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Date of Report 18/8/2022 PRISCA 5.1.0.17

					PRISCA	5.1.0.17	
Patient Data							
Name	MRS. NEHA KUMARI				Patient ID	012208170112	
Birthday	1/1/1993			}	Sample ID	11304122	
Age at delivery	31.01				Sample Date	17/08/2022	
Correction factors							
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown	
Weight in kg		Diabet	es	unknown	Pregnancies	unknown	
Smoker	Unknown	Origin		Asian			
Biochemical Data				Risks at sampl	ling date		
Parameter	Value	(	Corr MoM	Age Risk		1:764	
AFP	33.6 n	ng/ml	0.89	Trisomy 21 ris	sk	1:4582	
uE3	1.2 n	1.2 ng/ml 1.48 Neural tube defects risk		efects risk	1:8809		
hCG	18058.6  mIU/ml		0.63	Trisomy 18		<1:10000	
Ultrasound Data				Down's Syndre	ome Risk (Trisomy 2)	1 Screening)	
WOP Method	16+5 CRL (<>Robinson)			The calculated risk for Trisomy 21 is below the cut off which indicates a low risk.  After the result of the Trisomy 21 test it is expected			
1:10				that among 4582 women with the same data, there is one woman with a trisomy 21 pregnancy and 4581 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!			
1:100 1:250			Out off	Trisomy 18			
1:1 <mark>000</mark> 1:10000	1			indicates a low	d risk for Trisomy 18 : v risk Defect (NTD) Screen		
13 15 17 19 21 23			Age	The corrected MoM for AFP (0.89) is located in the low risk area for neural tube defects.			
The laboratory can n	ot be held resp	onsible fo	r their impaci	t on the risk asse	essment! Calculated va	lue has no diagnostic value!	