

Risk Above Cut Off

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Risk below Age risk

 Date of Report
 30-04-2024

 PRISCA
 5.2.0.13

Patient Data						
Name		ľ	MRS. SUNNY	Patient ID		12404280199
Birthday	01-01-1994			Sample ID		11823031
Age at Sample date	30.3					28-04-2024
Gestational age 13+6						
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	59	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	2	13+5
PAPP-A	6.8	mIU/ml	0.77	Method		CRL (<>Robinson)
fb-hCG	42.9	ng/ml	1.72	Scan date		27-04-2024
Risks at sampling date				Crown rump le	ength in mm	77
Age Risk			1:649	Nuchal translu	cency MoM	1.17
Biochemical T21 risk	al T21 risk		1:629	Nasal bone		present
Combined trisomy 21 risk			1:1570	Sonographer		DR. HARENDRA BHASKAR
Trisomy 13/18 + NT			<1:10000	Qualifications	in measuring NT	MBBS
Risk 1:10				Down's Syndr	ome Risk (Trisomy	21 Screening)
1:1000 1:1000 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 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 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 1570 women with the same data, there is one woman with a trisomy 21 pregnancy and 1569 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		
which indicates a low risk					- 0	

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