

 Date of Report
 13/6/2023

 PRISCA
 5.1.0.17

Patient Data						
Name	e MRS. ARCHNA					012306100279
Birthday			6/10/1996	Sample ID		11668133
Age at term			27.1	Sample Date		10/6/2023
Gestational age			12+2			
Correction factors		1				
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	51	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational age	2	12+2
PAPP-A	3.45	mIU/ml	0.58	Method		CRL (<>Robinson)
fb-hCG	27.7	ng/ml	0.66	Scan date		10/6/2023
Risks at sampling date				Nuchal translucency (NT) 1.2		
Age Risk	ge Risk 1:8			Nuchal translucency MoM 0.77		
Biochemical T21 risk	T21 risk			Nasal bone present		
Combined trisomy 21 risk			<1:10000			
Trisomy 13/18			<1:10000			
Risk				Down's Syndro	ome Risk (Trisomy 2	21 Screening)
				The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 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!		
Trisomy 13/18+NT The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk				The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values		
Risk Al	ıt Off		Risk above Ag	e Risk	Risk below Age risk	