

SONIYA

Patient data			
Name	Mrs. KEERTHI	Patient ID	0012202260274
Birthday	21/11/06	Sample ID	23279869
Age at sample date	15.3	Sample Date	26/02/22
Gestational age	11 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	70	diabetes	no
Smoker	no	Origin	Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	3.26 mIU/mL	1.41	Gestational age
fb-hCG	52.18 ng/mL	1.09	Method
Risks at sampling date			
Age risk	1:1091		CRL Robinson
Biochemical T21 risk	<1:10000		Scan date
Combined trisomy 21 risk	<1:10000		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		Nuchal translucency MoM
RISK			
1:10			Nasal bone
1:100			Sonographer
1:250			Qualifications in measuring NT
1:1000			MD
1:10000			Trisomy 21
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			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 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