## JITM Diagnostics

Gestational age  Correction factors  Fetuses 1   IVF   no diabetes   no Origin   Asian    Parameter   Value   Corr. MoM   Gestational age   12 + 6    PAPP-A   10.1 mIU/ml   2.41   Method   CRL Robinsor    For hCG   40.1 ng/ml   0.98   Risks at sampling date    Age risk   1:345   Nasal bone   presen    Combined trisomy 21 risk   1:8256   Sonographer   DR. PREETY SHARMA AGNIHOTR    Trisomy 13/18 + NT   <1:10000    Risk   1:10   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 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!    The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagnostic value!    The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagnostic value!    The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagnostic value!    The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagnostic value!    The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagnostic value!    The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagnostic value!    The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagnostic value!	Patient data						
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Gestational age  13 + 0  Correction factors  Fetuses 1   IVF   no of diabetes   no of the pregnancies   no of the pregnancies	Birthday	23/01/1991			Sample ID		2501005772/NOD
Gestational age  Tetuses  Tetu	Age at sample date	34.0			Sample Date	)	5/01/2025
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Biochemical data  Parameter Value Corr. MoM PAPP-A 10.1 mIU/ml 2.41 fb-hCG 40.1 ng/ml 0.98 Risks at sampling date  Age risk 1:345 Biochemical T21 risk 1:8256 Combined trisomy 21 risk 1:10000  Risk 1:1000  1:1000  Description of the information provided by the referring physician. Please note that risk calculations are statistical approaches and have no oil the risk assessment! Calculated risks have no the risk assessment! Calculated risks have no terms are tablest and the control of the impact on the risk assessment! Calculated risks have no	Weight	68	diabetes		no	pregnancies	
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fb-hCG 40.1 ng/ml 0.98  Risks at sampling date  Age risk 1:345  Biochemical T21 risk 1:8256  Combined trisomy 21 risk 1:1000  Risk 1:10  Risk 1:10  Risk 1:10  Cut off 1:10  1:1000  1:10000  Scan date 4/01/2025  Crown rump length in mm 67.58  Nuchal translucency MoM 0.74  Nasal bone present Sonographer DR. PREETY SHARMA AGNIHOTR Qualifications in measuring NT M.D.  Trisomy 13/18 + NT < 1:10000  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 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!  The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagnostic Value) The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagnostic Value) The laboratory can not be hold responsible for their impact on the risk assessment! Calculated risks have no	Parameter	Value Corr. MoM		Gestational age 12 + 6			
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Combined trisomy 21 risk  Trisomy 13/18 + NT  Sonographer  Qualifications in measuring NT  M.D.  Risk  1:10  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 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!  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 hold responsible for their impact on the risk assessment! Calculated risks have no	•	1:345			Nuchal translucency MoM 0.74		
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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