

Risk Above Cut Off

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Risk below Age risk

 Date of Report
 27-12-2023

 PRISCA
 5.2.0.13

Patient Data Name Birthday Age at Sample date Gestational age Correction factors Fetuses Weight in kg Smoker Biochemical Data Parameter PAPP-A	78.8 NO Value	IVF Diabetes Origin		Sample ID Sample Date unknown NO Asian	Previous trisomy 21 Pregnancies	012312260098 11819498 26-12-2023 unknown unknown
Birthday Age at Sample date Gestational age Correction factors Fetuses Weight in kg Smoker Biochemical Data Parameter	78.8 NO Value	IVF Diabetes Origin	08-05-1996 27.6	Sample ID Sample Date unknown NO Asian	·	11819498 26-12-2023 unknown
Age at Sample date Gestational age Correction factors Fetuses Weight in kg Smoker Biochemical Data Parameter	78.8 NO Value	Diabetes Origin	27.6	Sample Date unknown NO Asian	·	26-12-2023 unknown
Gestational age Correction factors Fetuses Weight in kg Smoker Biochemical Data Parameter	78.8 NO Value	Diabetes Origin		unknown NO Asian	·	unknown
Correction factors Fetuses Weight in kg Smoker Biochemical Data Parameter	78.8 NO Value	Diabetes Origin	12+2	unknown NO Asian	·	
Fetuses Weight in kg Smoker Biochemical Data Parameter	78.8 NO Value	Diabetes Origin		NO Asian	·	
Weight in kg Smoker Biochemical Data Parameter	78.8 NO Value	Diabetes Origin		NO Asian	·	
Smoker Biochemical Data Parameter	NO Value	Origin		Asian	Pregnancies	unknown
Biochemical Data Parameter	Value					
Parameter						
				Ultrasound Data		
PAPP-A	39		Corr Mom	Gestational age	2	12+2
	0.2	mIU/ml	0.90	Method		CRL (<>Robinson)
fb-hCG	76.5	ng/ml	2.08	Scan date		26-12-2023
Risks at sampling date				Crown rump le	ength in mm	57.2
Age Risk			1:813	Nuchal translu	cency MoM	1.20
Biochemical T21 risk			1:694	Nasal bone		PRESENT
Combined trisomy 21 risk			1:1531	Sonographer		DR.SANJEEV KUMAR
Trisomy 13/18 + NT			<1:10000	Qualifications :	in measuring NT	MD
Risk				Down's Syndro	ome Risk (Trisomy 21	Screening)
1:100 1:250 Cut off 1:1000 1:1000 1:1000 1:1000 1:1000 1:1000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000000				The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 1531 women with the same data, there is one woman with a trisomy 21 pregnancy and 1530 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 aapproaches and have no diagnostic value! The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998). The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		

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