MOLQ Express Lab
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Date of Report	21/3/2022
PRISCA	5.1.0.17

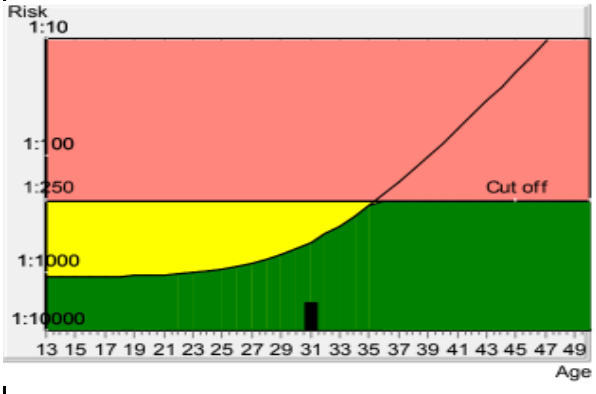
Patient Data			
Name	MRS. DHAVAL	Patient ID	012203190100
Birthday	12/2/1991	Sample ID	11261165
Age at term	31.07	Sample Date	19/3/2022
Gestational age	12+1		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	63	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+1
PAPP-A	2.18 mIU/ml	0.59	Method	CRL (<>Robinson)
fb-hCG	28.6 ng/ml	0.6	Scan date	19/3/2022
			Nuchal translucency	1.1
			Nuchal translucency MoM	0.76
			Nasal Bone	Presesnt

Risks at sampling date	
Age Risk	1:549
Biochemical T21 risk	1:3021
Combined trisomy 21 risk	<1:10000
Trisomy 13/18+NT	<1:10000

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 it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy 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

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