

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
Weight in kg	52	diabetes	no
Smoker	no	Origin	Asian
		Previous trisomy 21 pregnancies	unknown
Pregnancy data		Parameter	Value
Sample Date	08-11-2024	PAPP-A	11.99mIU/mL
		fb-hCG	30.51 ng/mL
			Corr. MoM
			1.74
			0.92
Risks at sampling date			
Age risk at sampling date	1:946	Trisomy 21	<1:10000
Overall population risk	1:600	Trisomy 13/18	<1:10000
Risk		Trisomy 21	
		<p>The calculated risk for Trisomy 21 is below the cut off which represents a low risk.</p> <p>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.</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>	
Trisomy 13/18			
<p>The calculated risk for trisomy 13/18 is < 1:10000, which indicates a low risk.</p>			

Sign of Physician

Prisca 5.1.0.17
Date of report: 10-11-2024

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
Weight	52	diabetes	no	
Smoker	no	Origin	Asian	
Biochemical data			Ultrasound data	
Parameter	Value	Corr. MoM	Gestational age	13 + 1
PAPP-A	11.99 mIU/mL	2.02	Method	CRL Robinson
fb-hCG	30.51 ng/mL	0.81	Scan date	08-11-2024
Risks at sampling date			Crown rump length in mm	
Age risk	1:933		72	
Biochemical T21 risk	<1:10000		Nuchal translucency MoM	
Combined trisomy 21 risk	<1:10000		0.12	
Trisomy 13/18 + NT	<1:10000		Nasal bone	
			present	
			Sonographer	
			N A	
			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