

Date of Report 08-05-2021  
PRISCA 5.0.2.37

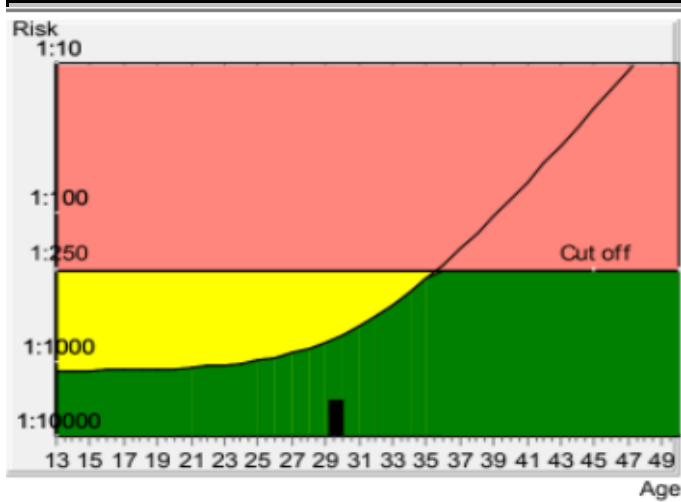
Patient Data			
Name	MRS SHALVI	Patient ID	012105070076
Birthday	16-09-1991	Sample ID	10905076
Age at delivery	29.6	Sample Date	09/05/2021
Gestational age	13+3		

Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	63	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	3.45 mIU/ml	0.59	Gestational age	13+3
fb-hCG	27.3 ng/ml	0.66	Method	CRL(<>Robinson
			Scan date	07-05-2021

Risks at sampling date				
Age Risk	1:696		Crown rump length in mm	73
Biochemical T21 risk	1:3128		Nuchal translucency MOM	0.61
Combined Trisomy 21 Risk	<1:10000		Nasal bone	Present
Trisomy 13/18 + NT	<1:10000		Sonographer	DR PRAKASH LALCHANDANI
			Qualification in measuring NT	MD



Down's Syndrome Risk (Trisomy 21 Screening)
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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 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

 Risk Above Cut Off

 Risk above Age Risk

 Risk below Age risk