

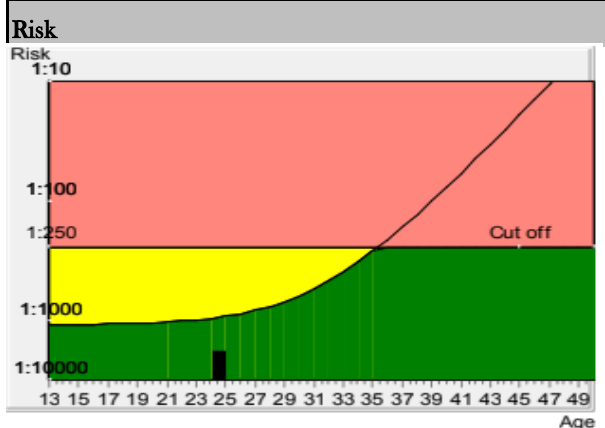
Date of Report 24/3/2022
PRISCA 5.1.0.17

Patient Data			
Name	MRS. RAJ KUMARI	Patient ID	012203200211
Birthday	19/09/1997	Sample ID	11246950
Age at term	24.11	Sample Date	20/3/2022
Gestational age	11+5		

Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	58	Diabetes	NO
Smoker	NO	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	11+3
PAPP-A	2.5 mIU/ml	0.74	Method	CRL (<>Robinson)
fb-hCG	34.5 ng/ml	0.67	Scan date	20/3/2022

Risks at sampling date			
Age Risk	1:953	Nuchal translucency	1.4
Biochemical T21 risk	1:7088	Nuchal translucency MoM	1.09
Combined trisomy 21 risk	<1:10000	Nasal Bone	Present
Trisomy 13/18+NT	<1:10000		



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 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 approaches 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