

| Patient data   |                  |  |                                   |
|--|------------------|--|-----------------------------------|
| Name   | MRS. GUNJAN GARG | Patient ID   |                                   |
| Birth day  | 8/06/1996        | Sample ID  | 2406039224/NOD                    |
| Age at sample date   | 28.0             | Sample Date  | 21/06/2024                        |
| Gestational age  | 13 + 3           |  |                                   |
| Correction factors   |                  |  |                                   |
| Fetuses  | 1                | IVF  | no                                |
| Weight   | 59               | diabetes   | no                                |
| Smoker   | no               | Origin   | Asian                             |
|  |                  | Previous trisomy 21 pregnancies  | no                                |
| Biochemical data   |                  | Ultrasound data  |                                   |
| Parameter  | Value            | Corr. MoM  |                                   |
| PAPP-A   | 5.1 mIU/ml       | 0.87   | Gestational age 13 + 3            |
| fb-hCG   | 47.1 ng/ml       | 1.16   | Method CRL Robinson               |
| Risks at sampling date   |                  |  | Scan date 21/06/2024              |
| Age risk   |                  | 1:818  | Crown rump length in mm 75.6      |
| Biochemical T21 risk   |                  | 1:2689   | Nuchal translucency MoM 0.70      |
| Combined trisomy 21 risk   |                  | <1:10000   | Nasal bone present                |
| Trisomy 13/18 + NT   |                  | <1:10000   | Sonographer ..                    |
|  |                  |  | Qualifications in measuring NT .. |
| 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 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> |                                   |
| 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

