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Date of Report 22-11-2024 PRISCA 5.2.0.13

				PRISCA	5.2.0.13
Patient Data				Tiuse/1	0.2.0.10
Name		MRS. DIMPI	Patient ID		12411200148
Birthday		21-10-2000	Sample ID		11960722
Age at Sample date		24.1	Sample Date		20-11-2024
Gestational age		13+6			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	66.9 Diabete	S	NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	e	13+2
PAPP-A	7.5 mIU/ml	0.99	Method		CRL (<>Robinson)
fb-hCG	21.4 ng/ml	0.89	Scan date		16-11-2024
Risks at sampling date			Crown rump l	ength in mm	75
Age Risk		1:1040	Nuchal translu	icency MoM	0.76
Biochemical T21 risk		1:8084	Nasal bone		PRESENT
Combined trisomy 21 risk		<1:10000	Sonographer		DR.
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MBBS
Risk			Down's Syndr	ome Risk (Trisomy 21	Screening)
1:10 1:100 1:250 Cut off			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.  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!		

## 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 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