Prisca	5.0.2.37
Date of report:	**18-07-2020**

Patient data			
Name	MRS. NEETU	Patient ID	2007220476/AMB
Birthday	07-01-2000	Sample ID	2007220476/AMB
Age at delivery	21.0	Sample Date	17-07-2020
Gestational age	12 + 5		

Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregancies	no
Weight	57.4	diabetes	no		
Smoker	no	Origin	Asian		

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 2
PAPP-A	1.1 mIU/ml	0.25	Method	CRL Robinson
fb-hCG	28.5 ng/ml	0.69	Scan date	14-07-2020
Risks at term			Crown rump length in mm	60.6
Age risk		1:1517	Nuchal translucency MoM	1.08
Biochemical T21 risk		1:520	Nasal bone	present
Combined trisomy 21 risk		1:2068	Sonographer	.
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 2068 women with the same data, there is one woman with a trisomy 21 pregnancy and 2067 women with not affected pregnancies.
The PAPP-A level is low.
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.

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Sign of Physician

🟩 below cut off	🟨 Below Cut Off, but above Age Risk	🟥 above cut off