



Patient data			
Name	MRS. RAMANPREET	Patient ID	
Birthday	24-03-1997	Sample ID	1805220059/AMB
Age at sample date	21.5	Sample Date	08-10-2018
Gestational age	13+2		
Correction factors			
Fetuses	1 IVF	unknown	Previous trisomy 21 unknown
Weight	72 diabetes	unknown	pregancies
Smoker	unknown Origin		Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	6.5 mIU/ml	1.23	13 + 2
fb-hCG	72.1 ng/ml	1.73	Method CRL Robinson
Risks at sampling date			Scan date 03-10-2018
Age risk		1:1000	Crown rump length in mm
Biochemical T21 risk		1:2603	Nuchal translucency MoM
Combined trisomy 21 risk		<1:10000	Nasal bone unknown
Trisomy 13/18 + NT Risk		<1:10000	Sonographer .
			Qualifications in measuring NT MD
			Trisomy 21
			The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
			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 pregnancy.
			The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.
			Please note that risk calculations are statistical approaches and have no diagnostic value!
			The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).
			The laboratory can not hold responsible for their impact on the risk assessment ! Calculated risks have no
			diagnostic value!

 below cut off	Trisomy 13/18 + NT	Below Cut Off, but above Age Risk	 above cut off
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The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low

risk.

Sign of Physician

