

Prisca 5.1.0.17  
Date of report: 30/01/26

| Patient data   |                    |                                 |                    |
|--|--------------------|---------------------------------|--------------------|
| Name   | Mrs. VAISHALI NAGE | Patient ID                      | 0672601290248      |
| Birthday   | 10/06/97           | Sample ID                       | Mrs. VAISHALI NAGE |
| Age at sample date   | 28.6               | Sample Date                     | 29/01/26           |
| Gestational age  | 13 + 0             |                                 |                    |
| Correction factors   |                    |                                 |                    |
| Fetuses  | 1                  | IVF                             | no                 |
| Weight   | 47                 | diabetes                        | no                 |
| Smoker   | no                 | Origin                          | Asian              |
|  |                    | Previous trisomy 21 pregnancies | unknown            |
| Biochemical data   |                    | Ultrasound data                 |                    |
| Parameter  | Value              | Corr. MoM                       |                    |
| PAPP-A   | 3.66 mIU/mL        | 0.58                            |                    |
| fb-hCG   | 36.95 ng/mL        | 0.91                            |                    |
| Risks at sampling date   |                    |                                 |                    |
| Age risk   | 1:763              | Gestational age                 | 12 + 6             |
| Biochemical T21 risk   | 1:1609             | Method                          | CRL Robinson       |
| Combined trisomy 21 risk   | 1:6942             | Scan date                       | 28/01/26           |
| Trisomy 13/18 + NT   | <1:10000           | Crown rump length in mm         | 67.2               |
|  |                    | Nuchal translucency MoM         | 1.00               |
|  |                    | Nasal bone                      | present            |
|  |                    | Sonographer                     | NA                 |
|  |                    | Qualifications in measuring NT  | Sonographer        |
| 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 6942 women with the same data, there is one woman with a trisomy 21 pregnancy and 6941 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**

below cut off
  Below Cut Off, but above Age Risk
  above cut off