

KOS DIAGNOSTIC LAB

6349/1 NICHOLSON ROAD, AMBALA CANTT

Prisca

5.1.0.17

Date of report:

15-07-2022

Patient data			
Name	MRS. PREETI SAIN	Patient ID	2207220415/AMB
Birthday	07-10-1991	Sample ID	2207220415/AMB
Age at sample date	30.8	Sample Date	14-07-2022
Gestational age	12 + 6		
Correction factors			
Fetuses	1	IVF	no
Weight	64	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	12.6 mIU/ml	2.96	Gestational age 12 + 3
fb-hCG	147 ng/ml	3.48	Method CRL Robinson
Risks at sampling date			Scan date 12-07-2022
Age risk		1:591	Crown rump length in mm 59.73
Biochemical T21 risk		1:578	Nuchal translucency MoM 0.80
Combined trisomy 21 risk		1:2848	Nasal bone present
Trisomy 13/18 + NT		<1:10000	Sonographer .
			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 2848 women with the same data, there is one woman with a trisomy 21 pregnancy and 2847 women with not affected pregnancies.</p> <p>The free beta HCG level is high.</p> <p>The PAPP-A level is high.</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