

JITM Diagnostics

| Patient data   |             |                                 |   |
|--|-------------|---------------------------------|---|
| Name   | MRS. POOJA  |                                 | Patient ID  |
| Birthday   | 25/06/1993  |                                 | Sample ID   |
| Age at sample date   | 31.6        |                                 | Sample Date   |
| Gestational age  | 13 + 1      |                                 |   |
| Correction factors   |             |                                 |   |
| Fetuses  | 1           | IVF                             | no  |
| Weight   | 96          | diabetes                        | no  |
| Smoker   | no          | Origin                          | Asian   |
|  |             | Previous trisomy 21 pregnancies | no  |
| Biochemical data   |             | Ultrasound data                 |   |
| Parameter  | Value       | Corr. MoM                       | Gestational age   |
| PAPP-A   | 1.38 mIU/ml | 0.48                            | 13 + 0  |
| fb-hCG   | 14.1 ng/ml  | 0.38                            | Method  |
| Risks at sampling date   |             |                                 | CRL Robinson  |
| Age risk   | 1:531       |                                 | Scan date   |
| Biochemical T21 risk   | 1:3851      |                                 | 17/01/2025  |
| Combined trisomy 21 risk   | <1:10000    |                                 | Crown rump length in mm   |
| Trisomy 13/18 + NT   | <1:10000    |                                 | 70.5  |
|  |             |                                 | Nuchal translucency MoM   |
|  |             |                                 | 0.87  |
|  |             |                                 | Nasal bone  |
|  |             |                                 | present   |
|  |             |                                 | Sonographer   |
|  |             |                                 | ..  |
|  |             |                                 | Qualifications in measuring NT  |
|  |             |                                 | ..  |
| 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 free beta HCG level is low.</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