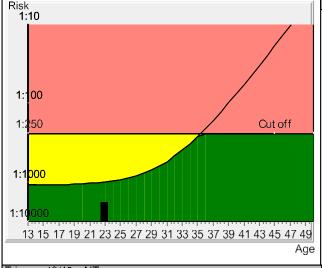
Prisca 5.1.0.17

Date of report: 01-06-2023

Patient data						
Name	MRS. KAVITA			Patient ID		
Birthday	03-08-2000			Sample ID		2305062230/NOD
Age at sample date	22.8			Sample Date		31-05-2023
Gestational age	12 + 0					
Correction factors						
Fetuses	1	IVF		no	Previous trisomy 21	no
Weight	58	diabetes		no	pregnancies	
Smoker	no	Origin		Asian		
Biochemical data				Ultrasound data		
Parameter	Value		Corr. MoM	Gestational	age	11 + 6
PAPP-A	2.66 mIU/mI		0.81	Method		CRL Robinson
fb-hCG	81.6 ng/ml 1.69		Scan date		30-05-2023	
Risks at sampling date			Crown rump length in mm		55	
Age risk			1:1016	Nuchal trans	slucency MoM	0.89
Biochemical T21 risk 1:1152			Nasal bone		present	
Combined trisomy 21 risk 1:5915			Sonographer			
Trisomy 13/18 + NT <1:10000			Qualifications in measuring NT MD			
Risk				Trisomy 21		



Trisomy 13/18 + NT

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

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 5915 women with the same data, there is one woman with a trisomy 21 pregnancy and 5914 women with not affected pregnancies.

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!