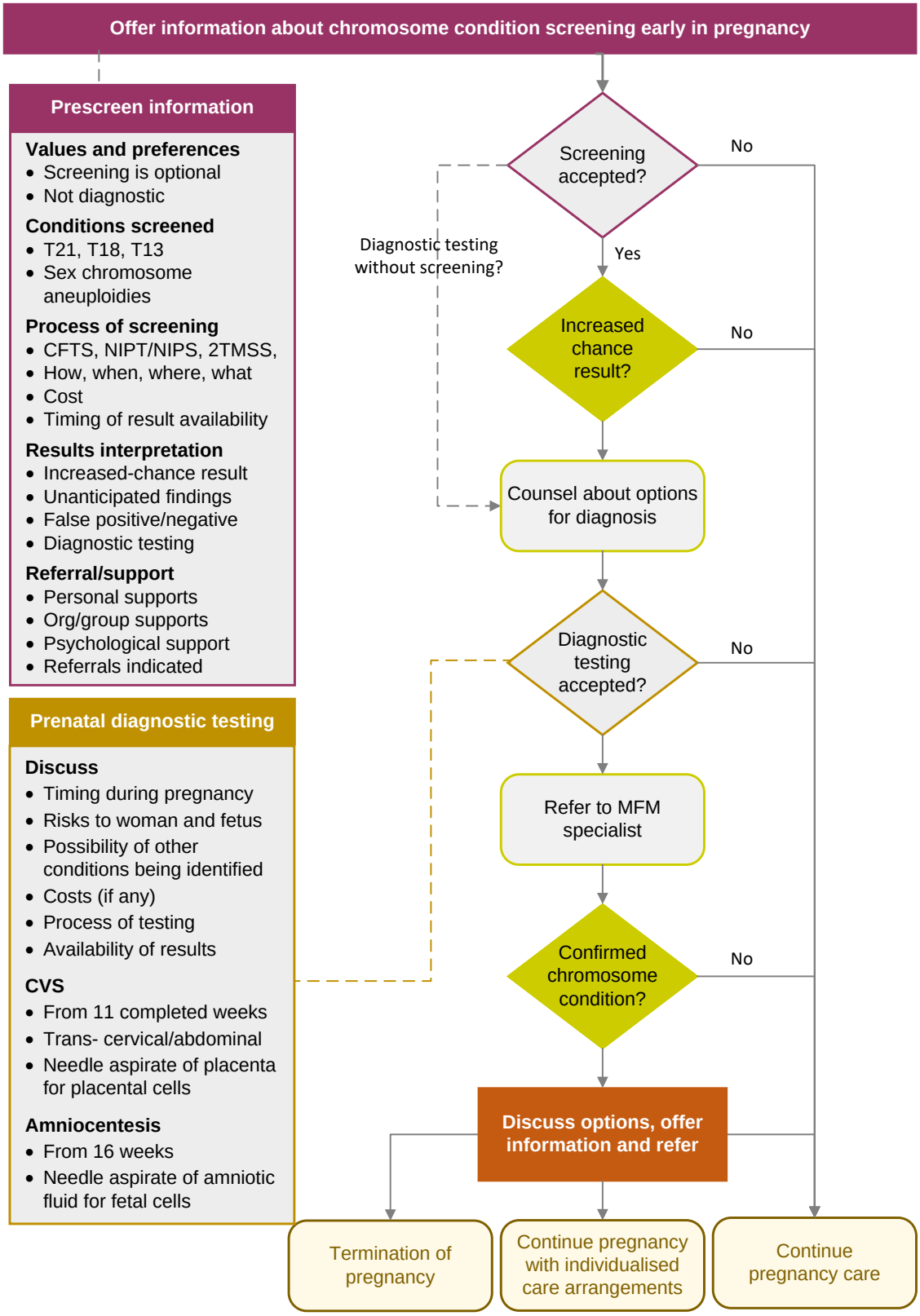
Chromosome condition screening during pregnancy



2TMSS: second trimester maternal serum screening, **CF:** cystic fibrosis, CFTS: combined first trimester screening, **CVS:** chorionic villus sampling, **FXS:** fragile X syndrome, **MFM:** maternal fetal medicine, **NIPT/NIPS**: non-invasive prenatal screening test, **NPV:** negative predictive value, **Org:** organisation, **PPV:** positive predictive value, **RGCS:** reproductive genetic carrier screen, **SMA:** spinal muscular atrophy, **T13:** Trisomy 13 (Patau syndrome) **T18:** Trisomy 18 (Edwards syndrome) **T21:** Trisomy 21 (Down syndrome)
Individualised care arrangements: may include (as relevant to individual circumstances) increased antenatal surveillance, increased ultrasound surveillance, preparation for palliative care, kinship or formal adoption, or foster care

Queensland Clinical Guideline. Preconception and prenatal genetic screening. Flowchart F24.36-3-V1-R29