Prisca   5.1.0.17

Date of report: **09-08-2019**

KOS DIAGNOSTIC LAB

Patient data			
Name	MRS. BALJEET	Patient ID	1908220500/AMB
Birthday	19-01-1997	Sample ID	1908220500/AMB
Age at delivery	23.1	Sample Date	08-08-2019
Gestational age	10 + 5		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	no
Weight	60	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	10 + 4
PAPP-A	1.1 mIU/ml	0.73	Method	CRL Robinson
fb-hCG	134 ng/ml	2.77	Scan date	07-08-2019
Risks at term			Crown rump length in mm	39
Age risk		1:1461	Nuchal translucency MoM	0.99
Biochemical T21 risk		1:368	Nasal bone	present
Combined trisomy 21 risk		1:1630	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 1630 women with the same data, there is one woman with a trisomy 21 pregnancy and 1629 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