Prisca 5.0.2.37

Date of report: **05-06-2018**

Patient data

Name	MRS. AMANPREET	Patient ID	1806220109/AMB
Birthday	14-01-1987	Sample ID	1806220109/AMB
Age at sample date	31.4	Sample Date	04-06-2018

Gestational age	11 + 4		
Correction factors			
Fetuses	1	IVF	yes	Previous trisomy 21	no
Weight	68	diabetes	no	pregancies	
Smoker	no	Origin	Asian		

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	11 + 4
PAPP-A	3.82 mIU/ml	1.91	Method	CRL Robinson
fb-hCG	105.2 ng/ml	2.62	Scan date	04-06-2018

Risks at sampling date			
Age risk	1:514	Crown rump length in mm	49.8
Biochemical T21 risk	1:1009	Nuchal translucency MoM	0.57
Combined trisomy 21 risk	1:4880	Nasal bone	present
Trisomy 13/18 + NT	<1:10000	Sonographer	DR. (MRS.) POONAM LOOMBA
		Qualifications in measuring NT	MD

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 4880 women with the same data, there is one woman with a trisomy 21 pregnancy and 4879 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!

1315 1719 212325 2729 313335 3739 414345 4749
Age

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

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