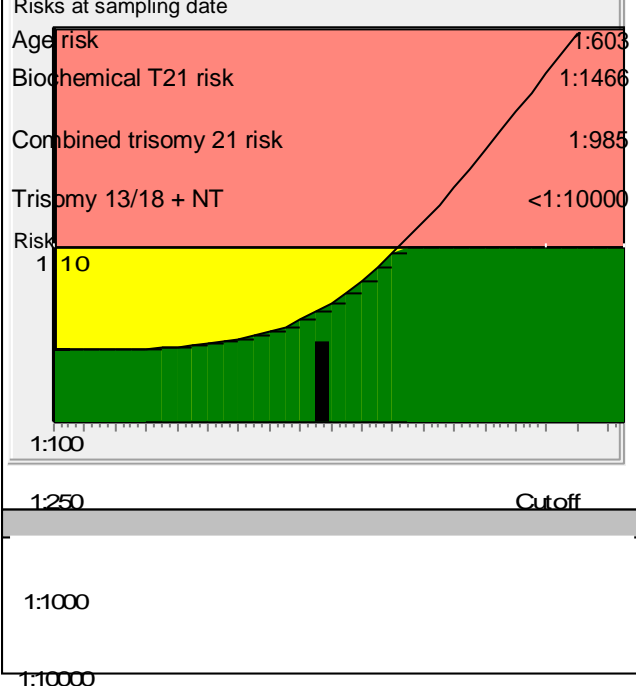


Prisca 5.0.2.37  
 Date of report: 08-11-2017

Patient data			
Name	MRS. SONALI	Patient ID	1711220123/AMB
Birthday	23-05-1987	Sample ID	1711220123/AMB
Age at sample date	30.5	Sample Date	07-11-2017
Gestational age	12 + 2		
Correction factors			
Fetuses	1 IVF	no	Previous trisomy 21 no
Weight	83 diabetes		no pregnancies
Smoker	no Origin		Asian
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age 12 + 0
PAPP-A	2.2 mIU/ml	1.04	Method CRL Robinson
fb-hCG	51.8 ng/ml	1.55	Scan date 05-11-2017
Risks at sampling date			Crown rump length in mm 55.3
Age risk		1:603	Nuchal translucency MoM 1.50
Biochemical T21 risk		1:1466	Nasal bone present
Combined trisomy 21 risk		1:985	Sonographer .
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT MD
Risk			Trisomy 21
1:10			<b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b>
1:100			After the result of the Trisomy 21 test (with NT) it is expected that among 985 women with the same data, there is one woman with a trisomy 21 pregnancy and 984 women with not affected pregnancies.
1:250			The laboratory 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)).
			The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!



1315 1719 212325 2729 313335 3739 414345 4749  
 Age



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	<input type="checkbox"/> above cut off
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