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				Date of Report PRISCA	14-09-2023 5.2.0.13
Patient Data				TRISCH	0.2.0.10
Name MRS ANJALI SHARMA			Patient ID		012309130129
Birthday 15-10-1991		Sample ID		11799866	
Age at Sample date 31.9		Sample Date		13-09-2023	
Gestational age 12+2			2		
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	52 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin	1	Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+2
PAPP-A	5.39 mIU/	ml 0.92	Method		CRL (<>Robinson)
fb-hCG	93.6 ng/ml	2.24	Scan date		13-09-2023
Risks at sampling date			Crown rump length in mm 57.7		
Age Risk		1:487	Nuchal translu	cency MoM	0.99
Biochemical T21 risk		1:364	Nasal bone		PRESENT
Combined trisomy 21 risk	X	1:1516	Sonographer		DR VIKAS GOYAL
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MBBS
Risk		Down's Syndrome Risk (Trisomy 21 Screening)			
1:100 1:250 1:1000 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:1000 1:1000 1:1000 1:1000 1:1000 1:250 1:25		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 1516 women with the same data, there is one woman with a trisomy 21 pregnancy and 1515 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 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			
	Above Cut Off	Risk above Ag	e Risk	Risk below Age risk	