

 Date of Report
 28/12/2022

 PRISCA
 5.1.0.17

Patient Data						
Name MRS. ROJI				Patient ID		012212260304
Birthday			24/10/1994	Sample ID		11507312
Age at term			28.08	Sample Date		26/12/2022
Gestational age 12+4						
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	1 unknown
Weight in kg	62	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom			12+3
PAPP-A	3.42	mIU/ml	0.64	Method		CRL (<>Robinson)
fb-hCG	34.5	ng/ml	0.94	Scan date		26/12/2022
Risks at sampling date				Nuchal translucency 1.1		
Age Risk			1:785	Nuchal translucency MoM 0.72		
Biochemical T21 risk			1:1993	Nasal bone Present		
Combined trisomy 21 risk	-		<1:10000			
Trisomy 13/18 + NT			<1:10000			
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:1000 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk				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 laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
Risk A	Above Cı	ıt Off		Risk above Ag	e Risk	Risk below Age risk