



HAL
open science

Premature termination codons in SOD1 causing Amyotrophic Lateral Sclerosis are predicted to escape the nonsense-mediated mRNA decay

Claire Guissart, Kevin Mouzat, Jovana Kantar, Baptiste Louveau, Paul Vilquin, Anne Polge, Cédric Raoul, Serge Lumbroso

► To cite this version:

Claire Guissart, Kevin Mouzat, Jovana Kantar, Baptiste Louveau, Paul Vilquin, et al.. Premature termination codons in SOD1 causing Amyotrophic Lateral Sclerosis are predicted to escape the nonsense-mediated mRNA decay. *Scientific Reports*, 2020, 10 (1), pp.20738. 10.1038/s41598-020-77716-5 . hal-04767217

HAL Id: hal-04767217

<https://hal.science/hal-04767217v1>

Submitted on 5 Nov 2024

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Distributed under a Creative Commons Attribution 4.0 International License



OPEN

Premature termination codons in *SOD1* causing Amyotrophic Lateral Sclerosis are predicted to escape the nonsense-mediated mRNA decay

Claire Guissart^{1,2}✉, Kevin Mouzat^{1,2}, Jovana Kantar^{1,2}, Baptiste Louveau³, Paul Vilquin³, Anne Polge¹, Cédric Raoul² & Serge Lumbroso^{1,2}

Amyotrophic lateral sclerosis (ALS) is the most common and severe adult-onset motoneuron disease and has currently no effective therapy. Approximately 20% of familial ALS cases are caused by dominantly-inherited mutations in the gene encoding Cu/Zn superoxide dismutase (*SOD1*), which represents one of the most frequent genetic cause of ALS. Despite the overwhelming majority of ALS-causing missense mutations in *SOD1*, a minority of premature termination codons (PTCs) have been identified. mRNA harboring PTCs are known to be rapidly degraded by nonsense-mediated mRNA decay (NMD), which limits the production of truncated proteins. The rules of NMD surveillance varying with PTC location in mRNA, we analyzed the localization of PTCs in *SOD1* mRNA to evaluate whether or not those PTCs can be triggered to degradation by the NMD pathway. Our study shows that all pathogenic PTCs described in *SOD1* so far can theoretically escape the NMD, resulting in the production of truncated protein. This finding supports the hypothesis that haploinsufficiency is not an underlying mechanism of *SOD1* mutant-associated ALS and suggests that PTCs found in the regions that trigger NMD are not pathogenic. Such a consideration is particularly important since the availability of *SOD1* antisense strategies, in view of variant treatment assignment.

Amyotrophic lateral sclerosis (ALS) is a fatal neurodegenerative disease characterized by the selective loss of both upper and lower motoneurons, leading to a progressive paralysis and death within 3–5 years¹.

About 20% of familial ALS cases are caused by mutations in the gene encoding the detoxifying copper-zinc superoxide dismutase (*SOD1*)². Currently, over 180 different mutations throughout the five exons of the *SOD1* gene (MIM 147450) have been described^{2,3}, the vast majority of which being missense point mutations resulting in a dominant mode of inheritance of ALS (with the exception of the D91A mutation) and spreading over the entire 154 amino acid sequence^{4,5}.

It has been well-established that *SOD1* mutants-mediated toxicity is caused by a gain-of-function rather than the loss of the detoxifying activity of *SOD1*² and that mutant *SOD1* can adopt multiple misfolded conformations that mediate toxicity². Moreover, mice with genetic ablation of *Sod1* do not recapitulate disease phenotype^{5–8}. Instead, *Sod1*-deficient mice show accelerated rate of muscle denervation, locomotor deficits and tremors, as well as increased vulnerability to stress. It is noteworthy that the 50% loss of *Sod1* activity described in heterozygous *Sod1*^{+/-} mice leads to an increased susceptibility to axonal injury, ischemia or glutamate-induced toxicity⁹.

Nonsense-mediated mRNA decay (NMD) is an eukaryotic quality control pathway that degrades mRNAs containing Premature termination codons (PTCs) caused by nonsense or frameshift mutations^{10,11}. It is important to note that some PTCs can escape NMD. This capability is governed by four rules¹²: (1) the 50 nucleotides rule: PTCs less than 50–55 nucleotides upstream of the last exon–exon junction typically do not trigger NMD; (2) the last exon rule: PTCs in the last exon of a gene also do not trigger NMD; (3) the long exon rule: exons

¹Laboratoire de Biochimie et Biologie Moléculaire, CHU Nîmes, University of Montpellier, Nîmes, France. ²The Neuroscience Institute of Montpellier, INM, INSERM, Univ Montpellier, Montpellier, France. ³Département de Pharmacologie et de Génomique des Tumeurs Solides, Hôpital Saint Louis, Assistance Publique Hôpitaux de Paris, Paris, France. ✉email: claire.guissart@inserm.fr

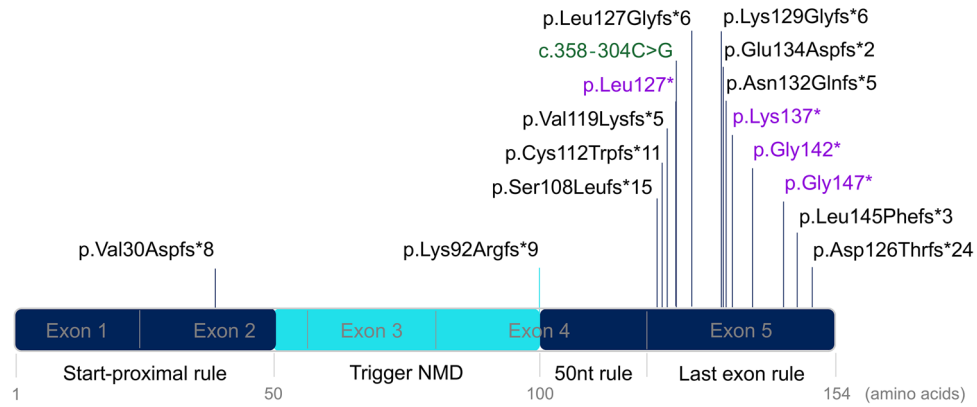


Figure 1. Schematic representation of the *SOD1* gene showing the localization of PTCs causing ALS and NMD escaping regions. Frameshift mutations and the deep intronic splicing mutation are placed according to the localization of their resulting PTCs (and not the mutation site itself). Nonsense mutations are indicated in purple, deep intronic splicing mutation in green and frameshift mutations in black. The region of *SOD1* where PTCs trigger NMD is represented in cyan. The regions that escape NMD are represented in dark blue.

greater than approximately 400 nucleotides inhibit NMD; (4) the start-proximal rule: PTCs located below 150 nucleotides from the start codon typically fail to trigger NMD.

PTCs that escape NMD in *SOD1* are thus expected to lead to the production of truncated *SOD1* protein, which can be highly unstructured with elevated toxicity as illustrated with the non-sense mutation p.Leu127*^{13,14}.

To evaluate the capacity of PTCs associated with ALS in *SOD1* to trigger NMD, we analyzed their localization through the gene.

Results

Regarding the NMD rules we estimated that the region of *SOD1* obeying the NMD is located between nucleotides 151–301, which correspond to amino acids 50–100 (Fig. 1). Accordingly, all PTCs located within this region are expected to trigger the NMD, resulting in the degradation of messenger RNAs, and leading to haploinsufficiency. Conversely, all PTCs located outside this region can result in the production of truncated *SOD1*, prone to induce misfolded protein. We found a total of 16 disease-associated-PTCs mutations in *SOD1* in the literature^{15,16}, including 4 nonsense mutations, 11 frameshift mutations and 1 deep intronic splicing mutation (Table 1, Fig. 1).

Among them, fourteen are predicted to escape the NMD according to the last exon rule and one (p.Val30Aspfs*8) is predicted to escape the NMD, obeying to the start proximal rule (Fig. 1). By faithfully following the 50 nucleotides rule, one frameshift mutation (p.Lys92Argfs*9) was found to introduce a PTC at the position 100, which is the last position predicted to trigger NMD. However, it is likely that at such borderline position, NMD is not completely activated and that truncated protein is at least partly produced.

To evaluate the impact of NMD on PTCs across the *SOD1* gene, we have analyzed experimental data from a large-scale analysis of approximately 80,000 matched tumor exomes and transcriptomes available on the cBio Cancer Genomics Portal^{17,18}. Among them, we identified 31 mutations in *SOD1* in 40 samples. 24 were missense mutations, 2 were translation start site mutations and 5 were truncating mutations (2 splicing, 2 frameshift and 1 non-sense mutation). The heterozygous non-sense mutation (p.Glu79*, E79*) identified in one sample by whole exome sequencing (WES) was almost not detected on the RNA sequencing (RNA-Seq) data from the same sample, suggesting a massive degradation of the *SOD1* mutated transcript by the NMD (Fig. 2). In contrast, heterozygous frameshift mutation (p.Lys137Aspfs*26, K137Dfs*26), located in the last exon of *SOD1*, was detected in both WES and RNA-Seq data, confirming NMD escape for this other PTC (Fig. 2).

Discussion

In this study, we have explored the impact of NMD on *SOD1* and shown that the activity of the NMD pathway is of broad importance for ALS caused by PTC in *SOD1*. Through a large-scale analysis of human cancer exomes and transcriptomes we were able to confirm that *SOD1* standardly obeys to the NMD pathway and its rules.

Our conclusion is supported by several arguments from the literature: (1) The presence of PTC in a region that escape NMD has been detected in mRNA extracted from immortalized lymphoblast cell lines from two patients harboring the c.358-304C>G mutation, thus confirming the impact on the protein level p.Val120Glnfs*8¹⁹; (2) Conformational changes of truncated proteins have been well characterized for PTC located in the region that escape NMD²⁰; (3) Heterozygous PTCs located in the region that trigger NMD seem to be more frequent in individuals from general population: in gnomAD database, for example, we found 6 individuals with a PTC theoretically triggering NMD (p.Glu50Glyfs*39, p.Leu68Gluys*19) versus 3 individuals aged between 40 and 65 years with a PTC theoretically escaping NMD (p.Val6Cysfs*4, p.Asp97Metfs*8).

Interestingly, the fact that all PTCs associated with ALS in *SOD1* can escape the NMD comforts the hypothesis that haploinsufficiency is not an underlying mechanism of the disease. Instead, the production of a misfolded truncated *SOD1* protein could cause a toxic gain-of-function. Therefore, even if we confirmed the massive

Mutation (GRCh37) ^a	Protein variant ^b	Resulting PTC position	References
Splicing mutation			
c.358-304C>G	p.Val120Glnfs*8	127	19
Non-sense mutations			
c.380 T>A	p.Leu127*	127	16
c.409A>T	p.Lys137*	137	15
c.424G>T	p.Gly142*	142	36
c.441 T>A	p.Cys147*	147	20,37
Frameshift mutations			
c.88_89insA	p.Val30Aspfs*8	37	20,38,39
c.275_276delAA	p.Lys92Argfs*9	100	40
c.320dupT	p.Ser108Leufs*15	122	41
c.335dupG	p.Cys112Trpfs*11	122	28,29
c.355delGinsAAAAC	p.Val119Lysfs*5	123	20,42
c.379_380delTT	p.Leu127Glyfs*6	132	43–45
c.380_383dupTGGG	p.Lys129Glyfs*6	134	20,46,47
c.401_402insTT	p.Glu134Aspfs*2	135	20,48
c.383_392dupGCAAAGGTGG	p.Asn132Glnfs*5	136	49
c.435delGinsCGTTTA	p.Leu145Phefs*3	147	50
c.376delG	p.Asp126Thrfs*24	149	51

Table 1. List of truncating mutations in *SOD1* associated with ALS. ^a Human genome variation society (HGVS) nomenclature V2.0 according to mRNA reference sequence GenBank: NM_000454.4. Nucleotide numbering uses + 1 as the A of the ATG translation initiation codon in the reference sequence, with the initiation codon as codon 1. ^b HGVS nomenclature according to protein reference sequence GenPept: NP_000445.1. Amino acid numbering uses p.1 as the Methionine corresponding to the initiation codon. This implies a 1-amino acid switch compared to former *SOD1* nomenclature (eg. L127X mutation was formerly known as L126X or, in some articles, L126Z).

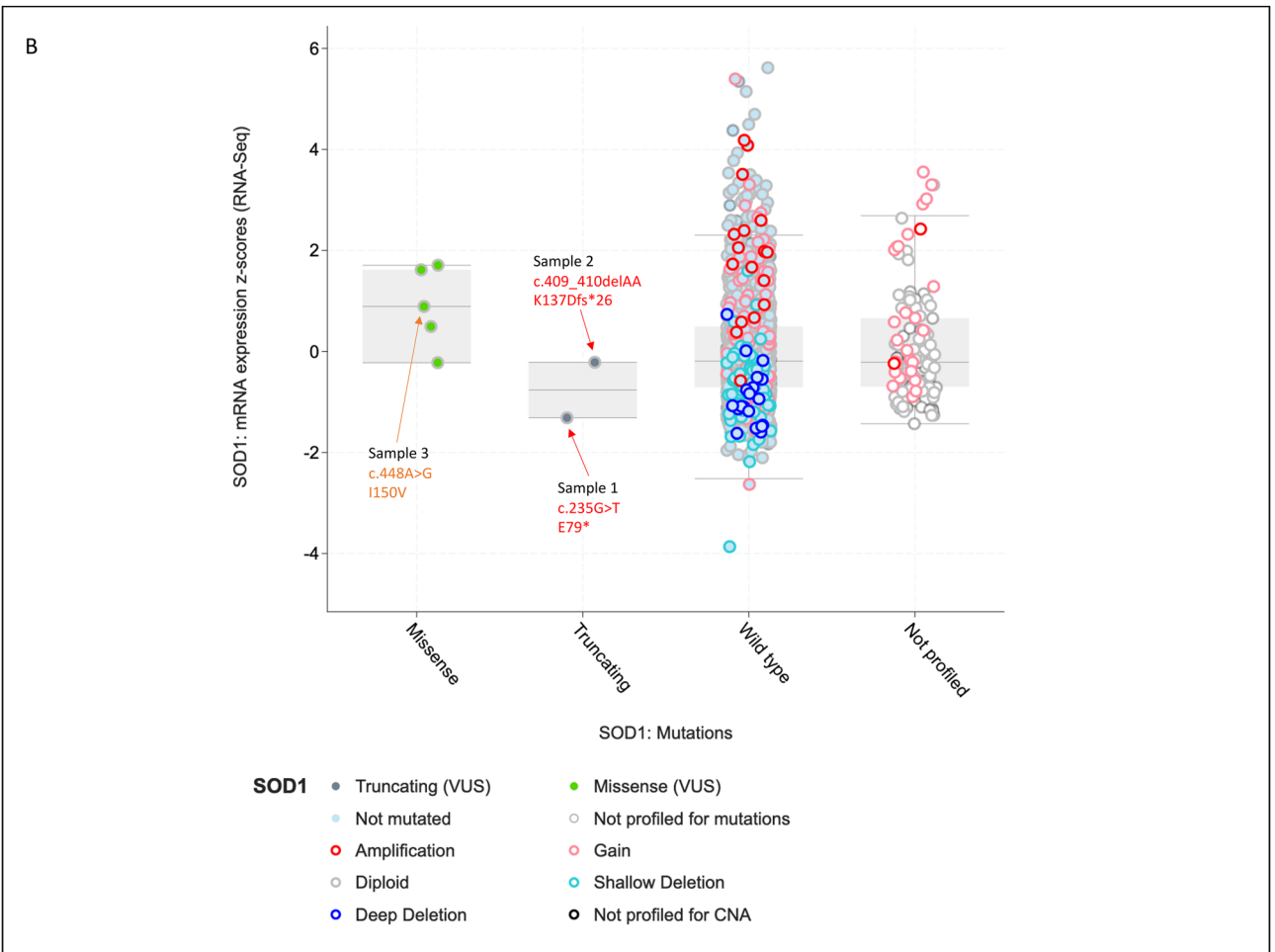
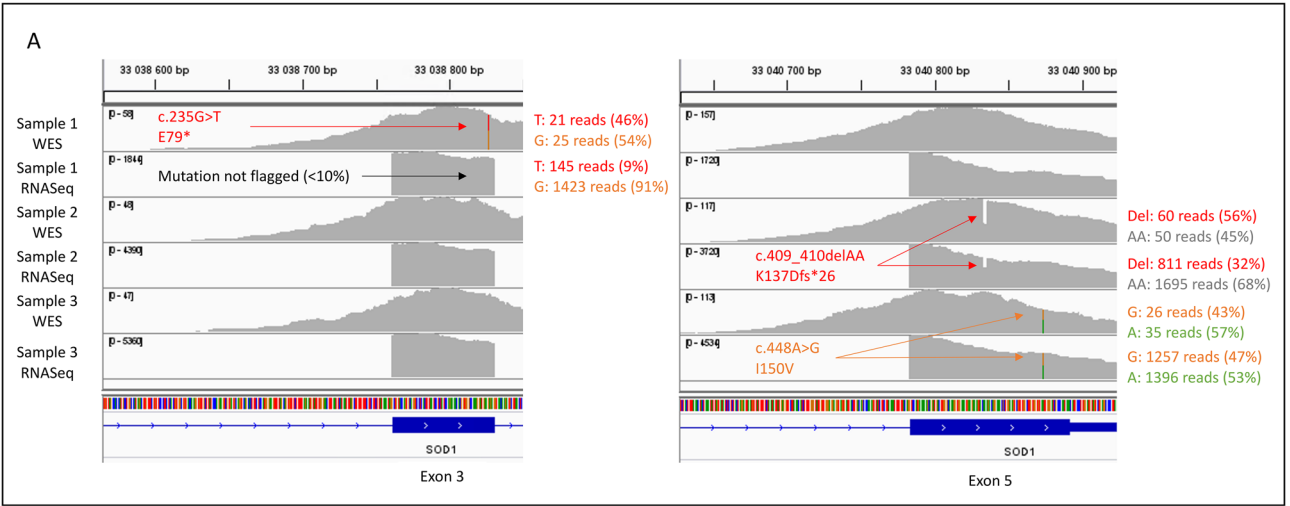
degradation of mRNA harboring a PTC in the region triggering NMD, as we could detect a small remaining amount of mutated mRNA (9% of reads on RNA-Seq data, see sample 1 on Fig. 2A), we cannot exclude a very late onset form of SLA in such situation.

Dimer destabilization, oligomerization and increased aggregation are the proposed mechanisms for mutant *SOD1* toxicity⁴. Recently it has been demonstrated that *SOD1* acts as a H₂O₂-responsive regulatory protein in the expression of ALS-linked genes. Both sequence preference and affinity of *SOD1* interactions with DNA depend on *SOD1* conformation²¹. Thereby, PTCs that escape NMD in *SOD1* are expected to cause toxic conformational changes. Indeed, some of the truncating mutations described here were proven to cause *SOD1* misfolding capable to interact with Derlin-1, triggering endoplasmic reticulum stress and contributing to motoneuron death (*i.e.* p.Val30Aspfs*8, p.Val119Lysfs*5, p.Leu127*, p.Glu134Aspfs*2, p.Gly142*)²⁰.

The toxic gain-of-function mechanism evidence provides a strong rationale for gene silencing as a therapy for *SOD1*-mediated ALS. Thus, clinically promising therapies, all aimed at enhancing specifically the degradation of the mutated *SOD1* RNA, such as anti-sense oligonucleotides (ASO) and RNA interference (RNAi) are being tested in preclinical and clinical studies^{22–25}. In the first clinical trial of ASO treatment in human beings, only ASO targeting missense mutations were developed. This trial had favorable safety outcomes, and a trial to assess the safety, tolerability and pharmacokinetics of a second generation *SOD1* ASO is currently in progress (ClinicalTrials.gov, NTC02623699)²⁶. To our knowledge, no ASO targeting a PTC has been investigated so far. This could be explained by the low proportion of patients carrying such mutations. For example, PTCs in *SOD1* are absent from MinE Database which includes 4366 whole genomes from ALS patients and 1832 whole genomes from controls, from different European ancestry²⁷.

Recently, complete loss of function of *SOD1* in human has been reported in a 2 years old girl with a homozygous truncating mutation and an absence of *SOD1* activity. The patient presented with axial hypotonia and loss of gross and fine motor function at 6 months of age, after which severe, progressive spastic tetraparesis developed and Babinski's sign was present in both feet. Atrophy, fasciculations, and other signs of lower motor neuron involvement were not noted. Her parents, both heterozygous for the mutation were healthy at the time of the report while the level of *SOD1* activity was half that of the normal level²⁸. Another report of the same homozygous truncating variant c.335dupG (p.Cys112Trpfs*11) in *SOD1* was identified in another patient with tetraspasticity. In contrast with Andersen et al. 2019 report, heterozygous carriers from this family had a markedly reduced enzyme activity when compared to wild-type controls but show no overt neurologic phenotype²⁹. Thus, while caution might be exercised regarding the use of gene therapies that may markedly depresses *SOD1* activity, reduction of *SOD1* appears to be well tolerated, as outlined by the favorable clinical trial safety outcomes.

Animal models, particularly *SOD1* rodent model³⁰, initially developed to investigate the complex processes occurring in ALS, had played a major role in performance evaluation of these silencing approaches²². More recently, other models like zebrafish³¹, *Drosophila*³² or patient-derived induced pluripotent stem cell³³ have also



◀**Figure 2.** (A) Evaluation of NMD effect on *SOD1* transcripts by comparison of matched tumor exomes and transcriptomes. Visualization with the Integrative Genome Viewer software (version 2.8.4, <http://software.broadinstitute.org/software/igv/>)³⁵ of WES and RNA-Seq alignments of 3 samples from the Cancer Cell Line Encyclopedia⁵². The missense mutation p.Ile150Val (I150V, sample 3) is used here as a control as it has no effect on NMD activity and is thus found heterozygous in both WES and RNA-Seq alignments. The non-sense mutation p.Glu79* located in exon 3 of *SOD1*, (E79*, sample 1) was detected on 9% of RNA-Seq reads versus 46% of WES reads, showing the degradation of the *SOD1* mutated transcript by NMD. In contrast, heterozygous frameshift mutation p.Lys137Aspfs*26 (K137Dfs*26, sample 2), located in the last exon of *SOD1* was detected in 32% of RNA-Seq reads versus 56% of WES reads, highlighting NMD escape. (B) *SOD1* mRNA expression correlation with *SOD1* mutations in 2029 samples. This plot was generated from the cBio Cancer Genomics Portal (<http://cbioportal.org>)^{17,18}. Although not statistically significant, sample 1 with the E79* mutation appears to have a lower *SOD1* mRNA expression (z -score = -1.31) compared to sample 2 with the K137Dfs*26 mutation (z -score = -0.21 , equivalent to the mean mRNA expression in the wild type group). Deep Deletion indicates a deep loss, possibly a homozygous deletion; Shallow Deletion indicates a shallow loss, possibly a heterozygous deletion; Gain indicates a low-level gain (a few additional copies, often broad); Amplification indicate a high-level amplification (more copies, often focal); Not profiled for CNA indicate the samples for which copy-number analysis was not performed. These levels are derived from copy-number analysis algorithms and indicate the copy-number level per gene.

been designed and tested to investigate the physiopathology of ALS. These models, particularly useful since the availability of *SOD1* antisense strategies, offer the possibility to study the pathogenicity of novel *SOD1* variants, especially complex intronic mutations that could either lead to an amino-acid(s) insertion or deletion and/or to the creation of a PTC.

In conclusion, we highlight that all described PTCs in *SOD1* causing ALS are predicted to escape the non-sense-mediated mRNA decay. More importantly, this observation suggests that truncating mutations found in the region of *SOD1* that trigger NMD may have no pathogenic significance. Such a consideration is particularly important since the availability of *SOD1* antisense strategies, in view of variant treatment assignment.

Methods

The PTCs of the human *SOD1* gene (NM_000454.4) resulting from nonsense, frameshift and splicing mutations that are associated with ALS were obtained from the Human gene mutation database (HGMD)³⁴, which provides systematic and in-depth qualitative and quantitative overviews of genetic research in both familial and sporadic ALS. Intronic mutations located outside the canonical sites and not confirmed by transcript analysis were excluded from this study.

The cBio Cancer Genomics Portal (<http://cbioportal.org>)^{17,18}, an open platform for exploring multidimensional cancer genomics data, was used to select tumor samples with PTC in *SOD1* for which whole exome sequencing (WES) and RNA-sequencing (RNA-seq) experimental data were performed. We generated plot from cBio Cancer Genomics Portal^{17,18} to analyze *SOD1* mRNA expression correlation with *SOD1* mutations.

Raw data (Bam files) from sample of interest were downloaded, when available, from NCBI Sequence Read Archive (SRA) and visualized with the Integrative Genome Viewer software (version 2.8.4, <http://software.broadinstitute.org/software/igv/>)³⁵. SRA accession numbers and detailed information about the samples are available in the Supplementary Table S1.

Received: 20 July 2020; Accepted: 12 November 2020

Published online: 26 November 2020

References

- Kiernan, M. C. *et al.* Amyotrophic lateral sclerosis. *Lancet (London, England)* **377**, 942–955 (2011).
- Huai, J. & Zhang, Z. Structural properties and interaction partners of familial ALS-associated *SOD1* mutants. *Front. Neurol.* **10**, 527 (2019).
- Broom, H. R., Rumpf, J. A. O. & Meiering, E. M. Many roads lead to Rome? Multiple modes of Cu, Zn superoxide dismutase destabilization, misfolding and aggregation in amyotrophic lateral sclerosis. *Essays Biochem.* **56**, 149–165 (2014).
- Pansarasa, O. *et al.* *SOD1* in amyotrophic lateral sclerosis: ‘ambivalent’ behavior connected to the disease. *Int. J. Mol. Sci.* **19**, 1345 (2018).
- Sangwan, S. & Eisenberg, D. S. Perspective on *SOD1* mediated toxicity in Amyotrophic Lateral Sclerosis. *Postepy Biochem.* **62**, 362–369 (2016).
- Rosen, D. R. *et al.* Mutations in Cu/Zn superoxide dismutase gene are associated with familial amyotrophic lateral sclerosis. *Nature* **362**, 59–62 (1993).
- Deng, H. X. *et al.* Amyotrophic lateral sclerosis and structural defects in Cu, Zn superoxide dismutase. *Science* **261**, 1047–1051 (1993).
- Reaume, A. G. *et al.* Motor neurons in Cu/Zn superoxide dismutase-deficient mice develop normally but exhibit enhanced cell death after axonal injury. *Nat. Genet.* **13**, 43–47 (1996).
- Saccon, R. A., Bunton-Stasyshyn, R. K. A., Fisher, E. M. C. & Fratta, P. Is *SOD1* loss of function involved in amyotrophic lateral sclerosis? *Brain* **136**, 2342–2358 (2013).
- Lykke-Andersen, S. & Jensen, T. H. Nonsense-mediated mRNA decay: an intricate machinery that shapes transcriptomes. *Nat. Rev. Mol. Cell Biol.* **16**, 665–677 (2015).
- Popp, M. W. & Maquat, L. E. Leveraging rules of nonsense-mediated mRNA decay for genome engineering and personalized medicine. *Cell* **165**, 1319–1322 (2016).

12. Lindeboom, R. G. H., Vermeulen, M., Lehner, B. & Supek, F. The impact of nonsense-mediated mRNA decay on genetic disease, gene editing and cancer immunotherapy. *Nat. Genet.* **51**, 1645–1651 (2019).
13. Lim, L., Lee, X. & Song, J. Mechanism for transforming cytosolic SOD1 into integral membrane proteins of organelles by ALS-causing mutations. *Biochim. Biophys. Acta - Biomembr.* **1848**, 1–7 (2015).
14. Lim, L. & Song, J. SALS-linked WT-SOD1 adopts a highly similar helical conformation as FALS-causing L126Z-SOD1 in a membrane environment. *Biochim. Biophys. Acta - Biomembr.* **1858**, 2223–2230 (2016).
15. Corrado, L. *et al.* SOD1 gene mutations in Italian patients with Sporadic Amyotrophic Lateral Sclerosis (ALS). *Neuromuscul. Disord.* **16**, 800–804 (2006).
16. Siddique, T. & Deng, H. X. Genetics of amyotrophic lateral sclerosis. *Hum. Mol. Genet.* **5**, 1465–1470 (1996).
17. Cerami, E. *et al.* The cBio cancer genomics portal: an open platform for exploring multidimensional cancer genomics data. *Cancer Discov.* **2**, 401–404 (2012).
18. Gao, J. *et al.* Integrative analysis of complex cancer genomics and clinical profiles using the cBioPortal complementary data sources and analysis options. *Sci Signal* **6**, 1–20 (2014).
19. Valdmanis, P. N. *et al.* A mutation that creates a pseudoexon in SOD1 causes familial ALS. *Ann. Hum. Genet.* **73**, 652–657 (2009).
20. Fujisawa, T. *et al.* A novel monoclonal antibody reveals a conformational alteration shared by amyotrophic lateral sclerosis-linked SOD1 mutants. *Ann. Neurol.* **72**, 739–749 (2012).
21. Li, X. *et al.* A new function of copper zinc superoxide dismutase: as a regulatory DNA-binding protein in gene expression in response to intracellular hydrogen peroxide. *Nucl. Acids Res.* **47**, 5074–5085 (2019).
22. Bravo-Hernandez, M. *et al.* Spinal subpial delivery of AAV9 enables widespread gene silencing and blocks motoneuron degeneration in ALS. *Nat. Med.* **26**, 118–130 (2020).
23. van Zundert, B. & Brown, R. H. Silencing strategies for therapy of SOD1-mediated ALS. *Neurosci. Lett.* **636**, 32–39 (2017).
24. Raoul, C. *et al.* Lentiviral-mediated silencing of SOD1 through RNA interference retards disease onset and progression in a mouse model of ALS. *Nat. Med.* **11**, 423–428 (2005).
25. McCampbell, A. *et al.* Antisense oligonucleotides extend survival and reverse decrement in muscle response in ALS models. *J. Clin. Investig.* **128**, 3558–3567 (2018).
26. Miller, T. M. *et al.* An antisense oligonucleotide against SOD1 delivered intrathecally for patients with SOD1 familial amyotrophic lateral sclerosis: a phase 1, randomised, first-in-man study. *Lancet Neurol.* **12**, 435–442 (2013).
27. Van Rheenen, W. *et al.* Project MinE: study design and pilot analyses of a large-scale whole-genome sequencing study in amyotrophic lateral sclerosis. *Eur. J. Hum. Genet.* **26**, 1537–1546 (2018).
28. Andersen, P. M. *et al.* Phenotype in an infant with SOD1 homozygous truncating mutation. *N. Engl. J. Med.* **381**, 486–488 (2019).
29. Park, J. H. *et al.* SOD1 deficiency: a novel syndrome distinct from amyotrophic lateral sclerosis. *Brain* **142**, 2230–2237 (2019).
30. Gurney, M. E. *et al.* Motor neuron degeneration in mice that express a human Cu, Zn superoxide dismutase mutation. *Science (80-)* **264**, 1772–1775 (1994).
31. Shaw, M. P. *et al.* Stable transgenic C9orf72 zebrafish model key aspects of the ALS/FTD phenotype and reveal novel pathological features. *Acta Neuropathol. Commun.* **6**, 125 (2018).
32. Walters, R., Manion, J. & Neely, G. G. Dissecting motor neuron disease with *Drosophila melanogaster*. *Front. Neurosci.* **13**, 331 (2019).
33. Fujimori, K. *et al.* Modeling sporadic ALS in iPSC-derived motor neurons identifies a potential therapeutic agent. *Nat. Med.* **24**, 1579–1589 (2018).
34. Stenson, P. D. *et al.* The Human Gene Mutation Database: towards a comprehensive repository of inherited mutation data for medical research, genetic diagnosis and next-generation sequencing studies. *Hum. Genet.* **136**, 665–677 (2017).
35. Robinson, J. T. *et al.* Integrative genome viewer. *Nat. Biotechnol.* **29**, 24–26 (2011).
36. Andersen, P. M. *et al.* Sixteen novel mutations in the Cu/Zn superoxide dismutase gene in amyotrophic lateral sclerosis: a decade of discoveries, defects and disputes. *Amyotroph. Lateral Scler. Other Motor Neuron Disord.* **4**, 62–73 (2003).
37. Wu, J., Shen, E., Shi, D., Sun, Z. & Cai, T. Identification of a novel Cys146X mutation of SOD1 in familial amyotrophic lateral sclerosis by whole-exome sequencing. *Genet. Med.* **14**, 823–826 (2012).
38. Hu, J. *et al.* A novel SOD1 mutation in amyotrophic lateral sclerosis with a distinct clinical phenotype. *Amyotroph. Lateral Scler.* **13**, 149–154 (2012).
39. Shi, S., Li, L., Chen, K. & Liu, X. Identification of the mutation of SOD1 gene in a familial amyotrophic lateral sclerosis. *Zhonghua Yi Xue Yi Chuan Xue Za Zhi* **21**, 149–152 (2004).
40. Tripolszki, K. *et al.* Genetic analysis of the SOD1 and C9ORF72 genes in Hungarian patients with amyotrophic lateral sclerosis. *Neurobiol. Aging* **53**(195), e1-195.e5 (2017).
41. Canosa, A. *et al.* A novel p.Ser108LeufsTer15 SOD1 mutation leading to the formation of a premature stop codon in an apparently sporadic ALS patient: insights into the underlying pathomechanisms. *Neurobiol. Aging* **72**(189), e11-189 (2018).
42. Jackson, M. *et al.* Copper/zinc superoxide dismutase 1 and sporadic amyotrophic lateral sclerosis: analysis of 155 cases and identification of a novel insertion mutation. *Ann. Neurol.* **42**, 803–807 (1997).
43. Pramatarova, A. *et al.* A two basepair deletion in the SOD1 gene causes familial amyotrophic lateral sclerosis. *Hum. Mol. Genet.* **3**, 2061–2062 (1994).
44. Watanabe, Y. *et al.* Absence of the mutant SOD1 in familial amyotrophic lateral sclerosis (FALS) with two base pair deletion in the SOD1 gene. *Acta Neurol. Scand.* **95**, 167–172 (1997).
45. Prudencio, M., Hart, P. J., Borchelt, D. R. & Andersen, P. M. Variation in aggregation propensities among ALS-associated variants of SOD1: correlation to human disease. *Hum. Mol. Genet.* **18**, 3217–3226 (2009).
46. Andersen, P. M. *et al.* Phenotypic heterogeneity in motor neuron disease patients with CuZn-superoxide dismutase mutations in Scandinavia. *Brain* **120**, 1723–1737 (1997).
47. Jonsson, P. A. *et al.* Minute quantities of misfolded mutant superoxide dismutase-1 cause amyotrophic lateral sclerosis. *Brain* **127**, 73–88 (2004).
48. Orrell, R. W. *et al.* Clinical and functional investigation of 10 missense mutations and a novel frameshift insertion mutation of the gene for copper-zinc superoxide dismutase in UK families with amyotrophic lateral sclerosis. *Neurology* **48**, 746–751 (1997).
49. Chen, S. *et al.* A novel 10-base pair insertion mutation in exon 5 of the SOD1 gene in a Chinese family with amyotrophic lateral sclerosis. *Neurobiol. Aging* **45**(212), e1-212.e4 (2016).
50. Kawamata, C., Morita, M., Shibata, N. & Nakano, I. Familial amyotrophic lateral sclerosis (FALS) with a novel SOD1 gene mutation: a clinicopathological study. *Rinsho Shinkeigaku* **47**, 211–216 (2007).
51. Brown, J. A. *et al.* SOD1, ANG, TARDBP and FUS mutations in amyotrophic lateral sclerosis: a United States clinical testing lab experience. *Amyotroph. Lateral Scler.* **13**, 217–222 (2012).
52. Ghandi, M. *et al.* Next-generation characterization of the cancer cell line encyclopedia. *Nature* **569**, 503–508 (2019).

Author contributions

Conceptualization, C.G. and S.L.; methodology, C.G.; formal analysis, C.G.; investigation, C.G.; resources, S.L.; data curation, A.P., B.L., C.G., C.R., J.K., K.M., P.V. and S.L.; writing—original draft preparation, C.G.; writing and editing, C.G., C.R., K.M. and S.L.; supervision, S.L.; All authors reviewed the manuscript.

Funding

This research did not receive any specific grant from funding agencies in the public, commercial, or not-for-profit sectors.

Competing interests

The authors declare no competing interests.

Additional information

Supplementary information is available for this paper at <https://doi.org/10.1038/s41598-020-77716-5>.

Correspondence and requests for materials should be addressed to C.G.

Reprints and permissions information is available at www.nature.com/reprints.

Publisher's note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.



Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit <http://creativecommons.org/licenses/by/4.0/>.

© The Author(s) 2020