Date of Report	03-02-19
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs Meenakshi**	Patient ID	0119020200293
Birthday	08-08-85	Sample ID	10396666
Age at delivery	33.5	Sample Date	02/02/2019
Gestational age	13+2		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	61	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
PAPP-A	3.89	mIU/ml	0.79
fb-hCG	21.56	ng/ml	0.53

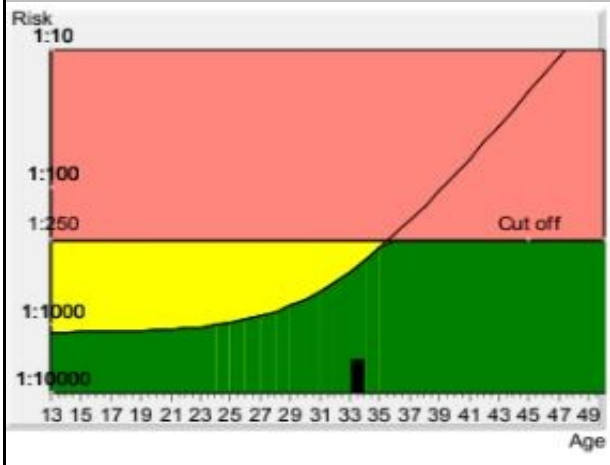
Ultrasound Data

Gestational age	13+2
Method	CRL (<>Hadlock)
Scan Date	02-02-19
CRL	70
Nuchal translucency MoM	0.68
Nasal Bone	Present
Sonographer	DR. PRAKASH LAL CHANDANI
Qualifications in measuring NT	MD

Risks at sampling date

Age Risk	1:382
Biochemical T21 Risk	1:5429
Combined Trisomy 21 Risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Risk



Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is <1:10000, which represents a low risk.

Down's Syndrome Risk (Trisomy 21 Screening)

The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 5429 women with the same data, there is one woman with a trisomy 21 pregnancy and 5428 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 risk calculations are statistical approach and have no diagnostic values!

The patient combined risk presumes the NT measurement was done according to accepted guidelines

The laboratory can not be held responsible for their impact on the risk assessment! Calculated value has no diagnostic value!

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