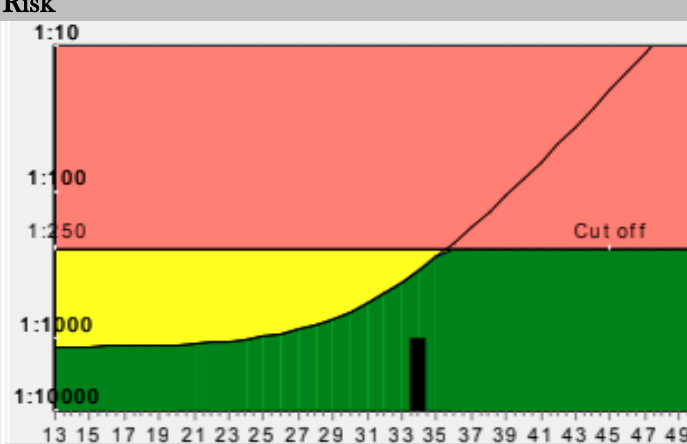




Date of Report 7/1/2024  
PRISCA 5.2.0.13

Patient Data				
Name	AKANKSHA AMAN WALIA		Patient ID	12401050119
Birthday	24/02/1990		Sample ID	11790935
Age at sample	33.9		Sample Date	5/1/2024
Gestational age	13+3			
Correction factors				
Fetuses	1	IVF	unknown	Previous trisomy 21 unknown
Weight in kg	50	Diabetes	NO	Pregnancies unknown
Smoker	NO	Origin	Asian	
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	13+2
PAPP-A	5.14 mIU/ml	0.55	Method	CRL (<>Robinson)
fb-hCG	50.3 ng/ml	1.65	Scan date	4/1/2024
Risks at sampling date			Crown rump length in mm	72.1
Age Risk		1:356	Nuchal translucency MoM	0.84
Biochemical T21 risk		1:171	Nasal bone	Present
Combined trisomy 21 risk		1:1026	Sonographer	DR.SHRUTI SANGWAN
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 1026 women with the same data, there is one woman with a trisomy 21 pregnancy and 1025 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