Prisca 5.1.0.17

Date of report: **24-08-2019**

Patient data

Name	MRS. RAJWANT KAUR	Patient ID	1908221426/AMB
Birthday	08-11-1990	Sample ID	1908221426/AMB
Age at delivery	29.3	Sample Date	23-08-2019
Gestational age	13 + 1		

Correction factors

Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	no
Weight	39.1	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

Parameter	Value	Corr. MoM
PAPP-A	3.5 mIU/ml	0.49
fb-hCG	70 ng/ml	1.88

Risks at term

Age risk	1:1039
Biochemical T21 risk	1:269
Combined trisomy 21 risk	1:1758
Trisomy 13/18 + NT	<1:10000

Ultrasound data

Gestational age	12 + 5
Method	CRL Robinson
Scan date	20-08-2019
Crown rump length in mm	65.77
Nuchal translucency MoM	0.65
Nasal bone	present
Sonographer	.
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 1758 women with the same data, there is one woman with a trisomy 21 pregnancy and 1757 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.

Sign of Physician

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