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 Date of Report
 11/10/2023

 PRISCA
 5.2.0.13

| | | | | | FMSCA | 3.2.0.13 | |
|----------------------|-------------------|-------------|-------------|---|--|---------------------------|--|
| Patient Data | Value | | | | | | |
| Name | | MRS JUI | I MUNNA | | Patient ID | 012310100236 | |
| Birthday | | | 17-11-2000 |) | Sample ID | 11478628 | |
| Age at delivery | | | 23.3 | } | Sample Date | 10/10/2023 | |
| Correction factors | , | | | | | | |
| Fetuses | 1 | IVF | | unknown | Previous trisomy 21 | unknown | |
| Weight in kg | 80 | Diabete | S | NO | Pregnancies | unknown | |
| Smoker | NO | Origin | | Asian | | | |
| Biochemical Data | | 9 | | Risks at sampl | | | |
| Parameter | Value | C | Corr MoM | Age Risk | | 1:1454 | |
| AFP | 55.4 | ng/ml | 1.39 | Biochemical T | risomy 21 Risk | 1:8499 | |
| uE3 | 1.37 | ng/ml | 1.25 | Neural Tube I | Defect Risk | Low risk area | |
| hCG | 11698.8 | mIU/ml | 0.62 | Trisomy 18 | | <1:10000 | |
| Inhibin | 214.7 | IU/ml | 1.68 | | | | |
| Ultrasound Data | | | | Down's Syndre | ome Risk (Trisomy 2 | 21 Screening) | |
| Gestational age | | 18+2 | | | l risk for Trisomy 21 | is below the cut off | |
| Method | BPD (♦ Hadlock) | | | which represents a low risk. After the result of the Trisomy 21 test it is expected that | | | |
| | | , | , | among 8499 w | omen with the same | data, there is one woman | |
| Risk | | | | with a trisomy affected pregna | 21 pregnancy and 849 | 98 women with not | |
| Risk 1:10 | | | | The calculated risk by PRISCA depends on the accuracy of | | | |
| 1.10 | | | | the information provided by the referring physician. Please | | | |
| | | | | | | atistical aapproaches and | |
| | | | | have no diagno | ostic value! | | |
| 1:100 | | / | | | | | |
| 1:250 Cut off | | | | Trisomy 18 | | | |
| | | | | The calculated risk for Trisomy 18 is <1:10000, which | | | |
| 1:1 <mark>p00</mark> | | | | indicates a low risk | | | |
| 1:10000 | | | | | Defect (NTD) Screen | ing | |
| 13 15 17 19 21 | 23 25 27 29 31 33 | 35 37 39 41 | 43 45 47 49 | | The corrected MoM for AFP (1.39) is located in the low | | |
| | | | | risk area for n | eural tube defects. | | |

value!

The laboratory can not be held responsible for their impact on the risk assessment! Calculated value has no diagnostic