MOLQ Express Lab
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Date of Report	16/1/2022
PRISCA	5.2.0.13

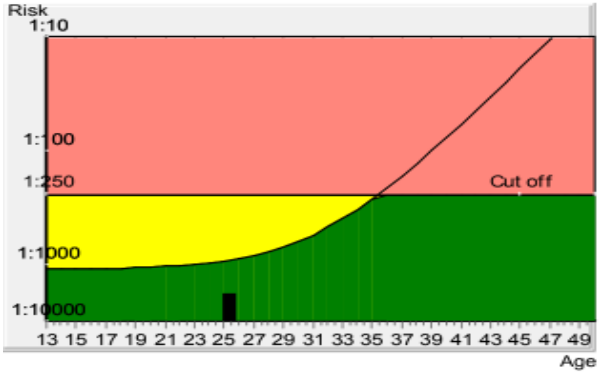
Patient Data			
Name	MRS. MEENAKSHI GURJAR	Patient ID	012201150002
Birthday	26/09/1996	Sample ID	11307529
Age at delivery	25.3	Sample Date	15/1/2022
Gestational age	12+0		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	69	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	11+5
PAPP-A	3.46 mIU/ml	1.11	Method	CRL(<>Robinson
fb-hCG	22.5 ng/ml	0.47	Scan date	13/1/2022
			Nuchal translucency	1.71
			Nuchal translucency MoM	1.3
			Nasal bone	present

Risks at sampling date	
Age Risk	1:930
Biochemical T21 risk	<1:10000
Combined T21 risk	<1:10000
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 21 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