

Date of report: 10-11-2024
Prisca 5.1.0.17

N A

Patient data		Ultrasound data							
Name	Mrs. AARTI SAHU TWIN A	Gestational age at sample date	13 + 4						
Birthday	02-11-1998	Method	Scan						
Age at sample date	26.0	Scan date	08-11-2024						
Patient ID	0382411090004								
Correction factors									
Fetuses	1	IVF	no	Previous trisomy 21 unknown pregnancies					
Weight in kg	52	diabetes	no						
Smoker	no	Origin	Asian						
Pregnancy data		Parameter	Value	Corr. MoM					
Sample Date		PAPP-A	11.99mIU/mL	1.74					
		fb-hCG	30.51 ng/mL	0.92					
Risks at sampling date									
Age risk at sampling date		Trisomy 21	<1:10000						
Overall population risk		Trisomy 13/18	<1:10000						
Risk									
		Trisomy 21 The calculated risk for Trisomy 21 is below the cut off which represents a low risk. After the result of the Trisomy 21 test it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. 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!							
Trisomy 13/18									
The calculated risk for trisomy 13/18 is < 1:10000, which indicates a low risk.									

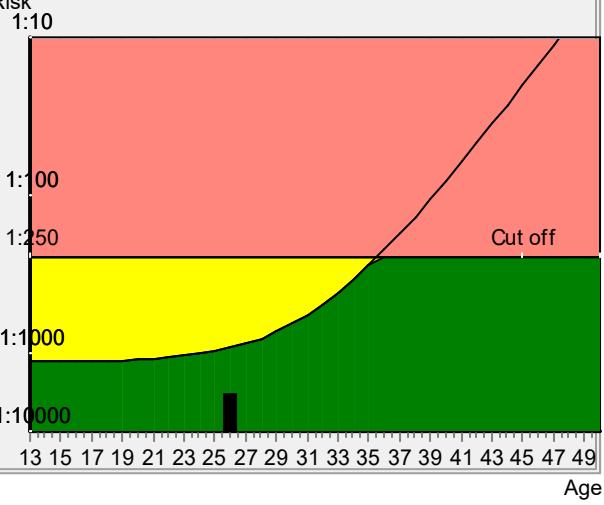
Sign of Physician

below cut off

Below Cut Off, but above Age Risk

above cut off

N A

Patient data								
Name	Mrs. AARTI SAHU TWIN B.	Patient ID	0382411090004.					
Birthday	02-11-1998	Sample ID	A1621729 TWIN B					
Age at sample date	26.0	Sample Date	08-11-2024					
Gestational age	13 + 1							
Correction factors								
Fetuses	1	IVF	no	Previous trisomy 21 pregnancies	unknown			
Weight	52	diabetes	no					
Smoker	no	Origin	Asian					
Biochemical data								
Parameter	Value	Corr. MoM	Ultrasound data					
PAPP-A	11.99 mIU/mL	2.02	Gestational age					
fb-hCG	30.51 ng/mL	0.81	13 + 1					
Risks at sampling date								
Age risk	1:933		Method					
Biochemical T21 risk		<1:10000	CRL Robinson					
Combined trisomy 21 risk		<1:10000	Scan date					
Trisomy 13/18 + NT		<1:10000	08-11-2024					
Risk								
1:10			Crown rump length in mm					
1:100			72					
1:250			Nuchal translucency MoM					
1:1000			0.12					
1:10000			Nasal bone					
13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49			present					
			Sonographer					
			N A					
			Qualifications in measuring NT					
			MD					
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 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								
The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.								

Sign of Physician

 below cut off

 Below Cut Off, but above Age Risk

 above cut off