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11394775

Date of Report 13/3/2022 PRISCA 5.1.0.17 ANKITA KHANDELWAL(F2) Patient ID 012203120021

6/7/1994 Sample ID Birthday Age at term 28.01 Sample Date 12/3/2022

Gestational age 13+6

Correction factors

Patient Data

Name

2 IVF Fetuses unknown Previous trisomy 21 unknown NO Pregnancies Weight in kg 73 Diabetes unknown NO Origin Smoker Asian

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+3
PAPP-A	$5.9~\mathrm{mIU/ml}$	0.57	Method	CRL (<>Robinson)
fb-hCG	32.4 ng/ml	0.4	Scan date	9/3/2022
Risks at sampling date			Nuchal translucency	1.2
Age Risk		1:853	Nuchal translucency MoM	0.66
Biochemical T21 risk		1:9318	Nasal Bone	Presesnt
Combined trisomy 21 risk		<1:10000		
Trisomy 13/18+NT		<1:10000		

Risk Down's Syndrome Risk (Trisomy 21 Screening)

Risk 1:10 1:100 Cut off 1:250 1:1000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49

The calculated risk for Trisomy 21 with NT is below the cut off, which represents a low risk.

After the result of the Trisomy 21 test it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy 9999 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 the risk calculations are statistical aapproaches and have no diagnostic value!

Trisomy 13/18+NT

The calculated risk for Trisomy 13/18 with NT is <1:10000, which indicates a low risk

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

Risk Above Cut Off

Risk above Age Risk

Risk below Age risk