MOLQ Express Lab
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Date of Report	9/9/2021
PRISCA	5.2.0.13

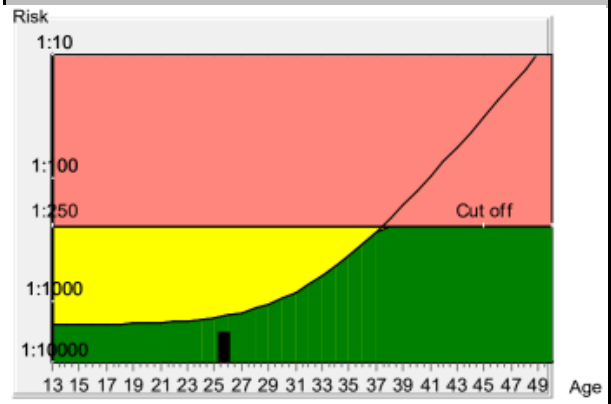
Patient Data			
Name	MRS. MAHIMA	Patient ID	012109080283
Birthday	23/06/1996	Sample ID	11090582
Age at delivery	25.74	Sample Date	8/9/2021
Gestational age	12+4		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	70	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+4
PAPP-A	3.02 mIU/ml	0.82	Method	CRL(<>Robinson
fb-hCG	61.2 ng/ml	1.5	Scan date	8/9/2021
			Nuchal translucency in mm	0.98

Risks at sampling date			
Age Risk	1:1336	Nuchal translucency MOM	0.59
Biochemical T21 risk	1:2069	Nasal bone	Present
Combined Trisomy 21 Risk	<1:10000	Qualification in measuring NT	CR
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 (with NT) 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

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