

 Date of Report
 21/11/2022

 PRISCA
 5.1.0.17

Patient Data					
Name MRS. MAMONI MUNDA			Patient ID		012211190265
Birthday 19/05/1995		Sample ID		11320870	
Age at term 27.11		Sample Date 20/11/2022			
Gestational age 13+2					
Correction factors					
Fetuses	1 IVF		unknown	Previous trisomy 21	unknown
Weight in kg	60 Diabetes	3	NO	Pregnancies	unknown
Smoker	NO Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age	2	13+1
PAPP-A	3.57 mIU/ml	0.5	Method		CRL (<>Robinson)
fb-hCG	19.3 ng/ml	0.64	Scan date		19/11/2022
Risks at sampling date			Nuchal translucency 1.6		
Age Risk		1:850	Nuchal translucency MoM 0.92		
Biochemical T21 risk		1:2523	Nasal bone Presen		
Combined trisomy 21 risk <1		<1:10000			
Trisomy 13/18 + NT <1:100		<1:10000			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:100 1:1000 1:1000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:1			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		