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Patient data					
Name	Mrs. NILIMA PRAFUL AHERWAR		Patient ID	0372501150080	
Birth day	03-03-1998		Sample ID	A1541165	
Age at sample date	26.9		Sample Date	15-01-2025	
Gestational age	12 + 6				
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	60	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	12 + 6	
PAPP-A	3.67 mIU/mL	0.81	Method	CRL Robinson	
fb-hCG	87.57 ng/mL	2.27	Scan date	15-01-2025	
Risks at sampling date			Crown rump length in mm	67	
Age risk		1:877	Nuchal translucency MoM	0.59	
Biochemical T21 risk		1:479	Nasal bone	present	
Combined trisomy 21 risk		1:2820	Sonographer	N A	
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD	
Risk			Trisomy 21		
			<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 2820 women with the same data, there is one woman with a trisomy 21 pregnancy and 2819 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>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>					

Sign of Physician