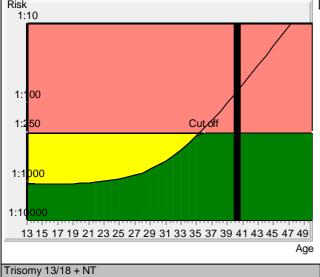
Prisca 5.2.0.13

Date of report: 10/05/2024

Patient data						
		1450	\/ANDAN::	ln		
Name				Patient ID		
Birthday	22/01/1984			Sample ID		2405220432/AMB
Age at sample date	40.3			Sample Date		9/05/2024
Gestational age	12 + 2					
Correction factors						
Fetuses	1	IVF		no	Previous trisomy 21	no
Weight	49.8	diabetes		no	pregnancies	
Smoker	no	Origin		Asian		
Biochemical data				Ultrasound data		
Parameter	Value Corr. MoN		Gestational age		12 + 0	
PAPP-A	1 mIU/ml 0.22		Method		CRL Robinson	
fb-hCG	365 ng/ml 7.42		Scan date		7/05/2024	
Risks at sampling date				Crown rump length in mm		55.3
Age risk 1:72			Nuchal translucency MoM		1.21	
Biochemical T21 risk >1:50			Nasal bone		present	
Combined trisomy 21 risk >1:50			Sonographer			
Trisomy 13/18 + NT 1:138			Qualifications in measuring NT			
Risk				Trisomy 21		



The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:138, which represents a low risk. The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.

After the result of the Trisomy 21 Test (with nuchal translucency), it is expected that among less than 50 pregnancies with the same data, there is one trisomy 21 pregnancy.

The free beta HCG level is high.

The PAPP-A level is low.

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!

The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).

The laboratory can not be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic value!

