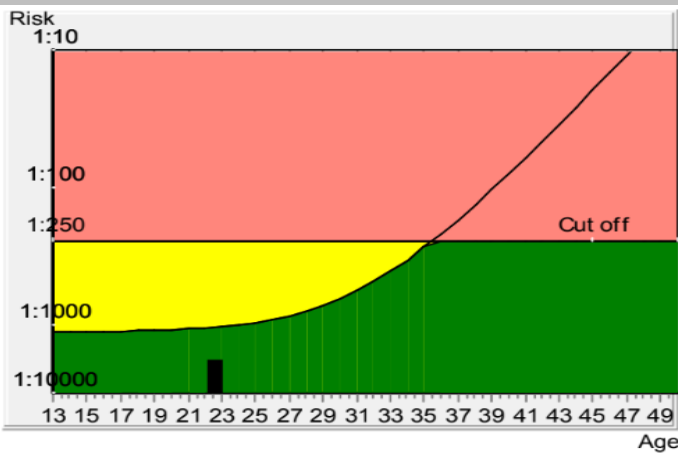


Date of Report 21/02/2020
PRISCA 5.0.2.37

Patient Data				
Name	Mrs Anita Devi	Patient ID	012002190256	
Birthdate	28/07/1997	Sample ID	10519624	
Age at delivery	22.6	Sample Date	19/02/2020	
Gestational age	12+5			
Correction factors				
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown
Weight in kg	50.8	Diabetes	no	Pregnancies
Smoker	no	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	11+5
PAPP-A	4.36 mIU/ml	0.75	Method	CRL (<>Robinson)
fb-hCG	41.8 ng/ml	0.86	Scan Date	13/02/2020
Risks at sampling date			Crown Rump Length (mm)	48.3
Age Risk	1:1048		Nuchal translucency MoM	1.60
Biochemical Trisomy 21 Risk	1:4637		Nasal Bone	present
Combined Trisomy 21 Risk	1:2295		Sonographer	DR. GULSHAN
Trisomy 13/18 + NT	<1:10000		Qualification in measuring NT	
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 2295 women with the same data, there is one woman with a trisomy 21 pregnancy and 2294 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