

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

which indicates a low risk

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The laboratory cannot be hold responsible for their impact on the

risk assessment! Calculated risks have no diagnostic values

Date of Report 18-09-2024

				PRISCA		5.2.0.13
Patient Data						
Name	RS. KALPANA DEVI W/O NEERAJ			Patient ID		12409160125
Birthday	20-01-1990			Sample ID		11966942
Age at Sample date	34.7			Sample Date		16-09-2024
Gestational age			11+1			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	65	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Da	ata	

Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	11+1	
PAPP-A	$3.4~\mathrm{mIU/ml}$	1.24	Method	CRL (<>Robinson)	
fb-hCG	55.8 ng/ml	1.1	Scan date	14-09-2024	
Risks at sampling date			Crown rump length in mm	42	
Age Risk		1:277	Nuchal translucency MoM	0.52	
Biochemical T21 risk		1:2156	Nasal bone	PRESENT	
Combined trisomy 21 risl	k	<1:10000	Sonographer	DR. RAVINDER KUMAR	
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NI	MBBS	
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:10		Cut off	The calculated risk for Trison cut off, which represents a low After the result of the Trisony 21 expected that among more than 10 there is one woman with a trisony The calculated risk by PRISCA de	risk. test (with NT) it is 1000 women with the same data, 21 pregnancy.	
1:1000 1:10000 13 15 17 19 21 23 25 2 Trisomy 13/18+NT	7 29 31 33 35 37 39	41 43 45 47 4	information provided by the referr the risk calculations are statistical a diagnostic value! The patient combined risk presum	ing physician. Please note that approaches and have no less that NT measurement was nes (Prenat Diagn 18:511-523;	