

Prisca 5.0.2.37
Date of report: 27-11-2017

Patient data			
Name	MRS. PARNEET	Patient ID	
Birthday	26-04-1986	Sample ID	1711220069/AMB
Age at sample date	32.0	Sample Date	27-11-2017
Gestational age	13 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	76	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies		no	
Biochemical data			Ultrasound data
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	4.56 mIU/ml	0.80	13 + 1
fb-hCG	23.4 ng/ml	0.86	Method
Risks at sampling date			CRL Robinson
Age risk		1:717	Scan date
Biochemical T21 risk		1:6905	27-11-2017
			Crown rump length in mm
			Nuchal translucency MoM
			Nasal bone
			present
Combined risk			
Trisomy 13/18 + NT		<1:10000	
Risk	1:10		
	1:100		
	1:250		
	1:1000		
	1:10000		
	Cutoff		
	Age		
	13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49		
Trisomy 13/18 + NT			

Sonographer .
Qualifications in measuring NT MD
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 2283 women with the same data, there is one woman with a trisomy 21 pregnancy and 2282 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!

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
---------------	--	-----------------------------------	---------------