MOLQ Express Lab
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Date of Report	02-02-2021
PRISCA	5.0.2.37

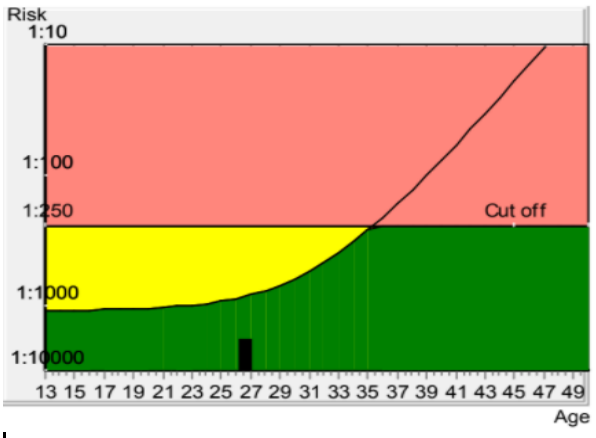
Patient Data			
Name	MRS BANDHAN DEEP KAUR	Patient ID	012101310008
Birthday	07-07-1994	Sample ID	10788079
Age at delivery	26.6	Sample Date	31/01/2021
Gestational age	11+5		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	51	Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown	Origin	Asian		

Biochemical Data		
Parameter	**Value**	**Corr Mom**
PAPP-A	3.65 mIU/ml	0.93
fb-hCG	40.2 ng/ml	0.74

Ultrasound Data	
Gestational age	11+4
Method	CRL(<>Robinson
Scan date	30-01-2021
Crown rump length in mm	47.4
Nuchal translucency MOM	0.85
Nasal bone	Present
Sonographer	DR.VIDYA SAGAR
Qualification in measuring NT	MD

Risks at sampling date	
Age Risk	1:858
Biochemical T21 risk	1:8639
Combined Trisomy 21 Risk	<1:10000
Trisomy 13/18 + NT	<1:10000



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