

| Patient data   |             |  |                 |
|--|-------------|--|-----------------|
| Name   | Ms.JASPAL   | Patient ID   | 1802220003/AMB  |
| Birthday   | 07-07-1990  | Sample ID  | 1802220003/AMB  |
| Age at sample date   | 29.0        | Sample Date  | 10-11-2018      |
| Gestational age  | 13 + 1      |  |                 |
| 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  | Gestational age |
| PAPP-A   | 1.63 mIU/ml | 0.60   | 13 + 1          |
| fb-hCG   | 33.8 ng/ml  | 0.77   | Method          |
|  |             |  | CRL Robinson    |
|  |             |  | Scan date       |
|  |             |  | 10-11-2018      |
| Risks at sampling date   |             | Crown rump length in mm  |                 |
| Age risk   | 1:676       | Nuchal translucency MoM  |                 |
| Biochemical T21 risk   | 1:2190      | Nasal bone   |                 |
| Combined trisomy 21 risk   | <1:10000    | Sonographer  |                 |
| Trisomy 13/18 + NT   | <1:10000    | 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