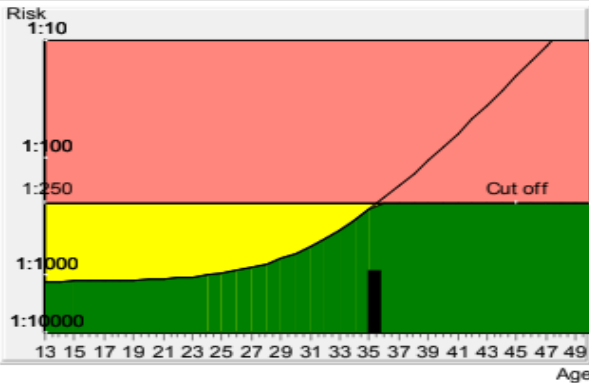



Date of Report 1/5/2022
PRISCA 5.1.0.17

Patient Data				
Name	MRS. MANITA DEVI	Patient ID	052204300038	
Birthday	27/12/1986	Sample ID	11459466	
Age at term	35.09	Sample Date	30/4/2022	
Gestational age	13+2			
Correction factors				
Fetuses	1 IVF	unknown	Previous trisomy 21	unknown
Weight in kg	71	Diabetes	NO	Pregnancies
Smoker	NO	Origin	Asian	unknown
Biochemical Data			Ultrasound Data	
Parameter	Value	Corr Mom	Gestational age	12+2
PAPP-A	3.4 mIU/ml	0.53	Method	CRL (<>Robinson)
fb-hCG	38.1 ng/ml	0.78	Scan date	23/4/2022
Risks at sampling date			Nuchal Translucency	2
Age Risk	1:260		Nuchal Translucency MoM	1.33
Biochemical T21 risk	1:609		Nasal Bone	Present
Combined T21 risk	1:935			
Trisomy 13/18+NT	1:6442			
Risk			Down's Syndrome Risk (Trisomy 21 Screening)	
			<p>The calculated risk for Trisomy 21 (with NT) is below the cut off, which represents a low risk.</p> <p>After the result of the Trisomy 21 test it is expected that among 935 women with the same data, there is one woman with a trisomy 21 pregnancy 934 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 the risk calculations are statistical approaches and have no diagnostic value!</p>	
Trisomy 13/18+NT			The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values	
The calculated risk for Trisomy 13/18 (with NT) is 1:6442 , which indicates a low risk				

 Risk Above Cut Off

 Risk above Age Risk

 Risk below Age risk