

Prisca

5.2.0.13

Date of report:

28/09/2024

Patient data			
Name	MRS. KASHISH	Patient ID	
Birthday	6/01/2004	Sample ID	2409060014/NOD
Age at sample date	20.7	Sample Date	28/09/2024
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	91	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.38 mIU/ml	1.59	Gestational age 12 + 5
fb-hCG	69.1 ng/ml	1.80	Method CRL Robinson
Risks at sampling date			Scan date 27/09/2024
Age risk		1:1090	Crown rump length in mm 65.4
Biochemical T21 risk		1:4159	Nuchal translucency MoM 0.84
Combined trisomy 21 risk		<1:10000	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer ..
			Qualifications in measuring NT ..
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 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