

Date of Report 06-09-2023  
PRISCA 5.2.0.13

Patient Data			
Name	MRS BHAWNA	Patient ID	12309050077
Birthday	13-11-1999	Sample ID	11059384
Age at Sample date	23.8	Sample Date	05-09-2023
Gestational age	13+2		

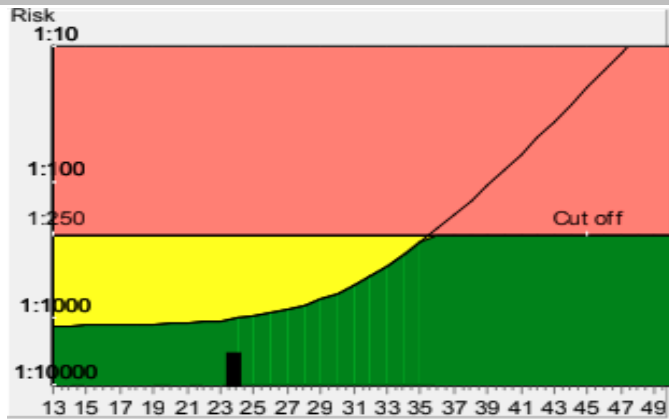
Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	62	Diabetes	NO
Smoker	NO	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Ultrasound Data	
------------------	--	--	-----------------	--

Parameter	Value	Corr Mom		
PAPP-A	4.87 mIU/ml	0.71	Gestational age	13+1
fb-hCG	61.2 ng/ml	2.06	Method	CRL (<>Robinson)
			Scan date	04-09-2023

Risks at sampling date			Ultrasound Data	
Age Risk		1:1032	Crown rump length in mm	69.5
Biochemical T21 risk		1:522	Nuchal translucency MoM	1.03
Combined trisomy 21 risk		1:2054	Nasal bone	PRESENT
Trisomy 13/18 + NT		<1:10000	Sonographer	DR RANJAN KUMAR
			Qualifications in measuring NT	MD

Risk	Down's Syndrome Risk (Trisomy 21 Screening)
------	---



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 2054 women with the same data, there is one woman with a trisomy 21 pregnancy and 2053 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).

Trisomy 13/18+NT	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