

1703 patients with  
hypertrophic  
phenotype  
cardiomyopathy

N excluded:  
- 6 patients as EF not available

Idiopathic or  
sarcomeric HCM  
n= 1298

Phenocopies  
n= 408

Genotype negative  
n= 531

Genotype positive  
n= 483

Not genetically tested  
n= 274

AL amyloidosis: n= 115  
Hereditary TTR amyloidosis: n= 66  
Wild-type or SSA: n= 48

AFD: n= 85  
CPT II deficiency: n= 1

GSD: n= 16\*

Mitochondrial diseases: n= 23

Noonan syndrome: n= 15  
LEOPARD syndrome: n= 7

Friedreich's ataxia: n= 11  
PFL1: n= 2