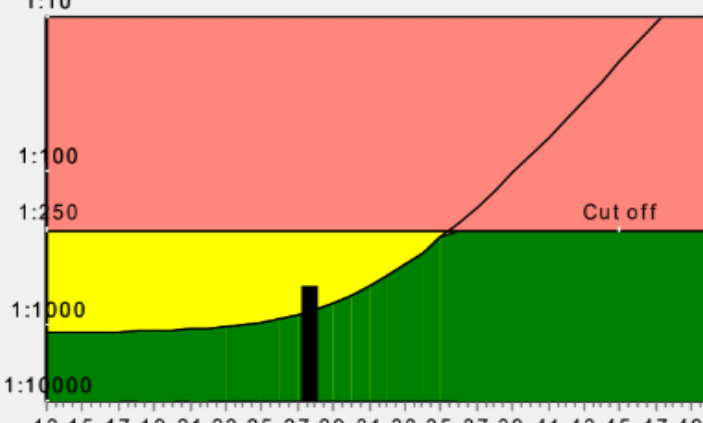


Date of Report 13-03-2024  
PRISCA 5.2.0.13

Patient Data				
Name	<b>MRS. MANISHA</b>		Patient ID	12403120079
Birthday	02-07-1996		Sample ID	11810109
Age at Sample date	27.7		Sample Date	12-03-2024
Gestational age	12+5			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21 unknown
Weight in kg	42.9	Diabetes	NO	Pregnancies unknown
Smoker	NO	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+4
PAPP-A	6.8 mIU/ml	0.79	Method	CRL (<>Robinson)
fb-hCG	162.5 ng/ml	4.05	Scan date	11-03-2024
Risks at sampling date			Crown rump length in mm	61.2
Age Risk	1:821		Nuchal translucency MoM	0.82
Biochemical T21 risk	1:93		Nasal bone	PRESENT
Combined trisomy 21 risk	1:572		Sonographer	DR.
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT	MBBS
Risk			Down's Syndrome Risk (Trisomy 21 Screening)	
			<p><b>The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.</b></p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 572 women with the same data, there is one woman with a trisomy 21 pregnancy and 571 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>	
Trisomy 13/18+NT			The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values	
<p>The calculated risk for Trisomy 13/18 (with NT) is &lt;1:10000 , which indicates a low risk</p>				

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