Date of report: **09-03-2019**

Prisca   5.0.2.37

KOS DIAGNOSTIC LAB

Patient data		
Name		MRS. CHANPREET
Birthday		22-10-1995
Age at delivery		23.9
Previous trisomy 21 pregancies		no

Correction factors			
Fetuses	1	diabetes	no
Weight	75	Origin	Asian
Smoker	no	IVF	no

Biochemical data		
Sample Date		08-03-2019
Gestational age at sample date		14 + 1

Parameter	Value	Corr. MoMs
AFP	26.5 ng/ml	1.05
HCG	25573 mIU/ml	0.69
uE3	0.2 ng/ml	1.08

Ultrasound data	
Gestational age	13 + 4
Method	CRL measurements
Crown rump length in mm	78.7
Date	04-03-2019
Nuchal translucency MoM	0.84
Nuchal translucency	1.60 mm
Nasal bone	present
Sonographer	.
Qualifications in measuring NT	MD

Risks at term	
Age risk	1:1432
Trisomy 21 risk	<1:10000
Combined trisomy 21 risk	<1:10000
Trisomy 18 risk	<1:10000

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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.

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 18

The calculated risk for trisomy 18 (with nuchal translucency) is < 1:10000, which represents a low risk.

Neural tube defects

The corrected MoM AFP (1.05) is located in the low risk area for neural tube defects.

🟩 below cut off   🟨 Below Cut Off, but above Age Risk   🟥 above cut off