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|   |                                 |   |   | Date of Report<br>PRISCA  | 26/07/2019<br>5.0.2.37 |  |
|---|---------------------------------|---|---|---|------------------------|--|
| Patient Data  |                                 |   |   | 1145 611  | 5787 <b>2</b> 787      |  |
| Name  | Mrs Jyoti (Fetus 2)             |   |   |   | 011907260052           |  |
| Birthday  |                                 | 7/10/1992   | Sample ID   |   | 010630896              |  |
| Age at delivery   |                                 | 26.8  | Sample Date   |   | 25/07/2019             |  |
| Gestational age   |                                 | 12+4  |   |   |                        |  |
| Correction factors  |                                 |   |   |   |                        |  |
| Fetuses   | 2 IVF                           |   | unknown   | Previous trisomy 21   | unknown                |  |
| Weight in kg  | 54 Diabetes                     |   | no  | Pregnancies   |                        |  |
| Smoker  | no Origin                       |   | Asian   |   |                        |  |
| Biochemical Data  |                                 |   | Ultrasound Data   |   |                        |  |
| Parameter   | Value                           | Corr Mom  | Gestational age   | e   | 12+3                   |  |
| PAPP-A  | 4.12  mIU/ml                    | 0.45  | Method  |   | CRI                    |  |
| fb-hCG  | 107.5  ng/ml                    | 1.02  | Scan Date   |   | 24/07/19               |  |
| Risks at sampling date  |                                 |   | CRL 59.2  |   |                        |  |
| Age Risk  |                                 | 1:873   | Nuchal translu  | cency MoM   | 0.71                   |  |
| Overall biochemical risk  |                                 | 1:624   | Nasal Bone  |   | present                |  |
| Trisomy 21  |                                 | 1:3900  | Sonographer   |   | DR.NITIN YADAV         |  |
| Trisomy 13/18   |                                 | <1:10000  | Qualification is  | n measuring NT  | MBBS,MD                |  |
| Risk  |                                 |   | Down's Syndrome Risk (Trisomy 21 Screening)   |   |                        |  |
| Risk<br>1:10  |                                 |   | The calculated risk for Trisomy 21 is below the cut off, which represents a low risk. |   |                        |  |
| 1:100<br>1:250<br>1:1000<br>1:10000<br>13 15 17 19 21 23 25 27                | Out off<br>1 43 45 47 49<br>Age | After the result of the Trisomy 21 test (with NT) it is expected that among 3900 women with the same data, there is one woman with a trisomy 21 pregnancy and 3899 women with not affected pregnancies. The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.  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! |   |   |                        |  |
| Trisomy 13/18 + NT  |                                 |   |   | The laboratory cannot be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic values |                        |  |
| The calculated risk for Trisomy 13/18 is <1:10000, which indicates a low risk |                                 |   |   | i: Calculated risks have no   | diagnosuc values       |  |

Risk above Age Risk

Risk below Age risk

Risk Above Cut Off