

Date of Report 28-01-2024
 PRISCA 5.2.0.13

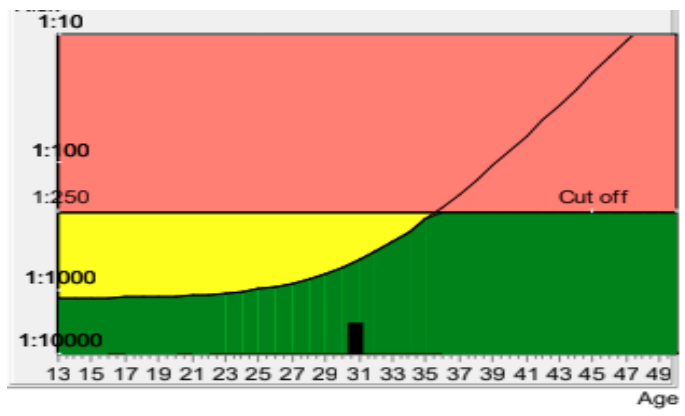
Patient Data	
Name	MRS SUMAN
Birth day	23-03-1993
Age at Sample date	30.8
Gestational age	13+4

Correction factors	
Fetuses	1 IVF
Weight in kg	65 Diabetes
Smoker	NO Origin

unknown	Previous trisomy 21	unknown
NO	Pregnancies	unknown
Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+3
PAPP-A	6.5 mIU/ml	0.91	Method	CRL (<>Robinson)
fb-hCG	31.2 ng/ml	1.17	Scan date	26-01-2024

Risks at sampling date			Ultrasound Data	
Age Risk	1:599		Crown rump length in mm	74
Biochemical T21 risk	1:2106		Nuchal translucency MoM	0.67
Combined trisomy 21 risk	<1:10000		Nasal bone	PRESENT
Trisomy 13/18 + NT	<1:10000		Sonographer	DR.PRAVEEN
			Qualifications in measuring NT	MD

Risk	Down's Syndrome Risk (Trisomy 21 Screening)
	<p>The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 women with not affected pregnancies.</p> <p>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!</p> <p>The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998).</p>

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