Priska   5.0.2.37

Date of report: **24-03-2018**

Patient data		
Name	MRS. REENA	Patient ID
Birthday	12-05-1994	Sample ID 1803220638/AMB
Age at sample date	23.9	Sample Date 23-03-2018
Gestational age	11 + 3	

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

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	11 + 2
PAPP-A	1.1 mIU/ml	0.51	Method	CRL Robinson
fb-hCG	21.1 ng/ml	0.49	Scan date	22-03-2018

Risks at sampling date		Crown rump length in mm	47.8
Age risk	1:964	Nuchal translucency MoM	1.31
Biochemical T21 risk	1:5071	Nasal bone	present
Combined trisomy 21 risk	1:8759	Sonographer	.
Trisomy 13/18 + NT	1:8666	Qualifications in measuring NT	MD

Risk

13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49
Age

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 8759 women with the same data, there is one woman with a trisomy 21 pregnancy and 8758 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 1:8666, 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