SagePath Labs Pvt Ltd.,

Prisca 5.1.0.17

Date of report: 19/02/21

Patient data			
Name	Mrs. CHANDANA	Patient ID	0012102190277
Birthday	05/09/99	Sample ID	21018381
Age at sample date	21.5	Sample Date	19/02/21
Gestational age	13 + 0		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	62	diabetes	no		
Smoker	no	Origin	Asian		

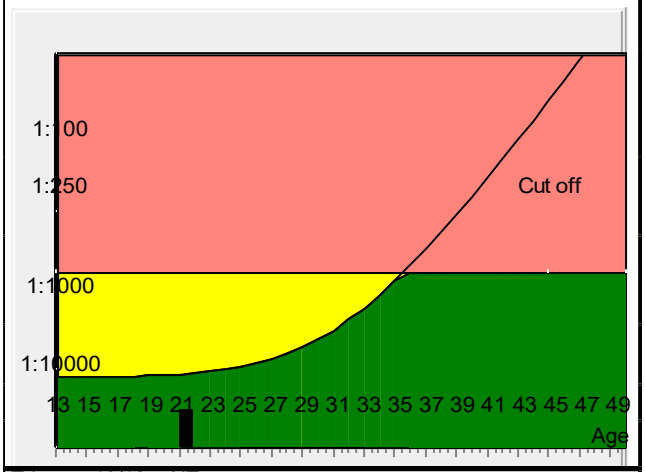
Biochemical data		
Parameter	Value	Corr. MoM
PAPP-A	6.78 mIU/mL	1.48
fb-hCG	43.62 ng/mL	1.19

Risks at sampling date	
Age risk	1:1083
Biochemical T21 risk	1:9946
Combined trisomy 21 risk	<1:10000
Trisomy 13/18 + NT	<1:10000

Ultrasound data	
Gestational age	13 + 0
Method	CRL Robinson
Scan date	19/02/21
Crown rump length in mm	69
Nuchal translucency MoM	0.71
Nasal bone	present
Sonographer	DR NA
Qualifications in measuring NT	MD

Risk

1:10



Trisomy 13/18 + NT

The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.

Trisomy 21

The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.

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.

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 hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!

Sign of Physician

below cut off | Below Cut Off, but above Age Risk | above cut off