

Date of Report 06-01-2025  
PRISCA 5.2.0.13

Patient Data			
Name	MRS. VARSHA	Patient ID	12501050125
Birthday	25-08-2002	Sample ID	11992319
Age at Sample date	22.4	Sample Date	05-01-2025
Gestational age	13+1		

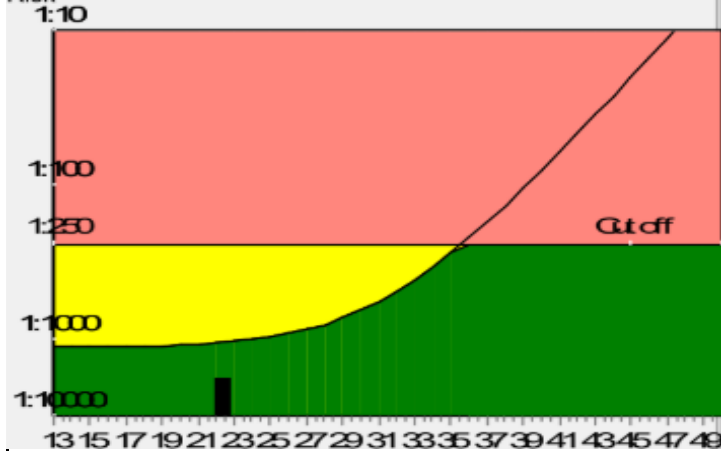
Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	51	Diabetes	NO
Smoker	NO	Origin	Asian
Previous trisomy 21		Pregnancies	unknown

Biochemical Data			Ultrasound Data	
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Parameter	Value	Corr Mom		
PAPP-A	6.3 mIU/ml	0.76	Gestational age	13+0
fb-hCG	12.5 ng/ml	0.38	Method	CRL (<>Robinson)
			Scan date	04-01-2025

Risks at sampling date			Ultrasound Data	
Age Risk		1:1068	Crown rump length in mm	67
Biochemical T21 risk		<1:10000	Nuchal translucency MoM	1.00
Combined trisomy 21 risk		<1:10000	Nasal bone	PRESENT
Trisomy 13/18 + NT		<1:10000	Sonographer	DR. HARENDER BHASKAR
			Qualifications in measuring NT	MBBS

Risk	Down's Syndrome Risk (Trisomy 21 Screening)
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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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. The free beta HCG level is low.

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).

<b>Trisomy 13/18+NT</b>	The laboratory cannot be hold 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