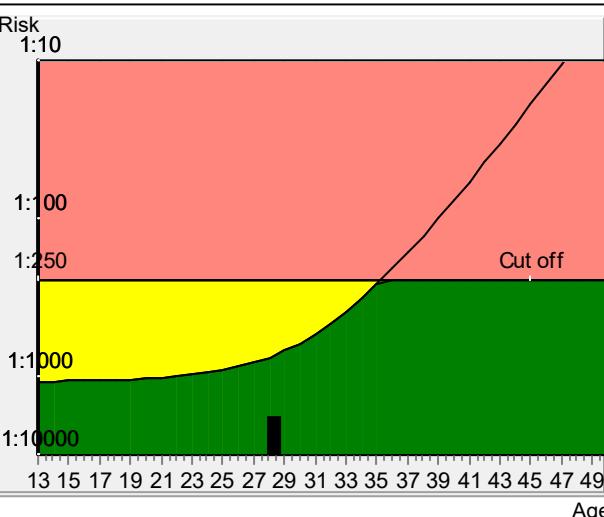


SAGEPATH LABS PVT LTD.

Prisca

5.1.0.17

Date of report: 19-01-2024

Patient data								
Name	W/O ANIL	Patient ID	0692401180133					
Birthday	12-10-1995	Sample ID	24968751					
Age at sample date	28.3	Sample Date	18-01-2024					
Gestational age	11 + 4							
Correction factors								
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown			
Weight	45	diabetes	no					
Smoker	no	Origin	Asian					
Biochemical data								
Parameter	Value	Corr. MoM	Ultrasound data					
PAPP-A	2.4 mIU/mL	0.66	Gestational age 11 + 3					
fb-hCG	49.5 ng/mL	0.87	Method CRL Robinson					
Risks at sampling date								
Age risk	1:749		Scan date 17-01-2024					
Biochemical T21 risk		1:2460	Crown rump length in mm 49					
Combined trisomy 21 risk		<1:10000	Nuchal translucency MoM 0.73					
Trisomy 13/18 + NT		<1:10000	Nasal bone unknown					
Sonographer N A								
Qualifications in measuring NT MD								
Trisomy 21								
<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <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								
<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>								

Sign of Physician

 below cut off

 Below Cut Off, but above Age Risk

 above cut off