

Prisca 5.1.0.17
Date of report: 02/03/25

NA

Patient data			
Name	Mrs. MANISHA	Patient ID	0012503010426
Birthday	04/11/00	Sample ID	a1839166
Age at sample date	24.3	Sample Date	01/03/25
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	48	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.32 mIU/mL	0.44	12 + 4
fb-hCG	44.76 ng/mL	1.00	Method
			CRL Robinson
			Scan date
			01/03/25
Risks at sampling date			Trisomy 21
Age risk	1:990		The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
Biochemical T21 risk	1:838		After the result of the Trisomy 21 test (with NT) it is expected that among 5163 women with the same data, there is one woman with a trisomy 21 pregnancy and 5162 women with not affected pregnancies.
Combined trisomy 21 risk	1:5163		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