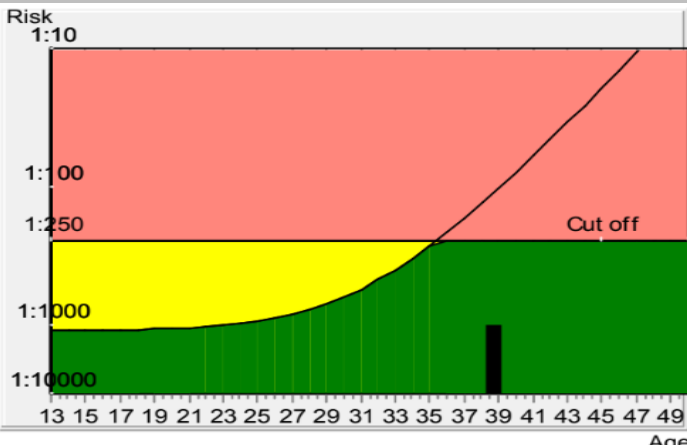


Date of Report 25/02/2020  
PRISCA 5.0.2.37

Patient Data				
Name	Mrs Sumera	Patient ID	012002240108	
Birthday	6/6/1981	Sample ID	10582900	
Age at delivery	38.7	Sample Date	24/02/2020	
Gestational age	12+1			
Correction factors				
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown
Weight in kg	68	Diabetes	no	Pregnancies
Smoker	no	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+1
PAPP-A	5.48 mIU/ml	1.64	Method	CRL (<>Robinson)
fb-hCG	102.4 ng/ml	2.19	Scan Date	24/02/2020
Risks at sampling date			Crown Rump Length (mm)	58
Age Risk		1:109	Nuchal translucency MoM	0.99
Biochemical Trisomy 21 Risk		1:267	Nasal Bone	present
Combined Trisomy 21 Risk		1:1014	Sonographer	DR. EKTA YADAV
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	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 1014 women with the same data, there is one woman with a trisomy 21 pregnancy and 1013 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