

*Free Home Sample Collection 9999 778 778

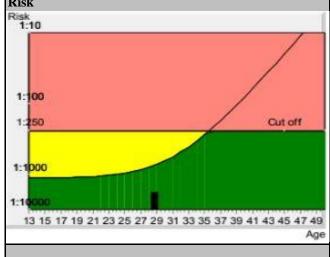


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Date of Report 27-12-18 **PRISCA** 5.0.2.37

| | | | | | TRISCA | 5.0.2.57 |
|---------------------------|----|----------|-----------|-------------|---------------------|--------------|
| Patient Data | | | | | | |
| Name | | | Mrs Jyoti | Patient ID | | 011812250171 |
| Birthday | | | 14-04-90 | Sample ID | | 10379929 |
| Age at delivery | | | 28.7 | Sample Date | | 25/12/2018 |
| Gestational age | | | 13+0 | | | |
| Correction factors | | | | | | |
| Fetuses | 1 | IVF | | unknown | Previous trisomy 21 | unknown |
| Weight in kg | 52 | Diabetes | | no | Pregnancies | |
| Smoker | no | Origin | | Asian | | |

| Biochemical Data | | | Ultrasound Data | | |
|--------------------------|--------------|----------|---|------------------------|--|
| Parameter | ameter Value | | Gestational age | 12+2 | |
| PAPP-A | 2.33 mIU/ml | 0.43 | Method | CRL (<>Hadlock) | |
| fb-hCG | 37.02 ng/ml | 0.83 | Scan Date | 21-12-18 | |
| Risks at sampling date | | | CRL | 57.3 | |
| Age Risk | | 1:758 | Nuchal translucency MoM | 0.93 | |
| Biochemical T21 Risk | | 1:901 | Nasal Bone | Present | |
| Combined Trisomy 21 Risk | k | 1:4701 | Sonographer | DR.SUBHASISH MOHAPATRA | |
| Trisomy 13/18 + NT | | <1:10000 | Qualifications in measuring NT | MBBS,MD | |
| Risk | | | Down's Syndrome Risk (Trisomy | 21 Screening) | |
| Risk | | - 1 | The calculated risk for Trisomy 21 (with nuchal | | |



Trisomy 13/18 + NTThe calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.

The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.

After the result of the Trisomy 21 test (with NT) it is expected that among 4701 women with the same data, there is one woman with a trisomy 21 pregnancy and 4700 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 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 held responsible for their impact on the risk assessment! Calculated value has no diagnostic value!