

Patient data			
Name	MRS. MANISHA	Patient ID	
Birthday	27-04-2001	Sample ID	2305060402/NOD
Age at sample date	22.1	Sample Date	30-05-2023
Gestational age		12 + 4	
Correction factors			
Fetuses	1	IVF	no
Weight	50	diabetes	no
Smoker		no	Origin
		Asian	
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.12 mIU/ml	0.62	12 + 3
fb-hCG	25.8 ng/ml	0.54	Method
Risks at sampling date			CRL Robinson
Age risk		1:1054	Scan date
Biochemical T21 risk		1:7727	29-05-2023
Combined trisomy 21 risk		<1:10000	Crown rump length in mm
Trisomy 13/18 + NT		<1:10000	60.8
Risk			Nuchal translucency MoM
1:10			0.82
1:100			Nasal bone
1:250			present
1:1000			Sonographer
1:10000			..
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			Qualifications in measuring NT
Age			MD
			Trisomy 21
			<b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b>
			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.**

	below risk		Below Cut Off, but above Age Risk		above cut off
--	------------	--	-----------------------------------	--	---------------

---

**Sign of Physician**

