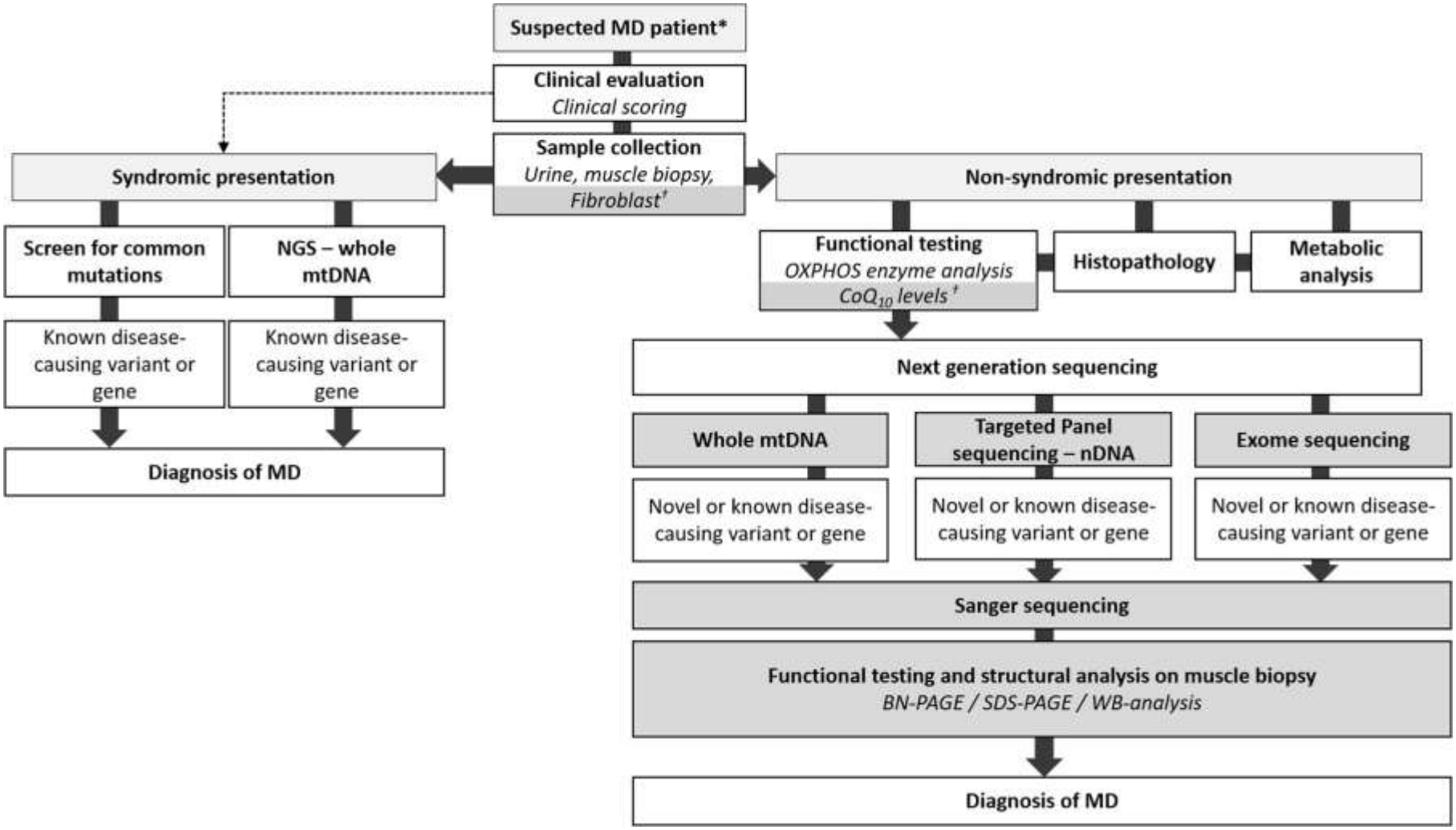

Supplemental Figure S1. *Diagnostic procedures followed in patients clinically suspected to have mitochondrial diseases (MDs). *Suggestive family history and suggestive signs and symptoms. †Procedures that are not routinely performed. BN-PAGE, blue native–polyacrylamide gel electrophoresis; CoQ_{10}, coenzyme Q_{10}; mtDNA, mitochondrial DNA; nDNA, nuclear DNA; NGS, next-generation sequencing; OXPHOS, oxidative phosphorylation; WB, Western blot analysis.*