

NA

Patient data			
Name	Mrs. ANKITA KHARE	Patient ID	0622509010059
Birthday	09/06/93	Sample ID	B3083057
Age at sample date	32.2	Sample Date	01/09/25
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	63	diabetes	no
Smoker	no	Origin	Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.4 mIU/mL	0.50	Gestational age 12 + 3
fb-hCG	35.16 ng/ml	0.98	Method CRL Robinson
Risks at sampling date			
Age risk	1:472		Scan date 29/08/25
Biochemical T21 risk	1:579		Crown rump length in mm 60.9
Combined trisomy 21 risk	1:88		Nuchal translucency MoM 1.86
Trisomy 13/18 + NT	1:856		Nasal bone present
Risk		Sonographer NA	
1:10		Qualifications in measuring NT MD	
1:100		Trisomy 21	
1:250		The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.	
1:1000		After the result of the Trisomy 21 test (with NT) it is expected that among 88 women with the same data, there is one woman with a trisomy 21 pregnancy and 87 women with not affected pregnancies.	
1:10000		The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.	
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49	Age	Please note that risk calculations are statistical approaches and have no diagnostic value!	
Trisomy 13/18 + NT		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 held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!	
The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:856, which represents a low risk.			

Sign of Physician

below cut off

Below Cut Off, but above Age Risk

above cut off