MOLQ Express Lab

*Free Home Sample Collection 9999 778 778

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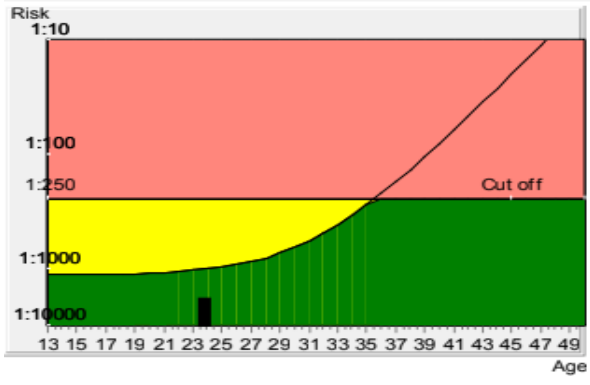
| | |
|---|---|
| Date of Report | 23/2/2023 |
| PRISCA | 5.1.0.17 |

| Patient Data | | | |
|---|---|---|---|
| Name | MRS. POOJA | Patient ID | 012302220098 |
| Birthday | 2/5/1999 | Sample ID | 11476899 |
| Age at term | 24.2 | Sample Date | 22/2/2023 |
| Gestational age | 13+1 | | |

| Correction factors | | | | | |
|---|---|---|---|---|---|
| Fetuses | 1 | IVF | unknown | Previous trisomy 21 | unknown |
| Weight in kg | 48 | Diabetes | NO | Pregnancies | unknown |
| Smoker | NO | Origin | Asian | | |

| Biochemical Data | | | Ultrasound Data | |
|---|---|---|---|---|
| Parameter | Value | Corr Mom | | 13+0 |
| PAPP-A | 3.89 mIU/ml | 0.44 | Method | CRL (<>Robinson) |
| fb-hCG | 25.9 ng/ml | 0.76 | Scan date | 21/2/2023 |
| **Risks at sampling date** | | | Nuchal translucency | 1.7 |
| Age Risk | | 1:1027 | Nuchal translucency MoM | 0.99 |
| Biochemical T21 risk | | 1:1526 | Nasal bone | Present |
| Combined trisomy 21 risk | | 1:6930 | | |
| Trisomy 13/18 + NT | | <1:10000 | | |

**Risk**




**Trisomy 13/18+NT**

**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 6930 women with the same data, there is one woman with a trisomy 21 pregnancy and 6929 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