

Prisca 5.1.0.17
Date of report: 19-11-2025

N A

Patient data			
Name	Mrs. SEJAL MIMROTH		Patient ID
Birthday	17-12-2003		Sample ID
Age at sample date	21.9		Sample Date
Gestational age	12 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	62	diabetes	no
Smoker	no	Origin	Asian
			Previous trisomy 21 pregnancies
			unknown
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.1 mIU/mL	0.28	12 + 0
fb-hCG	39.76 ng/mL	0.97	Method
			CRL Robinson
			Scan date
			14-11-2025
Risks at sampling date			Crown rump length in mm
			56
Age risk			Nuchal translucency MoM
1:1058			0.82
Biochemical T21 risk			Nasal bone
1:256			unknown
Combined trisomy 21 risk			Sonographer
1:1743			N A
Trisomy 13/18 + NT			Qualifications in measuring NT
<1:10000			MD
Risk			Trisomy 21
1:10			The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
1:100			After the result of the Trisomy 21 test (with NT) it is expected that among 1743 women with the same data, there is one woman with a trisomy 21 pregnancy and 1742 women with not affected pregnancies.
1:250			The PAPP-A level is low.
1:1000			The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.
1:10000			Please note that risk calculations are statistical approaches and have no diagnostic value!
Age			The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!
Cut off			
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