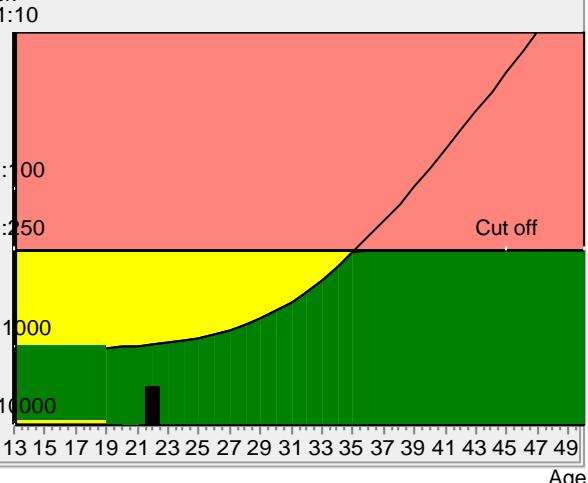


Date of report: 06-12-2022

Patient data			
Name	Mrs. NIDHI GAUR	Patient ID	0382212030038
Birthday	31-12-2000	Sample ID	24014373
Age at sample date	21.9	Sample Date	03-12-2022
Gestational age	10 + 5		
Correction factors			
Fetuses	1	IVF	no Previous trisomy 21 unknown
Weight	70	diabetes	no pregnancies
Smoker	no	Origin	Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	1.32 mIU/mL	0.95	Gestational age 10 + 5
fb-hCG	55.21 ng/mL	0.96	Method CRL Robinson
Risks at sampling date		Scan date 03-12-2022	
Age risk	1:985	Crown rump length in mm 40.04	
Biochemical T21 risk	1:6014	Nuchal translucency MoM 0.26	
Combined trisomy 21 risk	<1:10000	Nasal bone unknown	
Trisomy 13/18 + NT	<1:10000	Sonographer NA	
Qualifications in measuring NT NA		Trisomy 21	
RISK 1:10  1:100 1:250 1:1000 1:10000 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 Age		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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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 held 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

