

which indicates a low risk

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Date of Report 17-09-2024

					PRISCA	5.2.0.13
Patient Data						
Name	MRS. NEHA W/O TILAK RAJ			Patient ID		12409160125
Birthday	thday 12-07-2001					11966942
Age at Sample date 23.2				Sample Date 16-09-2024		
Gestational age 13+5				;		
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	44.9	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	e	13+5
PAPP-A	7.5 1	mIU/ml	0.65	Method		CRL (<>Robinson)
fb-hCG	24.5 1	ng/ml	0.85	Scan date		16-09-2024
Risks at sampling date				Crown rump length in mm 77		
Age Risk			1:1065	Nuchal translucency MoM 0.80		
Biochemical T21 risk			1:3526	Nasal bone PRESEN		PRESENT
Combined trisomy 21 risk <1:10000				Sonographer DR. AMEND		DR. AMENDA
Trisomy 13/18 + NT <1:10000			<1:10000	Qualifications in measuring NT MD		
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 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 than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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		
1:1(000				diagnostic value! The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523;		
13 15 17 19 21 23 25 Trisomy 13/18+NT	35 37 39	41 43 45 47 49	1998).			
The calculated risk for Trisomy 13/18 (with NT) is <1:10000,				The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		