

N A

Patient data			
Name	Mrs. SANJIVANI FUKE TWIN B	Patient ID	0672503170218
Birthday	06-01-2004	Sample ID	B2547579
Age at sample date	21.2	Sample Date	17-03-2025
Gestational age	13 + 3		
Correction factors			
Fetuses	2	IVF	no
Weight	44	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	3.016 mIU/mL	0.20	Gestational age 12 + 5
fb-hCG	40.5 ng/mL	0.51	Method CRL Robinson
Risks at sampling date			
Age risk	1:1102		Scan date 12-03-2025
Biochemical T21 risk	1:347		Crown rump length in mm 66.6
Combined trisomy 21 risk	1:2001		Nuchal translucency MoM 0.95
Trisomy 13/18 + NT	1:767		Nasal bone present
Sonographer N A			
Qualifications in measuring NT MD			
Trisomy 21			
The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. <p>After the result of the Trisomy 21 test (with NT) it is expected that among 2001 women with the same data, there is one woman with a trisomy 21 pregnancy and 2000 women with not affected pregnancies.</p> <p>The PAPP-A level is low.</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.</p> <p>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 held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>			
Trisomy 13/18 + NT			
The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:767, which represents a low risk.			

Sign of Physician