

\*Free Home Sample Collection 9999 778 778

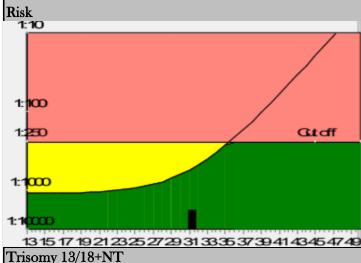


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Date of Report 31-12-2024 59019

					5.2.0.13	
Patient Data						
Name		MRS. KĄ	JAL RAI	Patient ID		12412300072
Birthday		0	7-09-1993	Sample ID		11860011
Age at Sample date		31.3		Sample Date		30-12-2024
Gestational age			12+2			
Correction factors						
Fetuses	1	IVF		unknown	Previous trisomy 21	unknown
Weight in kg	49	Diabetes		NO	Pregnancies	unknown
Smoker	NO	Origin		Asian		
Biochemical Data			Ultrasound Da	ata		

Diochemical Data			Cidasound Bata		
Parameter	Value	Corr Mom	Gestational age	12+2	
PAPP-A	4.5  mIU/ml	0.72	Method	CRL (<>Robinson)	
fb-hCG	18.7 ng/ml	0.44	Scan date	30-12-2024	
Risks at sampling date			Crown rump length in mm	59	
Age Risk		1:535	Nuchal translucency MoM	0.91	
Biochemical T21 risk		1:8648	Nasal bone	PRESENT	
Combined trisomy 21 risk		<1:10000	Sonographer	DR. DEEPIKA	
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD	
Risk			Down's Syndrome Risk (Trisomy 21 Screening)		
1:10					



## The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk

## The calculated risk for Trisomy 21(with NT) is below the cut off, which represents 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 the risk calculations are statistical aapproaches and have no diagnostic value!

The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523;

The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values