

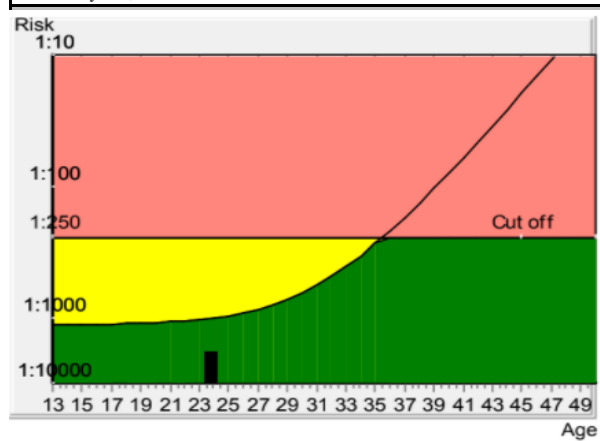
Date of Report 22-02-2021
PRISCA 5.0.2.37

Patient Data			
Name	MRS ANKITA	Patient ID	022102180013
Birthday	24-05-1997	Sample ID	10848783
Age at delivery	23.7	Sample Date	18/02/2021
Gestational age	12+5		

Correction factors			
Fetuses	1 IVF	unknown	Previous trisomy 21 unknown
Weight in kg	80.2	Diabetes	unknown
Smoker	Unknown	Origin	Asian
			Pregnancies unknown

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+4
PAPP-A	2.85 mIU/ml	0.84	Method	CRL(<>Robinson
fb-hCG	97.1 ng/ml	2.32	Scan date	18-02-2021

Risks at sampling date				
Age Risk	1:1015		Crown rump length in mm	62.3
Biochemical T21 risk	1:570		Nuchal translucency MOM	0.44
Combined Trisomy 21 Risk	1:3330		Nasal bone	Present
Trisomy 13/18 + NT	<1:10000		Sonographer	DR.MEENU SOLANKI
			Qualification in measuring NT	HMC



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 3330 women with the same data, there is one woman with a trisomy 21 pregnancy and 3329 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 held 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