

Prisca

5.1.0.17

Date of report:

23-11-2024

NA

Patient data					
Name	Mrs. POOJITHA		Patient ID	0352411220040	
Birthday	23-08-1996		Sample ID	A1811212	
Age at sample date	28.2		Sample Date	22-11-2024	
Gestational age	13 + 2				
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown
Weight	67	diabetes	no		
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	13 + 2	
PAPP-A	3.2 mIU/mL	0.59	Method	CRL Robinson	
fb-hCG	32 ng/ml	1.00	Scan date	22-11-2024	
Risks at sampling date			Crown rump length in mm	74	
Age risk	1:799		Nuchal translucency MoM	0.71	
Biochemical T21 risk	1:1467		Nasal bone	present	
Combined trisomy 21 risk	1:8361		Sonographer	NA	
Trisomy 13/18 + NT	<1:10000		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 8361 women with the same data, there is one woman with a trisomy 21 pregnancy and 8360 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

below cut off

Below Cut Off, but above Age Risk

above cut off