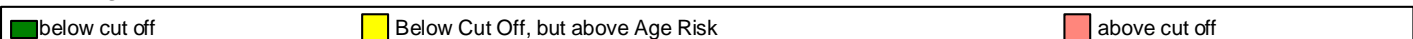


Patient data		MRS. WAHIDA		Patient ID	
Name		15/03/2000		Sample ID	
Birthday				231 1006614/NOD	
Age at sample date		23.6		Sample Date	
				4/11/2023	
Gestational age		11 + 5			
Correction factors					
Fetuses		1 IVF		no Previous trisomy 21	
Weight		65 diabetes		no pregnancies	
Smoker		no Origin		Asian	
Biochemical data				Ultrasound data	
Parameter		Value		Corr. MoM	
PAPP-A		1.97 mIU/ml		0.79	
fb-hCG		38.2 ng/ml		0.80	
				Gestational age	
				10 + 5	
				Method	
				CRL Robinson	
				Scan date	
				28/10/2023	
Risks at sampling date				Crown rump length in mm	
Age risk		1:982		40.6	
Biochemical T21 risk		1:5965		Nuchal translucency MoM	
Combined trisomy 21 risk		<1:10000		0.70	
Trisomy 13/18 + NT		<1:10000		Nasal bone	
Risk		1:10		present	
1:100				Sonographer	
1:250				..	
Cut off				Qualifications in measuring NT	
				..	
				Trisomy 21	
				<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></p> <p>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.</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					
Trisomy 13/18 + NT				Age	

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



Sign of Physician