

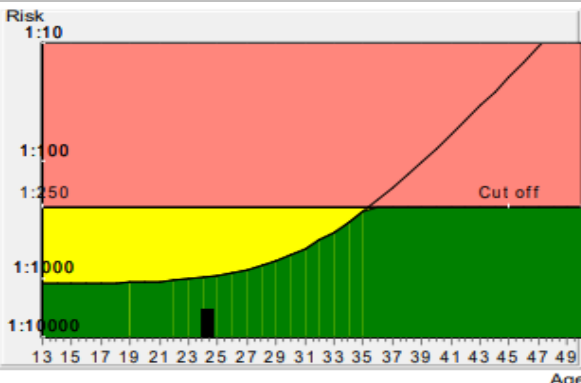
Date of Report 25/5/2023  
PRISCA 5.1.0.17

Patient Data	
Name	MRS. KOMAL W/O SANJAY
Birthdate	23/1/1999
Age at term	24.9
Gestational age	12+1

Correction factors	
Fetuses	1 IVF
Weight in kg	50.2
Smoker	NO
Diabetes	NO
Origin	Asian
Previous trisomy 21	unknown
Pregnancies	unknown

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+0
PAPP-A	4.26 mIU/ml	0.74	Method	CRL (<>Robinson)
fb-hCG	37.4 ng/ml	0.85	Scan date	23/05/2023

Risks at sampling date		Nuchal translucency (NT)	
Age Risk	1:974	Nuchal translucency MoM	1.1
Biochemical T21 risk	1:4377	Nasal bone	present
Combined trisomy 21 risk	<1:10000		
Trisomy 13/18	<1:10000		

Risk	Down's Syndrome Risk (Trisomy 21 Screening)
	<p><b>The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.</b></p> <p>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 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>

**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