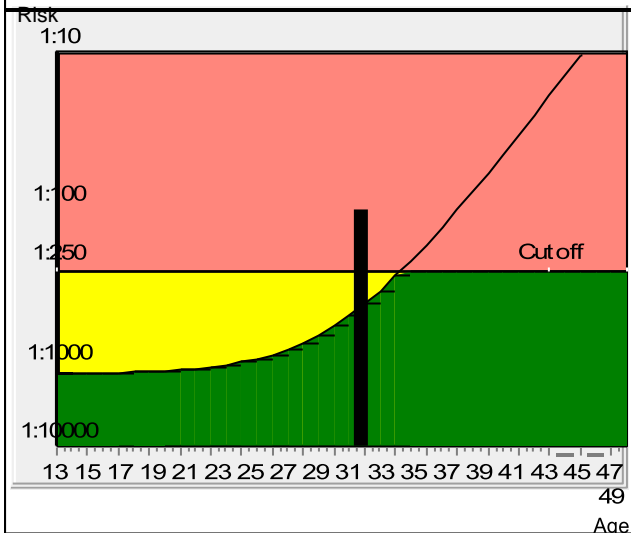


Prisca 5.0.2.37
Date of report: 29-05-2018

Patient data			
Name	MRS. POOJA	Patient ID	
Birthday	12-09-1985	Sample ID	1805220795/AMB
Age at sample date	32.7	Sample Date	29-05-2018
Gestational age	11 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	61	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown

Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	11 + 1
PAPP-A	4.8 mIU/ml	1.86	Method	CRL Robinson
fb-hCG	170.1 ng/ml	4.31	Scan date	24-05-2018
Risks at sampling date			Crown rump length in mm	46
Age risk		1:418	Nuchal translucency MoM	1.58
Biochemical T21 risk		1:245	Nasal bone	unknown
Combined trisomy 21 risk		1:100	Sonographer	.
Trisomy 13/18 + NT		<1:10000	Qualifications in measuring NT	MD



Trisomy 21
The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.
After the result of the Trisomy 21 test (with NT) it is expected that among 100 women with the same data, there is one woman with a trisomy 21 pregnancy and 99 women with not affected pregnancies.
The free beta HCG level is high.
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

Sign of Physician