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	Sample Collection 9999 778 778				

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				Date of Report PRISCA	01-12-2020 5.0.2.37
Patient Data					
Name	MRS AN	NCHAL DEVI	Patient ID		012011280113
Birthday		06-05-1997	Sample ID		10788998
Age at delivery		23.6	Sample Date		28/11/2020
Gestational age		12+3			
Correction factors	1				
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	53.2 Diabetes		unknown	Pregnancies	unknown
Smoker	Unknown Origin		Asian		
Biochemical Data			Ultrasound Da	ata	
Parameter	Value	Corr Mom	Gestational age	2	11+0
PAPP-A	2.98 mIU/ml	0.6	Method		CRL(<>Robinson
fb-hCG	36.1 ng/ml	0.73	Scan date		18-11-2020
Risks at sampling date			Crown rump le	ength in mm	40.4
Age Risk	ge Risk 1:1010		Nuchal translucency MOM 0.75		
Biochemical T21 risk		1:3658	Nasal bone		Present
Combined Trisomy 21	Risk	<1:10000	Sonographer		DR.ANKIT BHARGAVA
Trisomy 13/18 + NT		<1:10000	Qualification in	n measuring NT	MD
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:00 1:250 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:1000 1:1000 1:1000 1:1000 1:250	27 29 31 33 35 37 39 4	Out 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 and 9999 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 calculated risk for <1:10000, which indic		NT) is	on the risk assessment! Calculated risks have no diagnostic values		
Ris	sk Above Cut Off		Risk above Ag	e Risk	Risk below Age risk