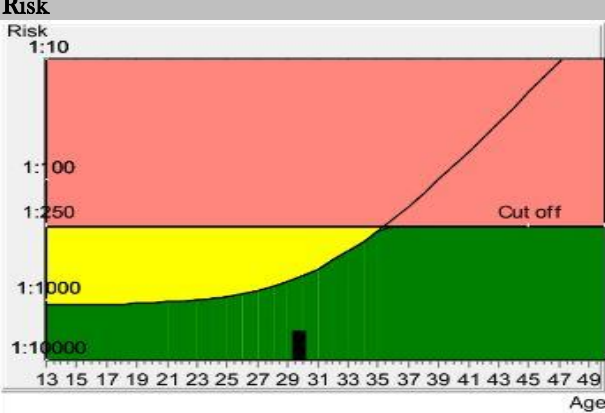



Date of Report 18-11-19
PRISCA 5.0.2.37

Patient Data					
Name	MRS MAMTA SAMANT		Patient ID	061911170001	
Birth day	07-03-90		Sample ID	80178394	
Age at delivery	29.7		Sample Date	17/11/2019	
Gestational age	12+0				
Correction factors					
Fetuses	1	IVF	unknown	Previous trisomy 21	unknown
Weight in kg	53	Diabetes	unknown	Pregnancies	unknown
Smoker	Unknown	Origin	Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	11+6	
PAPP-A	4.33 mIU/ml	1.02	Method	CRL (<>Robinson)	
fb-hCG	118.05 ng/ml	2.28	Scan Date	16-11-19	
Risks at sampling date			Crown rump length in mm	52	
Age Risk	1:657		Nuchal translucency MoM	0.86	
Biochemical T21 risk	1:579		Nasal Bone	Present	
Combined Trisomy 21 Risk	1:3031		sonographer	DR.RAJEEV SETHI	
Trisomy 13/18 + NT	<1:10000		Qualification in measuring NT	MBBS,DMRE	
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 more than 3031 women with the same data, there is one woman with a trisomy 21 pregnancy and 3030 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!</p>		
Trisomy 13/18 + NT			The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
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