

Prisca

5.1.0.17

Date of report:

12/02/21

SOWMYA REDDY

Patient data			
Name	Mrs. N. SANDHYA		Patient ID
Birthday	13/06/94	Sample ID	21007792
Age at sample date	26.7	Sample Date	10/02/21
Gestational age	11 + 1		
Correction factors			
Fetuses	1	IVF	no
Weight	68	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Biochemical data		Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age
PAPP-A	3.67 mIU/mL	2.02	11 + 1
fb-hCG	63.41 ng/mL	1.18	Method
			CRL Robinson
			Scan date
			10/02/21
			Crown rump length in mm
			45.3
			Nuchal translucency MoM
			0.80
			Nasal bone
			present
			Sonographer
			NA
			Qualifications in measuring NT
			NA
Risks at sampling date			
Age risk	1:833		
Biochemical T21 risk	<1:10000		
Combined trisomy 21 risk	<1:10000		
Trisomy 13/18 + NT	<1:10000		
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 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 held 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