Date of Report	22-10-18
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs Kanchan**	Patient ID	011810190117
Birthday	01-01-86	Sample ID	10371365
Age at delivery	32.8	Sample Date	19-10-18
Gestational age	12+4		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	67	Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
PAPP-A	5.26	mIU/ml	1.42
fb-hCG	147.4	ng/ml	3.31

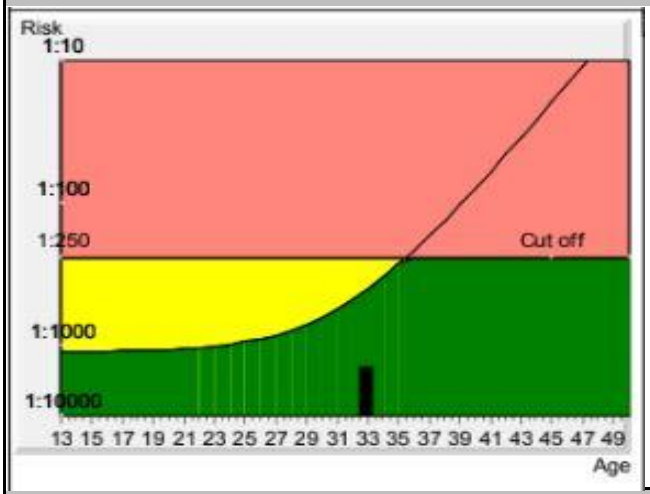
Risks at sampling date

Age Risk	1:423
Biochemical T21 Risk	1:265
Combined Trisomy 21 Risk	1:1423
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	11+5
Method	CRL (<>Robinson)
Scan Date	14-10-18
Crown rump length (mm)	52.7
Nuchal translucency MoM	0.57
Nasal Bone	Present
Sonographer	Dr. Harsha Sehgal
Qualifications in measuring NT	FMF, UK

Risk



Trisomy 13/18 + NT

The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates 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 represents a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 1423 women with the same data, there is one women with a trisomy 21 pregnancy and 1422 women with not affected pregnancies. The Free HCG level is high. 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 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