### Diagnostic Criteria

Patient has…
- [ ] DiGeorge defined genetically ONLY
- [ ] Classic clinical triad for complete DiGeorge syndrome
- [ ] Combination of genetic and phenotypic features

#### Defined Genetically Only

Select all that apply
- [ ] Chromosome 22q11.2 deletion
- [ ] TBX mutation
- [ ] Chromosome 10p13-14 deletion
- [ ] CHD7 mutation

#### Classical Clinical Triad

Select all that apply
- [ ] Conotruncal cardiac anomaly
- [ ] Hypocalcemia
- [ ] T cells < 500 /mm3 in the first 3 months of life

#### Combination of Features

Select all that apply
- [ ] Chromosome 22q11.2 deletion
- [ ] Conotruncal cardiac anomaly or cardiothoracic vascular anomaly
- [ ] Hypocalcemia
- [ ] Diminished T cell counts for age
- [ ] Hypoplastic thymus visualized
- [ ] Dysmorphic facies
- [ ] Tracheoesophageal fistula
- [ ] Coloboma Cleft palate (frank clefting or submucous cleft)
- [ ] Velopharyngeal insufficiency / hypernasal speech

### Other Questions

Fetal toxin exposure?
- [ ] Yes
- [ ] No
- [ ] Unknown

(if yes) Explain:
- [ ] Maternal diabetes
- [ ] Alcohol exposure
- [ ] Isotretinoin exposure
- [ ] Other: ______________________________