

Patient data			
Name Mrs. NIDA IRFAN MULLA TWIN A		Patient ID 0662401250113	
Birthday 02-03-2002		Sample ID A0125530	
Age at sample date 21.9		Sample Date 25-01-2024	
Gestational age 12 + 4			
Correction factors			
Fetuses 2	IVF no	Previous trisomy 21 pregnancies unknown	
Weight 52.6	diabetes no		
Smoker no	Origin Asian		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age 11 + 6
PAPP-A	15.12 mIU/mL	1.73	Method CRL Robinson
fb-hCG	41.23 ng/mL	0.44	Scan date 20-01-2024
Risks at sampling date			Crown rump length in mm 55
Age risk 1:1059			Nuchal translucency MoM 0.89
Biochemical T21 risk <1:10000			Nasal bone present
Combined trisomy 21 risk <1:10000			Sonographer NA
Trisomy 13/18 + NT <1:10000			Qualifications in measuring NT NA
Risk 1:10 1:100 1:250 1:1000 1:10000 Age			

Trisomy 21

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 risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.

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