

Prisca 5.1.0.17

**Date of report:** 06/12/24

N A

Patient data		Patient ID	
Name	Mrs. P KOTESHWARAMMA	Patient ID	0352412040003
Birthday	16/01/99	Sample ID	a1251667
Age at sample date	25.9	Sample Date	04/12/24
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	70	diabetes	no
Smoker	no	Origin	Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.08 mIU/mL	0.98	Gestational age
fb-hCG	13.03 ng/mL	0.38	Method
Risks at sampling date		Scan date	12 + 6 CRL Robinson
Age risk	1:940	Crown rump length in mm	02/12/24 67
Biochemical T21 risk	<1:10000	Nuchal translucency MoM	0.94
Combined trisomy 21 risk	<1:10000	Nasal bone	present
Trisomy 13/18 + NT	<1:10000	Sonographer	N A
		Qualifications in measuring NT	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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.	
1:250		The free beta HCG 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!	
		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!	
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