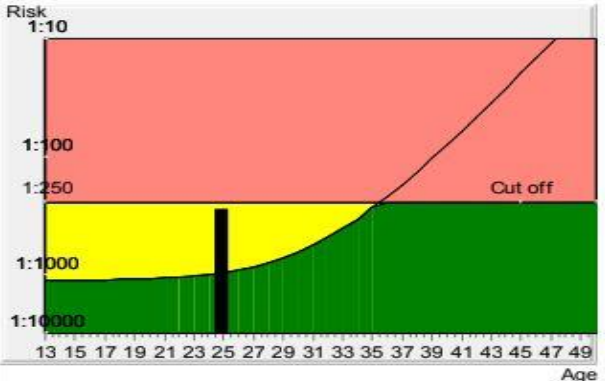



Date of Report 03-12-19  
PRISCA 5.0.2.37

Patient Data					
Name	<b>MRS LAXMI</b>	Patient ID	021912010013		
Birth day	10-02-95	Sample ID	10507378		
Age at delivery	24.8	Sample Date	1/12/2019		
Gestational age	12+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		
<b>Parameter</b>	<b>Value</b>	<b>Corr Mom</b>	Gestational age	12+4	
PAPP-A	5.13 mIU/ml	1.06	Method	CRL (<>Robinson)	
fb-hCG	28.2 ng/ml	0.62	Scan Date	01-12-19	
Risks at sampling date			Crown rump length in mm	62.2	
Age Risk	1:976		Nuchal translucency MoM	2.44	
Biochemical T21 risk	<1:10000		Nasal Bone	Present	
Combined Trisomy 21 Risk	1:279		sonographer	DR.VARUN SHARMA	
Trisomy 13/18 + NT	1:708		Qualification in measuring NT		
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 279 women with the same data, there is one woman with a trisomy 21 pregnancy and 278 women with not affected pregnancies. 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><b>Trisomy 13/18 + NT</b> The calculated risk for Trisomy 13/18 (with NT) is 1:708, which indicates a low risk</p>			<p>The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values</p>		

 Risk Above Cut Off

 Risk above Age Risk

 Risk below Age risk