

Patient data			
Name	MRS. MEENAKSHI	Patient ID	
Birthday	25/09/1979	Sample ID	2306220700/AMB
Age at sample date	43.8	Sample Date	27/06/2023
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	84	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.48 mIU/ml	0.73	Gestational age 12 + 4
fb-hCG	26.2 ng/ml	0.69	Method CRL Robinson
Risks at sampling date			Scan date 23/06/2023
Age risk		1:28	Crown rump length in mm 63
Biochemical T21 risk		1:189	Nuchal translucency MoM 1.23
Combined trisomy 21 risk		1:398	Nasal bone present
Trisomy 13/18 + NT		1:2924	Sonographer .
			Qualifications in measuring NT .
Risk			Trisomy 21
			<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 398 women with the same data, there is one woman with a trisomy 21 pregnancy and 397 women with not affected pregnancies.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>
Trisomy 13/18 + NT			
<p><b>The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:2924, which represents a low risk.</b></p>			

Sign of Physician

