KOS DIAGNOSTIC
LAB

Prisca 5.1.0.17

Date of report: 16/12/18

Patient data			
Name	Mrs. MANPREET	Patient ID	00261812150029
Birthday	26/08/95	Sample ID	H1857465
Age at delivery	23.8	Sample Date	14/12/18
Gestational age	11 + 3		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	45	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data

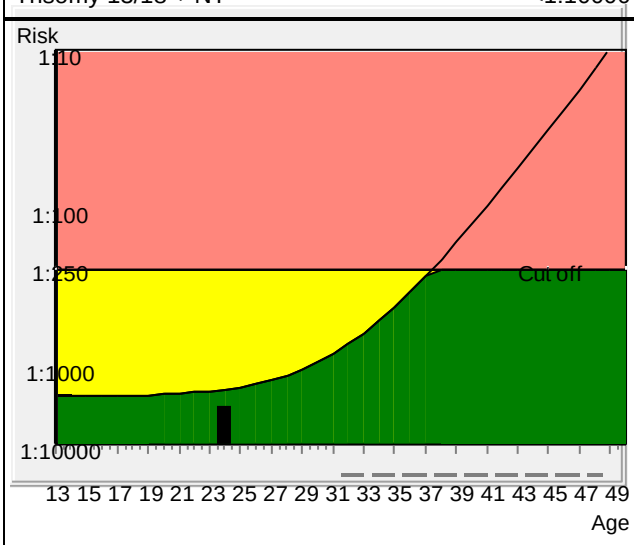
Parameter	Value	Corr. MoM
PAPP-A	1.98 mIU/ml	0.58
fb-hCG	53.2 ng/ml	1.05

Ultrasound data

Gestational age	11 + 2
Method	CRL Robinson
Scan date	13/12/18
Crown rump length in mm	46.6
Nuchal translucency MoM	1.25
Nasal bone	present
Sonographer	DR.BHUPINDER SINGH
Qualifications in measuring NT	MD

Risks at term

Age risk	1:1433
Biochemical T21 risk	1:2203
Combined trisomy 21 risk	1:4331
Trisomy 13/18 + NT	<1:10000



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 4331 women with the same data, there is one woman with a trisomy 21 pregnancy and 4330 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