

| Patient data   |                  |                                 |  |
|--|------------------|---------------------------------|--|
| Name   | MRS. SAPNA RAWAT | Patient ID                      |  |
| Birthday   | 22/12/94         | Sample ID                       | 2404042062/NOD   |
| Age at sample date   | 29.3             | Sample Date                     | 20/04/24   |
| Gestational age  | 11 + 4           |                                 |  |
| Correction factors   |                  |                                 |  |
| Fetuses  | 1                | IVF                             | no   |
| Weight   | 57               | diabetes                        | no   |
| Smoker   | no               | Origin                          | Asian  |
|  |                  | Previous trisomy 21 pregnancies | no   |
| Biochemical data   |                  | Ultrasound data                 |  |
| Parameter  | Value            | Corr. MoM                       |  |
| PAPP-A   | 2.98 mIU/ml      | 1.10                            | Gestational age 11 + 4   |
| fb-hCG   | 70.5 ng/ml       | 1.39                            | Method CRL Robinson  |
| Risks at sampling date   |                  |                                 | Scan date 20/04/24   |
| Age risk   |                  | 1:673                           | Crown rump length in mm 50.7   |
| Biochemical T21 risk   |                  | 1:2416                          | Nuchal translucency MoM 0.66   |
| Combined trisomy 21 risk   |                  | <1:10000                        | Nasal bone present   |
| Trisomy 13/18 + NT   |                  | <1:10000                        | Sonographer ..   |
|  |                  |                                 | Qualifications in measuring NT ..  |
| 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  |                  |                                 |  |
|  | 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

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