

Prisca 5.0.2.37  
Date of report: 22-11-2018

Patient data			
Name	MRS. JASPREET	Patient ID	1710220378/AMB
Birthday	16-04-1995	Sample ID	1710220378/AMB
Age at sample date	23.2	Sample Date	21-11-2018
Gestational age	13 + 4		
Correction factors			
Fetuses	1	IVF	no
Weight	59.2	diabetes	no
Smoker	no	Origin	Asian
Previous trisomy 21 pregnancies			no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	4.1 mIU/ml	0.74	Gestational age 13 + 4
fb-hCG	58.2 ng/ml	2.08	Method CRL Robinson
Risks at sampling date			Scan date 21-11-2018
Age risk		1:816	Crown rump length in mm 74
Biochemical T21 risk		1:456	Nuchal translucency MoM 0.83
Combined trisomy 21 risk		1:2641	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer .
			Qualifications in measuring NT MD
<p>Risk</p> <p>1:10 1:100 1:250 1:1000 1:10000</p> <p>13151719212325272931333537394143454749</p> <p>Age</p>		<p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 2641 women with the same data, there is one woman with a trisomy 21 pregnancy and 2640 women with not affected pregnancies.</p> <p>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!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>	
<p>Trisomy 13/18 + NT</p> <p><b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b></p>			

Sign of Physician

