Prisca   5.1.0.17

Date of report: **30-07-2019**

Patient data			
Name	MRS. JASWANT	Patient ID	
Birthday	09-09-1982	Sample ID	1907221581/AMB
Age at delivery	37.4	Sample Date	25-07-2019
Gestational age	13 + 4		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	no
Weight	88.6	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	13 + 4
PAPP-A	3.1 mIU/ml	1.00	Method	CRL Robinson
fb-hCG	26.1 ng/ml	0.99	Scan date	25-07-2019
Risks at term			Crown rump length in mm	78
Age risk		1:249	Nuchal translucency MoM	0.63
Biochemical T21 risk		1:1568	Nasal bone	present
Combined trisomy 21 risk		1:7808	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 7808 women with the same data, there is one woman with a trisomy 21 pregnancy and 7807 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 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