

N A

Patient data					
Name	W/O PRAVEE KUMAR PAL		Patient ID	0692411010052	
Birth day	08-07-1994		Sample ID	A1653824	
Age at sample date	30.3		Sample Date	31-10-2024	
Gestational age	12 + 6				
Correction factors					
Fetuses	1	IVF	no	Previous trisomy 21	unknown
Weight	52	diabetes	no	pregnancies	
Smoker	no	Origin	Asian		
Biochemical data			Ultrasound data		
Parameter	Value	Corr. MoM	Gestational age	12 + 4	
PAPP-A	8.01 mIU/mL	1.50	Method	CRL Robinson	
fb-hCG	37.07 ng/mL	0.92	Scan date	29-10-2024	
Risks at sampling date			Crown rump length in mm	63	
Age risk	1:628		Nuchal translucency MoM	0.80	
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