Prisca   5.0.2.37

Date of report: 22-03-2018

Patient data			
Name	MRS. PAYAL	Patient ID	1803220583/AMB
Birthday	18-07-1991	Sample ID	1803220583/AMB
Age at sample date	26.7	Sample Date	21-03-2018
Gestational age	11 + 5		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21	no
Weight	51	diabetes	no	pregancies	
Smoker	no	Origin	Asian		

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	11 + 4
PAPP-A	8.72 mIU/ml	2.92	Method	CRL Robinson
fb-hCG	48.1 ng/ml	1.12	Scan date	20-03-2018

Risks at sampling date			
		Crown rump length in mm	51.2
Age risk	1:852	Nuchal translucency MoM	1.45
Biochemical T21 risk	<1:10000	Nasal bone	present
Combined trisomy 21 risk	<1:10000	Sonographer	.
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT	MD

Risk

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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.
The PAPP-A 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