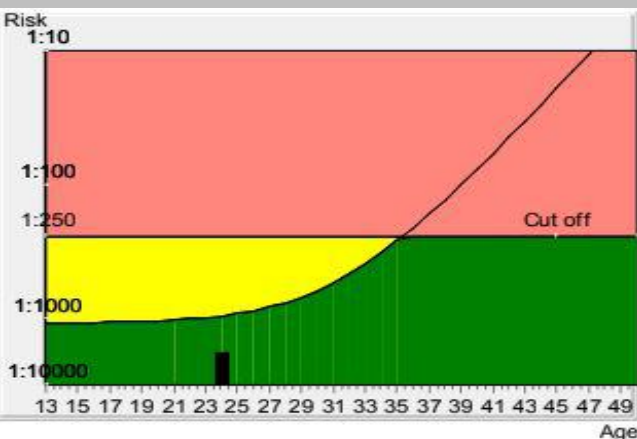


Date of Report
PRISCA

22/06/2019
5.0.2.37

Patient Data				
Name	Mrs. Ashiyana	Patient ID	011906210107	
Birthday	20/05/1995	Sample ID	10362239	
Age at delivery	24.1	Sample Date	21/06/2019	
Gestational age	11+5			
Correction factors				
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown
Weight in kg	60 Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown Origin	Asian		
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	11+5
PAPP-A	3.12 mIU/ml	0.96	Method	CRL (<>Robinson)
fb-hCG	53.8 ng/ml	1.05	Scan Date	21/06/2019
Risks at sampling date			Crown rump length in mm	50.1
Age Risk	1:967		Nuchal translucency MoM	0.96
Overall population risk	1:4878		Nasal bone	Present
Combined Trisomy 21 Risk	<1:10000		Sonographer	DR.SANJEEV KUMAR SINGHAL
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT	MBBS,PGDUS,DMRD
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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 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 NT) is <1:10000, which indicates a low risk				

 Risk Above Cut Off

 Risk above Age Risk

 Risk below Age risk