Date of Report 04-02-19
PRISCA 5.0.2.37

Patient Data			
Name	**Mrs NEHA PANJWANI**	Patient ID	011902020296
Birthday	19-08-89	Sample ID	10269862
Age at delivery	29.5	Sample Date	02/02/2019
Gestational age	12+5		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	73.5	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
PAPP-A	2.09	mIU/ml	0.65
fb-hCG	26.55	ng/ml	0.64

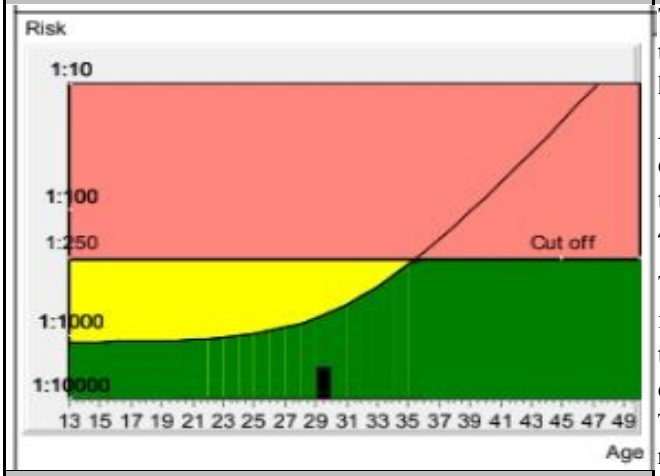
Risks at sampling date

Age Risk	1:693
Combined Trisomy 21 Risk	1:4066
Trisomy 13/18 + NT	1:7388

Ultrasound Data

Gestational age	12+5
Method	CRL (<>Hadlock)
Scan Date	24-12-18

Risk



Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is 1:7388, 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 4066 women with the same data, there is one woman with a trisomy 21 pregnancy and 4065 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