

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

Risk Above Cut Off

The calculated risk for Trisomy 13/18 (with NT) is

<1:10000, which indicates a low risk

Trisomy 13/18+NT



note that the risk calculations are statistical aapproaches and

The laboratory cannot be hold responsible for their impact

on the risk assessment! Calculated risks have no diagnostic

Risk below Age risk

have no diagnostic value!

Risk above Age Risk

values

Book a Test Online www.molq.in

				Date of Report PRISCA	3/5/2022 5.1.0.17
Patient Data					
Name	e MRS. PINKI NEGI				012205010115
Birthday		20/9/1989	Sample ID		11166440
Age at term	n 33				1/5/2022
Gestational age		13+6			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	69 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	13+4
PAPP-A	$3.82~\mathrm{mIU/ml}$	0.57	Method		CRL (<>Robinson)
fb-hCG	36.8 ng/ml	0.93	Scan date		30/04/2022
Risks at sampling date			Nuchal Translucency 1.5		
Age Risk		1:457	Nuchal Transl	ucency MoM	0.82
Biochemical T21 risk		1:871	Nasal Bone		Present
Combined T21 risk		<b>-</b> 1:4935			
Trisomy 13/18+NT		<1:10000			
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 it is expected that among 4935 women with the same data, there is one woman with a trisomy 21 pregnancy 4934 women with not affected pregnancies.  The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please		