

Patient data			
Name	MRS. SUFIA W/O SARFRAZ (A)	Patient ID	1709220048/AMB (A)
Birthday	16-07-1975	Sample ID	1709220048/AMB (A)
Age at sample date	42.1	Sample Date	04-09-2017
Gestational age	12 + 4		
Correction factors			
Fetuses	2	IVF	yes
Weight	70	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	20.5 mIU/ml	3.76	Gestational age 11 + 3
fb-hCG	190 ng/ml	2.63	Method CRL Robinson
Risks at sampling date			Scan date 27-08-2017
Age risk		1:44	Crown rump length in mm 48.7
Biochemical T21 risk		1:92	Nuchal translucency MoM 0.91
Combined trisomy 21 risk		1:396	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer DR. SAKSHI SHARMA
			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 396 women with the same data, there is one woman with a trisomy 21 pregnancy and 395 women with not affected pregnancies. The free beta HCG level is high. The PAPP-A level is high. The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs. 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!</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