

JITM Diagnostics

| Patient data   |                 |                                 |  |
|--|-----------------|---------------------------------|--|
| Name   | MRS. KANU PRIYA | Patient ID                      |  |
| Birth day  | 13/01/1995      | Sample ID                       | 2411025940/NOD   |
| Age at sample date   | 29.8            | Sample Date                     | 14/11/2024   |
| Gestational age  | 12 + 2          |                                 |  |
| Correction factors   |                 |                                 |  |
| Fetuses  | 1               | IVF                             | no   |
| Weight   | 60.8            | diabetes                        | no   |
| Smoker   | no              | Origin                          | Asian  |
|  |                 | Previous trisomy 21 pregnancies | no   |
| Biochemical data   |                 | Ultrasound data                 |  |
| Parameter  | Value           | Corr. MoM                       |  |
| PAPP-A   | 2.6 mIU/ml      | 0.73                            | Gestational age 12 + 0   |
| fb-hCG   | 31.1 ng/ml      | 0.68                            | Method CRL Robinson  |
| Risks at sampling date   |                 |                                 | Scan date 12/11/2024   |
| Age risk   |                 | 1:653                           | Crown rump length in mm 55.4   |
| Biochemical T21 risk   |                 | 1:4661                          | Nuchal translucency MoM 0.89   |
| Combined trisomy 21 risk   |                 | <1:10000                        | Nasal bone present   |
| Trisomy 13/18 + NT   |                 | <1:10000                        | Sonographer DR. B. CHAUHAN MBBS MD   |
|  |                 |                                 | 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 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