Date of Report 30/07/2019

PRISCA 5.0.2.37

Patient Data			
Name	Mrs Mehnaaz	Patient ID	011907280030
Birthday	19/06/1987	Sample ID	10604169
Age at delivery	32.1	Sample Date	28/07/19
Gestational age	13+6		

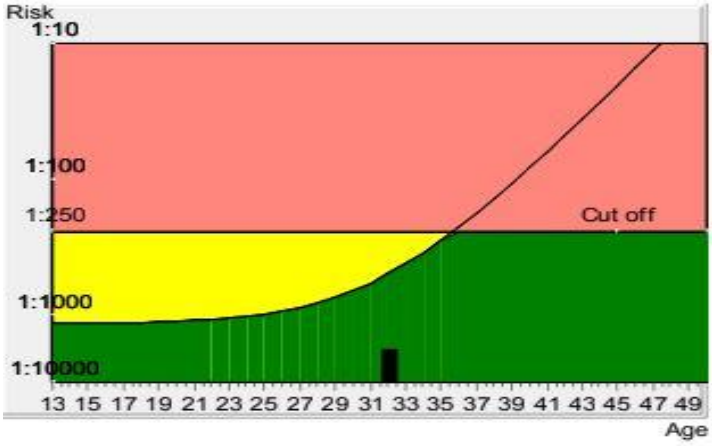
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	56	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data		
Parameter	**Value**	**Corr Mom**
PAPP-A	3.44 mIU/ml	0.45
fb-hCG	31.32 ng/ml	0.76

Risks at sampling date	
Age Risk	1:498
Biochemical Trisomy 21 Risk	1:791
Combined Trisomy 21 Risk	1:4338
Trisomy 13/18 + NT	<1:10000

Ultrasound Data	
Gestational age	13+5
Method	CRL (<>Robinson)
Scan Date	27/07/19
Crown Rump Length (mm)	79
Nuchal translucency MoM	0.89
Nasal Bone	present
Sonographer	Dr. Praveen Bhatia
Qualification in measuring NT	MBBS,MD

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)

TThe 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 4338 women with the same data, there is one woman with a trisomy 21 pregnancy and 4337 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 aapproaches and have no diagnostic value!

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