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Date of Report 1/5/2024
PRISCA 5.2.0.13

Patient Data				
Name	SONIA TANWAR		Patient ID	012404300065
Birthday	14/8/1986		Sample ID	11494133
Age at sample	37.7		Sample Date	30/04/2024
Gestational age	12+5			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21 unknown
Weight in kg	62	Diabetes	NO	Pregnancies unknown
Smoker	NO	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+4
PAPP-A	5.60 mIU/ml	1.00	Method	CRL (<>Robinson)
fb-hCG	33.5 ng/ml	0.95	Scan date	29/04/2024
Risks at sampling date			Crown rump length in mm	62.3
Age Risk		1:144	Nuchal translucency MoM	0.5
Biochemical T21 risk		1:1002	Nasal bone	Present
Combined trisomy 21 risk		1:4958	Sonographer	DR.
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 4958 women with the same data, there is one woman with a trisomy 21 pregnancy and 4957 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>	
			<p>Trisomy 13/18 + NT</p> <p>The calculated risk for Trisomy 13/18 (with NT) is < 1:10000, which indicates a low risk</p>	

Risk Above Cut Off

Risk above Age Risk

Risk below Age risk