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Date of Report 6/5/2024
PRISCA 5.2.0.13

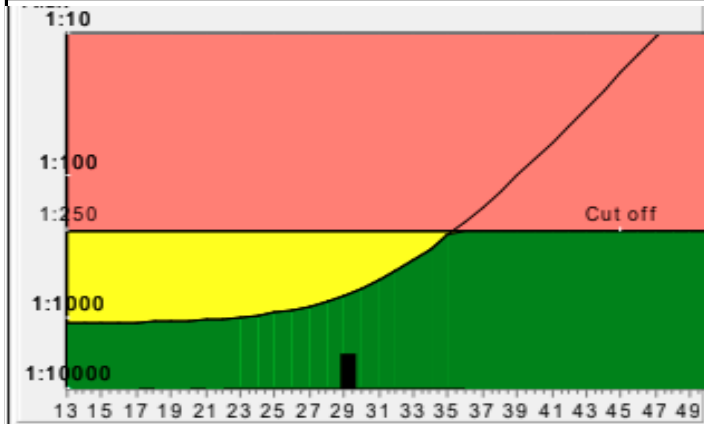
Patient Data	
Name	SUMAN
Birthdate	2/2/1995
Age at sample	29.3
Gestational age	11+6
Patient ID	D12405040142
Sample ID	11819430
Sample Date	4/5/2024

Correction factors	
Fetuses	1 IVF
Weight in kg	57 Diabetes
Smoker	NO Origin
unknown	Previous trisomy 21
NO	Pregnancies
Asian	unknown

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	11+5
PAPP-A	3.80 mIU/ml	0.86	Method	CRL (<>Robinson)
fb-hCG	31.2 ng/ml	0.69	Scan date	3/5/2024

Risks at sampling date		Down's Syndrome Risk (Trisomy 21 Screening)	
Age Risk	1:687	Crown rump length in mm	45
Biochemical T21 risk	1:6813	Nuchal translucency MoM	1.61
Combined trisomy 21 risk	1:3331	Nasal bone	Present
Trisomy 13/18 + NT	<1:10000	Sonographer	DR.
		Qualifications in measuring NT	MBBS

Risk



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 (with NT) it is expected that among 3331 women with the same data, there is one woman with a trisomy 21 pregnancy and 3330 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 approaches and have no diagnostic value! The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998). The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

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

Risk Above Cut Off
 Risk above Age Risk
 Risk below Age risk