

Supplementary Table. Overview of the clinical presentation in our patients. Age is provided in years. UL, upper limbs; LL, lower limbs; M, males; F, females; CN, unpublished candidate novel gene; WES, whole exome sequencing; FRDA, Friedreich ataxia; SCA expansion, dominant spinocerebellar atrophy due to repeats expansion; HSP panel, hereditary spastic paraparesis gene panel; no cosegreg; no cosegregation of the variant (absent in the patient); 0: sign absent; y:sign present; NA: not available/not assessed