

SAGEPATH LABS PVT LTD..

Prisca

5.1.0.17

Date of report:

19-10-2023

Patient data			
Name	Mrs. RUKMINI		Patient ID
Birthday	10-05-1998		0012310190344
Age at sample date	25.4		Sample ID
Gestational age	13 + 0		23998811
			Sample Date
			19-10-2023
Correction factors			
Fetuses	1	IVF	no
Weight	62.4	diabetes	no
Smoker	no	Origin	Asian
			Previous trisomy 21 pregnancies
			unknown
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	8.32 mIU/mL	1.83	13 + 0
fb-hCG	35.65 ng/mL	0.97	Method
Risks at sampling date			CRL Robinson
Age risk	1:957		Scan date
Biochemical T21 risk	<1:10000		19-10-2023
Combined trisomy 21 risk	<1:10000		Crown rump length in mm
Trisomy 13/18 + NT	<1:10000		70.5
			Nuchal translucency MoM
			0.45
			Nasal bone
			unknown
			Sonographer
			NA
			Qualifications in measuring NT
			MD
Risk			Trisomy 21
1:10			The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.
1:100			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.
1:250			The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.
1:1000			Please note that risk calculations are statistical approaches and have no diagnostic value!
1:10000			The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).
Age			The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			
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

