Date of Report	06-04-19
PRISCA	5.0.2.37

Patient Data			
Name	**Mrs Komal**	Patient ID	011904650205
Birthday	10-10-97	Sample ID	10421621
Age at delivery	21.5	Sample Date	05/04/2019
Gestational age by	13+1		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	53	Diabetes	no	Pregnancies	
Smoker	no	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
PAPP-A	2.65	mIU/ml	0.48
fb-hCG	25.6	ng/ml	0.59

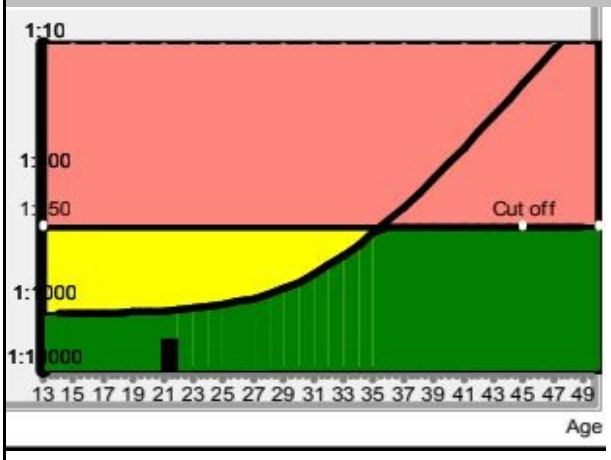
Risks at sampling date

Age Risk	1:1087
Biochemical T21 Risk	
Combined Trisomy 21 Risk	1:3496
Trisomy 13/18	1:3284

Ultrasound Data

Gestational age	13+1
Method	CRL (<>Hadlock)
Scan Date	18-03-19
CRL measurments	
Nuchal translucency MoM	
Nasal bone	Present
Sonographer	
Qualifications in measuring NT	

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 3496 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

Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is 1:3284, which represents a low risk.

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