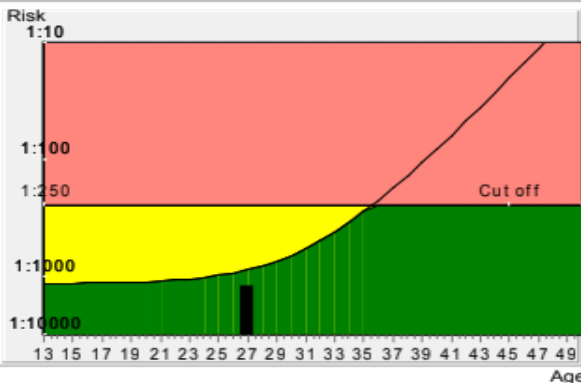



Date of Report 7/5/2023
PRISCA 5.2.0.13

Patient Data				
Name	MRS. SIMPI BEHRA	Patient ID	012305060205	
Birthday	6/6/1996	Sample ID	11651362	
Age at term	27.3	Sample Date	6/5/2023	
Gestational age	13+3			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21
Weight in kg	72	Diabetes	NO	Pregnancies
Smoker	NO	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	13+2	
PAPP-A	4.75 mIU/ml	0.79	Method	CRL (<>Robinson)
fb-hCG	85.2 ng/ml	3.14	Scan date	5/5/2023
Risks at sampling date			Nuchal translucency	
Age Risk	1:891		1.1	
Biochemical T21 risk	1:192		Nuchal translucency MoM	
Combined trisomy 21 risk	1:1191		0.61	
Trisomy 13/18 + NT	<1:10000		Nasal bone	
			Present	
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 with NT test it is expected that among 1191 women with the same data, there is one woman with a trisomy 21 pregnancy and 1190 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>	
<p>The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk</p>			<p>The laboratory cannot be held 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