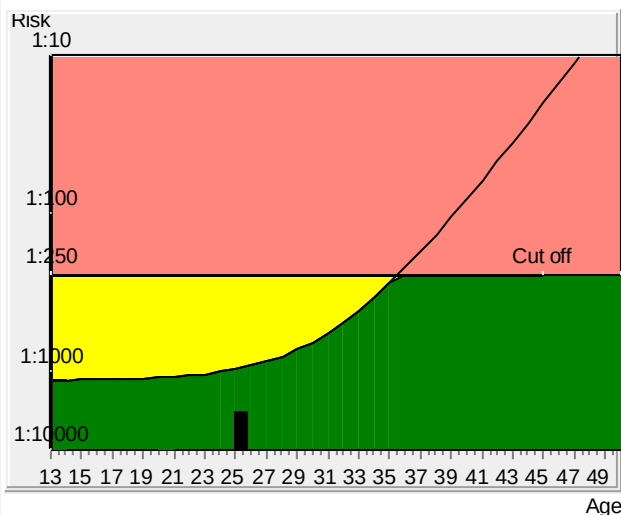


Patient data			
Name	MRS. SHIVANGI		Patient ID
Birthday	23/12/99	Sample ID	2504063338/NOD
Age at sample date	25.4	Sample Date	30/04/25
Gestational age	13 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	70.2	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies	no		
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	5.26 mIU/ml	1.17	Gestational age 12 + 5
fb-hCG	81.9 ng/ml	2.09	Method CRL Robinson
Risks at sampling date			Scan date 26/04/25
Age risk	1:970		Crown rump length in mm 66.7
Biochemical T21 risk	1:1412		Nuchal translucency MoM 0.83
Combined trisomy 21 risk	1:7358		Nasal bone present
Trisomy 13/18 + NT	<1:10000		Sonographer DR. RAJNEESH AGGARWAL
			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 7358 women with the same data, there is one woman with a trisomy 21 pregnancy and 7357 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