

KOS DIAGNOSTIC LAB

6349/1, NICHOLSON ROAD, AMBALA CANTT

Prisca

5.1.0.17

Date of report: 19-02-2022

| Patient data | | | |
|--|---------------|---|-----------------------------------|
| Name | MRS. NEHA (B) | Patient ID | 2202220425/AMB (B) |
| Birthday | 01-07-1996 | Sample ID | 2202220425/AMB (B) |
| Age at sample date | 25.6 | Sample Date | 17-02-2022 |
| Gestational age | 11 + 1 | | |
| Correction factors | | | |
| Fetuses | 2 | IVF | yes |
| Weight | 66 | diabetes | no |
| Smoker | no | Origin | Asian |
| | | Previous trisomy 21 pregnancies | no |
| Biochemical data | | Ultrasound data | |
| Parameter | Value | Corr. MoM | |
| PAPP-A | 7.8 mIU/ml | 2.33 | Gestational age 11 + 2 |
| fb-hCG | 123.5 ng/ml | 1.12 | Method CRL Robinson |
| Risks at sampling date | | | Scan date 18-02-2022 |
| Age risk | | 1:885 | Crown rump length in mm 47 |
| Biochemical T21 risk | | <1:10000 | Nuchal translucency MoM 0.78 |
| Combined trisomy 21 risk | | <1:10000 | Nasal bone present |
| Trisomy 13/18 + NT | | <1:10000 | Sonographer . |
| | | | Qualifications in measuring NT MD |
| Risk | | Trisomy 21 | |
| | | <p>The calculated risk for Trisomy 21 (with nuchal translucency) is below the cut off, which indicates a low risk.</p> <p>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.</p> <p>The risk for this twin pregnancy has been calculated for a singleton pregnancy with corrected MoMs.</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>The calculated risk for trisomy 13/18 (with nuchal translucency) is < 1:10000, which represents a low risk.</p> | | | |

Sign of Physician

below cut off

Below Cut Off, but above Age Risk

above cut off