

KOS DIAGNOSTIC LAB

Patient data			
Name	MRS. RAJINDER	Patient ID	
Birthday	15-09-1993	Sample ID	2005220220/AMB
Age at delivery	27.2	Sample Date	19-05-2020
Gestational age	13 + 3		
Correction factors			
Fetuses	1	IVF	no
Weight	53	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	6.25 mIU/ml	1.13	Gestational age 11 + 0
fb-hCG	184 ng/ml	5.85	Method CRL Robinson
Risks at term			Scan date 02-05-2020
Age risk		1:1234	Crown rump length in mm 44.2
Biochemical T21 risk		1:295	Nuchal translucency MoM 0.73
Combined trisomy 21 risk		1:1719	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer .
			Qualifications in measuring NT MD
Risk		Trisomy 21	
<p>Risk 1:10</p>		<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 1719 women with the same data, there is one woman with a trisomy 21 pregnancy and 1718 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

