

Prisca 5.1.0.17  
**Date of report:** 12/12/24

NA

Patient data			
Name	Mrs. POOJA BHARGAVA	Patient ID	0442412110008
Birthday	10/11/86	Sample ID	a1851959
Age at sample date	38.1	Sample Date	10/12/24
Gestational age	12 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	77	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	1.48 mIU/mL	0.59	Gestational age 12 + 1
fb-hCG	43.58 ng/mL	1.03	Method CRL Robinson
Risks at sampling date			
Age risk	1:128		Scan date 10/12/24
Biochemical T21 risk	1:220		Crown rump length in mm 58.6
Combined trisomy 21 risk	1:700		Nuchal translucency MoM 1.11
Trisomy 13/18 + NT	<1:10000		Nasal bone present
Sonographer NA			
Qualifications in measuring NT MD			
Trisomy 21			
<b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 700 women with the same data, there is one woman with a trisomy 21 pregnancy and 699 women with not affected pregnancies.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>			
Trisomy 13/18 + NT			
<b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b>			

Sign of Physician

below cut off

Below Cut Off, but above Age Risk

above cut off