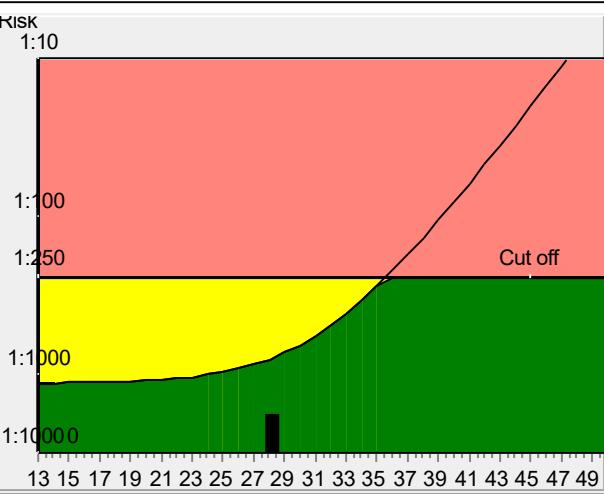


Prisca

5.1.0.17

Date of report: 31-08-2025

NA

Patient data			
Name	Mrs. NILIMA KANWAR	Patient ID	0622508300068
Birthday	04-06-1997	Sample ID	A1820336
Age at sample date	28.2	Sample Date	30-08-2025
Gestational age	13 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	64	diabetes	no
Smoker	no	Origin	Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.42 mIU/mL	0.60	Gestational age 13 + 1
fb-hCG	32 ng/ml	0.98	Method CRL Robinson
Risks at sampling date		Scan date 29-08-2025	
Age risk	1:799	Crown rump length in mm 71.3	
Biochemical T21 risk	1:1559	Nuchal translucency MoM 0.51	
Combined trisomy 21 risk	1:8846	Nasal bone present	
Trisomy 13/18 + NT	<1:10000	Sonographer NA	
Trisomy 21		Qualifications in measuring NT MD	
RISK 1:10 		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 8846 women with the same data, there is one woman with a trisomy 21 pregnancy and 8845 women with not affected pregnancies. 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 held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!	
Trisomy 13/18 + NT			
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.			

Sign of Physician

 below cut off

 Below Cut Off, but above Age Risk

 above cut off