## JITM Diagnostics

Patient data					
Name	MRS. SAPNA		Patient ID		
Birthday	20/02/1990		Sample ID 2502054717/N		
Age at sample date	35.0	Sample Date	e 28/02/20	)25	
Gestational age	12 + 5				
Correction factors					
Fetuses 1	IVF	no	Previous trisomy 21	no	
Weight 61	diabetes	no	pregnancies		
Smoker no	Origin	Asian			
Biochemical data		Ultrasound data			
Parameter Value	Value Corr. MoM Gesta		estational age 12 + 2		
PAPP-A 3.28 mIU/m	l 0.77	Method	CRL Robin	son	
fb-hCG 31.1 ng/ml	0.71	Scan date	25/02/2	025	
Risks at sampling date		Crown rump	b length in mm	59.2	
Age risk	1:272		Nuchal translucency MoM 1.88		
Biochemical T21 risk	emical T21 risk 1:1972		Nasal bone presen		
Combined trisomy 21 risk 1:304		Sonographer			
Trisomy 13/18 + NT	IT 1:1504		Qualifications in measuring NT		
Risk   1:10   1:00   1:250   1:10000   1:10000   1:10000   1:10000   1:10000   1:10000   1:10000   1:10000   1:10000   1:10000   1:10000   1:10000   1:10000   1:10000   1:10000   1:10000   1:10000   1:10000   1:10000   1:1504, which reported the set of the	Age 3/18 (with nuchal	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 304 women with the same data, there is one woman with a trisomy 21 pregnancy and 303 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