

Major Review

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The biophysics of superoxide dismutase-1 and amyotrophic lateral sclerosis

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Abstract

Few proteins have come under such intense scrutiny as superoxide dismutase-1 (SOD1). For almost a century, scientists have dissected its form, function and then later its malfunction in the neurodegenerative disease amyotrophic lateral sclerosis (ALS). We now know SOD1 is a zinc and copper metalloenzyme that clears superoxide as part of our antioxidant defence and respiratory regulation systems. The possibility of reduced structural integrity was suggested by the first crystal structures of human SOD1 even before deleterious mutations in the *sod1* gene were linked to the ALS. This concept evolved in the intervening years as an impressive array of biophysical studies examined the characteristics of mutant SOD1 in great detail. We now recognise how ALS-related mutations perturb the SOD1 maturation processes, reduce its ability to fold and reduce its thermal stability and half-life. Mutant SOD1 is therefore predisposed to monomerisation, non-canonical self-interactions, the formation of small misfolded oligomers and ultimately accumulation in the tell-tale insoluble inclusions found within the neurons of ALS patients. We have also seen that several post-translational modifications could push wild-type SOD1 down this toxic pathway. Recently we have come to view ALS as a prion-like disease where both the symptoms, and indeed SOD1 misfolding itself, are transmitted to neighbouring cells. This raises the possibility of intervention after the initial disease presentation. Several small-molecule and biologic-based strategies have been devised which directly target the SOD1 molecule to change the behaviour thought to be responsible for ALS. Here we provide a comprehensive review of the many biophysical advances that sculpted our view of SOD1 biology and the recent work that aims to apply this knowledge for therapeutic outcomes in ALS.

Introduction: the four ages of SOD1 discovery

There are roughly 500 new scientific publications relating to superoxide dismutase-1 (SOD1) every year. The deluge of information over the last few years is however relatively small in comparison with totality of knowledge accrued over the 80 years in which SOD1 has been a subject of scientific investigation. This level of interest reflects its prevalence and importance in our physiological processes, the relative ease with which one can work with SOD1 and its limitless capacity to retain enigma. The story of SOD1 recapitulates that of the golden age in Life Sciences which occurred over the last century. As new technologies have become available, SOD1 has proved an interesting and amenable test-case. Biophysics has been fruitful in providing answers to questions in SOD1 biology. In this section, we give a history of the discovery of SOD1, which can be divided into four distinct periods, and then describe how biophysics has shaped our view of SOD1 in relation to the neurodegenerative disease amyotrophic lateral sclerosis (ALS).

A source of copper in blood

The first age of SOD1 discovery began two centuries ago with the work of chemists, such as Christian Friedrich Bucholz, who found copper to be a constituent of plant and animal tissue. Subsequently, in the latter half of the 19th century, came the discovery of zinc in biological materials. The copper and zinc under investigation here comprised several different sources but a portion of it was present as co-factors for what would become known as haemocuprein and later SOD1.

In the late 1920s and 30s, a copper source was identified in the blood that could be freed from its organic component with the addition of hydrochloric or trichloroacetic acids but could not be removed by dialysis except at low pH (Warburg and Krebs, 1927; Locke *et al.*, 1932). Eisler *et al.* (1936) found that all of the copper in blood serum was bound to protein. In seminal papers, Thaddeus Mann and David Keilin described the purification of a 'distinctly blue' protein, which they named haemocuprein, from cow, sheep and horse blood together with the first of many methods for its crystallisation (Mann and Keilin, 1938a, 1938b).

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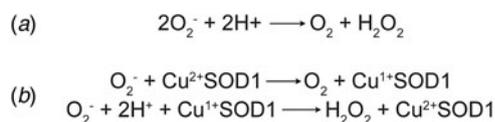


Fig. 1. Dismutation of superoxide anion. (a) The spontaneous dismutation of superoxide to oxygen and hydrogen peroxide (McCord and Fridovich, 1968). (b) Two steps in the SOD1 catalysed dismutation of superoxide which cyclically reduces then oxidises the copper centre (Klug *et al.*, 1972).

Calculation of its copper content indicated that one 35 kDa SOD1 molecule contained two copper atoms. Subsequent experiments on erythrocyte purified from human blood refined the molecular mass to 33 kDa and indicated it possessed high thermal stability (Markowitz *et al.*, 1959). Cerebrocuprein and hepatocuprein isolated from brain and liver, respectively, were found to be identical to the protein isolated from blood but no function could be ascribed (Mann and Keilin, 1938b; Porter *et al.*, 1957; Carrico and Deutsch, 1969). Variable copper metalation, dependent on source, and the ability to reversibly bind copper led to the hypothesis that erythrocyte acts as copper storage sink rather than an enzyme (Mohamed and Greenberg, 1954).

Concurrently with the initial characterisation of erythrocyte, Linus Pauling's investigation of the implications of quantum mechanics on covalent bonding had predicted the existence of three-electron bonds (Pauling, 1931). This led Edward Neuman to the discovery of the potassium superoxide state of the substance then known as potassium tetroxide. Superoxide was found to be paramagnetic with one unpaired electron (Neuman, 1934). Superoxide anion is the product of univalent reduction of dioxygen. Its primary reaction in aqueous solvents is the spontaneous, second-order dismutation to hydrogen peroxide and water (McClune and Fee, 1976). Understanding of the biological implications of superoxide generation began with biochemical characterisation of xanthine oxidase. Cytochrome *c* was found to be reduced by xanthine oxidase in the presence of its substrates. The rate of this reaction was increased under aerobic conditions and hypothesised to be mediated by superoxide (Horecker and Heppel, 1949; Rabani and Stein, 1962). Conversely, the reaction of xanthine oxidase with its substrate was shown to catalyse a chain reaction of aerobic auto-oxidation of sulphite via superoxide free radicals (Fridovich and Handler, 1958, 1961). Their observations of superoxide led Joe McCord and Irwin Fridovich to propose a scheme for the dismutation reaction (Fig. 1a) and write, 'once generated, this radical could be very damaging to the components of the cell by virtue of its high reactivity' (McCord and Fridovich, 1968).

A cupro-enzyme with dismutase activity

Discovery of erythrocyte's activity, and with it the beginning of the second age of SOD1 discovery, was a combination of many years of the rigorous biochemistry described above, and luck. McCord and Fridovich (1968) found that cytochrome *c* reduction by xanthine oxidase activity could be inhibited by carbonic anhydrase and myoglobin. On the verge of publication, the authors found that inhibition was due to an unknown impurity in both carbonic anhydrase and myoglobin preparations. From that position, the SOD activity of the impurity, erythrocyte, was proven and the centrality of its copper cofactor in the dismutation reaction (McCord and Fridovich, 1969; Klug *et al.*, 1972)

(Fig. 1b). In parallel, SOD1 was discovered to bind zinc in stoichiometric amounts to copper (Carrico and Deutsch, 1970).

Why does nature use superoxide dismutases?

'There is a bizarre enzymatic activity universally present in respiring cells. The substrate is an unstable free radical that can be present only in minuscule amounts at any instant, and the reaction catalyzed proceeds at a rapid rate even in the absence of the enzyme' (Fridovich, 1975).

When a SOD was first isolated from *Escherichia coli*, it was found to be pink and contained manganese rather than copper and zinc (Keele *et al.*, 1970). Iron SODs are related to these manganese SODs and another evolutionarily distinct form are the nickel SODs (Smith and Doolittle, 1992; Youn *et al.*, 1996). Copper/zinc superoxide dismutases (CuZnSOD) are found in the cytoplasm, intermembrane space of mitochondria, peroxisomes, nucleus, and the extracellular space of eukaryotes, and the periplasm of some prokaryotes. Manganese SODs are found in eukaryotic mitochondrial matrices as well as the prokaryotic cytoplasm, as are iron SODs. Nickel SODs are also largely prokaryotic. In almost every biological space, there is a SOD enzyme that operates within it. When oxygen is used in metabolism, SOD activity tends to be high and when organisms are exposed to oxygen, but it is not used in metabolism, SOD activity is low or inducible (Tally *et al.*, 1977). The exception which proves this rule is the obligate anaerobes which have very low or non-existent SOD activity (McCord *et al.*, 1971). Thus, enzymatic SOD activity is primarily a response to the presence or use of oxygen (Gerschman *et al.*, 1954) but may have evolved prior to Earth's great oxidation event, concomitantly with oxygenic photosynthesis. Please see excellent reviews of SOD evolution (Case, 2017) and structure (Perry *et al.*, 2010).

Superoxide is produced by many eukaryotic cellular processes and in different locations. The respiratory electron transport chain (ETC) found in the mitochondrial inner membrane provides the proton-motive force for ATP generation but contains at least 11 sites where electrons can be lost to the surrounding oxygen-rich environment (Wong *et al.*, 2017). Here superoxide is a Janus-faced component of the oxidative stress-redox signalling equilibrium; it is potentially damaging to any biological molecule, including mitochondrial DNA, while it also acts as a signal-transducer modulating respiration in a negative feedback loop. Electrons can be shed by the mitochondrial ETC into the matrix where it forms superoxide and is detoxified by manganese SODs. Conversely, the ETC complex III (cytochrome *bc1* complex) can shed electrons from its Q_o site into the intermembrane space (Han *et al.*, 2001; Bleier and Dröse, 2013) where it is likely to undergo dismutation catalysed by SOD1. Xanthine oxidase, inducible nitric oxide synthase and NADPH-oxidase enzymes all produce superoxide in various cellular compartments, and as a result, SOD1 is usually found in their environs. An example is peroxisomes, which are voracious oxygen consumers and prolific reactive oxygen species producers, and as a result, they too house SOD1 (Fransen *et al.*, 2012).

SOD1 structure discovery

Metal binding and disulphide bonding

With a function attributed to the previously mysterious erythrocyte (McCord and Fridovich, 1969), means of purification and enzymatic assay well described, the third age of SOD1

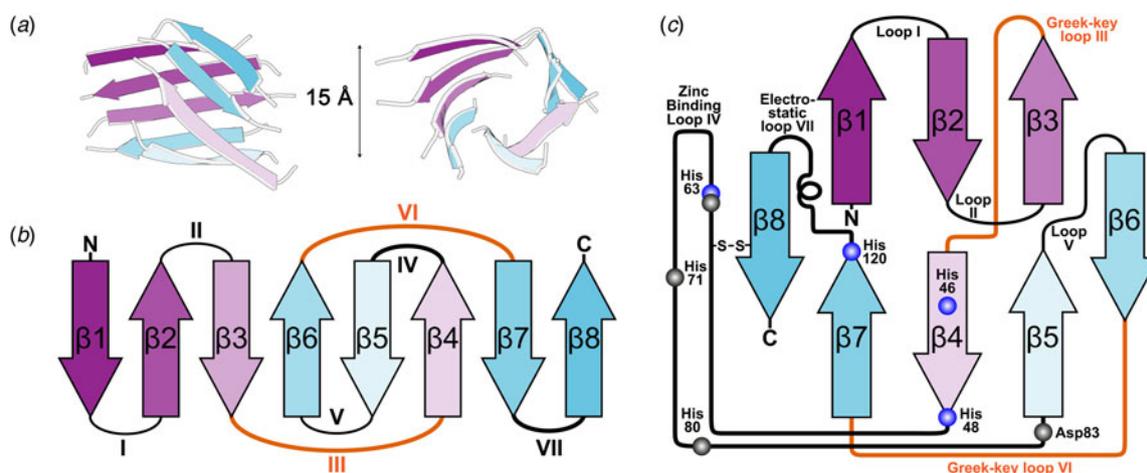


Fig. 2. The β -strand, loop and barrel structure of human SOD1. (a) β -barrel component of human SOD1 with approximately 15 Å diameter as described for bovine SOD1 (Thomas *et al.*, 1974). (b) Greek-key assembly of SOD1 β -strands showing loops which facilitate position swap of β -strands 4 and 6 in orange. (c) Arrangement of human SOD1 secondary structures within the β -barrel with residues involved in copper and zinc binding highlighted blue and grey respectively.

discovery began in earnest with biochemical and biophysical characterisation. Human and bovine SOD1 were each found to be composed of two sub-units of 16 kDa with strong interactions. This indicated that SOD1 is dimeric with high dimer affinity (Keele *et al.*, 1971; Hartz and Deutsch, 1972). Keele *et al.* (1971) found that disulphide reduction along with dodecyl sulphate denaturation was necessary to form monomeric SOD1. They concluded that a disulphide bond linked the SOD1 subunits. This was subsequently disproved by Hartz and Deutsch (1972), but these experiments form the first understanding of the role of metalation and disulphide state in the stability of SOD1. A SOD1 intra-subunit disulphide was later confirmed in the wheat CuZnSOD orthologue and a similar bond was shown to link bovine SOD1 Cys55 and Cys144 (Cys57 and Cys146 human numbering) (Beauchamp and Fridovich, 1973; Abernethy *et al.*, 1974). Analysing circular dichroism (CD) spectra from metalated, metal-apo and disulphide-reduced bovine SOD1, Wood *et al.* (1971) predicted the proximity of copper and zinc in the SOD1 structure and indicated the presence of one tryptophan per subunit. From electron paramagnetic resonance studies of SOD1 with the spectroscopically silent zinc replaced by cobalt, Rotilio *et al.* (1974) again demonstrated the proximity of the metal sites but also showed the zinc ion is not solvent exposed.

Wood *et al.* (1971) showed SOD1 has a β -sheet rich fold that is not significantly perturbed by metal removal. However, metal-free SOD1 is more susceptible to chaotrope-induced unfolding than the metalated form. They also ascertained that the disulphide bond has an impact on SOD1 tertiary structure. Forman and Fridovich (1973) found similar characteristics using enzyme activity and resistance to proteolysis as a measure of stability. With this approach, they gleaned that the zinc ion has a greater impact on stability than activity. The paradigm established here of metal loss and disulphide reduction contributing to SOD1 monomerisation and instability would become increasingly important following linkage with ALS two decades later.

The first crystallographic bovine SOD1 structure

The pursuit of a crystallographic structure of SOD1 ensued shortly after SOD catalytic activity was demonstrated. For the first decade, this work was dominated by Jane and David Richardson at Duke

University which yielded the structure of bovine SOD1 and invaluable insight on protein structure as a whole.

X-ray diffraction from SOD1 was first observed from crystals grown in phosphate buffer and 2-methyl-2,4-pentanediol which diffracted to 2 Å resolution using the precession method (Richardson *et al.*, 1972). With the revolution in synchrotron radiation crystallography still a decade away, data from multiple crystals were collected using a Picker 4-circle x-ray diffractometer based on a sealed tube. The structure was solved at 5.5 Å resolution by multiple isomorphous replacements using a combination of heavy atom derivatives soaked for up to 3 months with mercury substituted in the SOD1 zinc site (Thomas *et al.*, 1974). The authors noted a 15 Å diameter hollow passing through the centre of each monomer which they attributed to adoption of an antiparallel β -cylinder structure (Fig. 2a). A year later brought a refinement of this structure to 3 Å resolution and with it tracing the back-bone, assignment of $C\alpha$ positions and positioning of some side chains (PDB: 1SOD) (Richardson *et al.*, 1975a, 1975b). This joined the list of a dozen or so high-resolution structures at the time that included myoglobin, haemoglobin, lysozyme, ribonucleases A and S, carboxypeptidase, cytochrome c, chymotrypsin and immunoglobins.

The above research indicated that the SOD1 monomer possesses an eight stranded anti-parallel β -barrel (Fig. 2b), with the hydrophobic inner core, which constitutes roughly half of the protein backbone. The remainder of the structure is found in two loops (Fig. 2c). The 'external loop' (loop VII, amino acids 122–144 in the human enzyme and now commonly called the electrostatic loop), which was visible in the 5.5 Å resolution map, had unknown functional significance. A second loop (loop IV, amino acids 49–83, named the zinc loop) provides three coordinating ligands to the zinc ion and is held against the β -barrel by a disulphide bond but also forms part of the dimer interface along with the C-terminus and part of the β -barrel surface. Copper and zinc coordinating amino acids could be defined and the copper site geometry was categorised as distorted square planar with solvent accessibility to the copper ion (Fig. 3). The proposition that both subunits in the SOD1 dimer are conformationally identical was raised along with interface hydrogen bonding (Fig. 3).

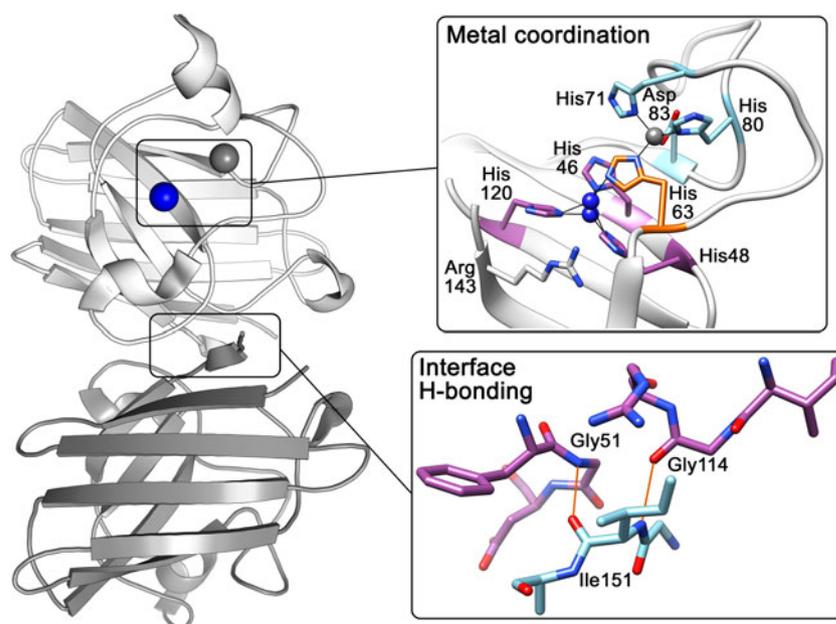


Fig. 3. Human SOD1 metal binding and dimerisation regions. Upper panel – Copper and zinc coordinating (purple and light blue respectively) sites are linked by His63 (orange). The copper ion (blue) can be found solvent exposed at two positions dependent on oxidation state while the zinc ion (grey) is internalised. Lower panel – Hydrogen bonding (orange) at the SOD1 dimer interface is mediated by backbone interactions of Ile151 with Gly51 and Gly114.

Immunoglobulin-like fold

Of particular interest was the similarity between the SOD1 β -barrel structure and other immunoglobulin domains despite sequence and functional diversity. Where SOD1 had disulphide, zinc binding and external loops connecting β -strands, hyper-variable loops could be found in immunoglobulin structures (Richardson *et al.*, 1975a). This led to the deduction of a nomenclature for β -sheet structures, the ‘Greek key’ sobriquet for the β -barrel arrangement typical of SOD1-like proteins, and a series of protein domain structure rules. The latter included conservation of β -structure over sequence, conservation of disulphide connectivity and the potential for insertions and deletions to occur in loops but not within secondary structure elements thus engineering functionality without detriment to stability (Richardson *et al.*, 1976; Richardson, 1977). It was also noted that similar domains may use different interaction surfaces for oligomerisation, a fact now easily observed by comparison of prokaryotic and eukaryotic CuZnSOD dimerisation (Bourne *et al.*, 1996).

Structure refinement and a catalytic mechanism

Working in the Richardsons’ laboratory, Elizabeth Getzoff and John Tainer were subsequently able to extend the limit of usable data to 2 Å and improve the bovine SOD1 model (PDB: 2SOD) with the stereochemistry and non-crystallographic symmetry restrained with Hendrickson–Konnert least-squares refinement (Tainer *et al.*, 1982). This allowed for determination of all non-hydrogen atom positions, placement of a single water molecule 3.2 Å from the copper ion and detailed assessment of hydrogen bonding interactions that hold the β -barrel and loops in place. Following further refinement and charge analysis, it became clear that loops IV and VII create both the active site and the electrostatic forces in and beyond the active site channel that speed diffusion of superoxide towards the positively charged copper, zinc and Arg141 (human Arg143) in the correct orientation to maximise productive interactions (Getzoff *et al.*, 1983). In the active site, superoxide displaces the copper bound water and hydrogen bonds with Arg141 side-chain guanidinium. The unpaired superoxide electron reduces copper (II) and oxygen is

released. The bond between copper and His61 (human His63) is broken and the histidine is protonated after which it hydrogen bonds with a second superoxide anion. Arg141 again hydrogen bonds with the substrate. Addition of another proton and bond rearrangement creates hydrogen peroxide and oxidises copper (I) (Tainer *et al.*, 1983). The structure of the fully reduced Cu (I) enzyme showing complete breakage of the Cu-His61 bond came two decades later at 1.15 Å resolution using a powerful synchrotron beam and cryogenically maintained crystals (Hough and Hasnain, 2003).

Recombinant human SOD1 crystallography

Taking advantage of advances in recombinant DNA technology throughout the 1970s, Getzoff and Tainer embarked on the biophysical characterisation of human SOD1. Expressed in *Saccharomyces cerevisiae* and crystallised with ammonium sulphate and poly-ethylene glycol precipitants, recombinant wild-type human SOD1 was found to form a variety of crystals which diffracted to at least 2.8 Å resolution on a rotating anode x-ray generator setup. Seeding could again be used to stimulate growth and wild-type crystals were used to grow a free sulphhydryl knock-out, thermostable mutant (Cys6Ala/Cys111Ser) following mutagenesis of the wild-type expression plasmid (Parge *et al.*, 1986). It was later shown that the removal of these free cysteines promotes refolding after thermal denaturation (Hallewell *et al.*, 1991). This work culminated in the crystal structures of recombinant wild-type and thermal-stable human SOD1 (Parge *et al.*, 1992). Diffraction data were now collected using synchrotron radiation on film at the Stanford Synchrotron Radiation Laboratory or on a Xenonics multi-wire proportional counting area detector; two developments that would revolutionise protein crystallography. Structural work continued from this point with increasing crystallographic resolution allowing ever greater detail to be revealed (Strange *et al.*, 2006) and nuclear magnetic resonance spectroscopy accessing solution states of SOD1 important in its journey from polypeptide to active protein (Banci *et al.*, 1998, 2002, 2006). In our laboratory, we routinely produce crystal structures of SOD1 in a high-throughput fashion. From initial

exposure of a crystal to x-rays through to seeing a 3D model of SOD1 now takes only hours rather than years of painstaking data collection, data reduction and model building. The combination of all of this work has given us an exquisitely detailed picture of SOD1 structure.

Amyotrophic lateral sclerosis

ALS is a neurodegenerative disease historically stated as only affecting the upper and lower motor neurons. Neuron death leads to a progressive inability to control the musculature and ultimately death usually within 5 years. We now understand that ALS is a motor system disorder affecting neurons and their support cells and forms a spectrum of clinical characteristics with frontotemporal lobar degeneration. Schizophrenia and cognitive impairment are also associated with ALS; however, this is rare for SOD1-related ALS. Following the first characterisation by Jean-Martin Charcot in 1869, conjecture on the aetiology of ALS varied between sporadic and familial (Kurland and Mulder, 1955).

SOD1 and familial ALS

The existence of SOD1 polymorphisms, prevalent within a specific geographical location and inherited from one generation to the next, that could change the physico-chemical properties of the enzyme was understood long before linkage with ALS was proven (Brewer, 1967; Beckman, 1973; Lepock *et al.*, 1990; Banci *et al.*, 1991). We now know that a portion of ALS incidence, roughly 20%, is attributed to inheritable genetic factors, termed familial ALS (fALS). A concerted effort to discover a chromosomal location associated with ALS came to fruition in 1991 with the discovery of the 21q22.1 locus (Siddique *et al.*, 1991). In 1993, the focus of SOD1 structure-function research shifted considerably with the knowledge that 11 amino acid substitutions in the SOD1 primary sequence were causative for ALS (Rosen *et al.*, 1993). In the following years, many new ALS-related mutations were discovered including the Asp90Ala identity of the polymorphism described by Beckman (1973). In the last decade and a half, SOD1 has been joined by a disparate group of genes which have varying modes of inheritance and degrees of ALS penetrance. Thus, ALS is a genetically heterogeneous disease. The year 1993 marked the beginning of the fourth age of SOD1. Here biophysics would play a fundamental role in our understanding of the transformation which turns this normally cytoprotective enzyme into a neurotoxin.

Biophysical experiments establish the nature of mutant SOD1

Prior to linkage with ALS, Parge *et al.* (1986) conjectured on the possibility that β -barrel and dimer interface mutations would affect SOD1 stability. Months after linkage with SOD1, a working hypothesis for toxicity was devised by mapping ALS mutations on the crystal structure of wild-type SOD1. Deng *et al.* (1993) then proposed that mutations to amino acids and conformations conserved in human, yeast and bovine SOD1 structures would destabilise β -barrel, dimer interface and loops. This hypothesis later became known as frame-work destabilisation (DiDonato *et al.*, 2003). Over the coming years, SOD1 cytoplasmic inclusions were found in the motor neurons and associated cells of post-mortem mutant SOD1 transgenic mice and human ALS patient tissues (Kato *et al.*, 1996; Shibata *et al.*, 1996a; Bruijn *et al.*, 1997, 1998). These observations were linked to framework destabilisation by *in vitro* biophysics experimentation which characterised SOD1 thermal instability (Rodriguez *et al.*, 2002; Tiwari and

Hayward, 2005), unfolding propensity (Cardoso *et al.*, 2002; Lindberg *et al.*, 2002; DiDonato *et al.*, 2003), dimer destabilisation (Hough *et al.*, 2004) and formation of high molecular mass soluble and insoluble structures (DiDonato *et al.*, 2003; Elam *et al.*, 2003a; Ray *et al.*, 2004). A consistent theme is the effect of SOD1 disulphide reduction and the inability to correctly bind copper and zinc on toxic properties. The structure of metal-deficient SOD1 shows loop disorder and exposure of the β -barrel (Strange *et al.*, 2003). *In vivo*, metal-free, disulphide-reduced SOD1 is over-represented within SOD1 aggregates indicating a defective post-translational modification (PTM) pathway (Jonsson *et al.*, 2006a; Karch *et al.*, 2009; Bourassa *et al.*, 2014). This brings the story of the pathogenesis of SOD1-related ALS back to the first experiments on the newly named SOD1 in the 1970s which showed that the presence of metal cofactors and the intra-subunit disulphide bond were prerequisites for stability and activity.

The pathogenesis of SOD1-related ALS

SOD1 mutations

There are more than 180 SOD1 polymorphisms associated with ALS, the majority of which are collated in the ALSod database (Abel *et al.*, 2012). Most are single amino acid substitutions, but truncations and frame-shift mutations that lead to truncations can also be found (Fig. 4a). The most common SOD1 polymorphisms are Ala4Val (Deng *et al.*, 1993), predominantly found in North America, and Asp90Ala (Andersen *et al.*, 1995) which is found in 5% of the Scandinavian population (Andersen *et al.*, 1996). These two polymorphisms reflect the diversity of SOD1 ALS mutations. They are found in different exons in the SOD1 gene (1 and 4, respectively) which code for amino acids with different properties in different parts of the SOD1 protein with different secondary structure. Ala4Val causes ALS with early onset and very short duration (Rosen *et al.*, 1994) whereas Asp90Ala leads to long survival times post presentation (Andersen *et al.*, 1996).

Much effort relating genotype, molecular characteristics and disease phenotype has yielded proposed relationships between SOD1 thermal stability, half-life, aggregation, K_m for hydrogen peroxide, hydrogen bond stability and free energy of wild-type/mutant heterodimerisation with disease course (Yim *et al.*, 1997; Sato *et al.*, 2005, 2005; Wang *et al.*, 2008; Prudencio *et al.*, 2009; Shi *et al.*, 2016). However, the small numbers of people which are known to have carried most SOD1 polymorphisms, which in some cases may be as low as one individual, and the wide distribution of longevities following the onset of symptoms significantly hamper the power of these investigations. Other predisposition genes, epigenetic and environmental factors are additionally all likely to contribute to the phenotype which eventually leads to symptom onset further complicating this task of unpicking causality in SOD1 ALS.

SOD1-ALS inheritance and penetrance

The majority of SOD1 mutations cause ALS in an autosomal dominantly inherited fashion. In the heterozygous state, Ala4Val SOD1 phenotypic penetrance is high. At age 70, 91% of Ala4Val allele-positive individuals will have developed ALS (Cudkowicz *et al.*, 1997) but examples do exist of Ala4Val carriers who remain symptom free (Weiss *et al.*, 2006). By contrast,

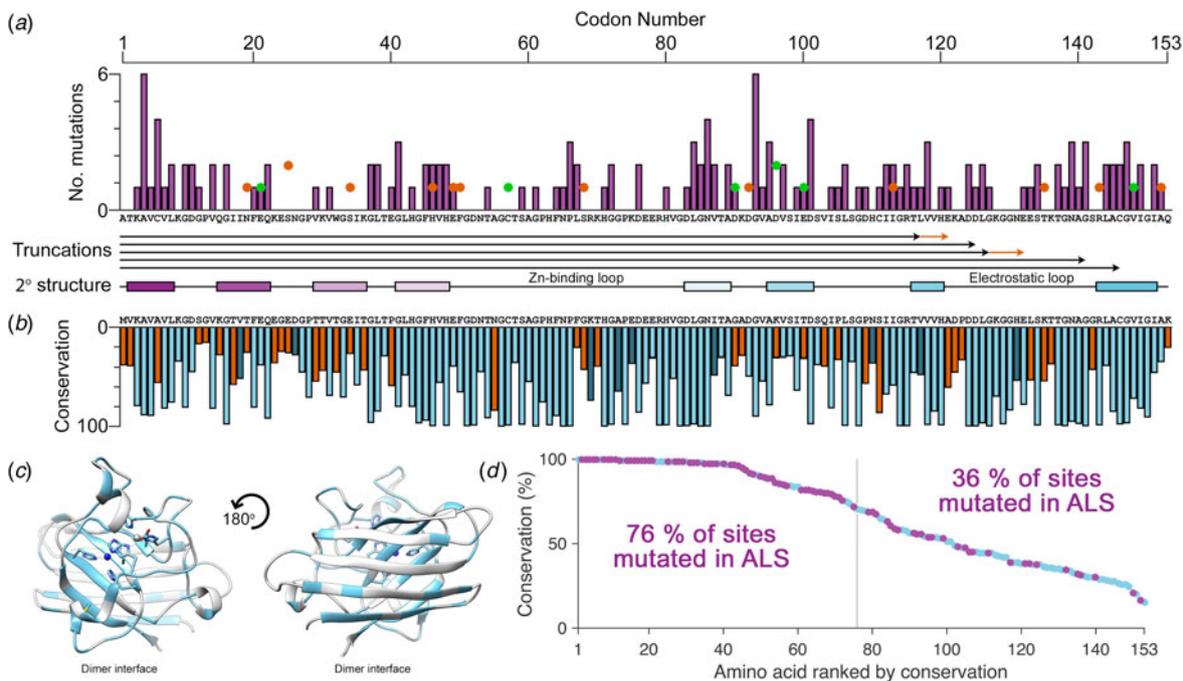


Fig. 4. The relationship between eukaryotic Cu/Zn superoxide dismutase codon conservation and human SOD1 ALS mutations. (a) The frequency of ALS mutations (purple bars), non-ALS polymorphisms (orange spots) and mutations with a disputed role in ALS (green spots) in the human SOD1 primary structure. Known truncation mutants are shown with black arrows and areas that differ from the wild-type sequence following a frame-shift mutation are shown in orange. The SOD1 secondary structure is also shown with β -strands coloured as in Fig. 2. (b) Consensus sequence of eukaryotic CuZnSODs. Light blue indicates identity between human SOD1 and the eukaryotic consensus, dark blue indicates conservative substitutions and orange indicates non-conservative substitutions. (c) Human SOD1 structure showing amino acids with >70% sequence conservation across eukaryotes highlighted in light blue. (d) Human SOD1 amino acids ranked by eukaryotic conservation (light blue). Sites that have at least one ALS-causing mutation are highlighted in purple. Sites that are highly conserved are more likely to have at least one ALS mutation.

Asp90Ala SOD1 generally results in recessively inherited ALS but has also been described as dominantly inherited with reduced penetrance in genetically distinct families (Andersen *et al.*, 1995; Robberecht *et al.*, 1996). There are several other examples of ALS mutant SOD1 homozygosity including a 13-year-old who had two copies of the Asn86Ser allele and died only 3.5 months after presentation. Heterozygotes with the same mutation have been noted to be asymptomatic or present with typical mutant SOD1 phenotype (Hayward *et al.*, 1998; Millecamps *et al.*, 2010). Individuals have also been found homozygous for a Leu126Ser substitution and a Δ Gly27/Pro28 exon 2 splicing-site variant (Kato *et al.*, 2001a; Zinman *et al.*, 2009). A further example describes three compound heterozygous Asp90Ala/Asp96Asn individuals who presented with early-onset ALS but very long disease duration (Hand *et al.*, 2001). These cases are indicative that a necessary load of toxic SOD1 species must be met before cell death ensues. As ever though SOD1 provides an enigmatic outlier; a homozygous Leu84Phe 48-year-old individual remained asymptomatic while an unrelated heterozygous individual presented with ALS at age 45 (Boukaftane *et al.*, 1998).

The Ile113Thr substitution is one of the most common ALS SOD1 mutations but the resulting phenotype can appear as sporadic ALS due to reduced penetrance (Rosen *et al.*, 1993; Suthers *et al.*, 1994). This high phenotypic variability also includes variable disease onset and duration (Orrell *et al.*, 1999; Lopate *et al.*, 2010). These characteristics are in stark contrast to Ala4Val SOD-related ALS despite the proximity of the two mutations; Ala4 and Ile113 side-chains are separated by only 3.7 Å when the protein is folded and mature. Other mutations have been noted to yield ALS with incomplete penetrance including

Gly61Arg (Conforti *et al.*, 2011), and Leu67Pro (del Grande *et al.*, 2011) however, in the absence of detailed immunohistochemistry, it is difficult to ascertain if these are truly ALS-causing SOD1 mutations or harmless polymorphisms co-incident with sporadic or low penetrance non-SOD1 fALS as has been noted for Asp90Ala and Glu100Lys families (Felbecker *et al.*, 2010). Indeed, SOD1 polymorphisms Ser25Thr, Ser25Asn, Ser34Ile, His46Tyr, Glu49Val, Phe50Cys, Ser68Phe, Asp92Gly, Ile113Met, Thr135Ile, Arg143Gly and Ala152Thr are known to exist in the wider population but, to date, are not associated with ALS in any way (Lek *et al.*, 2016). This is despite several occupying positions where other substitutions do cause ALS (His46Arg and Ile113Thr, e.g.) and several non-conservative substitutions at functionally important sites (Arg143 in the copper active site (Getzoff *et al.*, 1983) and Phe50 in the dimerisation region (Banci *et al.*, 1995)). Furthermore, several SOD1 polymorphisms linked to ALS have a disputed role in disease pathogenesis including Asn19Ser, Glu21Lys, Thr54Arg, Leu67Arg, Val148Ile, Asp90Ala, Glu100Lys, Asp96Asn and Asp96Val (Mayeux *et al.*, 2003; Andersen *et al.*, 2004; Felbecker *et al.*, 2010; Fujisawa *et al.*, 2015). Thus, not all SOD1 mutants cause ALS and those that do may do so irregularly. Felbecker *et al.* (2010) quantified this effect and estimated that only one-third of known SOD1 polymorphisms are actually pathogenic.

SOD1 mutations within the primary and tertiary structure

Eukaryotic CuZnSOD sequence conservation is high in functional parts of the molecule including interface, metal binding, β -barrel core and disulphide regions, while β -strands 2, 3 and 6 on the

opposing face of the β -barrel are divergent (Fig. 4b, c). Highly conserved SOD1 residues are more likely to have associated ALS mutations while poorly conserved residues are much less likely to harbour ALS mutations (Fig. 4d). Fifty-seven human SOD1 amino acids are not associated with any polymorphism, and while they are found sparsely through-out the primary sequence, they cluster in the β -strand 2–3 region and the zinc-binding loop. The zinc loop and disulphide sub-loop sub-structure within, overlaps with a region of very high sequence conservation extending through amino acids 43–86. This indicates that substitutions in the zinc loop have been sufficiently disadvantageous to be selected against over the whole course of eukaryotic evolution and recent human history. Conversely, β -strand 2–3 along with structurally adjacent β -strand 6 is an area of high sequence divergence. This non-functional side of the SOD1 β -barrel underwent change during early primate evolution that has been conserved in the Hominidae. The absence of ALS mutations in this region may reflect strong recent selection to inhibit SOD1 aggregation as is the case for other characteristics such as charge and thermostability (Dasmeh and Kepp, 2017) or indicate that substitutions are tolerated. The lack of sequence conservation and the existence of three non-ALS polymorphisms seems to support the latter. We predict that non-ALS allelic variation will be found to be high in this non-functional area of the SOD1 β -barrel as more sequence data become available.

Some SOD1 amino acids are more frequently mutated than others. For example, mutation of 4/4 cysteines but only 1/9 lysine sites is known to be causative for ALS. Glycine is the most common amino acid within the SOD1 sequence (16.2%). Of the 25 glycine residues within SOD1, 15 of those sites (60%) can harbour ALS mutations with a total of 29 different mutations. Gly93, for example, can be found mutated to serine, valine, alanine, cysteine, arginine and aspartic acid. Life expectancy following the onset of disease symptoms occupies a spectrum from 2 to 10 years which appears specific to each Gly93 substitution and also dictates the biophysical properties of the protein (Cudkowicz *et al.*, 1997; Pratt *et al.*, 2014).

Loss or gain of function?

SOD1 is abundant within motor neurons (Pardo *et al.*, 1995); however, reduced SOD activity of roughly 50% was quickly discovered following linkage between ALS and SOD1 (Bowling *et al.*, 1993; Deng *et al.*, 1993). This led investigators to suspect that SOD1-ALS was a loss of function disease. The resulting accumulation of superoxide radical, oxidative damage and mitochondrial pathology provided a logical route from SOD1 mutations to neuronal death (Bowling *et al.*, 1993). Progressing from this work, inhibition of SOD1 by metal chelation or antisense knock-down reduced the abundance of SOD1 protein along with SOD and choline acetyltransferase activities. The resulting oxidative stress, leading to apoptotic cell death, could be repressed with antioxidants (Rothstein *et al.*, 1994; Troy and Shelanski, 1994). However, the location of ALS mutations on the SOD1 structure ruled out a direct effect on catalysis (Deng *et al.*, 1993). Investigating how ALS mutations reduce SOD1 activity, Borchelt *et al.* (1994) concluded that a range of specific activities could be found for a variety of SOD1 mutants from zero up to 150% of wild-type for Gly85Arg and Gly37Arg, respectively. However, overall Gly37Arg SOD1 activity and stability were reduced in lymphoblasts from an fALS individual. This correlated with the reduction of SOD1 polypeptide half-life in a cell model

with Ala4Val and Gly85Arg having the shortest at 7.5 h as opposed to wild-type at roughly 30 h. The first biophysical characterisation of disease-relevant mutations on purified protein was performed by mapping human ALS substitutions onto *S. Cerevisiae* Cu/ZnSOD (Nishida *et al.*, 1994). On addition of copper and zinc to the metal-free protein, Gly85Arg was found to have 40% of wild-type activity while Gly93Ala had 80%. The latter could restore insensitivity to oxidative stress in SOD1 knock-out yeast. These results were the first indications that SOD1-ALS does not proceed by a loss of function mechanism.

A small proportion of dominantly inherited diseases are caused by haploinsufficiency; however, diseases with protein loss of function are usually inherited recessively. This led to questioning of the validity of the loss of function hypothesis. A dominant negative mechanism, where the mutant protein effects the functionality of the wild-type, formed the basis of a second hypothesis and, to this end, Gly37Arg SOD1 was shown to heterodimerise with wild-type (Borchelt *et al.*, 1994). However, Borchelt *et al.* (1995) found that co-expression of Gly41Asp and Gly85Arg SOD1 mutants with wild-type did not affect the activity or half-life of the wild-type form. Furthermore, high-level transgenic expression of Gly93Ala human SOD1 in mice led to an ALS-like phenotype at 3–4 months old whereas lower expression did not (Gurney *et al.*, 1994). This forms a parallel with the notion that high expression of SOD1 in spinal motor neurons predisposes them to mutant SOD1-mediated cell death (Pardo *et al.*, 1995), and SOD1 mutants induce apoptosis in neuronal culture where as the wild-type protein does not (Rabizadeh *et al.*, 1995). Borchelt *et al.* (1995) concluded ‘that it is unlikely that a partial loss of free radical scavenging activity underlies motor neuron disease in fALS’ and ‘the dominant character of SOD1-linked fALS is due to a gain of some injurious property of mutant SOD1 subunits’. Indeed, complete knock-out of SOD1 does not cause ALS-like symptoms (Reaume *et al.*, 1996). More recently, reduced expression resulting from a 50 bp deletion in the SOD1 promoter region was shown not to cause ALS in humans but also does not change the ALS phenotype in people harbouring mutant SOD1 (Ingre *et al.*, 2013). The possible implications of SOD1 loss of enzymatic activity in ALS have been excellently reviewed by Saccon *et al.* (2013) and will not be described further.

Aggregation, a toxic gain of function

While this work was ongoing, SOD1 was found to be a component of cytoplasmic hyaline inclusions and Lewy bodies in the spinal cord of SOD1 fALS patients (Shibata *et al.*, 1993, 1996a). Bruijn *et al.* (1997) were then able to demonstrate the formation of these aggregates in Gly85Arg SOD1 transgenic mice that express the mutant protein at a level comparable with native mouse SOD1 but which contributes little to cellular SOD activity. The human mutant SOD1 increased in abundance as the mice aged as opposed to the native mouse form which remained static. They found astrocytic SOD1 Lewy-like bodies prior to ALS-like symptom onset and they increased in number as the disease progressed. Within motor neurons, very few indistinct SOD1 aggregates were observed before symptoms which progressed to Lewy-like bodies and irregular inclusions concomitantly with the disease. This combined evidence from mice and humans of abnormal mutant SOD1 aggregation provided a putative gain of toxic function to complement dominant SOD1-related fALS

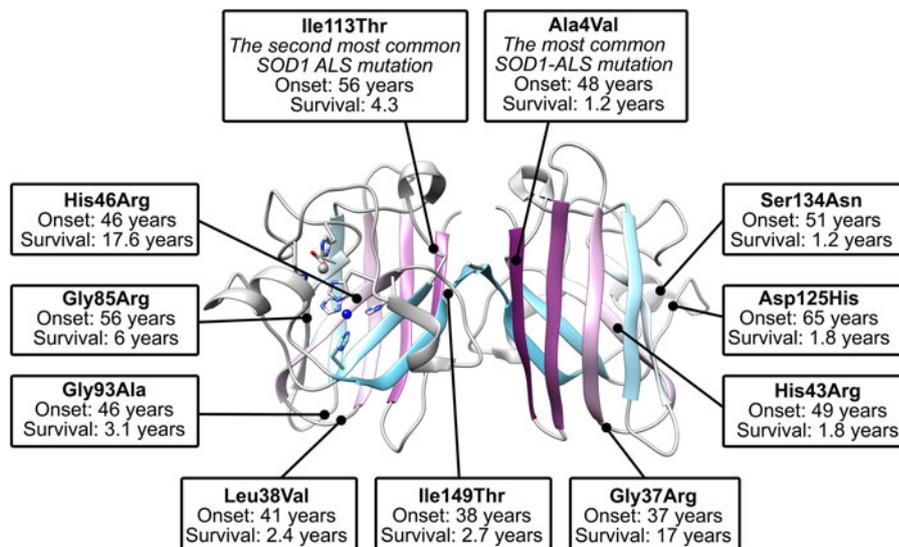


Fig. 5. The location of commonly studied ALS mutants within the SOD1 dimer structure. Shown are those mutants that have been described in crystal structures and have phenotypic data available from more than two cases (Wang *et al.*, 2008).

inheritance. A detailed description of SOD1 aggregates will follow but first we look at how ALS mutations affect the SOD1 structure.

Mutant SOD1 structure and phenotype

Of the 11 ALS SOD1 substitution mutations initially discovered by Rosen *et al.* (1993), six have been described by crystal structures. The complete library of crystallographic and NMR structures of SOD1 now comprises wild-type in various maturation states and mutants from all broad groups including dimer interface, β -barrel, metal-binding and loop regions. Mutants are found in metalation states dependent on mutant characteristics, expression platform, post-purification processing and crystallisation conditions. In some cases, mutant structures are very similar to the wild-type but often the destabilising nature of the resulting structural change is underestimated. The singular finding from this painstaking biophysical characterisation is that all ALS mutations frustrate molecular packing and reduce the likelihood that SOD1 will populate the canonical mature state. There is no single mechanism by which this destabilisation occurs, indeed every conceivable destabilisation strategy is represented; hydrogen bonds are weakened or broken, hydrophobic interactions are negated and salt-bridges removed. Most frequently however, the attractive and repulsive Van der Waals interactions which support the SOD1 structure are disturbed and new repulsive clashes are introduced. What follows is a dissection of the destabilising effects found in SOD1 ALS amino acid substitution mutants available in the Protein Data Bank (PDB) with correlation drawn to phenotypic data (Fig. 5). Where noted, mutations have been introduced into the thermostable SOD1 background (Cys6Ala/Cys111Ser) described by Parge *et al.* (1986).

Gly37Arg

The first structure of a mutant protein related to ALS as a whole was the crystallographic structure of Gly37Arg SOD1 (PDB: 1AZV) (Hart *et al.*, 1998). This mutation is found in loop III, adjacent to the Leu38 β -barrel plug and is conserved in 96% of eukaryotic CuZnSODs. The structure represents mature SOD1 with both metal ions and intra-subunit disulphide. The authors observed asymmetry in copper site geometry and subunit

B-factors. As with many ALS-related SOD1 mutant structures, the structural effect of the mutation is subtle. Comparison of wild-type and Gly37Arg SOD1 reveals a steric clash between the Arg37 and loop I Pro13 (Fig. 6). In the wild-type form, the closest distance between these two residues, Gly37 C α and Pro13 carbonyl oxygen, is 3.8 Å. However, substitution for arginine would reduce this nearest contact between C β and C γ side-chain atoms to 2.3 Å. This is well within the 3.2 Å sum of Van der Waals radii for carbon and oxygen atoms and would be energetically unfavourable. As a result, the backbone of loops I and III distort to accommodate the 3.3 Å separation found in the crystal structure. The Arg37 side-chain has comparatively high B-factors with the exception of the C β atom. While the side-chain is conformationally dynamic, the C β position is dictated by the position of loop III. Therefore, any mutation of Gly37, which by necessity incorporates an additional C β atom, will lead to SOD1 structural frustration and ALS. The reported Gly37Val cases led to a 14-month and 4.6-year disease progression (Kobayashi *et al.*, 2012; Bali *et al.*, 2017) in contrast to 16.5 years mean for Gly37Arg (Cudkowicz *et al.*, 1997; Bali *et al.*, 2017), possibly due to the increased repulsion presented by two conformationally restricted carbon atoms positioned between loops II and III rather than one.

Ala4Val

Crystallographic characterisation of Ala4Val mutant SOD1, the most frequently occurring ALS SOD1 mutation (Cudkowicz *et al.*, 1997; Wang *et al.*, 2008; Bali *et al.*, 2017), also indicated conservation of fold, metalation and disulphide bond but rearrangement around the mutation site situated close to the dimer interface. The valine side-chain is clearly defined by electron density in thermostable (PDB: 1N19) (Cardoso *et al.*, 2002) and wild-type (PDB: 1UXM) (Hough *et al.*, 2004) backgrounds as it is in the metal-free (PDB: 3GZQ) (Galaleldeen *et al.*, 2009) structure. This indicates it is conformationally restricted. Introducing two additional carbon atoms inside the tightly packed hydrophobic core of the β -barrel pushes surrounding Phe20, Leu106, Ile113, Ile149 and Ile151 away from the mutation site (Cardoso *et al.*, 2002) (Fig. 6). These permutations change inter-monomer orientation (Hough *et al.*, 2004). Mean interface Gly51-Ile151 and Ile151-Gly114 hydrogen bonding distances are decreased 2.04

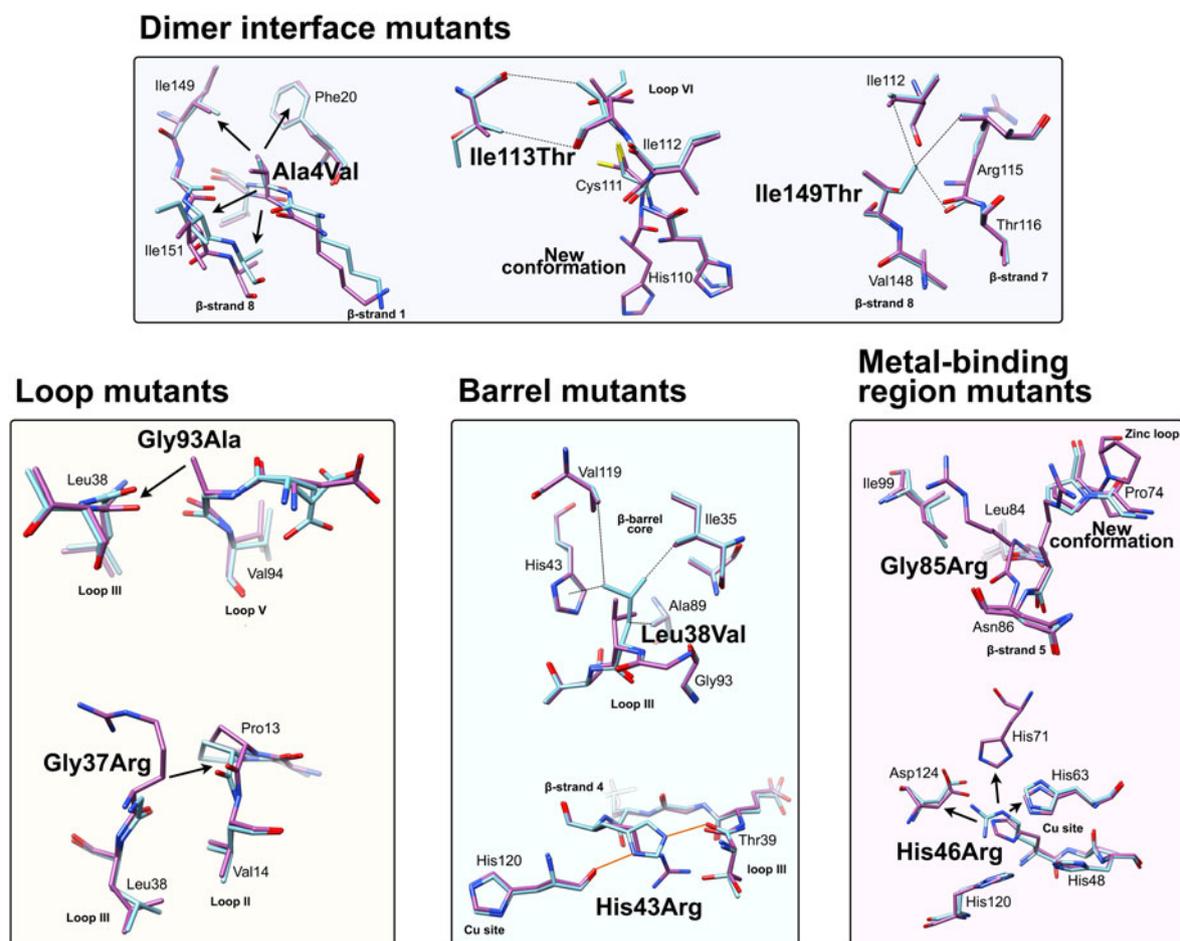


Fig. 6. Local structural destabilisation of SOD1 by ALS mutations in crystallography. Metal binding mutants; Ala4Val shows repulsion around the mutation site while Ile113Thr and Ile149Thr disrupt Van der Waals interactions at the dimer interface and the SOD1 core, respectively. Loop mutants; Gly93Ala causes repulsion between loop III and loop V and Gly37Arg causes repulsion between Loop III and Loop II. β -barrel mutants; Leu38Val disrupts Van der Waals interactions in the SOD1 core and His43Arg breaks a network of stabilising hydrogen bonds. Metal binding-region mutants; Gly85Arg causes repulsion and zinc loop disruption whereas His46Arg prevents copper binding and destabilises the electrostatic loop. Hydrogen bonds are shown in orange lines, repulsive interactions are shown as arrows, Van der Waals interactions are shown as dotted lines.

1.99 and 1.93 to 1.87 Å (calculated as acceptor–hydrogen distance or 2.78 to 2.75 and 2.87 to 2.83 Å calculated as acceptor–donor distance) with concomitant changes to bond angles $+1.7$ and -1.4° . Thus, on average, the Ile151–Gly114 hydrogen bond appears strengthened by the mutation. However, the variability of Ile151–Gly115 hydrogen bond distances and angles observed in Ala4Val SOD1 structures is far higher than those in wild-type structures; standard deviation 0.02–0.14 Å and 1.9–4.5°. This describes a frustrated molecule which is not able to settle into the tightly packed and stable structure occupied by wild-type SOD1. Alanine is found at this position in 88% of eukaryotic CuZnSODs; however, five other ALS-related mutations are found at codon 4, Ala4Ser, Ala4Thr, Ala4Pro, Ala4Asp (Naruse *et al.*, 2013) and Ala4Phe (Baek *et al.*, 2011). Each increases the number of side-chain atoms within the core of the SOD1 β -barrel. While these mutations cause variable onset of ALS, the disease course is invariably short, 7 months to 2 years, as a result.

Gly93Ala

Gly93 is found in loop V where it is distant from metal binding, disulphide and dimer interface regions but is conserved in 90% of eukaryotic CuZnSODs. Six different SOD1 codon 93 mutations

are known to cause ALS: Gly93Ala, Gly93Cys, Gly93Ser, Gly93Val, Gly93Asp and Gly93Arg with very similar mean age at onset (47.8 years) but high variance (± 14.8 years) and mutant-specific duration variability (Gly93Ala 2.2 ± 1.5 years, Gly93Cys 10.1 ± 6.1 years and Gly93Asp 10.5 ± 5.5 years) (Cudkowicz *et al.*, 1997). Comparison of metalated Gly93Ala SOD1 (PDB: 3GZP, 3GZO, 2WKO) with wild-type SOD1 shows that introduction of the alanine C β would cause a steric clash with the loop III Leu38 carbonyl (2.7 ± 0.1 Å). To mitigate this effect, the Leu38 peptide bond rotates towards the SOD1 core increasing this distance to 3.1 ± 0.1 Å. In addition, the peptide bonds of residues 92 and 93 rotate moving their carbonyls away from the core and shifting the overall position of loop V away from Leu38 in the loop III (Galalaldein *et al.*, 2009) (Fig. 6).

Leu38Val

Leu38 caps one end the β -barrel with its side-chain inserted into the hydrophobic core. This arrangement creates stabilising Van der Waals interactions with Val14, Ile35, His43, Ala89, Ala95, Val119 and Leu144. It is conserved in 84% of eukaryotic CuZnSODs. The atomic resolution crystal structure of Leu38Val SOD1 (PDB: 2WYT) shows interactions with Ile35, Ala95,

Val119 are lengthened by substitution for valine, and without rearrangement to accommodate the substitution, clashes would be created between Val38, His43, Ala89 side chains and Gly93 carbonyl (Antonyuk *et al.*, 2010). A favourable rotamer is selected to minimise clashes with His43; however, the backbone must distort slightly due to repulsion between the Val38 and Ala89 side-chains. In turn, this pushes Val14 away from the site of mutation (Fig. 6). The overall effect of Leu38Val is therefore to reduce structural cohesion within the core of the SOD1 β -barrel. In 10% of eukaryotic CuZnSODs, an aspartic acid substitutes for leucine underlining the importance of maintaining packing through Van der Waals interactions in the core of the molecule.

The Leu38Arg mutation is also associated with ALS. Here, the extra side-chain length must create highly destabilising clashes with the amino acids which normally form stabilising interactions with Leu38. Phenotypic data for Leu38Arg are sparse but show typical disease onset and progression (Millecamps *et al.*, 2010) while Leu38Val yields early-onset ALS with a very short disease course. These effects are comparable with mutation of Leu106 which caps the opposite end of the SOD1 β -barrel and forms hydrophobic Van der Waals interactions with Ala4, Phe20, Glu22, Ile112 and Ile113. Many of these interactions would be weakened or become repulsive on mutation to valine. Like Leu38Val, Leu104Val gives rise to ALS with early onset and short duration (Cudkowicz *et al.*, 1997). Leu106Phe increases the side-chain mass inside the SOD1 core like Leu38Arg, likely creating clashes with Phe20 and Val29. This causes a disease with typical age at onset but faster than usual progression (Battistini *et al.*, 2010).

Gly85Arg

Structural perturbations arising from substituting arginine for Gly85 are more pronounced in comparison with those described above. Within the SOD1 monomers found in seven available crystal structures (PDB: 2VR6, 2VR7, 2VR8, 2ZKW, 2ZKX, 3CQP, 3CQQ), the mutation site adopts two different backbone conformations. Eleven of 18 are found in a wild-type conformation with the arginine side-chain extending perpendicularly from β -strand 5 and clashing with zinc loop Pro74. Alternatively, seven of 18 monomers show a 180° rotation around the Leu84 C α -C ψ bond, which is accommodated by a further rotation around the Asn86 N-C α ϕ bond, and the arginine side-chain extends towards β -strand 6. In the latter case, zinc loop Pro74 is pushed away from the mutation site slightly changing the position of the upper zinc loop. In the former case, the steric clash between arginine and Pro74 is too severe to mitigate with a similar small shift and Pro74 rotates almost 90° (Fig. 6). This disrupts hydrogen bonding between Pro74 and Arg79, and destabilises or reorients the upper zinc-binding and electrostatic loops with knock-on effects on metal binding (Cao *et al.*, 2008). A Gly85Ser substitution has also been observed with early onset and fast disease progression (Takazawa *et al.*, 2010).

Ile113Thr

Ile113Thr is the second most common SOD1 ALS mutation (Bali *et al.*, 2017). Ile113 residues interact across the SOD1 dimer interface; Van der Waals interactions are present between the Ile113 side chain, the opposing Ile113 carbonyl oxygen and also Gly114 C α . These inter-molecular interactions are lengthened by the Ile113Thr substitution (PDB: 1UXL) (Fig. 6). In addition,

new repulsive interactions are created with Ile151 side-chain and Cys111 sulphhydryl. While there are no changes to interface hydrogen bonding in the crystal structure of Ile113Thr, the noted effects on interactions with Ile151 and Gly114 may affect the strength of these bonds in solution when higher energy states are accessible. As a result of the substitution, Cys111 adopts two side-chain rotamers and a new loop VI conformation, which extends up to Leu106, is stabilised in 30% of monomers. As noted by Hough *et al.* (2004), a hydrophobic contact with Ala4 is also lost leading to destabilisation compared to wild-type. These structural changes resulting from the mutation are subtle, and while Ile113 is conserved in 58% of eukaryotic CuZnSODs, it is often replaced by valine or leucine indicating a tolerance of these conservative substitutions. This is reflected in late disease onset, longer than average disease course and incomplete penetrance (Suthers *et al.*, 1994; Cudkowicz *et al.*, 1997).

Ile149Thr

The Ile149 side-chain C δ 1 has Van der Waals interactions with Val47, Ile112, Val148 and Leu117. The structure of Ile149Thr in a thermostable background (PDB: 4OH2) shows space is created in the core of the molecule on mutation to threonine and these hydrophobic interactions are lost. Ile112 relaxes into the newly available space as does the Arg115 peptide bond carbonyl which rotates, possibly weakening the hydrogen bond with Arg115 which links β -strands 7 and 8 (Fig. 6). Ile113 and Ile149 are foundation residues, they serve only structural purposes, there are no serious repulsive clashes generated by the conservative mutations described but a vacuum is created in the molecule which undermines structural cohesion.

His43Arg

His43 is found on β -strand 4 and forms strong hydrogen bonds with Thr39 in loop III and the copper coordinating ligand His120 in loop VII, the electrostatic loop. Held in this position, His43 provides support to Leu38 through Van der Waals interactions. The crystal structure of His43Arg, in a thermostable background (PDB: 1PTZ), shows the arginine maintaining hydrogen bonding with Thr39 and making a further hydrogen bond with Glu40 through its side-chain guanidinium group but loses the interaction with His120. To avoid a major steric clash with loop III, the Arg43 side-chain adopts a rotamer shifted away from Leu38 (Fig. 6). This reduces buried packing surface between the two residues fivefold and removes many of the Van der Waals interactions provided by His43 (DiDonato *et al.*, 2003).

His46Arg

His46 coordinates copper in the SOD1 active site and as a result is ubiquitously conserved across eukaryotic CuZnSODs. Large tertiary structure defects arising from His46Arg mutation are clear *in crystallo* (PDB: 1OZT and 1OEZ); in both metal-free and zinc-bound states, the electrostatic and zinc-binding loops are destabilised which enables non-native loop and β -barrel interactions between neighbouring molecules (Elam *et al.*, 2003a). This results from: (1) Lost His71-zinc coordination, changing both geometry, position and very likely affinity for zinc. (2) Clashes with copper and zinc coordinating His63 pushing it away from the site of mutation with similar effects. (3) Clashes with electrostatic loop Asp124, Thr137 and Ala140 (Fig. 6). His46Arg SOD1-related

ALS presents at the typical age for SOD1 ALS but has a very long disease course which can last more than 20 years (Aoki *et al.*, 1993; Bali *et al.*, 2017). Please see ‘Structural instability’ for a discussion of the relationship between His46Arg stability and phenotype.

Ser134Asn, Asp125His, Asp124Val and His80Arg

The electrostatic loop mutation Ser134Asn has much the same effect on SOD1 as His46Arg despite the very different location (Elam *et al.*, 2003a). While the Asn134 residue itself cannot be seen in the crystal structure (1OZU) due to disorder in the electrostatic loop, the reasons are clear; severe clashes with loop residues 125, 127–131 which prevent adoption of the canonical helical structure. Ser134 mutation to asparagine prevents adequate metal binding leading to hydrophobicity, abnormal intermolecular contacts and aggregation (Hayward *et al.*, 2002; Elam *et al.*, 2003a; Banci *et al.*, 2005; Marucci *et al.*, 2007). Counterintuitively however, the Ser134Asn phenotype is not aggressive despite the above molecular characteristics. Two heterozygous individuals developed late-onset ALS (63 and 52 years old) with variable progression (9 months and at least 2 years) while another heterozygous individual had no symptoms at 70 years old (Watanabe *et al.*, 1997; Marucci *et al.*, 2007). A fourth individual, who had trisomy 21 Down’s syndrome and two copies of the Ser134Asn SOD1 allele, developed ALS at 34 years of age which progressed over 3 years (Marucci *et al.*, 2007) and is reminiscent of the gene dosage effect observed in homozygous Asp90Ala ALS patients.

The crystal structure of Asp125His (PDB: 1P1V) (Elam *et al.*, 2003b) shows very similar destabilisation of the electrostatic loop between amino acids 129 and 135 as described for Ser134Asn. This is due to steric clashes with Gly126, Ser134 and Asn139 that prevent folding of the electrostatic loop helix. The clash with Ser134 causes the mutated His125 to flip rotamer and project into the solvent in two of three cases. In addition, a hydrogen bond which links Gly127 and the Asp125 side-chain is broken. Individuals with this mutation died of ALS at age 57 and 71, the latter following a 20-month disease duration (Enayat *et al.*, 1995).

Asp124Val also causes destabilisation of electrostatic loop residues 124–139 and upper zinc loop amino acids 66–79. The crystal structure of this mutant (PDB: 3H2P) shows the bonding network which links copper coordinating His46 through Asp124 to zinc coordinating His71 is lost on mutation to valine (Seetharaman *et al.*, 2010). This removes a vital stabilising link between the β -barrel core, disulphide sub-loop and electrostatic loop. Metal binding is consequently compromised.

His80Arg is the only verified *de novo* SOD1 mutation known to have caused ALS. Crystal structures of His80Arg SOD1 (PDB: 3QQD and 3H2Q) show destabilisation of the upper zinc and electrostatic loops due to complete loss of the zinc site and severe steric clashes with both His71 and Lys136 (Seetharaman *et al.*, 2010). His80Arg has been found in only one person who presented with ALS symptoms at 24 years of age and died 18 months later (Alexander *et al.*, 2002). This early disease onset would reduce the propagation of this mutation through future generations and illustrates that the majority of SOD1 mutations have been able to persist because they generally cause disease after the reproductive period of human life.

His46, His80, Ser134, Asp125 and Asp124 are ubiquitous across eukaryotic CuZnSODs. With the exception of His80, mutation of these residues gives rise to forms of ALS where the disease

presents late in life or with a protracted disease course. These mutations, to very highly conserved amino acids, often scramble or completely prevent important metal binding and manifest as phenotypically less severe than Leu38 and Leu106 mutant ALS. Mutations outside of the β -barrel core and dimerisation regions appear better tolerated by some molecular or cellular mechanism.

SOD1 maturation and ALS mutants

Following ribosomal translation, the progression from nascent and unfolded protein to the active and mature enzyme is crucially important in the production of SOD1. This post-translational maturation pathway includes folding, metal binding, intra-subunit disulphide formation and dimerisation. Appraisal of the mutant SOD1 crystal structures described above in conjunction with a wealth of biophysical, biochemical and cell biology research reveals common themes of structural instability, incomplete or aberrant metalation, disulphide reduction, dimerisation defects and propensity to exist in an unfolded state. Below we review the post-translational processes in the sequence they are likely to occur within cells and highlight how the different steps can be affected by ALS mutations paying particular attention to those SOD1 mutants which do not fit overall trends.

Folding

SOD1 folding can happen spontaneously or aided by the human copper chaperone for SOD1 (hCCS), but does not require ATP-dependent chaperones (Bruns and Kopito, 2007; Luchinat *et al.*, 2017). By analysing protease K-catalysed degradation following SOD1 translation, Bruns and Kopito (2007) showed that copper binding has no effect on folding. Conversely, removal of zinc from the folding environment prevents transformation into a highly protease-resistant form but not a moderately resistant intermediate. This is indicative of a two-step folding mechanism that comprises sequential zinc-independent β -barrel formation and zinc-dependent loop ordering. Analogously, using chaotropic unfolding, Lindberg *et al.* (2004, 2005) described a two-step folding mechanism for metal-free SOD1 with the denatured monomer folding independently through a transition state prior to a dimerisation event. Monomer folding is slow when compared to other two state proteins but in as expected for highly β -structure-rich proteins (Kayatekin *et al.*, 2012). Folding is the rate-limiting step and subsequent dimerisation is limited only by diffusion. Different SOD1 mutations perturb either one or both of the steps in this process and drive the protein towards immature states (Lindberg *et al.*, 2005).

Folding in the presence of zinc is faster than for the metal-free SOD1; however, the low affinity of the nascent protein for zinc may mean folding occurs slowly through metal-free species and zinc is bound later (Kayatekin *et al.*, 2008). Metal-free SOD1 monomer folding is nucleated by hydrophobic side-chain interactions within the core of the β -barrel on β -strands 1, 2, 3, 4 and 7 (Nordlund and Oliveberg, 2006) (Fig. 7). This folding nucleus operates independent of loop structures and metal binding with the enthalpic barrier to monomer folding related to the dehydration of these hydrophobic core residues (Kayatekin *et al.*, 2012; Yang *et al.*, 2018). The primary sequence distance between β -strands 4 and 7 (70 residues) offers a rationale for the relatively slow folding of β -rich SOD1 and the susceptibility for aggregation. The retention of N-terminal β -strands in the core of SOD1 aggregate structures could also be a facet of their swift assembly

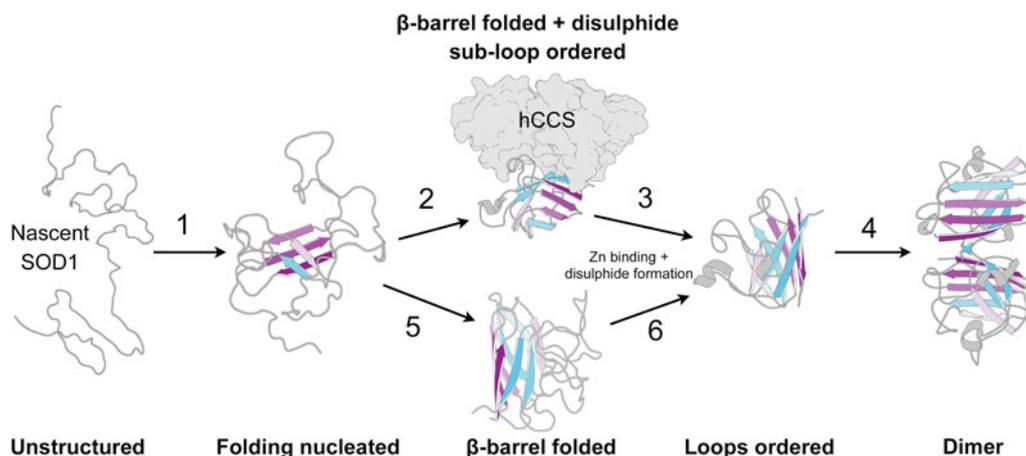


Fig. 7. The SOD1 folding pathway. Step 1: Nascent and completely unstructured SOD1 folding is nucleated by residues in β -strands 1, 2, 3, 4 and 7. Step 2: The molecular chaperone activity of hCCS then promotes folding of the remaining β -barrel structure and disulphide subloops. Step 3: SOD1 binds zinc and disulphide formation imparts stability on the disulphide sub-loop. This weakens hCCS-SOD1 heterodimer affinity. Step 4: Strong SOD1 interface Gly51-Ile151 hydrogen bonding and a stable dimer interface both promote SOD1 homodimerisation. Steps 5 and 6: hCCS mediated folding can be circumvented through spontaneous β -barrel organisation and zinc binding however the mechanism of disulphide formation, and therefore the formation of a stable dimer interface, through this hCCS-independent route is not entirely clear. SOD1 β -strands are coloured as in Fig. 2.

following translation while other strands involved in the transition state structure leave the ribosome much later.

Metal binding

SOD1 is thought to acquire free zinc from its surroundings while copper is acquired from hCCS or an hCCS-independent route (Subramaniam *et al.*, 2002; Carroll *et al.*, 2004). Zinc binding may occur in the free monomer state, or following chaperone-assisted folding when complexed with hCCS (Potter *et al.*, 2007; Kayatekin *et al.*, 2008; Leinartaitė *et al.*, 2010; Banci *et al.*, 2013; Boyd *et al.*, 2018). The canonical β -barrel fold of mature wild-type SOD1 is retained on the removal of metal cofactors under test-tube conditions (Wood *et al.*, 1971). In the metal-free state, the electrostatic and zinc loops have a high degree of disorder; however, lack of copper metalation does not significantly change the loop structure (Strange *et al.*, 2003; Sekhar *et al.*, 2015). The stabilising effect of zinc was discovered by Forman and Fridovich (1973), and our understanding of loop ordering on zinc binding reinforces that notion. However, zinc also positions the redox potential of the copper ion to enable catalytic cycling between Cu(I) and Cu(II) states (Nedd *et al.*, 2014). Furthermore, characterisation of *Candida albicans* SOD5 superoxide dismutase showed this enzyme has an intact copper site but ALS-like mutations to zinc site His71 and His80 (human numbering) which prevent zinc binding (Gleason *et al.*, 2014). As a result, *C. albicans* SOD5 has reduced copper affinity compared with other CuZnSODs (Robinett *et al.*, 2018). SOD5 is an extracellular enzyme which likely gains its metal cofactor through release of copper during host oxidative burst. Conversely, intracellular CuZnSODs, including human SOD1, must compete for copper with the high chelation capacity of the cellular interior making zinc binding a necessity. Thus, zinc binding serves roles in folding, stability, copper acquisition and catalysis.

The absence of free copper salts in biological systems was posited by Mann and Keilin (1938a) at the outset of SOD1 research. The intracellular chelation capacity of yeast, for example, means that of the estimated 390 000 copper atoms present in the cell, less than one ion exists free in solution (Rae *et al.*, 1999). To

service demand for the SOD1 copper cofactor, a relay system is necessary in order to prevent intra-cellular metal scavengers from limiting supply. Proteins responsible for the cellular trafficking of metals are termed metallochaperones and act primarily as sequestration and transport devices. Metallochaperones assist in the prevention of incorrect metal ion coordination resulting from the preference of metal binding proteins to coordinating metals at the high end of the Irving–Williams series. Sequestration also mitigates the hazard of reactive metals in solution, the case in point being free radical production by copper through redox chemistry.

Identification, structure and function of hCCS

The discovery of HAH1 and COX17 copper chaperones for the copper transporting P-type ATPases and cytochrome *c* oxidase, respectively, spurred the search for a protein that activated SOD1 with copper. Culotta *et al.* (1997) discovered the copper chaperone for SOD1 (CCS) through the similarity of SOD1 and LYS7 yeast knock-out strains' oxygen-dependent inability to synthesise lysine and methionine. hCCS was discovered by EST sequence comparison with LYS7 and found to be expressed in all human cell types. Within the central nervous system, hCCS was found to be relatively abundant within motor neurons, deep cerebellar neurons and cortical neurons overlapping the distribution of SOD1 (Rothstein *et al.*, 1999).

hCCS comprises two distinct domains which are connected by flexible linkers that allow a high degree of conformational freedom. In addition, an extended, largely unstructured region is found at the C-terminus (Wright *et al.*, 2011; Sala *et al.*, 2019). hCCS domain II (amino acids 86–234) is 47% identical to human SOD1 (Culotta *et al.*, 1997; Casareno *et al.*, 1998). Crystal structures of hCCS show that domain II has an anti-parallel Greek-key β -barrel and harbours a dimerisation interface very similar to that of SOD1 (Lamb *et al.*, 1999; Sala *et al.*, 2019). This SOD1-like domain mediates a direct interaction with SOD1 (Casareno *et al.*, 1998). While zinc binding is conserved between SOD1 and hCCS domain II, copper binding is not (Schmidt *et al.*, 1999). Furthermore, loss of zinc prevents hCCS homodimerisation and heterodimerisation with SOD1 thereby reducing its

ability to activate SOD1 (Endo *et al.*, 2000; Allen and Dennison, 2014; Wright *et al.*, 2016).

The first 85 amino acids, domain I, of hCCS and yCCS are homologous to the Atx1-like copper-binding proteins and contain the consensus MHCXXC copper (I)-binding motif which is present in HAH1, ATOX1, and both copper-transporting ATPases. Domain I binds copper (I) in a bis-cysteine arrangement with 10^{15} to 10^{18} M affinity (Brown *et al.*, 2004; Banci *et al.*, 2010, 2012a; Allen *et al.*, 2012b) without significant main chain or side chain perturbations by comparison of NMR structures PDB: 2CRL and 2RSQ produced by the Yokoyama and Banci groups, respectively. The affinity of domain I for copper is less than that of the tetra/tri-histidine SOD1 site, therefore copper flow along the hCCS trafficking route is uni-directional (Banci *et al.*, 2010). Heterodimer formation between SOD1 and hCCS along with the concerted action of domain I and a CSC motif in the C-terminal tail facilitates the delivery of both copper and the disulphide bond to the immature SOD1 substrate (Culotta *et al.*, 1997; Furukawa *et al.*, 2004).

The role of hCCS in SOD1-ALS

Mouse CCS knock-out does not change disease onset or life span in Gly37Arg, Gly93Ala and Gly85Arg SOD1 transgenic mice (Subramaniam *et al.*, 2002). Conversely, overexpression of hCCS in a mutant SOD1 background has a profound negative effect on disease; Gly93Ala SOD1/hCCS double transgenic mice have a mean survival of 36 days in comparison with 242 for single transgenic Gly93Ala SOD1 mice (Son *et al.*, 2007; Proescher *et al.*, 2008). Reduced life span is associated with quickly appearing vacuolar mitochondrial pathology. This is of note because CCS can direct SOD1 to the mitochondria where aggregation on membranes or within the intermembrane space is observed in ALS mutant cell and animal models (Deng *et al.*, 2006; Kawamata and Manfredi, 2008; Vande Velde *et al.*, 2008; Oladzad Abbasabadi *et al.*, 2013). In contrast, transgenic mice over expressing hCCS or wild-type SOD1/hCCS have normal life-span with no mitochondrial swelling (Son *et al.*, 2007). Supplementation of copper effectively confers wild-type-like survival of Gly93Ala SOD1/hCCS overexpressing mice (Williams *et al.*, 2016).

Incomplete metalation

Incomplete SOD1 metalation as a result of metal-binding ligand substitution was first observed by Banci *et al.* (1991), prior to the establishment of a link between SOD1 and ALS. One of the mutations described, Asp124Gly, was later shown to be associated with ALS (Andersen *et al.*, 2003). Following linkage with ALS, Deng *et al.* (1993) predicted that the Gly85Arg SOD1 mutation would disturb His46 and Asp83 metal site co-ordinating ligands. This was proven shortly after by incorporating the mutation into yeast CuZnSOD (Nishida *et al.*, 1994). Gly85Arg SOD1 activity was lost on addition of stoichiometric amounts of ethylenediaminetetraacetic acid, whereas wild-type and Gly93Ala retained their activity after the addition of more than 2000-fold excess. Visible light electronic absorption spectroscopy confirmed that zinc site geometry had indeed been compromised by the Gly85Arg substitution.

With the exception of the bridging ligand His63, every metal coordinating ligand (His63, His48, His71, His80, Asp83 and His120) has been found mutated in cases of ALS. On examination of the crystal structures of SOD1 where mutations are found in metal coordinating residues or close to metal sites, there is clear and rationalisable loss of metal cofactors. However, many

non-metal-binding region SOD1 mutations including Ala4Val, Leu39Val, Gly41Ser, Gly85Arg, Ile113Thr, Gly37Arg and Gly72Ser also have reduced metalation when expressed in heterologous systems (Hayward *et al.*, 2002; Luchinat *et al.*, 2014). Furthermore, metal-free, disulphide-reduced Gly85Arg, His46Arg, Val148Gly and Glu100Gly SOD1 variants do not adopt a transient electrostatic loop conformational sub-population found in 2% of the metal-free, disulphide-reduced wild-type protein (Sekhar *et al.*, 2015) that resembles the helix found in the mature protein. In the case of Ala4Val, Gly93Ala and Gly37Arg access to this conformation is partially restricted (Sekhar *et al.*, 2016). While this helix does not contain any of the zinc coordinating residues, it folds adjacent to the zinc site and may aid acquisition of zinc. On addition of zinc, the electrostatic loop loses its conformational plasticity coalescing into the conformation found in the mature form (Strange *et al.*, 2003; Culik *et al.*, 2018).

The dissociation constants for zinc and copper binding to wild-type SOD1 were determined to be 10^{-14} and 10^{-18} M⁻¹, respectively (Crow *et al.*, 1997). Eighteen to 30-fold reduction in zinc affinity for Ala4Val, Ala4Thr, Ile113Thr and Leu38Val SOD1 was demonstrated while the reduction in copper affinity was much less severe. Within cells, and even with supplementation of growth media with zinc, many of these mutants fail to adequately zinc metalate and remain in an unstructured state (Luchinat *et al.*, 2014). Absence of metals leads to the formation of high molecular mass SOD1 aggregates on incubation of purified protein at body temperature (Banci *et al.*, 2007; Wright *et al.*, 2013). Mutant SOD1 found in the soluble fraction of transgenic mouse spinal cords is metalated, albeit with perturbed Cu:Zn ratio. However, analysis of aggregates from mutant SOD1 transgenic mice spinal cord indicated no increase of copper or zinc content despite increased SOD1 content. Thus, the SOD1 that accumulates in aggregates is metal free (Lelie *et al.*, 2011). Additionally, several SOD1 conformation-specific antibodies, including the first described C4F6 that was raised against and specific for metal-free mutant SOD1, recognise their epitope in human ALS spinal tissue and lysates (Urushitani *et al.*, 2007; Brotherton *et al.*, 2012).

Disulphide bond

Wood *et al.* (1971) showed that removing metal cofactors and the disulphide bond from SOD1 led to the loss of secondary and tertiary structure. While SOD1 monomers will adopt their β -barrel fold in the absence of the intra-subunit disulphide, its presence stabilises the monomer and promotes dimerisation (Lindberg *et al.*, 2004). Conversely, formation of the disulphide can promote acquisition of structure from an otherwise globally unfolded precursor through an oxidative folding mechanism (Capper *et al.*, 2018). SOD1 must form and maintain its disulphide in the presence of reduced glutathione and the thioredoxin system which are purposed to reduce cytoplasmic thiols. When over-expressed in human cell culture, SOD1 does not naturally form the disulphide. Conversely, half of SOD1 monomers form the disulphide on co-expression with hCCS. Addition of copper to this system yields complete disulphide formation (Banci *et al.*, 2013).

The role the SOD1 intra-subunit disulphide plays in stabilisation and dimerisation is made clear in structures of the wild-type protein. Covalent bonding of Cys146, found in β -strand 8, to Cys57 anchors the zinc loop to the core of the protein. The disulphide sub-loop, a component of the zinc-binding loop IV, is defined by copper-coordinating His48 and metal-bridging ligand His63 at its N- and C-termini. The disulphide subloop forms 62%

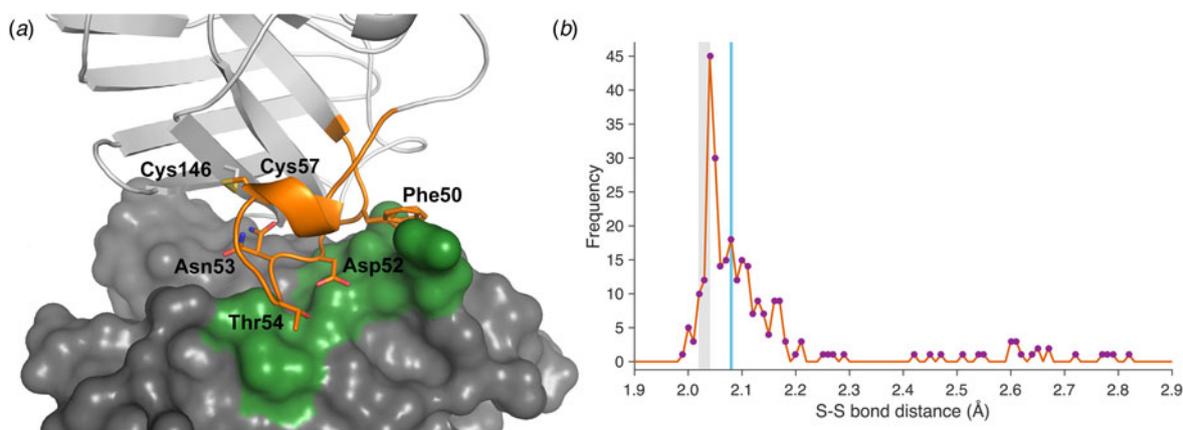


Fig. 8. The SOD1 disulphide sub-loop dimer interface. (a) The disulphide subloop forms extensive contacts across the SOD1 dimer interface. (b) S–S bond lengths found in all human SOD1 structures in the Protein Data Bank. The 2.02–2.04 Å average for all protein structures (Thornton, 1981; Petersen *et al.*, 1999) is shown in grey. By excluding bond lengths above 2.3 Å the mean SOD1 S–S bond length is found to be 2.08 Å, shown in blue.

of the buried surface area of the SOD1 dimer interface (Fig. 8a) and also contains Gly51 which creates two of the four interface hydrogen bonds (Fig. 3). We have recently shown disulphide sub-loop plasticity to also be a necessary precursor of heterodimerisation with hCCS (Sala *et al.*, 2019).

Lyons *et al.* (1996) showed that mature Ala4Val, Gly85Arg and Gly93Ala SOD1 are less resistant to intra-subunit disulphide reduction on exposure to ascorbate than the wild-type form. Indeed, all SOD1 mutants are more susceptible to disulphide loss in a reducing environment akin to the cytoplasm, promoting monomerisation and loss of structural cohesion. This effect is more pronounced for metal-binding region mutants and metal-free wild-type SOD1 (Tiwari and Hayward, 2003). In addition, Alvarez-Zaldiernas *et al.* (2016) showed that a reconstituted thio-redoxin system was capable of reducing Ala4Val and Gly93Ala disulphides. This disulphide-reduced SOD1 is observed in SOD1 transgenic mice, particularly in the brain and spinal cord (Jonsson *et al.*, 2006a), and is a major component of intracellular aggregates (Karch *et al.*, 2009).

The mean S–S length within a protein disulphide bond is 2.02–2.04 Å (Thornton, 1981; Petersen *et al.*, 1999). The S–S distance variation across every SOD1 half cystine pair found in the PDB shows a mode at 2.04 Å with a long tail stretching out to 2.82 Å (Fig. 8b). SOD1 intra-subunit S–S bond lengths above 2.3 Å are found in His46Arg/His48Gln double mutant (PDB: 3GQF) (Winkler *et al.*, 2009), Gly85Arg (PDB: 3CQP) (Cao *et al.*, 2008), Gly93Ala (PDB: 3GZO) (Galaledeen *et al.*, 2009) and His80Arg SOD1 crystallographic structures (PDB: 3H2Q and 3QQD) (Seetharaman *et al.*, 2010). These structures are well refined with resolutions of ~1.95 Å. It is very likely that these S–S bonds have been broken through exposure to x-rays as is the case for some disulphides found in the Ile113Thr SOD1 structure (PDB: 1UXL) (Hough *et al.*, 2004). This underscores the fragility of the mutant SOD1 disulphide bond and the importance of accounting for radiation damage in published crystal structures. Omitting these bonds gives a mean SOD1 disulphide bond S–S distance of 2.08 Å, similar to the 2.05 Å distance observed in bovine SOD structures (Hough *et al.*, 2000).

Dimerisation

SOD1 dimerises through portions of β -strands 1, 8, loop VI and the disulphide subloop section of loop IV. Due to primary

sequence separation of the N- and C-termini, SOD1 cannot dimerise before translation is complete. The interface is hydrophobic and excludes water. Ile151 residues create hydrogen bonds with Gly51 and Gly114 from the opposing monomer creating four interfacial hydrogen bonds. Overall, SOD1 dimer interface mutations disturb dimer interface packing but do not affect these bonds. Exceptions to this rule include hydrogen bond length variability for Ala4Val SOD1 which manifests as dimer reorientation, described in ‘Ala4Val’ section and Hough *et al.* (2004), and C-terminal truncations which remove Ile151 and therefore all interface hydrogen bonding.

As with the post-translational modifications described in the previous sections, homodimerisation increases SOD1 stability; obligate monomer SOD1 unfolds at lower chaotrope concentration than the dimer (Lindberg *et al.*, 2005). *In vivo*, dimerisation occurs after zinc binding and disulphide formation but in solution any state of wild-type SOD1 can be forced to dimerise by increasing its concentration (Arnesano *et al.*, 2004; Bruns and Kopito, 2007; Hörnberg *et al.*, 2007). When the SOD1 disulphide is reduced, the zinc-binding loop becomes dynamic and dimer affinity is reduced (Banci *et al.*, 1998). Conversely, SOD1 homodimer affinity is increased by progressively restricting zinc loop and disulphide subloop conformational sampling as SOD1 acquires post-translational modifications (Hörnberg *et al.*, 2007). The reciprocity of dimerisation and disulphide bonding is illustrated by crystal structures where both disulphide bonding cysteines are mutated to alanine. In these cases, promoting dimerisation through high concentration forces the disulphide subloop to adopt the conformation common to disulphide intact structures (Hörnberg *et al.*, 2007; Sala *et al.*, 2019).

The homodimer dissociation constant for mature wild-type SOD1 is in the low nanomolar to high picomolar range (10^{-8} to 10^{-10} M) (Khare *et al.*, 2004; McAlary *et al.*, 2013). Many SOD1 ALS mutants, including Gly93Ala, have dimer affinity equal to that of wild-type but dimer interface mutants Ala4Val, Val148Gly and Ile149Thr reduce this affinity (McAlary *et al.*, 2013; Capper *et al.*, 2018). The metal-free, disulphide-intact, wild-type SOD1 dimer dissociation constant is high nanomolar and every mutant tested by Broom *et al.* (2015a) was destabilised by up to 3-orders of magnitude with homodimer affinity ranging up to 100 μ M. As a result, reduced mutant homodimer formation, and indeed heterodimer formation with wild-type SOD1, is

observed within cells for all SOD1 mutants with the exception of Asp90Ala (Kim *et al.*, 2014). Additionally, an antibody raised against amino acids 143–151 in β -strand 8 that are only accessible when SOD1 is monomeric recognises its antigen in transgenic rodent and human ALS spinal cord samples. Immunoreactivity was highest for mitochondrial tissue fractions, particularly those from Gly85Arg mice, despite mitochondrial SOD1 being only a small component of total cellular SOD1. Tissues were stained increasingly through presymptomatic and initial symptom onset but declined at disease end stage (Rakhit *et al.*, 2007).

Reducing the affinity of metal-free SOD1 homodimerisation does not necessarily mean increased populations of unfolded monomers. Ala4Val, His46Arg and Val148Gly can all exist as folded monomers (Broom *et al.*, 2015b). The existence of monomeric immature SOD1 with a conformationally flexible disulphide loop is also a prerequisite for the formation of a transient heterodimer with hCCS (Sala *et al.*, 2019). We do not know if the affinity of hCCS for its substrate is altered by ALS SOD1 mutations but there is evidence that some mutants are not able to interact with hCCS while others form stable complexes (Son *et al.*, 2009). It is likely that mutation-dependent SOD1 homodimer affinity changes will also be reflected in the interaction with hCCS.

Structural instability

SOD1 zinc metalation, disulphide formation and dimerisation synergistically raise SOD1 thermal stability (Furukawa and O'Halloran, 2005). However, SOD1 nascent state stability is compromised by long, metal-binding loops. Shortening these functional structures increases metal-free, disulphide-reduced SOD1 stability (Yang *et al.*, 2018). This trade-off between functionality and stability is represented throughout nature. For example, proteins from thermophiles tend to have trimmed loops. Closer to our subject, *C. albicans* SOD5 and the *S. cerevisiae* CCS SOD-like domain have substantially shorter electrostatic loops in comparison with human SOD1 and the hCCS SOD-like domain, respectively, because the former does not bind zinc (Culotta *et al.*, 1997; Gleason *et al.*, 2014). Having shorter functional loops increases nascent-state stability but effective zinc binding has clearly outweighing functional benefits for SOD1.

The concept of reduced SOD1 metalation leading to structural instability was first described by Wood *et al.* (1971). ALS mutants are less resistant to chaotropic unfolding in both the metal-free and metalated states (Lindberg *et al.*, 2002; Stathopoulos *et al.*, 2003). This work led Lindberg *et al.* (2002) to propose that destabilisation leads to aggregation, while Stathopoulos *et al.* (2003) subsequently correlated decreased thermal stability, unfolding and aggregation.

Rodriguez *et al.* (2002) described reduced thermal stability of recombinant ALS SOD1 mutants in comparison with wild-type in metalation states as-isolated from insect cells. Furukawa and O'Halloran (2005) noted that the 42.9 °C melting temperature of nascent wild-type SOD1, lacking both disulphide and metal ions, is above physiological temperature and sequential addition of zinc (58.4 °C) and the Cys57-Cys146 disulphide (74.6 °C) incrementally increases thermal stability. Thus, wild-type SOD1 thermal stability increases as it progresses along the maturation pathway from metal-free, disulphide-reduced, monomeric protein to the fully metalated, disulphide intact, dimeric, active enzyme. Increased dimer affinity, reduced populations of excited states and increased thermal stability are reflected in increased enzymatic activity and half-life (Furukawa and O'Halloran, 2005; Culik *et al.*,

2018). Importantly, Gly93Ala and Ala4Val ALS substitutions reduced the melting temperature of the nascent protein below physiological temperature as ascertained by differential scanning calorimetry (DSC) experiments (Furukawa and O'Halloran, 2005). This is indicative that these species exist in a monomer unfolded state which was later observed by hydrogen/deuterium exchange and native mass spectrometry along with in-cell NMR for a range of mutants including Ala4Val and Gly93Ala (Shaw *et al.*, 2006; Luchinat *et al.*, 2014; McAlary *et al.*, 2016). Progression along the normal maturation pathway can stabilise SOD1 mutants *in vitro*, but frequently not to the same extent as wild-type, and this process is often ineffective within cells (Furukawa and O'Halloran, 2005; Luchinat *et al.*, 2014).

The majority of ALS-related SOD1 mutations have been shown to be thermally destabilising in all states where testing was possible (Rodriguez *et al.*, 2002; Stathopoulos *et al.*, 2003; Furukawa and O'Halloran, 2005; Münch *et al.*, 2011; Vassall *et al.*, 2011; Doyle *et al.*, 2016). The immature state is where the biggest stability deficits are found (Furukawa and O'Halloran, 2005; Kayatekin *et al.*, 2010) but destabilisation of the apo state is not the whole story. Metal site mutants are equally, if not more stable than wild-type, when metal free (Rodriguez *et al.*, 2005).

While metal-free, disulphide intact His46Arg SOD1 has been shown to have similar thermal stability compared to wild-type (Münch *et al.*, 2011; Abdolvahabi *et al.*, 2017). The metal-free, disulphide-reduced state is stabilised by 5 °C possibly affording it resistance to accumulation into large aggregates (Vassall *et al.*, 2011). His46Arg SOD1 is not unusual in this regard, Nordlund *et al.* (2009) showed that mutation of all four zinc coordinating residues stabilises the immature protein. Unlike non-metal-binding site ALS mutants, however, His46Arg and His48Gln cannot progress along the normal maturation pathway to reach full stability and activity (Hayward *et al.*, 2002; Bruns and Kopito, 2007). This strands His46Arg in a pseudo-stable state with a nearly 30 °C thermal stability deficit in comparison with mature, wild-type SOD1 perhaps enabling degradation. His46Arg SOD1-related ALS has a long disease course, possibly due to these characteristics.

The Val148Ile dimer interface substitution has been shown to thermally stabilise the metal-free, disulphide-reduced state by 5 °C (Vassall *et al.*, 2011) but conversely its homodimerisation affinity is decreased sevenfold (Broom *et al.*, 2015a). DSC experiments performed on mature Val148Ile SOD1 show no difference in melting temperature over a range of protein concentrations but amide proton chemical shift temperature coefficients describe stabilisation across the protein sequence (Doyle *et al.*, 2016). We also note that, like wild-type, Val148Ile SOD1 is not recognised by a misfolding-specific antibody targeting the Derlin-1 binding site in the SOD1 N-terminus possibly because of the above stabilisation (Fujisawa *et al.*, 2015). Total erythrocyte Val148Ile SOD1 activity was found to be ~58% of controls, indicative of decreased half-life in common with other destabilised ALS mutants (described in the following section). While only one individual is known to have carried the Val148Ile mutation (Ikeda *et al.*, 1995), inheritance in that individual's family was clearly dominant with disease onset very young with rapid progression (Sakuma *et al.*, 1995). Together, this is indicative of genuine SOD1-ALS. However, and as is often the case with SOD1-ALS mutations, drawing conclusions on causality from patients with very low-frequency alleles is challenging.

SOD1 like most proteins exists within a heterogenous soup of water, ions, small molecules and macromolecules within our cells.

This molecular crowding can have varied effects on proteins under investigation (Ma *et al.*, 2012). However, the SOD1 β -barrel is generally destabilised by the cellular environment when compared with purified protein in aqueous buffer. In some cases, this can reduce its folding transition temperature by 40% (Danielsson *et al.*, 2015; Bille *et al.*, 2019). The change in stability conferred by the environment occurs through interactions with β -strands 6 and 8 that stabilise the unfolded state (Bille *et al.*, 2019). As such, environmental effects are strongly determined by the primary sequence. SOD1 ALS-like mutations largely exacerbate the overall destabilising trend on the β -barrel with the exception of His46Arg which has very slightly increased stability (Gnutt *et al.*, 2019). This again highlights the cost-benefit relationship of introducing functionality. The conclusion of this work is that SOD1 species which appear folded under test-tube conditions may well be unfolded at physiological temperatures within cells. The adoption of a persistently unfolded state for many metal-free SOD1 ALS mutants in the cytoplasm is the result (Luchinat *et al.*, 2014).

Activity and degradation

The final step in SOD1 maturation yields a stable and active enzyme. However, erythrocyte SOD1 activity is almost always reduced in ALS patients (Saccon *et al.*, 2013; Keskin *et al.*, 2017). Exclusion of Asp90Ala mutants leads to a mean activity of roughly 50% of that found in non-ALS individuals. Clearly an inability to correctly metalate will reduce overall SOD1 activity but a second, related factor may also play a role. Sato *et al.* (2005) found erythrocytes taken from people heterozygous for an ALS SOD1 mutations had reduced abundance (0–58%) of the mutant SOD1 protein. Wild-type SOD1 is a long-lived protein with a 25-day half-life in cerebrospinal fluid. In transgenic rats, spinal cord wild-type SOD1 has a 15-day half-life whereas Gly93Ala SOD1 has a half-life of 9 days (Crisp *et al.*, 2015). Mutant SOD1 is also turned-over faster than wild-type in cell-based assays (Nakano *et al.*, 1996; Kabuta *et al.*, 2006). Mature red blood cells do not contain a nucleus and therefore cannot transcribe new mRNAs encoding cellular proteins. Thus, it appears that no new SOD1 is created but mutant SOD1 is preferentially degraded. Over the life-time of a red blood cell, this reduces the abundance of mutant SOD1. In general, SOD1 degradation proceeds by proteasomal or macroautophagic routes (Kabuta *et al.*, 2006; Keskin *et al.*, 2016). Ubiquitin E3 ligases dorphin, CHIP, mahogunin and cIAP have all been shown to ubiquitinate mutant SOD1 and signal its degradation (Choi *et al.*, 2004, 2016; Sone *et al.*, 2010; Chhangani *et al.*, 2016). However, SOD1 aggregation has been shown to sequester more than half of cellular ubiquitin thereby reducing the availability of free monomeric ubiquitin with downstream changes to the ubiquitination profile of more than 70 cellular proteins (Farrarwell *et al.*, 2018). Thus, ALS SOD1 mutants that cause the greatest structural disturbance and instability lead to the greatest reductions in activity and abundance but are also more likely to upset proteostatic mechanisms through aggregation and proteasomal sequestration (Sato *et al.*, 2005; McAlary *et al.*, 2016; Keskin *et al.*, 2017; Farrarwell *et al.*, 2018).

Cryptic mutations

The mutations described above induce structural changes to the SOD1 molecule that manifest as protein thermal instability, reduced homodimer affinity, inability to correctly metalate,

reduced activity, reduced half-life *in vivo* and increased aggregation propensity. However, several mutations display stability and aggregation characteristics very similar to wild-type when observed *in vitro*. Shaw *et al.* (2010) termed these SOD1 variants ‘cryptic mutations’ because it is not clear where the source of their toxicity lies. These include Asp90Ala, Asn86Ser, Asp101Asn, Asn139Lys and Glu100Lys (Shaw and Valentine, 2007). Cryptic mutations are usually surface exposed and involve the substitution of charged residues. While we do not have experimentally derived structures of these SOD1 mutants, analysis using the wild-type structure as a model provides us with an indication of their effect. In almost every case, a structural perturbation can be ascribed.

Surface mutations with mild phenotype

Asp90Ala

The Asp90Ala substitution does not affect dimer stability, metalation or activity (Andersen *et al.*, 1995; Marklund *et al.*, 1997; Rodriguez *et al.*, 2002; Lindberg *et al.*, 2005). The codon 90 position is not well conserved and the majority of eukaryotic CuZnSODs have a glycine residue at this site. Analysis of the wild-type SOD1 crystal structure shows the Asp90 side-chain making a series of hydrogen bonds within loop V with backbone amides of Asp92, Gly93 and Val94. Substitution with alanine would break these β -hairpin bonds but would not introduce steric repulsion within the loop, as is the case for Gly93 mutations. As a result, the Asp90Ala monomer is slightly destabilised (Marklund *et al.*, 1997; Rodriguez *et al.*, 2002; Lindberg *et al.*, 2005; Byström *et al.*, 2010). An alternative Asp90Val mutation creates further monomer destabilisation and a more aggressive phenotype (Morita *et al.*, 1998; Byström *et al.*, 2010).

Asn86Ser

The Asn86Ser mutation manifests ALS with a long disease course and incomplete penetrance but led to an extremely severe phenotype in a single homozygous individual (Hayward *et al.*, 1998; Sato *et al.*, 2005). Asn86 stabilises the electrostatic loop through a hydrogen bond with Asp124 which itself hydrogen bonds with zinc- and copper-coordinating His71 and His46. Conversion to serine would destabilise these interactions but would not introduce any repulsive contacts. There are no metalation studies on this mutant and the monomer does not appear effected but the Asn86Ser dimer is destabilised (Byström *et al.*, 2010).

Other examples

Other charged, surface mutations which potentially fall into this category include Glu40Gly (Bertolin *et al.*, 2014), Asp76Tyr (Andersen *et al.*, 1997), Asp76Val (Segovia-Silvestre *et al.*, 2002), Asp11Tyr (Georgouloupoulou *et al.*, 2010), Glu21Lys (Jones *et al.*, 1994a) and Glu121Gly (Canosa *et al.*, 2015). In each case, no bonding or Van der Waals interactions are likely to be disturbed or the substitution could be mitigated without changing the overall structure.

Surface mutations with typical phenotype

Asp101Asn

The Asp101Asn substitution (Jones *et al.*, 1994b) is also structurally conservative, removing only 1 Da of mass. However, it negates the electrostatic component of a salt-bridge formed by Asp101 and Arg79 which is conserved in more than 97% of eukaryotic CuZnSODs. This salt-bridge forms a strong

connection between the C-terminal end of β -strand 6 and the upper zinc loop including spatially proximal zinc coordinating residues His80 and Asp38. An alternative substitution for glycine removes both electrostatic, hydrogen bonding and Van der Waals interactions. In agreement with proposed structural changes, Asp101Asn was shown to have stability equal to that of wild-type in the metal-free state *in vitro* while Asp101Gly destabilises both metal-free monomer and dimer states (Rodriguez *et al.*, 2005; Byström *et al.*, 2010). A detailed analysis of aggregation kinetics in HEK293 cells showed that Asp101Gly exhibits a short lag phase before the formation of insoluble aggregates. By contrast, Asp101Asn SOD1 initially expresses as soluble species, but following a long lag phase, this pool is transferred into degradation-resistant insoluble aggregates (Prudencio *et al.*, 2009; Ayers *et al.*, 2014a). Given the structure of each mutation together with their biophysical and aggregation characteristics, it would be expected that Asp101Gly would yield a more severe phenotype relative to Asp101Asn. However, both mutations give rise to an early onset and rapidly progressing form of ALS (Wang *et al.*, 2008). A possible explanation for Asp101 mutation toxicity lies in their inability to correctly metalate and potential weakening of the Asp101-Arg79 salt-bridge. Ayers *et al.* (2014a) found that both Asp101Asn and Asp101Gly SOD1 fail to acquire copper or zinc in a cell-based assay. This is unsurprising given that Asp101 anchors the zinc loop to β -strand 6. Thus, like His46Arg described in 'Structural instability' section, the absence of a stability deficit in the apo state is over-shadowed by the inability of the mutant to progress along the PTM pathway. Indeed, Asp101Asn has also been shown to have delayed folding kinetics (Bruns and Kopito, 2007). A second contributory mechanism may also be the reduction of net negative charge, reviewed in 'SOD1 net charge' section.

ALS-associated mutations Asp101His and Asp101Try have also been observed and would create severe clashes with Arg79, Leu84, Val103 or Ile104 but result in a mild phenotype. Here the combination of severe structural instability together with an inability to metalate may create a protein that is easier to remove through proteostatic mechanisms or sequester thereby reducing the toxic protein load in comparison with Asp101Asn or Asp101Gly. It should be noted however that case numbers for the glycine, histidine and tyrosine substitutions are low making phenotype comparisons weak.

Other examples

Other examples of surface charge mutations which lead to a typical ALS phenotype include Arg110Gly (Kostrzewa *et al.*, 1994) which is likely to disturb the dimer interface and hydrogen bonding with loop VI helix, and Asn139Lys which is destabilised in the metal-free state (Rodriguez *et al.*, 2005; Münch and Bertolotti, 2010) because an internal electrostatic loop hydrogen bond is broken by the substitution. Little phenotype data are available for Asn139Lys but SOD1 activity is reduced 46.5% (Pramatarova *et al.*, 1995) and given its location is likely to reduce metalation. Like Asp101Asn described above, Asn139Lys has been shown to have delayed folding kinetics (Bruns and Kopito, 2007).

The pathogenesis of ALS due to surface mutations

We can summarise the above sections by saying that, a substitution that perturbs the structure little, that does not affect the non-covalent bonding configuration or does not inhibit maturation processes, leads to incomplete disease penetrance often with

later onset or longer disease course than mean. Conversely, large structural effects including breaking non-covalent interactions, introduction of repulsive interactions or inhibition of folding or PTMs lead to high penetrance and a natural history which is typical or comparatively severe for SOD1 ALS. As we have seen, however, there are exceptions to this dogma. The path from structure through stability and aggregation to disease phenotype cannot be unambiguously defined. Wang *et al.* (2008) found that stability and aggregation account for only 69% of the variability in mutant SOD1 phenotypes. There must therefore be other factors that influence disease outcomes for particular substitution mutations.

SOD1 net charge

With a preponderance of aspartate and glutamate residues over arginines and lysines, the SOD1 monomer carries a hypothetical net charge of -6 at neutral pH. This is the maximum charge found on any vertebrate SOD1 (Sandelin *et al.*, 2007). The structure surrounding ionisable side-chain groups together with the pH of the subcellular locale are important determinants of protein isoelectric point. Shi *et al.* (2013a) experimentally determined the net charge on metal-free, disulphide-reduced, monomeric, wild-type SOD1 at physiological pH to be 6.9. Formation of the SOD1 disulphide reduces the net charge to 6.05 per monomer. Binding of a zinc to disulphide intact SOD1 reduces the net charge to 4.3 and further addition of copper to create fully mature SOD1 decreases the net charge further to 3.7. Thus, the SOD1 PTM pathway increases stability but decreases monomer net charge.

ALS mutations and surface charge

ALS mutations have, on average, a propensity to reduce SOD1 net negative charge (Sandelin *et al.*, 2007). While 76% of negatively charged sites have been found mutated in ALS cases, only 13% of positively charged amino acids are known to harbour ALS mutations, Lys3 and Arg115 only. This disparity is thought to reflect a role of net repulsive charge in maintaining free movement in the intracellular environment (Mu *et al.*, 2017) and preventing aggregation; mutation of a positive residue increases net charge and therefore repulsion, whereas mutation of a negatively charged amino acid decreases net charge weakening repulsion and increasing aggregation propensity. In support of this, acetylation of SOD1 multiple lysine side-chains with aspirin (acetylsalicylic acid) increased fibrillation lag time and slowed growth (Abdolvahabi *et al.*, 2015). This could also be accomplished with a selection of charged anhydrides which modified only a single SOD1 lysine (Rasouli *et al.*, 2017). In both cases, inhibition of aggregation was not due to increased thermal stability as acylation ubiquitously decreased SOD1 melting temperature. However, inhibition of fibril growth was only observed in low ionic strength buffer indicating the charge effect may be limited under physiological conditions. Fibrillation inhibition also led to the accumulation of SOD1 into amorphous aggregates. This may be similar to the misfolding observed in the absence of large aggregates on cellular expression of SOD1 with poorly conserved lysines mutagenically removed (Crosby *et al.*, 2018).

It is possible to fit a simple linear model of instability and survival time following disease onset by omitting outlier mutants that change net charge. On inspection of these outliers, Byström *et al.* (2010) found that net charge is a modifier of disease phenotype. For example, Gly41Asp has a very similar relative stability and disease onset compared with Leu106Val, but increased net charge, and therefore repulsion, raises disease duration from 2 to 17 years.

The converse is also apparent; reducing net charge decreases survival time with Asp101Asn as the archetype example. Metal-free Asp101Asn stability is indistinguishable from wild-type but disease phenotype is similar to the very strongly destabilised Ala4Val (Byström *et al.*, 2010).

Glu100 mutations

The case of Glu100Lys is often cited as an example where net charge reduction leads to ALS. Glu100 is surface exposed at the C-terminal end of β -strand 6. Its negatively charged side-chain exists in multiple rotamers and forms no interactions with its surroundings. Glu100Lys, which reduces net charge to -4 , is likely to have limited structural effects, does not reduce erythrocyte SOD1 activity, can lead to a complicated phenotype with very long progression and was shown not to co-segregate with disease in two ALS families (Felbecker *et al.*, 2010; Yasser *et al.*, 2010; Keskin *et al.*, 2017). Conversely, the Glu100Gly mutation, which reduces SOD1 net charge only to -5 , leads to a typical ALS phenotype with high penetrance (Cudkowicz *et al.*, 1997; Aggarwal and Nicholson, 2005; Wang *et al.*, 2008). Glu100Gly decreases monomer stability (Rumfeldt *et al.*, 2006) and reduces overall thermal stability by 10 °C in the metal-free state in comparison with Glu100Lys which only creates a 4.4 °C stability deficit (Abdolvahabi *et al.*, 2017). A similar relationship is also observed when initial folding is observed; Glu100Lys acquires resistance to proteolysis only slightly slower than wild-type, whereas Glu100Gly folding is massively reduced and it behaves more like Ala4Val SOD1 (Bruns and Kopito, 2007).

SOD1 aggregation in ALS

Aggregation of protein is a common feature of neurodegenerative disease and normal ageing. This is typified by the predominantly α -synuclein-containing Lewy bodies seen in the post-mortem substantia nigra of individuals who suffered from Parkinson's disease. Bunina (1962) first described intracellular inclusions, later termed Bunina bodies, surrounded by an area of clear cytoplasm within the motor neurons of fALS cases. Single and multiple intracellular hyaline, i.e. glassy or transparent, regions often with Lewy body-like cores were subsequently described by Hirano *et al.* (1967), also in fALS neuronal tissues. While we do not know the content of these structures or the type of fALS under investigation in these original cases, we do see SOD1 accumulate in these structures within the cytoplasm of microglia, astrocytes and motor neurons of mutant SOD1 transgenic animal models and human ALS patients (Shibata *et al.*, 1993, 1996a; Kato *et al.*, 1996, 1997; Bruijn *et al.*, 1997, 1998). Together with mutations in the *sod1* gene, this is the central hallmark of SOD1-related ALS. Astrocytes have been observed with their cytoplasm almost completely replaced by 15–25 nm granule-coated fibrils. In other glia, fibrils form non-membrane-bound inclusions but are distinctly separated from the cytoplasmic constituents (Kato *et al.*, 1996, 1997). The appearance of insoluble SOD1 and gliosis in neuronal support cells appears to facilitate neuronal death (Beers *et al.*, 2006; Yamanaka *et al.*, 2008). SOD1 aggregation is also observed in Parkinson's disease and to a lesser degree in otherwise normal but aged neuronal tissue (Trist *et al.*, 2017).

Content of aggregates

Despite the relatively high turnover of many SOD1 mutants, it is a major component of aggregates in mutant SOD1 transgenic mice (Jonsson *et al.*, 2004). The SOD1 found in the amorphous and

fibrillar aggregates in the neural tissues of human and transgenic models has been shown to be partially or completely inactive, intra-subunit disulphide reduced, metal-free and misfolded but full-length (Jonsson *et al.*, 2006a; Shaw *et al.*, 2008; Karch *et al.*, 2009; Bergemalm *et al.*, 2010; Lelie *et al.*, 2011). Wild-type SOD1 is also present along with mutant protein and high molecular mass species can be linked by abnormal disulphide bonds (Jonsson *et al.*, 2004; Furukawa *et al.*, 2006; Karch *et al.*, 2009). Advanced glycation end products carboxymethyl-lysine, pyrroline and pentosidine co-localise with neuronal and astrocytic inclusions in human ALS and transgenic mice and are thought to be direct SOD1 modifications (Kato *et al.*, 1999, 2000; Shibata *et al.*, 1999). In mice expressing human Gly85Arg, SOD1 aggregates showed ubiquitin-positive staining only at their periphery (Bruijn *et al.*, 1997). However, in a cell model, ubiquitin was found to localise throughout SOD1 aggregates (Farrarwell *et al.*, 2018). A meta-analysis performed by Ciryam *et al.* (2017) has shown that five of the 15 proteins known to interact natively with SOD1 are drawn into inclusions but the majority of the 24 proteins that are found in aggregates do not have a known metabolic association with SOD1. SOD1 co-aggregators can be divided into four broad groups which offer a snap-shot of cellular activities in response to mutant SOD1: degradation machinery, neural-specific metabolism, neural cytoskeleton components and chaperones including metallochaperones. This includes ubiquitin, α B-crystallin, β -actin, metallothionein, glutamine synthetase, tubulins, tau, S-100, Hsp-27 and 70, proteasome, vimentin, neurofilament light chain, α -internexin, synaptophysin, enolase-2, protein-disulphide isomerase, glyceraldehyde-3-phosphate dehydrogenase and the copper chaperone for SOD1 (hCCS) (Kato *et al.*, 1997, 2000, 2001b; Watanabe *et al.*, 2001; Bergemalm *et al.*, 2010; Zetterström *et al.*, 2011).

Models of aggregation

Beginning with the Gly93Ala SOD1 transgenic mice described by Gurney *et al.* (1994), many strains have been produced that express various human wild-type (Wong *et al.*, 1995; Graffmo *et al.*, 2013) and SOD1 ALS mutants at different levels in different genetic backgrounds. These include, but are not limited to, Gly37Arg (Wong *et al.*, 1995), Gly85Arg (Bruijn *et al.*, 1997), His46Arg/His48Gln (Wang *et al.*, 2002) and Asp90Ala (Jonsson *et al.*, 2006b). When comparing wild-type and Gly93Ala SOD1 transgenic mice with equal transgene expression, Graffmo *et al.* (2013) found mean survival time is 367 ± 56 and 155 ± 9 days, respectively, compared to an average of 2 years for non-transgenic mice. Thus, the toxic load of human SOD1 expression appears cumulative in mice.

Mutant SOD1 transgenic mouse models were seen as a gold standard test for therapy development. The most popular is the Gly93Ala transgenic mouse created by Gurney *et al.* (1994), a year after the establishment of the link between SOD1 mutants and ALS (Rosen *et al.*, 1993). Gurney's mouse expresses human mutant SOD1 at a high level, up to 24-fold that found in humans. These mice have relatively short disease course which is convenient for many types of experiment. Despite many therapeutic approaches having been reported to positively affect disease-related characteristics in these mice, there has been a complete failure to translate these findings into new medicines (Benatar, 2007). This has raised concern about the validity of SOD1 transgenic mouse experiments as a whole. It remains an open question whether the limitations arise from the transgenic

models or the nature/protocols of the experiments and their limitations.

To further our understanding of the aggregation process, numerous cell and *in vitro* models have been developed. Mutant SOD1 aggregations are seen in cell culture models of ALS where they replicate some of the characteristics of post-mortem tissues (Durham *et al.*, 1997; Koide *et al.*, 1998). SOD1 was first observed to aggregate *in vitro* in response to oxidative modification (Rakhit *et al.*, 2002). Light scattering at 350 nm by SOD1 in the presence of soluble copper (II) showed that the metal-depleted enzyme readily forms high molecular mass species. The authors posed the question 'do these *in vitro* aggregates represent aggregates seen in ALS?'. This experiment illustrates the difficulty of recapitulating the disease state *in vitro*; any protein will aggregate if treated harshly enough, how is one to attach significance to that given that the nature of aggregates in living neuronal tissue is far from clear. Addressing this question, Lang *et al.* (2015) showed a resemblance between *in vitro* aggregates and those that appear in transgenic mouse models of ALS. Using antibody probes to characterise the surface of aggregates, they found that solvent-exposed structures are maintained *in vitro* and in mice. Similarly, with thioflavin T binding, they showed that the dynamics of growth in both systems can be equated.

Amyloid or not?

SOD1 aggregates have been shown to bind amyloid-sensitive dyes, such as congo red or thioflavin T and S, but yield less fluorescence than is observed for amyloid- β or α -synuclein (DiDonato *et al.*, 2003; Chattopadhyay *et al.*, 2008; Furukawa *et al.*, 2008; Oztug Durer *et al.*, 2009). Thioflavin S stains inclusions formed by the artificial SOD1 double mutant His46Arg/His48Gln and ALS mutants Gly37Arg, Gly85Arg and Gly93Ala in transgenic mouse spinal cord tissue (Wang *et al.*, 2002). The form of the fibres observed by Kato *et al.* (1996, 1997) seems to support the notion of amyloid fibrillation. However, Kerman *et al.* (2010) have shown that SOD1 aggregates in spinal cords are non-amyloid. Instead they contain monomeric and unfolded SOD1 in a highly amorphous arrangement. To our knowledge, there are no reports of SOD1 amyloid structures found in human post-mortem ALS tissues. On the contrary, they have been shown not to bind congo red and thioflavin S (Kato *et al.*, 1996). This does not mean the results of aggregation studies using thioflavin T fluorescence as a read-out for aggregation, for example, should be ignored. We should however expect the structure of SOD1 aggregates to differ from that of amyloid- β or α -synuclein. By extension, the structure or affinity of the dye-aggregate complexation will be altered and therefore the fluorescent properties of the dye itself.

Aggregate structure

For SOD1 aggregates, formation dictates form. This aphorism is true in two respects: Firstly, aggregation kinetics dictate morphology (Abdolvahabi *et al.*, 2016) (described in the following section), and secondly, the regions of the SOD1 molecule that mediate aggregation are sequestered within the aggregate structure. In the latter case, small hydrophobic side-chains occupy the core of the proto-fibril-like structure and exclude water in a manner reminiscent of the normal SOD1 folding nucleus (Nordlund and Oliveberg, 2006).

Insights on the structure of SOD1 fibrils can be gained from the observation of the immature state. In solution, wild-type metal-free SOD1 retains only one surface of the β -barrel (β -strands 1, 2, 3 and 6). The β -strands of the opposing face including loops constituting both metal sites, the active site channel, the disulphide sub-loop and a sizable portion of the dimer interface, all which are highly conserved across eukaryotic CuZnSODs (Fig. 4b), are significantly destabilised (Banci *et al.*, 2009). Simulation of aggregation *in silico* also shows β -strands 1 and 2 self-associate to form β -sheet structures (Bille *et al.*, 2013). Binding of conformation-specific antibodies to segmented epitopes along the SOD1 primary sequence confirms the presence of β -strands 1, 2, 3 in the core of aggregates along with strand 8 (Bergh *et al.*, 2015). β -strands 1, 2 and 6 along with strand 7 are found sequestered in aggregates as determined by protease digestion followed by mass spectrometry assignment; however, this varies along with mutation (Furukawa *et al.*, 2010).

Clues as to the structure of SOD1 aggregates can be gleaned from the ALS mutations themselves. C-terminal truncations at positions Leu117 (with four amino acids inserted before the stop codon), 127 (with five amino acids inserted before the stop codon), 125, 141 and 146 (Fig. 4a) are known to cause ALS in people and transgenic mice with intraneuronal SOD1-positive inclusions (Jackson *et al.*, 1997; Zu *et al.*, 1997; Jonsson *et al.*, 2004; Wu *et al.*, 2012; Nakamura *et al.*, 2015). Thus, the electrostatic loop and β -strand 8 including Cys146 are not required for aggregate formation.

Bergh *et al.* (2015) explored the variability of mutant SOD1 aggregate isoforms in transgenic mice and found that Asp90Ala aggregates had a distinct structure, with solvent exposure of β -strands 5, 6, 7, that was not seen in Gly93Ala, Gly85Arg and wild-type SOD1 isoforms. β -strand 4 and the disulphide sub-loop were however consistently found on the exterior of SOD1 aggregates and in a conformationally plastic state regardless of mutation. Notably, the Asp90Ala strain of SOD1 aggregation also appears in neuronal tissues of ALS patients with mutations in non-SOD1 genes whereas the aggregate strain common to other SOD1 mutations was not present (Forsberg *et al.*, 2019). The two strains of SOD1 aggregate may represent the amorphous and fibrillar aggregates seen in cell-free assays (Abdolvahabi *et al.*, 2016). Switching between deposition into each form can be controlled with the presence of calcium or the method used to induce aggregation (Stathopoulos *et al.*, 2003; Leal *et al.*, 2013). Solid-state NMR spectroscopy on fibrils formed by the SOD1 β -barrel core, stripped of electrostatic and zinc loops, indicates that β -strands 2, 3 and 5 retain their conformation from the free state. The rest of the molecule adopts a heterogeneous structure with multiple different conformations (Banci *et al.*, 2014). CD spectroscopy on fibrillar metal apo-SOD1 shows that its β -sheet content is increased with respect to soluble metalated and metal-free SOD1 (Banci *et al.*, 2007). Thus, some of the unstructured strands and loops present in the free state reconfigure during the fibrillation process. In cells, SOD1 aggregates appear porous. This facilitates the diffuse movement of soluble proteins and small molecules throughout the inclusion (Matsumoto *et al.*, 2005; Abdolvahabi *et al.*, 2015). SOD1 aggregates are therefore extensively β -structured, with the first three β -strands internalised but greatly heterogeneous in both constituents and conformation when compared to classically amyloid states (Fig. 9a).

Aggregate heterogeneity presents a problem for direct structural study, but much has been learnt by breaking the SOD1 molecule

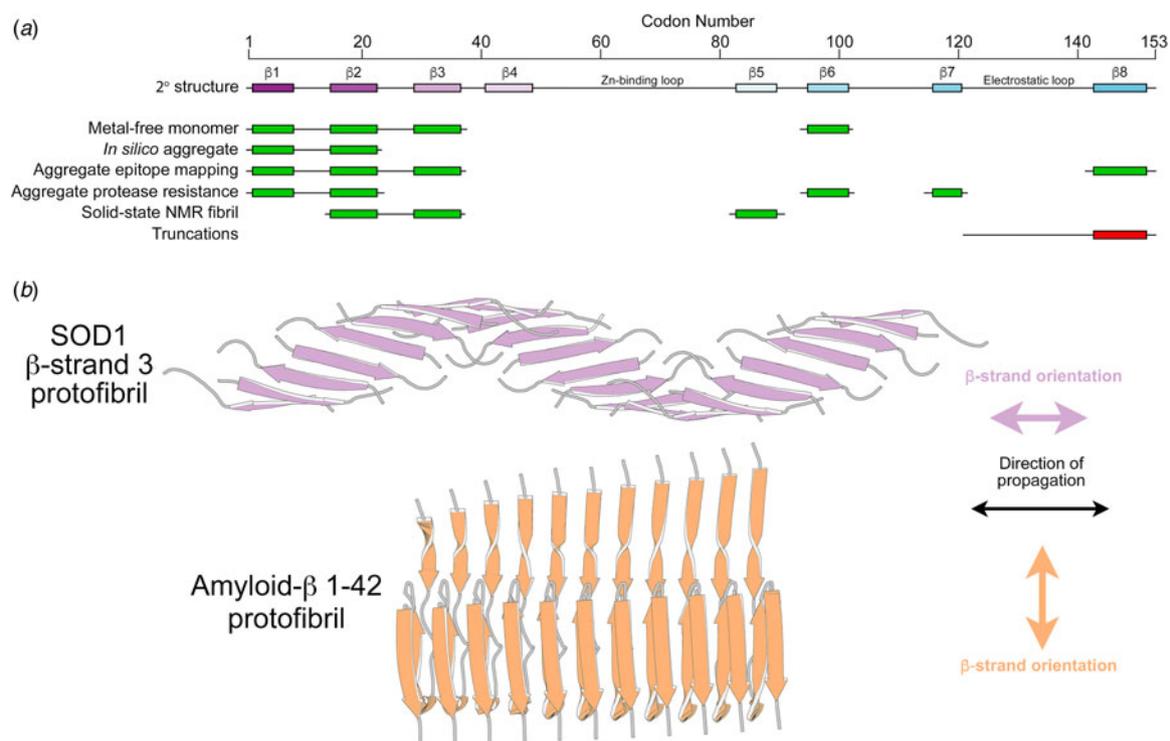


Fig. 9. The structure of SOD1 aggregates. (a) SOD1 β -strands in the N-terminal region of the protein are found to be structured in the metal-free state by NMR (Banci *et al.*, 2009). These are also commonly found to form the core of SOD1 aggregates as determined by modelling, antibody reactivity, resistance to proteolytic cleavage and solid-state NMR (green) while the absence of the C-terminal region in truncated forms of SOD1 negates its role in aggregation (red). (b) SOD1 β -strand 3 forms fibril structures with the alignment of each peptide parallel to the direction of strand elongation in contrast to traditional amyloid where the constituent protein or peptide assembles perpendicular to the direction of propagation.

down into peptides which self-assemble into fibrils amenable to crystallographic structure determination. Using this approach, segments from C-terminal β -strand 8 (residues 147–153) and loop VI (residues 101–107) were shown to form steric zippers through hydrophobic side-chain interactions (Ivanova *et al.*, 2014). Alternatively, a peptide representing β -strand 3 (residues 28–38) formed a twisted β -sheet structure through backbone hydrogen bonding with the individual β -strands aligned in the direction of polymerisation (Sangwan *et al.*, 2017). This contrasts markedly with information on amyloid commonly found in cross- β structures (Fig. 9b) and further explains the subdued response from tinctorial dyes charting SOD1 aggregation.

Aggregate formation

Cellular mechanisms exist to aid protein folding or to destroy proteins which have veered irretrievably off their folding pathway. Part of this functionality separates misfolded species into different cellular compartments (Kaganovich *et al.*, 2008). Intracellular stress granule formation occurs following heat-shock. These dynamic compartments harbour RNAs together with a collection of proteins with RNA-binding capacity coupled with the structural disorder. ALS mutant SOD1 has a tendency to accumulate into distinct domains around preformed stress granules with chaperones and ubiquitin. SOD1-containing stress granules are less dynamic than normal and can be transported along microtubules to JUNQ-like aggresomes (Johnston *et al.*, 2000; Mateju *et al.*, 2017). Localisation of mutant SOD1 to the JUNQ has been demonstrated in human ALS post-mortem spinal motor neurons and is toxic in cell models (Weisberg *et al.*, 2012).

SOD1 aggregates form and grow *in vitro* following a sigmoidal growth curve as any population does in a new environment with limited resources (Fig. 10). Aggregation must first be initiated, and this is marked by a lag period before a change in a parameter such as thioflavin T fluorescence is observed. This period before aggregation nucleation is highly variable and indicative of a stochastic process. Nucleation is followed by exponential growth which plateaus when the available substrate is exhausted (Banci *et al.*, 2007; Chattopadhyay *et al.*, 2008; Abdolvahabi *et al.*, 2016). However, *in vivo* SOD1 translation can supplement available substrate and indefinitely postpone slowing of growth within cells. In addition, a competition exists between recruitment of substrate protomers to fibrillar or amorphous aggregates (Fig. 10) (Abdolvahabi *et al.*, 2016).

Aggregation is a concentration-dependent process (Hofrichter *et al.*, 1974) and this is clearly seen in aggregate formation induced by variable expression levels of human wild-type and mutant SOD1 in transgenic mice (Wong *et al.*, 1995; Graffmo *et al.*, 2013). In humans, SOD1 is posited to be supersaturated based on calculated aggregation propensity and intracellular concentration (Ciryam *et al.*, 2017). The molecular environment also plays a role, with other solutes essentially increasing the effective concentration of SOD1 by molecular crowding thereby promoting fibrillation (Ma *et al.*, 2012). The solubility of SOD1 mutants and the ability of the cell to reduce the prevalence of toxic species through degradation are therefore a key. This may be reflected in the varying aggregation propensity of different SOD1 mutants and the lack of correlation between different experimental systems (McAlary *et al.*, 2016; Abdolvahabi *et al.*, 2017).

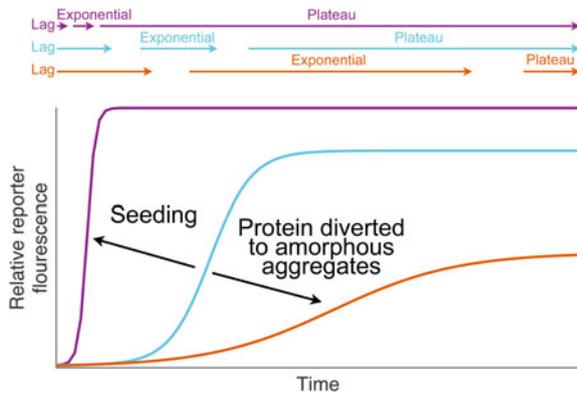


Fig. 10. *SOD1* aggregate growth. *SOD1* fibril growth is commonly assayed *in vitro* using a fluorescent dye such as thioflavin T. Fibril growth is preceded by a lag phase where little aggregation takes place. Aggregation is then nucleated in a stochastic process leading to exponential growth. Growth plateaus when all available *SOD1* has been incorporated into aggregates. Each process can be modified by changing the *SOD1* mutant under investigation, its concentration, adding aggregate seeds or sequestering protein in non-fibrillar aggregates.

Monomeric *SOD1* had been implicated in the pathogenesis of ALS (Hough *et al.*, 2004; Rakhit *et al.*, 2004; Ray *et al.*, 2004) partly in response to the first oxidation-mediated *SOD1* aggregation experiments, but the toxicity of a soluble, low molecular mass species was proven almost a decade later by Brotherton *et al.* (2013). A *SOD1* species existing as a monomeric, metal-free, disulphide-reduced, globally unfolded random coil and is enriched by *SOD1* ALS-related mutations (Luchinat *et al.*, 2014; Lang *et al.*, 2015). However, this completely unstructured monomer must still fold into β -structure-rich fibrils. Anzai *et al.* (2017) and Chattopadhyay *et al.* (2015) proposed a destabilised and aberrantly disulphide-bonded dimer as the protomer species, while Proctor *et al.* (2016) found, with the aid of atomic force microscopy, SEC and modelling, that *SOD1* forms a trimer with non-native dimer interfaces and very substantial structural rearrangement. Banci *et al.* (2007) had previously observed this phenomenon in wild-type *SOD1* and we have seen it occur to monomeric and dimeric Ala4Val *SOD1* (Wright *et al.*, 2013). By mutagenic stabilisation of the interfaces, Proctor *et al.* (2016) found they could trap this trimer on route to high molecular mass aggregates. They also found a correlation between the stability of the trimer and cell death. However, no single species has been definitively demonstrated as pathogenic or to nucleate aggregation, indeed each may exert toxicity, and together they form a continuum on the path to larger aggregates.

Prion-like behaviour of *SOD1*

Assays of *SOD1* aggregation often use mechanical agitation to speed fibrillation by fragmenting growing fibrils thus providing exponentially more termini for the incorporation of protomers. Initiation of fibrillation can also occur by seeding; transfer of wild-type or mutant fibrils from a mature reaction to a lag-phase reaction of the same or a different *SOD1* variant (Fig. 10) (Chia *et al.*, 2010). Importantly, transfer of spinal cord homogenate from mutant *SOD1* transgenic mice to an *in vitro* aggregation reaction has the same effect, demonstrating that these assays mimic *in vivo* aggregate structure and formation of kinetics.

Zilber *et al.* (1963) showed that spinal cord and medulla oblongata homogenates from post-mortem ALS individuals

could induce ALS-like symptoms in macaque monkeys. CNS homogenates or CSF fluid from these monkeys were also competent to induce ALS-like symptoms in other monkeys. Conversely, transfer of homogenates from human ALS patients to non-transgenic mice or Guinea pigs does not yield any ALS-like symptoms but transfer from human mutant *SOD1* patients to mutant *SOD1* transgenic mice reduces survival (Zilber *et al.*, 1963; Bidhendi *et al.*, 2018). Similarly, seeding with mouse mutant spinal cord homogenate reduces the age at which ALS-like symptoms are first observed in a transgenic mouse recipient and their age at the time of death (Ayers *et al.*, 2014b; Bidhendi *et al.*, 2016).

Intracellular *SOD1* is actively excreted into cerebrospinal fluid where it constitutes 75% of the total *SOD* activity (Jacobsson *et al.*, 2001). In addition, misfolded *SOD1* can be secreted by motor neurons and astrocytes and can traverse the extracellular space within exosomes (Urushitani *et al.*, 2006; Gomes *et al.*, 2007). Aggregated *SOD1* is also effectively imported by micropinocytosis facilitating transfer from dead cells to those nearby. This pathway facilitates the transfer of aggregated *SOD1* between cells without direct physical contact and seeds cytoplasmic accumulation of mutant *SOD1* in recipient cells which may not yet have initiated aggregation (Münch *et al.*, 2011; Grad *et al.*, 2014). Misfolded wild-type *SOD1* can be a component of the aggregates transferred from one cell to another, but once misfolding has been seeded, wild-type *SOD1* can continue this function without the necessity of mutant *SOD1* (Grad *et al.*, 2014).

Transgenic mouse experiments have shown that the site of seed inoculation largely dictates the site of initial *SOD1* aggregation and the onset of disease symptoms. This is followed by the spread of aggregates along the neuroaxis and into the brainstem possibly across synaptic junctions (Ayers *et al.*, 2016; Bidhendi *et al.*, 2016). This parallels the human ALS phenotype where symptoms of the disease arise in a single region of the body and spread as the disease progresses (Ravits and La Spada, 2009). Molecular and clinical characteristics of *SOD1*-ALS are therefore strongly reminiscent of the prion diseases where protein misfolding is templated and pathogenic. By implication, a single misfolding or aggregation event is sufficient to trigger the disease and this may account for variability in age of onset and reduced penetrance of some *SOD1* mutants. The propagated misfolding paradigm also opens the somewhat remote possibility of person-to-person transmission as has been observed for amyloid- β (Jaunmuktane *et al.*, 2015).

Impact on disease

Intuitively one would expect the molecular characteristics of mutant *SOD1* to manifest at some level in the disease phenotype in light of the relatively simple inheritance and disease penetrance of many *SOD1* mutants. Armed with a cause and effect hypothesis such as increased aggregation propensity leads to shorter disease duration (Prudencio *et al.*, 2009), and the ability to define this process *in vitro* or in cells, how good are we at predicting the disease? Lindberg *et al.* (2002) first related the stability of mutant, immature *SOD1* to patient survival time after initial diagnosis. Wang *et al.* (2008) developed this using the Chiti-Dobson equation and found that 69% of the variability in disease duration could be accounted for by the synergistic effect of mutant *SOD1* instability and aggregation. This was subsequently disputed in light of DSC and dynamic light-scattering characterisation of several *SOD1* mutants (Vassall *et al.*, 2011). Given the stochasticity

in relatively simple microplate fluorescence assays for SOD1 aggregation (Abdolvahabi *et al.*, 2016), the small sample sizes of each SOD1 ALS mutation together with human genetic, physical and environmental heterogeneity, and the inherent differences in SOD1 mutations themselves sometimes resulting in incomplete penetrance, it will always be difficult to predict clinical milestones. In an effort to combat the latter point, a small angle X-scattering study focussed only on ALS substitutions at the Gly93 site was able to discern differences in their aggregation paralleling disease course following diagnosis (Pratt *et al.*, 2014). A second option is to increase the number of data points and restrict experimental conditions by looking at the relationship between *in vitro* and cell or transgenic models. Lang *et al.* (2015) showed a good inverse linear relationship between mutant transgenic mouse survival times and the ratio of unfolded aggregation precursor to total SOD1 protein. A similar effect was seen in cultured neuronal cells (McAlary *et al.*, 2016).

Are aggregates toxic?

By benchmarking our model systems against the disease, it should be possible to find entities that inhibit disease characteristics in the model system and then apply that knowledge to the disease in humans. This philosophy only functions if we study the toxic species. Aggregates clearly co-segregate with SOD1 mutations and by extension SOD1-related ALS. Furthermore, their presence augurs rapid cell death (Matsumoto *et al.*, 2005). However, true insoluble aggregates only appear at late stages in transgenic mouse models, after or disconnected from the onset of ALS-like symptoms, indicating they may be a response to toxicity or innocuous bi-product rather than a cause. Highlighting this point, Gill *et al.* (2019) found that the amount of aggregated SOD1 in transgenic mouse spinal cord was inversely proportional to disease progression as measured by NeuroScore. Thus, mice fair better if they sequester more SOD1 in aggregates. The authors also found that regions most affected by disease displayed higher soluble misfolded to total soluble SOD1 ratios (Gill *et al.*, 2019). These low molecular mass, soluble misfolded SOD1 species, free of all PTMs are present throughout life (Jonsson *et al.*, 2004; Rakhit *et al.*, 2007; Zetterström *et al.*, 2007; Lee *et al.*, 2015; Gill *et al.*, 2019) and may therefore be a better candidate for the toxic species than inert aggregates.

Mitochondrial toxicity

The pathway from SOD1 mutations to destabilisation, misfolding and aggregation is well delineated. This does not however, tell us why neuronal cells die in SOD1 ALS. Accumulation of ubiquitinated SOD1 into aggregates which also sequester chaperone proteins is a potential source of toxicity as proteostatic activities become impaired. Another possible source of toxicity is the degeneration of mitochondria.

Transgenic mice expressing Gly37Arg human SOD1 develop intraneuronal vacuoles that displace cytoskeletal structures. The vacuoles appear to be mitochondria with intermembrane spaces swollen to the point of membrane structure breakdown (Wong *et al.*, 1995). Vacuolisation appears in the presymptomatic stage and increases in prevalence as disease symptoms manifest and progress but their occurrence is largely restricted to axons and dendrites (Kong and Xu, 1998). SOD1 transported into the mitochondrial IMS in the disulphide-reduced, metal-free state and is retained through hCCS-catalysed maturation. Mutant and wild-

type SOD1 aggregate within mitochondria (Pasinelli *et al.*, 2004; Vijayvergiya *et al.*, 2005; Deng *et al.*, 2006; Cozzolino *et al.*, 2009) leading to membrane disruption (Oladzad Abbasabadi *et al.*, 2013; Salehi *et al.*, 2015; Watanabe *et al.*, 2016). Progressive vacuolisation may be a consequence of this aberrant accumulation. The combined effect of these changes is defective respiration, electron transport and ATP synthesis (Bowling *et al.*, 2006; Magrané *et al.*, 2012).

Membrane association

Misfolded SOD1 is found at high levels around vacuoles in transgenic mouse neurons in what appears to remain of the IMS (Wong *et al.*, 1995; Jaarsma *et al.*, 2001; Graffmo *et al.*, 2013). SOD1 associates with the intracellular face of the plasma membrane but misfolded SOD1 is also found on the outer mitochondrial membrane in ALS spinal cord cells (Petkau and Copps, 1979; Vande Velde *et al.*, 2008). Accumulation is progressive and, importantly, begins just prior to the onset of ALS (Pickles *et al.*, 2013). SOD1, particularly the metal-free state, incorporates within biological membranes and disorders lipid packing (Petkau and Chelack, 1976; Petkau and Copps, 1979; Lepock *et al.*, 1981). Chng and Strange (2014) used molecular dynamics simulations to show how residues in the conformationally dynamic electrostatic and zinc-binding loops bind octanol and mediate interactions with bilayer surfaces. A C-terminal SOD1 truncation at Leu126 facilitates the adoption of a membrane-penetrant helical structure with several transmembrane helices (Lim *et al.*, 2015), and similarly, Ala4Val SOD1 self-association created a tetramer capable of functioning as an integral membrane ion channel (Allen *et al.*, 2012a).

Association of metal-free wild-type with mono- or polyunsaturated fatty acids including arachidonic, oleic and linoleic acid cause SOD1 aggregation (Kim *et al.*, 2005). When incubated with phospholipids, Rasouli *et al.* (2018) found SOD1 aggregation was influenced by the charge of the head group, lipid concentration and the SOD1 variant. For example, anionic phospholipids greatly stimulated metal-free wild-type SOD1 fibril formation over a broad range of concentrations whereas zwitterionic did not. Cationic phospholipids did catalyse SOD1 aggregation but over a much smaller range of concentration than anionic lipids. Thioflavin T binding indicated that His46Arg did not form fibrils and unusually Ala4Val also had limited fibrillation but wild-type, Gly37arg and Asp90Ala did.

Like hCCS, macrophage inhibitory factor (MIF) has a direct and high affinity interaction with both immature wild-type and mutant SOD1 (Israelson *et al.*, 2015; Shvil *et al.*, 2018). MIF inhibits SOD1 deposition onto the outer mitochondrial membrane and chaperones SOD1 away from misfolding within cells and in a cell-free environment (Israelson *et al.*, 2015). MIF overexpression in neuronal cells increases viability while MIF knock-out hastens disease onset and death in SOD1 mutant transgenic mice. MIF has low abundance within motor neurons when compared with other cell types, thus this offers a rationale for motor neuron-selective toxicity despite ubiquitous SOD1 expression.

Secondary SOD1 post-translational modifications

Four PTMs contribute to the stability and function of SOD1. They are, in the sequential order in which they likely occur in the cytosol: zinc and copper acquisition, intra-subunit disulphide bond formation and dimerisation. However, SOD1 is also known to undergo a comprehensive range of secondary PTMs that are

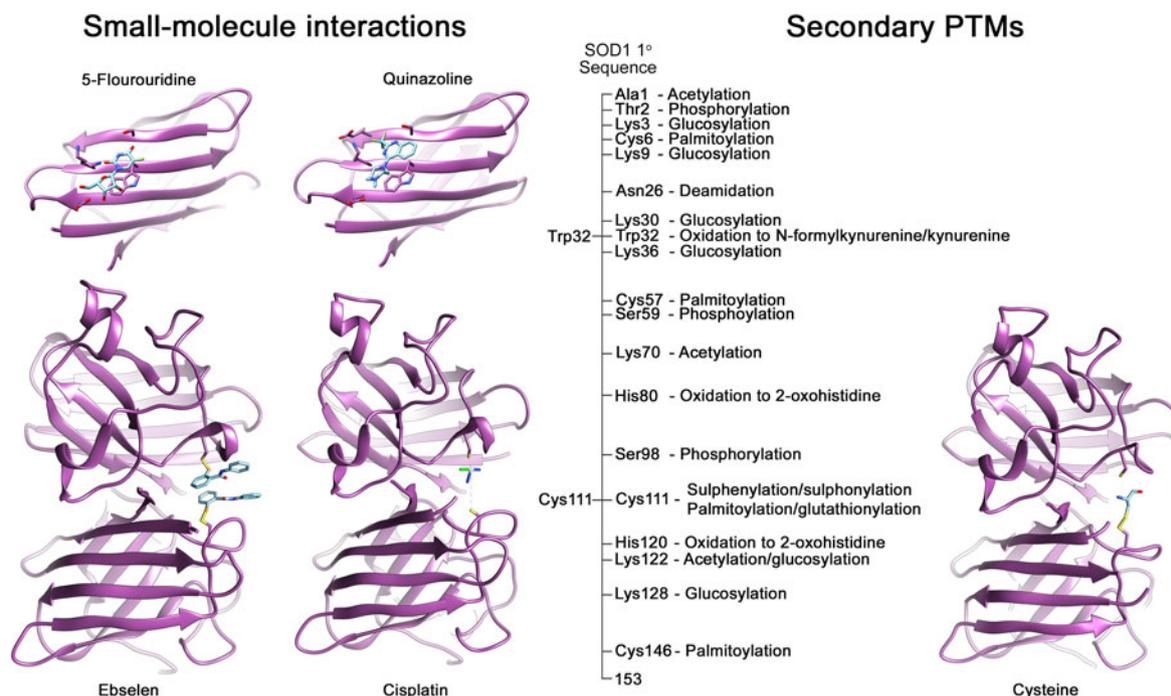


Fig. 11. Secondary post-translational modifications and drug molecule interactions with human SOD1. Post-translational modifications observed *ex vivo* or *in vitro* are listed along with several structures of drug molecule interactions which have been validated crystallographically at Trp32 and Cys111 sites. Cysteinylation of Cys111 is the only native SOD1 post-translational modification which has been crystallographically characterised and was found to be destabilising (Auclair *et al.*, 2013a, 2013b).

not necessary for the formation of the functional dimeric molecule (Fig. 11). These non-inheritable modifications can change the chemical and biophysical properties of SOD1 as do ALS-related mutations and so may have implications for the pathogenesis or treatment of ALS. Here we review only those PTMs that have an association with ALS.

Acetylation

Like many human proteins, the SOD1 N-terminus is cleaved to form acetylalanine (Barra *et al.*, 1980; Jabusch *et al.*, 1980). Covalent addition of an acetyl group can also take place at accessible serine or lysine residues and changes the overall surface charge of a protein. Lysine acetylation is a global regulator of protein activity and signalling. SOD1 Lys70, which is solvent exposed on the surface of the zinc-binding loop, has been observed to be acetylated in cell culture and reduces the amount of soluble SOD1 in a manner similar to ALS mutations. Addition of deacetylase inhibitors increased the prevalence of this modification and decreased SOD1 activity by inhibiting hCCS-dependent activation. Sirtuin-1 is known to act as a deacetylase for SOD1 Lys70 and restores activation by hCCS (Lin *et al.*, 2015). Acetylation of Lys122, on the N-terminus of the electrostatic loop, does not change SOD1 activity but is found only in specific cell types and regions within the CNS indicating functional importance (Kaliszewski *et al.*, 2016). Non-specific lysine acetylation of wild-type, Asp90Ala and Ala4Val SOD1 with aspirin *in vitro* inhibited the nucleation rate and extension of aggregates overall. As the number of acetyl groups increased, the thermal stability of metal-free SOD1 decreased. Destabilisation of acetylated fibril aggregates was also observed and thought to occur by Coulombic repulsion (Abdolvahabi *et al.*, 2015).

Glycation

Protein glycation is the irreversible, stochastic, non-enzymatic addition of a sugar or dicarbonyl to protein amino groups. In practice, addition is usually limited to arginine and lysine side chains. Initial addition is followed by a series of reactions which lead to heterogeneous advanced glycation end-product (AGE) formation. AGEs negatively affect proteins by inhibiting function, crosslinking, decreasing solubility and increasing protease resistance. As a result, they are associated with neurodegenerative diseases such as Alzheimer's and Parkinson's diseases (Li *et al.*, 2012). Arai *et al.* (1987a) found glucose-modified SOD1 to constitute roughly 20% of the erythrocytic SOD1 pool and also found differences in glucosylation between SOD1 from young and old cells. *In vitro* glucosylation occurs at Lys3, Lys9, Lys30, Lys36, Lys122 and Lys128. These modifications reduce SOD1 activity (Arai *et al.*, 1987a, 1987b). Modification of metalated and metal-free wild-type SOD1 with the advanced glycation end-product precursor methylglyoxal induces partial unfolding and, in the case of apoSOD1, the formation of high molecular mass oligomers (Sirangelo *et al.*, 2016). In addition, ALS SOD1 mutant proteins are more easily glucosylated and fructosylated *in vitro* with respect to wild-type (Takamiya *et al.*, 2003). These *in vitro* experiments are of importance because AGE-modified SOD1 is a major component of aggregates found in post-mortem ALS neural tissues (Kato *et al.*, 1999, 2000; Shibata *et al.*, 1999).

Oxidation

Oxidative stress has long been associated with neurodegenerative disease and ALS is a typical example. SOD1 was first observed to aggregate in response to oxidative damage by Rakhit *et al.* (2002). Oxidation of wild-type SOD1 by hydrogen peroxide causes

misfolding and aggregation *in vitro* and in cells. Furthermore, oxidation promotes association with ubiquitin and Hsp70, as is the case in human mutant SOD1 aggregates, and is toxic to motor neuron cultures (Ezzi *et al.*, 2007). There are three sites at which SOD1 is known to be oxidised: Cys111, Trp32 and the copper site histidines. Efficient formation of oxo-histidine-118 bovine SOD1 (His120 human numbering) by hydrogen peroxide has been observed *in vitro* (Uchida and Kawakishi, 1994). Multiple human SOD1 copper site histidines are oxidised by ascorbic acid-CuCl₂, again *in vitro* (Rakhit *et al.*, 2002). There is no evidence of copper site modification in the cellular environment or an impact on ALS but Cys111 and Trp32 both have implications for disease pathogenesis.

Human SOD1 Cys111 residues are solvent exposed but oppose each other 9 Å apart across the sub-unit interface when the protein is in the dimeric form. Their side-chain sulphhydryl groups are reactive (Calabrese *et al.*, 1975). Cysteine is used sparingly in nature due to its reactivity and a cysteine at position 111 is uncommon in the primary sequence of SOD1 orthologues. Its persistence in different phylogenetic groups however could point to a functional role, possibly environmental redox sensing (Go *et al.*, 2015). Cys111 is known to be a key modulator of SOD1 expression level, folding, stability, aggregation and toxicity (Jabusch *et al.*, 1980; Lepock *et al.*, 1990; Hallewell *et al.*, 1991; Suzuki *et al.*, 2011; Chen *et al.*, 2012; Zhang *et al.*, 2017). A potential mechanism for this may be found in the asymmetry and hydrogen bonding network of SOD1 loop VI within which Cys111 is found (Ihara *et al.*, 2012). The Cys111 side-chain sulphhydryl is amenable to a host of post-translational modification reactions, the simplest of which is reversible oxidation to sulphenic acid (Xu *et al.*, 2018). Successive addition of oxygen by hydrogen peroxide or incubation in the air creates sulphinylation and sulphonylation of SOD1. Sulphonylation of wild-type SOD1 Cys111 predisposes it to adopt a Gly93Ala mutant-like conformation and inhibits fast axonal transport (Bosco *et al.*, 2010). The sulphonylated form increases in prevalence as the disease progresses and it also localises to hyaline inclusions found within spinal cord neurons of mutant SOD1 transgenic mice (Fujiwara *et al.*, 2007; Chen *et al.*, 2012; Xu *et al.*, 2018). Finally, Bosco *et al.* (2010) observed immunoreactivity of a conformation-specific antibody which recognises sulphonylated wild-type SOD1 in four of nine sporadic ALS cases tested but not in control individuals or ALS cases with mutations in other genes. This highlights the potentially disastrous effect of irreversible oxidation on SOD1 behaviour.

Mass spectrometry analysis of erythrocytic SOD1 from Gly93Ala and wild-type transgenic mice together with native human wild-type SOD1 showed Trp32 to be post-translationally oxidised (Taylor *et al.*, 2007). Like Cys111, the presence of tryptophan in the mid-point of β -strand 3, where it is unusually solvent exposed for a large non-polar amino acid, is uncommon and again suggests some functional importance. *In vitro* Trp32 is oxidised by hydrogen peroxide in the presence of bicarbonate to N-formylkynurenine and kynurenine or a di-tryptophan species. This promotes misfolding and non-disulphide, covalently linked dimerisation ultimately leading to aggregation (Zhang *et al.*, 2003, 2004; Medinas *et al.*, 2010; Coelho *et al.*, 2014). Trp32 also facilitates the conversion of wild-type SOD1 to a misfolded form by mutant SOD1; a process which is hindered by the substitution of serine at position 32 (Grad *et al.*, 2011). Similarly, substitution of phenylalanine inhibits SOD1 aggregation and prolongs survival of SOD1 mutant cultured neurons (Zhang *et al.*, 2003; Taylor *et al.*, 2007).

Thiolation

S-thiolation is the reversible formation of mixed disulphide bonds between free protein sulphhydryls and low molecular mass cellular thiols such as glutathione or cysteine. These modifications are known to protect cysteine from irreversible oxidation to sulphinate and sulphonate. Modification of SOD1 with cysteine or glutathione results in addition to at most one Cys111 per dimer likely due to lack of space in the inter-dimer groove (Wilcox *et al.*, 2009; Auclair *et al.*, 2013a). The preponderance of glutathionylation *versus* cysteinylation *in vivo* has been disputed. Furthermore, sample preparation under oxidative conditions has been shown to increase the extent of thiolation (Auclair *et al.*, 2014). S-glutathionylation can be spontaneous or glutathione-S-transferase catalysed and often results from oxidative stress. Covalent linkage of glutathione to SOD1 was first observed in human erythrocyte SOD1 with the Asp90Ala mutation (Marklund *et al.*, 1997). The site of glutathione adduct formation was subsequently found to be Cys111 (Wilcox *et al.*, 2009). Glutathionylation of mutant and wild-type SOD1 reduces dimer affinity (Redler *et al.*, 2011; McAlary *et al.*, 2013). Analysing wild-type SOD1 purified from nerve tissue Auclair *et al.*, 2013b found roughly 40% to be cysteinylated at Cys111 in comparison with little cysteinylation in red blood cell SOD1 (Auclair *et al.*, 2014). This modification protected Cys111 from sulphinylation and sulphonylation in response to hydrogen peroxide but decreased the protein's overall thermal stability by 5 °C (Auclair *et al.*, 2013a, 2013b).

Acylation

S-acylation is the addition of a lipid group to a protein cysteine. Palmitoylation is the most common S-acylation and involves the catalysed formation of a covalent but reversible thioester bond between a free sulphhydryl and the fatty acid palmitate. As a result, the hydrophobicity of the target protein is increased and this can facilitate partitioning of otherwise soluble proteins with lipid membranes. Disulphide-reduced SOD1 has been shown to be palmitoylated in cells, transgenic spinal cord lysate and human spinal cord. Modification takes place primarily at Cys6 but to a lesser extent at Cys111 and even Cys57 and Cys146. SOD1 mutants were also shown to have a higher degree of palmitoylation which may be a consequence of their propensity to exist in the disulphide-reduced state. In addition, wild-type SOD1 also showed a higher degree of palmitoylation in sporadic ALS cases. The degree of palmitoylation correlated with the amount of SOD1 associated with cell membranes (Antinone *et al.*, 2013, 2017). Further analysis found hCCS to be palmitoylated in the opposite manner to SOD1 with decreasing levels in mutant SOD1-related ALS and sporadic ALS (Antinone *et al.*, 2017). Reversible palmitoylation could therefore play a role in reducing the three-dimensional search space of hCCS for zinc-bound, disulphide-reduced SOD1 by facilitating membrane association prior to copper and disulphide transfer.

Phosphorylation

SOD1 was first observed to be phosphorylated in myeloid cells after exposure to granulocyte-colony stimulating factor (Csar *et al.*, 2001). It has subsequently been shown to be phosphorylated at Thr2, Ser59 and Ser98 (Wilcox *et al.*, 2009; Tsang *et al.*, 2014). Ser59 and Ser98 are phosphorylated through a direct interaction with Dun1 in response to oxidative stress and this directs SOD1

to the nucleus where it acts as a transcription factor for genes involved in the oxidative stress response (Tsang *et al.*, 2014). While production of phosphorylated SOD1 in amounts necessary for biophysical characterisation is challenging, Fay *et al.* (2016) constructed and crystallised a SOD1 Thr2Asp phosphomimetic mutant. The wild-type SOD1 structure is maintained by the Thr2Asp substitution but strengthens the dimer interaction and slows dissociation rates even when the Ala4Val mutation was present along with glutathionylation at Cys111. In addition, the authors predicted that Thr2Asp destabilises a non-native trimeric structure. As a result, the Thr2Asp substitution reduces cell death resulting from Ala4Val SOD1 expression (Fay *et al.*, 2016).

Deamidation

Deamidation is the removal of the side-chain amide from a glutamine or asparagine residue. Under physiological conditions, the reaction is not enzymatic but slow and may act as a molecular clock (Robinson and Robinson, 2001). The sequential mechanism leads to a mixture of aspartic and isoaspartic acids. Deamidation therefore recapitulates a single amino acid missense mutation. SOD1 deamidation has been observed in erythrocytic SOD1 and occurs naturally to a SOD1 truncation comprising β -strands 1–4 and the disulphide sub-loop (amino acids 1–61 incorporating Cys6/57Ser mutations) *in vitro* during protein purification at Asn26 (Shi *et al.*, 2013b; Bierzynska-Krzysik *et al.*, 2014). Asn26 is positioned in SOD1 loop II within a tripeptide, Ser-Asn-Gly, known to predispose the central residue to deamidation (Wright, 1991). Furthermore, single and compound deamidation replicating Asn/Asp amino acid substitutions created modest reductions in SOD1 thermal stability and increased rate of aggregation (Shi *et al.*, 2013b).

Targeting the SOD1 molecule for therapeutic benefit

Small molecules

Small molecules, produced synthetically or derived from natural sources, formed the mainstay of a revolution in drug discovery that occurred in the latter half of the 20th century and still constitute roughly $\frac{3}{4}$ of new chemical entities registered by the US Food & Drug Administration. Like ALS mutations or natural amino acid modifications, interactions with exogenous small molecules could have positive or negative effects on SOD1 behaviours associated with ALS. Several small-molecule strategies have been developed to address the underlying folding, metalation, stability and aggregation characteristics associated with ALS mutant SOD1 and target-specific sites on the SOD1 molecule.

The Trp32 site

Human SOD1 has a solvent-exposed tryptophan residue on the surface of β -strand 3. This residue is poorly conserved with threonine, serine, asparagine, glutamate, lysine and arginine all preferred across other eukaryotic CuZnSOD analogues. Indeed, only a few simian species and the African elephant are known to incorporate tryptophan at this position while a conservative phenylalanine substitution is found at the equivalent position in human extracellular CuZnSOD (SOD3) indicating hydrophobicity at this site has some functionality. Accordingly, Pokrishevsky *et al.* (2018) showed that substitution of Trp32 for the more frequently found serine decreases mutant and wild-type SOD1 stability and resistance to proteolytic digestion.

Despite reduced stability, the Trp32Ser substitution confers a protective effect on SOD1 aggregation. This includes slowed aggregation kinetics, an inability to propagate Trp32 SOD1 seeded aggregation, and a propensity to form amorphous rather than fibrillar aggregates (Grad *et al.*, 2011; Pokrishevsky *et al.*, 2018). β -strand 3 has been found in the core of SOD1 aggregates (Bergh *et al.*, 2015) so removing a hydrophobic interface may hinder fibril formation. All likely contribute to reduced loss of motor function on the expression of Trp32Ser human SOD1 in zebra fish in comparison with human wild-type (DuVal *et al.*, 2019). Thus, inhibiting abnormal SOD1 self-interactions at Trp32 is a possible strategy to prevent aggregation and toxicity.

Several small molecules have been shown crystallographically to bind at or near the Trp32 site. This includes a series of aromatics which exploit a hydrophobic interaction with the Trp32 indole ring system (Fig. 11) (Antonyuk *et al.*, 2010; Kershaw *et al.*, 2013; Ramu *et al.*, 2018) and four catecholamines that interact with a groove in the SOD1 surface created by loop II close to Trp32 (Wright *et al.*, 2013). A fragment-linking approach created one compound which was able to bind the Trp32-loop II site with low micromolar affinity and inhibit Trp32 oxidation (Manjula *et al.*, 2018), a post-translational modification known to facilitate aggregation (Taylor *et al.*, 2007).

5-Fluorouridine was also shown crystallographically to interact with the SOD1 Trp32 site (Fig. 11) (Wright *et al.*, 2013). Inclusion of 5-fluorouridine in cell culture media prior to exposure to human SOD1-ALS spinal cord homogenates could significantly reduce induced SOD1 aggregation (Pokrishevsky *et al.*, 2017). 5-Fluorouridine, uridine and telbivudine have also been shown to have positive effects in cell and animal model systems including postponing disease onset and increasing survival in Gly93Ala transgenic mice (Pokrishevsky *et al.*, 2018; DuVal *et al.*, 2019; Rando *et al.*, 2019).

The Cys111 site

Like Trp32, SOD1 Cys111 is poorly conserved and oxidation of its sulphhydryl has a role in aggregation (Chen *et al.*, 2012; Xu *et al.*, 2018). Opposing Cys111 residues are separated by a short distance across the SOD1 dimer interface. Covalent tethering of Cys111 side-chain sulphhydryls by maleimide cross-linking was shown to substantially increase mutant SOD1 thermal stability (Auclair *et al.*, 2010). This experiment is reminiscent of covalent tethering of the SOD1 dimers by the introduction of a Val148Cys substitution performed by Ray *et al.* (2004) which, taking cues from lysozyme and thymidylate synthase research, was able to reinforce Ala4Val SOD1 dimer affinity and prevent *in vitro* aggregation. We were able to mimic these effects by binding the small, antioxidant molecule ebselen to opposing Cys111 residues where they increased dimer affinity of Ala4Val SOD1 nearly 60-fold. In a second mode of action, ebselen was also an efficient catalyst for SOD1 disulphide bond formation which restored normal folding to Ala4Val and Gly93Ala mutants within cells (Capper *et al.*, 2018).

Cisplatin addition to Cys111 prevents and even reverses aggregation of metal-free SOD1 in cell-free and cell-based models. The affinity of this interaction was found to be 37 μ M and cisplatin was able to increase metal-free SOD1 thermal stability by 8 °C (Banci *et al.*, 2012b). Crystallographic determination of the SOD1-cisplatin structure shows the ligand positioned at Cys111 (Fig. 11) but full ligand occupancy is not possible due to steric clashes across the dimer interface and cisplatin geometry is significantly distorted (Banci *et al.*, 2012b; Shabalina *et al.*, 2015).

Metal-coordinating sites

An inability to correctly metalate is a recurring characteristic of SOD1 mutants and increasing the availability of zinc, copper or both is a logical response. High doses of zinc are toxic to Gly93Ala transgenic mice but doses up to 60 mg kg⁻¹ day⁻¹ were tolerated. A higher dose of 90 mg kg⁻¹ day⁻¹ was also tolerated when supplemented with 0.3 mg kg⁻¹ day⁻¹ copper. In each case, mouse survival was increased by roughly 10 days on average (Ermilova *et al.*, 2005). A subsequent 2011 ALS clinical trial assessed a similar mixture of zinc and copper in humans for toxicity but details of progression beyond phase I/II are sparse.

Taking the alternative approach of increasing copper availability, diacetylbis(N(4)-methylthiosemicarbazonato) copper (II) (CuATSM) was shown to increase survival in Gly93Ala, Gly37Arg and Gly93Ala/hCCS overexpressing transgenic mice in a dose-dependent fashion (Soon *et al.*, 2011; McAllum *et al.*, 2013; Williams *et al.*, 2016; Vieira *et al.*, 2017). In a cell model, CuATSM increases SOD1 abundance and activity while reducing aggregation and cell death but only for non-metal-binding site mutants (Farrarwell *et al.*, 2019). CuATSM has recently completed a phase I clinical trials with a further phase I dose escalation trial underway and a phase II trial began on 30 September with completion date of 30 December 2020.

Promoting production of stable SOD1 over unfolded or aggregated species with pharmaceutical chaperones has been one of the two most explored avenues for SOD1-specific therapeutic development. What holds these strategies back as effective treatments for SOD1 ALS are combinations of toxicity, specificity, affinity and delivery; problems often associated with drug discovery but especially difficult to overcome when the target is in the central nervous system.

Biologics

Biologics, including antibodies and engineered proteins, offer a solution to the specificity problem above. The SOD1 interactome is relatively small but several interaction partners have the potential for therapeutic repurposing. This repertoire has been broadened by surface display techniques, directed evolution and novel antibody generation providing us with many proteins which are able to specifically interact with disulphide-reduced, metal-free, aggregated or soluble misfolded SOD1.

Antibodies

The production of antibodies which react specifically with SOD1 structural conformations commonly found in ALS has revolutionised our understanding of the SOD1 structure–misfunction relationship in the last decade or so. These antibodies are often termed misfolding-specific because they preferentially bind epitopes that are not accessible when SOD1 is mature, either through internalisation or reduced conformational dynamics. The first such antibody was a polyclonal generated against a peptide found in β -strand 8 including the disulphide-forming Cys146 and specifically recognised monomeric SOD1 (Rakhit *et al.*, 2007). Together with analytical or diagnostic use, conformation or mutant-specific antibodies may also have therapeutic applications. While SOD1 aggregates are found within cells, there is strong evidence that aggregates or small misfolded species can traverse the extracellular space and seed aggregation in surrounding cells. If this toxic transport could be inhibited, the progressive nature of ALS could be arrested. Justification for this comes in part from work showing the presence of native SOD1 reactive

antibodies in sporadic ALS sufferers with longer than mean survival (van Blitterswijk *et al.*, 2011).

The D3H5 IgG antibody was generated against recombinant, metal-free Gly93Ala SOD1 and when delivered directly to the cerebral ventricles was able to postpone weight loss and increase life span by small amounts. Increasing the period over which the antibody was administered and starting the treatment earlier increased survival from +6 to +9 days compared to control Gly93Ala transgenic mice (Gros-Louis *et al.*, 2010).

Active immunisation with peptides designed to create a native immunogenic response specific for misfolded SOD1 has also been shown to increase life-span and slow disease progression in transgenic mice (Urushitani *et al.*, 2007; Liu *et al.*, 2012). However, selection of the immunising peptide has to be carefully considered to not induce an immune response which in itself can be life threatening (Sábado *et al.*, 2015).

Single-chain recognition proteins

High-affinity antibodies can now be generated through traditional protocols cheaply and efficiently. However, their size, flexibility and quaternary structure present a problem for structural study, rational redesign and intracellular delivery. Single-chain recognition domains circumvent these problems and have found interesting uses in relation to SOD1.

Patents have been filed for SOD1-reactive nanobodies; small, single-chain antibodies often generated by immunisation of camelids. Micro-nanomolar binding affinity for wild-type and mutant SOD1 has been described which was able to prevent the formation of large Ala4Val SOD1 fibrils *in vitro* with some protein forming small amorphous aggregates. Expression of the nanobody within cells reduced SOD1 aggregation and a single direct delivery of nanobody to Gly93Ala SOD1 transgenic mouse brain delayed disease onset and prolonged survival (Robberecht *et al.*, 2016).

A thermophilic variant of the immunoglobulin-binding protein G B1 domain (HTB1) was optimised for SOD1 binding by directing mutations to putative stable regions on the surface of the protein followed by yeast surface display (Banerjee *et al.*, 2017). Unmodified HTB1 could not bind SOD1 while the product of this process was a binding domain with 300 and 674 nM dissociation constants for Gly93Ala and Gly37Arg, respectively. *In vitro* aggregation in the presence of modified HTB1 showed SOD1 forming small amorphous aggregates rather than large fibrils, reduced intracellular SOD1 misfolding and increased cell viability. The versatility of this technique is illustrated by the successful retargeting of HTB1 to bind amyloid- β peptide with low micromolar affinity with positive effects on aggregation and toxicity (Banerjee *et al.*, 2018).

Using phage display in bacteria, Ghadge *et al.* (2013) created two single chain variable fragments (scFv) from a pre-existing library which were able to interact with wild-type, Ala4Val and Val148Gly SOD1 but not Gly93Ala SOD1 possibly indicating Gly93 forms part of the protein–protein interface. When co-expressed with Ala4Val SOD1 in neuronal-like cells, SOD1 aggregation was reduced and cell viability was increased. Ghadge *et al.* (2018) then showed that scFV adeno-associated viral (AAV) delivery to Gly93Ala transgenic mice increased survival when administered neonatally and just prior to the onset of symptoms. On the whole, single chain recognition domains do not replicate the very high target affinity that can be found with well-selected antibodies. However, when combined with reliable, efficient and targeted gene delivery by AAV vectors, these

small proteins are a very promising new strategy to engage and change the fate of many cellular proteins including SOD1.

Conclusion

After more than 50 years of concerted effort, we have gained a very detailed understanding of SOD1 biology. Despite this body of work, new SOD1 functions and behaviour are regularly discovered including, in recent years, a role in signalling, stress granule formation and transcriptional regulation. Perhaps the most enigmatic behaviour of SOD1 discovered over these years is its propensity to aggregate and the ramifications this has on ALS.

We now understand how ALS mutations cause aggregation in SOD1-ALS. Generally, mutations to conserved residues in the metal-binding, disulphide, dimer interface half of the SOD1 β -barrel prevent adequate folding and acquisition of a stable structure. These unfolded structures predispose mutant SOD1 to degradation but also lead to the accumulation of the non-conserved part of the barrel into non-amyloid fibrils or amorphous aggregates. While the exact structure of these aggregates remains unresolved, the research community is developing the tools with which to arrest their formation. Small-molecule therapies including CuATSM and ebselen confer order on the functional part of the SOD1 β -barrel while Trp32-targeting compounds bind in the region known to form the core of SOD1 aggregates. What currently limits the usefulness of these compounds as therapeutics, either singly or as a cocktail, is a combination of affinity, specificity and delivery. Honing delivery of zinc and copper, the specificity of disulphide formation and the affinity of anti-aggregation compounds will be necessary for this strategy to succeed in the clinic.

The presence of SOD1 aggregates in mutant SOD1 cases of ALS is unquestionable but there is also strong evidence that wild-type SOD1 is a component of aggregates in at least some sporadic ALS cases (Shibata *et al.*, 1994, 1996b). There are also many reports of misfolded SOD1 in sporadic ALS cases (Bosco *et al.*, 2010; Forsberg *et al.*, 2010; Pokrishevsky *et al.*, 2012; Grad *et al.*, 2014; Paré *et al.*, 2018). Some of the secondary post-translational modifications described in 'Secondary SOD1 post-translational modifications' section may be responsible for destabilising wild-type SOD1 and lead to misfolding (Bosco *et al.*, 2010). This topic is however contentious; Da Cruz *et al.* (2017) found no SOD1 misfolding in sporadic ALS samples. If differences in detection are eliminated, this discrepancy must arise from the tissue source, inherent variability of patient samples and subjective data interpretation.

Heterogeneity is common in ALS and we should not be surprised that some sporadic cases, or tissues therein, exhibit SOD1 misfolding and aggregation while others do not. The size of aggregates, the abundance of misfolded SOD1 and the proportion of cells that display these characteristics is also key here. Discovery of misfolded SOD1 in non-SOD1 fALS cases, Parkinson's disease and aged but healthy individuals supports the notion that SOD1 aggregates are a commonality in ageing and neurodegenerative disease (Nishiyama *et al.*, 1995; Trist *et al.*, 2017; Forsberg *et al.*, 2019). In each case, the size and number of these inclusions was lower than is common in SOD1-ALS patients, thus it may be that investigators expect to see large mutant SOD1-like aggregates when much smaller, punctate forms are missed in non-SOD1 ALS tissue samples.

If SOD1 aggregation is common to mutant SOD1-ALS, non-SOD1 ALS, neurodegenerative disease in general and ageing,

is it toxic? The toxicity of protein or peptide aggregates has been questioned recently across neurodegenerative disease research. Low molecular mass misfolded species including SOD1 trimers with non-native secondary, tertiary and quaternary structure have been posited to be toxic (Zhu *et al.*, 2018). These species may exist within cells but there is currently no clear rationale or mechanistic understanding as to their toxicity. Basic science examining this concept is of the upmost importance if we are to create new, viable therapeutics to specifically address these species.

The linkage of SOD1 with ALS in 1993 stimulated a proliferation in research and understanding of SOD1 biology. Throughout, biophysics has been at the forefront, and as we find answers to the questions highlighted above, biophysics will also doubtless continue to be instrumental.

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References

- Abdolvahabi A, Shi Y, Rhodes NR, Cook NP, Martí AA and Shaw BF (2015) Arresting amyloid with Coulomb's law: acetylation of ALS-linked SOD1 by aspirin impedes aggregation. *Biophysical Journal* **108**, 1199–1212.
- Abdolvahabi A, Shi Y, Chuprin A, Rasouli S and Shaw BF (2016) Stochastic formation of fibrillar and amorphous superoxide dismutase oligomers linked to amyotrophic lateral sclerosis. *ACS Chemical Neuroscience* **7**, 799–810.
- Abdolvahabi A, Shi Y, Rasouli S, Croom CM, Aliyan A, Marti AA and Shaw BF (2017) Kaplan-Meier meets chemical kinetics: intrinsic rate of SOD1 amyloidogenesis decreased by subset of ALS mutations and cannot fully explain age of disease onset. *ACS Chemical Neuroscience* **8**, 1378–1389.
- Abel O, Powell JF, Andersen PM and Al-Chalabi A (2012) ALSod: a user-friendly online bioinformatics tool for amyotrophic lateral sclerosis genetics. *Human Mutation* **33**, 1345–1351.
- Abernethy JL, Steinman HM and Hill RL (1974) Bovine erythrocyte superoxide dismutase. Subunit structure and sequence location of the intrasubunit disulfide bond. *Journal of Biological Chemistry* **249**, 7339–7347.
- Aggarwal A and Nicholson G (2005) Age dependent penetrance of three different superoxide dismutase 1 (sod 1) mutations. *The International Journal of Neuroscience* **115**, 1119–1130.
- Alexander MD, Traynor BJ, Miller N, Corr B, Frost E, McQuaid S, Brett FM, Green A and Hardiman O (2002) 'True' sporadic ALS associated with a novel SOD-1 mutation. *Annals of Neurology* **52**, 680–683.
- Allen S and Dennison C (2014) The importance of Zn(ii) binding by the human copper metallochaperone for Cu,Zn-superoxide dismutase. *RSC Advances* **4**, 22542–22544.
- Allen MJ, Lacroix JJ, Ramachandran S, Capone R, Whitlock JL, Ghadge GD, Arnsdorf MF, Roos RP and Lal R (2012a) Mutant SOD1

- forms ion channel: implications for ALS pathophysiology. *Neurobiology of Disease* **45**, 831–838.
- Allen S, Badarau A and Dennison C** (2012b) Cu(I) affinities of the domain 1 and 3 sites in the human metallochaperone for Cu,Zn-superoxide dismutase. *Biochemistry* **51**, 1439–1448.
- Alvarez-Zaldiernas C, Lu J, Zheng Y, Yang H, Blasi J, Solsona C and Holmgren A** (2016) Cellular redox systems impact the aggregation of Cu-Zn superoxide dismutase linked to familial amyotrophic lateral sclerosis. *Journal of Biological Chemistry* **291**, 17197–17208.
- Andersen PM, Nilsson P, Ala-Hurula V, Keränen ML, Tarvainen I, Haltia T, Nilsson L, Binzer M, Forsgren L and Marklund SL** (1995) Amyotrophic lateral sclerosis associated with homozygosity for an Asp90Ala mutation in CuZn-superoxide dismutase. *Nature Genetics* **10**, 61–66.
- Andersen PM, Nilsson P, Keränen ML, Forsgren L, Hägglund J, Karlsborg M, Ronnevi LO, Gredal O and Marklund SL** (1997) Phenotypic heterogeneity in motor neuron disease patients with CuZn-superoxide dismutase mutations in scandinavia. *Journal of Brain and Neurology* **120**, 1723–1737.
- Andersen PM, Restagno G, Stewart HG and Chiò A** (2004) Disease penetrance in amyotrophic lateral sclerosis associated with mutations in the SOD1 gene. *Annals of Neurology* **55**, 298–299.
- Andersen PM, Forsgren L, Binzer M, Nilsson P, Ala-Hurula V, Keränen ML, Bergmark L, Saarinen A, Haltia T, Tarvainen I, Kinnunen E, Udd B and Marklund SL** (1996) Autosomal recessive adult-onset amyotrophic lateral sclerosis associated with homozygosity for Asp90Ala CuZn-superoxide dismutase mutation. A clinical and genealogical study of 36 patients. *Journal of Brain and Neurology* **119**, 1153–1172.
- Andersen PM, Sims KB, Xin WW, Kiely R, O'Neill G, Ravits J, Pioro E, Harati Y, Brower RD, Levine JS, Heinicke HU, Seltzer W, Boss M and Brown RH Jr** (2003) Sixteen novel mutations in the Cu/Zn superoxide dismutase gene in amyotrophic lateral sclerosis: a decade of discoveries, defects and disputes. *Amyotrophic Lateral Sclerosis* **4**, 62–73.
- Antinone SE, Ghadge GD, Lam TT, Wang L, Roos RP and Green WN** (2013) Palmitoylation of superoxide dismutase 1 (SOD1) is increased for familial ALS-linked SOD1 mutants. *Journal of Biological Chemistry* **288**, 21606–21617.
- Antinone SE, Ghadge GD, Ostrow LW, Roos RP and Green WN** (2017) S-acylation of SOD1, CCS, and a stable SOD1-CCS heterodimer in human spinal cords from ALS and non-ALS subjects. *Scientific Reports* **7**, 41141.
- Antonyuk S, Strange RW and Hasnain SS** (2010) Structural discovery of small molecule binding sites in Cu-Zn human superoxide dismutase familial amyotrophic lateral sclerosis mutants provides insights for lead optimization. *Journal of Medicinal Chemistry* **53**, 1402–1406.
- Anzai I, Tokuda E, Mukaiyama A, Akiyama S, Endo F, Yamanaka K, Misawa H and Furukawa Y** (2017) A misfolded dimer of Cu/Zn-superoxide dismutase leading to pathological oligomerization in amyotrophic lateral sclerosis. *Protein Science* **26**, 484–496.
- Aoki M, Ogasawara M, Matsubara Y, Narisawa K, Nakamura S, Itoyama Y and Abe K** (1993) Mild ALS in Japan associated with novel SOD mutation. *Nature Genetics* **5**, 323–324.
- Arai K, Iizuka S, Tada Y, Oikawa K and Taniguchi N** (1987a) Increase in the glycosylated form of erythrocyte Cu-Zn-superoxide dismutase in diabetes and close association of the nonenzymatic glycosylation with the enzyme activity. *Biochimica et Biophysica Acta* **924**, 292–296.
- Arai K, Maguchi S, Fujii S, Ishibashi H, Oikawa K and Taniguchi N** (1987b) Glycation and inactivation of human Cu-Zn-superoxide dismutase. Identification of the in vitro glycosylated sites. *Journal of Biological Chemistry* **262**, 16969–16972.
- Aresano F, Banci L, Bertini I, Martinelli M, Furukawa Y and O'Halloran TV** (2004) The unusually stable quaternary structure of human Cu,Zn-superoxide dismutase 1 is controlled by both metal occupancy and disulfide status. *Journal of Biological Chemistry* **279**, 47998–48003.
- Auclair JR, Boggio KJ, Petsko GA, Ringe D and Agar JN** (2010) Strategies for stabilizing superoxide dismutase (SOD1), the protein destabilized in the most common form of familial amyotrophic lateral sclerosis. *Proceedings of the National Academy of Sciences of the USA* **107**, 21394–21399.
- Auclair JR, Brodtkin HR, D'Aquino JA, Petsko GA, Ringe D and Agar JN** (2013a) Structural consequences of cysteinylolation of Cu/Zn-superoxide dismutase. *Biochemistry* **52**, 6145–6150.
- Auclair JR, Johnson JL, Liu Q, Salisbury JP, Rotunno M, Petsko GA, Ringe D, Brown Jr RH, Bosco DA and Agar JN** (2013b) Post-Translational modification by cysteine protects Cu/Zn-superoxide dismutase from oxidative damage. *Biochemistry* **52**, 6137–6144.
- Auclair JR, Salisbury JP, Johnson JL, Petsko GA, Ringe D, Bosco DA, Agar NYR, Santagata S, Durham HD and Agar JN** (2014) Artifacts to avoid while taking advantage of top-down mass spectrometry based detection of protein S-thiolation. *Proteomics* **14**, 1152–1157.
- Ayers J, Lelie H, Workman A, Prudencio M, Brown H, Fromholt S, Valentine J, Whitelegge J and Borchelt D** (2014a) Distinctive features of the D101N and D101 G variants of superoxide dismutase 1; two mutations that produce rapidly progressing motor neuron disease. *Journal of Neurochemistry* **128**, 305–314.
- Ayers JI, Fromholt S, Koch M, DeBosier A, McMahon B, Xu G and Borchelt DR** (2014b) Experimental transmissibility of mutant SOD1 motor neuron disease. *Acta Neuropathologica* **128**, 791–803.
- Ayers JI, Fromholt SE, O'Neal VM, Diamond JH and Borchelt DR** (2016) Prion-like propagation of mutant SOD1 misfolding and motor neuron disease spread along neuroanatomical pathways. *Acta Neuropathologica* **131**, 103–114.
- Baek W, Koh S-H, Park JS, Kim YS, Kim HY, Kwon MJ, Ki C-S and Kim SH** (2011) A novel codon4 mutation (A4F) in the SOD1 gene in familial amyotrophic lateral sclerosis. *Journal of the Neurological Sciences* **306**, 157–159.
- Bali T, Self W, Liu J, Siddique T, Wang LH, Bird TD, Ratti E, Atassi N, Boylan KB, Glass JD, Maragakis NJ, Caress JB, McCluskey LE, Appel SH, Wymer JP, Gibson S, Zinman L, Mozaffar T, Callaghan B, McVey AL, Jockel-Balsarotti J, Allred P, Fisher ER, Lopate G, Pestronk A, Cudkowicz ME and Miller TM** (2017) Defining SOD1 ALS natural history to guide therapeutic clinical trial design. *Journal of Neurology Neurosurgery & Psychiatry* **88**, 99–105.
- Banci L, Bertini I, Cabelli DE, Hallewell RA, Tung JW and Viezzoli MS** (1991) A characterization of copper/zinc superoxide dismutase mutants at position 124. Zinc-deficient proteins. *European Journal of Biochemistry FEBS* **196**, 123–128.
- Banci L, Bertini I, Chiu CY, Mullenbach GT and Viezzoli MS** (1995) Synthesis and characterization of a monomeric mutant Cu/Zn superoxide dismutase with partially reconstituted enzymic activity. *European Journal of Biochemistry* **234**, 855–860.
- Banci L, Benedetto M, Bertini I, Del Conte R, Piccioli M and Viezzoli MS** (1998) Solution structure of reduced monomeric Q133M2 copper, zinc superoxide dismutase (SOD). Why is SOD a dimeric enzyme? *Biochemistry* **37**, 11780–11791.
- Banci L, Bertini I, Cramaro F, Del Conte R and Viezzoli MS** (2002) The solution structure of reduced dimeric copper zinc superoxide dismutase. The structural effects of dimerization. *European Journal of Biochemistry FEBS* **269**, 1905–1915.
- Banci L, Bertini I, D'Amelio N, Gaggelli E, Libralesso E, Matecko I, Turano P and Valentine JS** (2005) Fully metallated S134N Cu, Zn-superoxide dismutase displays abnormal mobility and intermolecular contacts in solution. *Journal of Biological Chemistry* **280**, 35815–35821.
- Banci L, Bertini I, Cantini F, D'Amelio N and Gaggelli E** (2006) Human SOD1 before harboring the catalytic metal: solution structure of copper-depleted, disulfide-reduced form. *Journal of Biological Chemistry* **281**, 2333–2337.
- Banci L, Bertini I, Durazo A, Giroto S, Gralla EB, Martinelli M, Valentine JS, Vieru M and Whitelegge JP** (2007) Metal-free superoxide dismutase forms soluble oligomers under physiological conditions: a possible general mechanism for familial ALS. *Proceedings of the National Academy of Sciences of the USA* **104**, 11263–11267.
- Banci L, Bertini I, Boca M, Calderone V, Cantini F, Giroto S and Vieru M** (2009) Structural and dynamic aspects related to oligomerization of apo SOD1 and its mutants. *Proceedings of the National Academy of Sciences of the USA* **106**, 6980–6985.

- Banci L, Bertini I, Ciofi-Baffoni S, Kozyreva T, Zovo K and Palumaa P (2010) Affinity gradients drive copper to cellular destinations. *Nature* **465**, 645–648.
- Banci L, Bertini I, Cantini F, Kozyreva T, Massagni C, Palumaa P, Rubino JT and Zovo K (2012a) Human superoxide dismutase 1 (hSOD1) maturation through interaction with human copper chaperone for SOD1 (hCCS). *Proceedings of the National Academy of Sciences of the USA* **109**, 13555–13560.
- Banci L, Barbieri L, Bertini I, Luchinat E, Secci E, Zhao Y and Aricescu AR (2013) Atomic-resolution monitoring of protein maturation in live human cells by NMR. *Nature Chemical Biology* **9**, 297–299.
- Banci L, Blaževič O, Cantini F, Danielsson J, Lang L, Luchinat C, Mao J, Oliveberg M and Ravera E (2014) Solid-state NMR studies of metal-free SOD1 fibrillar structures. *Journal of Biological Inorganic Chemistry JBIC* **19**, 659–666.
- Banci L, Bertini I, Blaževič O, Calderone V, Cantini F, Mao J, Trapananti A, Vieru M, Amori I, Cozzolino M and Carri MT (2012b) Interaction of cisplatin with human superoxide dismutase. *Journal of the American Chemical Society* **134**, 7009–7014.
- Banerjee V, Oren O, Ben-Zeev E, Taube R, Engel S and Papo N (2017) A computational-combinatorial approach identifies a protein inhibitor of superoxide dismutase 1 misfolding, aggregation, and cytotoxicity. *Journal of Biological Chemistry* **292**, 15777–15788.
- Banerjee V, Oren O, Dagan B, Taube R, Engel S and Papo N (2018) An engineered variant of the B1 domain of protein G suppresses the aggregation and toxicity of intra- and extracellular A β 42. *ACS Chemical Neuroscience* **10**, 1488–1496.
- Barra D, Martini F, Bannister JV, Schinina ME, Rotilio G, Bannister WH and Bossa F (1980) The complete amino acid sequence of human Cu/Zn superoxide dismutase. *FEBS Letters* **120**, 53–56.
- Battistini S, Ricci C, Lotti EM, Benigni M, Gagliardi S, Zucco R, Bondavalli M, Marcello N, Ceroni M and Cereda C (2010) Severe familial ALS with a novel exon 4 mutation (L106F) in the SOD1 gene. *Journal of the Neurological Sciences* **293**, 112–115.
- Beauchamp CO and Fridovich I (1973) Isozymes of superoxide dismutase from wheat germ. *Biochimica et Biophysica Acta (BBA) – Protein Structure* **317**, 50–64.
- Beckman G (1973) Population studies in northern Sweden. VI. Polymorphism of superoxide dismutase. *Hereditas* **73**, 305–310.
- Beers DR, Henkel JS, Xiao Q, Zhao W, Wang J, Yen AA, Siklos L, McKercher SR and Appel SH (2006) Wild-type microglia extend survival in PU.1 knockout mice with familial amyotrophic lateral sclerosis. *Proceedings of the National Academy of Sciences of the USA* **103**, 16021–16026.
- Benatar M (2007) Lost in translation: treatment trials in the SOD1 mouse and in human ALS. *Neurobiology of Disease* **26**, 1–3.
- Bergemalm D, Forsberg K, Srivastava V, Graffmo KS, Andersen PM, Brännström T, Wingsle G and Marklund SL (2010) Superoxide dismutase-1 and other proteins in inclusions from transgenic amyotrophic lateral sclerosis model mice. *Journal of Neurochemistry* **114**, 408–418.
- Bergh J, Zetterström P, Andersen PM, Brännström T, Graffmo KS, Jonsson PA, Lang L, Danielsson J, Oliveberg M and Marklund SL (2015) Structural and kinetic analysis of protein-aggregate strains *in vivo* using binary epitope mapping. *Proceedings of the National Academy of Sciences of the USA* **112**, 4489–4494.
- Bertolin C, D'Ascenzo C, Querin G, Gaiani A, Boaretto F, Salvaro C, Vazza G, Angelini C, Cagnin A, Pegoraro E, Sorarù G and Mostacciolo ML (2014) Improving the knowledge of amyotrophic lateral sclerosis genetics: novel SOD1 and FUS variants. *Neurobiology of Aging* **35**, 1212.e7–1212.e10.
- Bidhendi EE, Bergh J, Zetterström P, Andersen PM, Marklund SL and Brännström T (2016) Two superoxide dismutase prion strains transmit amyotrophic lateral sclerosis-like disease. *Journal of Clinical Investigation* **126**, 2249–2253.
- Bidhendi EE, Bergh J, Zetterström P, Forsberg K, Pakkenberg B, Andersen PM, Marklund SL and Brännström T (2018) Mutant superoxide dismutase aggregates from human spinal cord transmit amyotrophic lateral sclerosis. *Acta Neuropathologica* **136**, 939–953.
- Bierczyńska-Krzysik A, Łopaciuk M, Pawlak-Morka R and Stadnik D (2014) Investigation of asparagine deamidation in a SOD1-based biosynthetic human insulin precursor by MALDI-TOF mass spectrometry. *Acta Biochimica Polonica* **61**, 349–357.
- Bille A, Jónsson SÆ, Akke M and Irbäck A (2013) Local unfolding and aggregation mechanisms of SOD1: a monte carlo exploration. *Journal of Physical Chemistry B* **117**, 9194–9202.
- Bille A, Jensen KS, Mohanty S, Akke M and Irbäck A (2019) Stability and local unfolding of SOD1 in the presence of protein crowders. *The Journal of Physical Chemistry. B* **123**, 1920–1930.
- Bleier L and Dröse S (2013) Superoxide generation by complex III: from mechanistic rationales to functional consequences. *Biochimica et Biophysica Acta* **1827**, 1320–1331.
- Borchelt DR, Lee MK, Slunt HS, Guarnieri M, Xu ZS, Wong PC, Brown RH, Price DL, Sisodia SS and Cleveland DW (1994) Superoxide dismutase 1 with mutations linked to familial amyotrophic lateral sclerosis possesses significant activity. *Proceedings of the National Academy of Sciences of the USA* **91**, 8292–8296.
- Borchelt DR, Guarnieri M, Wong PC, Lee MK, Slunt HS, Xu ZS, Sisodia SS, Price DL and Cleveland DW (1995) Superoxide dismutase 1 subunits with mutations linked to familial amyotrophic lateral sclerosis do not affect wild-type subunit function. *Journal of Biological Chemistry* **270**, 3234–3238.
- Bosco DA, Morfini G, Karabacak NM, Song Y, Gros-Louis F, Pasinelli P, Goolsby H, Fontaine BA, Lemay N, McKenna-Yasek D, Frosch MP, Agar JN, Julien JP, Brady ST and Brown RH Jr (2010) Wild-type and mutant SOD1 share an aberrant conformation and a common pathogenic pathway in ALS. *Nature Neuroscience* **13**, 1396–1403.
- Boukaftane Y, Khoris J, Moulard B, Salachas F, Meininger V, Malafosse A, Camu W and Rouleau GA (1998) Identification of six novel SOD1 gene mutations in familial amyotrophic lateral sclerosis. *Canadian Journal of Neurological Sciences* **25**, 192–196.
- Bourassa MW, Brown HH, Borchelt DR, Vogt S and Miller LM (2014) Metal-deficient aggregates and diminished copper found in cells expressing SOD1 mutations that cause ALS. *Frontiers in Aging Neuroscience* **6**, 110.
- Bourne Y, Redford SM, Steinman HM, Lepock JR, Tainer JA and Getzoff ED (1996) Novel dimeric interface and electrostatic recognition in bacterial Cu,Zn superoxide dismutase. *Proceedings of the National Academy of Sciences* **93**, 12774–12779.
- Bowling AC, Schulz JB, Brown RH and Beal MF (1993) Superoxide dismutase activity, oxidative damage, and mitochondrial energy metabolism in familial and sporadic amyotrophic lateral sclerosis. *Journal of Neurochemistry* **61**, 2322–2325.
- Bowling AC, Schulz JB, Brown RH and Beal MF (2006) Superoxide dismutase activity, oxidative damage, and mitochondrial energy metabolism in familial and sporadic amyotrophic lateral sclerosis. *Journal of Neurochemistry* **61**, 2322–2325.
- Boyd SD, Liu L, Bulla L and Winkler DD (2018) Quantifying the interaction between copper-zinc superoxide dismutase (Sod1) and its copper chaperone (Ccs1). *Journal of Proteomics & Bioinformatics* **11**, 1–5.
- Brewer GJ (1967) Achromatic regions of tetrazolium stained starch gels: inherited electrophoretic variation. *American Journal of Human Genetics* **19**, 674–680.
- Broom HR, Rumpfält JAO, Vassall KA and Meiering EM (2015a) Destabilization of the dimer interface is a common consequence of diverse ALS-associated mutations in metal free SOD1. *Protein Science* **24**, 2081–2089.
- Broom HR, Vassall KA, Rumpfält JAO, Doyle CM, Tong MS, Bonner JM and Meiering EM (2015b) Combined isothermal titration and differential scanning calorimetry define 3-state thermodynamics of fALS-associated mutant Apo SOD1 dimers and increased population of folded monomer. *Biochemistry* **55**, 519–533.
- Brotherton TE, Li Y, Cooper D, Gearing M, Julien J-P, Rothstein JD, Boylan K and Glass JD (2012) Localization of a toxic form of superoxide dismutase 1 protein to pathologically affected tissues in familial ALS. *Proceedings of the National Academy of Sciences of the USA* **109**, 5505–5510.
- Brotherton TE, Li Y and Glass JD (2013) Cellular toxicity of mutant SOD1 protein is linked to an easily soluble, non-aggregated form *in vitro*. *Neurobiology of Disease* **49**, 49–56.

- Brown NM, Torres AS, Doan PE and O'Halloran TV (2004) Oxygen and the copper chaperone CCS regulate posttranslational activation of Cu,Zn superoxide dismutase. *Proceedings of the National Academy of Sciences of the USA* **101**, 5518–5523.
- Bruijn LI, Houseweert MK, Kato S, Anderson KL, Anderson SD, Ohama E, Reaume AG, Scott RW and Cleveland DW (1998) Aggregation and motor neuron toxicity of an ALS-linked SOD1 mutant independent from wild-type SOD1. *Science* **281**, 1851–1854.
- Bruijn LI, Becher MW, Lee MK, Anderson KL, Jenkins NA, Copeland NG, Sisodia SS, Rothstein JD, Borchelt DR, Price DL and Cleveland DW (1997) ALS-linked SOD1 mutant G85R mediates damage to astrocytes and promotes rapidly progressive disease with SOD1-containing inclusions. *Neuron* **18**, 327–338.
- Bruns CK and Kopito RR (2007) Impaired post-translational folding of familial ALS-linked Cu, Zn superoxide dismutase mutants. *EMBO Journal* **26**, 855–866.
- Bunina TL (1962) On intracellular inclusions in familial amyotrophic lateral sclerosis. *S.S. Korsakov Neuropathology and Psychiatry Magazine* **3**, 1292–1298.
- Byström R, Andersen PM, Gröbner G and Oliveberg M (2010) SOD1 mutations targeting surface hydrogen bonds promote amyotrophic lateral sclerosis without reducing Apo-state stability. *Journal of Biological Chemistry* **285**, 19544–19552.
- Calabrese L, Federici G, Bannister WH, Bannister JV, Rotilio G and Finazzi-Agrò A (1975) Labile sulfur in human superoxide dismutase. *European Journal of Biochemistry* **56**, 305–309.
- Canosa A, Calvo A, Moglia C, Barberis M, Brunetti M, Cammarosano S, Manera U, Ilardi A, Restagno G and Chiò A (2015) A novel p.E121 G heterozygous missense mutation of SOD1 in an apparently sporadic ALS case with a 14-year course. *Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration* **16**, 127–128.
- Cao X, Antonyuk SV, Seetharaman SV, Whitson LJ, Taylor AB, Holloway SP, Strange RW, Doucette PA, Valentine JS, Tiwari A, Hayward LJ, Padua S, Cohlberg JA, Hasnain SS and Hart PJ (2008) Structures of the G85R variant of SOD1 in familial amyotrophic lateral sclerosis. *Journal of Biological Chemistry* **283**, 16169–16177.
- Capper MJ, Wright GSA, Barbieri L, Luchinat E, Mercatelli E, McAlary L, Yerbury JJ, O'Neill PM, Antonyuk SV, Banci L and Hasnain SS (2018) The cysteine-reactive small molecule ebselen facilitates effective SOD1 maturation. *Nature Communications* **9**, 1693.
- Cardoso RME, Thayer MM, DiDonato M, Lo TP, Bruns CK, Getzoff ED and Tainer JA (2002) Insights into Lou Gehrig's disease from the structure and instability of the A4 V mutant of human Cu,Zn superoxide dismutase. *Journal of Molecular Biology* **324**, 247–256.
- Carrico RJ and Deutsch HF (1969) Isolation of human hepatocypreïn and cerebrocypreïn their identity with erythrocyte. *Journal of Biological Chemistry* **244**, 6087–6093.
- Carrico RJ and Deutsch HF (1970) The presence of zinc in human cytochrome and some properties of the apoprotein. *Journal of Biological Chemistry* **245**, 723–727.
- Carroll MC, Girouard JB, Ulloa JL, Subramaniam JR, Wong PC, Valentine JS and Culotta VC (2004) Mechanisms for activating Cu- and Zn-containing superoxide dismutase in the absence of the CCS Cu chaperone. *Proceedings of the National Academy of Sciences of the USA* **101**, 5964–5969.
- Casareno RL, Waggoner D and Gitlin JD (1998) The copper chaperone CCS directly interacts with copper/zinc superoxide dismutase. *Journal of Biological Chemistry* **273**, 23625–23628.
- Case AJ (2017) On the origin of superoxide dismutase: an evolutionary perspective of superoxide-mediated redox signaling. *Antioxidants* **6**, 82.
- Chattopadhyay M, Durazo A, Sohn SH, Strong CD, Gralla EB, Whitelegge JP and Valentine JS (2008) Initiation and elongation in fibrillation of ALS-linked superoxide dismutase. *Proceedings of the National Academy of Sciences of the USA* **105**, 18663–18668.
- Chattopadhyay M, Nwadiibia E, Strong CA, Gralla EB, Valentine JS and Whitelegge JP (2015) The disulfide bond, but not zinc or dimerization, controls initiation and seeded growth in amyotrophic lateral sclerosis-linked Cu-Zn superoxide dismutase (SOD1) fibrillation. *Journal of Biological Chemistry* **290**, 30624–30636.
- Chen X, Shang H, Qiu X, Fujiwara N, Cui L, Li X-M, Gao T-M and Kong J (2012) Oxidative modification of cysteine 111 promotes disulfide bond-independent aggregation of SOD1. *Neurochemical Research* **37**, 835–845.
- Chhanganani D, Endo F, Amanullah A, Upadhyay A, Watanabe S, Mishra R, Yamanaka K and Mishra A (2016) Mahogunin ring finger 1 confers cytoprotection against mutant SOD1 aggregates and is defective in an ALS mouse model. *Neurobiology of Disease* **86**, 16–28.
- Chia R, Tattum MH, Jones S, Collinge J, Fisher EMC and Jackson GS (2010) Superoxide dismutase 1 and tgSOD1 mouse spinal cord seed fibrils, suggesting a propagative cell death mechanism in amyotrophic lateral sclerosis. *PLoS ONE* **5**, e10627.
- Chng C-P and Strange RW (2014) Lipid-associated aggregate formation of superoxide dismutase-1 is initiated by membrane-targeting loops. *Proteins Structure Function and Bioinformatics* **82**, 3194–3209.
- Choi J-S, Cho S, Park SG, Park BC and Lee DH (2004) Co-chaperone CHIP associates with mutant Cu/Zn-superoxide dismutase proteins linked to familial amyotrophic lateral sclerosis and promotes their degradation by proteasomes. *Biochemical and Biophysical Research Communications* **321**, 574–583.
- Choi JS, Kim K, Lee DH, Cho S, Ha JD, Park BC, Kim S, Park SG and Kim J-H (2016) cIAPs promote the proteasomal degradation of mutant SOD1 linked to familial amyotrophic lateral sclerosis. *Biochemical and Biophysical Research Communications* **480**, 422–428.
- Ciryam P, Lambert-Smith IA, Bean DM, Freer R, Cid F, Tartaglia GG, Saunders DN, Wilson MR, Oliver SG, Morimoto RI, Dobson CM, Vendruscolo M, Favrin G and Yerbury JJ (2017) Spinal motor neuron protein supersaturation patterns are associated with inclusion body formation in ALS. *Proceedings of the National Academy of Sciences* **114**, E3935–E3943.
- Coelho FR, Iqbal A, Linares E, Silva DF, Lima FS, Cuccovia IM and Augusto O (2014) Oxidation of the tryptophan 32 residue of human superoxide dismutase 1 caused by its bicarbonate-dependent peroxidase activity triggers the non-amyloid aggregation of the enzyme. *Journal of Biological Chemistry* **289**, 30690–30701.
- Conforti FL, Barone R, Fermo SL, Giliberto C, Patti F, Gambardella A, Quattrone A and Zappia M (2011) Sporadic motor neuron disease in a familial novel SOD1 mutation: incomplete penetrance or chance association? *Amyotrophic Lateral Sclerosis* **12**, 220–222.
- Cozzolino M, Pesaresi MG, Amori I, Crosio C, Ferri A, Nencini M and Carrì MT (2009) Oligomerization of mutant SOD1 in mitochondria of motoneuronal cells drives mitochondrial damage and cell toxicity. *Antioxidants and Redox Signaling* **11**, 1547–1558.
- Crisp MJ, Mawuenyega KG, Patterson BW, Reddy NC, Chott R, Self WK, Wehl CC, Jockel-Balsarotti J, Varadhachary AS, Bucelli RC, Yarasheski KE, Bateman RJ and Miller TM (2015) *In vivo* kinetic approach reveals slow SOD1 turnover in the CNS. *Journal of Clinical Investigation* **125**, 2772–2780.
- Crosby K, Crown AM, Roberts BL, Brown H, Ayers JI and Borchelt DR (2018) Loss of charge mutations in solvent exposed Lys residues of superoxide dismutase 1 do not induce inclusion formation in cultured cell models. *PLoS ONE* **13**, e0206751.
- Crow JP, Sampson JB, Zhuang Y, Thompson JA and Beckman JS (1997) Decreased zinc affinity of amyotrophic lateral sclerosis-associated superoxide dismutase mutants leads to enhanced catalysis of tyrosine nitration by peroxynitrite. *Journal of Neurochemistry* **69**, 1936–1944.
- Csar XF, Wilson NJ, Strike P, Sparrow L, McMahon KA, Ward AC and Hamilton JA (2001) Copper/zinc superoxide dismutase is phosphorylated and modulated specifically by granulocyte-colony stimulating factor in myeloid cells. *Proteomics* **1**, 435–443.
- Cudkovic ME, McKenna-Yasek D, Sapp PE, Chin W, Geller B, Hayden DL, Schoenfeld DA, Hosler BA, Horvitz HR and Brown RH (1997) Epidemiology of mutations in superoxide dismutase in amyotrophic lateral sclerosis. *Annals of Neurology* **41**, 210–221.
- Culik RM, Sekhar A, Nagesh J, Deol H, Rumpfolt JAO, Meiering EM and Kay LE (2018) Effects of maturation on the conformational free-energy landscape of SOD1. *Proceedings of the National Academy of Sciences* **115**, E2546–E2555.
- Culotta VC, Klomp LW, Strain J, Casareno RL, Krebs B and Gitlin JD (1997) The copper chaperone for superoxide dismutase. *Journal of Biological Chemistry* **272**, 23469–23472.

- Da Cruz S, Bui A, Saberi S, Lee SK, Stauffer J, McAlonis-Downes M, Schulte D, Pizzo DP, Parone PA, Cleveland DW and Ravits J (2017) Misfolded SOD1 is not a primary component of sporadic ALS. *Acta Neuropathologica* **124**, 97–111.
- Danielsson J, Mu X, Lang L, Wang H, Binolfi A, Theillet F-X, Bekei B, Logan DT, Selenko P, Wennerström H and Oliveberg M (2015) Thermodynamics of protein destabilization in live cells. *Proceedings of the National Academy of Sciences* **112**, 12402–12407.
- Dasmeh P and Kepp KP (2017) Superoxide dismutase 1 is positively selected to minimize protein aggregation in great apes. *Cellular and Molecular Life Sciences CMLS* **74**, 3023–3037.
- del Grande A, Luigetti M, Conte A, Mancuso I, Lattante S, Marangi G, Stipa G, Zollino M and Sabatelli M (2011) A novel L67P SOD1 mutation in an Italian ALS patient. *Amyotrophic Lateral Sclerosis* **12**, 150–152.
- Deng HX, Hentati A, Tainer JA, Iqbal Z, Cayabyab A, Hung WY, Getzoff ED, Hu P, Herzfeldt B, Roos RP, Warner C, Deng G, Soriano E, Smyth C, Parge HE, Ahmed A, Roses AD, Hallewell RA, Pericak-Vance MA and Siddique T (1993) Amyotrophic lateral sclerosis and structural defects in Cu,Zn superoxide dismutase. *Science* **261**, 1047–1051.
- Deng H-X, Shi Y, Furukawa Y, Zhai H, Fu R, Liu E, Gorrie GH, Khan MS, Hung W-Y, Bigio EH, Lukas T, Dal Canto MC, O'Halloran TV and Siddique T (2006) Conversion to the amyotrophic lateral sclerosis phenotype is associated with intermolecular linked insoluble aggregates of SOD1 in mitochondria. *Proceedings of the National Academy of Sciences of the USA* **103**, 7142–7147.
- DiDonato M, Craig L, Huff ME, Thayer MM, Cardoso RME, Kassmann CJ, Lo TP, Bruns CK, Powers ET, Kelly JW, Getzoff ED and Tainer JA (2003) ALS mutants of human superoxide dismutase form fibrous aggregates via framework destabilization. *Journal of Molecular Biology* **332**, 601–615.
- Doyle CM, Rumpfheldt JAO, Broom HR, Sekhar A, Kay LE and Meiering EM (2016) Concurrent increases and decreases in local stability and conformational heterogeneity in Cu, Zn superoxide dismutase variants revealed by temperature-dependence of amide chemical shifts. *Biochemistry* **55**, 1346–1361.
- Durham HD, Roy J, Dong L and Figlewicz DA (1997) Aggregation of mutant Cu/Zn superoxide dismutase proteins in a culture model of ALS. *Journal of Neuropathology & Experimental Neurology* **56**, 523–530.
- DuVal MG, Hinge VK, Snyder N, Kanyo R, Bratvold J, Pokrishevsky E, Cashman NR, Blinov N, Kovalenko A and Allison WT (2019) Tryptophan 32 mediates SOD1 toxicity in a *in vivo* motor neuron model of ALS and is a promising target for small molecule therapeutics. *Neurobiology of Disease* **124**, 297–310.
- Eisler B, Rosdahl KG and Theorell H (1936) Untersuchungen über die zustandsform des kupfers im blutserum mit hilfe der kataphorese. *Biochemische Zeitschrift* **286**, 435–438.
- Elam JS, Malek K, Rodríguez JA, Doucette PA, Taylor AB, Hayward LJ, Cabelli DE, Valentine JS and Hart PJ (2003b) An alternative mechanism of bicarbonate-mediated peroxidation by copper-zinc superoxide dismutase: rates enhanced via proposed enzyme-associated peroxycarbonate intermediate. *Journal of Biological Chemistry* **278**, 21032–21039.
- Elam JS, Taylor AB, Strange R, Antonyuk S, Doucette PA, Rodriguez JA, Hasnain SS, Hayward LJ, Valentine JS, Yeates TO and Hart PJ (2003a) Amyloid-like filaments and water-filled nanotubes formed by SOD1 mutant proteins linked to familial ALS. *Natural Structural Biology* **10**, 461–467.
- Enayat ZE, Orrell RW, Claus A, Ludolph A, Bachus R, Brockmüller J, Ray-Chaudhuri K, Radunovic A, Shaw C and Wilkinson J (1995) Two novel mutations in the gene for copper zinc superoxide dismutase in UK families with amyotrophic lateral sclerosis. *Human Molecular Genetics* **4**, 1239–1240.
- Endo T, Fujii T, Sato K, Taniguchi N and Fujii J (2000) A pivotal role of Zn-binding residues in the function of the copper chaperone for SOD1. *Biochemical and Biophysical Research Communications* **276**, 999–1004.
- Ermilova IP, Ermilov VB, Levy M, Ho E, Pereira C and Beckman JS (2005) Protection by dietary zinc in ALS mutant G93A SOD transgenic mice. *Neuroscience Letters* **379**, 42–46.
- Ezzi SA, Urushitani M and Julien J-P (2007) Wild-type superoxide dismutase acquires binding and toxic properties of ALS-linked mutant forms through oxidation. *Journal of Neurochemistry* **102**, 170–178.
- Farrarwell NE, Lambert-Smith I, Mitchell K, McKenna J, McAlary L, Ciryam P, Vine KL, Saunders DN and Yerbury JJ (2018) SOD1^{A4V} aggregation alters ubiquitin homeostasis in a cell model of ALS. *Journal of Cell Science* **131**, jcs209122.
- Farrarwell NE, Yerbury MR, Plotkin SS, McAlary L and Yerbury JJ (2019) CuATSM protects against the *in vitro* cytotoxicity of wild type-like SOD1 mutants but not mutants that disrupt metal binding. *ACS Chemical Neuroscience* **10**, 1555–1564.
- Fay JM, Zhu C, Proctor EA, Tao Y, Cui W, Ke H and Dokholyan NV (2016) A phosphomimetic mutation stabilizes SOD1 and rescues cell viability in the context of an ALS-associated mutation. *Structure* **24**, 1898–1906.
- Felbecker A, Camu W, Valdmanis PN, Sperfeld A-D, Waibel S, Steinbach P, Rouleau GA, Ludolph AC and Andersen PM (2010) Four familial ALS pedigrees discordant for two SOD1 mutations: are all SOD1 mutations pathogenic? *Journal of Neurology Neurosurgery & Psychiatry* **81**, 572–577.
- Forman HJ and Fridovich I (1973) On the stability of bovine superoxide dismutase. The effects of metals. *Journal of Biological Chemistry* **248**, 2645–2649.
- Forsberg K, Jonsson PA, Andersen PM, Bergemalm D, Graffmo KS, Hultdin M, Jacobsson J, Rosquist R, Marklund SL and Brännström T (2010) Novel antibodies reveal inclusions containing non-native SOD1 in sporadic ALS patients. *PLoS ONE* **5**, e11552.
- Forsberg K, Graffmo K, Pakkenberg B, Weber M, Nielsen M, Marklund S, Brännström T and Andersen PM (2019) Misfolded SOD1 inclusions in patients with mutations in C9orf72 and other ALS/FTD-associated genes. *Journal of Neurology, Neurosurgery, and Psychiatry* **90**, 861–869.
- Fransen M, Nordgren M, Wang B and Apanasets O (2012) Role of peroxisomes in ROS/RNS-metabolism: implications for human disease. *Biochimica et Biophysica Acta (BBA) – Molecular Basis of Disease* **1822**, 1363–1373.
- Fridovich I (1975) Superoxide dismutases. *Annual Review of Biochemistry* **44**, 147–159.
- Fridovich I and Handler P (1958) Xanthine oxidase: III. Sulfite oxidation as an ultra sensitive assay. *Journal of Biological Chemistry* **233**, 1578.
- Fridovich I and Handler P (1961) Detection of free radicals generated during enzymic oxidations by the initiation of sulfite oxidation. *Journal of Biological Chemistry* **236**, 1836–1840.
- Fujisawa T, Yamaguchi N, Kadowaki H, Tsukamoto Y, Tsuburaya N, Tsubota A, Takahashi H, Naguro I, Takahashi Y, Goto J, Tsuji S, Nishitoh H, Homma K and Ichijo H (2015) A systematic immunoprecipitation approach reinforces the concept of common conformational alterations in amyotrophic lateral sclerosis-linked SOD1 mutants. *Neurobiology of Disease* **82**, 478–486.
- Fujiwara N, Nakano M, Kato S, Yoshihara D, Ookawara T, Eguchi H, Taniguchi N and Suzuki K (2007) Oxidative modification to cysteine sulfonic acid of Cys111 in human copper-zinc superoxide dismutase. *Journal of Biological Chemistry* **282**, 35933–35944.
- Furukawa Y and O'Halloran TV (2005) Amyotrophic lateral sclerosis mutations have the greatest destabilizing effect on the apo- and reduced form of SOD1, leading to unfolding and oxidative aggregation. *Journal of Biological Chemistry* **280**, 17266–17274.
- Furukawa Y, Torres AS and O'Halloran TV (2004) Oxygen-induced maturation of SOD1: a key role for disulfide formation by the copper chaperone CCS. *EMBO Journal* **23**, 2872–2881.
- Furukawa Y, Fu R, Deng H-X, Siddique T and O'Halloran TV (2006) Disulfide cross-linked protein represents a significant fraction of ALS-associated Cu, Zn-superoxide dismutase aggregates in spinal cords of model mice. *Proceedings of the National Academy of Sciences of the USA* **103**, 7148–7153.
- Furukawa Y, Kaneko K, Yamanaka K, O'Halloran TV and Nukina N (2008) Complete loss of post-translational modifications triggers fibrillar aggregation of SOD1 in the familial form of amyotrophic lateral sclerosis. *Journal of Biological Chemistry* **283**, 24167–24176.
- Furukawa Y, Kaneko K, Yamanaka K and Nukina N (2010) Mutation-dependent polymorphism of Cu,Zn-superoxide dismutase aggregates in the familial form of amyotrophic lateral sclerosis. *Journal of Biological Chemistry* **285**, 22221–22231.
- Galeledeen A, Strange RW, Whitson LJ, Antonyuk SV, Narayana N, Taylor AB, Schuermann JP, Holloway SP, Hasnain SS and Hart PJ (2009) Structural and biophysical properties of metal-free pathogenic

- SOD1 mutants A4 V and G93A. *Archives of Biochemistry and Biophysics* **492**, 40–47.
- Georgouloupoulou E, Gellera C, Bragato C, Sola P, Chiari A, Bernabei C and Mandrioli J (2010) A novel SOD1 mutation in a young amyotrophic lateral sclerosis patient with a very slowly progressive clinical course. *Muscle & Nerve* **42**, 596–597.
- Gerschman R, Gilbert DL, Nye SW, Dwyer P and Fenn WO (1954) Oxygen poisoning and X-irradiation: a mechanism in common. *Science* **119**, 623–626.
- Getzoff ED, Tainer JA, Weiner PK, Kollman PA, Richardson JS and Richardson DC (1983) Electrostatic recognition between superoxide and copper, zinc superoxide dismutase. *Nature* **306**, 287–290.
- Ghadge GD, Pavlovic J, Parthasarathy SK, Kay BK and Roos RP (2013) Single chain variable fragment antibodies block aggregation and toxicity induced by familial ALS-linked mutant forms of SOD1. *Neurobiology of Disease* **56**, 74–78.
- Ghadge GD, Kay BK, Drigotas C and Roos RP (2018) Single chain variable fragment antibodies directed against SOD1 ameliorate disease in mutant SOD1 transgenic mice. *Neurobiology of Disease* **121**, 131–137.
- Gill C, Phelan JP, Hatzipetros T, Kidd JD, Tassinari VR, Levine B, Wang MZ, Moreno A, Thompson K, Maier M, Grimm J, Gill A and Vieira FG (2019) SOD1-positive aggregate accumulation in the CNS predicts slower disease progression and increased longevity in a mutant SOD1 mouse model of ALS. *Scientific Reports* **9**, 6724.
- Gleason JE, Galaldeen A, Peterson RL, Taylor AB, Holloway SP, Waninger-Saroni J, Cormack BP, Cabelli DE, Hart PJ and Culotta VC (2014) Candida albicans SOD5 represents the prototype of an unprecedented class of Cu-only superoxide dismutases required for pathogen defense. *Proceedings of the National Academy of Sciences of the USA* **111**, 5866–5871.
- Gnutt D, Timr S, Ahlers J, König B, Manderfeld E, Heyden M, Sterpone F and Ebbinghaus S (2019) Stability effect of quinary interactions reversed by single point mutations. *Journal of the American Chemical Society* **141**, 4660–4669.
- Go Y-M, Chandler JD and Jones DP (2015) The cysteine proteome. *Free Radical Biology & Medicine* **84**, 227–245.
- Gomes C, Keller S, Altevogt P and Costa J (2007) Evidence for secretion of Cu,Zn superoxide dismutase via exosomes from a cell model of amyotrophic lateral sclerosis. *Neuroscience Letters* **428**, 43–46.
- Grad LI, Guest WC, Yanai A, Pokrishevsky E, O'Neill MA, Gibbs E, Semchenko V, Yousefi M, Wishart DS, Plotkin SS and Cashman NR (2011) Intermolecular transmission of superoxide dismutase 1 misfolding in living cells. *Proceedings of the National Academy of Sciences of the USA* **108**, 16398–16403.
- Grad LI, Yerbury JJ, Turner BJ, Guest WC, Pokrishevsky E, O'Neill MA, Yanai A, Silverman JM, Zeineddine R, Corcoran L, Kumita JR, Luheshi LM, Yousefi M, Coleman BM, Hill AF, Plotkin SS, Mackenzie IR and Cashman NR (2014) Intercellular propagated misfolding of wild-type Cu/Zn superoxide dismutase occurs via exosome-dependent and -independent mechanisms. *Proceedings of the National Academy of Sciences of the USA* **111**, 3620–3625.
- Graffmo KS, Forsberg K, Bergh J, Birve A, Zetterström P, Andersen PM, Marklund SL and Brännström T (2013) Expression of wild-type human superoxide dismutase-1 in mice causes amyotrophic lateral sclerosis. *Human Molecular Genetics* **22**, 51–60.
- Gros-Louis F, Soucy G, Larivière R and Julien J-P (2010) Intracerebroventricular infusion of monoclonal antibody or its derived Fab fragment against misfolded forms of SOD1 mutant delays mortality in a mouse model of ALS. *Journal of Neurochemistry* **113**, 1188–1199.
- Gurney ME, Pu H, Chiu AY, Dal Canto MC, Polchow CY, Alexander DD, Caliendo J, Hentati A, Kwon YW and Deng HX (1994) Motor neuron degeneration in mice that express a human Cu,Zn superoxide dismutase mutation. *Science* **264**, 1772–1775.
- Hallewell RA, Imlay KC, Lee P, Fong NM, Gallegos C, Getzoff ED, Tainer JA, Cabelli DE, Tekamp-Olson P, Mullenbach GT and Cousins LS (1991) Thermostabilization of recombinant human and bovine CuZn superoxide dismutases by replacement of free cysteines. *Biochemical and Biophysical Research Communications* **181**, 474–480.
- Han D, Williams E and Cadenas E (2001) Mitochondrial respiratory chain-dependent generation of superoxide anion and its release into the intermembrane space. *Biochemical Journal* **353**, 411–416.
- Hand CK, Mayeux-Portas V, Khoris J, Briolotti V, Clavelou P, Camu W and Rouleau GA (2001) Compound heterozygous D90A and D96N SOD1 mutations in a recessive amyotrophic lateral sclerosis family. *Annals of Neurology* **49**, 267–271.
- Hart PJ, Liu H, Pellegrini M, Nersissian AM, Gralla EB, Valentine JS and Eisenberg D (1998) Subunit asymmetry in the three-dimensional structure of a human CuZnSOD mutant found in familial amyotrophic lateral sclerosis. *Protein Science Publication Protein Society* **7**, 545–555.
- Hartz JW and Deutsch HF (1972) Subunit structure of human superoxide dismutase. *Journal of Biological Chemistry* **247**, 7043–7050.
- Hayward C, Brock DJ, Minns RA and Swingle RJ (1998) Homozygosity for Asn86Ser mutation in the CuZn-superoxide dismutase gene produces a severe clinical phenotype in a juvenile onset case of familial amyotrophic lateral sclerosis. *Journal of Medical Genetics* **35**, 174.
- Hayward LJ, Rodriguez JA, Kim JW, Tiwari A, Goto JJ, Cabelli DE, Valentine JS and Brown Jr RH (2002). Decreased metallation and activity in subsets of mutant superoxide dismutases associated with familial amyotrophic lateral sclerosis. *Journal of Biological Chemistry* **277**, 15923–15931.
- Hirano A, Kurland LT and Sayre GP (1967) Familial amyotrophic lateral sclerosis. A subgroup characterized by posterior and spinocerebellar tract involvement and hyaline inclusions in the anterior horn cells. *Archives of Neurology* **16**, 232–243.
- Hofrichter J, Ross PD and Eaton WA (1974) Kinetics and mechanism of deoxyhemoglobin S gelation: a new approach to understanding sickle cell disease. *Proceedings of the National Academy of Sciences of the USA* **71**, 4864–4868.
- Horecker BL and Heppel LA (1949) The reduction of cytochrome c by milk xanthine oxidase. *Journal of Biological Chemistry* **178**, 683–690.
- Hörnberg A, Logan DT, Marklund SL and Oliveberg M (2007) The coupling between disulphide status, metallation and dimer interface strength in Cu/Zn superoxide dismutase. *Journal of Molecular Biology* **365**, 333–342.
- Hough MA and Hasnain SS (2003) Structure of fully reduced bovine copper zinc superoxide dismutase at 1.15 Å. *Structure* **11**, 937–946.
- Hough MA, Grossmann JG, Antonyuk SV, Strange RW, Doucette PA, Rodriguez JA, Whitson LJ, Hart PJ, Hayward LJ, Valentine JS and Hasnain SS (2004) Dimer destabilization in superoxide dismutase may result in disease-causing properties: structures of motor neuron disease mutants. *Proceedings of the National Academy of Sciences of the USA* **101**, 5976–5981.
- Hough MA, Strange RW and Hasnain SS (2000) Conformational variability of the Cu site in one subunit of bovine CuZn superoxide dismutase: the importance of mobility in the Glu119-Leu142 loop region for catalytic function. *Journal of Molecular Biology* **304**, 231–241.
- Ihara K, Fujiwara N, Yamaguchi Y, Torigoe H, Wakatsuki S, Taniguchi N and Suzuki K (2012) Structural switching of Cu,Zn-superoxide dismutases at loop VI: insights from the crystal structure of 2-mercaptoethanol-modified enzyme. *Bioscience Reports* **32**, 539–548.
- Ikeda M, Abe K, Aoki M, Ogasawara M, Kameya T, Watanabe M, Shoji M, Hirai S and Itoyama Y (1995) A novel point mutation in the Cu/Zn superoxide dismutase gene in a patient with familial amyotrophic lateral sclerosis. *Human Molecular Genetics* **4**, 491–492.
- Ingre C, Pinto S, Birve A, Press R, Danielsson O, de Carvalho M, Gudmundsson G and Andersen PM (2013) No association between VAPB mutations and familial or sporadic ALS in Sweden, Portugal and Iceland. *Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration* **14**, 620–627.
- Israelson A, Ditsworth D, Sun S, Song S, Liang J, Hruska-Plochan M, McAlonis-Downes M, Abu-Hamad S, Zoltsman G, Shani T, Maldonado M, Bui A, Navarro M, Zhou H, Marsala M, Kaspar BK, Da Cruz S and Cleveland DW (2015) Macrophage migration inhibitory factor as a chaperone inhibiting accumulation of misfolded SOD1. *Neuron* **86**, 218–232.
- Ivanova MI, Sievers SA, Guenther EL, Johnson LM, Winkler DD, Galaldeen A, Sawaya MR, Hart PJ and Eisenberg DS (2014) Aggregation-triggering segments of SOD1 fibril formation support a common pathway for familial and sporadic ALS. *Proceedings of the National Academy of Sciences of the USA* **111**, 197–201.

- Jaarsma D, Rognoni F, van Duijn W, Verspaget HW, Haasdijk ED and Holstege JC (2001) Cuzn superoxide dismutase (SOD1) accumulates in vacuolated mitochondria in transgenic mice expressing amyotrophic lateral sclerosis-linked SOD1 mutations. *Acta Neuropathologica* **102**, 293–305.
- Jabusch JR, Farb DL, Kerschensteiner DA and Deutsch HF (1980) Some sulfhydryl properties and primary structure of human erythrocyte superoxide dismutase. *Biochemistry* **19**, 2310–2316.
- Jackson M, Al-Chalabi A, Enayat ZE, Chioza B, Leigh PN and Morrison KE (1997) Copper/zinc superoxide dismutase 1 and sporadic amyotrophic lateral sclerosis: analysis of 155 cases and identification of a novel insertion mutation. *Annals of Neurology* **42**, 803–807.
- Jacobsson J, Jonsson PA, Andersen PM, Forsgren L and Marklund SL (2001) Superoxide dismutase in CSF from amyotrophic lateral sclerosis patients with and without CuZn-superoxide dismutase mutations. *Journal of Brain and Neurology* **124**, 1461–1466.
- Jaunmuktane Z, Mead S, Ellis M, Wadsworth JDF, Nicoll AJ, Kenny J, Launchbury F, Linehan J, Richard-Loendt A, Walker AS, Rudge P, Collinge J and Brandner S (2015) Evidence for human transmission of amyloid- β pathology and cerebral amyloid angiopathy. *Nature* **525**, 247–250.
- Johnston JA, Dalton MJ, Gurney ME and Kopito RR (2000) Formation of high molecular weight complexes of mutant Cu,Zn-superoxide dismutase in a mouse model for familial amyotrophic lateral sclerosis. *Proceedings of the National Academy of Sciences* **97**, 12571–12576.
- Jones CT, Swingle RJ and Brock DJ (1994a) Identification of a novel SOD1 mutation in an apparently sporadic amyotrophic lateral sclerosis patient and the detection of Ile113Thr in three others. *Human Molecular Genetics* **3**, 649–650.
- Jones CT, Shaw PJ, Chari G and Brock DJH (1994b) Identification of a novel exon 4 SOD1 mutation in a sporadic amyotrophic lateral sclerosis patient. *Molecular and Cellular Probes* **8**, 329–330.
- Jonsson PA, Ernhill K, Andersen PM, Bergemalm D, Brännström T, Gredal O, Nilsson P and Marklund SL (2004) Minute quantities of misfolded mutant superoxide dismutase-1 cause amyotrophic lateral sclerosis. *Journal of Brain and Neurology* **127**, 73–88.
- Jonsson PA, Graffmo KS, Andersen PM, Brännström T, Lindberg M, Oliveberg M and Marklund SL (2006a) Disulphide-reduced superoxide dismutase-1 in CNS of transgenic amyotrophic lateral sclerosis models. *Journal of Brain and Neurology* **129**, 451–464.
- Jonsson PA, Graffmo KS, Brännström T, Nilsson P, Andersen PM and Marklund SL (2006b) Motor neuron disease in mice expressing the wild type-like D90A mutant superoxide dismutase-1. *Journal of Neuro pathology & Experimental Neurology* **65**, 1126–1136.
- Kabuta T, Suzuki Y and Wada K (2006) Degradation of amyotrophic lateral sclerosis-linked mutant Cu,Zn-superoxide dismutase proteins by macroautophagy and the proteasome. *Journal of Biological Chemistry* **281**, 30524–30533.
- Kaganovich D, Kopito R and Frydman J (2008) Misfolded proteins partition between two distinct quality control compartments. *Nature* **454**, 1088.
- Kaliszewski M, Kennedy AK, Blaes SL, Shaffer RS, Knott AB, Song W, Hauser HA, Bossy B, Huang T-T and Bossy-Wetzel E (2016) SOD1 lysine 123 acetylation in the adult central nervous system. *Frontiers in Cellular Neuroscience* **10**, 287.
- Karch CM, Prudencio M, Winkler DD, Hart PJ and Borchelt DR (2009) Role of mutant SOD1 disulfide oxidation and aggregation in the pathogenesis of familial ALS. *Proceedings of the National Academy of Sciences of the USA* **106**, 7774–7779.
- Kato S, Shimoda M, Watanabe Y, Nakashima K, Takahashi K and Ohama E (1996) Familial amyotrophic lateral sclerosis with a two base pair deletion in superoxide dismutase 1: gene multisystem degeneration with intracytoplasmic hyaline inclusions in astrocytes. *Journal of Neuro pathology & Experimental Neurology* **55**, 1089–1101.
- Kato S, Hayashi H, Nakashima K, Nanba E, Kato M, Hirano A, Nakano I, Asayama K and Ohama E (1997) Pathological characterization of astrocytic hyaline inclusions in familial amyotrophic lateral sclerosis. *American Journal of Pathology* **151**, 611–620.
- Kato S, Horiuchi S, Nakashima K, Hirano A, Shibata N, Nakano I, Saito M, Kato M, Asayama K and Ohama E (1999) Astrocytic hyaline inclusions contain advanced glycation endproducts in familial amyotrophic lateral sclerosis with superoxide dismutase 1 gene mutation: immunohistochemical and immunoelectron microscopical analyses. *Acta Neuropathologica* **97**, 260–266.
- Kato M, Aoki M, Ohta M, Nagai M, Ishizaki F, Nakamura S and Itoyama Y (2001a) Marked reduction of the Cu/Zn superoxide dismutase polypeptide in a case of familial amyotrophic lateral sclerosis with the homozygous mutation. *Neuroscience Letters* **312**, 165–168.
- Kato S, Sumi-Akamaru H, Fujimura H, Sakoda S, Kato M, Hirano A, Takikawa M and Ohama E (2001b) Copper chaperone for superoxide dismutase co-aggregates with superoxide dismutase 1 (SOD1) in neuronal Lewy body-like hyaline inclusions: an immunohistochemical study on familial amyotrophic lateral sclerosis with SOD1 gene mutation. *Acta Neuropathologica* **102**, 233–238.
- Kato S, Horiuchi S, Liu J, Cleveland DW, Shibata N, Nakashima K, Nagai R, Hirano A, Takikawa M, Kato M, Nakano I and Ohama E (2000) Advanced glycation endproduct-modified superoxide dismutase-1 (SOD1)-positive inclusions are common to familial amyotrophic lateral sclerosis patients with SOD1 gene mutations and transgenic mice expressing human SOD1 with a G85R mutation. *Acta Neuropathologica* **100**, 490–505.
- Kawamata H and Manfredi G (2008) Different regulation of wild-type and mutant Cu,Zn superoxide dismutase localization in mammalian mitochondria. *Human Molecular Genetics* **17**, 3303–3317.
- Kayatekin C, Zitzewitz JA and Matthews CR (2008) Zinc binding modulates the entire folding free energy surface of human Cu,Zn superoxide dismutase. *Journal of Molecular Biology* **384**, 540–555.
- Kayatekin C, Zitzewitz JA and Matthews CR (2010) Disulfide-reduced ALS variants of Cu, Zn superoxide dismutase exhibit increased populations of unfolded species. *Journal of Molecular Biology* **398**, 320–331.
- Kayatekin C, Cohen NR and Matthews CR (2012) Enthalpic barriers dominate the folding and unfolding of the human Cu,Zn superoxide dismutase monomer. *Journal of Molecular Biology* **424**, 192–202.
- Keele BB, McCord JM and Fridovich I (1970) Superoxide dismutase from *Escherichia coli* B. A new manganese-containing enzyme. *Journal of Biological Chemistry* **245**, 6176–6181.
- Keele BB, McCord JM and Fridovich I (1971) Further characterization of bovine superoxide dismutase and its isolation from bovine heart. *Journal of Biological Chemistry* **246**, 2875–2880.
- Kerman A, Liu H-N, Croul S, Bilbao J, Rogaeva E, Zinman L, Robertson J and Chakrabarty A (2010) Amyotrophic lateral sclerosis is a non-amyloid disease in which extensive misfolding of SOD1 is unique to the familial form. *Acta Neuropathologica* **119**, 335–344.
- Kershaw NM, Wright GSA, Sharma R, Antonyuk SV, Strange RW, Berry NG, O'Neill PM and Hasnain SS (2013) X-ray crystallography and computational docking for the detection and development of protein-ligand interactions. *Current Medicinal Chemistry* **20**, 569–575.
- Keskin I, Forsgren E, Lange DJ, Weber M, Birve A, Synofzik M, Gilthorpe JD, Andersen PM and Marklund SL (2016) Effects of cellular pathway disturbances on misfolded superoxide dismutase-1 in fibroblasts derived from ALS patients. *PLoS ONE* **11**, e0150133.
- Keskin I, Birve A, Berdyski M, Hjertkvist K, Rofougaran R, Nilsson TK, Glass JD, Marklund SL and Andersen PM (2017) Comprehensive analysis to explain reduced or increased SOD1 enzymatic activity in ALS patients and their relatives. *Amyotrophic Lateral Sclerosis and Frontotemporal Degeneration* **0**, 1–7.
- Khare SD, Caplow M and Dokholyan NV (2004) The rate and equilibrium constants for a multistep reaction sequence for the aggregation of superoxide dismutase in amyotrophic lateral sclerosis. *Proceedings of the National Academy of Sciences of the USA* **101**, 15094–15099.
- Kim Y-J, Nakatomi R, Akagi T, Hashikawa T and Takahashi R (2005) Unsaturated fatty acids induce cytotoxic aggregate formation of amyotrophic lateral sclerosis-linked superoxide dismutase 1 mutants. *Journal of Biological Chemistry* **280**, 21515–21521.
- Kim J, Lee H, Lee JH, Kwon DY, Genovesio A, Fenistein D, Ogier A, Brondani V and Grailhe R (2014) Dimerization, oligomerization, and aggregation of human amyotrophic lateral sclerosis Cu/Zn-superoxide dismutase 1 mutant forms in live cells. *Journal of Biological Chemistry* **289**, 15094–15103.

- Klug D, Rabani J and Fridovich I (1972) A direct demonstration of the catalytic action of superoxide dismutase through the use of pulse radiolysis. *Journal of Biological Chemistry* **247**, 4839–4842.
- Kobayashi J, Kuroda M, Kawata A, Mochizuki Y, Mizutani T, Komori T, Ikeuchi T and Koide R (2012) Novel G37 V mutation of SOD1 gene in autopsied patient with familial amyotrophic lateral sclerosis. *Amyotrophic Lateral Sclerosis* **13**, 570–572.
- Koide T, Igarashi S, Kikugawa K, Nakano R, Inuzuka T, Yamada M, Takahashi H and Tsuji S (1998) Formation of granular cytoplasmic aggregates in COS7 cells expressing mutant Cu/Zn superoxide dismutase associated with familial amyotrophic lateral sclerosis. *Neuroscience Letters* **257**, 29–32.
- Kong J and Xu Z (1998) Massive mitochondrial degeneration in motor neurons triggers the onset of amyotrophic lateral sclerosis in mice expressing a mutant SOD1. *Journal of Neuroscience* **18**, 3241–3250.
- Kostrzewa M, Burck-Lehmann U and Müller U (1994) Autosomal dominant amyotrophic lateral sclerosis: a novel mutation in the Cu/Zn superoxide dismutase-1 gene. *Human Molecular Genetics* **3**, 2261–2262.
- Kurland LT and Mulder DW (1955) Epidemiologic investigations of amyotrophic lateral sclerosis. 2. Familial aggregations indicative of dominant inheritance. I. *Neurology* **5**, 182–196.
- Lamb AL, Wernimont AK, Pufahl RA, Culotta VC, O'Halloran TV and Rosenzweig AC (1999) Crystal structure of the copper chaperone for superoxide dismutase. *Natural Structural Biology* **6**, 724–729.
- Lang L, Zetterström P, Brännström T, Marklund SL, Danielsson J and Oliveberg M (2015) SOD1 aggregation in ALS mice shows simplistic test tube behavior. *Proceedings of the National Academy of Sciences of the USA* **112**, 9878–9883.
- Leal SS, Cardoso I, Valentine JS and Gomes CM (2013) Calcium ions promote superoxide dismutase 1 (SOD1) aggregation into non-fibrillar amyloid. *Journal of Biological Chemistry* **288**, 25219–25228.
- Lee J-Y, Kawaguchi Y, Li M, Kapur M, Choi SJ, Kim H-J, Park S-Y, Zhu H and Yao T-P (2015) Uncoupling of protein aggregation and neurodegeneration in a mouse amyotrophic lateral sclerosis model. *Neurodegenerative Diseases* **15**, 339–349.
- Leinartaite L, Saraboji K, Nordlund A, Logan DT and Oliveberg M (2010) Folding catalysis by transient coordination of Zn²⁺ to the Cu ligands of the ALS-associated enzyme Cu/Zn superoxide dismutase 1. *Journal of the American Chemical Society* **132**, 13495–13504.
- Lek M, Karczewski KJ, Minikel EV, Samocha KE, Banks E, Fennell T, O'Donnell-Luria AH, Ware JS, Hill AJ, Cummings BB, Tukiainen T, Birnbaum DP, Kosmicki JA, Duncan LE, Estrada K, Zhao F, Zou J, Pierce-Hoffman E, Berghout J, Cooper DN, DeFlaux N, DePristo M, Do R, Flannick J, Fromer M, Gauthier L, Goldstein J, Gupta N, Howrigan D, Kiezun A, Kurki MI, Moonshine AL, Natarajan P, Orozco L, Peloso GM, Poplin R, Rivas MA, Ruano-Rubio V, Rose SA, Ruderfer DM, Shakir K, Stenson PD, Stevens C, Thomas BP, Tiao G, Tusie-Luna MT, Weisburd B, Won HH, Yu D, Altshuler DM, Ardissino D, Boehnke M, Danesh J, Donnelly S, Elosua R, Florez JC, Gabriel SB, Getz G, Glatt SJ, Hultman CM, Kathiresan S, Laakso M, McCarroll S, McCarthy MI, McGovern D, McPherson R, Neale BM, Palotie A, Purcell SM, Saleheen D, Scharf JM, Sklar P, Sullivan PF, Tuomilehto J, Tuang MT, Watkins HC, Wilson JG, Daly MJ, MacArthur DG and Exome Aggregation Consortium (2016) Analysis of protein-coding genetic variation in 60,706 humans. *Nature* **536**, 285–291.
- Lelie HL, Liba A, Bourassa MW, Chattopadhyay M, Chan PK, Gralla EB, Miller LM, Borchelt DR, Valentine JS and Whitelegge JP (2011) Copper and zinc metallation status of copper-zinc superoxide dismutase from amyotrophic lateral sclerosis transgenic mice. *Journal of Biological Chemistry* **286**, 2795–2806.
- Lepock JR, Arnold LD, Petkau A and Kelly K (1981) Interaction of superoxide dismutase with phospholipid liposomes. An uptake, spin label and calorimetric study. *Biochimica et Biophysica Acta* **649**, 45–57.
- Lepock JR, Frey HE and Hallewell RA (1990) Contribution of conformational stability and reversibility of unfolding to the increased thermostability of human and bovine superoxide dismutase mutated at free cysteines. *Journal of Biological Chemistry* **265**, 21612–21618.
- Li J, Liu D, Sun L, Lu Y and Zhang Z (2012) Advanced glycation end products and neurodegenerative diseases: mechanisms and perspective. *Journal of the Neurological Sciences* **317**, 1–5.
- Lim L, Lee X and Song J (2015) Mechanism for transforming cytosolic SOD1 into integral membrane proteins of organelles by ALS-causing mutations. *Biochimica et Biophysica Acta (BBA) – Biomembranes* **1848**, 1–7.
- Lin C, Zeng H, Lu J, Xie Z, Sun W, Luo C, Ding J, Yuan S, Geng M and Huang M (2015) Acetylation at lysine 71 inactivates superoxide dismutase 1 and sensitizes cancer cells to genotoxic agents. *Oncotarget* **6**, 20578–20591.
- Lindberg MJ, Tibell L and Oliveberg M (2002) Common denominator of Cu/Zn superoxide dismutase mutants associated with amyotrophic lateral sclerosis: decreased stability of the apo state. *Proceedings of the National Academy of Sciences of the USA* **99**, 16607–16612.
- Lindberg MJ, Normark J, Holmgren A and Oliveberg M (2004) Folding of human superoxide dismutase: disulfide reduction prevents dimerization and produces marginally stable monomers. *Proceedings of the National Academy of Sciences* **101**, 15893–15898.
- Lindberg MJ, Byström R, Boknäs N, Andersen PM and Oliveberg M (2005) Systematically perturbed folding patterns of amyotrophic lateral sclerosis (ALS)-associated SOD1 mutants. *Proceedings of the National Academy of Sciences of the USA* **102**, 9754–9759.
- Liu H-N, Tjostheim S, Dasilva K, Taylor D, Zhao B, Rakhit R, Brown M, Chakraborty A, McLaurin J and Robertson J (2012) Targeting of monomer/misfolded SOD1 as a therapeutic strategy for amyotrophic lateral sclerosis. *The Journal of Neuroscience* **32**, 8791–8799.
- Locke A, Main ER and Rosbash DO (1932) The copper and non-hemoglobinous iron contents of the blood serum in disease. *Journal of Clinical Investigation* **11**, 527.
- Lopate G, Baloh RH, Al-Lozi MT, Miller TM, Fernandes Filho JA, Ni O, Leston A, Florence J, Schierbecker J and Allred P (2010) Familial ALS with extreme phenotypic variability due to the I113 T SOD1 mutation. *Amyotrophic Lateral Sclerosis* **11**, 232–236.
- Luchinat E, Barbieri L, Rubino JT, Kozyreva T, Cantini F and Banci L (2014) In-cell NMR reveals potential precursor of toxic species from SOD1 fALS mutants. *Nature Communications* **5**, 1–10.
- Luchinat E, Barbieri L and Banci L (2017) A molecular chaperone activity of CCS restores the maturation of SOD1 fALS mutants. *Scientific Reports* **7**, 17433.
- Lyons TJ, Liu H, Goto JJ, Nersissian A, Roe JA, Graden JA, Café C, Ellerby LM, Bredesen DE, Gralla EB and Valentine JS (1996) Mutations in copper-zinc superoxide dismutase that cause amyotrophic lateral sclerosis alter the zinc binding site and the redox behavior of the protein. *Proceedings of the National Academy of Sciences of the USA* **93**, 12240–12244.
- Ma Q, Fan J-B, Zhou Z, Zhou B-R, Meng S-R, Hu J-Y, Chen J and Liang Y (2012) The contrasting effect of macromolecular crowding on amyloid fibril formation. *PLoS ONE* **7**, e36288.
- Magrané J, Sahawneh MA, Przedborski S, Estévez ÁG and Manfredi G (2012) Mitochondrial dynamics and bioenergetic dysfunction is associated with synaptic alterations in mutant SOD1 motor neurons. *The Journal of Neuroscience* **32**, 229–242.
- Manjula R, Wright GSA, Strange RW and Padmanabhan B (2018) Assessment of ligand binding at a site relevant to SOD1 oxidation and aggregation. *FEBS Letters* **592**, 1725–1737.
- Mann T and Keilin D (1938a) Haemocuprein, a copper-protein compound of red blood corpuscles. *Nature* **142**, 148.
- Mann T and Keilin D (1938b) Haemocuprein and hepatocuprein, copper-protein compounds of blood and liver in mammals. *Proceedings of the Royal Society of London B Biological Sciences* **126**, 303–315.
- Marklund SL, Andersen PM, Forsgren L, Nilsson P, Ohlsson PI, Wikander G and Oberg A (1997) Normal binding and reactivity of copper in mutant superoxide dismutase isolated from amyotrophic lateral sclerosis patients. *Journal of Neurochemistry* **69**, 675–681.
- Markowitz H, Cartwright GE and Wintrobe MM (1959) Studies on copper metabolism XXVII. The isolation and properties of an erythrocyte cuproprotein (Erythrocuprein). *Journal of Biological Chemistry* **234**, 40–45.

- Marucci G, Morandi L, Bartolomei I, Salvi F, Pession A, Righi A, Lauria G and Foschini MP (2007) Amyotrophic lateral sclerosis with mutation of the Cu/Zn superoxide dismutase gene (SOD1) in a patient with down syndrome. *Neuromuscular Disorders* **17**, 673–676.
- Mateju D, Franzmann TM, Patel A, Kopach A, Boczek EE, Maharana S, Lee HO, Carra S, Hyman AA and Alberti S (2017) An aberrant phase transition of stress granules triggered by misfolded protein and prevented by chaperone function. *EMBO Journal* **36**, 1669–1687.
- Matsumoto G, Stojanovic A, Holmberg CI, Kim S and Morimoto RI (2005) Structural properties and neuronal toxicity of amyotrophic lateral sclerosis-associated Cu/Zn superoxide dismutase 1 aggregates. *Journal of Cell Biology* **171**, 75–85.
- Mayeux V, Corcia P, Besson G, Jafari-Schluep H-F, Briolotti V and Camu W (2003) N19s, a new SOD1 mutation in sporadic amyotrophic lateral sclerosis: no evidence for disease causation. *Annals of Neurology* **53**, 815–818.
- McAlary L, Yerbury JJ and Aquilina JA (2013) Glutathionylation potentiates benign superoxide dismutase 1 variants to the toxic forms associated with amyotrophic lateral sclerosis. *Scientific Reports* **3**, 3275.
- McAlary L, Aquilina JA and Yerbury JJ (2016) Susceptibility of mutant SOD1 to form a destabilized monomer predicts cellular aggregation and toxicity but not *in vitro* aggregation propensity. *Frontiers in Neuroscience* **10**, 499.
- McAllum EJ, Lim NK-H, Hickey JL, Paterson BM, Donnelly PS, Li Q-X, Liddell JR, Barnham KJ, White AR and Crouch PJ (2013) Therapeutic effects of CuII(atm) in the SOD1-G37R mouse model of amyotrophic lateral sclerosis. *Amyotroph Lateral Scler Frontotemporal Degener* **14**, 586–590.
- McClune GJ and Fee JA (1976) Stopped flow spectrophotometric observation of superoxide dismutation in aqueous solution. *FEBS Letters* **67**, 294–298.
- McCord JM and Fridovich I (1968) The reduction of cytochrome c by milk xanthine oxidase. *Journal of Biological Chemistry* **243**, 5753–5760.
- McCord JM and Fridovich I (1969) Superoxide dismutase. An enzymic function for erythrocyte (hemocuprein). *Journal of Biological Chemistry* **244**, 6049–6055.
- McCord JM, Keele BB and Fridovich I (1971) An enzyme-based theory of obligate anaerobiosis: the physiological function of superoxide dismutase. *Proceedings of the National Academy of Sciences of the USA* **68**, 1024–1027.
- Medinas DB, Gozzo FC, Santos LFA, Iglesias AH and Augusto O (2010) A ditryptophan cross-link is responsible for the covalent dimerization of human superoxide dismutase 1 during its bicarbonate-dependent peroxidase activity. *Free Radical Biology & Medicine* **49**, 1046–1053.
- Millecamps S, Salachas F, Cazeneuve C, Gordon P, Bricka B, Camuzat A, Guillot-Noël L, Russaouen O, Bruneteau G, Pradat P-F, Le Forestier N, Vandenberghe N, Danel-Brunaud V, Guy N, Thauvin-Robinet C, Lacomblez L, Couratier P, Hannequin D, Seilhean D, Le Ber I, Corcia P, Camu W, Brice A, Rouleau G, LeGuern E and Meininger V (2010) SOD1, ANG, VAPB, TARDBP, and FUS mutations in familial amyotrophic lateral sclerosis: genotype–phenotype correlations. *Journal of Medical Genetics* **47**, 554–560.
- Mohamed MS and Greenberg DM (1954) Isolation of purified copper protein from horse liver. *The Journal of General Physiology* **37**, 433–439.
- Morita M, Abe K, Takahashi M, Onodera Y, Okumura H, Niino M, Tashiro K, Nakano I and Itoyama Y (1998) A novel mutation Asp90Val in the SOD1 gene associated with Japanese familial ALS. *European Journal of Neurology* **5**, 389–392.
- Mu X, Choi S, Lang L, Mowray D, Dokholyan NV, Danielsson J and Oliveberg M (2017) Physicochemical code for quinary protein interactions in *Escherichia coli*. *Proceedings of the National Academy of Sciences* **114**, E4556–E4563.
- Münch C and Bertolotti A (2010) Exposure of hydrophobic surfaces initiates aggregation of diverse ALS-causing superoxide dismutase-1 mutants. *Journal of Molecular Biology* **399**, 512–525.
- Münch C, O'Brien J and Bertolotti A (2011) Prion-like propagation of mutant superoxide dismutase-1 misfolding in neuronal cells. *Proceedings of the National Academy of Sciences of the USA* **108**, 3548–3553.
- Nakamura M, Bieniek KF, Lin W-L, Graff-Radford NR, Murray ME, Castanedes-Casey M, Desaro P, Baker MC, Rutherford NJ, Robertson J, Rademakers R, Dickson DW and Boylan KB (2015) A truncating SOD1 mutation, p.Gly141X, is associated with clinical and pathologic heterogeneity, including frontotemporal lobar degeneration. *Acta Neuropathologica* **130**, 145–157.
- Nakano R, Inuzuka T, Kikugawa K, Takahashi H, Sakimura K, Fujii J, Taniguchi N and Tsuji S (1996) Instability of mutant Cu/Zn superoxide dismutase (Ala4Thr) associated with familial amyotrophic lateral sclerosis. *Neuroscience Letters* **211**, 129–131.
- Naruse H, Iwata A, Takahashi Y, Ichihara K, Kamei S, Yamatoku M, Hirayama T, Suzuki N, Aoki M, Miyagawa T, Shimizu J, Tsuji S and Goto J (2013) Familial amyotrophic lateral sclerosis with novel A4D SOD1 mutation with late age at onset and rapid progressive course. *Neurology and Clinical Neuroscience* **1**, 45–47.
- Nedd S, Redler RL, Proctor EA, Dokholyan NV and Alexandrova AN (2014) Cu,Zn-superoxide dismutase without Zn is folded but catalytically inactive. *Journal of Molecular Biology* **426**, 4112–4124.
- Neuman EW (1934) Potassium superoxide and the three-electron bond. *Journal of Chemical Physics* **2**, 31–33.
- Nishida CR, Gralla EB and Valentine JS (1994) Characterization of three yeast copper-zinc superoxide dismutase mutants analogous to those coded for in familial amyotrophic lateral sclerosis. *Proceedings of the National Academy of Sciences of the USA* **91**, 9906–9910.
- Nishiyama K, Murayama S, Shimizu J, Ohya Y, Kwak S, Asayama K and Kanazawa I (1995) Cu/Zn superoxide dismutase-like immunoreactivity is present in Lewy bodies from Parkinson disease: a light and electron microscopic immunocytochemical study. *Acta Neuropathologica* **89**, 471–474.
- Nordlund A and Oliveberg M (2006) Folding of Cu/Zn superoxide dismutase suggests structural hotspots for gain of neurotoxic function in ALS: parallels to precursors in amyloid disease. *Proceedings of the National Academy of Sciences* **103**, 10218–10223.
- Nordlund A, Leinartaitė L, Saraboji K, Aisenbrey C, Gröbner G, Zetterström P, Danielsson J, Logan DT and Oliveberg M (2009) Functional features cause misfolding of the ALS-provoking enzyme SOD1. *Proceedings of the National Academy of Sciences* **106**, 9667–9672.
- Oladzad Abbasabadi A, Javanian A, Nikkhal M, Meratan AA, Ghiasi P and Nemat-Gorgani M (2013) Disruption of mitochondrial membrane integrity induced by amyloid aggregates arising from variants of SOD1. *International Journal of Biological Macromolecules* **61**, 212–217.
- Orrell RW, Habgood JJ, Malaspina A, Mitchell J, Greenwood J, Lane RJ and deBellerocche JS (1999) Clinical characteristics of SOD1 gene mutations in UK families with ALS. *Journal of the Neurological Sciences* **169**, 56–60.
- Oztug Durer ZA, Cohlberg JA, Dinh P, Padua S, Ehrenclo K, Downes S, Tan JK, Nakano Y, Bowman CJ, Hoskins JL, Kwon C, Mason AZ, Rodriguez JA, Doucette PA, Shaw BF and Valentine JS (2009) Loss of metal ions, disulfide reduction and mutations related to familial ALS promote formation of amyloid-like aggregates from superoxide dismutase. *PLoS ONE* **4**, e5004.
- Pardo CA, Xu Z, Borchelt DR, Price DL, Sisodia SS and Cleveland DW (1995) Superoxide dismutase is an abundant component in cell bodies, dendrites, and axons of motor neurons and in a subset of other neurons. *Proceedings of the National Academy of Sciences of the USA* **92**, 954–958.
- Paré B, Lehmann M, Beaudin M, Nordström U, Saikali S, Julien J-P, Giltthorpe JD, Marklund SL, Cashman NR, Andersen PM, Forsberg K, Dupré N, Gould P, Brännström T and Gros-Louis F (2018) Misfolded SOD1 pathology in sporadic amyotrophic lateral sclerosis. *Scientific Reports* **8**, 14223.
- Parge HE, Getzoff ED, Scandella CS, Hallewell RA and Tainer JA (1986) Crystallographic characterization of recombinant human CuZn superoxide dismutase. *Journal of Biological Chemistry* **261**, 16215–16218.
- Parge HE, Hallewell RA and Tainer JA (1992) Atomic structures of wild-type and thermostable mutant recombinant human Cu,Zn superoxide dismutase. *Proceedings of the National Academy of Sciences of the USA* **89**, 6109–6113.
- Pasinelli P, Belford ME, Lennon N, Bacskai BJ, Hyman BT, Trotti D and Brown RH (2004) Amyotrophic lateral sclerosis-associated SOD1 mutant proteins bind and aggregate with Bcl-2 in spinal cord mitochondria. *Neuron* **43**, 19–30.
- Pauling L (1931) The nature of the chemical bond. ii. the one-electron bond and the three-electron bond. *Journal of the American Chemical Society* **53**, 3225–3237.

- Perry JJP, Shin DS, Getzoff ED and Tainer JA (2010) The structural biochemistry of the superoxide dismutases. *Biochimica et Biophysica Acta (BBA) – Proteins and Proteomics* **1804**, 245–262.
- Petersen MTN, Jonson PH and Petersen SB (1999) Amino acid neighbours and detailed conformational analysis of cysteines in proteins. *Protein Engineering, Design & Selection* **12**, 535–548.
- Petkau A and Chelack WS (1976) Radioprotective effect of superoxide dismutase on model phospholipid membranes. *Biochimica et Biophysica Acta (BBA) – Biomembranes* **433**, 445–456.
- Petkau A and Copps TP (1979) Distribution of superoxide dismutase in the erythrocyte membrane. *Biophysical Journal* **25**, 190a.
- Pickles S, Destroismans L, Peyrard SL, Cadot S, Rouleau GA, Brown RH, Julien J-P, Arbour N and Vande Velde C (2013) Mitochondrial damage revealed by immunoselection for ALS-linked misfolded SOD1. *Human Molecular Genetics* **22**, 3947–3959.
- Pokrishevsky E, Grad LI, Yousefi M, Wang J, Mackenzie IR and Cashman NR (2012) Aberrant localization of FUS and TDP43 is associated with misfolding of SOD1 in amyotrophic lateral sclerosis. *PLoS ONE* **7**, e35050.
- Pokrishevsky E, Hong RH, Mackenzie IR and Cashman NR (2017) Spinal cord homogenates from SOD1 familial amyotrophic lateral sclerosis induce SOD1 aggregation in living cells. *PLoS ONE* **12**, e0184384.
- Pokrishevsky E, McAlary L, Farrowell NE, Zhao B, Sher M, Yerbury JJ and Cashman NR (2018) Tryptophan 32-mediated SOD1 aggregation is attenuated by pyrimidine-like compounds in living cells. *Scientific Reports* **8**, 15590.
- Porter H, Folch J, Beggrovs J and Ainsworth S (1957) Cerebrocuprein I. a copper-containing protein isolated from brain. *Journal of Neurochemistry* **1**, 260–271.
- Potter SZ, Zhu H, Shaw BF, Rodriguez JA, Doucette PA, Sohn SH, Durazo A, Faull KF, Gralla EB, Nersissian AM and Valentine JS (2007) Binding of a single zinc ion to one subunit of copper–zinc superoxide dismutase apoprotein substantially influences the structure and stability of the entire homodimeric protein. *Journal of the American Chemical Society* **129**, 4575–4583.
- Pramatarova A, Figlewicz DA, Krizus A, Han FY, Ceballos-Picot I, Nicole A, Dib M, Meininger V, Brown RH and Rouleau GA (1995) Identification of new mutations in the Cu/Zn superoxide dismutase gene of patients with familial amyotrophic lateral sclerosis. *American Journal of Human Genetics* **56**, 592–596.
- Pratt AJ, Shin DS, Merz GE, Rambo RP, Lancaster WA, Dyer KN, Borbat PP, Poole FL, Adams MWW, Freed JH, Crane BR, Tainer JA and Getzoff ED (2014) Aggregation propensities of superoxide dismutase G93 hotspot mutants mirror ALS clinical phenotypes. *Proceedings of the National Academy of Sciences of the USA* **111**, E4568–E4576.
- Proctor EA, Fee L, Tao Y, Redler RL, Fay JM, Zhang Y, Lv Z, Mercer IP, Deshmukh M, Lyubchenko YL and Dokholyan NV (2016) Nonnative SOD1 trimer is toxic to motor neurons in a model of amyotrophic lateral sclerosis. *Proceedings of the National Academy of Sciences* **113**, 614–619.
- Proescher JB, Son M, Elliott JL and Culotta VC (2008) Biological effects of CCS in the absence of SOD1 enzyme activation: implications for disease in a mouse model for ALS. *Human Molecular Genetics* **17**, 1728–1737.
- Prudencio M, Hart PJ, Borchelt DR and Andersen PM (2009) Variation in aggregation propensities among ALS-associated variants of SOD1: correlation to human disease. *Human Molecular Genetics* **18**, 3217–3226.
- Rabani J and Stein G (1962) The radiation chemistry of aqueous solutions of cytochrome c. *Radiation Research* **17**, 327–340.
- Rabizadeh S, Gralla EB, Borchelt DR, Gwinn R, Valentine JS, Sisodia S, Wong P, Lee M, Hahn H and Bredesen DE (1995) Mutations associated with amyotrophic lateral sclerosis convert superoxide dismutase from an anti-apoptotic gene to a proapoptotic gene: studies in yeast and neural cells. *Proceedings of the National Academy of Sciences of the USA* **92**, 3024–3028.
- Rae TD, Schmidt PJ, Pufahl RA, Culotta VC and O'Halloran TV (1999) Undetectable intracellular free copper: the requirement of a copper chaperone for superoxide dismutase. *Science* **284**, 805–808.
- Rakhit R, Cunningham P, Furtos-Matei A, Dahan S, Qi X-F, Crow JP, Cashman NR, Kondejewski LH and Chakrabarty A (2002) Oxidation-induced misfolding and aggregation of superoxide dismutase and its implications for amyotrophic lateral sclerosis. *Journal of Biological Chemistry* **277**, 47551–47556.
- Rakhit R, Crow JP, Lepock JR, Kondejewski LH, Cashman NR and Chakrabarty A (2004) Monomeric Cu,Zn-superoxide dismutase is a common misfolding intermediate in the oxidation models of sporadic and familial amyotrophic lateral sclerosis. *Journal of Biological Chemistry* **279**, 15499–15504.
- Rakhit R, Robertson J, Vande Velde C, Horne P, Ruth DM, Griffin J, Cleveland DW, Cashman NR and Chakrabarty A (2007) An immunological epitope selective for pathological monomer-misfolded SOD1 in ALS. *Nature Medicine* **13**, 754–759.
- Ramu M, Unni S, Wright GSA, Srinivas BMM and Padmanabhan B (2018) Rational discovery of a SOD1 tryptophan oxidation inhibitor with therapeutic potential for amyotrophic lateral sclerosis. *Journal of Biomolecular Structure & Dynamics* **37**, 3936–3946.
- Rando A, de la Torre M, Martinez-Muriana A, Zaragoza P, Musaro A, Hernández S, Navarro X, Toivonen JM and Osta R (2019) Chemotherapeutic agent 5-fluorouracil increases survival of SOD1 mouse model of ALS. *PLoS ONE* **14**, e0210752.
- Rasouli S, Abdolvahabi A, Croom CM, Plewman DL, Shi Y, Ayers JI and Shaw BF (2017) Lysine acylation in superoxide dismutase-1 electrostatically inhibits formation of fibrils with prion-like seeding. *Journal of Biological Chemistry* **292**, 19366–19380.
- Rasouli S, Abdolvahabi A, Croom CM, Plewman D, Shi Y and Shaw BF (2018) Glycerolipid headgroups control rate and mechanism of SOD1 aggregation and accelerate fibrillization of slowly aggregating ALS mutants. *ACS Chemical Neuroscience* **9**, 1743–1756.
- Ravits JM and La Spada AR (2009) ALS motor phenotype heterogeneity, focality, and spread: deconstructing motor neuron degeneration. *Neurology* **73**, 805–811.
- Ray SS, Nowak RJ, Strokovich K, Brown RH, Walz T and Lansbury PT (2004) An intersubunit disulfide bond prevents in vitro aggregation of a superoxide dismutase-1 mutant linked to familial amyotrophic lateral sclerosis. *Biochemistry* **43**, 4899–4905.
- Reaume AG, Elliott JL, Hoffman EK, Kowall NW, Ferrante RJ, Siwek DE, Wilcox HM, Flood DG, Beal MF, Brown RH, Scott RW and Snider WD (1996) Motor neurons in Cu/Zn superoxide dismutase-deficient mice develop normally but exhibit enhanced cell death after axonal injury. *Nature Genetics* **13**, 43–47.
- Redler RL, Wilcox KC, Proctor EA, Fee L, Caplow M and Dokholyan NV (2011) Glutathionylation at Cys-111 induces dissociation of wild type and FALS mutant SOD1 dimers. *Biochemistry* **50**, 7057–7066.
- Richardson JS (1977) beta-Sheet topology and the relatedness of proteins. *Nature* **268**, 495–500.
- Richardson DC, Bier CJ and Richardson JS (1972) Two crystal forms of bovine superoxide dismutase. *Journal of Biological Chemistry* **247**, 6368–6369.
- Richardson J, Thomas KA, Rubin BH and Richardson DC (1975a) Crystal structure of bovine Cu,Zn superoxide dismutase at 3 Å resolution: chain tracing and metal ligands. *Proceedings of the National Academy of Sciences of the USA* **72**, 1349–1353.
- Richardson JS, Thomas KA and Richardson DC (1975b) Alpha-carbon coordinates for bovine Cu, Zn superoxide dismutase. *Biochemical and Biophysical Research Communications* **63**, 986–992.
- Richardson JS, Richardson DC, Thomas KA, Silverton EW and Davies DR (1976) Similarity of three-dimensional structure between the immunoglobulin domain and the copper, zinc superoxide dismutase subunit. *Journal of Molecular Biology* **102**, 221–235.
- Robberecht W, Aguirre T, Van den Bosch L, Tilkin P, Cassiman JJ and Matthijs G (1996) D90a heterozygosity in the SOD1 gene is associated with familial and apparently sporadic amyotrophic lateral sclerosis. *Neurology* **47**, 1336–1339.
- Robberecht W, Rousseau F and Schymkowitz J (2016) Single domain antibodies against sod1 and their use in medicine. *Patents US20160115245A1 and WO2014191493A1*.
- Robinett NG, Culbertson EM, Peterson RL, Sanchez H, Andes D, Nett JE and Culotta VC (2018) Exploiting the vulnerable active site of a copper-only superoxide dismutase to disrupt fungal pathogenesis. *Journal of Biological Chemistry* **294**, 2700–2713.

- Robinson NE and Robinson AB** (2001) Molecular clocks. *Proceedings of the National Academy of Sciences of the USA* **98**, 944–949.
- Rodriguez JA, Valentine JS, Eggers DK, Roe JA, Tiwari A, Brown RH and Hayward LJ** (2002) Familial amyotrophic lateral sclerosis-associated mutations decrease the thermal stability of distinctly metallated species of human copper/zinc superoxide dismutase. *Journal of Biological Chemistry* **277**, 15932–15937.
- Rodriguez JA, Shaw BF, Durazo A, Sohn SH, Doucette PA, Nersissian AM, Faull KF, Eggers DK, Tiwari A, Hayward LJ and Valentine JS** (2005) Destabilization of apoprotein is insufficient to explain Cu,Zn-superoxide dismutase-linked ALS pathogenesis. *Proceedings of the National Academy of Sciences* **102**, 10516–10521.
- Rosen DR, Siddique T, Patterson D, Figlewicz DA, Sapp P, Hentati A, Donaldson D, Goto J, O'Regan JP and Deng HX** (1993) Mutations in Cu/Zn superoxide dismutase gene are associated with familial amyotrophic lateral sclerosis. *Nature* **362**, 59–62.
- Rosen DR, Bowling AC, Patterson D, Usdin TB, Sapp P, Mezey E, McKenna-Yasek D, O'Regan J, Rahmani Z and Ferrante RJ** (1994) A frequent ala 4 to val superoxide dismutase-1 mutation is associated with a rapidly progressive familial amyotrophic lateral sclerosis. *Human Molecular Genetics* **3**, 981–987.
- Rothstein JD, Bristol LA, Hosler B, Brown RH and Kuncel RW** (1994) Chronic inhibition of superoxide dismutase produces apoptotic death of spinal neurons. *Proceedings of the National Academy of Sciences of the USA* **91**, 4155–4159.
- Rothstein JD, Dykes-Hoberg M, Corson LB, Becker M, Cleveland DW, Price DL, Culotta VC and Wong PC** (1999) The copper chaperone CCS is abundant in neurons and astrocytes in human and rodent brain. *Journal of Neurochemistry* **72**, 422–429.
- Rotilio G, Calabrese L, Mondovi B and Blumberg WE** (1974) Electron paramagnetic resonance studies of cobalt-copper bovine superoxide dismutase. *Journal of Biological Chemistry* **249**, 3157–3160.
- Rumfeldt JAO, Stathopoulos PB, Chakraborty A, Lepock JR and Meiering EM** (2006) Mechanism and thermodynamics of guanidinium chloride-induced denaturation of ALS-associated mutant Cu,Zn superoxide dismutases. *Journal of Molecular Biology* **355**, 106–123.
- Sábado J, Casanovas A, Rodrigo H, Arqué G and Esquerda JE** (2015) Adverse effects of a SOD1-peptide immunotherapy on SOD1(G93A) mouse slow model of amyotrophic lateral sclerosis. *Neuroscience* **310**, 38–50.
- Saccon RA, Bunton-Stasyshyn RKA, Fisher EMC and Fratta P** (2013) Is SOD1 loss of function involved in amyotrophic lateral sclerosis? *Journal of Brain and Neurology* **136**, 2342–2358.
- Sakuma R, Abe K, Aoki M, Ikeda M, Okita N, Hiwatari M, Sakurai M and Itoyama Y** (1995) A clinical variance in familial amyotrophic lateral sclerosis with a point mutation in Cu/Zn superoxide dismutase gene. *European Journal of Neurology* **2**, 369–374.
- Sala FA, Wright GSA, Antonyuk SV, Garratt RC and Hasnain SS** (2019) Molecular recognition and maturation of SOD1 by its evolutionarily destabilised cognate chaperone hCCS. *PLoS Biology* **17**, e3000141.
- Salehi M, Nikkhal M, Ghasemi A and Arab SS** (2015) Mitochondrial membrane disruption by aggregation products of ALS-causing superoxide dismutase-1 mutants. *International Journal of Biological Macromolecules* **75**, 290–297.
- Sandelin E, Nordlund A, Andersen PM, Marklund SSL and Oliveberg M** (2007) Amyotrophic lateral sclerosis-associated copper/zinc superoxide dismutase mutations preferentially reduce the repulsive charge of the proteins. *Journal of Biological Chemistry* **282**, 21230–21236.
- Sangwan S, Zhao A, Adams KL, Jayson CK, Sawaya MR, Guenther EL, Pan AC, Ngo J, Moore DM, Soriaga AB, Do TD, Goldschmidt L, Nelson R, Bowers MT, Koehler CM, Shaw DE, Novitch BG and Eisenberg DS** (2017) Atomic structure of a toxic, oligomeric segment of SOD1 linked to amyotrophic lateral sclerosis (ALS). *Proceedings of the National Academy of Sciences of the USA* **114**, 8770–8775.
- Sato T, Nakanishi T, Yamamoto Y, Andersen PM, Ogawa Y, Fukada K, Zhou Z, Aoike F, Sugai F, Nagano S, Hirata S, Ogawa M, Nakano R, Ohi T, Kato T, Nakagawa M, Hamasaki T, Shimizu A and Sakoda S** (2005) Rapid disease progression correlates with instability of mutant SOD1 in familial ALS. *Neurology* **65**, 1954–1957.
- Schmidt PJ, Ramos-Gomez M and Culotta VC** (1999) A gain of superoxide dismutase (SOD) activity obtained with CCS, the copper metallochaperone for SOD1. *Journal of Biological Chemistry* **274**, 36952–36956.
- Seetharaman SV, Winkler DD, Taylor AB, Cao X, Whitson LJ, Doucette PA, Valentine JS, Schirf V, Demeler B, Carroll MC, Culotta VC and Hart PJ** (2010) Disrupted zinc-binding sites in structures of pathogenic SOD1 variants D124V and H80R. *Biochemistry* **49**, 5714–5725.
- Segovia-Silvestre T, Andreu AL, Vives-Bauza C, Garcia-Arumi E, Cervera C and Gamez J** (2002) A novel exon 3 mutation (D76 V) in the SOD1 gene associated with slowly progressive ALS. *Amyotrophic Lateral Sclerosis and Other Motor Neuron Disorders* **3**, 69–74.
- Sekhar A, Rumfeldt JA, Broom HR, Doyle CM, Bouvignies G, Meiering EM and Kay LE** (2015) Thermal fluctuations of immature SOD1 lead to separate folding and misfolding pathways. *ELife* **4**, e07296.
- Sekhar A, Rumfeldt JAO, Broom HR, Doyle CM, Sobering RE, Meiering EM and Kay LE** (2016) Probing the free energy landscapes of ALS disease mutants of SOD1 by NMR spectroscopy. *Proceedings of the National Academy of Sciences of the USA* **113**, E6939–E6945.
- Shabalin I, Dauter Z, Jaskolski M, Minor W and Wlodawer A** (2015) Crystallography and chemistry should always go together: a cautionary tale of protein complexes with cisplatin and carboplatin. *Acta Crystallographica. Section D, Biological Crystallography* **71**, 1965–1979.
- Shaw BF and Valentine JS** (2007) How do ALS-associated mutations in superoxide dismutase 1 promote aggregation of the protein? *Trends in Biochemical Sciences* **32**, 78–85.
- Shaw BF, Durazo A, Nersissian AM, Whitelegge JP, Faull KF and Valentine JS** (2006) Local unfolding in a destabilized, pathogenic variant of superoxide dismutase 1 observed with H/D exchange and mass spectrometry. *Journal of Biological Chemistry* **281**, 18167–18176.
- Shaw BF, Moustakas DT, Whitelegge JP and Faull KF** (2010) Taking charge of proteins: from neurodegeneration to industrial biotechnology. *Advances in Protein Chemistry and Structural Biology* **79**, 127–164.
- Shaw BF, Lelie HL, Durazo A, Nersissian AM, Xu G, Chan PK, Gralla EB, Tiwari A, Hayward LJ, Borchelt DR, Valentine JS and Whitelegge JP** (2008) Detergent-insoluble aggregates associated with amyotrophic lateral sclerosis in transgenic mice contain primarily full-length, unmodified superoxide dismutase-1. *Journal of Biological Chemistry* **283**, 8340–8350.
- Shi Y, Mowery RA and Shaw BF** (2013a) Effect of metal loading and subcellular pH on net charge of superoxide dismutase-1. *Journal of Molecular Biology* **425**, 4388–4401.
- Shi Y, Rhodes NR, Abdolvahabi A, Kohn TP, Cook NP, Martí AA and Shaw BF** (2013b) Deamidation of asparagine to aspartate destabilizes Cu, Zn superoxide dismutase, accelerates fibrillization and mirrors ALS-linked mutations. *Journal of the American Chemical Society* **135**, 15897–15908.
- Shi Y, Acerson MJ, Abdolvahabi A, Mowery RA and Shaw BF** (2016) Gibbs energy of superoxide dismutase heterodimerization accounts for Variable survival in amyotrophic lateral sclerosis. *Journal of the American Chemical Society* **138**, 5351–5362.
- Shibata N, Hirano A, Kobayashi M, Asayama K, Umahara T and Ikemoto A** (1993) Immunohistochemical demonstration of Cu/Zn superoxide dismutase in the spinal cord of patients with familial amyotrophic lateral sclerosis. *Acta Histochemica et Cytochemica* **26**, 619–622.
- Shibata N, Hirano A, Kobayashi M, Sasaki S, Kato T, Matsumoto S, Shiozawa Z, Komori T, Ikemoto A and Umahara T** (1994) Cu/Zn superoxide dismutase-like immunoreactivity in Lewy body-like inclusions of sporadic amyotrophic lateral sclerosis. *Neuroscience Letters* **179**, 149–152.
- Shibata N, Hirano A, Kobayashi M, Siddique T, Deng HX, Hung WY, Kato T and Asayama K** (1996a) Intense superoxide dismutase-1 immunoreactivity in intracytoplasmic hyaline inclusions of familial amyotrophic lateral sclerosis with posterior column involvement. *Journal of Neuropathology & Experimental Neurology* **55**, 481–490.
- Shibata N, Asayama K, Hirano A and Kobayashi M** (1996b) Immunohistochemical study on superoxide dismutases in spinal cords from autopsied patients with amyotrophic lateral sclerosis. *Developmental Neuroscience* **18**, 492–498.
- Shibata N, Hirano A, Kato S, Nagai R, Horiuchi S, Komori T, Umahara T, Asayama K and Kobayashi M** (1999) Advanced glycation endproducts are deposited in neuronal hyaline inclusions: a study on familial amyotrophic

- lateral sclerosis with superoxide dismutase-1 mutation. *Acta Neuropathologica* **97**, 240–246.
- Shvil N, Banerjee V, Zoltsman G, Shani T, Kahn J, Abu-Hamad S, Papo N, Engel S, Bernhagen J and Israelson A** (2018) MIF inhibits the formation and toxicity of misfolded SOD1 amyloid aggregates: implications for familial ALS. *Cell death & disease* **9**, 107.
- Siddique T, Figlewicz DA, Pericak-Vance MA, Haines JL, Rouleau G, Jeffers AJ, Sapp P, Hung W-Y, Bebout J, McKenna-Yasek D, Deng G, Horvitz HR, Gusella JF, Brown RH Jr and Roses AD** (1991) Linkage of a gene causing familial amyotrophic lateral sclerosis to chromosome 21 and evidence of genetic-locus heterogeneity. *New England Journal of Medicine* **324**, 1381–1384.
- Sirangelo I, Vella FM, Irace G, Manco G and Iannuzzi C** (2016) Glycation in demetallated superoxide dismutase 1 prevents amyloid aggregation and produces cytotoxic ages adducts. *Frontiers in Molecular Biosciences* **3**, 55.
- Smith MW and Doolittle RF** (1992) A comparison of evolutionary rates of the two major kinds of superoxide dismutase. *Journal of Molecular Evolution* **34**, 175–184.
- Son M, Puttapparthi K, Kawamata H, Rajendran B, Boyer PJ, Manfredi G and Elliott JL** (2007) Overexpression of CCS in G93A-SOD1 mice leads to accelerated neurological deficits with severe mitochondrial pathology. *Proceedings of the National Academy of Sciences of the USA* **104**, 6072–6077.
- Son M, Fu Q, Puttapparthi K, Matthews CM and Elliott JL** (2009) Redox susceptibility of SOD1 mutants is associated with the differential response to CCS over-expression *in vivo*. *Neurobiology of Disease* **34**, 155–162.
- Sone J, Niwa J, Kawai K, Ishigaki S, Yamada S, Adachi H, Katsuno M, Tanaka F, Doyu M and Sobue G** (2010) Dorfin ameliorates phenotypes in a transgenic mouse model of amyotrophic lateral sclerosis. *Journal of Neuroscience Research* **88**, 123–135.
- Soon CPW, Donnelly PS, Turner BJ, Hung LW, Crouch PJ, Sherratt NA, Tan J-L, Lim NK-H, Lam L, Bica L, Lim S, Hickey JL, Morizzi J, Powell A, Finkelstein DI, Culvenor JG, Masters CL, Duce J, White AR, Barnham KJ and Li QX** (2011) Diacetylbis(N(4)-methylthiosemicarbazono) copper(II) (CuII(atm)) protects against peroxynitrite-induced nitrosative damage and prolongs survival in amyotrophic lateral sclerosis mouse model. *Journal of Biological Chemistry* **286**, 44035–44044.
- Stathopoulos PB, Rumpfolt JAO, Scholz GA, Irani RA, Frey HE, Hallewell RA, Lepock JR and Meiering EM** (2003) Cu/Zn superoxide dismutase mutants associated with amyotrophic lateral sclerosis show enhanced formation of aggregates *in vitro*. *Proceedings of the National Academy of Sciences of the USA* **100**, 7021–7026.
- Strange RW, Antonyuk S, Hough MA, Doucette PA, Rodriguez JA, Hart PJ, Hayward LJ, Valentine JS and Hasnain SS** (2003) The structure of holo and metal-deficient wild-type human Cu, Zn superoxide dismutase and its relevance to familial amyotrophic lateral sclerosis. *Journal of Molecular Biology* **328**, 877–891.
- Strange RW, Antonyuk SV, Hough MA, Doucette PA, Valentine JS and Hasnain SS** (2006) Variable metallation of human superoxide dismutase: atomic resolution crystal structures of Cu-Zn, Zn-Zn and as-isolated wild-type enzymes. *Journal of Molecular Biology* **356**, 1152–1162.
- Subramaniam JR, Lyons WE, Liu J, Bartnikas TB, Rothstein J, Price DL, Cleveland DW, Gitlin JD and Wong PC** (2002) Mutant SOD1 causes motor neuron disease independent of copper chaperone-mediated copper loading. *Nature Neuroscience* **5**, 301–307.
- Suthers G, Laing N, Wilton S, Dorosz S and Waddy H** (1994) ‘Sporadic’ motoneuron disease due to familial SOD1 mutation with low penetrance. *Lancet* **344**, 1773.
- Suzuki M, Yasui K, Ishikawai H, Nomura M, Watanabe T, Mikami H, Yamano T and Ono S** (2011) Familial amyotrophic lateral sclerosis with Cys111Tyr mutation in Cu/Zn superoxide dismutase showing widespread Lewy body-like hyaline inclusions. *Journal of the Neurological Sciences* **300**, 182–184.
- Tainer JA, Getzoff ED, Beem KM, Richardson JS and Richardson DC** (1982) Determination and analysis of the 2 A-structure of copper, zinc superoxide dismutase. *Journal of Molecular Biology* **160**, 181–217.
- Tainer JA, Getzoff ED, Richardson JS and Richardson DC** (1983) Structure and mechanism of copper, zinc superoxide dismutase. *Nature* **306**, 284–287.
- Takamiya R, Takahashi M, Myint T, Park YS, Miyazawa N, Endo T, Fujiwara N, Sakiyama H, Misonou Y, Miyamoto Y, Fujii J and Taniguchi N** (2003) Glycation proceeds faster in mutated Cu, Zn-superoxide dismutases related to familial amyotrophic lateral sclerosis. *FASEB Journal* **17**, 938–940.
- Takazawa T, Ikeda K, Hirayama T, Kawabe K, Nakamura Y, Ito H, Kano O, Yoshii Y, Tanaka F, Sobue G and Iwasaki Y** (2010) Familial amyotrophic lateral sclerosis with a novel G85S mutation of superoxide dismutase 1 gene: clinical features of lower motor neuron disease. *Internal Medicine (Tokyo, Japan)* **49**, 183–186.
- Tally FP, Goldin BR, Jacobus NV and Gorbach SL** (1977) Superoxide dismutase in anaerobic bacteria of clinical significance. *Infection and Immunity* **16**, 20–25.
- Taylor DM, Gibbs BF, Kabashi E, Minotti S, Durham HD and Agar JN** (2007) Tryptophan 32 potentiates aggregation and cytotoxicity of a copper/zinc superoxide dismutase mutant associated with familial amyotrophic lateral sclerosis. *Journal of Biological Chemistry* **282**, 16329–16335.
- Thomas KA, Rubin BH, Bier CJ, Richardson JS and Richardson DC** (1974) The crystal structure of bovine Cu₂Zn₂ superoxide dismutase at 5.5-Å resolution. *Journal of Biological Chemistry* **249**, 5677–5683.
- Thornton JM** (1981) Disulphide bridges in globular proteins. *Journal of Molecular Biology* **151**, 261–287.
- Tiwari A and Hayward LJ** (2003) Familial amyotrophic lateral sclerosis mutants of copper/zinc superoxide dismutase are susceptible to disulfide reduction. *Journal of Biological Chemistry* **278**, 5984–5992.
- Tiwari A and Hayward LJ** (2005) Mutant SOD1 instability: implications for toxicity in amyotrophic lateral sclerosis. *Neurodegenerative Diseases* **2**, 115–127.
- Trist BG, Davies KM, Cottam V, Genoud S, Ortega R, Roudeau S, Carmona A, De Silva K, Wasinger V, Lewis SJG, Sachdev P, Smith B, Troakes C, Vance C, Shaw C, Al-Sarraj S, Ball HJ, Halliday GM, Hare DJ and Double KL** (2017) Amyotrophic lateral sclerosis-like superoxide dismutase 1 proteinopathy is associated with neuronal loss in Parkinson’s disease brain. *Acta Neuropathologica* **134**, 113–127.
- Troy CM and Shelanski ML** (1994) Down-regulation of copper/zinc superoxide dismutase causes apoptotic death in PC12 neuronal cells. *Proceedings of the National Academy of Sciences* **91**, 6384–6387.
- Tsang CK, Liu Y, Thomas J, Zhang Y and Zheng XFS** (2014) Superoxide dismutase 1 acts as a nuclear transcription factor to regulate oxidative stress resistance. *Nature Communications* **5**, 3446.
- Uchida K and Kawakishi S** (1994) Identification of oxidized histidine generated at the active site of Cu,Zn-superoxide dismutase exposed to H₂O₂. Selective generation of 2-oxo-histidine at the histidine 118. *Journal of Biological Chemistry* **269**, 2405–2410.
- Urushitani M, Sik A, Sakurai T, Nukina N, Takahashi R and Julien J-P** (2006) Chromogranin-mediated secretion of mutant superoxide dismutase proteins linked to amyotrophic lateral sclerosis. *Nature Neuroscience* **9**, 108–118.
- Urushitani M, Ezzi SA and Julien J-P** (2007) Therapeutic effects of immunization with mutant superoxide dismutase in mice models of amyotrophic lateral sclerosis. *Proceedings of the National Academy of Sciences of the USA* **104**, 2495–2500.
- van Blitterswijk M, Gulati S, Smoot E, Jaffa M, Maher N, Hyman BT, Ivinson AJ, Scherzer CR, Schoenfeld DA, Cudkowicz ME, Brown RH Jr and Bosco DA** (2011) Anti-superoxide dismutase antibodies are associated with survival in patients with sporadic amyotrophic lateral sclerosis. *Amyotrophic Lateral Sclerosis* **12**, 430–438.
- Vande Velde C, Miller TM, Cashman NR and Cleveland DW** (2008) Selective association of misfolded ALS-linked mutant SOD1 with the cytoplasmic face of mitochondria. *Proceedings of the National Academy of Sciences of the USA* **105**, 4022–4027.
- Vassall KA, Stubbs HR, Primmer HA, Tong MS, Sullivan SM, Sobering R, Srinivasan S, Briere L-AK, Dunn SD, Colón W and Meiering EM** (2011) Decreased stability and increased formation of soluble aggregates by immature superoxide dismutase do not account for disease severity in ALS. *Proceedings of the National Academy of Sciences of the USA* **108**, 2210–2215.
- Vieira FG, Hatzipetros T, Thompson K, Moreno AJ, Kidd JD, Tassinari VR, Levine B, Perrin S and Gill A** (2017) CuATSM efficacy is independently replicated in a SOD1 mouse model of ALS while

- unmetallated ATSM therapy fails to reveal benefits. *IBRO Reports* 2, 47–53.
- Vijayvergiya C, Beal MF, Buck J and Manfredi G (2005) Mutant superoxide dismutase 1 forms aggregates in the brain mitochondrial matrix of amyotrophic lateral sclerosis mice. *Journal of Neuroscience* 25, 2463–2470.
- Wang J, Xu G, Gonzales V, Coonfield M, Fromholt D, Copeland NG, Jenkins NA and Borchelt DR (2002) Fibrillar inclusions and motor neuron degeneration in transgenic mice expressing superoxide dismutase 1 with a disrupted copper-binding site. *Neurobiology of Disease* 10, 128–138.
- Wang Q, Johnson JL, Agar NYR and Agar JN (2008) Protein aggregation and protein instability govern familial amyotrophic lateral sclerosis patient survival. *PLoS Biology* 6, e170.
- Warburg O and Krebs H (1927) Über locker gebundenes kupfer und eisen in blutserum. *Biochem Z* 190, 143.
- Watanabe M, Dykes-Hoberg M, Culotta VC, Price DL, Wong PC and Rothstein JD (2001) Histological evidence of protein aggregation in mutant SOD1 transgenic mice and in amyotrophic lateral sclerosis neural tissues. *Neurobiology of Disease* 8, 933–941.
- Watanabe S, Ilieva H, Tamada H, Nomura H, Komine O, Endo F, Jin S, Mancias P, Kiyama H and Yamanaka K (2016) Mitochondria-associated membrane collapse is a common pathomechanism in SIGMAR1- and SOD1-linked ALS. *EMBO Molecular Medicine* 8, 1421–1437.
- Watanabe M, Aoki M, Abe K, Shoji M, Iizuka T, Ikeda Y, Hirai S, Kurokawa K, Kato T, Sasaki H and Itoyama Y (1997) A novel missense point mutation (S134N) of the Cu/Zn superoxide dismutase gene in a patient with familial motor neuron disease. *Human Mutation* 9, 69–71.
- Weisberg SJ, Lyakhovetsky R, Werdiger A, Gitler AD, Soen Y and Kaganovich D (2012) Compartmentalization of superoxide dismutase 1 (SOD1G93A) aggregates determines their toxicity. *Proceedings of the National Academy of Sciences of the USA* 109, 15811–15816.
- Weiss MD, Ravits JM, Schuman N and Carter GT (2006) A4 V superoxide dismutase mutation in apparently sporadic ALS resembling neuralgic amyotrophy. *Amyotrophic Lateral Sclerosis* 7, 61–63.
- Wilcox KC, Zhou L, Jordon J, Huang Y, Yu Y, Redler RL, Chen X, Caplow M and Dokholyan NV (2009) Modifications of superoxide dismutase (SOD1) in human erythrocytes: a possible role in amyotrophic lateral sclerosis (ALS). *Journal of Biological Chemistry* 284, 13940–13947.
- Williams JR, Trias E, Beilby PR, Lopez NI, Labut EM, Samuel Bradford C, Roberts BR, McAllum EJ, Crouch PJ, Rhoads TW, Pereira C, Son M, Elliott JL, Franco MC, Estévez AG, Barbeito L and Beckman JS (2016) Copper delivery to the CNS by CuATSM effectively treats motor neuron disease in SOD(G93A) mice co-expressing the copper-chaperone-for-SOD. *Neurobiology of Disease* 89, 1–9.
- Winkler DD, Schuermann JP, Cao X, Holloway SP, Borchelt DR, Carroll MC, Proescher JB, Culotta VC and Hart PJ (2009) Structural and biophysical properties of the pathogenic SOD1 variant H46R/H48Q. *Biochemistry* 48, 3436–3447.
- Wong PC, Pardo CA, Borchelt DR, Lee MK, Copeland NG, Jenkins NA, Sisodia SS, Cleveland DW and Price DL (1995) An adverse property of a familial ALS-linked SOD1 mutation causes motor neuron disease characterized by vacuolar degeneration of mitochondria. *Neuron* 14, 1105–1116.
- Wong H-S, Dighe PA, Mezera V, Montnerier P-A and Brand MD (2017) Production of superoxide and hydrogen peroxide from specific mitochondrial sites under different bioenergetic conditions. *Journal of Biological Chemistry* 292, 16804–16809.
- Wood E, Dagleish D and Bannister W (1971) Bovine erythrocyte cupro-zinc protein. *European Journal of Biochemistry* 18, 187–193.
- Wright HT (1991) Sequence and structure determinants of the nonenzymatic deamidation of asparagine and glutamine residues in proteins. *Protein Engineering* 4, 283–294.
- Wright GSA, Hasnain SS and Grossmann JG (2011) The structural plasticity of the human copper chaperone for SOD1: insights from combined size-exclusion chromatographic and solution X-ray scattering studies. *Biochemical Journal* 439, 39–44.
- Wright GSA, Antonyuk SV, Kershaw NM, Strange RW and Samar Hasnain S (2013) Ligand binding and aggregation of pathogenic SOD1. *Nature Communications* 4, 1758.
- Wright GSA, Antonyuk SV and Hasnain SS (2016) A faulty interaction between SOD1 and hCCS in neurodegenerative disease. *Scientific Reports* 6, 27691.
- Wu J, Shen E, Shi D, Sun Z and Cai T (2012) Identification of a novel Cys146X mutation of SOD1 in familial amyotrophic lateral sclerosis by whole-exome sequencing. *Genetics in Medicine* 14, 823–826.
- Xu W-C, Liang J-Z, Li C, He Z-X, Yuan H-Y, Huang B-Y, Liu X-L, Tang B, Pang D-W, Du H-N, Yang Y, Chen J, Wang L, Zhang M and Liang Y (2018) Pathological hydrogen peroxide triggers the fibrillization of wild-type SOD1 via sulfenic acid modification of Cys-111. *Cell death & disease* 9, 67.
- Yamanaka K, Chun SJ, Boillee S, Fujimori-Tonou N, Yamashita H, Gutmann DH, Takahashi R, Misawa H and Cleveland DW (2008) Astrocytes as determinants of disease progression in inherited amyotrophic lateral sclerosis. *Nature Neuroscience* 11, 251–253.
- Yang F, Wang H, Logan DT, Mu X, Danielsson J and Oliveberg M (2018) The cost of long catalytic loops in folding and stability of the ALS-associated protein SOD1. *Journal of the American Chemical Society* 140, 16570–16579.
- Yasser S, Fecto F, Siddique T, Sheikh KA and Athar P (2010) An unusual case of familial ALS and cerebellar ataxia. *Amyotrophic Lateral Sclerosis* 11, 568–570.
- Yim HS, Kang JH, Chock PB, Stadtman ER and Yim MB (1997) A familial amyotrophic lateral sclerosis-associated A4 V Cu, Zn-superoxide dismutase mutant has a lower Km for hydrogen peroxide. Correlation between clinical severity and the Km value. *Journal of Biological Chemistry* 272, 8861–8863.
- Youn HD, Kim EJ, Roe JH, Hah YC and Kang SO (1996) A novel nickel-containing superoxide dismutase from *Streptomyces* spp. *Biochemical Journal* 318, 889–896.
- Zetterström P, Stewart HG, Bergemalm D, Jonsson PA, Graffmo KS, Andersen PM, Brännström T, Oliveberg M and Marklund SL (2007) Soluble misfolded subfractions of mutant superoxide dismutase-1s are enriched in spinal cords throughout life in murine ALS models. *Proceedings of the National Academy of Sciences of the USA* 104, 14157–14162.
- Zetterström P, Graffmo KS, Andersen PM, Brännström T and Marklund SL (2011) Proteins that bind to misfolded mutant superoxide dismutase-1 in spinal cords from transgenic amyotrophic lateral sclerosis (ALS) model mice. *Journal of Biological Chemistry* 286, 20130–20136.
- Zhang H, Andrekopoulos C, Joseph J, Chandran K, Karoui H, Crow JP and Kalyanaraman B (2003) Bicarbonate-dependent peroxidase activity of human Cu,Zn-superoxide dismutase induces covalent aggregation of protein: intermediacy of tryptophan-derived oxidation products. *Journal of Biological Chemistry* 278, 24078–24089.
- Zhang H, Joseph J, Crow J and Kalyanaraman B (2004) Mass spectral evidence for carbonate-anion-radical-induced posttranslational modification of tryptophan to kynurenine in human Cu, Zn superoxide dismutase. *Free Radical Biology & Medicine* 37, 2018–2026.
- Zhang K, Zhang Y, Zi J, Xue X and Wan Y (2017) Production of human Cu, Zn SOD with higher activity and lower toxicity in *E. coli* via mutation of free cysteine residues. *Biomed Research International* 2017, 4817376.
- Zhu C, Beck MV, Griffith JD, Deshmukh M and Dokholyan NV (2018) Large SOD1 aggregates, unlike trimeric SOD1, do not impact cell viability in a model of amyotrophic lateral sclerosis. *Proceedings of the National Academy of Sciences of the USA* 115, 4661–4665.
- Zil'ber LA, Bajdakova ZL, Gardaš'jan AN, Kononov NV, Bunina TL and Barabadze EM (1963) Study of the etiology of amyotrophic lateral sclerosis. *Bulletin of the World Health Organization* 29, 449–456.
- Zinman L, Liu HN, Sato C, Wakutani Y, Marvelle AF, Moreno D, Morrison KE, Mohlke KL, Bilbao J, Robertson J and Rogava E (2009) A mechanism for low penetrance in an ALS family with a novel SOD1 deletion. *Neurology* 72, 1153–1159.
- Zu JS, Deng HX, Lo TP, Mitsumoto H, Ahmed MS, Hung WY, Cai ZJ, Tainer JA and Siddique T (1997) Exon 5 encoded domain is not required for the toxic function of mutant SOD1 but essential for the dismutase activity: identification and characterization of two new SOD1 mutations associated with familial amyotrophic lateral sclerosis. *Neurogenetics* 1, 65–71.