

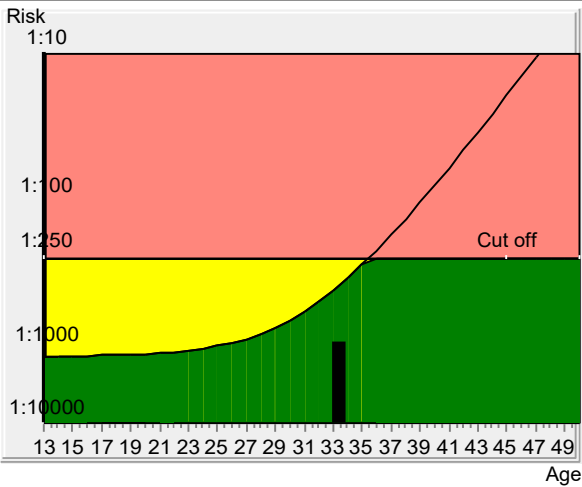
JITM SKILLS PVT LTD
D-87, SECTOR-2, NOIDA (U.P)

Prisca

5.1.0.17

Date of report: 01-05-2023

| Patient data | | | |
|--------------------------|--------------------|---------------------------------|---|
| Name | MRS. NIDHI AGARWAL | Patient ID | |
| Birthday | 25-12-1989 | Sample ID | 2304058372/MJT |
| Age at sample date | 33.3 | Sample Date | 30-04-2023 |
| Gestational age | 12 + 4 | | |
| Correction factors | | | |
| Fetuses | 1 | IVF | no |
| Weight | 71 | diabetes | no |
| Smoker | no | Origin | Asian |
| | | Previous trisomy 21 pregnancies | no |
| Biochemical data | | Ultrasound data | |
| Parameter | Value | Corr. MoM | Gestational age |
| PAPP-A | 1.1 mIU/ml | 0.33 | 11 + 6 |
| fb-hCG | 43.1 ng/ml | 1.02 | Method |
| | | | CRL Robinson |
| | | | Scan date |
| | | | 25-04-2023 |
| Risks at sampling date | | | Crown rump length in mm |
| Age risk | | 1:382 | 53.6 |
| Biochemical T21 risk | | 1:133 | Nuchal translucency MoM |
| Combined trisomy 21 risk | | 1:889 | 0.70 |
| Trisomy 13/18 + NT | | <1:10000 | Nasal bone |
| | | | present |
| | | | Sonographer |
| | | | DR. NAYAB FAKHIR |
| | | | Qualifications in measuring NT |
| | | | M.D |
| | | | 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 889 women with the same data, there is one woman with a trisomy 21 pregnancy and 888 women with not affected pregnancies. |
| | | | The PAPP-A level is low. |
| | | | 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