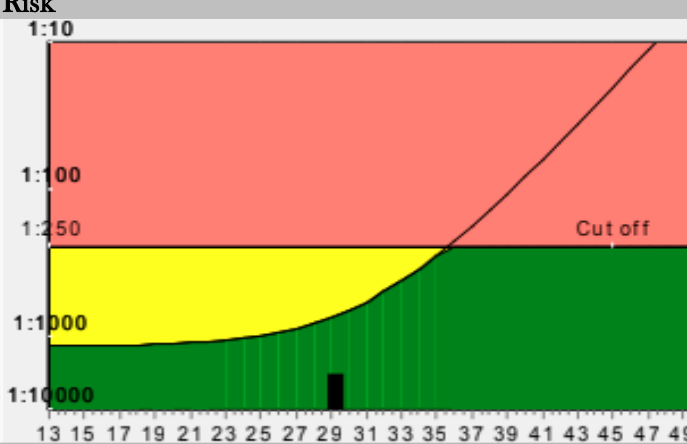


Date of Report 29/09/2023
PRISCA 5.2.0.13

Patient Data				
Name	SHEETAL	Patient ID	012309280225	
Birth day	26/07/1994	Sample ID	11781119	
Age at sample	29.2	Sample Date	28/09/2023	
Gestational age	13+6			
Correction factors				
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown
Weight in kg	60 Diabetes	NO	Pregnancies	unknown
Smoker	NO Origin	Asian		
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+6
PAPP-A	4.35 mIU/ml	0.50	Method	CRL (<>Robinson)
fb-hCG	16.8 ng/ml	0.68	Scan date	28/09/2023
Risks at sampling date			Crown rump length in mm	81
Age Risk	1:743		Nuchal translucency MoM	0.96
Biochemical T21 risk	1:2028		Nasal bone	Present
Combined trisomy 21 risk	1:9530		Sonographer	DR SHRUTI SANGWAN
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT	MBBS
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 9530 women with the same data, there is one woman with a trisomy 21 pregnancy and 9529 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 that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998).</p> <p>The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values</p>	
Trisomy 13/18 + NT				
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