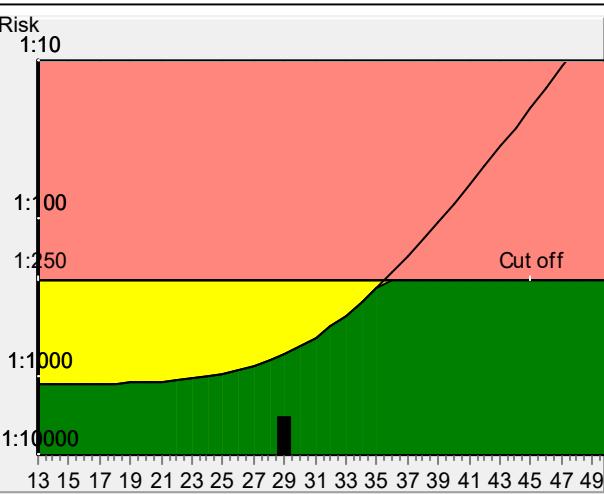


Prisca 5.1.0.17
Date of report: 18/01/26

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
Weight	64	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	2.47 mIU/mL	0.30	Gestational age 12 + 5
fb-hCG	37 ng/mL	0.47	Method CRL Robinson
Risks at sampling date			
Age risk	1:736		Scan date 14/01/26
Biochemical T21 risk	1:939		Crown rump length in mm 66
Combined trisomy 21 risk	1:5755		Nuchal translucency MoM 0.65
Trisomy 13/18 + NT	1:3614		Nasal bone present
		Sonographer NA	Sonographer NA
		Qualifications in measuring NT	Sonographer
Trisomy 21			
The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. <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 held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p>			
 <p>Trisomy 13/18 + NT</p> <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

Prisca 5.1.0.17
Date of report: 18/01/26

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
Weight	63	diabetes	no
Smoker	no	Origin	Asian
Biochemical data			
Parameter	Value	Corr. MoM	Ultrasound data
PAPP-A	2.47 mIU/mL	0.31	Gestational age 12 + 4
fb-hCG	37 ng/mL	0.45	Method CRL Robinson
Risks at sampling date			
Age risk	1:732		Scan date 14/01/26
Biochemical T21 risk	1:1126		Crown rump length in mm 64
Combined trisomy 21 risk	1:6791		Nuchal translucency MoM 0.73
Trisomy 13/18 + NT	1:3814		Nasal bone present
		Sonographer NA	Sonographer NA
		Qualifications in measuring NT	Sonographer
Trisomy 21			
The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.			
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.			
The PAPP-A level is low.			
The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.			
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!			
The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).			
The laboratory can not be held responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!			
Trisomy 13/18 + NT			
The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:3814, which represents a low risk.			

Sign of Physician

below cut off

Below Cut Off, but above Age Risk

above cut off