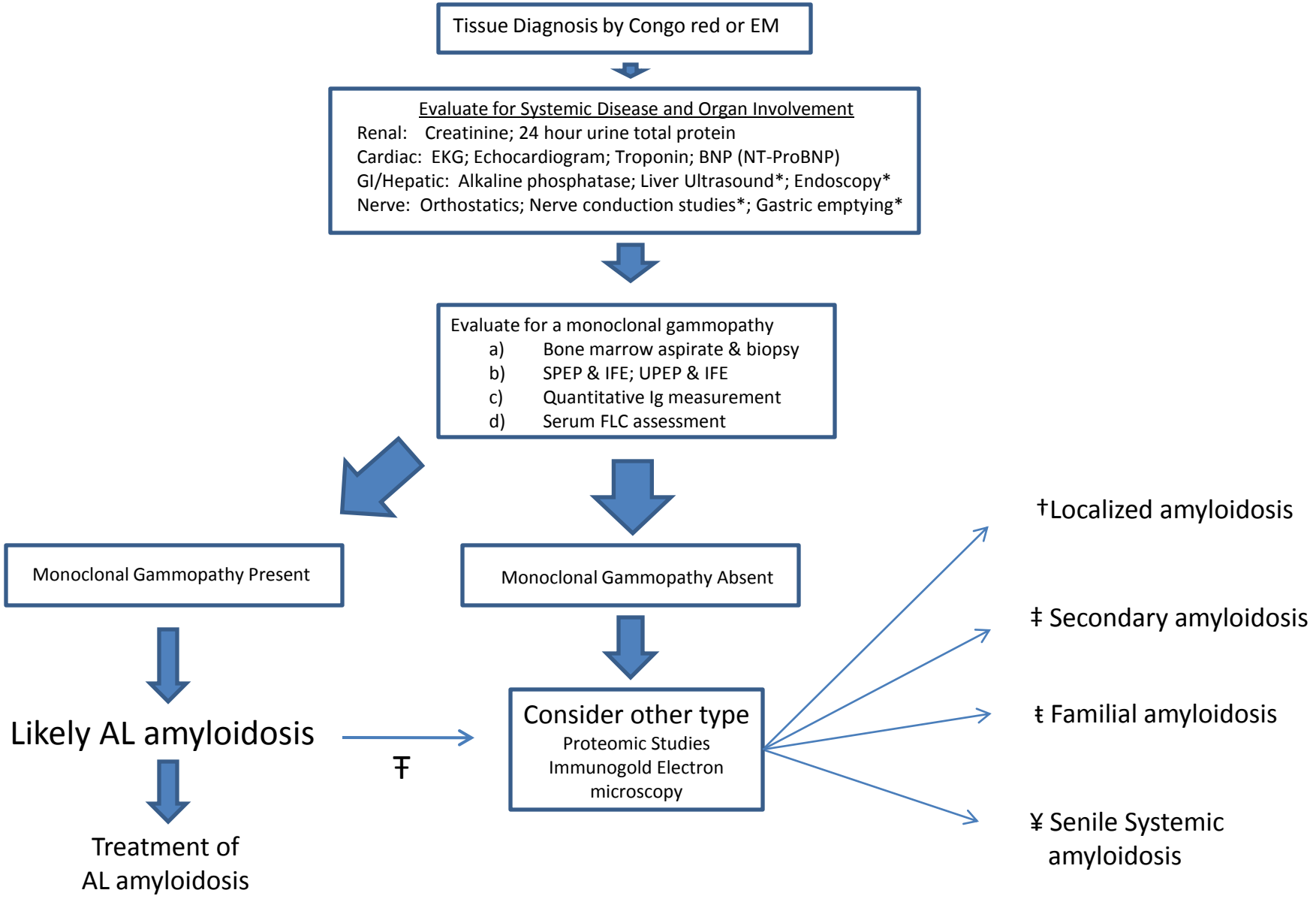

* If clinically indicated

Ŧ Indications for familial testing with or w/out a monoclonal gammopathy[7]

1. African Americans for mutant TTR
2. Dominant peripheral nervous system involvement
3. Isolated renal involvement and no amyloid in bone marrow; screen for fibrinogen Aα
4. Family history concerning for familial amyloidosis

†: kappa or lambda precursor protein; refer to system specific specialist

‡: Consider when clinically indicated i.e. chronic inflammatory condition

ŧ : Most likely transthyretin (TTR) mutation but consider other hereditary forms

¥ : Wild type TTR; often isolated cardiac involvement