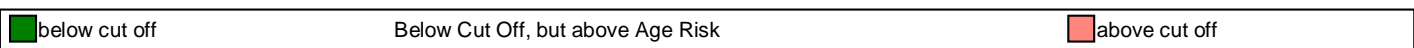


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Patient data			
Name	MRS. ARTI	Patient ID	
Birthday	20/08/96	Sample ID	2305042714/NOD
Age at sample date	26.8	Sample Date	22/05/23
Gestational age		12 + 2	
Correction factors			
Fetuses	1	IVF	no
Weight	53.2	diabetes	no
		Previous trisomy 21 pregnancies	no
Smoker	no	Origin	Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.98 mIU/ml	0.48	Gestational age 12 + 0
fb-hCG	40.2 ng/ml	0.84	Method CRL Robinson
Risks at sampling date		Scan date	20/05/23
Age risk		Crown rump length in mm	56.1
Biochemical T21 risk		Nuchal translucency MoM	0.68
Combined trisomy 21 risk		Nasal bone	present
Trisomy 13/18 + NT Risk		Sonographer	DR. R.S. CHAUHAN
1:10		Qualifications in measuring NT	MD
1:100		Trisomy 21	
1:250		<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 7793 women with the same data, there is one woman with a trisomy 21 pregnancy and 7792 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 risk calculations are statistical approaches and have no diagnostic value!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>	
1:1000			
1:10000			
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			
Age			

Trisomy 13/18 + NT
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.



Sign of Physician

