

Prisca 5.1.0.17
Date of report: 20/03/25

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Patient data			
Name	A1860189	Patient ID	0482503190256
Birthday	30/12/94	Sample ID	A1860189
Age at sample date	30.2	Sample Date	19/03/25
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	67	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	4.08 mIU/mL	0.93	12 + 6
fb-hCG	34.41 ng/mL	1.00	Method
			CRL Robinson
			Scan date
			17/03/25
Risks at sampling date			Trisomy 21
Age risk	1:642		The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
Biochemical T21 risk	1:3422		After the result of the Trisomy 21 test (with NT) it is expected that among 9996 women with the same data, there is one woman with a trisomy 21 pregnancy and 9995 women with not affected pregnancies.
Combined trisomy 21 risk	1:9996		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	<1:10000		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 < 1:10000, which represents a low risk.			

Sign of Physician

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