MOLQ Express Lab
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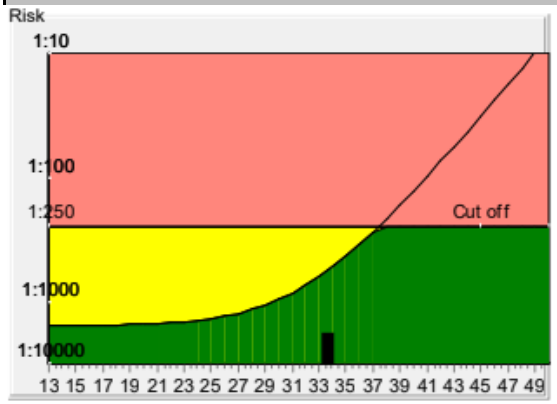
Date of Report	29/9/2021
PRISCA	5.2.0.13

Patient Data			
Name	MRS. MOUMITA DAS	Patient ID	012109270223
Birthday	30/07/1988	Sample ID	11153219
Age at delivery	33.68	Sample Date	28/9/2021
Gestational age	13+2		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	44	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+0
PAPP-A	4.1 mIU/ml	0.51	Method	CRL(<>Robinson
fb-hCG	31.1 ng/ml	0.72	Scan date	26/9/2021
			Nuchal translucency in mm	1.36
Risks at sampling date			Nuchal translucency MOM	0.77
Age Risk		1:556	Nasal bone	Present
Biochemical T21 risk		1:1392		
Combined Trisomy 21 Risk		1:7873		
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	CR

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 21 test (with NT) it is expected that among 7873 women with the same data, there is one woman with a trisomy 21 pregnancy and 7872 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