

Prisca 5.2.0.13
Date of report: 18/07/2024

Patient data			
Name	MRS. SARABJIT KAUR	Patient ID	
Birthday	8/10/1988	Sample ID	2407220815/AMB
Age at sample date	35.8	Sample Date	17/07/2024
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	51	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.1 mIU/ml	0.53	Gestational age
fb-hCG	41.1 ng/ml	0.91	Method
			CRL Robinson
			Scan date
			15/07/2024
Risks at sampling date			Trisomy 21
Age risk		1:233	The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.
Biochemical T21 risk		1:380	After the result of the Trisomy 21 test (with NT) it is expected that among 132 women with the same data, there is one woman with a trisomy 21 pregnancy and 131 women with not affected pregnancies.
Combined trisomy 21 risk		1:132	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!
Trisomy 13/18 + NT		1:1236	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			
The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:1236, which represents a low risk.			

Sign of Physician

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