

KOS DIAGNOSTIC LAB

6349/1, NICHOLSON ROAD,AMBALA CANTT

Prisca

5.1.0.17

Date of report: 19-02-2022

Patient data			
Name	MRS. NEHA (A)	Patient ID	2202220425/AMB (A)
Birthday	01-07-1996	Sample ID	2202220425/AMB (A)
Age at sample date	25.6	Sample Date	17-02-2022
Gestational age	11 + 0		
Correction factors			
Fetuses	2	IVF	yes
Weight	66	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	7.8 mIU/ml	2.53	11 + 1
fb-hCG	123.5 ng/ml	1.10	Method
			CRL Robinson
			Scan date
			18-02-2022
Risks at sampling date			Crown rump length in mm
Age risk		1:880	45.6
Biochemical T21 risk		<1:10000	Nuchal translucency MoM
Combined trisomy 21 risk		<1:10000	1.20
Trisomy 13/18 + NT		<1:10000	Nasal bone
			present
			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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</p> <p>The PAPP-A level is high.</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