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Supplemental Data

Mutations in *TPRN* Cause a Progressive Form of Autosomal-Recessive Nonsyndromic Hearing Loss

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Table S1. Genes Located in the Critical Region and Tested by Direct Sequencing

Gene	Position	No. coding exons
OLFM1	137967103-137989951	7
CI116	138387026-138391761	3
MRPS2	138392483-138396518	4
PAEP	138453604-138458622	7
GLT6D1	138515502-138531386	4
SOHLH1	138585257-138591374	7
KCNT1	138594031-138684992	31
CAMSAP1	138700334-138799005	17
UBADC1	138824816-138853226	10
BTBD14A	138903264-138942370	5
LHX3	139088098-139096955	6
QSOX2	139098182-139137687	12
GPSM1	139221932-139236287	9
DNLZ	139256352-139258241	3
CARD9	139261698-139268133	13
SNAPC4	139270030-139292889	22
PMPCA	139305116-139318212	13
INPP5E	139323075-139334256	10
SEC16A	139334549-139377507	32
EGFL7	139557382-139567129	8
AGPAT2	139567595-139581911	6
FAM69B	139607024-139619169	5
TMEM141	139685777-139687768	5
C9orf86	139702381-139735639	15
PHPT1	139743256-139745488	4
MAMDC4	139746819-139755249	27
EDF1	139757233-139760738	4
TRAF2	139780965-139821066	10
FBXW5	139834888-139839173	8
C9orf141	not listed in UCSC anymore	4
PTGDS	139871956-139876193	6
C9orf142	139886870-139888427	7
CLIC3	139889060-139891024	6
ABCA2	139901686-139922740	49
C9orf139	139921916-139931234	2
FUT7	139924626-139927292	2

NPDC1	139933909-139940676	9
ENTPD2	139942553-139948505	9
C9orf140	139956581-139965028	6
UAP1L1	139971953-139978989	9
MAN1B1	139997800-140003639	13
DPP7	140004993-140009195	13
GRIN1	140033609-140063207	20
LRRC26	140063212-140064491	2
SSNA1	140083054-140084822	3
TPRN (C9orf75)	140086069-140094980	3
TMEM203	140098537-140100090	1
NDOR1	140100119-140113812	14
RNF208	140114707-140115775	1
SLC34A3	140126113-140131006	12
TUBB2C	140135711-140138159	4
COBRA1	140149759-140167999	13
C9orf167	140172280-140177092	1
NOXA1	140317847-140328857	14
ENTPD8	140328817-140335901	9
MRPL41	140446309-140447006	1
WDR85	140449361-140473387	11
ZMYND19	140476531-140484937	6
ARRDC1	140500096-140509811	8
NP_116326	140509785-140513308	1
EHMT1	140513444-140730576	27
CACNA1B	140772241-141019075	46

Genomic positions of genes are given in accordance to Human GRCh37 (hg19) of the UCSC Genome Browser.