MOLQ Express Lab
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| | |
|---|---|
| Date of Report | 17/3/2023 |
| PRISCA | 5.2.0.13 |

| Patient Data | | | |
|---|---|---|---|
| Name | MRS. AARTI AGNIHOTRI | Patient ID | 012303160239 |
| Birthday | 5/1/1993 | Sample ID | 11476276 |
| Age at term | 30.2 | Sample Date | 16/3/2023 |
| Gestational age | 13+5 | | |

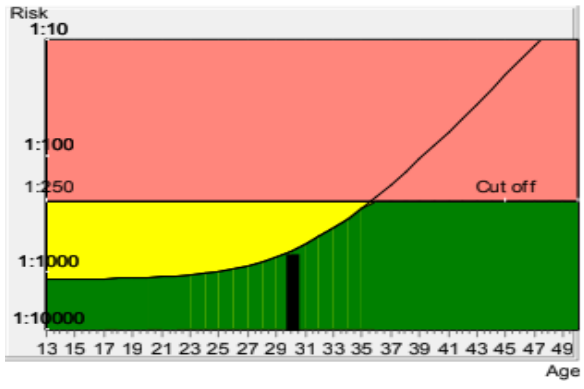
| Correction factors | | | | | |
|---|---|---|---|---|---|
| Fetuses | 1 | IVF | unknown | Previous trisomy 21 | unknown |
| Weight in kg | 47 | Diabetes | NO | Pregnancies | unknown |
| Smoker | NO | Origin | Asian | | |

| Biochemical Data | | |
|---|---|---|
| **Parameter** | **Value** | **Corr Mom** |
| PAPP-A | 5.82 mIU/ml | 0.53 |
| fb-hCG | 67.2 ng/ml | 2.37 |

| Risks at sampling date | |
|---|---|
| Age Risk | 1:657 |
| Biochemical T21 risk | 1:115 |
| Combined trisomy 21 risk | 1:713 |
| Trisomy 13/18 + NT | <1:10000 |

| Ultrasound Data | |
|---|---|
| | 12+6 |
| Method | CRL (<>Robinson) |
| Scan date | 10/3/2023 |
| Nuchal translucency | 1.4 |
| Nuchal translucency MoM | 0.85 |
| Nasal bone | Present |

**Risk**




**The calculated risk for Trisomy 13/18 (with NT) is <1:10000 , which indicates a low risk**

**Down's Syndrome Risk (Trisomy 21 Screening)**

**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 713 women with the same data, there is one woman with a trisomy 21 pregnancy and 712 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 Above Cut Off  Risk above Age Risk  Risk below Age risk