Prisca 5.2.0.13

Date of report: 29/10/2024

## JITM Diagnostics

Patient data			
Name	MRS. NIKITA	Patient ID	
Birthday	19/05/2003	Sample ID	2410053727/NOD
Age at sample date	21.4	Sample Date	29/10/2024
Gestational age	13 + 6		
Correction factors			
Fetuses 1	IVF	no	Previous trisomy 21 no
Weight 37	diabetes	no	pregnancies
Smoker no	Origin	Asian	
Biochemical data		Ultrasound data	
Parameter Value	Corr. MoM	Gestational	age 13 + 5
PAPP-A 8.79 mIU/n	nl 0.76	Method CRL Robinson	
fb-hCG 26.4 ng/ml	0.57	Scan date	28/10/2024
Risks at sampling date		Crown rump length in mm 80.5	
Age risk	1:1112	1:1112 Nuchal translucency MoM 1.01	
Biochemical T21 risk	<1:10000	Nasal bone present	
Combined trisomy 21 risk	<1:10000 Sonographer DR. KHEMENDRA KUMAR		
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT M.D	
Risk 1:10 1:250 Cut off 1:10000 1:11000 1:110000 1		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 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