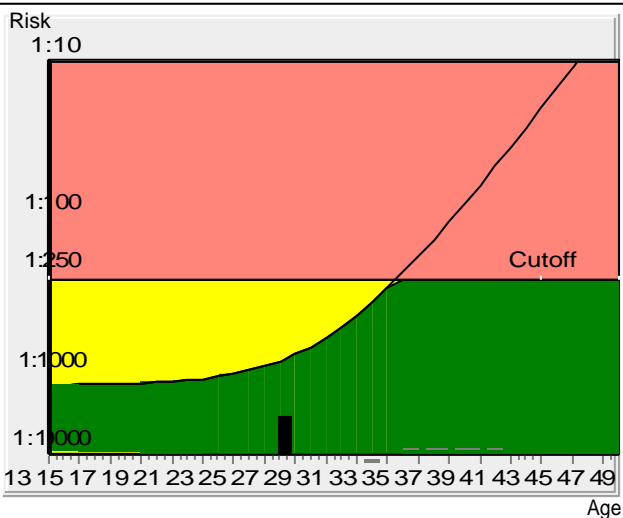


Patient data		
Name	MRS. RUBEL DHIMAN	Patient ID
Birthday	18-11-1989	Sample ID
Age at sample date	28.9	Sample Date
Gestational age	12 + 6	
Correction factors		
Fetuses	1 IVF	unknown Previous trisomy 21
Weight	55 diabetes	unknown pregnancies
Smoker	unknown Origin	Asian
Biochemical data		Ultrasound data
Parameter	Value	Corr. MoM
PAPP-A	2.8 mIU/ml	0.69
fb-hCG	35.1 ng/ml	1.17
Risks at sampling date		Gestational age
Age risk	1:790	12 + 6
Biochemical T21 risk	1:1460	Method
Combined trisomy 21 risk	1:2641	LMP
Trisomy 13/18 + NT	<1:10000	Scan date
		Crown rump length in mm
		Nuchal translucency MoM
		Nasal bone
		Sonographer
		Qualifications in measuring NT
		MD
		Trisomy 21
		The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
		After the result of the Trisomy 21 test (with NT) it is expected that among 2641 women with the same data, there is one woman with a trisomy 21 pregnancy and 2640 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 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.



Sign of Physician