

Date of Report 30-05-2019
 PRISCA 5.0.2.37

Patient Data	
Name Mrs SHRISHTY	Patient ID 011905280053
Birth day 1/11/1991	Sample ID 10610880
Age at sample date 27.6	Sample Date 28/05/2019
Gestational age by CRL 12+1	

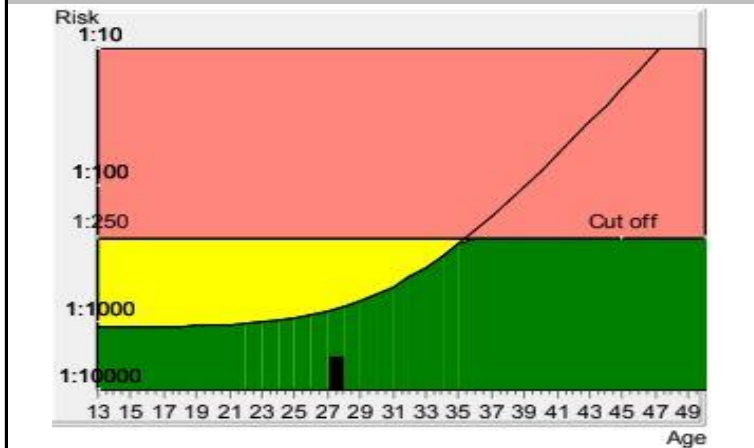
Correction factors			
Fetuses 1 IVF	unknown	Previous trisomy 21	unknown
Weight in kg 50	Diabetes	unknown	Pregnancies unknown
Smoker Unknown	Origin	Asian	

Biochemical Data	Ultrasound Data
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Parameter	Value	Corr Mom	
PAPP-A	3.56 mIU/ml	0.74	Method CRL (<>Robinson)
fb-hCG	46.32 ng/ml	0.89	Scan Date 27/05/2019

Risks at sampling date	
Age Risk 1:812	Crown rump length (mm) 52.6
Biochemical T21 Risk 1:3305	Nuchal translucency MoM 0.64
Combined Trisomy 21 Risk <1:10000	Nasal Bone Present
Trisomy 13/18 + NT <1:10000	Sonographer DR. PUNAM BAJAJ
	Qualifications in measuring NT MBBS.DNB

Risk	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.

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

