

which indicates a low risk

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Date of Report 26-11-2024 PRISCA 5 9 0 13

			PRISCA		5.2.0.13
Patient Data					
Name		MRS. NEHA YADAV	Patient ID		12411250105
Birthday		02-02-1990	Sample ID		11889178
Age at Sample date		34.8	Sample Date		25-11-2024
Gestational age		12+1			
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	59	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Sillokei	NO Oligili		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	12+1	
PAPP-A	$3.9~\mathrm{mIU/ml}$	0.82	Method	CRL (<>Robinson)	
fb-hCG	45.2 ng/ml	1.09	Scan date	25-11-2024	
Risks at sampling date			Crown rump length in mm 56.1		
Age Risk		1:279	Nuchal translucency MoM	0.72	
Biochemical T21 risk		1:916	Nasal bone	PRESENT	
Combined trisomy 21 risk		1:4861	Sonographer	DR. INDRAJEET	
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD	
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:10 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 4861 women with the same data, there is one woman with a trisomy 21 pregnancy and 4860 women with not affected pregnancies.		
1:1 <mark>0000</mark> 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 4			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 Trison	my 13/18 (with NT)	is <1.10000	The laboratory cannot be hold responsible for their impact on the		
The calculated risk for Trisomy 13/18 (with NT) is <1:10000,			risk assessment! Calculated risks have no diagnostic values		