

Patient data			
Name	Mrs. S SANDHYA		Patient ID
0012302160517			
Birthday	28-02-1998		Sample ID
24077510			
Age at sample date	25.0		Sample Date
16-02-2023			
Gestational age	12 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	52	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		unknown	
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.94 mIU/mL	0.87	12 + 4
fb-hCG	47.87 ng/mL	1.07	Method
Risks at sampling date			CRL Robinson
Age risk	1:960		Scan date
Biochemical T21 risk	1:3863		17-02-2023
Combined trisomy 21 risk	<1:10000		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		64
			Nuchal translucency MoM
			0.97
			Nasal bone
			unknown
			Sonographer
			NA
			Qualifications in measuring NT
			MD
Risk			Trisomy 21
1:10			The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
1:100			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.
1:250			The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.
1:1000			Please note that risk calculations are statistical approaches and have no diagnostic value!
1:10000			The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).
Age			The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			
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