Prisca
 5.2.0.13

 Date of report:
 17/02/2025

## JITM Diagnostics

Patient data				
Name	MRS. RASHI	Patient ID		
Birthday	19/12/1989	Sample ID		2502030776/NOD
Age at sample date	35.2	Sample Date	9	16/02/2025
Gestational age	13 + 1			
Correction factors				
Fetuses 1	IVF	no	Previous trisomy 21	no
Weight 44.2	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data			Ultrasound data	
Parameter Value	Corr. MoM	Gestational age		13 + 0
PAPP-A 10.1 mlU/m	l 1.38	Method CRL Robinsor		
fb-hCG 79.1 ng/ml	1.69	Scan date		15/02/2025
Risks at sampling date		Crown rump length in mm 70		70.21
Age risk	1:268	Nuchal trans	slucency MoM	0.80
Biochemical T21 risk	1:915	Nasal bone		present
Combined trisomy 21 risk			r	DR. SHITIJ BUDHIRAJA
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT		M.D
Risk 1:10         1:10         1:100         1:250         Cut off         1:1000         1:1000         1:1000         1:1000         1:1000         1:1000         1:1000         1:1000         1:10000         1:10000         1:10000         1:10000         1:10000         Age         Trisomy 13/18 + NT         The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.		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 4525 women with the same data, there is one woman with a trisomy 21 pregnancy and 4524 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!		

Sign of Physician