MOLQ Express Lab
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Date of Report	27/11/2021
PRISCA	5.2.0.13

Patient Data			
Name	MRS. MANISHA W/O SOHAN	Patient ID	012111250104
Birthday	8/3/1991	Sample ID	11272969
Age at delivery	31.26	Sample Date	25/11/2021
Gestational age	12+0		

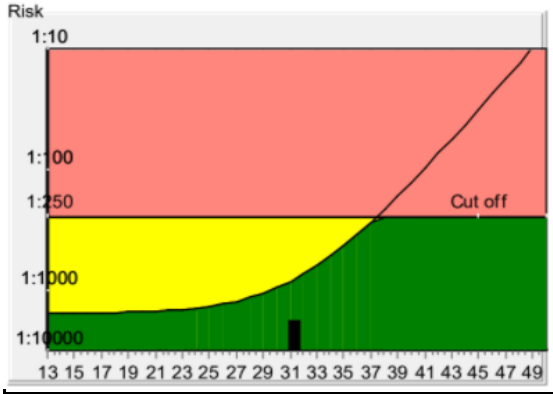
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	40.9	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			
Parameter	**Value**		**Corr Mom**
PAPP-A	9.78	mIU/ml	1.73
fb-hCG	121	ng/ml	2.25

Risks at sampling date	
Age Risk	1:824
Biochemical T21 risk	1:2057
Combined T21 risk	1:9990
Trisomy 13/18+NT	<1:10000

Ultrasound Data	
Gestational age	11+5
Method	CRL(<>Robinson
Scan date	24/11/2021
Nuchal translucency	0.9
Nuchal translucency MoM	0.64
Nasal bone	Present

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 9990 women with the same data, there is one woman with a trisomy 21 pregnancy and 9989 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