MOLQ Express Lab
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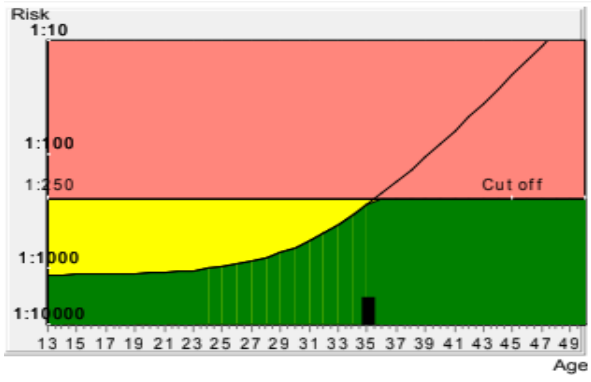
Date of Report	7/5/2023
PRISCA	5.2.0.13

Patient Data			
Name	**MRS. PUSHPA**	Patient ID	012305060069
Birthday	29/4/1988	Sample ID	11552294
Age at term	35.6	Sample Date	6/5/2023
Gestational age	13+2		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	68	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	**Value**	**Corr Mom**		12+6
PAPP-A	7.24 mIU/ml	1.18	Method	CRL (<>Robinson)
fb-hCG	23.8 ng/ml	0.83	Scan date	3/5/2023
Risks at sampling date			Nuchal translucency	1.56
Age Risk		1:279	Nuchal translucency MoM	0.94
Biochemical T21 risk		1:3733	Nasal bone	Present
Combined trisomy 21 risk		<1:10000		
Trisomy 13/18 + NT		<1:10000		

Risk



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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 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