

 Date of Report
 18/9/2022

 PRISCA
 5.1.0.17

n : n					
Patient Data					
Name MRS. BHAWNA				012209150250	
Birthday			Sample ID		11245912
Age at term		27.07	Sample Date		15/9/2022
Gestational age		12+2	2		
Correction factors	<u> </u>				
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	46 Diabete	S	NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	12+1
PAPP-A	$3.24~\mathrm{mIU/m}$	0.48	Method		CRL (<>Robinson)
fb-hCG	29.7 ng/ml	0.68	Scan date		15/9/2022
Risks at sampling date			Nuchal translucency 1.4		
Age Risk		1:846	Nuchal translucency MoM 0.9		
Biochemical T21 risk		1:2048	Nasal bone Presen		
Combined trisomy 21 r	1:9965				
Trisomy 13/18 + NT	<1:10000				
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:250 1:1000 1:10000 13 15 17 19 21 23 25 Trisomy 13/18+NT The calculated risk for <1:10000, which indica	Trisomy 13/18 (wi	Age	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 with NT test it is expected that among 9965 women with the same data, there is one woman with a trisomy 21 pregnancy and 9964 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		
The calculated risk for <1:10000 , which indicates		th NT) is	on the risk ass	essment! Calculated risks	