MOLQ Express Lab
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Date of Report	13/3/2023
PRISCA	5.2.0.13

Patient Data			
Name	**MRS. PINKI DEVI**	Patient ID	012303120232
Birthday	30/5/1985	Sample ID	11408009
Age at term	38.2	Sample Date	12/3/2023
Gestational age	12+5		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	70	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	5.1 mIU/ml	1.05
fb-hCG	29.4 ng/ml	0.87

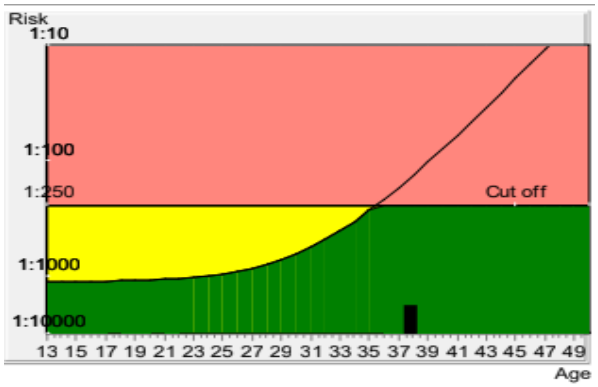
Risks at sampling date

Age Risk	1:142
Biochemical T21 risk	1:1342
Combined trisomy 21 risk	1:6492
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

	12+5
Method	CRL (<>Robinson)
Scan date	12/3/2023
Nuchal translucency	1.3
Nuchal translucency MoM	0.79
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 6492 women with the same data, there is one woman with a trisomy 21 pregnancy and 6491 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