MOLQ Express Lab
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Date of Report	27-04-2021
PRISCA	5.0.2.37

Patient Data			
Name	MS. SHAILJA	Patient ID	012104200071
Birthday	23-06-1989	Sample ID	10943697
Age at delivery	31.8	Sample Date	20/04/2021
Gestational age	12+2		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	72.5	Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
PAPP-A	4.19	mIU/ml	1.28
fb-hCG	77.7	ng/ml	1.72

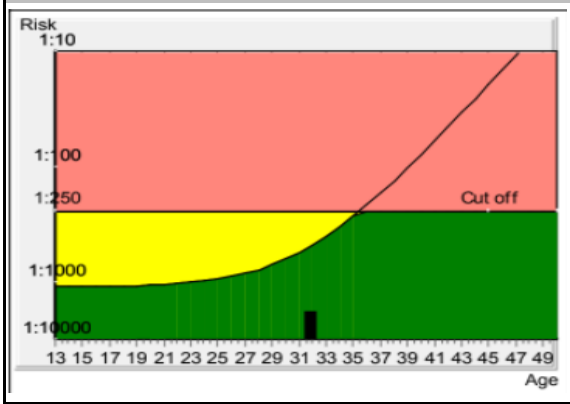
Ultrasound Data

Gestational age	12+0
Method	CRL(<>Robinson
Scan date	19-04-2021
Crown rump length in mm	53.6
Nuchal translucency MOM	0.88
Nasal bone	Present
Sonographer	Dr Anand Singh
Qualification in measuring NT	MBBS

Risks at sampling date

Age Risk	1:494
Biochemical T21 risk	1:1416
Combined Trisomy 21 Risk	1:6697
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 (with NT) it is expected that among more than 6697 women with the same data, there is one woman with a trisomy 21 pregnancy and 6696 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