

Risk Above Cut Off

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Risk below Age risk

				Date of Report PRISCA	17/6/2023 5.2.0.13
Patient Data				1140 011	3,2,3,13
Name MRS.NEETU KUMARI			Patient ID		012306160151
Birthday		4/11/1998	Sample ID		11654661
Age at term		25.2	Sample Date		16/6/2023
Gestational age		13+2			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	70 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			13+2
PAPP-A	$3.59~\mathrm{mIU/ml}$	0.46	Method		CRL (<>Robinson)
fb-hCG	20.4 ng/ml	0.59	Scan date		16/6/2023
Risks at sampling date			Crown rump length in mm 70.6		
Age Risk 1:1002		Nuchal translucency MoM 0.62			
Biochemical T21 risk 1:2761		1:2761	Nasal bone Present		
Combined trisomy 21 risk <1:10000					
Trisomy 13/18 + NT		< 1:10000			
Risk			Down's Syndro	ome Risk (Trisomy 21	Screening)
1:100 1:250 Cut 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 with NT test it is expected that among 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 women with not affected pregnancies.		
1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			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 laboratory cannot be hold responsible for their impact		
The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk			on the risk assessment! Calculated risks have no diagnostic values		

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