

Prisca

5.1.0.17

Date of report:

06-06-2022

Patient data			
Name	MRS. NEHA	Patient ID	2206220111/AMB
Birth day	08-01-1998	Sample ID	2206220111/AMB
Age at sample date	24.4	Sample Date	04-06-2022
Gestational age	12 + 2		
Correction factors			
Fetuses	1	IVF	no
Weight	40.5	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	
PAPP-A	5.32 mIU/ml	0.94	Gestational age 12 + 0
fb-hCG	294 ng/ml	5.51	Method CRL Robinson
			Scan date 02-06-2022
Risks at sampling date		Crown rump length in mm 56	
Age risk	1:977	Nuchal translucency MoM	1.29
Biochemical T21 risk	1:161	Nasal bone	present
Combined trisomy 21 risk	1:243	Sonographer	.
Trisomy 13/18 + NT	<1:10000	Qualifications in measuring NT	MD
Trisomy 21			
		<p>The calculated risk for Trisomy 21 (with nuchal translucency) is above the cut off, which indicates an increased risk.</p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 243 women with the same data, there is one woman with a trisomy 21 pregnancy and 242 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.</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 hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>	
		<p>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</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

 below cut off	 Below Cut Off, but above Age Risk	 above cut off
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