

Supplemental Data

Mutations in *C12orf65*

in Patients with Encephalomyopathy

and a Mitochondrial Translation Defect

Hana Antonicka, Elsebet Østergaard, Florin Sasarman, Woranontee Weraarpachai, Flemming Wibrand, Anne Marie B. Pedersen, Richard J. Rodenburg, Marjo S. van der Knaap, Jan A.M. Smeitink, Zofia M. Chrzanowska-Lightowlers, and Eric A. Shoubridge

Table S1. Homozygous Regions Larger than 5 Mb in the DNA of Patient 1

Chromosome	SNPs	Interval (Mb)
1	rs1347539 - rs1414904	76.69–95.34
1	rs10495094 - rs1570105	214.51–230.74
1	rs1072320 - rs1892445	236.31–qter
3	rs1112189 - rs1421165	144.51–179.11
6	rs6911727 - rs449159	9.06–14.97
8	rs10488368 - rs10503691	Pter–20.67
11	rs1486911 - rs10502234	110.26–119.41
12	rs1106349 - rs10492049	117.35–125.52
15	rs1532517 - rs1672407	22.08–29.23
16	rs10500362 - rs1898680	8.14–13.18
16	rs10521238 - rs10514487	50.12–78.74
17	rs10521195 - rs63953	12.10–28.68
20	rs4142363 - rs4811374	10.98–50.71
21	rs2835256 - rs10483083	36.34–qter