

Prisca 5.1.0.17  
Date of report: 25-10-2024

N A

Patient data			
Name	W/O ANSUL KUMAR		Patient ID
Birthday	13-07-1992	Sample ID	
Age at sample date	32.3	Sample Date	
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	77	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.57 mIU/mL	0.69	
fb-hCG	43.1 ng/mL	1.30	
Risks at sampling date			
Age risk	1:473		
Biochemical T21 risk	1:700		
Combined trisomy 21 risk	1:3973		
Trisomy 13/18 + NT	<1:10000		
		Gestational age	13 + 1
		Method	CRL Robinson
		Scan date	23-10-2024
		Crown rump length in mm	71
		Nuchal translucency MoM	0.79
		Nasal bone	unknown
		Sonographer	N A
		Qualifications in measuring NT	MD
Risk		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 3973 women with the same data, there is one woman with a trisomy 21 pregnancy and 3972 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 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