Date of Report	14-01-2024
PRISCA	5.2.0.13

Patient Data			
Name	MRS ANISHA	Patient ID	012401130030
Birthday	30-03-2000	Sample ID	11802752
Age at Sample date	23.8	Sample Date	13-01-2024
Gestational age	12+4		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	73	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+4
PAPP-A	4.82 mIU/ml	1.10	Method	CRL (<>Robinson)
fb-hCG	69.5 ng/ml	2	Scan date	13-01-2024

Risks at sampling date		Crown rump length in mm	60.4
Age Risk	1:1008	Nuchal translucency MoM	0.64
Biochemical T21 risk	1:1472	Nasal bone	PRESENT
Combined trisomy 21 risk	1:7873	Sonographer	DR.RAHUL
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT	MBBS

Risk

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

Down's Syndrome Risk (Trisomy 21 Screening)
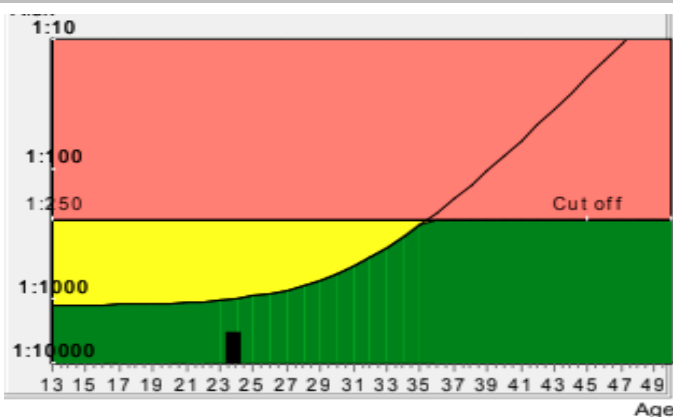
The calculated risk for Trisomy 21(with NT) is below the cut off, which represents a low risk.
After the result of the Trisomy 21 test (with NT) it is expected that among 7873 women with the same data, there is one woman with a trisomy 21 pregnancy and 7873 women with not affected pregnancies.
The calculated 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

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