

Date of Report 22-01-20  
PRISCA 5.0.2.37

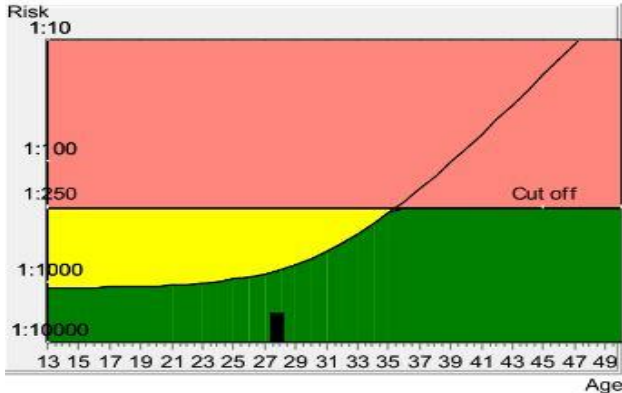
Patient Data			
Name	MRS PRIYA KUMARI	Patient ID	012001210093
Birthday	15-03-92	Sample ID	10510112
Age at delivery	27.9	Sample Date	21/1/2020
Gestational age	12+4		

Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	78	Diabetes	unknown
Smoker	Unknown	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+1
PAPP-A	4.65 mIU/ml	1.4	Method	CRL(<>Robinson
fb-hCG	123.5 ng/ml	2.88	Scan date	18-01-20

Risks at sampling date			Ultrasound Data	
Age Risk	1:807		Crown rump length in mm	55
Biochemical T21 risk	1:718		Nuchal translucency MOM	0.62
Combined Trisomy 21 Risk	1:3797		Nasal bone	Present
Trisomy 13/18 + NT	<1:10000		Sonographer	DR.SAHIL LOOMBA
			Qualification in measuring NT	MBBS,DNB

**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 3797 women with the same data, there is one woman with a trisomy 21 pregnancy and 3796 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