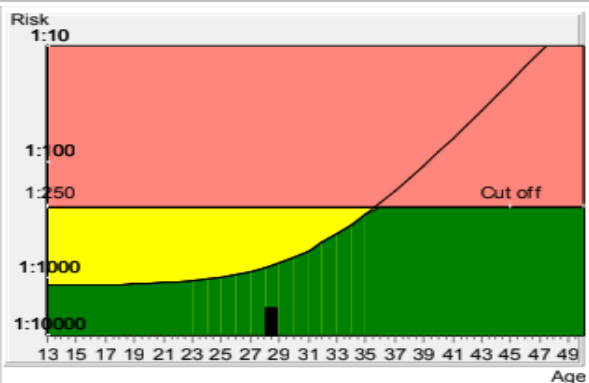


Date of Report 15/9/2022
PRISCA 5.1.0.17

Patient Data				
Name	MRS. BALJEET KAUR	Patient ID	012209140237	
Birthday	6/4/1994	Sample ID	11565712	
Age at term	28.1	Sample Date	14/9/2022	
Gestational age	13+6			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21
Weight in kg	55	Diabetes	NO	Pregnancies
Smoker	NO	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+0
PAPP-A	4.65 mIU/ml	0.49	Method	CRL (<>Robinson)
fb-hCG	20.6 ng/ml	0.81	Scan date	8/9/2022
Risks at sampling date			Nuchal translucency	1.2
Age Risk	1:799		Nuchal translucency MoM	0.69
Biochemical T21 risk	1:1377		Nasal bone	Present
Combined trisomy 21 risk	1:8028			
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 with NT test it is expected that among more than 8028 women with the same data, there is one woman with a trisomy 21 pregnancy and 8027 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 laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values	
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