Prisca   5.1.0.17

Date of report: **22/03/22**

TRIVENI MS(O,B,G)

Patient data			
Name	Mrs. SOUMYA W/O. RAMESH	Patient ID	0312203210101
Birthday	03/03/98	Sample ID	23284128
Age at sample date	24.0	Sample Date	21/03/22
Gestational age	11 + 4		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	43	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	11 + 4
PAPP-A	1.98 mIU/mL	0.52	Method	CRL Robinson
fb-hCG	57.63 ng/mL	0.99	Scan date	21/03/22
Risks at sampling date			Crown rump length in mm	51
Age risk		1:963	Nuchal translucency MoM	0.95
Biochemical T21 risk		1:1266	Nasal bone	present
Combined trisomy 21 risk		1:6225	Sonographer	DR NA
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD

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 6225 women with the same data, there is one woman with a trisomy 21 pregnancy and 6224 women with not affected pregnancies.

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!

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

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