

Prisca 5.2.0.13
Date of report: 12/01/25

Patient data			
Name	MRS. NARESHU	Patient ID	
Birthday	19/08/99	Sample ID	2501220250/AMB
Age at sample date	25.4	Sample Date	11/01/25
Gestational age	12 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	58	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.08 mIU/ml	0.31	Gestational age 12 + 0
fb-hCG	62.5 ng/ml	1.32	Method CRL Robinson
			Scan date 10/01/25
Risks at sampling date			Trisomy 21
Age risk		1:931	The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
Biochemical T21 risk		1:147	After the result of the Trisomy 21 test (with NT) it is expected that among 1039 women with the same data, there is one woman with a trisomy 21 pregnancy and 1038 women with not affected pregnancies.
Combined trisomy 21 risk		1:1039	The PAPP-A level is low.
Trisomy 13/18 + NT		<1:10000	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!
			The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).
			The laboratory can not hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!
Trisomy 13/18 + NT			Sign of Physician
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			

