

 Date of Report
 20/3/2023

 PRISCA
 5.2.0.13

Patient Data					
Name	ame MRS. RASHI		Patient ID		012303180301
Birthday		11/2/1996	Sample ID		11551771
Age at term		27.7	Sample Date		18/3/2023
Gestational age		12+0			
Correction factors	<u> </u>				
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	73 Diabetes		NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom			11+6
PAPP-A	$2.65~\mathrm{mIU/ml}$	0.76	Method		CRL (<>Robinson)
fb-hCG	23.7 ng/ml	0.59	Scan date		18/3/2023
Risks at sampling date			Nuchal translucency 1		
Age Risk		1:837	Nuchal translucency MoM		0.72
Biochemical T21 risk		1:8709	Nasal bone Prese		Present
Combined trisomy 21 ris	sk	<1:10000			
Trisomy 13/18 + NT		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 Cut off 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000			The calculated risk for Trisomy 21(with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21with NT test it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy and 9999 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 the risk calculations are statistical aapproaches and have no diagnostic value!		
The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk			on the risk assessment! Calculated risks have no diagnostic values		
Risk	Above Cut Off		Risk above Ag	e Risk	Risk below Age risk