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Date of Report 6/5/2024  
PRISCA 5.2.0.13

Patient Data			
Name	DEEPTI W/O SANDEEP	Patient ID	012405040176
Birthday	30/4/1993	Sample ID	12004509
Age at sample	31	Sample Date	4/5/2024
Gestational age	12+1		

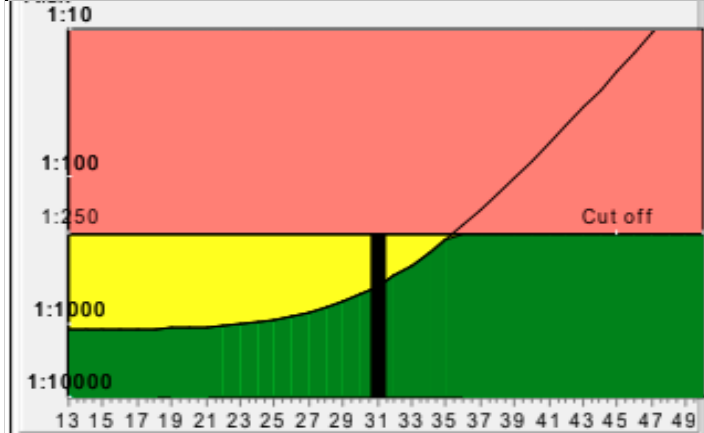
Correction factors			
Fetuses	1	IVF	unknown
Weight in kg	71	Diabetes	NO
Smoker	NO	Origin	Asian
		Previous trisomy 21	unknown
		Pregnancies	unknown

Biochemical Data	Ultrasound Data
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Parameter	Value	Corr Mom	
PAPP-A	4.90 mIU/ml	1.28	Gestational age 11+6
fb-hCG	91.4 ng/ml	2.33	Method CRL (<>Robinson)
			Scan date 2/5/2024

Risks at sampling date		Ultrasound Data	
Age Risk	1:556	Crown rump length in mm	46.6
Biochemical T21 risk	1:735	Nuchal translucency MoM	1.65
Combined trisomy 21 risk	1:251	Nasal bone	Present
Trisomy 13/18 + NT	<1:10000	Sonographer	DR.
		Qualifications in measuring NT	MBBS

Risk	Down's Syndrome Risk (Trisomy 21 Screening)
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The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 251 women with the same data, there is one woman with a trisomy 21 pregnancy and 250 women with not affected pregnancies. 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! The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998). The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

**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