

Patient data			
Name	MRS. RAMANDEEP KAUR	Patient ID	
Birthday	14/01/1993	Sample ID	2404220045/AMB
Age at sample date	31.2	Sample Date	2/04/2024
Gestational age	11 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	57	diabetes	no
Smoker	no	Origin	Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	0.9 mIU/ml	0.29	11 + 5
fb-hCG	36 ng/ml	0.73	Method
Risks at sampling date			CRL Robinson
Age risk Biochemical		1:534	Scan date
T21 risk		1:249	1/04/2024
Combined trisomy 21 risk		1:74	Crown rump length in mm
Trisomy 13/18 + NT		1:115	51.9
Risk			Nuchal translucency MoM
1:10			1.73
1:100			Nasal bone
1:250			present
1:1000			Sonographer
1:10000			..
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			Qualifications in measuring NT
Age			Trisomy 21

Trisomy 13/18 + NT  
**The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:115, which represents a low risk.**

**The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.**  
 After the result of the Trisomy 21 test (with NT) it is expected that among 74 women with the same data, there is one woman with a trisomy 21 pregnancy and 73 women with not affected pregnancies.  
 The PAPP-A level is low.  
 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 be held responsible for their impact on the risk assessment! Calculated risks have no diagnostic value!

