



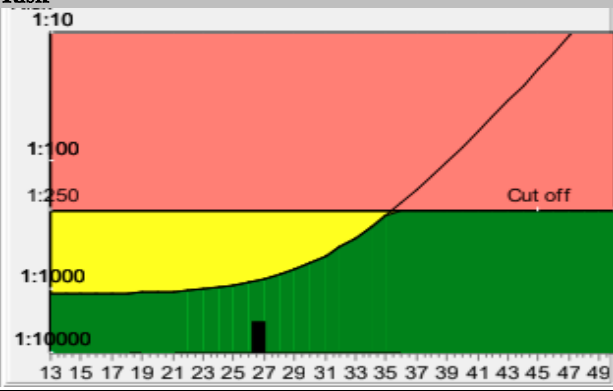
Date of Report 5/2/2024
PRISCA 5.2.0.13

Patient Data			
Name	KIRAN YADAV	Patient ID	012402030103
Birthday	8/7/1997	Sample ID	11530868
Age at sample	26.6	Sample Date	3/2/2024
Gestational age	12+1		

Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	57	Diabetes	NO
Smoker	NO	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+1
PAPP-A	3.90 mIU/ml	0.79	Method	CRL (<>Robinson)
fb-hCG	22.8 ng/ml	0.54	Scan date	3/2/2024

Risks at sampling date			
Age Risk	1:872	Crown rump length in mm	57.7
Biochemical T21 risk	<1:10000	Nuchal translucency MoM	0.73
Combined trisomy 21 risk	<1:10000	Nasal bone	Present
Trisomy 13/18 + NT	<1:10000	Sonographer	DR.DEEPIKA
		Qualifications in measuring NT	MD

Risk	Down's Syndrome Risk (Trisomy 21 Screening)
	<p>The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</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> <p>The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998).</p> <p>The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values</p>
<p>Trisomy 13/18 + NT</p> <p>The calculated risk for Trisomy 13/18 (with NT) is < 1:10000, which indicates a low risk</p>	

Risk Above Cut Off
 Risk above Age Risk
 Risk below Age risk