

Prisca 5.1.0.17
Date of report: 18/01/26

Patient data				
Name	Mrs. JYOTI MASANE TWIN A		Patient ID	0662601160025
Birthday	16/01/97		Sample ID	B4187912 TWIN A
Age at sample date	29.0		Sample Date	16/01/26
Gestational age	13 + 0			
Correction factors				
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies
Weight	64	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 5
PAPP-A	2.47 mIU/mL	0.30	Method	CRL Robinson
fb-hCG	37 ng/mL	0.47	Scan date	14/01/26
Risks at sampling date			Crown rump length in mm	
Age risk	1:736		66	
Biochemical T21 risk	1:939		Nuchal translucency MoM	
Combined trisomy 21 risk	1:5755		0.65	
Trisomy 13/18 + NT	1:3614		Nasal bone	
			present	
			Sonographer	
			NA	
			Qualifications in measuring NT	
			Sonographer	
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 5755 women with the same data, there is one woman with a trisomy 21 pregnancy and 5754 women with not affected pregnancies.</p> <p>The PAPP-A level is low.</p> <p>The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.</p> <p>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.</p> <p>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 1:3614, which represents a low risk.</p>				

Sign of Physician

below cut off
 Below Cut Off, but above Age Risk
 above cut off

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Patient data				
Name	Mrs. JYOTI MASANE TWIN B		Patient ID	0662601160025
Birthday	16/01/97		Sample ID	B4187912 TWIN B
Age at sample date	29.0		Sample Date	16/01/26
Gestational age	12 + 6			
Correction factors				
Fetuses	2	IVF	no	Previous trisomy 21 pregnancies
Weight	63	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	12 + 4
PAPP-A	2.47 mIU/mL	0.31	Method	CRL Robinson
fb-hCG	37 ng/mL	0.45	Scan date	14/01/26
Risks at sampling date			Crown rump length in mm	
Age risk	1:732		64	
Biochemical T21 risk	1:1126		Nuchal translucency MoM	
Combined trisomy 21 risk	1:6791		0.73	
Trisomy 13/18 + NT	1:3814		Nasal bone	
			present	
			Sonographer	
			NA	
			Qualifications in measuring NT	
			Sonographer	
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 6791 women with the same data, there is one woman with a trisomy 21 pregnancy and 6790 women with not affected pregnancies.</p> <p>The PAPP-A level is low.</p> <p>The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.</p> <p>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.</p> <p>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 1:3814, which represents a low risk.</p>				

Sign of Physician