MOLQ Express Lab

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Date of Report	29/5/2023
PRISCA	5.1.0.17

Patient Data			
Name	MRS. SANGEETA	Patient ID	012305250096
Birthday	24/11/1987	Sample ID	11659350
Age at term	36.1	Sample Date	25/5/2023
Gestational age	13+1		

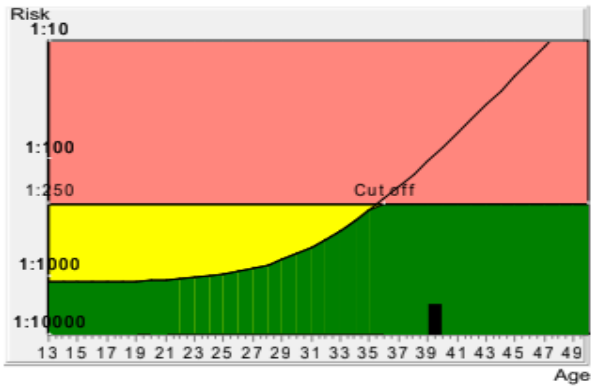
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	55	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			
Parameter	**Value**		**Corr Mom**
PAPP-A	3.69	mIU/ml	0.49
fb-hCG	18.9	ng/ml	0.59

Risks at sampling date	
Age Risk	1:249
Biochemical T21 risk	1:848
Combined trisomy 21 risk	1:4707
Trisomy 13/18	<1:10000

Ultrasound Data	
Gestational age	13+1
Method	CRL (<>Robinson)
Scan date	25/5/2023
Nuchal translucency (NT)	0.8
Nuchal translucency MoM	0.51
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 4707 women with the same data, there is one woman with a trisomy 21 pregnancy and 4706 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