

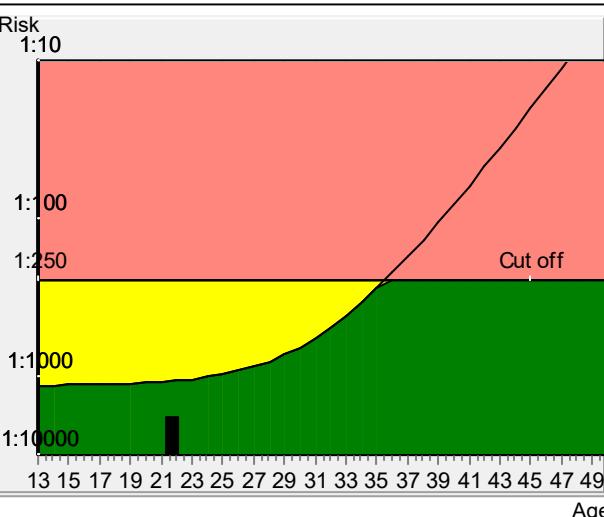
SAGEPATH LABS PVT LTD.

Prisca

5.1.0.17

Date of report: 15/02/24

R.UMA

Patient data			
Name	Mrs. CH MADAVI	Patient ID	0012402150085
Birthday	12/06/02	Sample ID	A0285801
Age at sample date	21.7	Sample Date	15/02/24
Gestational age	13 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	33	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	10.24 mIU/mL	1.05	Gestational age 13 + 0
fb-hCG	33.1 ng/mL	0.78	Method CRL Robinson
Risks at sampling date			
Age risk	1:1088		Scan date 13/02/24
Biochemical T21 risk	<1:10000		Crown rump length in mm 70
Combined trisomy 21 risk	<1:10000		Nuchal translucency MoM 0.63
Trisomy 13/18 + NT	<1:10000		Nasal bone present
			Sonographer R.UMA
			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