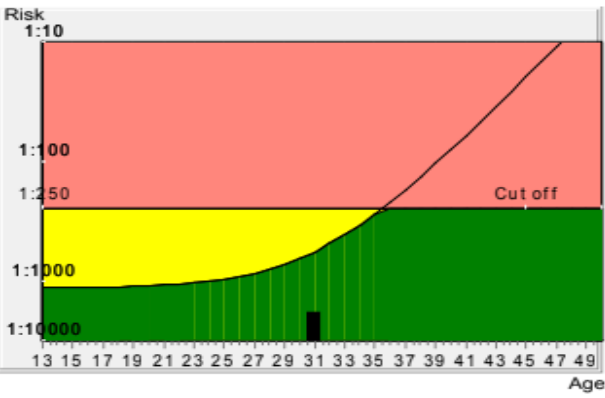


Date of Report 5/7/2023
PRISCA 5.1.0.17

Patient Data		
Name	MRS. KUMARI ANJU	Patient ID 012307040186
Birthday	14/07/1992	Sample ID 11674269
Age at term	31.6	Sample Date 4/7/2023
Gestational age	12+6	
Correction factors		
Fetuses	1 IVF	unknown Previous trisomy 21 unknown
Weight in kg	53 Diabetes	NO Pregnancies unknown
Smoker	NO Origin	Asian
Biochemical Data		Ultrasound Data
Parameter	Value	Corr Mom
PAPP-A	5.38 mIU/ml	0.76
fb-hCG	70.5 ng/ml	1.98
Risks at sampling date		Gestational age 12+6
Age Risk	1:574	Method CRL (<>Robinson)
Biochemical T21 risk	1:376	Scan date 4/7/2023
Combined trisomy 21 risk	1:2214	Nuchal translucency (NT) 1.25
Trisomy 13/18	<1:10000	Nuchal translucency MoM 0.76
		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 2214 women with the same data, there is one woman with a trisomy 21 pregnancy and 2213 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