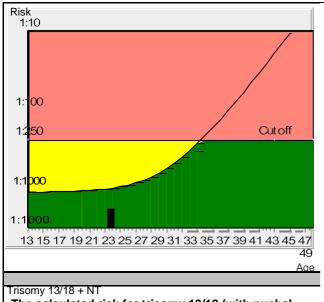
Date of report:

24-11-2018

Patient data						
Name		MRS. RENU	Patient ID		1806220134/AMB	
Birthday		20-04-199	7 Sample ID		1806220134/AMB	
Age at sample date	21		Sample Date		23-11-2018	
Gestational age	12 + 4					
Correction factors						
	1					
Fetuses		IVF		Previous trisomy 21	no	
Weight	48	diabetes		pregancies		
Smokernour data	no l	Origin		.ata		
Parameter	Value	Corr. Mo	M Gestational	l age	12 + 4	
PAPP-A	3.8 mIU/ml 1.28		Method		CRL Robinson	
fb-hCG	37.1 ng/ml 1.07		7 Scan date			
Risks at sampling date			Crown rum	Crown rump length in mm		
Age risk 1:1006			Nuchal tran	Nuchal translucency MoM		
Biochemical T21 risk 1:8960			Nasal bone	Nasal bone presen		
Combined trisomy 21 risk <1:10000) Sonograph	Sonographer .		
Trisomy 13/18 + NT <1:10000			Qualificatio	Qualifications in measuring NT MD		



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

Trisomy 2

The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.

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.

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 NTmeasurement 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!

Sign of Physician