

| Patient data   |                    |  |                                |
|--|--------------------|--|--------------------------------|
| Name   | DR. DEEPTI MANOCHA | Patient ID   | 1703220202/AMB                 |
| Birthday   | 13-12-1975         | Sample ID  | 1703220202/AMB                 |
| Age at sample date   | 41.2               | Sample Date  | 08-03-2017                     |
| Gestational age  | 12 + 3             |  |                                |
| Correction factors   |                    |  |                                |
| Fetuses  | 1 IVF              | no   | Previous trisomy 21            |
| Weight   | 87 diabetes        | no   | pregancies                     |
| Smoker   | no                 | Origin   | Asian                          |
| Biochemical data   |                    | Ultrasound data  |                                |
| Parameter  | Value              | Corr. MoM  | Gestational age                |
| PAPP-A   | 2.2 mIU/ml         | 1.04   | 12 + 3                         |
| fb-hCG   | 37.5 ng/ml         | 1.16   | Method                         |
| Risks at sampling date   |                    |  | CRL Robinson                   |
| Age risk   | 1:56               |  | Scan date                      |
| Biochemical T21 risk   | 1:270              |  | 08-03-2017                     |
| Combined trisomy 21 risk   | 1:1361             |  | Crown rump length in mm        |
| Trisomy 13/18 + NT   | <1:10000           |  | 61.9                           |
|  |                    |  | Nuchal translucency MoM        |
|  |                    |  | 0.71                           |
|  |                    |  | Nasal bone                     |
|  |                    |  | present                        |
|  |                    |  | Sonographer                    |
|  |                    |  | DR. POONAM LOOMBA              |
|  |                    |  | 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 1361 women with the same data, there is one woman with a trisomy 21 pregnancy and 1360 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