MOLQ Express Lab

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Date of Report	30-08-2023
PRISCA	5.1.0.17

Patient Data

Name	MRS.PRIYANKA MANDAL	Patient ID	12308290133
Birthday	31-07-1993	Sample ID	11792883
Age at Sample	30.1	Sample Date	29-08-2023
Gestational age	12+2		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	69.3	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	3.26 mIU/ml	0.78
fb-hCG	42.3 ng/ml	1.11

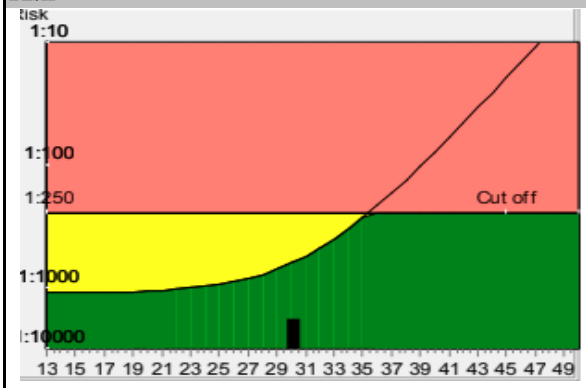
Risks at sampling date

Age Risk	1:634
Biochemical T21 risk	1:1795
Combined trisomy 21 risk	1:9661
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	12+2
Method	CRL (<>Robinson)
Scan date	29-08-2023
Crown rump length in mm	56.6
Nuchal transiucency MoM	0.54
Nasal Bone	Present
Sonographer	DR
Qualifications in measuring NT	MBBS

Risk



Trisomy 13/18 +NT

The calculated risk for Trisomy 13/18 (with nuchal translucency) 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 9661 women with the same data, there is one woman with a trisomy 21 pregnancy and 9660 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 risk calculations are statistical approaches and have no diagnostic value!
The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).

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