Prisca 5.2.0.13

Date of report: 9/08/2024

Patient data					
Name	MRS. CHERCHI HELEN				
Birthday	27/07/1998				2408015627/NOD
Age at sample date	26.0			e	9/08/2024
Gestational age 12 + 1					
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21	no
Weight	87	diabetes	no	pregnancies	
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter Val	Value Corr. MoM		Gestational age 11 + 5		
PAPP-A 2.6 m	nIU/m	1.22	Method		CRL Robinson
fb-hCG 118 n	g/ml	2.80	Scan date		6/08/2024
Risks at sampling date			Crown rump length in mm 51.65		
Age risk				Nuchal translucency MoM 0.5	
Biochemical T21 risk 1:663			Nasal bone present		
Combined trisomy 21 risk 1:3618			Sonographer		
risomy 13/18 + NT <1:10000			Qualifications in measuring NT		
Risk 1:10			Trisomy 21 The calculated risk for Trisomy 21 (with nuchal		
1:100 1:250 1:1000 1:10000 13 15 17 19 21 23 25 27 29 3	Cut off 5 37 39 41 43 45 47 49 Age	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 3618 women with the same data, there is one woman with a trisomy 21 pregnancy and 3617 women with not affected pregnancies. The free beta HCG level is high. 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!			

Sign of Physician

Trisomy 13/18 + NT

risk.

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