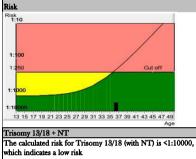




Date of Report 21-05-2019

					PRISCA	5.0.2.37
Patient Data						
Name			Mrs Harshe	Patient ID		011905200018
Birthday			11/1/1983	Sample ID		10420175
Age at sample date			36.4	Sample Date		20/05/2019
Gestational age by CRL			13+6			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	73.6	Diabetes		unknown	Pregnancies	unknown
Smoker	Unknown	Origin		Asian		

Smoker	Unknown Origin		Asian		
Biochemical Data			Ultrasound Data		
Parameter	Value	Corr Mom	Gestational age by CRL	13+0	
PAPP-A	$5.86~\mathrm{mIU/ml}$	1.07	Method	CRL (◇Robinson)	
fb-hCG	24.6 ng/ml	0.65	Scan Date	15/05/2019	
Risks at sampling date			Crown rump length (mm)	67.5	
Age Risk		1:210	Nuchal translucency MoM	0.58	
Biochemical T21 Risk		1:3759	Nasal Bone	Present	
Combined Trisomy 21 Risk		<1:10000	Sonographer	DR. RITU JAIN	
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT		
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
Risk 1:10			The calculated risk for Trisomy cut off, which represents a low r		



After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21

same data, mere is one woman with a unsoing 2.1
pregnancy.
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





