MOLQ Express Lab
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Date of Report	3/11/2022
PRISCA	5.1.0.17

Patient Data			
Name	MRS. MANITA CHAUHAN	Patient ID	012210310154
Birthday	20/1/1990	Sample ID	11245864
Age at term	33.02	Sample Date	2/11/2022
Gestational age	12+6		

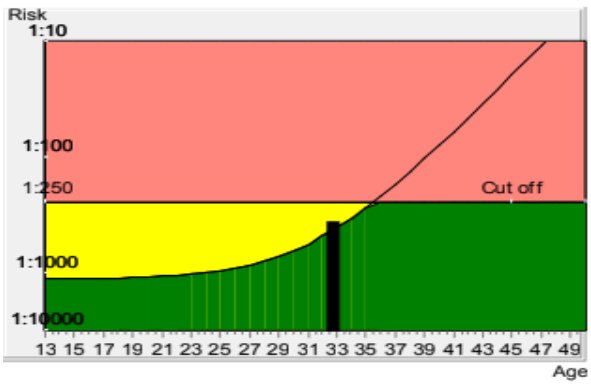
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	48	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data		
Parameter	**Value**	**Corr Mom**
PAPP-A	6.82 mIU/ml	0.85
fb-hCG	191.2 ng/ml	5.18

Risks at sampling date	
Age Risk	1:428
Biochemical T21 risk	1:57
Combined trisomy 21 risk	1:359
Trisomy 13/18 + NT	<1:10000

Ultrasound Data	
Gestational age	12+6
Method	CRL (<>Robinson)
Scan date	2/11/2022
Nuchal translucency	1.3
Nuchal translucency MoM	0.78
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 21with NT test it is expected that among 359 women with the same data, there is one woman with a trisomy 21 pregnancy and 358 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