MOLQ Express Lab
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Date of Report	24-11-18
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs Tripti Rai**	Patient ID	011811220133
Birthday	17-05-89	Sample ID	10316226
Age at delivery	29.5	Sample Date	22/11/18
Gestational age	12+2		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	64.7	Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
PAPP-A	5.16	mIU/ml	1.61
fb-hCG	101.26	ng/ml	2.25

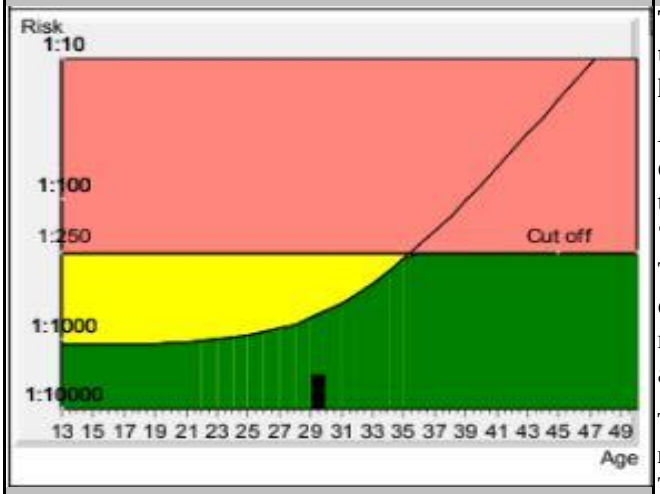
Risks at sampling date

Age Risk	1:678
Biochemical T21 Risk	1:1488
Combined Trisomy 21 Risk	1:7357
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	12+1
Method	CRL (<>Hadlock)
Scan Date	22-11-18
CRL	55
Nuchal translucency MoM	0.69
Nasal Bone	Present
Sonographer	DR.PRAKASH LALCHANDANI
Qualifications in measuring NT	MD

Risk



Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.

Down's Syndrome Risk (Trisomy 21 Screening)

The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 7357 women with the same data, there is one woman with a trisomy 21 pregnancy and 7356 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
The laboratory can not be held responsible for their impact on the risk assessment! Calculated value has no diagnostic value!

Risk Above Cut Off Risk above Age Risk Risk below Age risk