

| Patient data   |             |                                 |  |
|--|-------------|---------------------------------|--|
| Name   | MRS. SWETA  | Patient ID                      |  |
| Birthday   | 12/11/1995  | Sample ID                       | 2409041483/NOD   |
| Age at sample date   | 28.9        | Sample Date                     | 19/09/2024   |
| Gestational age  | 12 + 4      |                                 |  |
| Correction factors   |             |                                 |  |
| Fetuses  | 1           | IVF                             | no   |
| Weight   | 65          | diabetes                        | no   |
| Smoker   | no          | Origin                          | Asian  |
|  |             | Previous trisomy 21 pregnancies | no   |
| Biochemical data   |             | Ultrasound data                 |  |
| Parameter  | Value       | Corr. MoM                       |  |
| PAPP-A   | 4.25 mIU/ml | 1.14                            | Gestational age 12 + 4   |
| fb-hCG   | 20.6 ng/ml  | 0.47                            | Method CRL Robinson  |
| Risks at sampling date   |             |                                 | Scan date 19/09/2024   |
| Age risk   |             | 1:736                           | Crown rump length in mm 64.5   |
| Biochemical T21 risk   |             | <1:10000                        | Nuchal translucency MoM 0.73   |
| Combined trisomy 21 risk   |             | <1:10000                        | Nasal bone present   |
| Trisomy 13/18 + NT   |             | <1:10000                        | Sonographer DR. (MRS.) NEERJA CHOPRA   |
|  |             |                                 | Qualifications in measuring NT MD  |
| Risk   |             |                                 | Trisomy 21   |
| 1:10   |             |                                 | <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 more than 10000 women with the same data, there is one woman with a trisomy 21 pregnancy.</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> |
| 1:100  |             |                                 |  |
| 1:250  |             |                                 |  |
| 1:1000   |             |                                 |  |
| 1:10000  |             |                                 |  |
| 13 15 17 19 21 23 25 27 29 31 33 35 37 39 41 43 45 47 49   | Age         |                                 |  |
| 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

