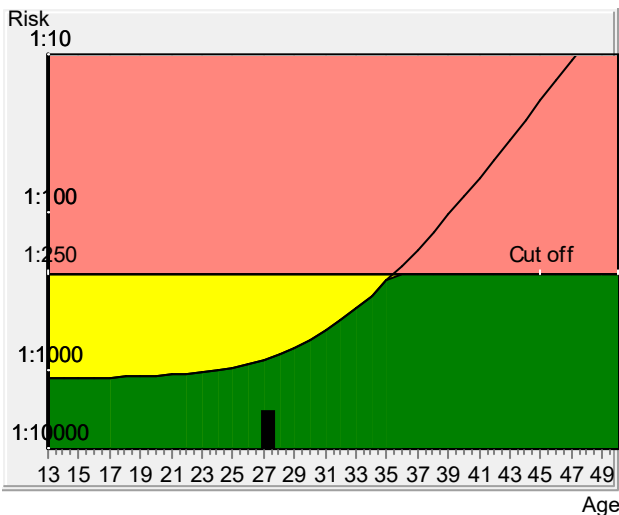


Patient data			
Name	Mrs. P ARUNA	Patient ID	0012306060276
Birth day	03-03-1996	Sample ID	24338558
Age at sample date	27.3	Sample Date	06-06-2023
Gestational age	12 + 5		
Correction factors			
Fetuses	1	IVF	no
Weight	41	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	5.27 mIU/mL	0.80	Gestational age 12 + 4
fb-hCG	45.65 ng/mL	0.99	Method CRL Robinson
Risks at sampling date			Scan date 05-06-2023
Age risk		1:849	Crown rump length in mm 64.1
Biochemical T21 risk		1:3244	Nuchal translucency MoM 0.55
Combined trisomy 21 risk		<1:10000	Nasal bone unknown
Trisomy 13/18 + NT		<1:10000	Sonographer NA
			Qualifications in measuring NT MD
Trisomy 21			
<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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</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>			
Trisomy 13/18 + NT			
<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			



Sign of Physician