

Prisca 5.2.0.13
Date of report: 03/02/22

Patient data			
Name	PRIANKA DAS NATH.		Patient ID
Birthday	23/07/92		Sample ID
Age at sample date	29.5		Sample Date
Gestational age	12 + 2		
Correction factors			
Fetuses	1	IVF	unknown
Weight	68	diabetes	unknown
Smoker	unknown	Origin	Asian
Previous trisomy 21		unknown	
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	5.25 mIU/mL	1.44	12 + 1
fb-hCG	7.18 ng/mL	0.20	Method
Risks at sampling date			CRL Robinson
Age risk	1:677		Scan date
Biochemical T21 risk	<1:10000		31/01/22
Combined trisomy 21 risk	<1:10000		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		58.3
			Nuchal translucency MoM
			1.19
			Nasal bone
			unknown
			Sonographer
			DR. PRIMAL MANDAL.
			Qualifications in measuring NT
			M.B.B.S.
Risk			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 free beta HCG level is low. 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