

Date of Report 09-11-2023
 PRISCA 5.2.0.13

Patient Data	
Name MRS AMRITA	Patient ID 012311080130
Birthday 07-02-1990	Sample ID 11854581
Age at Sample date 33.7	Sample Date 08-11-2023
Gestational age 12+4	

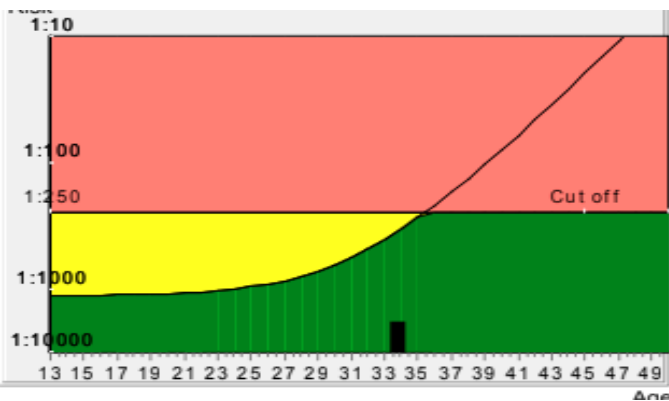
Correction factors			
Fetuses 1	IVF unknown	Previous trisomy 21 unknown	
Weight in kg 63	Diabetes NO	Pregnancies unknown	
Smoker NO	Origin Asian		

Biochemical Data	Ultrasound Data
------------------	-----------------

Parameter	Value	Corr Mom	
PAPP-A	3.87 mIU/ml	0.74	Method CRL (<>Robinson)
fb-hCG	56.7 ng/ml	1.56	Scan date 07-11-2023

Risks at sampling date	Ultrasound Data
Age Risk 1:353	Crown rump length in mm 60
Biochemical T21 risk 1:398	Nuchal translucency MoM 0.64
Combined trisomy 21 risk 1:2278	Nasal bone PRESENT
Trisomy 13/18 + NT <1:10000	Sonographer DR.INDERJEET
	Qualifications in measuring NT MD

Risk	Down's Syndrome Risk (Trisomy 21 Screening)
------	---



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 2278 women with the same data, there is one woman with a trisomy 21 pregnancy and 2277 women with not affected pregnancies.
 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!
 The patient combined risk presumes that NT measurement was done according to accepted guidelines (Prenat Diagn 18:511-523; 1998).

The calculated risk for Trisomy 13/18 (with NT) is <1:10000, which indicates a low risk
 The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values

Risk Above Cut Off
 Risk above Age Risk
 Risk below Age risk