

Risk Above Cut Off

\*Free Home Sample Collection 9999 778 778



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Risk below Age risk

Date of Report 12-09-2023 PRISCA 5.2.0.13

1 IVF 48 Diabetes NO Origin  Value	24.7 11+6	Sample ID Sample Date  unknown Previous trisomy 2  NO Pregnancies  Asian	12309110093 11795071 11-09-2023 21 unknown unknown
1 IVF 48 Diabetes NO Origin	31-12-1998 24.7 11+6	Sample ID Sample Date  unknown Previous trisomy 2  NO Pregnancies  Asian	11795071 11-09-2023 21 unknown
48 Diabetes NO Origin	24.7 11+6	unknown Previous trisomy 2  NO Pregnancies  Asian	11-09-2023 21 unknown
48 Diabetes NO Origin	11+6	unknown Previous trisomy 2  NO Pregnancies  Asian	21 unknown
48 Diabetes NO Origin		unknown Previous trisomy 2  NO Pregnancies  Asian	
48 Diabetes NO Origin		NO Pregnancies Asian	
48 Diabetes NO Origin		NO Pregnancies Asian	
NO Origin		Asian	unknown
, ,			
Value			
Value		Ultrasound Data	
	Corr Mom	Gestational age	11+5
$4.65~\mathrm{mIU/ml}$	0.86	Method	CRL (<>Robinson)
56.2  ng/ml	1.17	Scan date	10-09-2023
		Crown rump length in mm	53.7
	1:950	Nuchal translucency MoM	1.05
	1:2982	Nasal bone	PRESENT
	<1:10000	Sonographer	DRDEEPIKA
	<1:10000	Qualifications in measuring NT	MD
		Down's Syndrome Risk (Trisomy	y 21 Screening)
1:100 1:250 Cut off 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49  Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk		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 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 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	
	4.65 mIU/ml 56.2 ng/ml	4.65 mIU/ml 0.86 56.2 ng/ml 1.17  1:950 1:2982 <1:10000 <1:10000	Value Corr Mom  4.65 mIU/ml 0.86 Method  56.2 ng/ml 1.17 Scan date  Crown rump length in mm  1:950 Nuchal translucency MoM  1:2982 Nasal bone  Sonographer  <1:10000 Qualifications in measuring NT  Down's Syndrome Risk (Trisomy 2 cut off, which represents a low ris After the result of the Trisomy 21 test expected that among 10000 women we there is one woman with a trisomy 21 19999 women with not affected pregnather information provided by the referring the risk calculations are statistical aappeliagnostic value!  The patient combined risk presumes done according to accepted guidelines 1998).  The laboratory cannot be hold response.

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