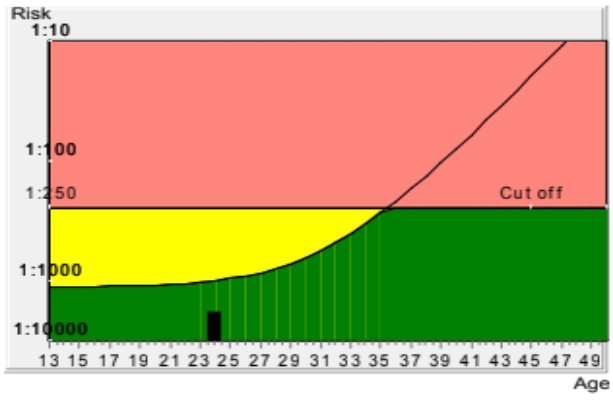


Date of Report 3/7/2023
PRISCA 5.1.0.17

Patient Data				
Name	MRS. JYOTI KANTI BEHRA	Patient ID	012307020101	
Birthday	22/7/1999	Sample ID	011640652	
Age at term	24.3	Sample Date	2/7/2023	
Gestational age	12+4			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21
Weight in kg	80	Diabetes	NO	Pregnancies
Smoker	NO	Origin	Asian	unknown
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+4
PAPP-A	3.24 mIU/ml	0.83	Method	CRL (<>Robinson)
fb-hCG	23.5 ng/ml	0.69	Scan date	2/7/2023
Risks at sampling date			Nuchal translucency (NT)	1.2
Age Risk		1:1003	Nuchal translucency MoM	0.75
Biochemical T21 risk		1:9158	Nasal bone	present
Combined trisomy 21 risk		<1:10000		
Trisomy 13/18		<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 test 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.</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 calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk			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