

JITM Diagnostics

| Patient data  |                    |                                 |   |
|---|--------------------|---------------------------------|---|
| Name  | MRS. LALITA GULATI | Patient ID                      |   |
| Birthday  | 12/05/1990         | Sample ID                       | 2411032331/NOD  |
| Age at sample date  | 34.5               | Sample Date                     | 17/11/2024  |
| Gestational age   | 13 + 3             |                                 |   |
| Correction factors  |                    |                                 |   |
| Fetuses   | 1                  | IVF                             | no  |
| Weight  | 58.45              | diabetes                        | no  |
| Smoker  | no                 | Origin                          | Asian   |
|   |                    | Previous trisomy 21 pregnancies | no  |
| Biochemical data  |                    | Ultrasound data                 |   |
| Parameter   | Value              | Corr. MoM                       |   |
| PAPP-A  | 10.9 mIU/ml        | 1.84                            | Gestational age   |
| fb-hCG  | 46.2 ng/ml         | 1.13                            | Method  |
|   |                    |                                 | CRL Robinson  |
|   |                    |                                 | Scan date   |
|   |                    |                                 | 16/11/2024  |
| Risks at sampling date  |                    |                                 | Trisomy 21  |
| Age risk  |                    | 1:311                           | <b>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</b>  |
| Biochemical T21 risk  |                    | 1:4681                          | 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. |
| Combined trisomy 21 risk  |                    | <1:10000                        | The calculated risk by PRISCA depends on the accuracy of the information provided by the referring physician.   |
| Trisomy 13/18 + NT  |                    | <1:10000                        | Please note that risk calculations are statistical approaches and have no diagnostic value!   |
|   |                    |                                 | The patient combined risk presumes the NT measurement was done according to accepted guidelines (Prenat Diagn 18: 511-523 (1998)).                                    |
|   |                    |                                 | The laboratory can not be hold responsible for their impact on the risk assessment ! Calculated risks have no diagnostic value!                                       |
|   |                    |                                 |   |
| Trisomy 13/18 + NT  |                    |                                 |   |
| <b>The calculated risk for trisomy 13/18 (with nuchal translucency) is &lt; 1:10000, which represents a low risk.</b> |                    |                                 |   |

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**Sign of Physician**