MOLQ Express Lab
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Date of Report	28/7/2021
PRISCA	5.1.0.17

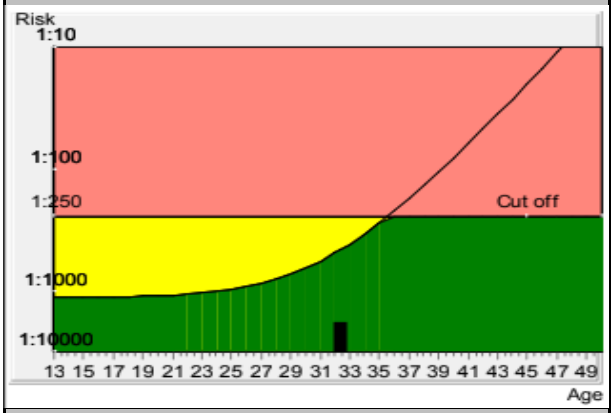
Patient Data			
Name	MRS. SANGAM SHARMA	Patient ID	012107240053
Birthday	1/3/1989	Sample ID	727305
Age at delivery	32.4	Sample Date	24/07/2021
Gestational age	13+0		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	57	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+6
PAPP-A	3.9 mIU/ml	0.69	Method	CRL(<>Robinson
fb-hCG	64.1 ng/ml	1.43	Scan date	23/07/2021
			Crown rump length in mm	64.6

Risks at sampling date			
Age Risk	1:461	Nuchal translucency MOM	0.61
Biochemical T21 risk	1:541	Nasal bone	UNKNOWN
Combined Trisomy 21 Risk	1:3119	Sonographer	DR.
Trisomy 13/18 + NT	<1:10000	Qualification in measuring NT	MD

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 21 test (with NT) it is expected that among 3119 women with the same data, there is one woman with a trisomy 21 pregnancy and 3118 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