



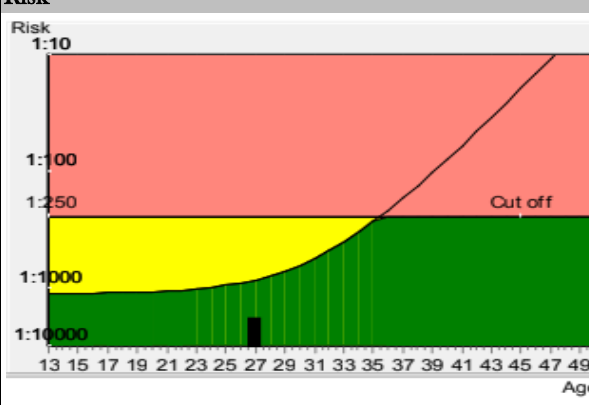
Date of Report 20/3/2023  
PRISCA 5.2.0.13

Patient Data			
Name	MRS. MANDEEP KAUR	Patient ID	102303190017
Birthday	24/4/1996	Sample ID	11556103
Age at term	27.3	Sample Date	19/3/2023
Gestational age	12+4		

Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	60	Diabetes	NO
Smoker	NO	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom		12+2
PAPP-A	3.89 mIU/ml	0.7	Method	CRL (<>Robinson)
fb-hCG	34.4 ng/ml	0.93	Scan date	17/3/2023

Risks at sampling date			
Age Risk	1:867	Nuchal translucency	1.3
Biochemical T21 risk	1:2827	Nuchal translucency MoM	0.87
Combined trisomy 21 risk	<1:10000	Nasal bone	Present
Trisomy 13/18 + NT	<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 with NT 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>

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