

<1:10000, which indicates a low risk

Risk Above Cut Off

Risk below Age risk

					Date of Report PRISCA	3/8/2022 5.1.0.17
Patient Data						
Name	:	MRS. SU	NIL KUMARI	Patient ID		012208020117
Birthday			30/12/1986	Sample ID		11594749
Age at term			36	Sample Date		2/8/2022
Gestational age			11+2			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	59	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data				Ultrasound Data		
Parameter	Value		Corr Mom	Gestational ago	e	11+2
PAPP-A	2.15	mIU/ml	0.65	Method		CRL (<>Robinson)
fb-hCG	26.4	ng/ml	0.52	Scan date		2/8/2022
Risks at sampling date				Nuchal translucency 1.3		
Age Risk			1:227	Nuchal translu	icency MoM	1.02
Biochemical T21 risk			1:2076	Nasal bone		Present
Combined trisomy 21 risk			1:8087			
Trisomy 13/18 + NT			<1:10000			
Risk				Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10 1:250 Cut off 1:1000 1:1000 1:3 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49				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 8087 women with the same data, there is one woman with a trisomy 21 pregnancy and 8086 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				The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic		

values

Risk above Age Risk