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Date of Report 21-01-19
PRISCA 5.0.2.37

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Patient Data					
Name	N	Irs Moni Singh	Patient ID		011901190107
Birthday		15-11-93	Sample ID		10421987
Age at delivery		25.6	Sample Date		19/01/19
Gestational age		20+4			
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	78 Diabetes	S	no	Pregnancies	
Smoker	no Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	e	12+6
AFP	46.1 ng/ml	0.74	Method		CRL (⇔Hadlock)
uE3	2.41 mIU/ml	1.55	Scan Date		27-11-18
hCG	13616.5 ng/ml	0.96	CRL measurm	ents	70.0 mm
Risks at sampling date			NT Translucer	ncy	1.6
Age Risk		1:1348	NT Mom		0.91
Biochemical T21 Risk		1:3462	Nasal Bone		Present
Combined T21 Risk	<1:10000		Sonographer		DR.SANJEEV KUMAR SINGHAL
Trisomy 18	Frisomy 18 <1:10000		Qualifications in measuring NT		MBBS,PGDUS,DMRD
Risk Risk			Down's Syndrome Risk (Trisomy 21 Screening) The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.		
			After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.		
1:1000 1:10000 13 15 17 19 21 23 25	27 29 31 33 35 37 3	9 41 43 45 47 49	The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that risk calculations are statistical approach and have no diagnostic values! The patient combined risk presumes the NT		
		Ace	1		

The calculated risk for trisomy 18 is <1:10000, which

Trisomy 13/18 + NT

represents a low risk.

measurement was done according to accepted guidelines
The laboratory can not be held responsible for their impact on the risk

AFP (0.74) is located in the low risk area for neural tube defects.

assessment! Calculated value has no diagnostic value!The corrected MoM