![MOLQ Express Lab logo]

*Free Home Sample Collection 9999 778 778 | Download "MOLQ" App on (iOS, Android, Windows) | Book a Test Online www.molq.in

Date of Report		15/2/2022
PRISCA		5.2.0.13

Patient Data

Name	MRS. POOJA SHARMA	Patient ID	052202120057
Birthday	6/7/1995	Sample ID	11248950
Age at delivery	26.6	Sample Date	12/02.2022
Gestational age	12+4		

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	2.68	mIU/ml	0.7
fb-hCG	25.9	ng/ml	0.59

Risks at sampling date

Age Risk	1:884
Biochemical T21 risk	1:7722
Combined T21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	12+3
Method	CRL(<>Robinson
Scan date	12/2/2022
Nuchal translucency	1.4
Nuchal translucency MoM	0.91
Nasal Bone	Present
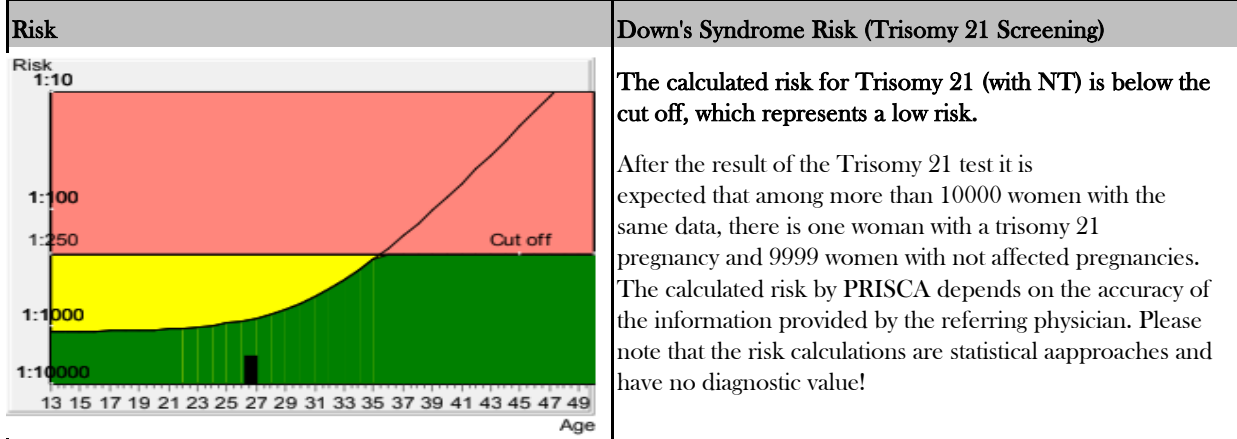
Risk

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 <1:10000, which indicates a low risk

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