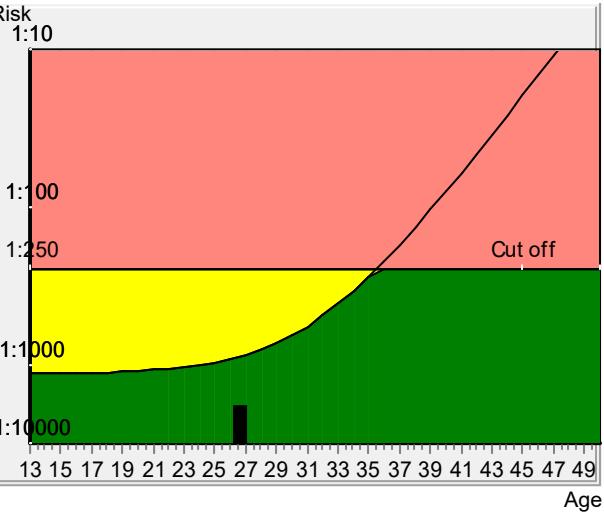


Prisca 5.1.0.17
Date of report: 25-10-2025

NA

Patient data			
Name	Mrs. PUTUL KUMARI W/O PARTH SA TWIN B	Patient ID	0012510250239
Birthday	06-03-1999	Sample ID	B3804920 TWIN B
Age at sample date	26.6	Sample Date	22-10-2025
Gestational age	12 + 6		
Correction factors			
Fetuses	2	IVF	no
Weight	68.3	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	4.94 mIU/mL	0.69	Gestational age 12 + 5
fb-hCG	172.93 ng/mL	2.16	Method CRL Robinson
Risks at sampling date			
Age risk	1:891		Scan date 21-10-2025
Biochemical T21 risk	1:375		Crown rump length in mm 64.9
Combined trisomy 21 risk	1:2291		Nuchal translucency MoM 0.72
Trisomy 13/18 + NT	<1:10000		Nasal bone present
Sonographer NA			
Qualifications in measuring NT NA			
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 2291 women with the same data, there is one woman with a trisomy 21 pregnancy and 2290 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 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