

*Free Home Sample Collection 9999 778 778



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Date of Report 25-11-2024 PRISCA 5.2.0.13

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Patient Data						
Name	MRS. MONIKA			Patient ID	12411240015	
Birthday			17-12-1990	Sample ID		11895110
Age at Sample date		33.9			Sample Date	
Gestational age			12+5			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	62.9	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		

Weight in kg 62.9 Diabetes		NC		Pregnancies	unknown
Smoker NO Origin			Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+4
PAPP-A	APP-A 5.7 mIU/ml		Method		CRL (<>Robinson)
fb-hCG	29.3 ng/ml	0.84	Scan date		23-11-2024
Risks at sampling date			Crown rump le	ength in mm	61.4
Age Risk		1:342	Nuchal translucency MoM		1.13
Biochemical T21 risk	1:3373	Nasal bone PRES		PRESENT	
Combined trisomy 21 risk 1:9304			Sonographer DR.		
Trisomy 13/18 + NT <1:10000			Qualifications	in measuring NT	MBBS
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off			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 9304 women with the same data, there is one woman with a trisomy 21 pregnancy and 9303 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 aapproaches and have no diagnostic value!		

Trisomy 13/18+NT

1:10000

The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk

13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49

done according to accepted guidelines (Prenat Diagn 18:511-523; 1998).

The patient combined risk presumes that NT measurement was

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values