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| Patient Data Name MRS. MISHIKA GULATI Patient ID 0122062501 Birthday 12/1/1992 Sample ID 114283 Age at term 30.1 Sample Date 25/6/20 Gestational age 12+2 Correction factors 1000 Fetuses 1 IVF unknown Previous trisony 21 unknown Weight in kg 66.5 Diabetes NO Pregnancies unknown Smoker NO Origin Asian 1000 Previous trisony 21 unknown Biochemical Data Ultrasound Data 11 | | | | | | Date of Report PRISCA | 28/6/2022 5.1.0.17 |
|--|-------------------------------------|------------------------|------------|-----------|--|--|---|
| Birthday 12/1/1992 Sample ID 114283 Age at term 30.1 Sample Date 25/6/20 Gestational age 12+2 Correction factors Fetuses 1 IVF unknown Previous trisomy 21 unkno Weight in kg 66.5 Diabetes NO Smoker NO Origin Asian Biochemical Data Ultrasound Data Parameter Value Corr Mom PAPP-A 5.82 mIU/ml 1.33 Bi-hCG 71.5 ng/ml 1.86 Scan date 22/6/20 Risks at sampling date Nuchal translucency Age Risk 1:606 Biochemical T21 risk 1:1541 Combined trisomy 21 risk 1:2485 Trisomy 13/18 + NT <1:10000 Risk Pressource of the result of the Trisomy 21 (with NT) test it is expected that among 21/85 works. After the result of the Trisomy 21 (with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 (with NT) test it is expected that among 2485 women with the same data, there is one woman with a trisomy 21 pregnancies one woman with a trisomy 21 pregnancies and 2484 women with not affected pregnancies. The calculated risk for Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 with NT is The baboratory cannot be hold responsible for their impact on the risk assessment! Calculated risk have no diagnostic value 1.5 hor Tisomy 13/18 with NT is | Patient Data | | | | | | |
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| Combined trisomy 21 risk1:2485Trisomy 13/18 + NT<1:10000 | Age Risk | | | 1:606 | Nuchal translu | cency MoM | 1.28 |
| Trisomy 13/18 + NT <1:10000 | Biochemical T21 risk | ochemical T21 risk | | | Nasal bone | | Present |
| RiskDown's Syndrome Risk (Trisomy 21 Screening)RiskThe calculated risk for Trisomy 21 (withNT) is below the cut off, which represents a low risk. After the result of the Trisomy 21 (with NT) test it is expected that among 2485 women with the same data, there is one woman with a trisomy 21 pregnancy and 2484 women with not affected pregnancies. The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that the risk calculations are statistical aapproaches and have no diagnostic value!Trisomy 13/18+NT The calculated risk for Trisomy 13/18 with NT isThe laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic | Combined trisomy 21 risk | | | 1:2485 | | | |
| Risk. The calculated risk for Trisomy 21 (withNT) is below the cut off, which represents a low risk. 1:100 After the result of the Trisomy 21 (with NT) test it is expected that among 2485 women with the same data, there is one woman with a trisomy 21 pregnancy and 2484 women with not affected pregnancies. 1:1000 The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that the risk calculations are statistical aapproaches and have no diagnostic value! Trisomy 13/18+NT The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic | Trisomy 13/18 + NT | | | <1:10000 | | | |
| 1:100Cut off1:250Cut off1:1000Cut off1:1000Cut off1:1000Firstony 21 (with NT) test it is expected that among 2485 women with the same data, there is one woman with a trisomy 21 pregnancy and 2484 women with not affected pregnancies.1:1000The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that the risk calculations are statistical aapproaches and have no diagnostic value!Trisomy 13/18+NTThe laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic | Risk | | | | Down's Syndr | ome Risk (Trisomy 21 | Screening) |
| Trisomy 13/18+NT The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic | 1:100 1:250 1:1000 1:10000 | 29 31 33 | 35 37 39 4 | | cut off, which After the resul expected that a same data, the pregnancy and The calculated the informatio note that the r | represents a low risk. t of the Trisomy 21 (wit among 2485 women with re is one woman with a 2484 women with not a l risk by PRISCA depen n provided by the referr isk calculations are statis | h NT) test it is h the trisomy 21 affected pregnancies. ids on the accuracy of ing physician. Please |
| | Trisomy 13/18+NT | | | | | | |