Date of Report		05-11-2020
PRISCA		5.0.2.37

Patient Data

Name	**MRS HEMA**	Patient ID	012011020012
Birthday	29-09-1986	Sample ID	10738387
Age at delivery	34.1	Sample Date	02/11/2020
Gestational age	12+4		

Correction factors

Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	71	Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown	Origin	Asian		

Biochemical Data

Parameter	Value		Corr Mom
PAPP-A	1.75	mIU/ml	0.47
fb-hCG	21.9	ng/ml	0.5

Risks at sampling date

Age Risk	1:330
Biochemical T21 risk	1:1376
Combined Trisomy 21 Risk	1:7567
Trisomy 13/18 + NT	<1:10000

Ultrasound Data

Gestational age	12+2
Method	CRL(<>Robinson
Scan date	31-10-2020
Crown rump length in mm	57.3
Nuchal translucency MOM	0.8
Nasal bone	Present
Sonographer	DR.GIRISH PANDIT
Qualification in measuring NT	MD
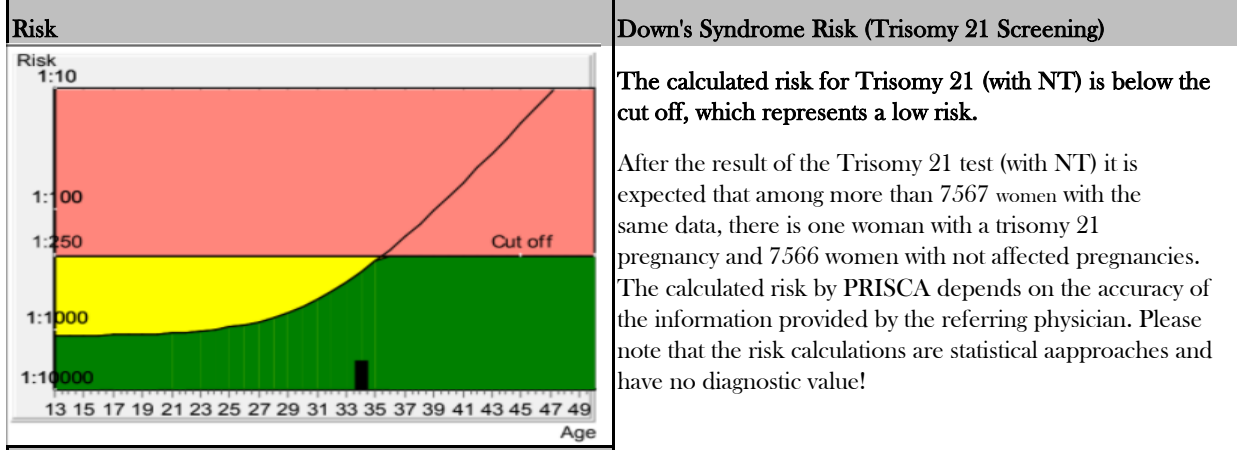
Risk

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 21 test (with NT) it is expected that among more than 7567 women with the same data, there is one woman with a trisomy 21 pregnancy and 7566 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