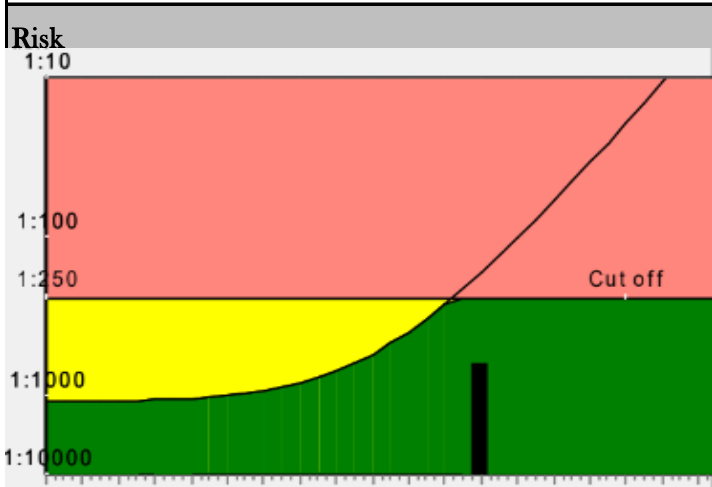





Date of Report 31-03-2024  
PRISCA 5.2.0.13

Patient Data					
Name	MRS. SNEHA		Patient ID	12403290298	
Birthday	25-04-1987		Sample ID	11844457	
Age at Sample date	36.9		Sample Date	29-03-2024	
Gestational age	12+1				
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	69	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			
PAPP-A	4.9 mIU/ml	1.24	Gestational age	12+0	
fb-hCG	81.8 ng/ml	2.07	Method	CRL (<>Robinson)	
Risks at sampling date			Scan date	29-03-2024	
Age Risk	1:171		Crown rump length in mm	54	
Biochemical T21 risk	1:288		Nuchal translucency MoM	1.18	
Combined trisomy 21 risk	1:642		Nasal bone	PRESENT	
Trisomy 13/18 + NT	<1:10000		Sonographer	DR. RUBY RAHUL	
Risk			Qualifications in measuring NT		
			Down's Syndrome Risk (Trisomy 21 Screening)		
<b>Trisomy 13/18+NT</b> The calculated risk for Trisomy 13/18 (with NT) is <1:10000 , which indicates a low risk			The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 642 women with the same data, there is one woman with a trisomy 21 pregnancy and 641 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! The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998). The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		

 Risk Above Cut Off     
  Risk above Age Risk     
  Risk below Age risk