Priska   5.1.0.17

Date of report: **01-10-2019**

KOS DIAGNOSTIC LAB

Patient data			
Name	MRS. NEHA	Patient ID	1909221836/AMB
Birthday	18-09-1998	Sample ID	1909221836/AMB
Age at delivery	21.6	Sample Date	30-09-2019
Gestational age	11 + 2		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21	no
Weight	66	diabetes	no	pregnancies	
Smoker	no	Origin	Asian		

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	11 + 0
PAPP-A	1.8 mIU/ml	1.00	Method	CRL Robinson
fb-hCG	168.5 ng/ml	3.96	Scan date	28-09-2019
Risks at term			Crown rump length in mm	43.2
Age risk		1:1505	Nuchal translucency MoM	0.83
Biochemical T21 risk		1:286	Nasal bone	present
Combined trisomy 21 risk		1:1651	Sonographer	.
Trisomy 13/18 + NT		<1:10000	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 1651 women with the same data, there is one woman with a trisomy 21 pregnancy and 1650 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!

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

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