

Prisca                      5.0.2.37  
**Date of report:        09-05-2017**

Patient data			
Name	MRS. SUKHVEER KAUR	Patient ID	1705220247/AMB
Birthday	18-10-1990	Sample ID	1705220247/AMB
Age at sample date	26.6	Sample Date	08-05-2017
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	78	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.26 mIU/ml	1.76	Gestational age
fb-hCG	48.8 ng/ml	1.47	Method
			Scan date
Risks at sampling date			Crown rump length in mm
Age risk		1:882	Nuchal translucency MoM
Biochemical T21 risk		1:6583	Nasal bone
Combined trisomy 21 risk		<1:10000	Sonographer
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT
			MD
Trisomy 21			
<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></p> <p>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.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>			
Trisomy 13/18 + NT			
<p><b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b></p>			

**Sign of Physician**

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