

Date of Report 16-10-2023
PRISCA 5.2.0.13

Patient Data			
Name	MRS SHABANA	Patient ID	012310140189
Birthday	10-02-1995	Sample ID	11670488
Age at Sample date	28.7	Sample Date	14-10-2023
Gestational age	12+2		

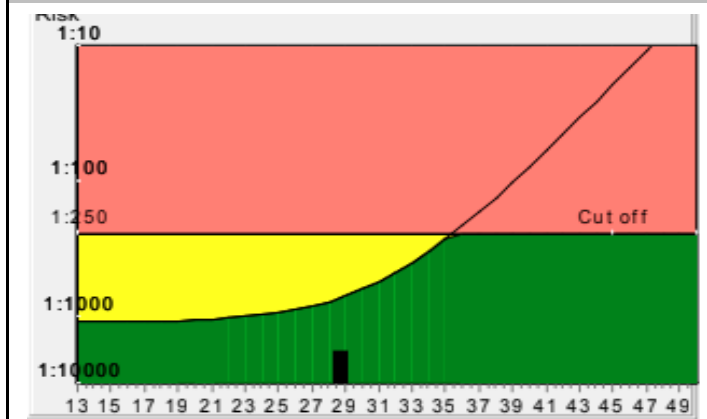
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	68	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data		
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Parameter	Value	Corr Mom			
PAPP-A	3.12 mIU/ml	0.73	Gestational age	11+6	
fb-hCG	16.8 ng/ml	0.44	Method	CRL (<>Robinson)	
			Scan date	12-10-2023	

Risks at sampling date			Ultrasound Data		
Age Risk	1:741		Crown rump length in mm	52.4	
Biochemical T21 risk	<1:10000		Nuchal translucency MoM	0.57	
Combined trisomy 21 risk	<1:10000		Nasal bone	PRESENT	
Trisomy 13/18 + NT	<1:10000		Sonographer	DR PONAM	
			Qualifications in measuring NT		

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 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