MOLQ Express Lab

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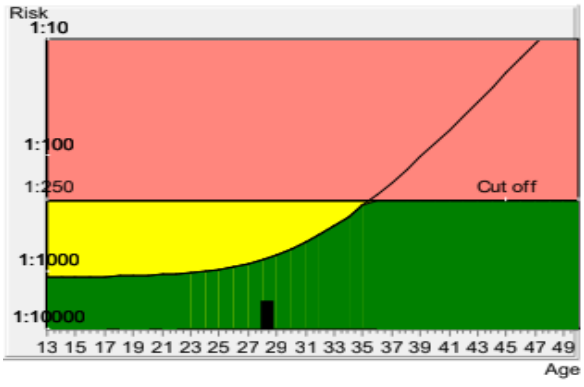
Date of Report	7/8/2022
PRISCA	5.1.0.17

Patient Data			
Name	MRS. PRIYANKA	Patient ID	102208070007
Birthday	18/3/1994	Sample ID	011567223
Age at term	28.1	Sample Date	7/8/2022
Gestational age	12+5		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	62.5	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+2
PAPP-A	3.6 mIU/ml	0.65	Method	CRL (<>Robinson)
fb-hCG	42.5 ng/ml	1.21	Scan date	2/8/2022
			Nuchal translucency	1.2
Risks at sampling date			Nuchal translucency MoM	0.78
Age Risk		1:774	Nasal bone	Present
Biochemical T21 risk		1:1138		
Combined trisomy 21 risk		1:6516		
Trisomy 13/18 + NT		<1:10000		

Risk



Trisomy 13/18+NT

The calculated risk for Trisomy 13/18 (with NT) is <1:10000 , which indicates a low risk

Down's Syndrome Risk (Trisomy 21 Screening)

The calculated risk for Trisomy 21(with NT) is below the cut off, which represents a low risk.
After the result of the Trisomy 21with NT test it is expected that among 6516 women with the same data, there is one woman with a trisomy 21 pregnancy and 6515 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 laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

Risk Above Cut Off Risk above Age Risk Risk below Age risk