Date of Report	27-02-19
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs Champa**	Patient ID	011902260068
Birthday	01-01-97	Sample ID	10433699
Age at delivery	22.2	Sample Date	26/02/19
Gestational age by	13+2		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	55.4	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
PAPP-A	4.26	mIU/ml	0.77
fb-hCG	73.15	ng/ml	1.73

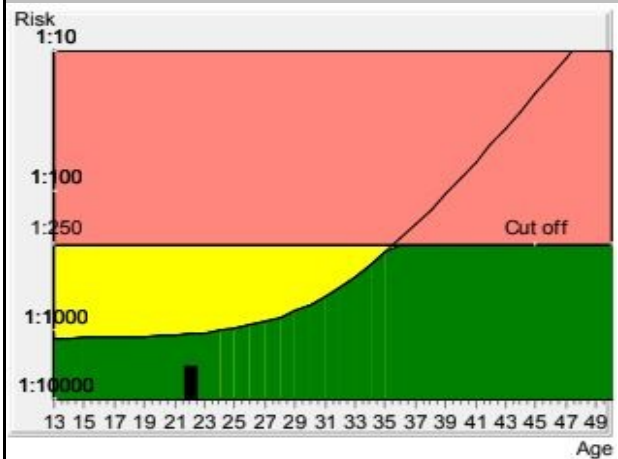
Ultrasound Data

Gestational age	13+2
Method	CRL (<>Hadlock)
Scan Date	26-02-19
CRL measurments	70.6
Nuchal translucency MoM	0.68
Nasal bone	Present
Sonographer	DR.SAHIL LOOMBA
Qualifications in measuring NT	MBBS,DNB

Risks at sampling date

Age Risk	1:1078
Biochemical T21 Risk	1:1041
Combined Trisomy 21 Risk	1:5979
Trisomy 13/18	<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 5979 women with the same data, there is one woman with a trisomy 21 pregnancy and 5978 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

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