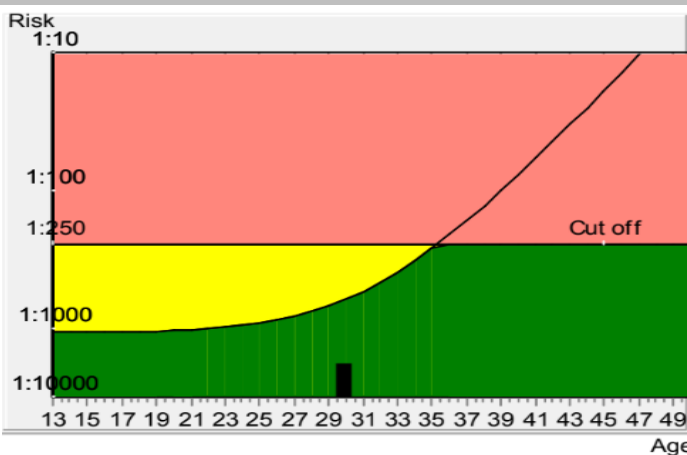


Date of Report 18/02/2020
PRISCA 5.0.2.37

Patient Data				
Name	Mrs Sweccha Mishra	Patient ID	062002160055	
Birthday	24/04/1990	Sample ID	10658549	
Age at delivery	29.8	Sample Date	16/02/2020	
Gestational age	11+3			
Correction factors				
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown
Weight in kg	56.3	Diabetes	no	Pregnancies
Smoker	no	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	11+0
PAPP-A	3.15 mIU/ml	1.01	Method	CRL (<>Robinson)
fb-hCG	22.9 ng/ml	0.42	Scan Date	14/02/2020
Risks at sampling date			Crown Rump Length (mm)	40
Age Risk		1:633	Nuchal translucency MoM	0.84
Biochemical Trisomy 21 Risk		<1:10000	Nasal Bone	present
Combined Trisomy 21 Risk		<1:10000	Sonographer	DR. N.S. PIPLANI
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	FIAMS, PCMS
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 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