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Date of Report 11-10-2024 PRISCA 5.2.0.13

				PRISCA	5.2.0.13	
	N	IRS. ANJALI	Patient ID		12410090175	
16-10-2002			Sample ID		11863909	
Age at Sample date 22.0			Sample Date		09-10-2024	
		11+6				
1	IVF		unknown	Previous trisomy 21	unknown	
74	Diabetes		NO	Pregnancies	unknown	
NO	Origin		Asian			
Biochemical Data				Ultrasound Data		
Value		Corr Mom	Gestational age	2	11+6	
5.1	mIU/ml	1.58	Method		CRL (<>Robinson)	
114.4	ng/ml	2.76	Scan date		09-10-2024	
Risks at sampling date				ength in mm	52	
		1:1030	Nuchal translu	cency MoM	0.86	
		1:1287	Nasal bone		PRESENT	
		1:6226	Sonographer		DR. SANJEEV KUMAR	
		<1:10000	Qualifications	in measuring NT	MD	
1:1000 1:10000 1:10000				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 6225 women with the same data, there is one woman with a trisomy 21 pregnancy and 6225 women with not affected pregnancies. The free beta HCG level is high. 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 patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523;		
	74 NO Value 5.1 114.4	1 IVF 74 Diabetes NO Origin Value 5.1 mIU/ml 114.4 ng/ml	16-10-2002 22.0 11+6 1 IVF 74 Diabetes NO Origin Value Corr Mom 5.1 mIU/ml 1.58 114.4 ng/ml 2.76 1:1030 1:1287 1:6226 <1:10000	NO Origin Asian Value Corr Mom 5.1 mIU/ml 1.58 Method 114.4 ng/ml 2.76 Scan date Crown rump le 1:1030 Nuchal translut 1:1287 Nasal bone 1:6226 Sonographer <1:10000 Qualifications Down's Syndry The calculated cut off, which After the result of expected that an woman with a traffected pregnant Cut off Cut off The calculated reinformation profits risk calculating diagnostic value! The patient condone according	MRS. ANJALI 16-10-2002 22.0 22.0 11+6 1 IVF	

The calculated risk for Trisomy 13/18 (with NT) is <1:10000,

Trisomy 13/18+NT

which indicates a low risk

The laboratory cannot be hold responsible for their impact on the

risk assessment! Calculated risks have no diagnostic values