

| Patient data   |                   |           |  |                                 |
|--|-------------------|-----------|--|---------------------------------|
| Name   | MRS. SHILPI SINGH |           | Patient ID   | 2103044890/DHJ                  |
| Birthday   | 01-04-1990        |           | Sample ID  | 2103044890/DHJ                  |
| Age at sample date   | 31.0              |           | Sample Date  | 23-03-2021                      |
| Gestational age  | 12 + 6            |           |  |                                 |
| Correction factors   |                   |           |  |                                 |
| Fetuses  | 1                 | IVF       | no   | Previous trisomy 21 pregnancies |
| Weight   | 73                | diabetes  | no   |                                 |
| Smoker   | no                | Origin    | Asian  |                                 |
| Biochemical data   |                   |           | Ultrasound data  |                                 |
| Parameter  | Value             | Corr. MoM | Gestational age  | 12 + 1                          |
| PAPP-A   | 2.1 mIU/ml        | 0.58      | Method   | CRL Robinson                    |
| fb-hCG   | 22.5 ng/ml        | 0.55      | Scan date  | 18-03-2021                      |
| Risks at sampling date   |                   |           | Crown rump length in mm  | 58.8                            |
| Age risk   | 1:574             |           | Nuchal translucency MoM  | 1.17                            |
| Biochemical T21 risk   | 1:3408            |           | Nasal bone   | present                         |
| Combined trisomy 21 risk   | 1:9056            |           | Sonographer  | ..                              |
| Trisomy 13/18 + NT   | <1:10000          |           | Qualifications in measuring NT   | MD                              |
| Risk   |                   |           | Trisomy 21   |                                 |
|  |                   |           | <p><b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b></p> <p>After the result of the Trisomy 21 test (with NT) it is expected that among 9056 women with the same data, there is one woman with a trisomy 21 pregnancy and 9055 women with not affected pregnancies.</p> <p>The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician. Please note that risk calculations are statistical approaches and have no diagnostic value!</p> <p>The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).</p> <p>The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!</p> |                                 |
| Trisomy 13/18 + NT   |                   |           |  |                                 |
| <p><b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b></p> |                   |           |  |                                 |

Sign of Physician

below cut off
  Below Cut Off, but above Age Risk
  above cut off