

Date of Report 10-05-2021
PRISCA 5.0.2.37

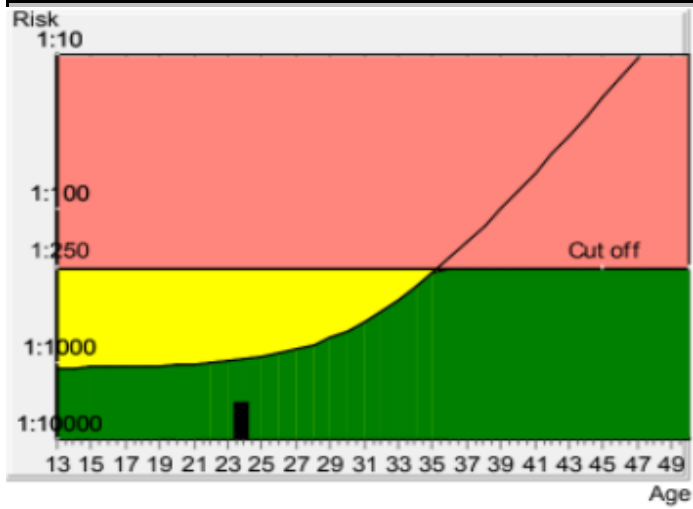
Patient Data			
Name	MRS MAMTA RANI	Patient ID	012105080014
Birthday	30-07-1997	Sample ID	10913117
Age at delivery	23.8	Sample Date	08/05/2021
Gestational age	11+4		

Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	64	Diabetes	unknown
Smoker	Unknown	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Ultrasound Data	
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Parameter	Value	Corr Mom		
PAPP-A	1.75 mIU/ml	0.62	Gestational age	11+2
fb-hCG	50.3 ng/ml	0.99	Method	CRL(<>Robinson
			Scan date	07-05-2021

Risks at sampling date				
Age Risk		1:972	Crown rump length in mm	44.8
Biochemical T21 risk		1:2008	Nuchal translucency MOM	0.97
Combined Trisomy 21 Risk		1:9143	Nasal bone	Present
Trisomy 13/18 + NT		<1:10000	Sonographer	DR.JYOTIKA
			Qualification in measuring NT	



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 9143 women with the same data, there is one woman with a trisomy 21 pregnancy and 9142 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 approaches and have no diagnostic value!

Trisomy 13/18 + NT	
The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk	The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values