Prisca

5.2.0.13

Date of report: 18/02/2025

JITM Diagnostics

Patient data					
Name	MRS. RIMPY (A) Patient ID			2502031234/NOD (A)	
Birthday	29/08/1997			2502031234/NOD (A)	
Age at sample date	27.5	Sample Date		16/02/2025	
Gestational age	12 + 5				
Correction factors		•			
Fetuses 2	IVF	no	Previous trisomy 2	1 no	
Weight 55	diabetes	no	pregnancies		
Smoker no	Origin	Asian			
Biochemical data		Ultrasound da	ata		
Parameter Value	Corr. MoM	Gestational	age	12 + 3	
PAPP-A 5.38 mIU/m	ol 0.60	Method		CRL Robinso	
fb-hCG 98.9 ng/ml	1.01				
Risks at sampling date		Crown rump length in mm		6	
Age risk	1:836	Nuchal translucency MoM			
Biochemical T21 risk	1:1551	Nasal bone		preser	
Combined trisomy 21 risk	1:6270	51		DR. GAGANDEEP KAUF	
Trisomy 13/18 + NT	<1:10000	Qualification	ns in measuring NT	M.[
1:10 1:10 1:250 1:1000 1:250 1:10000 1:10000 1:3 15 17 19 21 23 25 27 29 31 33 3 Trisomy 13/18 + NT The calculated risk for trisomy 13 translucency) is < 1:10000, which risk.	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 6270 women with the same data, there is one woman with a trisomy 21 pregnancy and 6269 women with not affected pregnancies. 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 measuremen was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)). The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!				

Sign of Physician

Prisca

5.2.0.13

Date of report: 18/02/2025

JITM Diagnostics

Patient data				
Name	MRS. RIMPY (B) Pa			2502031234/NOD (B)
Birthday	29/08/1997	Sample ID		2502031234/NOD (B)
Age at sample date	27.5	Sample Date		16/02/2025
Gestational age	13 + 1			
Correction factors			-	
Fetuses 2	IVF	no	Previous trisomy 2	1 no
Weight 55	diabetes	no	pregnancies	
Smoker no	Origin	Asian		
Biochemical data		Ultrasound da	ata	
Parameter Value	Corr. MoM	Gestational	age	13 +
PAPP-A 5.38 mIU/m	l 0.51	Method		CRL Robinso
fb-hCG 98.9 ng/ml	1.06	Scan date 16/02/202		
Risks at sampling date		Crown rump length in mm		7
Age risk	1:848	Nuchal translucency MoM		0.8
Biochemical T21 risk	1:902	Nasal bone pres		
Combined trisomy 21 risk	1:4943	51		DR. GAGANDEEP KAUF
Trisomy 13/18 + NT	<1:10000	Qualification	is in measuring NT	M.[
1:10 1:10 1:250 1:1000 1:250 1:1000 1:10000 1:3 15 17 19 21 23 25 27 29 31 33 35 Trisomy 13/18 + NT The calculated risk for trisomy 13 translucency) is < 1:10000, which risk.	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 4943 women with the same data, there is one woman with a trisomy 21 pregnancy and 4942 women with not affected pregnancies. 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 Diag 18: 511-523 (1998)). The laboratory can not be hold responsible for their impac on the risk assessment ! Calculated risks have no diagnostic value!			

Sign of Physician