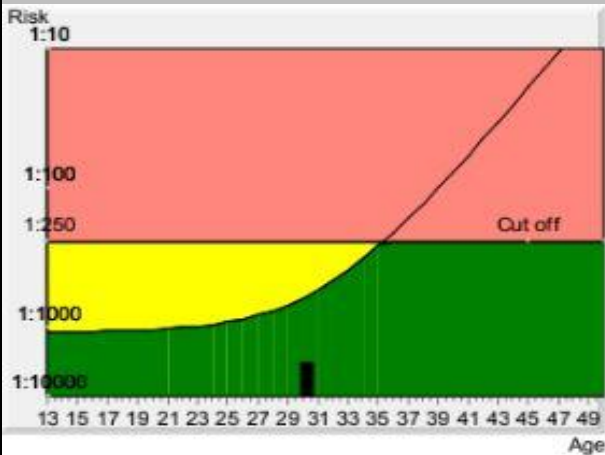


Date of Report 02-12-18
PRISCA 5.0.2.37

Patient Data				
Name	Mrs Suman w/o Gorav		Patient ID	011811300210
Birthday	12-09-88		Sample ID	10422477
Age at delivery	30.2		Sample Date	30/11/18
Gestational age	12+0			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21 unknown
Weight in kg	52.4	Diabetes	no	Pregnancies
Smoker	no	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+5
PAPP-A	3.15 mIU/ml	0.96	Method	CRL (<>Hadlock)
fb-hCG	47.69 ng/ml	0.93	Scan Date	29-11-18
Risks at sampling date			CRL	46.4
Age Risk		1:609	Nuchal translucency MoM	0.47
Biochemical T21 Risk		1:4164	Nasal Bone	Present
Combined Trisomy 21 Risk		<1:10000	Sonographer	DR.ANITA NARULA
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 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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</p> <p>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!</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 nuchal translucency) is < 1:10000, which represents a low risk.</p>				

Risk Above Cut Off

Risk above Age Risk

Risk below Age risk