

Prisca 5.0.2.37
Date of report: 15/07/18

Patient data		
Name	MRS. KARAMJEET	Patient ID 1807220397/AMB
Birthday	02/03/84	Sample ID 1807220397/AMB
Age at sample date	34.4	Sample Date 14/07/18
Gestational age	13 + 0	
Correction factors		
Fetuses	1 IVF	no Previous trisomy 21 no
Weight	65 diabetes	no pregnancies
Smoker	no Origin	Asian
Biochemical data		Ultrasound data
Parameter	Value	Corr. MoM
PAPP-A	3.27 mIU/ml	0.87
fb-hCG	33.1 ng/ml	1.05
Risks at sampling date		Gestational age 12 + 5
Age risk	1:317	Method CRL Robinson
Biochemical T21 risk	1:1317	Scan date 12/07/18
Combined trisomy 21 risk	1:301	Crown rump length in mm 66.7
Trisomy 13/18 + NT	1:8290	Nuchal translucency MoM 1.77
Risk		Nasal bone present
		Sonographer .
1:1000		Qualifications in measuring NT MD
1:10000		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 301 women with the same data, there is one woman with a trisomy 21 pregnancy and 300 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 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 be hold responsible for their impact on the risk assessment ! Calculated risks have no
		diagnostic value!
1315 1719 212325 2729 313335 3739 414345 4749		
Age		

below cut off
 Below Cut Off, but above Age Risk
 above cut off

The calculated risk for Trisomy 13/18 (with nuchal

translucency) is 1:8290, which represents a low risk.

Sign of Physician

