

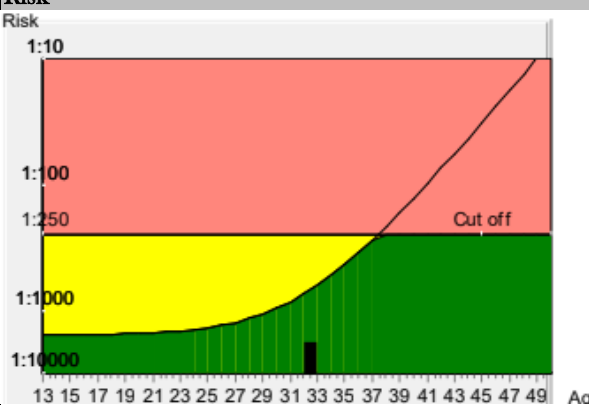
Date of Report 19/10/2021
PRISCA 5.2.0.13

Patient Data			
Name	MRS. POOJA	Patient ID	012110170010
Birthday	27/10/1989	Sample ID	11056802
Age at delivery	32.51	Sample Date	17/10/2021
Gestational age	12+0		

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	Gestational age	11+3
PAPP-A	2.86 mIU/ml	0.79	Method	CRL(<>Robinson
fb-hCG	59.1 ng/ml	1.27	Scan date	14/10/2021

Risks at sampling date				
Age Risk		1:681	Nuchal translucency	1.2
Biochemical T21 risk		1:1441	Nuchal translucency MoM	0.91
Combined T21 risk		1:7108	Nasal bone	Absent
Trisomy 13/18 + NT		<1:10000	Qualification in measuring NT	CR

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 7108 women with the same data, there is one woman with a trisomy 21 pregnancy and 7107 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>

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