

| Patient data   |             |  |                                |
|--|-------------|--|--------------------------------|
| Name   | MRS. SHEELY | Patient ID   | 2109220706/AMB                 |
| Birthday   | 15-07-1991  | Sample ID  | 2109220706/AMB                 |
| Age at sample date   | 30.2        | Sample Date  | 22-09-2021                     |
| Gestational age  | 11 + 1      |  |                                |
| Correction factors   |             |  |                                |
| Fetuses  | 1           | IVF  | no                             |
| Weight   | 58.2        | diabetes   | no                             |
| Smoker   | no          | Origin   | Asian                          |
|  |             | Previous trisomy 21 pregnancies  | no                             |
| Biochemical data   |             | Ultrasound data  |                                |
| Parameter  | Value       | Corr. MoM  | Gestational age                |
| PAPP-A   | 2.38 mIU/ml | 1.14   | 11 + 1                         |
| fb-hCG   | 33.7 ng/ml  | 0.64   | Method                         |
|  |             |  | CRL Robinson                   |
|  |             |  | Scan date                      |
|  |             |  | 22-09-2021                     |
| Risks at sampling date   |             |  | Crown rump length in mm        |
| Age risk   |             | 1:597  | 45                             |
| Biochemical T21 risk   |             | <1:10000   | Nuchal translucency MoM        |
| Combined trisomy 21 risk   |             | <1:10000   | 0.48                           |
| Trisomy 13/18 + NT   |             | <1:10000   | Nasal bone                     |
|  |             |  | present                        |
|  |             |  | Sonographer                    |
|  |             |  | .                              |
|  |             |  | 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



below cut off

Below Cut Off, but above Age Risk

above cut off