



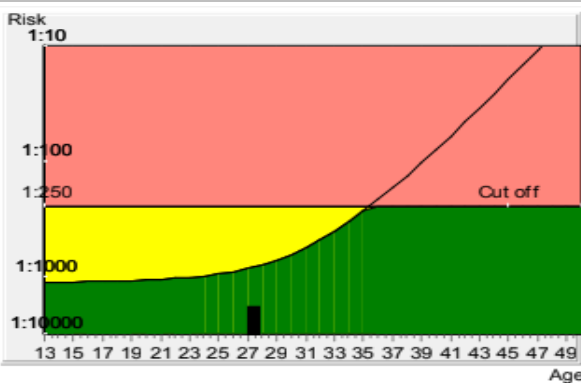
Date of Report 2/2/2023  
PRISCA 5.1.0.17

Patient Data			
Name	MRS. VARSHA	Patient ID	012301310096
Birthday	21/8/1995	Sample ID	11408275
Age at term	27.1	Sample Date	31/1/2023
Gestational age	12+3		

Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	61	Diabetes	NO
Smoker	NO	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom		12+2
PAPP-A	5.72 mIU/ml	1.11	Method	CRL (<>Robinson)
fb-hCG	95.2 ng/ml	2.49	Scan date	30/01/2023

Risks at sampling date		Nuchal translucency	
Age Risk	1:829	Nuchal translucency	1.4
Biochemical T21 risk	1:696	Nuchal translucency MoM	0.89
Combined trisomy 21 risk	1:3436	Nasal bone	Present
Trisomy 13/18 + NT	<1:10000		

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 with NT test it is expected that among 3436 women with the same data, there is one woman with a trisomy 21 pregnancy and 3435 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 calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk

The laboratory cannot be held 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