

KOS DIAGNOSTIC LAB

Patient data			
Name	MRS. POONAM	Patient ID	
Birthday	27-11-1995	Sample ID	1909220317/AMB
Age at delivery	24.3	Sample Date	06-09-2019
Gestational age	13 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	55	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	5.8 mIU/ml	1.27	Gestational age 12 + 4
fb-hCG	90.1 ng/ml	2.69	Method CRL Robinson
Risks at term			Scan date 03-09-2019
Age risk		1:1414	Crown rump length in mm 63.18
Biochemical T21 risk		1:1253	Nuchal translucency MoM 0.70
Combined trisomy 21 risk		1:6732	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer .
			Qualifications in measuring NT MD
Risk	1:10		Trisomy 21
			<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 6732 women with the same data, there is one woman with a trisomy 21 pregnancy and 6731 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><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

