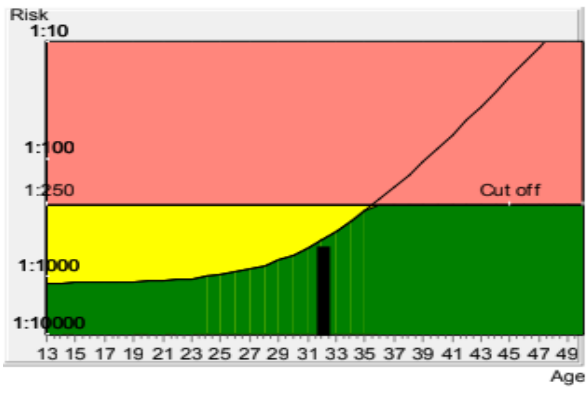
Date of Report	31/8/2022
PRISCA	5.1.0.17

Patient Data			
Name	**MRS. MANDVI**	Patient ID	012208280091
Birthday	20/07/1990	Sample ID	11594511
Age at term	32.07	Sample Date	28/8/2022
Gestational age	13+2		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	64	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	**Value**	**Corr Mom**	Gestational age	13+0
PAPP-A	6.48 mIU/ml	1.01	Method	CRL (<>Robinson)
fb-hCG	170.2 ng/ml	5.84	Scan date	27/8/2022
Risks at sampling date			Nuchal translucency	0.9
Age Risk		1:489	Nuchal translucency MoM	0.53
Biochemical T21 risk		1:94	Nasal bone	Present
Combined trisomy 21 risk		1:561		
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 561 women with the same data, there is one woman with a trisomy 21 pregnancy and 560 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