Prisca 5.2.0.13

Date of report: 11/06/2024

| Patient data | | | | | |
|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|---------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|------------------------------------|----------------|--|
| Name | MRS. PRIYA | | | | |
| Birthday | 5/05/1991 | | | 2406018437/NOD | |
| Age at sample date | 33.1 | | Sample Date 10/06/2024 | | |
| Gestational age | 12 + 6 | | | | |
| Correction factors | | | | | |
| Fetuses 1 | IVF | no | Previous trisomy 21 | no | |
| Weight 54 | diabetes | no | pregnancies | | |
| Smoker no | Origin | Asian | | | |
| Biochemical data | | Ultrasound da | ata | | |
| Parameter Value | Corr. MoM | Gestational | age | 12 + 6 | |
| PAPP-A 1.18 mIU/m | 0.23 | Method CRL Robinson | | | |
| fb-hCG 41.1 ng/ml | 0.92 | Scan date 10/06/2024 | | | |
| Risks at sampling date | | | Crown rump length in mm 67 | | |
| Age risk | 1:404 | | Nuchal translucency MoM 0.76 | | |
| Biochemical T21 risk | ochemical T21 risk 1:54 | | Nasal bone presen | | |
| Combined trisomy 21 risk 1:398 | | Sonographer DR. NEERJA CHOPRA | | | |
| Trisomy 13/18 + NT | Trisomy 13/18 + NT 1:1628 | | Qualifications in measuring NT M.D | | |
| Risk 1:10 | Trisomy 21 | - | | | |
| 1:1000 1:10000 1:10000 1:10000 1:10000 1:10000 Age Trisomy 13/18 + NT The calculated risk for Trisomy 13/18 (with nuchal translucency) is 1:1628, which represents a low risk. | | 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 398 women with the same data, there is one woman with a trisomy 21 pregnancy and 397 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! | | | |