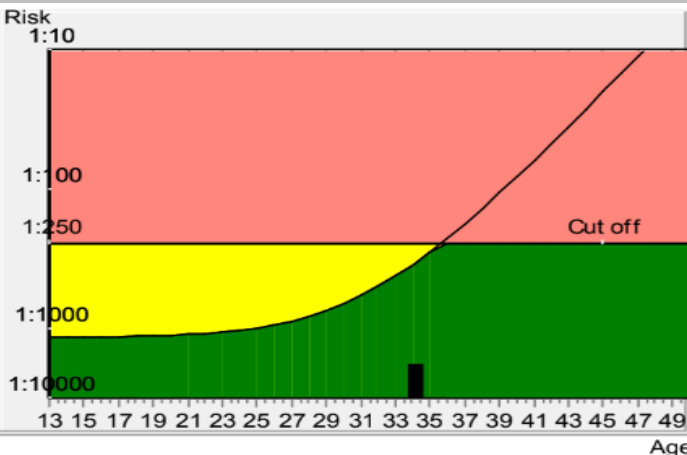


Date of Report 15/01/2020
PRISCA 5.0.2.37

Patient Data				
Name	Mrs Anju Verma	Patient ID	012001140165	
Birthday	14/11/1985	Sample ID	10632986	
Age at delivery	34.2	Sample Date	14/01/2020	
Gestational age	13+5			
Correction factors				
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown
Weight in kg	70	Diabetes	no	Pregnancies
Smoker	no	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+4
PAPP-A	2.65 mIU/ml	0.47	Method	CRL (<>Robinson)
fb-hCG	18.95 ng/ml	0.49	Scan Date	13/01/2020
Risks at sampling date			Crown Rump Length (mm)	74
Age Risk	1:338		Nuchal translucency MoM	0.66
Biochemical Trisomy 21 Risk	1:1513		Nasal Bone	present
Combined Trisomy 21 Risk	1:8240		Sonographer	DR. ROHIT CHANDAK
Trisomy 13/18 + NT	<1:10000		Qualification in measuring NT	MBBS, MD
Risk			Down's Syndrome Risk (Trisomy 21 Screening)	
			<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 8240 women with the same data, there is one woman with a trisomy 21 pregnancy and 8239 women with not affected pregnancies.</p> <p>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 approaches and have no diagnostic value!</p>	
Trisomy 13/18 + NT			The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values	
The calculated risk for trisomy 13/18 (with nuchal translucency) is <1:10000, which represents a low risk.				



Risk Above Cut Off



Risk above Age Risk



Risk below Age risk