MOLQ Express Lab
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Date of Report	21/8/2021
PRISCA	5.1.0.17

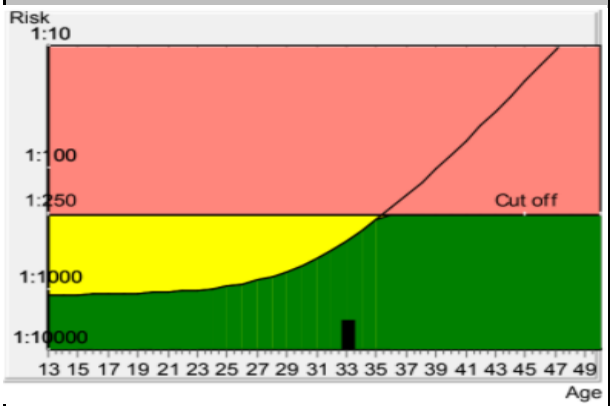
Patient Data			
Name	MRS. NEERU	Patient ID	012108200243
Birthday	26/07/1988	Sample ID	11211803
Age at delivery	33.1	Sample Date	20/8/2021
Gestational age	12+3		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	72	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+2
PAPP-A	4.5 mIU/ml	1.29	Method	CRL(<>Robinson
fb-hCG	39.8 ng/ml	0.89	Scan date	20/08/2021
			Crown rump length in mm	56

Risks at sampling date			
Age Risk	1:394	Nuchal translucency MOM	0.74
Biochemical T21 risk	1:5348	Nasal bone	Present
Combined Trisomy 21 Risk	<1:10000	Sonographer	DR. KIRTI RAJ
Trisomy 13/18 + NT	<1:10000	Qualification in measuring NT	MBBS, MD

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 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

Trisomy 13/18 + NT

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

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