Prisca 5.2.0.13

Date of report: 30/09/2024

## JITM Diagnostics

Patient data					
Name MR	MRS. JASPREET KAUR				
Birthday	13/04/1990	Sample ID		2409063344/NOD	
Age at sample date	34.5		e	29/09/2024	
Gestational age	11 + 0				
Correction factors					
Fetuses 1	IVF	no	Previous trisomy 21	no	
Weight 64	diabetes	no	pregnancies		
Smoker no	Origin	Asian			
Biochemical data			Ultrasound data		
Parameter Value	Corr. MoM	<u> </u>			
PAPP-A 1.85 mIU/m	1.07	Method CRL Robinson			
fb-hCG 54.1 ng/ml	1.03				
Risks at sampling date			Crown rump length in mm 40.		
Age risk	1:286	Nuchal translucency MoM Nasal bone		0.87	
Biochemical T21 risk			_	present	
Combined trisomy 21 risk			er	DR. RAHUL KUMAR	
Trisomy 13/18 + NT	<1:10000	Trisomy 21	is in measuring NT	M.D	
1:10  1:250  1:10000  1:3 15 17 19 21 23 25 27 29 31 33 33  Trisomy 13/18 + NT  The calculated risk for trisomy 13 translucency) is < 1:10000, which 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 9177 women with the same data, there is one woman with a trisomy 21 pregnancy and 9176 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!				