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				Date of Report PRISCA	14-11-2023 5.2.0.13
Patient Data					
Name		BISWABARA	Patient ID		012311130057
Birthday		12-03-1994	Sample ID		11805978
Age at Sample date		29.7	Sample Date		13-11-2023
Gestational age		12+6			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	62 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+6
PAPP-A	3.78 mIU/ml	0.64	Method		CRL (<>Robinson)
fb-hCG	35.2 ng/ml	1.04	Scan date		13-11-2023
Risks at sampling date			Crown rump length in mm 72		
Age Risk		1:680	Nuchal translu	cency MoM	0.56
Biochemical T21 risk		1:1362	Nasal bone		PRESENT
Combined trisomy 21 risk 1:7664		1:7664	Sonographer DR PRAKAS		
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MD
Risk			Down's Syndro	ome Risk (Trisomy 21	Screening)
1:10 1:100 1:250 1:1000 1:1000 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 1:1250 1:1000 1:1250 1:1000 1:1250 1:1000 1:1250 1:1000 1:1250 1:1000 1:1250 1:1000 1:1250 1:1000 1:1250 1:1000 1:1250 1:1000 1:1250 1:1000 1:1250 1:1000 1:1250 1:1000 1:1250 1:1000 1:1250 1:10000 1:10000	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 7664 women with the same data, there is one woman with a trisomy 21 pregnancy and 7663 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				
	k Above Cut Off		Risk above Ag	e Risk	Risk below Age risk