

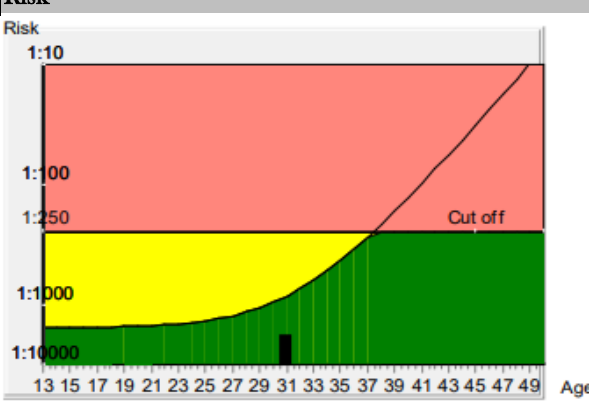
Date of Report 4/1/2022
PRISCA 5.2.0.13

Patient Data			
Name	MRS. VANDNA SHARMA	Patient ID	012201010028
Birthday	18/08/1991	Sample ID	11271085
Age at delivery	30.9	Sample Date	1/1/2022
Gestational age	12+4		


Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	77.9	Diabetes	NO
Smoker	NO	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown


Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+3
PAPP-A	2.42 mIU/ml	0.75	Method	CRL(<>Robinson
fb-hCG	50.1 ng/ml	1.27	Scan date	31/12/2021

Risks at sampling date				
Age Risk		1:864	Nuchal translucency	1.7
Biochemical T21 risk		1:1624	Nuchal translucency MoM	1.06
Combined T21 risk		1:5747	Nasal bone	present
Trisomy 13/18+NT		<1:10000		

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 it is expected that among 5747 women with the same data, there is one woman with a trisomy 21 pregnancy and 5746 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!</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>	<p>The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values</p>

 Risk Above Cut Off

 Risk above Age Risk

 Risk below Age risk