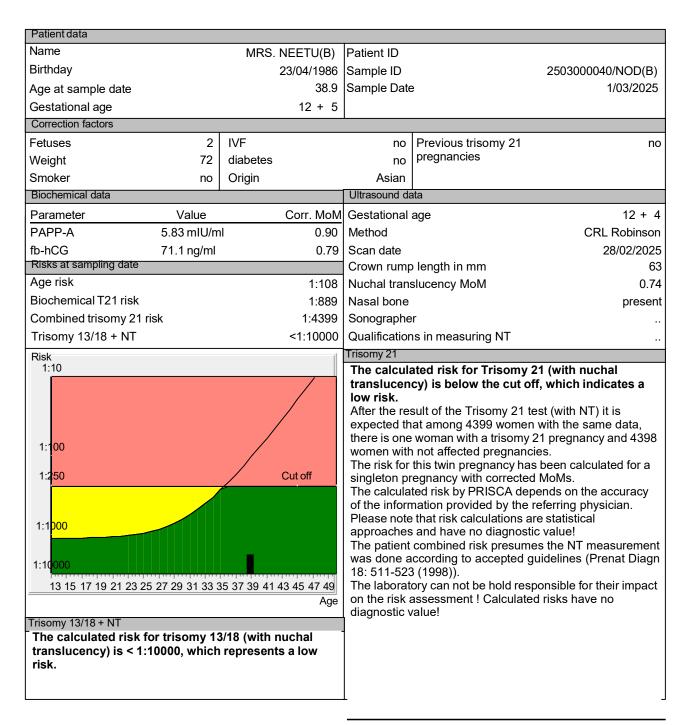
Prisca 5.2.0.13

Date of report: 2/03/2025

Patient data						
Name		MRS. NEETU(A)				
Birthday		23/04/1986				2503000040/NOD(A)
Age at sample date	38.9			Sample Date 1/03/2025		
Gestational age			13 + 1			
Correction factors						
Fetuses	2	IVF		no	Previous trisomy 21	no
Weight	72	diabetes		no	pregnancies	
Smoker	no	Origin		Asian		
Biochemical data				Ultrasound da	ta	
Parameter	Value		Corr. MoM	Gestational	age	13 + 0
PAPP-A	5.83 mIU/m	ıl	0.76	Method CRL Robinson		
fb-hCG	71.1 ng/ml		0.83	Scan date 28/02/2025		
Risks at sampling date			Crown rump length in mm 69			
Age risk 1:109			Nuchal translucency MoM 0.86			
Biochemical T21 risk 1:546			Nasal bone present			
Combined trisomy 21 risk 1:2769			3			
Trisomy 13/18 + NT <1:10000			Qualifications in measuring NT			
Risk 1:10 1:1000 1:1000 1:100000 1:100000 1:100000 1:100000 1:100000 1:100000 1:1000			Trisomy 21 The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. After the result of the Trisomy 21 test (with NT) it is expected that among 2769 women with the same data, there is one woman with a trisomy 21 pregnancy and 2768 women with not affected pregnancies. The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs. 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! The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)). The laboratory can not be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic value!			
The calculated risk translucency) is < ?						-

Prisca 5.2.0.13

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Sign of Physician