

Prisca 5.1.0.17  
**Date of report: 28-03-2023**

Patient data			
Name	MRS. SHANTA DEVI	Patient ID	
Birthday	01-01-1991	Sample ID	2303220607/AMB
Age at sample date	32.2	Sample Date	27-03-2023
Gestational age	12 + 2		
Correction factors			
Fetuses	1	IVF	yes
Weight	52	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.48 mIU/ml	1.05	Gestational age 11 + 4
fb-hCG	48 ng/ml	0.99	Method CRL Robinson
Risks at sampling date			Scan date 22-03-2023
Age risk		1:462	Crown rump length in mm 50.62
Biochemical T21 risk		1:3269	Nuchal translucency MoM 0.49
Combined trisomy 21 risk		<1:10000	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer ..
			Qualifications in measuring NT MD
Trisomy 21			
<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></p> <p>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.</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 hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>			
Trisomy 13/18 + NT			
<p><b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b></p>			

**Sign of Physician**

below cut off
  Below Cut Off, but above Age Risk
  above cut off