

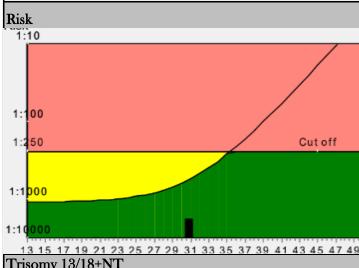
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					Date of Report PRISCA	25-08-2024 5.2.0.13
Patient Data						
Name	MRS. AYUSHI JAIN			Patient ID		12408240163
Birthday			25-11-1993	Sample ID		11485099
Age at Sample date		30.7				24-08-2024
Gestational age	11+6					
Correction factors						
Fetuses	1 I	VF		unknown	Previous trisomy 21	unknown
Weight in kg	65 I	Diabetes		NO	Pregnancies	unknown
Smoker	NO (	Origin		Asian		
Biochemical Data				Ultrasound D	ata	
Parameter	Value Co		Corr Mom	Gestational age		11+4
PAPP-A	3.8 mIU/ml 1.01		1.01	Method		CRL (<>Robinson)
fb-hCG	22.5 ng/ml 0.52		0.52	Scan date		22-08-2024
Risks at sampling date				Crown rump l	ength in mm	47
Age Risk	1:571		Nuchal translucency MoM		0.62	
Biochemical T91 risk			<1.10000	Nasal bone		PRESENT

## Nasal bone Biochemical T21 risk <1:10000 PRESENT Combined trisomy 21 risk <1:10000 Sonographer DR. SAURABH Trisomy 13/18 + NT <1:10000 Qualifications in measuring NT Down's Syndrome Risk (Trisomy 21 Screening)



## Trisomy 13/18+NT 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.

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 diagnostic value!

The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523;

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