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Date of Report 11-04-19 PRISCA 5.0.2.37

| | | | | | PRISCA | 5.0.2.37 |
|-----------------------------|-------|----------|-------------|---|------------------------|----------------|
| Patient Data | | | | | | |
| Name | | | Mrs Sonia | Patient ID | | 011904100197 |
| Birthday | | | 21-10-92 | Sample ID | | 10396160 |
| Age at delivery | | | 26.5 | Sample Date | | 10-04-2019 |
| Gestational age by | | | 12+3 | | | |
| Correction factors | | | | | | |
| Fetuses | 1 | IVF | | unknown | Previous trisomy 21 | unknown |
| Weight in kg | 44 | Diabetes | | no | Pregnancies | |
| Smoker | no | Origin | | Asian | | |
| Biochemical Data | | | | Ultrasound Data | | |
| Parameter | Value | | Corr Mom | Gestational ag | e | 12+2 |
| PAPP-A | 4.15 | mIU/ml | 0.78 | Method | | CRL (⇔Hadlock) |
| fb-hCG | 64.2 | ng/ml | 1.27 | Scan Date | | 09-04-19 |
| | | | | CRL measurm | ents | 58 |
| Risks at sampling date | | | | Nuchal translucency MoM 0.65 | | |
| Age Risk | | | 1:887 | Nasal bone | | Present |
| Biochemical T21 Risk 1:1867 | | 1:1867 | Sonographer | | DR.PRAKASH LALCHANDANI | |
| Combined Trisomy 21 R | isk | | <1:10000 | Qualifications | in measuring NT | MD |
| Trisomy 13/18 | | | <1:10000 | | | |
| Risk 1:10 | | | | Down's Syndrome Risk (Trisomy 21 Screening) The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk. | | |
| 1:1000 1:1000 1:10000 | | | | After the result of the Trisomy 21 test (with NT) it is expected that among more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy. The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that risk calculations are statistical approach and have no diagnostic values! The patient combined risk presumes the NT measurement was done according to accepted guidelines The laboratory can not be hold responsible for their impact on the risk assessment! Calculated risks have no diagnostic value! | | |