

Prisca

5.1.0.17

Date of report:

01-06-2023

Patient data			
Name	MRS. KAVITA	Patient ID	
Birth day	03-08-2000	Sample ID	2305062230/NOD
Age at sample date	22.8	Sample Date	31-05-2023
Gestational age	12 + 0		
Correction factors			
Fetuses	1	IVF	no
Weight	58	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	no
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	2.66 mIU/ml	0.81	11 + 6
fb-hCG	81.6 ng/ml	1.69	Method
			CRL Robinson
			Scan date
			30-05-2023
Risks at sampling date			Crown rump length in mm
Age risk	1:1016		55
Biochemical T21 risk	1:1152		Nuchal translucency MoM
Combined trisomy 21 risk	1:5915		0.89
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 5915 women with the same data, there is one woman with a trisomy 21 pregnancy and 5914 women with not affected pregnancies.</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