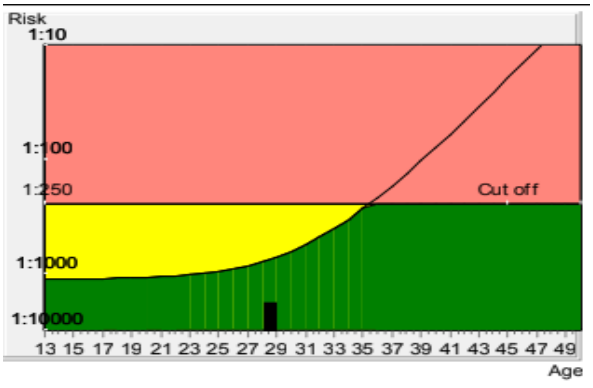
Date of Report	3/8/2022
PRISCA	5.1.0.17

Patient Data			
Name	MRS. MEENAKSHI VATS	Patient ID	052208020010
Birthday	2/1/1994	Sample ID	11561654
Age at term	28.1	Sample Date	2/8/2022
Gestational age	13+1		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	60	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	**Value**	**Corr Mom**	Gestational age	12+4
PAPP-A	6.48 mIU/ml	1.1	Method	CRL (<>Robinson)
fb-hCG	101.5 ng/ml	2.86	Scan date	1/8/2022
			Nuchal translucency	1.1
Risks at sampling date			Nuchal translucency MoM	0.7
Age Risk		1:760	Nasal bone	Present
Biochemical T21 risk		1:442		
Combined trisomy 21 risk		1:2475		
Trisomy 13/18 + NT		<1:10000		

Risk



Trisomy 13/18+NT

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

Down's Syndrome Risk (Trisomy 21 Screening)

The calculated risk for Trisomy 21(with NT) is below the cut off, which represents a low risk.
After the result of the Trisomy 21with NT test it is expected that among 2475 women with the same data, there is one woman with a trisomy 21 pregnancy and 2474 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 laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

Risk Above Cut Off Risk above Age Risk Risk below Age risk