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				Date of Report PRISCA	21-07-2024 5.2.0.13
Patient Data					
Name	MRS. US	SHA KUMARI	Patient ID		12407200088
Birthday		17-03-2004	Sample ID		11877792
Age at Sample date	20.3		Sample Date		20-07-2024
Gestational age		13+0			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	46 Diabetes	3	NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	13+0
PAPP-A	5.6 mIU/ml	0.64	Method		CRL(<>Robinson)
fb-hCG	17.6 ng/ml	0.49	Scan date		20-07-2024
Risks at sampling date			Crown rump length in mm 66.8		
Age Risk 1:1101		Nuchal translucency MoM 0.53			
Biochemical T21 risk	<u>Γ21 risk</u> <1:10000		Nasal bone		present
Combined trisomy 21 risk <1:10000		Sonographer DR. RAHU			
Trisomy 13/18 + NT		<1:10000	Qualifications	in measuring NT	MBBS
Risk Tien 1:10			Down's Syndrome Risk (Trisomy 21 Screening)		
1: 00 1:250 1:1000 1:1000 1:10000 13 15 17 19 21 23 25 27 Trisomy 13/18 + NT		The calculated risk for Trisomy 21 test (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 amog 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! 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			
The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk				Calculated risks have no o	liagnostic values



Risk Above Cut Off

