

Prisca 5.1.0.17
Date of report: 27-12-2019

KOS DIAGNOSTIC LAB

Patient data			
Name	MRS. LALSA (B)	Patient ID	
Birthday	01-01-1990	Sample ID	1912220940/AMB
Age at delivery	30.5	Sample Date	24-12-2019
Gestational age	12 + 0		
Correction factors			
Fetuses	2	IVF	yes
Weight	55.9	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	5.94 mIU/ml	1.05	
fb-hCG	51.1 ng/ml	0.60	
Risks at term			
Age risk		1:908	
Biochemical T21 risk		<1:10000	
Combined trisomy 21 risk		<1:10000	
Trisomy 13/18 + NT		<1:10000	
		Gestational age	11 + 6
		Method	CRL Robinson
		Scan date	23-12-2019
		Crown rump length in mm	54.7
		Nuchal translucency MoM	0.83
		Nasal bone	present
		Sonographer	DR. SAKSHI SHARMA
		Qualifications in measuring NT	MD
Risk 1:10		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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</p> <p>The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.</p> <p>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.</p> <p>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

