method Scan date Nuchal translucency Nuchal translucency MoM 1:622 Nuchal translucency MoM Nasal bone <1:10000 VI:10000 The calculated risk for Trisomy 21 (wit cut off, which represents a low risk. After the result of the Trisomy 21 (wit cut off, which represents a low risk. After the result of the Trisomy 21 (wit cut off, which represents a low risk. The calculated risk by PRISCA depend the information provided by the referring note that the risk calculations are statistical have no diagnostic value! The laboratory cannot be hold responsion the risk assessment! Calculated risks values	