"unspecified" Heritable Disorder of Connective Tissue

- Marfan Syndrome
- Loeys –Dietz Syndrome
- Ehlers Danlos Syndromes
- Osteogenesis imperfecta

Ophthamology Assessment

- Cardiology review with echocardiogram

(Re) Consider the underlying Genetic Diagnosis

- Assessment of bone health

Hypermobile child
Beighton ≥ 4/9

- Review clinical progress and evolution of phenotype
  - Annually to biannually

Check for functional symptoms of "joint hypermobility syndrome" and formulate a management plan

- Anxiety
- Disabling Fatigue
- Joint instability episodes
- Soft tissue injuries
- Widespread musculoskeletal pain
- Osteopenia