Prisca   5.1.0.17

Date of report: **13-10-2022**

KOS DIAGNOSTIC LAB

Patient data			
Name	MRS. SUNITA	Patient ID	2210220214/AMB
Birthday	14-05-1985	Sample ID	2210220214/AMB
Age at sample date	37.4	Sample Date	12-10-2022
Gestational age	11 + 3		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	no
Weight	70	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	10 + 5
PAPP-A	0.31 mIU/ml	0.16	Method	CRL Robinson
fb-hCG	54.5 ng/ml	1.12	Scan date	07-10-2022
Risks at sampling date			Crown rump length in mm	41
Age risk		1:148	Nuchal translucency MoM	0.95
Biochemical T21 risk		>1:50	Nasal bone	present
Combined trisomy 21 risk		1:51	Sonographer	.
Trisomy 13/18 + NT		1:72	Qualifications in measuring NT	MD

Risk

Trisomy 21

The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.
After the result of the Trisomy 21 test (with NT) it is expected that among 51 women with the same data, there is one woman with a trisomy 21 pregnancy and 50 women with not affected pregnancies.
The PAPP-A level is low.
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 test (with nuchal translucency) is 1:72, which represents an increased risk.

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Sign of Physician

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