

Prisca 5.2.0.13
Date of report: 09/01/22

Patient data			
Name	PAPIYA DWEAN		Patient ID
Birthday	10/09/93		Sample ID
Age at sample date	28.3		Sample Date
Gestational age	11 + 3		
Correction factors			
Fetuses	1	IVF	unknown
Weight	54	diabetes	unknown
Smoker	unknown	Origin	Asian
Previous trisomy 21		unknown	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.03 mIU/mL	0.61	11 + 2
fb-hCG	36.84 ng/mL	0.76	Method
			CRL Robinson
			Scan date
			04/01/22
Risks at sampling date			
Age risk			Crown rump length in mm
1:741			47
Biochemical T21 risk			Nuchal translucency MoM
1:2665			0.81
Combined trisomy 21 risk			Nasal bone
<1:10000			unknown
Trisomy 13/18 + NT			Sonographer
<1:10000			DR.S.K.SARKAR.
			Qualifications in measuring NT
			MBBS.
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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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!			
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