

Prisca                      5.1.0.17  
**Date of report:        03-12-2022**

Patient data			
Name	MRS. RAJWINDER KAUR	Patient ID	
Birthday	17-01-2001	Sample ID	2212220047/AMB
Age at sample date	21.9	Sample Date	02-12-2022
Gestational age	12 + 0		
Correction factors			
Fetuses	1	IVF	yes
Weight	61	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.83 mIU/ml	0.59	Gestational age 11 + 5
fb-hCG	29.4 ng/ml	0.62	Method CRL Robinson
			Scan date 30-11-2022
Risks at sampling date			Trisomy 21
Age risk		1:1038	<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 1191 women with the same data, there is one woman with a trisomy 21 pregnancy and 1190 women with not affected pregnancies.</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>
Biochemical T21 risk		1:5211	
Combined trisomy 21 risk		1:1191	
Trisomy 13/18 + NT		1:2568	
			Qualifications in measuring NT MD
Trisomy 13/18 + NT			
<p><b>The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:2568, which represents a low risk.</b></p>			

**Sign of Physician**