

Prisca 5.1.0.17  
 Date of report: 20/12/25

Patient data			
Name	Mrs. JYOTI BHALSING FETUS-B	Patient ID	0662512190058
Birthday	19/10/99	Sample ID	B3948791 FETUS-B
Age at sample date	26.2	Sample Date	19/12/25
Gestational age	12 + 0		
Correction factors			
Fetuses	2	IVF	no
Weight	62.9	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.19 mIU/mL	0.57	11 + 6
fb-hCG	45.26 ng/mL	0.45	Method
			CRL Robinson
			Scan date
			18/12/25
			Crown rump length in mm
			54.3
			Nuchal translucency MoM
			0.83
			Nasal bone
			present
			Sonographer
			NA
			Qualifications in measuring NT
			Sonographer
Risks at sampling date			Trisomy 21
Age risk		1:889	<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 risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.</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>
Biochemical T21 risk		1:7638	
Combined trisomy 21 risk		<1:10000	
Trisomy 13/18 + NT		<1:10000	
<p><b>Trisomy 13/18 + NT</b></p> <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