| Date of Report | 20/3/2022 |
| PRISCA | 5.1.0.17 |
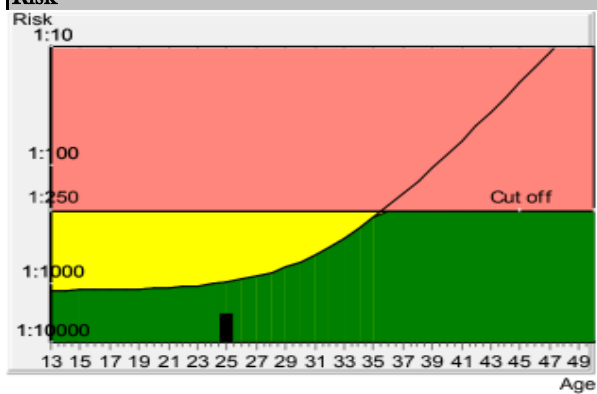
Patient Data	
Name	**MRS. PRIYA KUMARI**
Birthday	20/04/1997
Age at term	25.03
Gestational age	13+2

Patient ID	012203180001
Sample ID	11242742
Sample Date	18/3/2022

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	48	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data		
Parameter	Value	Corr Mom
PAPP-A	5.4 mIU/ml	0.71
fb-hCG	34.2 ng/ml	0.74

Ultrasound Data	
Gestational age	12+5
Method	CRL (<>Robinson)
Scan date	15/3/2022
Nuchal translucency	1.2
Nuchal translucency MoM	0.73
Nasal Bone	Presesnt

Risks at sampling date	
Age Risk	1:990
Biochemical T21 risk	1:5461
Combined trisomy 21 risk	<1:10000
Trisomy 13/18+NT	<1:10000

Risk

Down's Syndrome Risk (Trisomy 21 Screening)

The calculated risk for Trisomy 21 with NT is below the cut off, which represents a low risk.

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. 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 aapproaches and have no diagnostic value!

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

Trisomy 13/18+NT

The calculated risk for Trisomy 13/18 with NT is <1:10000 , which indicates a low risk

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