Prisca   5.0.2.37

Date of report: **16-06-2019**

Patient data			
Name	MRS. MUNISH PAL	Patient ID	1906220664/AMB
Birthday	19-09-1989	Sample ID	1906220664/AMB
Age at sample date	29.7	Sample Date	15-06-2019
Gestational age	13 + 0		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregancies	no
Weight	52.8	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 6
PAPP-A	5.36 mIU/ml	1.12	Method	CRL Robinson
fb-hCG	186 ng/ml	4.65	Scan date	14-06-2019
Risks at sampling date			Crown rump length in mm	67
Age risk		1:678	Nuchal translucency MoM	0.71
Biochemical T21 risk		1:159	Nasal bone	present
Combined trisomy 21 risk		1:928	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 928 women with the same data, there is one woman with a trisomy 21 pregnancy and 927 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