Date of Report	06-01-19
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs Pooja**	Patient ID	011901050061
Birthday	25-10-85	Sample ID	10408642
Age at delivery	33.2	Sample Date	05/01/2019
Gestational age	12+0		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	102	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
PAPP-A	3.15	mIU/ml	1.95
fb-hCG	25.37	ng/ml	0.62

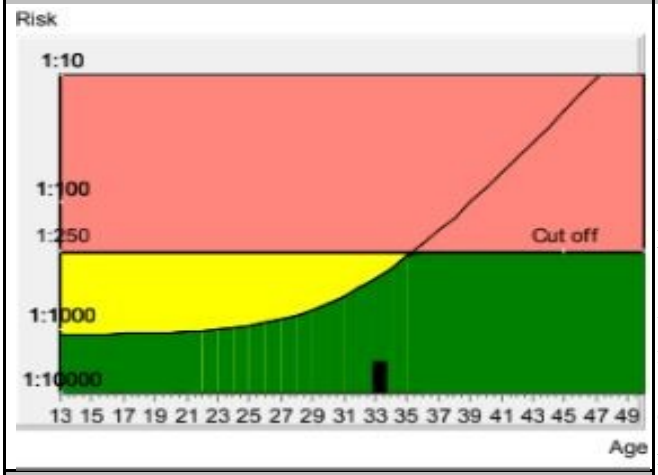
Risks at sampling date

Age Risk	1:384
Biochemical T21 Risk	<1:10000
Combined Trisomy 21 Risk	
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	12+0
Method	CRL (<>Hadlock)
Scan Date	28-11-18
CRL	
Nuchal translucency MoM	
Nasal Bone	Present
Sonographer	
Qualifications in measuring NT	

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 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 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