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 Date of Report
 5/5/2024

 PRISCA
 5.2.0.13

ent Data ne JYOTI		Patient ID		012405030229
				12001852
	22.1	Sample Date		3/5/2024
estational age 13+3				
1 IVF		unknown	Previous trisomy 21	unknown
53 Diabetes		NO	Pregnancies	unknown
NO Origin		Asian		
		Ultrasound Data		
Value	Corr Mom	Gestational age 12+1		
7.50 mIU/ml	0.86	Method		CRL (<>Robinson)
113.5 ng/ml	3.79	Scan date		24/04/2024
Risks at sampling date		Crown rump length in mm 56		
	1:1084	Nuchal translucency MoM		1.22
	1:172	Nasal bone		Present
sk	1:342	Sonographer DR. HARENI		DR. HARENDRA BHASKAR
	<1:10000	Qualifications in measuring NT		MBBS
Risk		Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off 1:10000 1:10000 1:10000 1:10000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 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 342 women with the same data, there is one woman with a trisomy 21 pregnancy and 341 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 approaches and have no diagnostic value! The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998). The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
	53 Diabetes NO Origin Value 7.50 mIU/ml 113.5 ng/ml sk	1/4/2002 22.1 13+3 1 IVF 53 Diabetes NO Origin Value Corr Mom 7.50 mIU/ml 0.86 113.5 ng/ml 3.79 1:1084 1:172 sk 1:342 <1:10000 Cut off Cut off	IVF unknown 53 Diabetes NO NO Origin Asian Ultrasound Day Value Corr Mom 7.50 mIU/ml 0.86 Method 113.5 ng/ml 3.79 Scan date Crown rump letter Nuchal transluture in Section of Secti	1 IVF unknown Previous trisomy 21 1 IVF Unknown Previous trisomy 21 1 Diabetes NO Origin Asian Value Corr Mom Gestational age 7.50 mIU/ml 0.86 Method 113.5 ng/ml 3.79 Scan date Crown rump length in mm 1:1084 Nuchal translucency MoM 1:172 Nasal bone sk 1:342 Sonographer <1:10000 Qualifications in measuring NT Down's Syndrome Risk (Trisomy 21 with off, which represents a low risk. After the result of the Trisomy 21 test (with Nava and the same data, there is one pregnancy and 341women with not affected plevel is high. The calculated risk by PRISCA depends information provided by the referring phrisk calculations are statistical approaches value! The patient combined risk presumes that according to accepted guidelines (Prenat The laboratory cannot be hold responsible assessment! Calculated risks have no diagent according to accepted guidelines (Prenat The laboratory cannot be hold responsible assessment! Calculated risks have no diagent according to accepted guidelines (Prenat The laboratory cannot be hold responsible assessment! Calculated risks have no diagent according to accepted guidelines (Prenat The laboratory cannot be hold responsible assessment! Calculated risks have no diagent according to accepted guidelines (Prenat The laboratory cannot be hold responsible assessment! Calculated risks have no diagent according to accepted guidelines (Prenat The laboratory cannot be hold responsible assessment! Calculated risks have no diagent according to accepted guidelines (Prenat The laboratory cannot be hold responsible assessment! Calculated risks have no diagent according to accepted guidelines (Prenat The laboratory cannot be hold responsible assessment! Calculated risks have no diagent according to accepted guidelines (Prenat The laboratory cannot be hold responsible assessment! Calculated risks have no diagent according to accepted guidelines (Prenat The laboratory cannot be hold responsible according to accepted guidelines (Prenat The laboratory cannot be hold responsible according to accepted g