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1.11

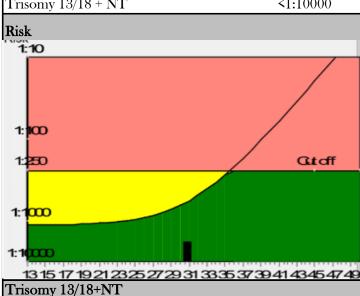
					Date of Report	05-12-2024
					PRISCA	5.2.0.13
Patient Data						
Name	MRS. KAVITA W/O RAKESH			Patient ID		12412020123
Birthday	02-02-1994			Sample ID		11947075
Age at Sample date	30.8			Sample Date		02-12-2024
Gestational age	12+6					
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	57	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ago	e	12+5
PAPP-A	<b>6.</b> 3	mIU/ml	0.96	Method		CRL (<>Robinson)
fb-hCG	49.5	ng/ml	1.43	Scan date		02-12-2024
Risks at sampling date				Crown rump l	ength in mm	62.7

## Age Risk 1:585 Nuchal translucency MoM

Biochemical T21 risk Nasal bone 1:1477 present

Combined trisomy 21 risk 1:4316 Sonographer DR. AMENDA

Trisomy 13/18 + NT <1:10000 Qualifications in measuring NT MD



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

which indicates a low risk

## The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.

Down's Syndrome Risk (Trisomy 21 Screening)

After the result of the Trisomy 21 test (with NT) it is expected that among 4316 women with the same data, there is one woman with a trisomy 21 pregnancy and 4315 women with not affected pregnancies.

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).

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values