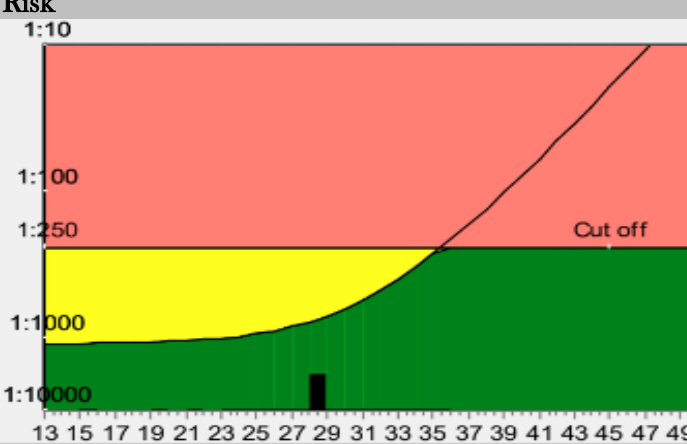




Date of Report 4/9/2023  
PRISCA 5.2.0.13

Patient Data					
Name	ASHA	Patient ID	012309030208		
Birth day	18/03/1995	Sample ID	11781247		
Age at sample	28.5	Sample Date	3/9/2023		
Gestational age	12+3				
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	47	Diabetes	NO	Pregnancies	unknown
Smoker	NO	Origin	Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	12+2	
PAPP-A	4.5 mIU/ml	0.65	Method	CRL (<>Robinson)	
fb-hCG	48.2 ng/ml	1.15	Scan date	2/9/2023	
Risks at sampling date			Crown rump length in mm	58	
Age Risk	1:760		Nuchal translucency MoM	1.18	
Biochemical T21 risk	1:1260		Nasal bone	Present	
Combined trisomy 21 risk	1:3100		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 3100 women with the same data, there is one woman with a trisomy 21 pregnancy and 3099 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