

Prisca 5.0.2.37  
Date of report: 20-09-2017

Patient data			
Name	MRS. NEELAM	Patient ID	1709220425/AMB
Birthday	07-10-1988	Sample ID	1709220425/AMB
Age at sample date	28.9	Sample Date	18-09-2017
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	54	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.6 mIU/ml	0.56	Gestational age 12 + 3
fb-hCG	26.6 ng/ml	0.79	Method CRL Robinson
Risks at sampling date			Scan date 14-09-2017
Age risk		1:740	Crown rump length in mm 62.7
Biochemical T21 risk		1:1916	Nuchal translucency MoM 0.74
Combined trisomy 21 risk		<1:10000	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer .
			Qualifications in measuring NT MD
		<b>Trisomy 21</b> <b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b> 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 be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!	
Trisomy 13/18 + NT			
<b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b>			

Sign of Physician

