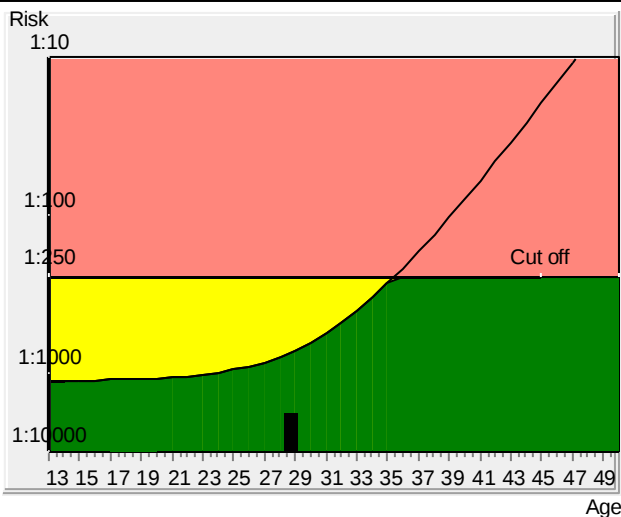


JITM Diagnostics

| Patient data             |             |              |                                |                     |
|--------------------------|-------------|--------------|--------------------------------|---------------------|
| Name                     |             | MRS. SHEETAL | Patient ID                     |                     |
| Birthday                 |             | 23/09/1996   | Sample ID                      |                     |
| Age at sample date       |             | 28.7         | 2506012445/NOD                 |                     |
| Gestational age          |             | 12 + 4       | Sample Date                    |                     |
|                          |             |              | 8/06/2025                      |                     |
| Correction factors       |             |              |                                |                     |
| Fetuses                  | 1           | IVF          | no                             | Previous trisomy 21 |
| Weight                   | 66          | diabetes     | no                             | pregnancies         |
| Smoker                   | no          | Origin       | Asian                          |                     |
| Biochemical data         |             |              | Ultrasound data                |                     |
| Parameter                | Value       | Corr. MoM    | Gestational age                |                     |
| PAPP-A                   | 6.38 mIU/ml | 1.75         | 12 + 3                         |                     |
| fb-hCG                   | 40.6 ng/ml  | 0.94         | Method                         |                     |
| Risks at sampling date   |             |              | CRL Robinson                   |                     |
| Age risk                 |             |              | Scan date                      |                     |
| 1:746                    |             |              | 7/06/2025                      |                     |
| Biochemical T21 risk     |             |              | Crown rump length in mm        |                     |
| <1:10000                 |             |              | 62.7                           |                     |
| Combined trisomy 21 risk |             |              | Nuchal translucency MoM        |                     |
| <1:10000                 |             |              | 1.18                           |                     |
| Trisomy 13/18 + NT       |             |              | Nasal bone                     |                     |
| <1:10000                 |             |              | present                        |                     |
| Risk                     |             |              | Sonographer                    |                     |
|                          |             |              | DR. ANKIT KHANDELWAL MBBS DNB  |                     |
|                          |             |              | Qualifications in measuring NT |                     |
|                          |             |              | MD                             |                     |
|                          |             |              | Trisomy 21                     |                     |



**The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.**

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.

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!

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

**The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.**

Sign of Physician