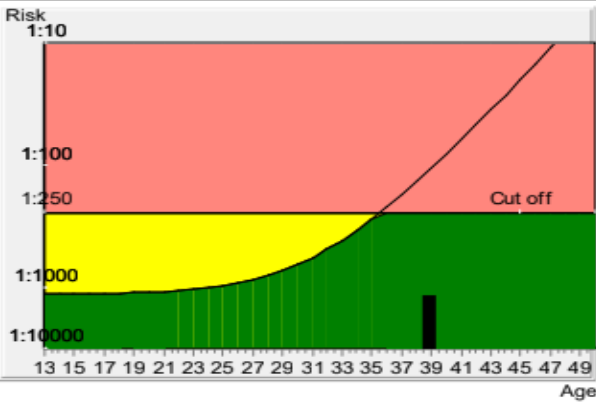





Date of Report 20/8/2023  
PRISCA 5.1.0.17

Patient Data		
Name	MRS. POONAM BANSAL	Patient ID 012308190126
Birthday	25/09/1984	Sample ID 11672440
Age at term	39.3	Sample Date 19/8/2023
Gestational age	13+0	
Correction factors		
Fetuses	1 IVF	unknown Previous trisomy 21 unknown
Weight in kg	82 Diabetes	NO Pregnancies unknown
Smoker	NO Origin	Asian
Biochemical Data		Ultrasound Data
Parameter	Value	Corr Mom
PAPP-A	6.6 mIU/ml	1.49
fb-hCG	59.7 ng/ml	2
Risks at sampling date		Gestational age 13+0
Age Risk	1:108	Method CRL (<>Robinson)
Biochemical T21 risk	1:280	Scan date 19/8/2023
Combined trisomy 21 risk	1:1176	Nuchal translucency (NT) 1.6
Trisomy 13/18	<1:10000	Nuchal translucency MoM 0.95
		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 test it is expected that among 1176 women with the same data, there is one woman with a trisomy 21 pregnancy and 1175 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