MOLQ Express Lab
*Free Home Sample Collection 9999 778 778
Download "MOLQ" App on
Book a Test Online www.molq.in

Date of Report	24/2/2022
PRISCA	5.2.0.13

Patient Data

Name	MRS. CHANDANI	Patient ID	012202230084
Birthday	15-1-1990	Sample ID	11261189
Age at term	32.1	Sample Date	23/02.2022
Gestational age	13+0		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	73.5	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	1.97 mIU/ml	0.47
fb-hCG	10.2 ng/ml	0.25

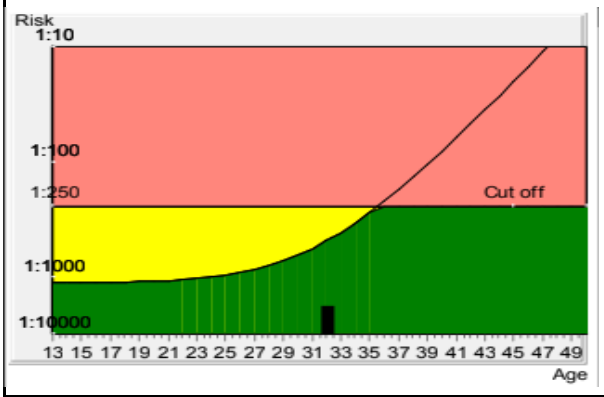
Risks at sampling date

Age Risk	1:484
Biochemical T21 risk	1:4409
Combined T21 risk	<1:10000
Trisomy 13/18+NT	1:2655

Ultrasound Data

Gestational age	13+0
Method	CRL (<>Robinson)
Scan date	23/2/2022
Nuchal translucency	1.5
Nuchal translucency MoM	0.89
Nasal Bone	Present

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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 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

Trisomy 13/18+NT

The calculated risk for Trisomy 13/18 with NT is 1:2655 , which indicates a low risk

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