

| Patient data   |                    |           |  |                          |
|--|--------------------|-----------|--|--------------------------|
| Name   | MRS. SALONI JINDAL |           | Patient ID   | 2503063964/NOD           |
| Birthday   | 30/05/1994         |           | Sample ID  | 2503063964/NOD           |
| Age at sample date   | 30.8               |           | Sample Date  | 31/03/2025               |
| Gestational age  | 12 + 4             |           |  |                          |
| Correction factors   |                    |           |  |                          |
| Fetuses  | 1                  | IVF       | no   | Previous trisomy 21      |
| Weight   | 60                 | diabetes  | no   | pregnancies              |
| Smoker   | no                 | Origin    | Asian  |                          |
| Biochemical data   |                    |           | Ultrasound data  |                          |
| Parameter  | Value              | Corr. MoM | Gestational age  | 12 + 4                   |
| PAPP-A   | 2.6 mIU/ml         | 0.64      | Method   | CRL Robinson             |
| fb-hCG   | 53.7 ng/ml         | 1.20      | Scan date  | 31/03/2025               |
| Risks at sampling date   |                    |           | Crown rump length in mm  | 64.1                     |
| Age risk   | 1:579              |           | Nuchal translucency MoM  | 0.73                     |
| Biochemical T21 risk   | 1:833              |           | Nasal bone   | present                  |
| Combined trisomy 21 risk   | 1:4784             |           | Sonographer  | DR. (MRS.) NEERJA CHOPRA |
| Trisomy 13/18 + NT   | <1:10000           |           | Qualifications in measuring NT   | MD                       |
|  |                    |           | 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 4784 women with the same data, there is one woman with a trisomy 21 pregnancy and 4783 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