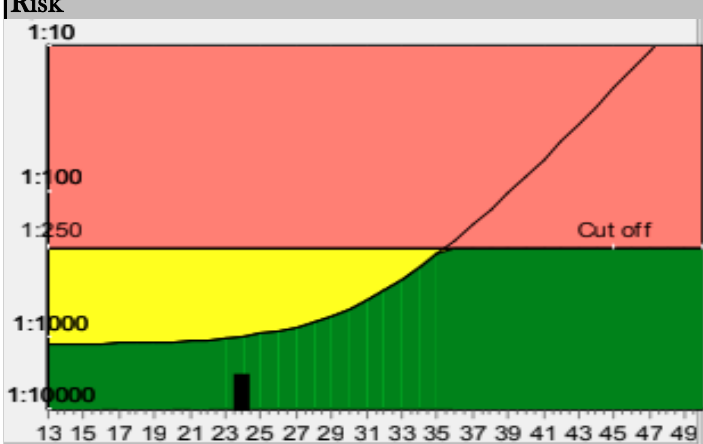


Date of Report 21/8/2023  
PRISCA 5.2.0.13

Patient Data				
Name	POOJA	Patient ID	012309200147	
Birth day	14/11/1999	Sample ID	11781154	
Age at sample	23.80	Sample Date	20/8/2023	
Gestational age	12+4			
Correction factors				
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown
Weight in kg	45 Diabetes	NO	Pregnancies	unknown
Smoker	NO Origin	Asian		
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+3
PAPP-A	5.51 mIU/ml	0.71	Method	CRL (<>Robinson)
fb-hCG	64.6 ng/ml	1.58	Scan date	19/09/2023
Risks at sampling date			Crown rump length in mm	59
Age Risk	1:1006		Nuchal translucency MoM	1.10
Biochemical T21 risk	1:1012		Nasal bone	Present
Combined trisomy 21 risk	1:3179		Sonographer	Dr Harendra bhaskar
Trisomy 13/18 + NT	<1:10000		Qualifications in measuring NT	MBBS
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 (with NT) it is expected that among 3179 women with the same data, there is one woman with a trisomy 21 pregnancy and 3178 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 patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998).</p> <p>The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values</p>	
Trisomy 13/18 + NT				
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