Date of Report	26-09-2023
PRISCA	5.2.0.13
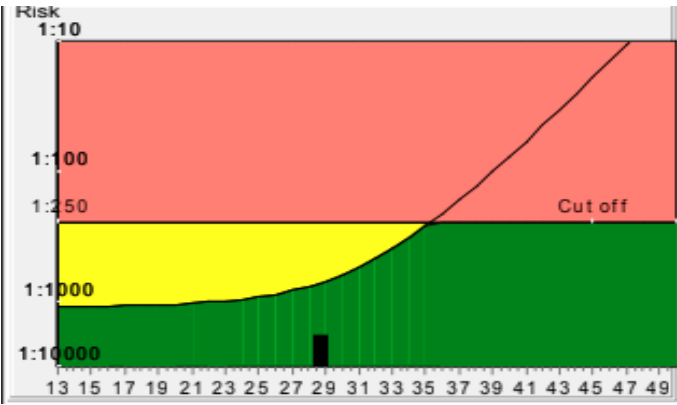
Patient Data

Name	**MRS ARCHNA**	Patient ID	012309250155
Birthday	01-01-1995	Sample ID	11806224
Age at Sample date	28.7	Sample Date	25-09-2023
Gestational age	11+5		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	58	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data

Parameter	Value	Corr Mom
PAPP-A	3.46 mIU/ml	0.85
fb-hCG	62.5 ng/ml	1.35

Risks at sampling date

Age Risk	1:721
Biochemical T21 risk	1:1576
Combined trisomy 21 risk	1:8451
Trisomy 13/18 +NT	<1:10000

Ultrasound Data

Gestational age	11+5
Method	CRL (<>Robinson)
Scan date	25-09-2023
Crown rump length in mm	49.7
Nuchal translucency MoM	0.82
Nasal bone	PRESENT
Sonographer	DR. NAVIN KUMAR
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(with NT) test it is expected that among 8451 women with the same data, there is one woman with a trisomy 21 pregnancy and 8450 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 the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).
The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no

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