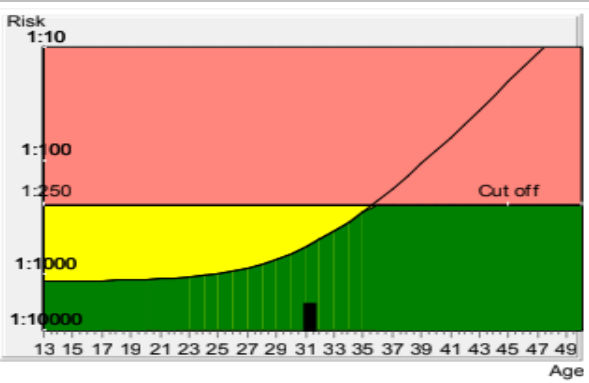




Date of Report 29/11/2022
PRISCA 5.1.0.17

Patient Data				
Name	MRS. SHWETA GOYAL	Patient ID	022211290008	
Birthday	7/8/1991	Sample ID	11516722	
Age at term	31.09	Sample Date	27/11/2022	
Gestational age	13+5			
Correction factors				
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown
Weight in kg	46	Diabetes	NO	Pregnancies
Smoker	NO	Origin	Asian	unknown
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+4
PAPP-A	5.26 mIU/ml	0.47	Method	CRL (<>Robinson)
fb-hCG	25.5 ng/ml	0.89	Scan date	28/11/2022
Risks at sampling date			Nuchal translucency	1.5
Age Risk	1:562		Nuchal translucency MoM	0.82
Biochemical T21 risk	1:713		Nasal bone	Present
Combined trisomy 21 risk	1:4221			
Trisomy 13/18 + NT	<1:10000			
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 with NT test it is expected that among 4221 women with the same data, there is one woman with a trisomy 21 pregnancy and 4220 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 held 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