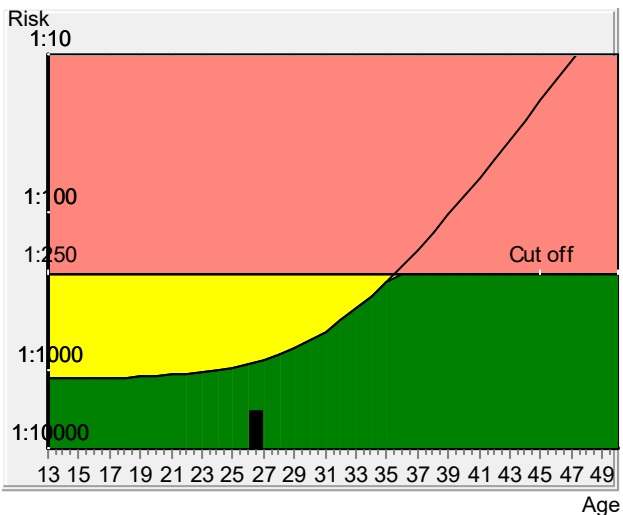


Prisca 5.1.0.17
 Date of report: 25/12/20

Patient data			
Name	Mrs. T.ANUSHA	Patient ID	0012012250027
Birthday	13/07/94	Sample ID	20121769
Age at sample date	26.5	Sample Date	25/12/20
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	37.3	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	2.98 mIU/mL	0.38	Gestational age 12 + 5
fb-hCG	45.36 ng/mL	0.98	Method CRL Robinson
Risks at sampling date			Scan date 24/12/20
Age risk		1:901	Crown rump length in mm 65.2
Biochemical T21 risk		1:530	Nuchal translucency MoM 0.90
Combined trisomy 21 risk		1:3030	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer NA
			Qualifications in measuring NT NA
Trisomy 13/18 + NT			Trisomy 21
<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 3030 women with the same data, there is one woman with a trisomy 21 pregnancy and 3029 women with not affected pregnancies.</p> <p>The PAPP-A level is low.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>



Sign of Physician