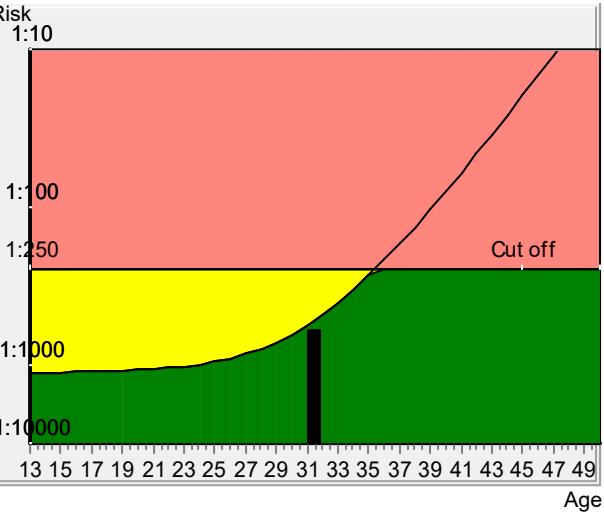


N A

Patient data								
Name		Mrs. A.HARIKA		Patient ID	0902509180018			
Birthday		04/05/94		Sample ID	A1977481			
Age at sample date		31.4		Sample Date	18/09/25			
Gestational age		12 + 3						
Correction factors								
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown			
Weight	61	diabetes	no					
Smoker	no	Origin	Asian					
Biochemical data								
Parameter	Value	Corr. MoM	Ultrasound data					
PAPP-A	1.41 mIU/mL	0.38	Gestational age					
fb-hCG	42.52 ng/mL	1.00	12 + 3					
Risks at sampling date								
Age risk		1:533	Method					
Biochemical T21 risk		1:286	CRL Robinson					
Combined trisomy 21 risk		1:603	Scan date					
Trisomy 13/18 + NT		1:6036	18/09/25					
Crown rump length in mm								
Nuchal translucency MoM								
Nasal bone								
Sonographer								
Qualifications in measuring NT								
MD								
Trisomy 21								
<b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b>								
After the result of the Trisomy 21 test (with NT) it is expected that among 603 women with the same data, there is one woman with a trisomy 21 pregnancy and 602 women with not affected pregnancies.								
The PAPP-A level is low.								
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!								
Risk								
								
<b>Trisomy 13/18 + NT</b>								
<b>The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:6036, which represents a low risk.</b>								

Sign of Physician

 below cut off

 Below Cut Off, but above Age Risk

 above cut off