Prisca   5.0.2.37

Date of report: **16-10-2017**

Patient data			
Name	MRS. DEEPIKA	Patient ID	1707220362/AMB
Birthday	11-09-1983	Sample ID	1707220362/AMB
Age at sample date	34.8	Sample Date	14-10-2017
Gestational age	13 + 3		

Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregancies
Weight	60	diabetes	no	no
Smoker	no	Origin	Asian	

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	13 + 3
PAPP-A	4.79 mIU/ml	1.75	Method	CRL Robinson
fb-hCG	122.5 ng/ml	3.05	Scan date	14-10-2017
Risks at sampling date			Crown rump length in mm	68.1
Age risk		1:276	Nuchal translucency MoM	1.02
Biochemical T21 risk		1:400	Nasal bone	present
Combined trisomy 21 risk		1:1565	Sonographer	DR. (MRS.) ADITI MAHAJAN
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD

Risk
1:10

Trisomy 21

The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
After the result of the Trisomy 21 test (with NT) it is expected that among 1565 women with the same data, there is one woman with a trisomy 21 pregnancy and 1564 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.

Sign of Physician

below cut off   Below Cut Off, but above Age Risk   above cut off