

Date of Report 17-04-2021
PRISCA 5.0.2.37

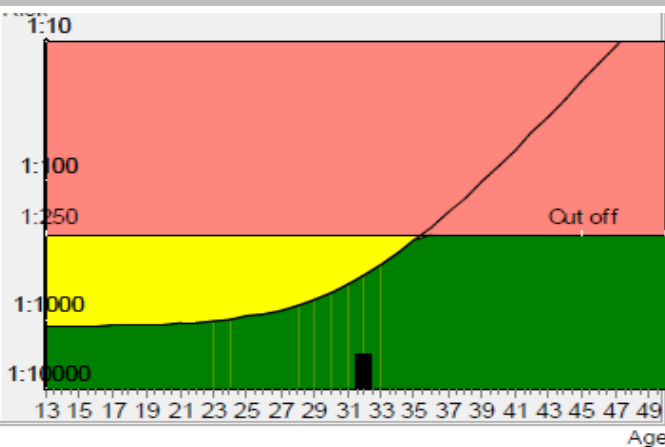
Patient Data			
Name	MRS. KRISHNA	Patient ID	012104140199
Birth day	24-05-1989	Sample ID	10893590
Age at delivery	31.9	Sample Date	14/04/2021
Gestational age	12+2		

Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	52	Diabetes	unknown
Smoker	Unknown	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+2
PAPP-A	3.24 mIU/ml	0.6	Method	CRL(<>Robinson
fb-hCG	44.6 ng/ml	0.92	Scan date	14-04-2021
Risks at sampling date			Crown rump length in mm	84

Age Risk	1:493	Nuchal translucency MOM	0.42
Biochemical T21 risk	1:1128	Nasal bone	Present
Combined Trisomy 21 Risk	1:6331	Sonographer	Dr Shruti Sangwan
Trisomy 13/18 + NT	<1:10000	Qualification in measuring NT	MD

Risk **Down's Syndrome Risk (Trisomy 21 Screening)**



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 more than 6331 women with the same data, there is one woman with a trisomy 21 pregnancy and 6330 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