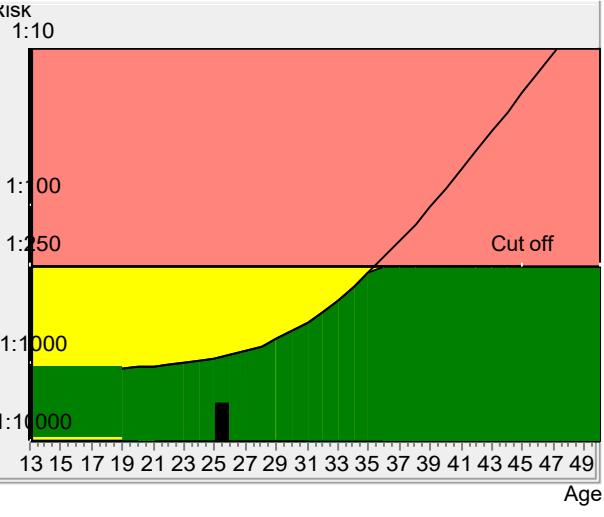


MA DYPATHI SHILPA

Patient data			
Name	Mrs. THRIBHUVANA	Patient ID	0012102220480
Birthday	01/09/95	Sample ID	21002089
Age at sample date	25.5	Sample Date	22/02/21
Gestational age	12 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	62.75	diabetes	no
Smoker	no	Origin	Asian
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.56 mIU/mL	1.34	Gestational age
fb-hCG	61.36 ng/mL	1.41	Method
Risks at sampling date		CRL Robinson	
Age risk	1:932	Scan date	22/02/21
Biochemical T21 risk	1:4735	Crown rump length in mm	59
Combined trisomy 21 risk	<1:10000	Nuchal translucency MoM	1.17
Trisomy 13/18 + NT	<1:10000	Nasal bone	present
		Sonographer	DR NA
		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.			
<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 held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>			
			
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

 below cut off

 Below Cut Off, but above Age Risk

 above cut off