Oxidation resistance 1 regulates post-translational modifications of peroxiredoxin 2 in the cerebellum

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Abstract

Protein aggregation, oxidative and nitrosative stress are etiological factors common to all major neurodegenerative disorders. Therefore, identifying proteins that function at the crossroads of these essential pathways may provide novel targets for therapy. Oxidation resistance 1 (Oxr1) is a protein proven to be neuroprotective against oxidative stress, although the molecular mechanisms involved remain unclear. Here, we demonstrate that Oxr1 interacts with the multifunctional protein, peroxiredoxin 2 (Prdx2), a potent antioxidant enzyme highly expressed in the brain that can also act as a molecular chaperone. Using a combination of in vitro assays and two animal models, we discovered that expression levels of Oxr1 regulate the degree of oligomerization of Prdx2 and also its post-translational modifications (PTMs), specifically suggesting that Oxr1 acts as a functional switch between the antioxidant and chaperone functions of Prdx2. Furthermore, we showed in the Oxr1 knockout mouse that Prdx2 is aberrantly modified by overoxidation and S-nitrosylation in the cerebellum at the pre-symptomatic stage; this in-turn affected the oligomerization of Prdx2, potentially impeding its normal functions and contributing to the specific cerebellar neurodegeneration in this mouse model.

1. Introduction

Neurodegenerative disorders are becoming more prevalent with increased global life expectancy and a better understanding of the underlying disease mechanisms will be essential for designing new therapeutic approaches to tackle this growing health burden [1–3]. Aberrant proteostasis and increased oxidative and nitrosative (O/N) stress have been strongly linked with the neurodegenerative process, irrespective of the etiology [4,5]. O/N stress arises from an excessive accumulation of reactive oxygen species (ROS) and reactive nitrogen species (RNS) resulting from an imbalance between free radical production and detoxification by antioxidant proteins [6]. Elevated levels of O/N stress markers such as oxidised lipids and nitrated proteins have been reported in post mortem tissue of individuals affected by major neurodegenerative disorders such as Alzheimer’s disease (AD), Parkinson’s disease (PD), and amyotrophic lateral sclerosis (ALS) [7–9]. Importantly, these markers are not only observed at the end-stage of the disease but are also detected in peripheral blood of patients during the course of the disease and pre-symptomatically in animal models of these conditions [10–14]. For example, mouse models of ALS show elevated levels of carbonylated proteins, a marker of ROS-induced protein damage, before any motor neuron pathology or clinical features are observed, suggesting that O/N stress is an early event of the neurodegenerative process [15].

Protein aggregation is another important feature consistently observed in neurodegenerative disease. Indeed, the formation of...
neurotoxic protein aggregates has been reported in brain and spinal cord regions affected by AD, PD, and ALS [16]. In normal conditions, chaperone proteins are employed to fold misfolded proteins and disassemble protein aggregates, and thus play a key role in the global cellular proteostasis [17]. In neurodegenerative disease, however, the chaperone activity in cells is disrupted. For instance, mutations in the gene encoding for the mitochondrial chaperone mtHsp70 have been identified in patients with PD, and have been shown to disrupt mtHsp70 activity and to lead to apoptosis [18,19]. In addition, PTMs, such as S-nitrosylation and sulfonylation, have been associated with dysregulation of protein function. For instance, aberrant S-nitrosylation of the chaperone protein disulfide-isomerase (PDI) inhibits its activity, leading to neurodegeneration [20,21]. Therefore, identifying proteins that function in both the O/N stress and proteostasis pathways may produce valuable insights into the underlying pathogenic mechanisms of neurodegenerative conditions.

Oxidation resistance 1 (OXR1) was originally identified as a gene capable of rescuing bacterial oxidative DNA damage [22]. Subsequently, Oxr1 has been shown to act as a potent regulator of oxidative stress in mammalian neurons that can also delay neurodegeneration in vivo. Oxr1 possesses a highly conserved C-terminal TLDc domain shared with four other proteins in mammals, several of which have been proven to be important for brain function [23]. Mutations in the TLDc domain of TBC1 domain family member 24 (TBC1D24) influence its role in neuronal cell development and function and are associated with a range of familial neurological disorders, including epilepsy and deafness, onychodystrophy, osteodystrophy and mental retardation with seizures (DOORS) syndrome [23–25]. Moreover, mice lacking all Oxr1 isoforms present a progressive ataxic phenotype concomitant with degeneration of granule neurons in the cerebellum from postnatal day 19 (P19) [26–28]. Over-expression of Oxr1 in vivo is able to delay motor neuron degeneration and motor dysfunction in a mouse model of ALS [29]. In vitro studies in mammalian cells also demonstrated that levels of Oxr1 can modulate the sensitivity of neurons to oxidative stress [26,28]: knock-down of Oxr1 leads to oxidative stress-induced cell death while over-expression of Oxr1 reduces ROS levels and apoptosis of cells subjected to oxidative stress [26,28]. Furthermore, studies in a range of experimental systems have demonstrated a dysregulation of the expression of a number of antioxidant genes when Oxr1 is disrupted. For example, glutathione peroxidase 2 (GPX2), heme oxygenase-1 (HO-1), catalase (CAT) and peroxiredoxin 4 are reportedly induced in patients with PD, and have been shown to disrupt mtHsp70, thus taken together, these data suggest that the TLDc proteins have the ability to reduce O/N stress. For cell death assay, cells were treated with 500 µM H2O2 for 1 h. Fluorescent microscope Axioplan2 Imaging (Carl Zeiss) was used for cell death assay to visualise the pyknotic nuclei. For S-nitrosocysteine (SNOC) treatment of SH-SY5Y cells, a fresh stock solution of SNOC was prepared immediately before each use by mixing 200 mM cysteine (Sigma) with 200 mM NaNO2 (Sigma) supplemented with 200 mM β-mercaptoethanol (Sigma) supplemented with HCl (0.5 N). Cells were treated with 200 µM SNOC for 30 min. Control cells were treated with a vehicle solution (HCl diluted in water).

2. Materials and methods

2.1. Reagents

Reagents used were as follows: Insulin (I0516, Sigma), citrate synthase (C2620, Sigma), protease inhibitor (5871S, Cell Signaling), protease/phosphatase inhibitor (5872S, Cell Signaling), protease inhibitor (11836170001, Roche). Antibodies used were: Prdx2 (10545–2-AP, Proteintech), Prdx-So2/3 (ab16830, Abcam), β-actin (ab8226, Abcam), HA (H6908, Sigma), α-tubulin (Sigma), MYC (Sigma), Oxr1 [28].

2.2. Animals

Constitutive Oxr1 knockout (Oxr11/1) mice maintained on a C57BL6/J background have been previously described [27]. Oxr1 Tg mice, where a full-length cDNA transgene of Oxr1-FL is expressed from a ubiquitous actin-derived promoter, were also maintained on a C57BL6/J background and have been previously described [28]. All experiments were conducted in adherence with the guidelines set forth by the UK Home Office regulations, and with the approval of the University of Oxford Ethical Review Panel.

2.3. Expression constructs

Mouse Prdx2 cDNA (NM.001317385.1) was cloned from total brain cDNA extract into pET-22b vector with or without introducing thrombin cleavage site (Thr) in frame with a His6 tag: F′– 5′-GAGACAGTATGGCCCTCGGCAACGCGCAAATC-3′, R′– 5′-GAGACCTGGAGATGATGATGATGATGAT-3′; for Oxr1, 5′–GAGACTCGAGGTGTTTGGAGAAGTATTCC-3′, respectively. Mouse Oxr1-C (NM_001130164) was cloned from total brain cDNA into pET-22b backbone in frame with His6 tag (F′– 5′-GAGACATATGCTCCGGTCTTATTTT-3′, R′– 5′-GAGACTCGAGGTGTTTGGAGAAGTATTCC-3′, respectively. Mouse Oxr1-C was used as an internal control for transfection experiments.

2.4. Cell culture, transfection, and treatment

Neuro2a (N2a) and SH-SY5Y cells were cultured in DMEM with GlutaMAX (Gibco) and Ham’s F12:DMEM (Gibco) respectively, supplemented with 10% fetal bovine serum (Gibco) and 1% penicillin-streptomycin solution (Gibco) in an incubator at 37 °C in 5% CO2 atmosphere. Cells were seeded 24 h prior to transfection. Cells were transfected using Fugene 6 (Promega) diluted in Opti-MEM (Gibco) according to manufacturer’s protocol. 24 h after transfection culture media was changed to media supplemented with desired concentrations of H2O2. For cell death assay, cells were treated with 500 µM H2O2 for 5 h, and for Prdx2 overoxidation experiments, cells were treated with 300 µM H2O2 for 1 h. Fluorescent microscope Axiosplan2 Imaging (Carl Zeiss) was used for cell death assay to visualise the pyknotic nuclei. For S-nitrosocysteine (SNOC) treatment of SH-SY5Y cells, a fresh stock solution of SNOC was prepared immediately before each use by mixing 200 mM cysteine (Sigma) with 200 mM NaNO2 (Sigma) supplemented with HCl (0.5 N). Cells were treated with 200 µM SNOC for 30 min. Control cells were treated with a vehicle solution (HCl diluted in water).

2.5. Immunoprecipitation and western blotting

SDS-PAGE: Tissue samples or cells were lysed in cold RIPA buffer supplemented with protease inhibitor (Cell signaling) and 1% Triton X-100 or 1% CHAPS in a tissue Precellys homogenizer (Bertin Corp.) (for tissue) or by trituration with a syringe (for cells). Protein concentration was assessed by bichoninic acid (BCA) assay (Novagen). For co-


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immunoprecipitation, protein extracts were incubated overnight at 4 °C with either Prdx2 antibody, EZView MYC beads (Sigma) or G protein sepharose beads (Sigma) only, as a control. Protein-antibody complexes were pulled down using G protein sepharose beads (Sigma) and washed with cold RIPA buffer. Immunoprecipitated proteins were incubated with Laemmli loading buffer (Bio-Rad) or NuPAGE loading buffer (Life Technologies) supplemented with 5% β-mercaptoethanol (Sigma) at 100 °C for 5 min.

Blue Native-PAGE (BN-PAGE)- Brain and cerebella tissue samples and cells were homogenised in cold PBS buffer supplemented with protease/phosphatase inhibitors (New England Biolabs) using Precellys homogenizer (only for tissue samples) and lysed by repeated freezing and thawing (for tissue and cells). After measuring protein concentration using BCA assay (Novagen), samples were mixed with 5x loading buffer (62.5 mM Tris-HCl pH 6.8, 40% glycerol, 0.01% bromophenol blue) and were resolved on 10% Bis-Tris native gel.

After samples were run on either SDS-PAGE or BN-PAGE, proteins were transferred to PVDF membrane (Amersham). The membrane was then blocked with 5% skim milk in PBST at 1 h at room temperature and incubated overnight with primary antibodies at 4 °C. The membranes were washed with PBST and incubated with secondary HRP-conjugated antibodies and signal was detected using ECL or ECL Prime Western Blotting detection kit (both Amersham) using an ImageQuant LAS 4000 (GE Healthcare). Western blot signal was quantified using ImageJ.

2.6. Protein expression and purification

Prdx2-His6 and Prdx2-Thr-His6 proteins were produced in E.coli strain BL21(DE3) (Invitrogen) and purified according to protocol published by Yan et al. [37]. Briefly, protein expression was induced at OD600 = 0.7–0.8 by addition of isopropyl β-D-1-thiogalactopyranoside (IPTG) (Sigma) to final concentration of 0.8 mM. After incubation on a shaker for 5 h at 25 °C, bacterial cells were collected by centrifugation and resuspended in 500 mM NaCl PBS supplemented with protease inhibitors. After sonication of the bacterial cultures and centrifugation for 30 min at 4 °C, recombinant His6-tagged protein was incubated with 1 ml TALON metal affinity resin (Clontech) for 1 h at 4 °C. Proteins were washed using increasing concentration of imidazole, and eluted with 500 mM imidazole. Eluted samples were dialysed overnight against PBS buffer at 4 °C using Slide-A-Lyzer dialysis cassette (Thermo Fisher). The His6 tag was cleaved from Prdx2-Thr-His6 using the Thrombin Clean-Cleave Kit (Sigma Aldrich) at 17 °C for 4 h, as per the manufacturer’s protocol and dialysed against PBS buffer following cleavage. Oxr1-C was produced as described before [28]. Briefly, protein expression was induced at OD600 ~ 0.8 by addition of IPTG to a final concentration of 0.8 mM. Bacterial cultures were incubated on a shaker for 16 h at 18 °C and bacterial cells were collected by centrifugation. Protein was purified as described above. Full-length human OXR1 recombinant protein expressed in E.coli was produced from Origene (PTP60961). Purified proteins were analysed by western blotting with anti-Prdx2 and anti-Oxr1 antibodies. Protein concentrations were measured using NanoDrop spectrophotometer with extinction coefficients of 21,555 M⁻¹ cm⁻¹ for Prdx2 and 38,055 M⁻¹ cm⁻¹ for Oxr1-C and. DNA fragments encoding TBC1D24 full-length (aa 1–553) and TLDc domain (aa 336–553) were subcloned into the pEF-Bicistronic vector for insect S9 cell expression and pNIC28-Bsa4 for E. coli expression, respectively. Recombinant full-length TBC1D24 was expressed in S9 cells, while the TBC1D24 TLDc-domain was expressed in E.coli and purified as above, followed by size exclusion chromatography. The purity of the recombinant protein preparation was ascertained by coomassie staining (Supplementary Fig. 1).

2.7. Peroxiredoxin activity assay

Prdx2 activity was assessed using ferrous oxidation-xylene orange (FOX) assay [38]. The assay solution contained 100 µM dithiothreitol (DTT, Sigma Aldrich), 50 µM H2O2 (Sigma Aldrich), 2.2 µg recombinant uncleaved purified Prdx2, with or without 0.95 µg Oxr1-C or 4.5 µg OXR1-FL. The reaction was initiated by addition of H2O2 and quenched at appropriate time points by addition of 200 µl of FOX reagent (1 part of FOX A reagent (25 mM ammonium ferrous sulfate (Sigma) in 2.5 M H2SO4 (Sigma)) and 100 parts of FOX B reagent (100 mM sorbitol (Sigma) with 125 µM xylene orange (Sigma) in water)). Samples were incubated at room temperature for at least 30 min before reading absorbance at 560 nm with Fluorostar Omega plate reader.

2.8. Chaperone activity assay

In vitro holdase activity of cleaved Prdx2 and Oxr1-C was assessed in chemically- and thermally-induced aggregation assays. The ability of Prdx2 and Oxr1-C to protect insulin from DTT-induced aggregation was measured as change of absorbance at 650 nm at 25 °C. The reaction solution contained 30 µg insulin (Sigma Aldrich) with or without 20 µg Prdx2, with 22 µg (6 µM final concentration), 33 µg (9 µM) Oxr1-C, 22 µg Tbc1d24, 22 µg Tbc1d24 TLDc domain or 85 µg (6 µM) OXR1-FL and was initiated by addition of DTT to a final concentration of 6.66 mM. Ability of Prdx2 and Oxr1-C to protect citrate synthase (CS) from aggregating was monitored by changes in absorbance at 360 nm during incubation at 44 °C. Reaction solution contained 10 µg of CS and 10 µg of Prdx2, Oxr1-C or both in 50 mM HEPES pH 8.0.

2.9. Biotin switch assay

The biotin switch assay was performed as previously described using the reagents from the S-nitrosylated protein detection kit (Cayman) [39]. For tissue, snap-frozen brains or cerebella were homogenised in buffer A supplemented with protease inhibitors (Cell Signaling) using a tissue Precellys homogenizer (Bertin Corp.). After centrifugation at maximum speed for 30 min at 4 °C, protein concentration was quantified using a BCA assay (Thermo Scientific) and an equal amount of protein was used per sample. All subsequent steps were performed protected from direct light and in amber tubes. Blocking reagent provided in the kit was added, and samples were incubated for 30 min at 50 °C with regular vortexing. Proteins were subsequently precipitated using cold acetone to remove any trace of blocking reagent, and incubated at —20 °C for at least 30 min, followed by centrifugation at 3000g for 10 min at 4 °C. Protein pellets were resuspended in the labelling/reducing reagent provided and prepared as per the manufacturer’s instructions. A combined protein sample was incubated with diluted dimethylformamide only as a negative control. Samples were incubated for 1 h at room temperature on a shaker. Proteins were acetone precipitated as above. Protein pellets were resuspended in wash buffer provided; a fraction of the protein extracts was kept aside as input to determine total Prdx2 levels. To pull down S-nitrosylated proteins, protein extracts were incubated overnight at 4 °C on a shaker with 50 µl EZview streptavidin beads (Sigma). Beads were washed three times with wash buffer and resuspended in NuPAGE loading buffer supplemented with β-mercaptoethanol, and boiled for 5 min. Total protein (input) and S-nitrosylated proteins were run on pre-cast NuPAGE gels as per standard procedures. Results were expressed as intensity of signal for SNO-Prdx2 over total level of Prdx2 per sample.

For cells, cells transfected for 48 h and treated with either vehicle or SNOC were washed with PBS and scraped from the dish. After centrifugation at 1000 rpm for 5 min, cell pellets were resuspended directly in blocking reagent diluted in buffer A and incubated at 50 °C for 30 min. After centrifugation and subsequent acetone precipitation, protein pellets were resuspended in labelling/reducing reagent and incubated at room temperature for 1 h. After acetone precipitation, proteins were resuspended in wash buffer and quantified by BCA assay (Thermo Scientific). The subsequent steps are identical as for tissue above.
2.10. Organomercury resin capture (ORC)

ORC from brain and cerebellum tissue was performed as previously described [40]. Briefly, samples were homogenised in homogenization buffer (250 mM HEPES free acid, 1 mM DTPA, 0.1 mM neocuproine, 1% Triton X-100, pH 7.7) using a tissue Precellys homogenizer (Bertin Corp.). Samples were clarified by centrifugation at full speed for 10 min at 4 °C and protein concentration was quantified by BCA assay (Thermo Scientific). An equal amount of protein was used per sample. As a negative control a combined protein sample was incubated with the reducing agent DTT (10 mM) for 15 min at room temperature. Proteins were acetone precipitated and resuspended in blocking buffer (250 mM HEPES acid free, 1 mM DTPA, 0.1 mM neocuproine, 2.5% SDS, pH 7.7) supplemented with 50 mM blocking agent MMTS. Protein extracts were incubated for 30 min at 50 °C, with frequent vortexing, followed by acetone precipitation. Protein pellets were resuspended in loading
buffer (250 mM MES, 1 mM DTPA, 1% SDS, pH 6.0) and ready to be loaded on to the activated resin in glass columns; a fraction was kept aside as input. Organomercury resin (produced by the laboratory of Professor Harry Ischiropoulos University of Pennsylvania, USA and provided by Dr Mark Crabtree, University of Oxford) was washed and activated by consecutive washes with isopropanol, water, 0.1 M sodium bicarbonate pH 8.8, and equilibration buffer (50 mM NaCl, 50 mM MES, 1 mM DTPA, pH 6.0). After the resin was drained, samples were added and capped with 0.5 ml loading buffer and incubated at room temperature for 1 h protected from light to allow the reaction between S-nitrosylated proteins and the organomercury resin. The S-nitrosylated proteins cross-linked to the resin were then washed with consecutive 50 bed volumes of buffer A (50 mM Tris-HCl, 0.3 mM NaCl, 0.5% SDS, pH 7.5), buffer B (50 mM Tris-HCl, 0.3 mM NaCl, 0.05% SDS, pH 7.5), buffer C (50 mM Tris-HCl, 0.3 mM NaCl, 1% Triton X-100, 1 M urea, pH 7.5), and buffer D (50 mM Tris-HCl, 0.3 mM NaCl, 0.1% Triton X-100, 0.1 M urea, pH 7.5), followed by three washes of 200 bed volumes of water. Bound S-nitrosylated protein were then eluted by incubating the resin with 5% performic acid for 45 min at room temperature. The eluted proteins were precipitated by adding one volume of 100% TCA to four volumes of protein sample and incubated overnight at 4 °C. After centrifugation at full speed for 30 min at 4 °C, protein pellets were washed in cold acetone and resuspended in NuPAGE loading buffer supplemented with β-mercaptoethanol, and boiled for 5 min. Total protein (input) and S-nitrosylated proteins were run on pre-cast NuPAGE gels as per standard procedures. Results were expressed as intensity of signal for SNO-Prdx2 over total level of Prdx2 per sample.

2.11. Statistical analyses

Statistical analysis was performed using Prism software using one-way ANOVA or unpaired t-test. Data are presented as mean ± SEM.

3. Results

3.1. Oxr1 modulates the oligomerization state of the antioxidant enzyme Prdx2

It has been shown recently that Oxr1 regulates the activity and degree of oligomerization of the multifunctional enzyme glucose-6-phosphate isomerase (GPI/Gpi1) via protein-protein interaction [27]. Therefore, seeing that Oxr1 can influence ROS levels as well as protein multimer formation and aggregation, we tested the hypothesis that Oxr1 is neuroprotective against O/N stress by regulating the function of essential proteins within these two pathways [26,28,36,41]. As a promising candidate interactor for Oxr1, we selected peroxiredoxin 2 (Prdx2) because it acts both as a chaperone and an antioxidant as well as being highly expressed in neurons [42–44]. Indeed, the expression patterns of Prdx2 and Oxr1 overlap in the CNS, including the granule cell layer of the cerebellum [26,42,45]. Furthermore, similar to the knockout of Oxr1, loss of Prdx2 in mice leads to accumulation of ROS and oxidative DNA damage, although this occurs in the CA1 region of the hippocampus as opposed to the cerebellum of Oxr1 knockout mutants [46].

Given that the multiple functions of Prdx2 are regulated by its degree of oligomerization, we first tested whether Oxr1 levels would affect Prdx2 oligomerization in the brain [47,48]. Using non-reducing native PAGE to preserve protein complexes, we assessed Prdx2 oligomerization in tissues from a constitutive Oxr1 knockout mouse in which all isoforms of the protein are disrupted (Oxr1<sup>Δ/Δ</sup>) [26–28]. Tissue samples were collected at a pre-symptomatic timepoint (P18), prior to the onset of neurodegeneration in this model. In wild-type (Oxr1<sup>+/+</sup>) brain tissue, we observed a dimeric form of Prdx2 in addition to HMW complexes with an apparent molecular weight of approximately 400 kDa, as previously described (Fig. 1A) [48–50]. Interestingly, there was a significant decrease in these HMW Prdx2 complexes in the brain of Oxr1<sup>Δ/Δ</sup> mice compared to wild-type controls, both in the brain without the cerebellum (Fig. 1A) and in the cerebellum alone (Fig. 1B).

To confirm that Oxr1 levels regulate Prdx2 oligomerization, we next assessed the levels of Prdx2 multimers in brains from adult mice overexpressing an HA-tagged full-length Oxr1 (Oxr1-FL) cDNA transgene (Oxr1 Tg) [28]. Conversely to the observations from Oxr1<sup>Δ/Δ</sup> knockout mice, levels of the HMW Prdx2 complexes were significantly increased in Oxr1 Tg tissue compared to wild-type controls (Fig. 1C). Together, these data suggest that the levels of Oxr1 modulate Prdx2 oligomerization in vivo. To validate that higher levels of Oxr1 indeed favour higher-order Prdx2 complexes, we next analysed the degree of oligomerization of Prdx2 in cells over-expressing Oxr1 and upon treatment with H<sub>2</sub>O<sub>2</sub> to promote assembly of HMW Prdx2 complexes via Prdx2 overoxidation [48,50,51]. Interestingly, in cells over-expressing Oxr1-FL, specific HMW Prdx2 complexes were detected - namely its decameric species at the molecular weight of approximately 220 kDa - while this was not present in control vector-transfected cells (Fig. 1D). Overall, our data suggest that Oxr1 is able to modulate Prdx2 oligomerization.

3.2. Oxr1 interacts with the antioxidant enzyme Prdx2

To understand the mechanism by which Oxr1 regulates the oligomerization of Prdx2, we next tested whether Oxr1 and Prdx2 could interact. To assess binding between Prdx2 and Oxr1, we performed co-immunoprecipitation experiments in neuronal N2A cells co-expressing Prdx2 and either Oxr1-FL or the shortest Oxr1 isoform (Oxr1-C). The latter isoform primarily consists of the TLDc domain, but is highly expressed in the brain and has a proven function as a neuroprotective protein [26,28]. Both Oxr1 isoforms were found to co-immunoprecipitate with Prdx2 (Fig. 2A). The TLDc domain is common between both Oxr1 isoforms, thus we went on to investigate whether this interaction would be maintained with other TLDc-containing proteins. We discovered that Prdx2 also interacts with full-length nuclear receptor coactivator 7 (Ncoa7-FL), and to a lesser extent with the shorter TLDc domain-containing isoform (Ncoa7-B) (Fig. 2A). We also observed binding to Tbc1d24, suggesting that Prdx2 is a common binding partner for most of the TLDc proteins (Fig. 2A). To investigate whether the Oxr1 interaction occurs in vivo, we performed co-immunoprecipitation from wild-type mouse brain tissue and detected an interaction between endogenous Oxr1 and Prdx2, confirming our <i>in vitro</i> data (Fig. 2B).

3.3. Oxr1 and Prdx2 do not synergistically regulate H<sub>2</sub>O<sub>2</sub>-induced cell death

To investigate the significance of the Oxr1:Prdx2 interaction, we next tested systematically how Oxr1 and Prdx2 might influence each other's functional roles. Both Oxr1 and Prdx2 have been shown to be neuroprotective under oxidative stress; therefore, we quantified H<sub>2</sub>O<sub>2</sub>-induced cell death in a neuronal cell line expressing either Oxr1, Prdx2 or both proteins [26,52]. Consistent with previous reports, over-expression of Oxr1-FL, Oxr1-C or Prdx2 significantly reduced cell death under oxidative stress conditions; however, Prdx2 was significantly less protective than Oxr1 in this assay (Fig. 2C) [26,28,53]. Co-expression of Prdx2 together with Oxr1 did not alter significantly the neuroprotective properties of Oxr1.

Considering that Prdx2 can reduce H<sub>2</sub>O<sub>2</sub> at a high kinetic rate, we next investigated whether the neuroprotection against oxidative stress that is conferred by Oxr1 could be via regulation of Prdx2 activity [54,55]. To test this, we carried out a peroxide activity assay that measures reduction of H<sub>2</sub>O<sub>2</sub> using purified recombinant proteins, and as expected, recombinant Prdx2 was able to dramatically reduce H<sub>2</sub>O<sub>2</sub> levels (Fig. 2D) [37]. Recombinant Oxr1-C and Oxr1-F1 FL were not, however, able to reduce H<sub>2</sub>O<sub>2</sub> levels in this assay, similar to what has been observed previously (Fig. 2D) [28]. When Prdx2 and Oxr1-C were...
incubated together, no enhancement in the rate of H$_2$O$_2$ decay was observed as compared to the rate observed for Prdx2 alone (Fig. 2D), suggesting that Prdx2 and Oxr1 do not interact synergistically to regulate cell death under oxidative stress or to reduce H$_2$O$_2$ levels.

3.4. Oxr1 and Prdx2 both modulate protein aggregation in vitro

As well as being an antioxidant enzyme, Prdx2 also possesses holdase activity [48,56]. Holdases are a class of chaperones that bind unfolded or misfolded molecules and prevent them from aggregating in an ATP-independent manner [17]. To investigate any functional interactions between Oxr1 and Prdx2 that may modulate Prdx2 chaperone activity, we used a well-established in vitro aggregation assay with recombinant proteins. Insulin was induced to form large aggregates by incubation with DTT; its aggregation correlates with an increased turbidity of the reaction solution, which can be quantified by spectrophotometry [57]. Using this system, we observed a reduction in the aggregation of insulin in the presence of recombinant Prdx2, as expected, while lysozyme, a non-chaperone control, did not affect aggregation (Fig. 3A-C). Importantly, when Prdx2 was incubated with recombinant Oxr1-C, this led to a further significant reduction in insulin aggregation rate and in the aggregation at endpoint as compared to Prdx2 alone; this suggests that Oxr1-C may enhance the chaperone activity of Prdx2 (Fig. 3A-C). We confirmed these data by using a
second aggregation assay that utilizes thermally-induced aggregation of citrate synthase (CS). In agreement with the insulin aggregation data, the presence of Prdx2 reduced the thermally-induced aggregation of CS, and the rate of aggregation was significantly reduced further when Prdx2 was co-incubated with Oxr1-C (Fig. 3D-F).

Surprisingly, when these assays were performed with Oxr1-C only, we detected a significant reduction of approximately 50% in both the chemically- and thermally-induced aggregation rate of insulin and CS (Fig. 3D-I). Given that these assays are cell-free, these data are not reliant on other proteins, thus suggesting that Oxr1-C possesses a potential holdase activity. To confirm this observation, we tested whether increasing Oxr1-C concentration would lead to an increase in protective activity against chemically-induced protein aggregation. We did observe a significant decrease in substrate aggregation rate as well as a slight reduction in the aggregation at end point with an increased concentration of Oxr1-C, further supporting a chaperone activity for Oxr1-C (Fig. 3G-I).

As the Oxr1-C isoform is mainly comprised of the TLDc domain, our data indicate that the chaperone activity we observe may be conserved in other TLDc-containing proteins. To test this hypothesis, we quantified the chaperone activity of full-length OXR1 (OXR-FL) and TBC1D24 using the chemically-induced protein aggregation assay. Using a recombinant OXR1-FL, we detected a significant reduction in both the aggregation rate and aggregation at end point compared to the control, suggesting that OXR1-FL also possesses holdase activity (Fig. 3G-I). Interestingly, we did not detect any reduction in the formation of insulin aggregates when incubated with full-length TBC1D24, suggesting that TBC1D24 does not possess holdase activity (Fig. 3G-I). We performed a similar experiment using a recombinant protein constituted of only the C-terminal TLDc domain of TBC1D24. Similar to full-length TBC1D24, the TLDc domain alone did not show any chaperone activity (Fig. 3G-I). Therefore, we have identified a novel activity for Oxr1 as a...
potential holdase that is likely not conserved with the related TLDC protein TBC1D24.

3.5. Oxr1 modulates Prdx2 post-translational modifications (PTMs)

Prdx2 function is regulated by its level of oligomerization and through PTMs such as overoxidation and S-nitrosylation, in addition to conformational changes [48,58-60]. In particular, reversible oxidation of catalytic cysteine residues occurs during the catalytic cycle of Prdx2: in the presence of low levels of H₂O₂, Prdx2 is oxidised and reactivated, while under high levels of H₂O₂, its cysteine residues are overoxidized, forming sulfenic and sulfonic acid (Prdx-SO₂⁻⁻, SO₃⁻⁻), which inactivates its antioxidant activity but stabilizes the HMW complexes which drive its chaperone activity [48,51,61]. Given our findings that Oxr1 levels modulate Prdx2 oligomerization, we next investigated whether Oxr1 could affect Prdx2 overoxidation. We first assessed overoxidation of endogenous Prdx2 in vitro by quantifying the levels of Prdx2-SO₂⁻⁻ over the total level of Prdx2 from H₂O₂-treated cells. Using an antibody that recognizes overoxidized Prdx1, Prdx2, Prdx3 and Prdx4, we confirmed previously reported overoxidation of the aforementioned Prdxs induced by H₂O₂ treatment [62]. Interestingly, under these conditions, levels of Prdx2 overoxidation were significantly higher in cells overexpressing Oxr1-FL (Fig. 4A) and a similar effect was observed in cells over-expressing Oxr1-C (Fig. 4B). This suggests that Oxr1 modulates, directly or indirectly, Prdx2 overoxidation. We next tested if levels of Oxr1 would similarly affect overoxidation of Prdx2 in vivo by using the cerebellum and the remaining brain tissue from Oxr1 Tg mice or control littermates. Similar to what we observed in vitro in Oxr1 over-expressing cells, a significant increase in Prdx2 overoxidation levels in brain of transgenic animals were detectable as compared to wild-type mice (Fig. 4C). There was also a significant increase in Prdx2-SO₂⁻⁻ levels in cerebellar tissue from Oxr1 Tg mice (Fig. 4D). To further investigate the role of Oxr1 in regulating Prdx2 overoxidation, we analysed the levels of Prdx-SO₂⁻⁻ in brains from Oxr1 mutant mice as compared to controls at the presymptomatic P18 timepoint. We did not observe any significant changes in Prdx2 overoxidation in the brain (Fig. 4E); however, levels of Prdx2-SO₂⁻⁻ were significantly reduced in cerebella from Oxr1 mutant animals as compared to controls (Fig. 4F). These data suggest that Oxr1 regulates levels of Prdx2 overoxidation; interestingly, this occurs differentially in the cerebellum – the site of neurodegeneration in the Oxr1 mutant – as compared to the rest of the brain.

S-nitrosylation is another essential PTM regulating Prdx2 function. Indeed, addition of an NO group at its cysteine residues has been shown to reduce the antioxidant activity of Prdx2 [58]. To test whether Oxr1 would have an effect on Prdx2 S-nitrosylation in vitro, we used a neuronal cell line SH-SY5Y treated with the physiological cell-permeable NO-donor, S-nitroscysteine (SNOC), to induce S-nitrosylation of proteins. Using a biotin switch assay, we quantified levels of S-nitrosylated endogenous Prdx2 (SNO-Prdx2) and found that control cells treated with a NO-donor, S-nitrosocysteine (SNOC), which inactivates its antioxidant activity but stabilizes the HMW complexes which drive its chaperone activity [48,51,61]. Given our findings that Oxr1 levels modulate Prdx2 oligomerization, we next investigated whether Oxr1 could affect Prdx2 overoxidation. We first assessed overoxidation of endogenous Prdx2 in vitro by quantifying the levels of Prdx2-SO₂⁻⁻ over the total level of Prdx2 from H₂O₂-treated cells. Using an antibody that recognizes overoxidized Prdx1, Prdx2, Prdx3 and Prdx4, we confirmed previously reported overoxidation of the aforementioned Prdxs induced by H₂O₂ treatment [62]. Interestingly, under these conditions, levels of Prdx2 overoxidation were significantly higher in cells overexpressing Oxr1-FL (Fig. 4A) and a similar effect was observed in cells over-expressing Oxr1-C (Fig. 4B). This suggests that Oxr1 modulates, directly or indirectly, Prdx2 overoxidation. We next tested if levels of Oxr1 would similarly affect overoxidation of Prdx2 in vivo by using the cerebellum and the remaining brain tissue from Oxr1 Tg mice or control littermates. Similar to what we observed in vitro in Oxr1 over-expressing cells, a significant increase in Prdx2 overoxidation levels in brain of transgenic animals were detectable as compared to wild-type mice (Fig. 4C). There was also a significant increase in Prdx2-SO₂⁻⁻ levels in cerebellar tissue from Oxr1 Tg mice (Fig. 4D). To further investigate the role of Oxr1 in regulating Prdx2 overoxidation, we analysed the levels of Prdx-SO₂⁻⁻ in brains from Oxr1 mutant mice as compared to controls at the presymptomatic P18 timepoint. We did not observe any significant changes in Prdx2 overoxidation in the brain (Fig. 4E); however, levels of Prdx2-SO₂⁻⁻ were significantly reduced in cerebella from Oxr1 mutant animals as compared to controls (Fig. 4F). These data suggest that Oxr1 regulates levels of Prdx2 overoxidation; interestingly, this occurs differentially in the cerebellum – the site of neurodegeneration in the Oxr1 mutant – as compared to the rest of the brain.

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4. Discussion

Our current study sheds new light on the molecular mode of action of Oxr1 as a functional switch for the essential antioxidant and chaperone Prdx2. Indeed, we show that Oxr1 modulates oligomerization of HMW Prdx2 chaperone complexes and key PTMs regulating the dual functions of Prdx2.

Our study is in accordance with our previous observations that Oxr1 reduces aggregation of the mutated ALS-associated TDP-43 and FUS proteins in cellular models of ALS; this process relies on the interaction between Oxr1 and TDP-43 or FUS, but is independent of the proteasome or autophagy [36]. Based on our current findings, we hypothesise that Oxr1 may reduce protein aggregation first through its chaperone activity and second by favoring the formation of HMW Prdx2 chaperone complexes. Another key function described for Oxr1 is its ability to reduce levels of ROS and oxidative stress markers in oxidative stress-treated neurons as well as in SOD1G93A ALS mouse model [26,29]. However, Oxr1 has no direct catalytic activity, thus it has remained unclear how Oxr1 could carry out this role. Here, we showed that Oxr1 regulates the activity of the antioxidant protein Prdx2, in particular, by modulating two key PTMs of Prdx2, its overoxidation and S-nitrosylation: loss of Oxr1 increases SNO-Prdx2 and reduces its overoxidation, while increasing Oxr1 levels leads to the opposite effect. This is significant because these PTMs regulate Prdx2 antioxidant activity and are intertwined: S-nitrosylation of Prdx2 reduces its H₂O₂-induced over-oxidation and inhibits its antioxidant activity [58]. By affecting both the degree of oligomerization and the PTMs of Prdx2, Oxr1 could potentially fine-tune the amount of peroxidases versus chaperones in cells. A slight shift in this proportion could have a major impact on the global cell function given that Prdxs represent 0.2-0.8% of the total soluble protein fraction in cells and tissues, and that Prdx2 is highly expressed in neurons [42-44,64]. Therefore, Oxr1 may indirectly regulate the levels of oxidative stress in cells by modulating the PTMs of Prdx2 and subsequently fine-tuning its antioxidant activity. This newly described function is likely to run in concert with the proposed role for Oxr1 as a regulator of antioxidant gene expression, this latter regulatory hypothesis being supported by a number of studies that have combined OXR1 knock-down with genome-wide or targeted transcriptomics [30,33]. Further work is required to understand the molecular mechanisms involved, although there is evidence that gene modulation via p53-associated pathways is more plausible than Oxr1 acting upstream of the antioxidant Nrf2 signaling cascade [30,31].

The current study also provides an insight into the link between loss of Oxr1 and cellular degeneration in the cerebellum of the Oxr1 mutant mouse [28]. We observed higher levels of SNO-Prdx2 and reduced levels of Prdx2-SO₂⁻⁻ in the neurodegenerative Oxr1 knockout mouse at a presymptomatic stage, specifically in the cerebellum - the only brain region affected in this mutant - suggesting that aberrant PTM of Prdx2 may participate in the neurodegeneration observed in this mouse. Importantly, SNO-Prdx2 breaks the normal redox cycle that regenerates Prdx2, inhibiting Prdx2 antioxidant activity and leading to high levels of oxidative stress in cells [58]. Increased levels of SNO-Prdx2 have been observed in the brain of the P/Q/MB-exposed PD mouse model, as well as in dopaminergic neurons differentiated from induced pluripotent stem cells from PD patients and in post mortem brains of PD.
patients, suggesting that SNO-Prdx2 participates in the neurodegenerative process [58,65]. It was also shown in vitro that S-nitrosylation of Prdx2 led to a reduction of Prdx2-SO2/3 and of its neuroprotective function against H2O2-induced apoptosis [58]. Therefore, our current model is that loss of Oxr1 in the knockout mouse model impairs Prdx2 chaperone activity through reduction of HMW Prdx2 chaperone complexes and reduction of the complexes-stabilising PTM, Prdx2-SO2/3. In addition, Prdx2 antioxidant activity is also inactivated by the increase in S-nitrosylation of Prdx2. These changes in PTMs being cerebellum-specific, this may lead to a detrimental increase in protein aggregation and oxidative stress in the cerebellum, precipitating the degeneration of the cerebellar granule cells in Oxr1 knockout animals. Of note, we focused here on overoxidation and S-nitrosylation of Prdx2; however, Prdx2 function can be affected by other PTMs, such as phosphorylation and nitration, which have also been involved in neurodegeneration [66].

Using two in vitro assay systems, we have discovered that Oxr1-C and OXR1-FL possess a potential holdase activity. Because Oxr1-C is almost exclusively constituted of the TLDc domain, we tested whether this function was shared with another TLDc-containing protein. Surprisingly, however, TBC1D24 did not possess any holdase activity; neither the full-length protein nor the TLDc domain alone. This interesting finding indicates that, although all TLDc-containing proteins are neuroprotective, they may also carry out their own specific functions [23]. This may be driven in part by key residues in the TLDc domain that are yet to be identified. Importantly, we also showed that Tbc1d24 interacts with Prdx2, suggesting that the holdase activity is independent of the ability to bind Prdx2. Here, we have focused on the interplay between Oxr1 and Prdx2; yet the fact that Prdx2 has the ability to bind other TLDc-containing proteins suggests that functional compensation and competition with other TLDc proteins may play a part, and would require further investigation.

5. Conclusions

In summary, our study not only discovered a new function for Oxr1 as a potential holdase, it also identified a novel mode of action for Oxr1, acting on the oxidative stress response and protein aggregation by modulating the oligomerization and PTMs of the fundamental antioxidant enzyme Prdx2. Our study also places Oxr1 at the intersection...
between two key pathways involved in the neurodegenerative process.

Acknowledgments

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Fig. 5. Oxr1 regulates S-nitrosylation of Prdx2 in vitro and in vivo in the cerebellum. (A) SH-SY5Y cells were treated for 30 min with 200 μM SNOC or vehicle (veh) and processed for the biotin switch assay to quantify the levels of S-nitrosylated endogenous Prdx2 (SNO-PRDX2) by western blot in the presence of the constructs as indicated. (N = 3 independent repeats). (B-C) Representative western blots of SNO-PRDX2 levels in brain (B) or cerebellum (C) as determined by biotin switch from mice over-expressing Oxr1 (Oxr1 Tg) compared to wild-type controls (Oxr1+/+). (N = 3 animals per group). (D-E) SNO-PRDX2 levels in brain (D-F) or cerebellum (E-G) determined by biotin switch (D-E) or organomercury resin capture (ORC, F-G) from Oxr1 knockout (Oxr1−/−) mice compared to wild-type controls (N = 3 animals per group). In all panels, quantification is shown as the ratio of SNO-PRdx2 to total Prdx2 with α-tubulin used as a loading control. Data presented as mean ± SEM. Negative controls (−ve Ctrl) were combined proteins diluted in dimethylformamide without labelling/reducing reagent (for biotin switch) or proteins incubated with 10 mM DTT for 15 min (for ORC). Panel A: 1-way ANOVA (comparison of each construct between vehicle and SNOC-treated conditions). Panels B-G: t-test: **p < 0.05, ***p < 0.01, ****p < 0.001 as compared to empty-vector-transfected or to control mice, *p < 0.05 as compared to vehicle treated cells.
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