MOLQ Express Lab
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Date of Report	19/7/2021
PRISCA	5.1.0.17

Patient Data			
Name	MRS. KAJAL	Patient ID	012107150094
Birthday	3/4/2003	Sample ID	11042375
Age at delivery	18.3	Sample Date	16/07/2021
Gestational age	12+6		

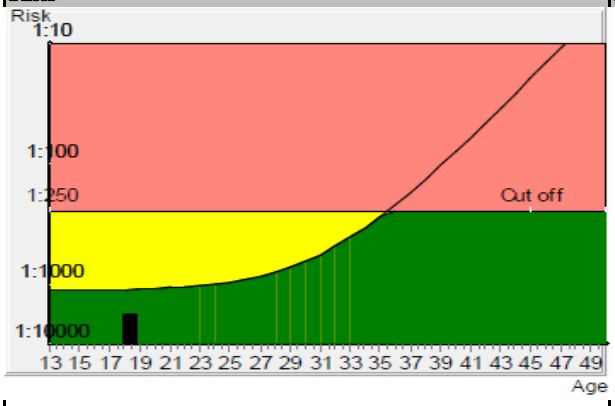
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	53	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data		
Parameter	**Value**	**Corr Mom**
PAPP-A	2.77 mIU/ml	0.47
fb-hCG	25.6 ng/ml	0.55

Risks at sampling date	
Age Risk	1:1119
Biochemical T21 risk	1:3988
Combined Trisomy 21 Risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound Data	
Gestational age	12+4
Method	CRL(<>Robinson
Scan date	14/07/2021
Crown rump length in mm	62.2
Nuchal translucency MOM	0.5
Nasal bone	Present
Sonographer	DR. BIPUL BISWAS
Qualification in measuring NT	MBBS MD

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 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