

Date of Report 25-01-2024  
PRISCA 5.2.0.13

Patient Data			
Name	MRS NIKITA	Patient ID	012401230175
Birthday	01-09-1997	Sample ID	11819963
Age at Sample date	26.4	Sample Date	23-01-2024
Gestational age	12+6		

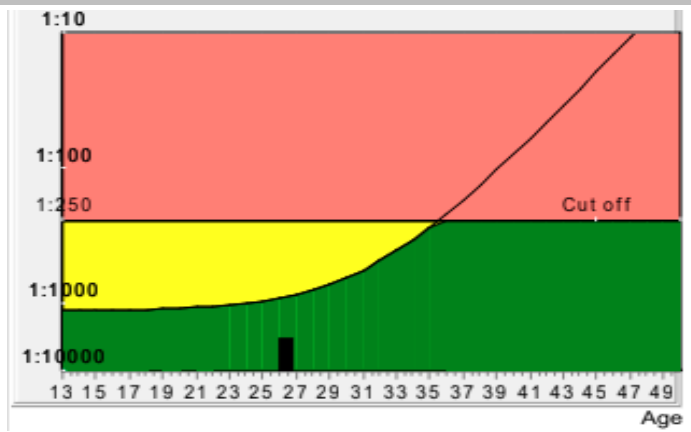
Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	67	Diabetes	NO
Smoker	NO	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data	Ultrasound Data
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Parameter	Value	Corr Mom		
PAPP-A	4.9 mIU/ml	0.91	Gestational age	12+5
fb-hCG	28.1 ng/ml	0.85	Method	CRL (<>Robinson)
			Scan date	23-01-2024

Risks at sampling date		Ultrasound Data	
Age Risk	1:904	Crown rump length in mm	63
Biochemical T21 risk	1:6484	Nuchal translucency MoM	0.68
Combined trisomy 21 risk	<1:10000	Nasal bone	PRESENT
Trisomy 13/18 + NT	<1:10000	Sonographer	DR DHRUV TANEJA
		Qualifications in measuring NT	MD

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 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 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 calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk

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