

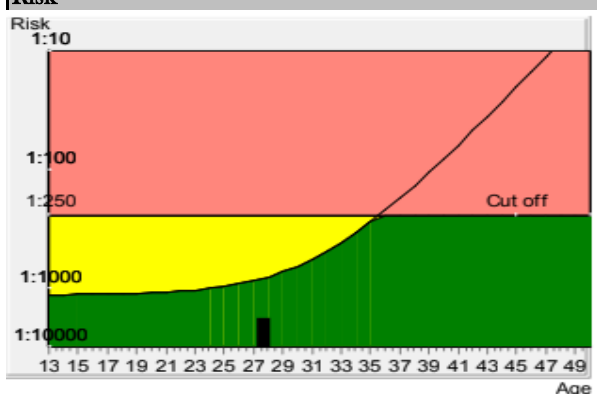
Date of Report 13/3/2022
PRISCA 5.1.0.17

Patient Data			
Name	ANKITA KHANDELWAL(F1)	Patient ID	012203120021
Birthday	6/7/1994	Sample ID	11394775
Age at term	28.01	Sample Date	12/3/2022
Gestational age	13+2		

Correction factors			
Fetuses	2	IVF	unknown
Weight in kg	73	Diabetes	NO
Smoker	NO	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+6
PAPP-A	5.9 mIU/ml	0.68	Method	CRL (<>Robinson)
fb-hCG	32.4 ng/ml	0.37	Scan date	9/3/2022

Risks at sampling date			
Age Risk	1:838	Nuchal translucency	1.2
Biochemical T21 risk	<1:10000	Nuchal translucency MoM	0.72
Combined trisomy 21 risk	<1:10000	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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy 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