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Date of Report 07/01/2024 **PRISCA** 5.2.0.13

| | | | | | PRISCA | 5.2.0.13 |
|--|-----------------|--------|----------|---|------------------------|--------------------------|
| Patient Data | Value | | | | | |
| Name | | MRS. | SUREKHA | | Patient ID | 012401050069 |
| Birthday | 01-07-1997 | | | 7 | Sample ID | 11834493 |
| Age at delivery | 26.9 | | |) | Sample Date | 05/01/2024 |
| Correction factors | 5 | | | | | |
| Fetuses | 1 | IVF | | unknown | Previous trisomy 21 | unknown |
| Weight in kg | 47 | Diabet | es | NO | Pregnancies | unknown |
| Smoker | NO | Origin | | Asian | | |
| Biochemical Data | | 9 | | Risks at sampl | ing date | |
| Parameter | Value | | Corr MoM | Age Risk | | 1:1256 |
| AFP | 92.7 | ng/ml | 1.19 | Biochemical T | risomy 21 R isk | 1:3904 |
| uE3 | 1.98 | ng/ml | 1.13 | Neural Tube I | Defect Risk | Low risk area |
| hCG | 39494.2 | mIU/ml | 1.87 | Trisomy 18 | | <1:10000 |
| Inhibin | 241.2 | IU/ml | 1.27 | | | |
| Ultrasound Data | | | | Down's Syndr | ome Risk (Trisomy | 21 Screening) |
| Gestational age | | 20+0 | | | • | 1 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 3904 w | omen with the same | data, there is one woman |
| Risk | | | | with a trisomy affected pregna | | 903 women with not |
| 1:10 | | | | 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 diagno | ostic value! | |
| 1:100 | | / | | | | |
| 1:250 | | | Cut off | | | |
| COLOTT | | | | Trisomy 18 | | |
| | | | | The calculated risk for Trisomy 18 is <1:10000, which | | |
| 1:1 <mark>000</mark> | | | | indicates a low risk | | |
| | | | | Neural Tube Defect (NTD) Screening | | |
| 1:10000 | | | | | | |
| 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49 | | | | The corrected MoM for AFP (1.19) is located in the low risk area for neural tube defects. | | |
| <i>a</i> | | | | pisk area for n | emai tube defects. | |

value!

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