

which indicates a low risk

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Date of Report 01-07-2024 PRISCA 5.2.0.13

				PRISCA	5.2.0.13
Patient Data					
Name	MRS. K.ANITA		Patient ID		12406300021
Birthday		06-06-2001	Sample ID		11814067
Age at Sample date	23.1		Sample Date		30-06-2024
Gestational age		13+2			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	53 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian	ı	
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational ag	re	13+1
PAPP-A	6.1  mIU/ml	0.74	Method		CRL (<>Robinson)
fb-hCG	26.5 ng/ml	0.85	Scan date		29-06-2024
Risks at sampling date			Crown rump length in mm 67.9		
Age Risk		1:1055	Nuchal translu	icency MoM	0.58
Biochemical T21 risk 1:4746		1:4746	Nasal bone PRF		PRESENT
Combined trisomy 21 risk <1:10000		<1:10000	Sonographer		DR.
Trisomy 13/18 + NT <1:10000		Qualifications in measuring NT MB		MBBS	
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:10 1:100			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.		
1:1000 1:1(000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			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!  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 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		