

Prisca 5.0.2.37
Date of report: 06-09-2017

Patient data			
Name	MRS. NARINDER	Patient ID	1709220097/AMB
Birthday	17-06-1982	Sample ID	1709220097/AMB
Age at sample date	35.2	Sample Date	05-09-2017
Gestational age	13 + 3		
Correction factors			
Fetuses	1	IVF	yes
Weight	75	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	1.12	Gestational age 12 + 0
fb-hCG	140 ng/ml	4.97	Method CRL Robinson
Risks at sampling date			Scan date 26-08-2017
Age risk		1:268	Crown rump length in mm 55.9
Biochemical T21 risk		1:63	Nuchal translucency MoM 0.68
Combined trisomy 21 risk		1:368	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer DR. H.S. MANN
			Qualifications in measuring NT MD
Risk		Trisomy 21	
		<p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 368 women with the same data, there is one woman with a trisomy 21 pregnancy and 367 women with not affected pregnancies.</p> <p>The free beta HCG level is high.</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>	
Trisomy 13/18 + NT			
<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p>			

Sign of Physician

below cut off	Below Cut Off, but above Age Risk	above cut off
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