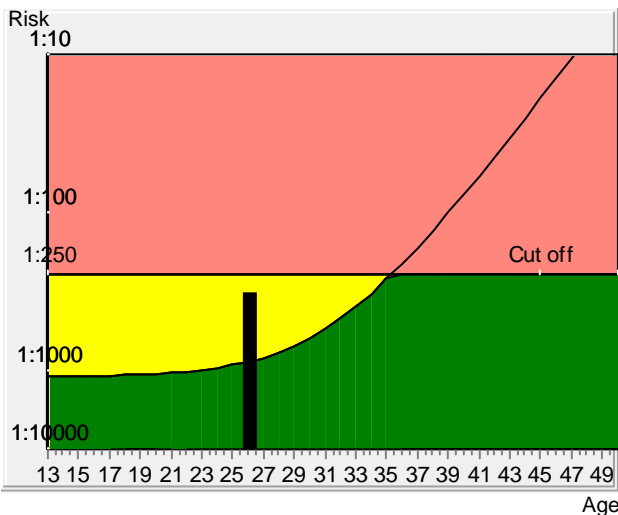


Prisca 5.2.0.13
Date of report: 23/02/2024

Patient data			
Name	MRS. SEEMA	Patient ID	
Birthday	1/01/1998	Sample ID	2402220595/AMB
Age at sample date	26.1	Sample Date	22/02/2024
Gestational age	11 + 6		
Correction factors			
Fetuses	1	IVF	yes
Weight	58	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.35 mIU/ml	0.44	Gestational age 11 + 5
fb-hCG	59.7 ng/ml	1.22	Method CRL Robinson
			Scan date 21/02/2024
Risks at sampling date		Trisomy 21	
Age risk	1:886	Crown rump length in mm	51.8
Biochemical T21 risk	1:473	Nuchal translucency MoM	1.51
Combined trisomy 21 risk	1:328	Nasal bone	present
Trisomy 13/18 + NT	1:6763	Sonographer	..
		Qualifications in measuring NT	..
Trisomy 13/18 + NT		Trisomy 21	
<p>The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:6763, 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 328 women with the same data, there is one woman with a trisomy 21 pregnancy and 327 women with not affected pregnancies.</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