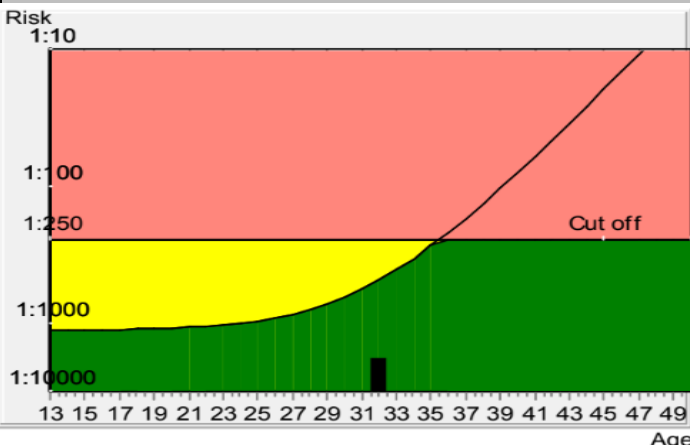


Date of Report 18/02/2020  
PRISCA 5.0.2.37

Patient Data				
Name	Mrs Twinky Khetarpal	Patient ID	012002160127	
Birthday	24/03/1988	Sample ID	DPLTA00089770	
Age at delivery	31.9	Sample Date	16/02/2020	
Gestational age	12+5			
Correction factors				
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown
Weight in kg	67	Diabetes	no	Pregnancies
Smoker	no	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+3
PAPP-A	4.5 mIU/ml	1.07	Method	CRL (<>Robinson)
fb-hCG	28.2 ng/ml	0.64	Scan Date	15/02/2020
Risks at sampling date			Crown Rump Length (mm)	59.1
Age Risk	1:496		Nuchal translucency MoM	0.62
Biochemical Trisomy 21 Risk	1:9272		Nasal Bone	present
Combined Trisomy 21 Risk	<1:10000		Sonographer	DR. DIVYA KANT
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 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