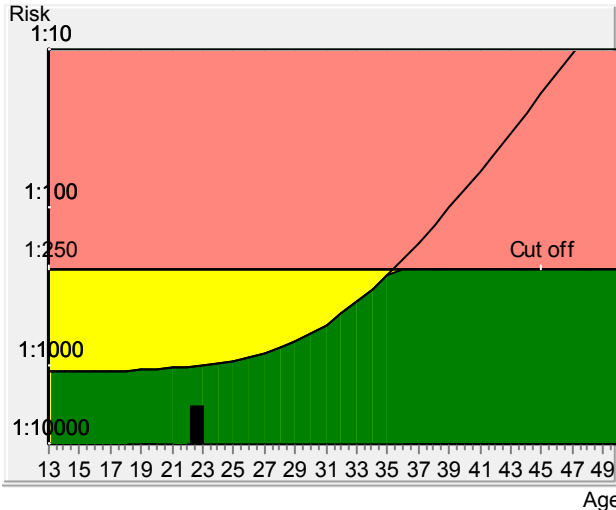


Patient data						
Name		MRS. BHAWNA SINGHAL		Patient ID	2104059995/DHJ	
Birthday		27-09-1998		Sample ID	2104059995/DHJ	
Age at sample date		22.6		Sample Date	27-04-2021	
Gestational age		12 + 0				
Correction factors						
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	no	
Weight	51	diabetes	no			
Smoker	no	Origin	Asian			
Biochemical data			Ultrasound data			
Parameter	Value	Corr. MoM	Gestational age			11 + 6
PAPP-A	4.1 mIU/ml	1.08	Method			CRL Robinson
fb-hCG	46.1 ng/ml	0.92	Scan date			26-04-2021
Risks at sampling date			Crown rump length in mm			54.1
Age risk		1:1022	Nuchal translucency MoM			0.84
Biochemical T21 risk		1:9062	Nasal bone			present
Combined trisomy 21 risk		<1:10000	Sonographer			..
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT			MD
Risk			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 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 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.						

 below cut off