Date of Report	15/9/2021
PRISCA	5.2.0.13

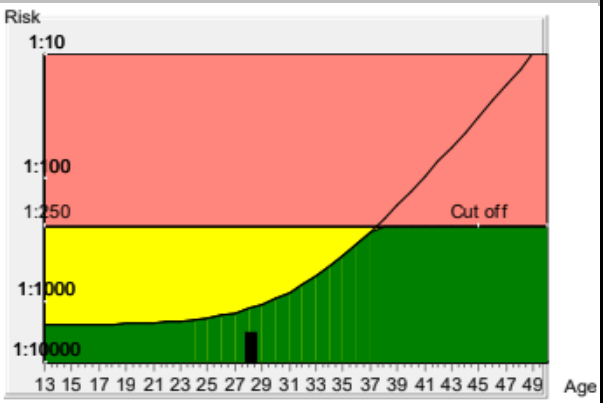
Patient Data			
Name	MRS.NAJBUL	Patient ID	012109140085
Birthday	1/1/1994	Sample ID	11329981
Age at delivery	28.24	Sample Date	14/9/2021
Gestational age	12+0		

Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	46	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		

Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	11+6
PAPP-A	1.62 mIU/ml	0.33	Method	CRL(<>Robinson
fb-hCG	21.8 ng/ml	0.44	Scan date	13/9/2021
			Nuchal translucency in mm	1.2

Risks at sampling date			
Age Risk	1:1143	Nuchal translucency MOM	0.82
Biochemical T21 risk	1:2274	Nasal bone	Present
Combined Trisomy 21 Risk	<1:10000		
Trisomy 13/18 + NT	<1:10000	Qualification in measuring NT	CR

Risk



Trisomy 13/18 + NT

The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low 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 (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 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

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