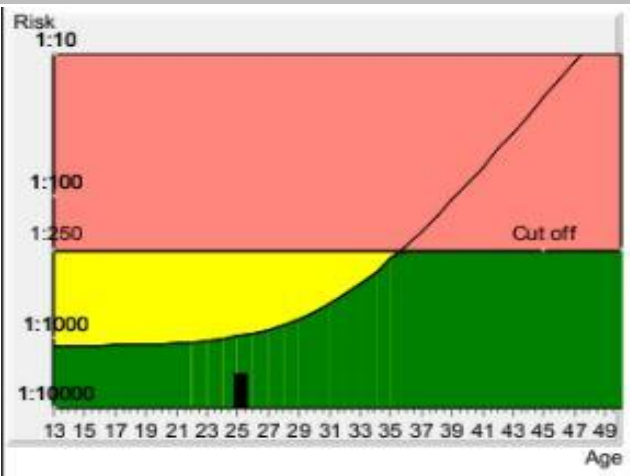



Date of Report 31-10-18
PRISCA 5.0.2.37

Patient Data				
Name	Mrs Pooja W/o Sandeep Featus 2	Patient ID	011810310055	
Birthday	24-07-93	Sample ID	010337532	
Age at delivery	25.3	Sample Date	29/10/18	
Gestational age	13+4			
Correction factors				
Fetuses	2	IVF	unknown	Previous trisomy 21
Weight in kg	65.3	Diabetes	unknown	Pregnancies
Smoker	Unknown	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+3
PAPP-A	3.87 mIU/ml	0.42	Method	CRL (<>Robinson)
fb-hCG	36.86 ng/ml	0.44	Scan Date	29-10-18
Risks at sampling date			Crown rump length (mm)	73
Age Risk		1:983	Nuchal translucency MoM	0.5
Biochemical T21 Risk		1:3790	Nasal Bone	Present
Combined Trisomy 21 Risk		<1:10000	Sonographer	DR.SURENDER SINGH
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MBBS,MD
Risk			Down's Syndrome Risk (Trisomy 21 Screening)	
			<p>The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 3790 with the same data, there is one women with a trisomy 21 pregnancy and 3789 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> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines.</p>	
Trisomy 13/18 + NT			The laboratory can not be held responsible for their impact on the risk assessment! Calculated value has no diagnostic value!	
<p>The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk</p>				

 Risk Above Cut Off

 Risk above Age Risk

 Risk below Age risk