

 Date of Report
 22/12/2022

 PRISCA
 5.1.0.17

| Patient Data | | | | | |
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| Name MRS. SWAPNA AMAL | | | Patient ID | | 012212210132 |
| Birthday 19/04/1995 | | | Sample ID | | 11360864 |
| Age at term | | 28.01 | Sample Date | | 21/12/2022 |
| Gestational age 12+5 | | | | | |
| Correction factors | | | | | |
| Fetuses | 1 IVF | | unknown | Previous trisomy 21 | unknown |
| Weight in kg | 55 Diabetes | | NO | Pregnancies | unknown |
| Smoker | NO Origin | | Asian | | |
| Biochemical Data | | | Ultrasound Data | | |
| Parameter | Value | Corr Mom | | | 12+5 |
| PAPP-A | 6.6 mIU/ml | 1.02 | Method | | CRL (<>Robinson) |
| fb-hCG | 112.5 ng/ml | 3.08 | Scan date | | 21/12/2022 |
| Risks at sampling date | | | Nuchal translucency 1.23 | | |
| Age Risk | | 1:823 | Nuchal translucency MoM 0.3 | | 0.75 |
| Biochemical T21 risk | | 1:331 | Nasal bone Presen | | |
| Combined trisomy 21 risk | 1:1912 | | | | |
| Trisomy 13/18 + NT | | <1:10000 | | | |
| Risk | | Down's Syndrome Risk (Trisomy 21 Screening) | | | |
| 1:100 1:250 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:10000 1:1 | | | The calculated risk for Trisomy 21(with NT) is below the cut off, which represents a low risk. After the result of the Trisomy 21with NT test it is expected that among 1912 women with the same data, there is one woman with a trisomy 21 pregnancy and 1911 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! The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values | | |
| Risk A | Above Cut Off | | Risk above Ag | e Risk | Risk below Age risk |