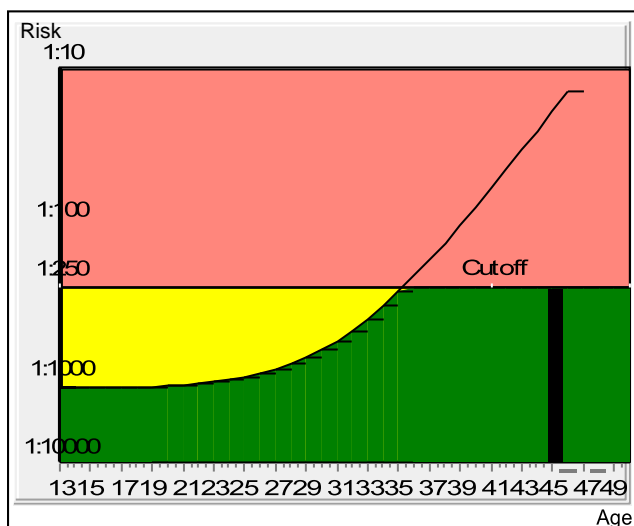


Patient data			
Name	MRS. NEELAM	Patient ID	1808220517/AMB
Birthday	01-07-1973	Sample ID	1808220517/AMB
Age at sample date	45.1	Sample Date	20-08-2018
Gestational age	11 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	67	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Risks at sampling date			
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	1.5 mIU/ml	0.79	11 + 1
fb-hCG	47.2 ng/ml	1.14	Method
			CRL Robinson
			Scan date
			18-08-2018
			Crown rump length in mm
			45.2
			Nuchal translucency MoM
			0.67
			Nasal bone
			present
			Sonographer
			DR. POONAM LOOMBA
			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 256 women with the same data, there is one woman with a trisomy 21 pregnancy and 255 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 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

