

Prisca 5.1.0.17
Date of report: 30-05-2024

Patient data				
Name	Mrs. PALLAVI SONKAMBLE		Patient ID	0672405290160
Birthday	23-06-2000		Sample ID	A0761355
Age at sample date	23.9		Sample Date	29-05-2024
Gestational age	11 + 3			
Correction factors				
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies
Weight	40	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	11 + 1
PAPP-A	4.31 mIU/mL	1.11	Method	CRL Robinson
fb-hCG	53.98 ng/mL	0.88	Scan date	27-05-2024
Risks at sampling date			Crown rump length in mm	45
Age risk	1:962		Nuchal translucency MoM	0.89
Biochemical T21 risk	<1:10000		Nasal bone	present
Combined trisomy 21 risk	<1:10000		Sonographer	N A
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 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 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