Prisca   5.0.2.37

Date of report: **06-09-2017**

Patient data

Name	MRS. URMILA	Patient ID	1709220109/AMB
Birthday	08-08-1972	Sample ID	1709220109/AMB
Age at sample date	45.1	Sample Date	05-09-2017
Gestational age	13 + 1		

Correction factors

Fetuses	1	IVF	yes	Previous trisomy 21	no
Weight	49	diabetes	no	pregancies	
Smoker	no	Origin	Asian		

Biochemical data / Ultrasound data

Parameter	Value	Corr. MoM	Ultrasound data	
PAPP-A	11.2 mIU/ml	2.04	Gestational age	11 + 4
fb-hCG	180.1 ng/ml	5.29	Method	CRL Robinson
			Scan date	25-08-2017

Risks at sampling date

Age risk	1:19	Crown rump length in mm	50.3
Biochemical T21 risk	>1:50	Nuchal translucency MoM	0.66
Combined trisomy 21 risk	1:64	Nasal bone	present
Trisomy 13/18 + NT	<1:10000	Sonographer	DR. H.S. MANN
		Qualifications in measuring NT	MD

Risk

Trisomy 21

The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 64 women with the same data, there is one woman with a trisomy 21 pregnancy and 63 women with not affected pregnancies.
The free beta HCG level is high.
The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.
Please note that risk calculations are statistical approaches 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 diagnostic value!

Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.

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Sign of Physician

🟩 below cut off	🟨 Below Cut Off, but above Age Risk	🟥 above cut off