Dyclonine rescues frataxin deficiency in animal models and buccal cells of patients with Friedreich’s ataxia

Sunil Sahdeo1, Brian D. Scott1, Marissa Z. McMackin1, Mittal Jasoliya1, Brandon Brown2, Heike Wulff2, Susan L. Perlman3, Mark A. Pook4 and Gino A. Cortopassi1,*

1Department of Molecular Biosciences, School of Veterinary Medicine, 2Department of Pharmacology, School of Medicine, University of California, Davis, CA 95616, USA, 3Department of Neurology, University of California School of Medicine, Los Angeles, CA 90095, USA and 4School of Health Sciences and Social Care, Brunel University, Uxbridge UB8 3PH, UK

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Inherited deficiency in the mitochondrial protein frataxin (FXN) causes the rare disease Friedreich’s ataxia (FA), for which there is no successful treatment. We identified a redox deficiency in FA cells and used this to model the disease. We screened a 1600-compound library to identify existing drugs, which could be of therapeutic benefit. We identified the topical anesthetic dyclonine as protective. Dyclonine increased FXN transcript and FXN protein dose-dependently in FA cells and brains of animal models. Dyclonine also rescued FXN-dependent enzyme deficiencies in the iron–sulfur enzymes, aconitase and succinate dehydrogenase. Dyclonine induces the Nrf2 [nuclear factor (erythroid-derived 2)-like 2] transcription factor, which we show binds an upstream response element in the FXN locus. Additionally, dyclonine also inhibited the activity of histone methyltransferase G9a, known to methylate histone H3K9 to silence FA chromatin. Chronic dosing in a FA mouse model prevented a performance decline in balance beam studies. A human clinical proof-of-concept study was completed in eight FA patients dosed twice daily using a 1% dyclonine rinse for 1 week. Six of the eight patients showed an increase in buccal cell FXN levels, and fold induction was significantly correlated with disease severity. Dyclonine represents a novel therapeutic strategy that can potentially be repurposed for the treatment of FA.

INTRODUCTION

Friedreich’s ataxia (FA) is a severe neurodegenerative disease that is the most common autosomal recessive inherited movement disorder (1). There is no cure or effective treatment for FA. The disease causes degeneration and demyelination in dorsal root ganglion (DRG) neurons and spinocerebellar tracts, resulting in movement and speech disorders (2,3). Symptoms begin between the ages of 5–15 years, and patients are often wheelchair bound within 10–15 years of diagnosis. Early death is common and usually occurs from cardiac complications (4). FA is caused by a decrease in the mitochondrial protein frataxin (FXN gene), which has been shown to have roles in iron–sulfur cluster synthesis, iron transfer and antioxidant defense (1,5–8). The decrease in FXN stems from an accumulation of GAA triplet repeats in the first intron of the gene through inheritance (9–11). Here, we report on screening and identification of a potential therapeutic for FA. We previously identified antioxidant defects in DRG cells of an FA mouse model (12). Using this to develop a specific hypothesis for disease pathogenesis, we designed a cellular disease model for FA in order to identify a potential treatment. The cell model identified dyclonine, and we have shown that it induces FXN in cells, animal models and FA patients through a novel mechanism. Dyclonine is an oral anesthetic used to provide topical anesthesia to mucous membranes through sodium channel inhibition (13). It is the active ingredient in Sucrets, an over-the-counter throat lozenge, and has been in use for over 50 years (14). Additionally, dyclonine is used during dental examinations and procedures to numb mucous membranes of the mouth (15).

*To whom correspondence should be addressed. Tel: +1 5307549665; Fax: +1 5307549342; Email: gcortopassi@ucdavis.edu

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Figure 1. High-throughput screening reveals that dyclonine protects FA patient fibroblasts from diamide stress. (A) Effect of antioxidant inhibitors on 50B11 cell viability with FXN knockdown. Eleven inhibitors of thiol-related antioxidants were tested in an siRNA-mediated, FXN-deficient 50B11 DRG cell line [10 μM antimycin A, 1 μM auranofin, 100 μM BSO, 100 μM carmustine, 10 μM diamide, 0.1% diethyl maleate (DEM), 0.1% ethanol, 0.03% H2O2, 1 mM L-glutathione (L-GSH), 0.1% phenethyl isothiocyanate (PEITC), 100 μM dichloronitrobenzene (DCNB) and 1 mM N-methyl protoporphyrin (NMP)]. Cell viability was measured with Calcein-AM after 24 h and normalized to untreated control (n = 3). Increased sensitivity to cell death was induced by inhibitors of thiol-related antioxidants diamide and auranofin in FXN knockdown cells compared with AllStars non-targeting siRNA negative control. (B) FXN-dependent sensitivity to diamide is dose-dependent in 50B11 cells. Cell viability was measured with Calcein-AM after 24 h of treatment with 3–300 μM diamide and normalized to untreated control (n = 3). (C) Friedreich’s patient cells are sensitive to diamide. To confirm these effects in patient cells with low FXN, we tested 100 μM diamide in fibroblasts and
RESULTS

FXN-deficient cells are sensitive to the thioredoxin-oxidant diamide

There is ample support for dysregulated antioxidant defenses in FA (12,16–18). We screened that 12 inhibitors of thiol antioxidants were screened in FXN-deficient DRG neural cells, chosen since the DRGs are the primary site of disease pathogenesis (Fig. 1A and B). Measuring viable cells with Calcein-AM staining, the thioredoxin-oxidant diamide produced the greatest sensitivity (19). To confirm this phenotype in patient cells, we found that FXN-deficient FA fibroblasts and lymphoblasts were also clearly more sensitive to diamide treatment than healthy controls (Fig. 1C). Diamide is a specific thiol oxidant that causes oxidative damage and results in cell death (20,21). In the FA patient fibroblasts, a robust high-throughput screening assay based on diamide sensitivity was developed, with an average Z’-score of 0.7. We then screened a 1600-compound library of drugs that have been approved for use in human trials, and identified protective drugs (Fig. 1D). After the initial screen, multiple rounds of hit confirmation, and then concentration response curves, 33 drugs were found that reproducibly protected the patient cells from the thiol stress induced by diamide (Supplementary Material, Table S1). Of the drugs found to protect FA fibroblasts from diamide-induced oxidative stress, dyclonine was one of the most potent (Fig. 1E).

Dyclonine induces FXN in patient cells and in animal model tissues

Multiple mechanisms of action of protection from diamide were possible. One is reactivation of FXN expression, since its deficiency mediated the sensitivity to diamide. Thus, all 33 confirmed that screening hits including dyclonine were tested for their ability to induce FXN protein levels in the absence of diamide (Supplementary Material, Table S1). Of the compounds found to increase FXN in FA patient cells, we eliminated some because of safety concerns (i.e. cotinine and nifursol), and others because of their primary use is in veterinary medicine (i.e. oxendazole) (22–24). We chose to follow-up with dyclonine because of its potency of diamide protection, reproducibility of FXN induction and safe use in humans for decades. FA patient lymphoblasts treated with dyclonine show clear induction of FXN protein after 48 h of dyclonine exposure (Fig. 2A). Additionally, FXN mRNA transcript measured by RT-PCR is also dose-dependently increased after 24 h drug treatment in FA lymphoblasts (Fig. 2B). We observed induction of FXN in multiple animal models after intraperitoneal or oral dosing. FA-YG8 hFXN+/− with FXN (GAA)190 expansion; mFXN−/− transgenic mice (8) dosed for 1 week once daily with dyclonine i.p. or p.o. showed a clear induction of FXN protein in cerebellar lysate (Fig. 2C).

In addition, to explore the effects of dyclonine in an additional model with chronic dosing, we treated FA-PandKIKO mice [mFXN+/− with FXN (GAA)230 expansion; mFXN−/−] with 25 mg/kg dyclonine for 4 weeks p.o. This higher dose was chosen in an attempt to ascertain the maximum induction of FXN within the limits of dyclonine solubility. Thus, showing dyclonine can reverse the in vivo FXN protein defect in an additional FA model at a higher dose and chronic duration (Fig. 2D). These chronic studies resulted in no observable changes in sedation, feeding behavior or gross pathology. We observed these effects on FXN in multiple additional tissues, including heart and liver (Supplementary Material, Fig. S1).

Dyclonine activates the Nrf2 pathway

Of the 33 drugs that protected FA patient cells from diamide stress, some drugs were reported antioxidant response element (ARE)/nuclear factor (erythroid-derived 2)-like 2 (Nrf2) inducers in literature such as ebselen (25). Nrf2 is a transcription factor that responds to oxidative and thiol stress by binding AREs and driving the expression of multiple antioxidant and anti-inflammatory target genes including heme oxygenase 1 (HMOX1), NAD(P)H dehydrogenase (quinone) (NQO1) and glutathione peroxidase (GPX4) (26,27). We explored the activity of all 33 drugs as ARE/Nrf2 inducers for their ability to stimulate ARE-luciferase reporter system activity (Supplementary Material, Table S1), including dyclonine which dose-dependently drove induction of ARE-luciferase (Fig. 3A). Remarkably, multiple drugs that increased ARE-luciferase expression were also found to be FXN inducers (Supplementary Material, Table S1). In addition, after oral dyclonine dosing in the FA-YG8 mouse model of FA drives a dose-dependent increase in known Nrf2-target proteins in cerebellar lysate including heme oxygenase (HO1), NQO1 and GPX4 (Fig. 3B). Thus, dyclonine is an activator of the ARE/Nrf2 pathway.

Evolutionarily conserved Nrf2-binding sites (AREs) exist in the FXN gene and are functional Nrf2-binding sites

We studied the mechanism of induction of FXN by dyclonine with regard to Nrf2. If dyclonine induces FXN through the Nrf2 pathway, then there should be active Nrf2-binding sites, i.e. AREs (28) within the FXN locus that are triggered by dyclonine. We studied the mechanism of induction of FXN by dyclonine in multiple animal models after intraperitoneal or oral dosing. FA-YG8 hFXN+/− with FXN (GAA)190 expansion; mFXN−/− transgenic mice (8) dosed for 1 week once daily with dyclonine i.p. or p.o. showed a clear induction of FXN protein in cerebellar lysate (Fig. 2C).

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Figure 2. Dyclonine induces FXN in cultured FA patient cells and FA mouse model cerebellum in vivo. (A) Dyclonine increases FXN protein expression in FA patient lymphoblast cell lines. To test if the mechanism of protection from diamide toxicity for dyclonine was through an increase in FXN protein levels, FA lymphoblasts were treated with 0.3–10 μM dyclonine or vehicle control (0.1% DMSO). Total protein was collected after 48 h, and lysates were probed by western blot analysis for FXN expression and normalized to β-actin. The plotted data represent the mean fold change in FXN protein in drug-treated cells, normalized to vehicle control. Error bars represent SEM. **P < 0.01, t-test (n = 2–15). A representative blot is shown for FA patient lymphoblast line 15850 treated with 10 μM dyclonine for 48 h. (B) Dyclonine increases FXN transcript levels in FA patient lymphoblasts. FA lymphoblast cell line 14518 was treated with 3–30 μM dyclonine or vehicle control (0.1% DMSO). RNA was analyzed after 24 h by RT-PCR with expression with primers for FXN normalized to β-actin. The plotted data represent the mean fold change in FXN transcript in drug-treated cells normalized to vehicle control. Error bars represent SEM. *P < 0.05, **P < 0.01, (n = 4), t-test. (C) Dyclonine increases FXN...
in vivo protein concentration in FA-YG8 mouse cerebellum. To determine the ability of dyclonine to reverse the in vivo FXN protein defect, FA-YG8 transgenic mice [bFXN \textsuperscript{+/−} with FXN (GAA)\textsubscript{190} expansion; mFXN \textsuperscript{−/−}] were treated with 1–10 mg/kg dyclonine for 1 week i.p. or p.o. Cerebellar lysates were probed by western blot analysis for FXN expression and normalized to β-actin. The plotted data represent the mean fold change in FXN protein in drug-treated mice normalized to vehicle control. Error bars represent SEM. \textsuperscript{∗} P < 0.05, \textsuperscript{∗∗} P < 0.01, t-test (n = 3–10 and 11–12 months of age). A representative blot is shown for cerebellum from FA-YG8 mice treated with 10 mg/kg dyclonine for 1 week. FA-YG8 [bFXN \textsuperscript{+/−} with FXN (GAA)\textsubscript{190} expansion; mFXN \textsuperscript{−/−}] mice with two copies of transgene used as a control. (D) Dyclonine increases FXN protein levels in FA-PandKIKO mouse cerebellum. To determine the ability of dyclonine to reverse the in vivo FXN protein defect in an additional FA model at a higher dose and duration, FA-PandKIKO mice [mFXN \textsuperscript{−/−} with FXN (GAA)\textsubscript{230} expansion; mFXN \textsuperscript{−/−}] were treated with 25 mg/kg dyclonine for 1 or 4 weeks p.o. Cerebellar lysates were probed by western blot analysis for FXN expression and normalized to β-actin. The plotted data represent the mean fold change in FXN protein in drug-treated mice normalized to vehicle control. Error bars represent SEM. \textsuperscript{∗} P < 0.01, t-test (n = 3–17 and 8–10 months of age). A representative blot is shown for cerebellum from FA-YG8 mice treated with 25 mg/kg dyclonine for 1 week. Wild-type C57BL/6 mice used as a control.
Figure 3. Dyclonine drives an induction of Nrf2 through the AREs in FXN gene. (A) Dyclonine increases the expression of ARE-luciferase reporter gene. To explore the mechanism of FXN induction by dyclonine, effects on the nrf2-target ARE were evaluated in a reporter HeLa cell line transduced with ARE-luciferase. Cells were treated with 1.25–10 μM dyclonine or vehicle control (0.1% DMSO). After 24 h, cells were lysed and luciferase activity was measured on a plate reader. The plotted data represent the mean fold change in luminescence in drug-treated cells normalized to vehicle control. (+) control = 5 μM sulforaphane. Error bars represent SEM. *P < 0.01, t-test (n = 3). (B) Dyclonine increases nrf2-target protein expression in the FA mouse cerebellum. FA-YG8 transgenic mice [hFXN(+/−) with FXN(GAA)190 expansion; mFxn(−/−)] were treated with 1–10 mg/kg dyclonine for 1 week p.o. Cerebellar lysates were probed by western blot analysis for nrf2-target proteins HO1, NQO1, and GPX4 expression and normalized to β-actin. The plotted data represent the mean fold change in FXN protein in drug-treated mice normalized to vehicle control. Error bars represent SEM. *P < 0.05, **P < 0.01, t-test. (n = 3). A representative blot is shown for cerebellum from FA-YG8 mice treated with 10 mg/kg dyclonine for 1 week. (C) Dyclonine induces nrf2 binding to ARE sites in Fxn and Hmox1. Multiple ARE sites were found upstream of FXN gene (Supplementary Material, Fig. S2). The top two candidates 5597 and 16722 bp upstream were selected from position weight. ChIP assays were performed to determine the binding of Nrf2 to these sites and the promoter of Hmox1 as a positive control. FA lymphoblasts (GM14518, GM15850 and GM16220) were treated with vehicle (0.1% DMSO), 5 μM dyclonine or 5 μM of the nrf2-inducer dimethyl-fumarate for 24 h. Chromatin was immunoprecipitated with anti-nrf2 antibody or anti-IgG negative control and by PCR for target loci. The plotted data represent the mean fold enrichment of PCR product with Nrf2 pulldown compared with IgG control. This shows increased amplification of regions of DNA for both the ARE sites, 16 722 bp upstream of Fxn and Hmox1 after dyclonine treatment. Error bars represent SEM. *P < 0.05, **P < 0.01, t-test (n = 5 individual pulldowns per condition, n = 3 per experiments).
Table 1. ARE sites in human FXN gene

<table>
<thead>
<tr>
<th>Position</th>
<th>Position relative to FXN</th>
<th>Direction</th>
<th>DNA strand</th>
<th>Sequence</th>
<th>PWM score</th>
<th>Evolutionarily conserved (95% CI)</th>
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<td>Template</td>
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<tr>
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<td>−13 792</td>
<td>Antisense</td>
<td>Non-template</td>
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<tr>
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<td>Template</td>
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</table>

FXN gene starts at 71650479 and ends at 71715094 (64 615 bp). GAA-repeat insertion is 1731 bp downstream of FXN exon-1 start site. Position weight matrix (PWM) score from Wang et al. (28). Searched for the regular expression of ARE consensus sequence: **TGAC[ACTG][3]GC** from 20 kb upstream to 5 kb downstream of FXN.

receive dyclonine. Patients or caregivers dosed with dyclonine by an oral rinse for 6–7 days twice daily, and then came in for a final cheek swabbing for buccal cells. These blinded samples were then sent for analysis. FXN protein was analyzed by the clinically validated dipstick immunoassay (44,45). We modified these methods by further by normalizing FXN to mitochondrial cytochrome oxidase expression, i.e. FXN/Complex IV (CIV). Four repeats of the dipstick immunoassay were performed per patient sample. Of the 8 patients who received dyclonine, six experienced FXN increase (ranging from 60 to 480% induction), and two experienced no induction (Table 2). Interestingly, patients with greater neurological impairment as measured by the Friedreich’s ataxia rating scale (FARS) (46,47), experienced greater FXN induction (Fig. 6A). Other correlations that were also significant included the Functional Disability Score (FDS) and Z2 ataxia functional composite score (Fig. 6B and C) (48). While the correlation of FXN response with GAA repeat length was not significant (Supplementary Material, Fig. 6D), this trend could underlay the differential response seen in Figure 4A.

**DISCUSSION**

FA is caused by inheritance of (GAA)$_n$ expansions in intron 1 of the nuclear FXN gene that decreases its expression to about 20% of normal. All consequences of Friedreich’s are thought to result from this epigenetic silencing of FXN (5,6,11,49). There is currently no approved or effective therapy for this ultimately lethal disease that currently affects approximately 6000 in the USA and 20 000 in Europe (1). We identified a biochemical deficiency in Friedreich’s patient cells to design and perform a drug screen, and have identified a potential treatment for the disease that rescues FXN deficiency as well as downstream consequences in cells, animal models and in FA patients.

Our previous microarray of Friedreich’s mouse model DRGs indicated defects in thioredoxin reductase, thiol-antioxidant transcripts and a deficiency in Nrf2 activity in FXN knockdown cells (12). Also, a deficiency in Nrf2 activation was noted in Friedreich’s fibroblasts (18). Taken together, these data suggested that FXN deficiency may cause deficiency in Nrf2 activation or the related mitochondrial thioredoxin reductase pathway (50), which leads to decreased mitochondrial antioxidant protection, increased reactive oxygen species, inflammation and neurodegeneration (51).

Given the data on thiol antioxidant deficiency in mouse DRG (12), and previous data on antioxidant deficiency in FA fibroblasts (12,16,17), we tested poisons of thiol antioxidants in a FXN-deficient DRG cell line (Fig. 1A). In these cells, diamide and auranofin were more toxic to FXN knockdown cells compared with controls, whereas hydrogen peroxide (H$_2$O$_2$) and buthionine sulfoxamine (BSO) displayed little to no difference. This was interesting because diamide oxidizes thioredoxin (52), and auranofin is an inhibitor of thioredoxin reductase (53). These data support the idea that, in the context of the dorsal root ganglia, thioredoxin reductase is an important antioxidant system. Others have shown that, in neural mitochondria, thioredoxin reductase is the dominant antioxidant system (54,55). We examined FA patient cells and also observed increased cell death after diamide treatment which was reversed by the reductant dithiothreitol (DTT), confirming a thiol-oxidant mechanism (Fig. 1C).

To expedite the search for clinical candidates, we took a repurposing approach and examined a collection of 1600 drugs with mostly known pharmacokinetic and safety profiles. Our initial screen identified 100 protective hits in a range of drug classes, of which 33 were found to be reproducibly dose-dependent, for a hit rate of ~2.5% (Supplementary Material, Table S1). One of the strongest hit in terms of efficacy and potency at protecting from diamide stress was dyclonine, the active agent in Sucrets. Dyclonine was initially characterized as an anti-epileptic (56), but through its sodium channel inhibitory properties it has been used for anesthesia since 1955 (13). Dyclonine’s protective effect could potentially be due to multiple mechanisms, including increased mitochondrial biogenesis, Fe/S cluster biogenesis, iron binding or others. Since all FA consequences are the result of deficient FXN expression, we explored whether dyclonine was able to induce FXN in FA patient cells and the animal model (Fig. 2). We discovered an induction of FXN protein, and this was supported further with dose-dependent increases in FXN mRNA by RT-PCR. This has implications for therapy, since dyclonine rescues the protein deficiency that causes the disease in FA lymphoblasts and fibroblasts. In animal models of FA, we observed a clear 2-fold induction in FXN protein in two mouse models, in multiple tissues including cerebellum, liver and heart (Fig. 2C and D, and Supplementary Material, Fig. S1), two of which (cerebellum and heart) are major sites of disease pathogenesis. Our animal studies included intraperitoneal and oral dosing, with
drug exposures from 4 days to 4 weeks. In all, we have done 12 independent in vivo studies with dyclonine with between 3 and 6 mice per group, and each study supported an induction in FXN by dyclonine. Since FA patients are asymptomatic when their FXN levels are 40–50% of normal levels, an induction of 2-fold should boost patient FXN levels into this range, and alleviate consequences of FXN deficiency (57).

Early on in our development efforts, we noted that a few of the drugs we found to protect from diamide were also Nrf2 inducers in literature. Nrf2 translocates to the nucleus in response to oxidative stress (and especially thiol stress) and binds to AREs in target genes (27). Dyclonine had never been described to induce Nrf2, and we demonstrated a dose-dependent stimulation, and that multiple Nrf2-target proteins are elevated after treatment with dyclonine in both FA patient cells and animal model tissues (Fig. 3B). To investigate if the induction of FXN was Nrf2-mediated, we used bioinformatics to identify ARE sequences in the FXN locus, finding three ARE sites upstream of the transcription start site of FXN that are completely conserved across ape evolution (Table 1 and Supplementary Material, Fig. S2). The functionality of ARE sites was verified by ChIP and pull down with an Nrf2 antibody (Fig. 3C). Thus the induction of FXN is likely the result of recruitment of Nrf2 to adjacent ARE sites. Data also suggest that other Nrf2 inducers will reactivate FXN expression.

We observe that dyclonine has a differential effect on FXN re-expression from alleles of different expansion length. Specifically, longer GAA alleles, which experience greater epigenetic silencing, are more ‘reactivatable’ by dyclonine in cells (Fig. 4A), and this is consistent with what is observed in patients (Fig. 6C). There is precedent for Nrf2 activation having epigenetic effects on genes (35, 58–60). There is also recent precedent that the classical Nrf2-inducer sulforaphane may be working through epigenetic unsilencing of the Nrf2 locus (35, 36). Dyclonine has been reported to have inhibitory activity on histone lysine methyltransferase G9a, which is known to methylate histone H3K9 (PubChem, assay ID 504332). Inhibitory methylation of histone H3K9 is a known epigenetic mark at the FXN locus (31–34). And we observed experimentally that dyclonine inhibited histone methyltransferase G9 in lymphoblast nuclear extracts (Fig. 4B). These data support either of two mechanisms, i.e. that dyclonine induces Nrf2, and Nrf2 through its epigenetic unsilencing effects reactivates FXN; or that dyclonine suppresses histone methyltransferase G9, which in turn reactivates both the FXN locus and the Nrf2 locus.

Since dyclonine is an anesthetic whose pain ameliorating effects are mediated through sodium channel inhibition, there could be concern of the sedative potential of inhibiting neuronal NaV1.2 sodium channels (61). Additionally, sodium channel blockers are used to treat cardiac arrhythmias and alteration of heart rate would be a concern prior to treatment FA patients with systemic dyclonine. Since there were no data in the literature about the specific concentrations at which this occurred, we examined inhibition of NaV1.2 currents using patch-clamp electrophysiology. As expected, there was clear dose-dependent inhibition with an IC50 of ~6 μM. This IC50 is 30-fold higher than our EC50 for protection from diamide in FA fibroblasts, and we observe FXN induction in FA lymphoblasts with 1 μM dyclonine. Additionally, after 4 weeks of chronic daily dosing in mice, no sedation or adverse phenotype was observed.

To verify that there was no systemic peripheral anesthesia from ingestion of a topical anesthetic at the doses that induce FXN reactivation, we measured hot plate response and found no delay. Finally, we tested ~20 sodium channel inhibitors or anesthetics including lidocaine, bupivicaine and propofol and none were significantly active in the diamide protection assay. Taken together, this reduces the concern that the protective...
actions of dyclonine are mediated through sodium channel blockade, since the doses required for FXN activation are lower than those for sodium channel engagement.

The most well-defined downstream function for mammalian FXN is in iron–sulfur cluster biogenesis (40,42,62), and dyclonine treatment induced iron–sulfur cluster enzyme activity in animal and cell models (Fig. 5A–C). We also examined several neurobehavioral parameters in multiple FA mouse models, including open field activity, grip strength, treadscan and motor coordination on the level beam task. The most sensitive behavioral test was time to cross a level beam. Cerebellar defects in FA patients lead to motor coordination difficulties, and the level beam test mimics the walking score in the FARS ataxia score (47). We observed a significant improvement in this parameter after 4 weeks of oral dyclonine dosing.

Since dyclonine is FDA-approved as an oral rinse, its prescription and dispensing was allowed in an IRB-approved trial. Ten patients were recruited, but 2 did not complete the treatment and stopped 2–3 days into the study, likely because of the taste or mouth numbing that persisted for up to 1 h after each dosing. Of the remaining patients, 6 of the 8 showed an increase in FXN after 1 week. Additionally, both healthy volunteers responded to dyclonine with an increase in FXN (Table 2). The correlations with neurological scores are highly significant, and suggest that dyclonine produces a greater induction in patients with more severe FXN deficiency (Fig. 6).

FXN deficiency causes oxidative stress and subsequent cell death in dorsal root ganglia and cardiac tissue (3). Oxidative stress is suspected to be important in multiple neurodegenerative diseases (63–65). Here, we describe a well-known anesthetic drug dyclonine that protects FA patient cells from diamide-induced oxidative stress, increases FXN and rescues primary biochemical end points of the disease in cell and animal models, and appears to do so involving the Nrf2/ARE pathway. Limitations of this work include translation of beneficial effects seen in mice to humans, as well as the hope that FXN engagement after topical exposure with a rinse solution of dyclonine will model results after a systemic dosing regimen in FA patients. Although there are a number of questions that need to be addressed for these findings to be clinically applied to patients, including a better understanding of pharmacokinetics, optimal dosing formulation for oral bioavailability and cardiovascular toxicology, dyclonine represents a potential novel treatment strategy for the disease.

MATERIALS AND METHODS

Cell lines

Human control and FA patient fibroblasts and lymphoblasts were obtained from Coriell Institute for Medical Research repository. Patient lymphoblast cell lines used: GM16214, GM04079, GM16205, GM16243, GM15850, GM16220 and GM16197. Patient fibroblast lines used: 1133, 1134.

Diamide screening assay

Patient fibroblasts were grown in MEM media (Life Technologies Corp.) supplemented with 15% fetal bovine serum in t225 tissue culture flasks and kept below 70% confluency. Cells were trypsinized and density determined using a Vi-Cell counter (Beckman Coulter Corp.). Five thousand fibroblast cells were then aliquoted into 96-well poly-d-lysine-coated black/clear culture plates (Becton Dickinson Corp.) in growth media without antibiotics in a volume of 180 μl. Cells were allowed to adhere for 3–4 h at 37°C. Drugs (10 μm stock in dimethyl sulfoxide, DMSO) were dispensed into assay plate wells after an intermediate dilution in PBS, giving a final DMSO concentration of 0.1% using an electronic multichannel pipetter (BioHit, Sartorius Corp.). Pharmakon drug library consisted of 1600 compounds (Microsource Discovery Systems, Inc.) in 96 well plates at stock concentration of 10 μM in DMSO. Test compounds were all be tested at 10 μM final assay concentration in primary screen. Eight wells each of 300 μM dithiothreitol (Sigma-Aldrich Corp.) or 0.1% DMSO were used as positive or negative controls, respectively. Cells were then incubated at 37°C with 5% CO₂ overnight. After 24 h, diamide (Sigma-Aldrich Corp.) was added to all wells at a final concentration of 125 μM from a 100 μM stock solution prepared in DMSO and allowed to incubate at 37°C with 5% CO₂ overnight (14–18 h). Plates were then washed with PBS, supplemented with 1 μM Calcein-AM (Molecular Probes, Invitrogen Corp.) and incubated at room temperature for 45 min. Cells were then washed with PBS to remove residual dye, and read on BMG PolarStar Optima with 485 excitation and 520 emission wavelengths (BMG LabTech).

Western blot analysis

Cells were lysed using lysis buffer (Promega Corp.) supplemented with a complete protease inhibitor cocktail (Roche Applied Science) and phenylmethylsulfonyl fluoride (Sigma-Aldrich Corp.). Tissues were further homogenized using 0.5 mm glass beads in a Bullet Blender high-throughput homogenizer (Next Advance, Inc.). After pelleting cellular debris by spinning at 16 000 rpm at 4°C for 15 min, protein was quantified by Bradford assay (66). For western blotting, 40–50 μg protein was added per lane of 4–12% Bis-Tris gels (Invitrogen Corp.). Primary antibodies were diluted in Odyssey blocking buffer (LI-COR Biosciences). Antibodies used included: anti-FXN (provided by Franco Taroni M.D., Istituto Besta), anti-HO1 (#sc-10789, Santa Cruz Biotechnology, Inc.), anti-NQO1 (#3187, Cell Signaling Technologies), anti-Gpx4 (#ab125066, AbCam), anti-nrf2 (sc-25820, Santa Cruz) and anti-actin (#A2668, Sigma). Direct conjugated secondary antibodies (anti-rabbit IRDye800CW and anti-mouse IRDye680 from LI-COR) were used to detect and quantify the signal of primary antibodies and imaged using a LI-COR Odyssey.

Bioinformatics and phylogenetic analysis of FXN locus

A position weight matrix was used to identify ARE sequences in FXN based on previously described functional ARE sequences (28). A pre-filter was used with the consensus sequence of nnnnnnTGACnnnnn to identify potential ARE sites. The position weight matrix was used to calculate the score for each of the pre-filtered hits. Phylogenetic footprinting was then done using Clustalw2, and a multiple alignment was done between the great apes (Homo sapiens, Pan troglodytes,
Figure 5. Dyclonine recovers downstream effects of FXN deficiency in vitro and in vivo. (A) Dyclonine increases aconitase activity in FA patient lymphoblast cell lines. FA lymphoblasts (GM16205, GM16197 and GM16243) were treated with 10 μM dyclonine or vehicle control (0.1% DMSO). Cell pellets were collected after 48 h, and aconitase activity was measured in lysates over 90 min to ensure linear range was captured. A healthy volunteer lymphoblast cell line was used as a control. The plotted data represent the mean rate change in fluorescence (dF/dt) from 25 to 30 min. Error bars represent SEM. *P < 0.05, t-test. (n = 3). (B) Dyclonine increases aconitase activity in FA mouse model cerebellum in vivo. FA-YG8 transgenic mice [hFXN+/− with FXN (GAA)190 expansion; mFXN−/−] were treated with 5 mg/kg dyclonine for 1 week p.o. Aconitase activity was measured in cerebellar lysates over 90 min to ensure linear range was captured. Positive controls were vehicle-treated FA-YG8 transgenic mice [hFXN+/+ with FXN (GAA)190 expansion; mFXN−/−]. The plotted data represent the mean rate change in fluorescence (dF/dt) from 15 to 20 min. Error bars represent SEM. *P < 0.05, t-test (n = 4 mice per group, 11–12 months of age). (C) Dyclonine increases succinate dehydrogenase activity in FA mouse model liver in vivo. FA-PandKIKO mice [mFXN+/− with FXN (GAA)230 expansion; mFXN−/−] were treated with 25 mg/kg dyclonine for 1 week p.o. Succinate dehydrogenase activity was measured in liver lysates over 90 min. Positive controls were vehicle-treated WT C57/Bl6 mice. The plotted data represent
follows: FXNFWD 5′- ATCTTCTCCATCCAGTGGACCT-3′; ARE 16223FWD 5′- CCTGCCGTACTCGATCCCTC-3′ and ARE 16223REV 5′- CCACCTCGGTACTGTCGTG-3′; NQO1FWD 5′- CCCTTTTGAAGCCTTGGCACGAAA-3′ and NQO1REV 5′- TGCACCAGGAAAGTTGTTGTAT-3′; HO1FWD 5′- CCCTGCTGATATACCTTTCCGAA-3′ and HO1REV 5′- ATGTCGCCACTCCAGACTCCA-3′ (67).

ARE reporter assay

HeLa cells were treated with Cignal Lentivirus Reporter ARE-luciferase lentiviral reagent (SABiosciences, Cat # CLS-2020L) and underwent selection with 1 μg/ml of puromycin to generate stable cell line. Hela-ARE stable cells were washed with PBS and 1 × EDTA, and plated in phenol-free DMEM media supplemented with 10% FBS in white wall/bottom 96-well plates in 90 μl. Plates were incubated at 37°C for 2 h to allow cells to adhere. Drugs or vehicle were added to all wells (10 μl/well). Cell plates were incubated at 37°C for 24 h. About 75 μl/well of cell lysis/luciferase Bright-Glo reagent (Promega) were then added to all wells. Wells were mixed with a biotech electronic pipettor (100 μl 2×). After 5 min without shaking, luminescence was read on BMG Polarstar.

Aconitase assay

Aconitase activity was measured in patient lymphoblast and mouse model cerebellar lysate using the Aconitase Assay Kit (Cayman chemicals, Cat # 700600) and performed according to the manufacturer’s instructions.

Succinate dehydrogenase assay

Succinate dehydrogenase activity was measured (67) in mouse model liver lysate using the Complex II Enzyme Activity Microplate Assay Kit (Abcam, Cat # ab109908) and performed according to the manufacturer’s instructions.

Histone methyltransferase assay

G9 methyltransferase activity was measured in healthy lymphoblast nuclear extracts (Epipik Nuclear Extraction Kit, Epigentek, Cat # OP-0002) and using the EpiQuik H3K9 Histone Methyltransferase Activity/Inhibition Assay Kit (Epigentek, Cat # P-3003) and performed according to the manufacturer’s instructions.

Animal procedures and behavioral assessments

All animal procedures were approved by the Institutional Animal Care and Use Committee at the University of California, Davis and the UC Davis Animal Care and Use Committee. The mean latency to cross a beam post-dosing minus the time to cross pre-dosing for each individual mouse and serve as a change from baseline measurement. As a change from baseline, a negative number means that the animal was faster after dosing than it was before dosing. Positive controls were vehicle-treated WT C57/B6 mice, *P < 0.05, t-test (n = 7–9 mice per group, 12–14 months of age).

<table>
<thead>
<tr>
<th>Patient</th>
<th>GAA1</th>
<th>GAA2</th>
<th>Fold induction</th>
<th>P-value</th>
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</thead>
<tbody>
<tr>
<td>FRDA4522</td>
<td>600</td>
<td>500</td>
<td>0.4 ± 0.1</td>
<td>0.01</td>
</tr>
<tr>
<td>FRDA0055</td>
<td>1178</td>
<td>826</td>
<td>4.8 ± 1.9</td>
<td>0.01</td>
</tr>
<tr>
<td>FRDA0171</td>
<td>600</td>
<td>600</td>
<td>3.6 ± 1.7</td>
<td>0.02</td>
</tr>
<tr>
<td>FRDA0157</td>
<td>850</td>
<td>850</td>
<td>2.6 ± 1.4</td>
<td>0.01</td>
</tr>
<tr>
<td>FRDA0235</td>
<td>1122</td>
<td>230</td>
<td>0.6 ± 0.3</td>
<td>0.02</td>
</tr>
<tr>
<td>FRDA4539</td>
<td>1045</td>
<td>670</td>
<td>1.6 ± 0.5</td>
<td>0.07</td>
</tr>
<tr>
<td>FRDA4579</td>
<td>1135</td>
<td>71</td>
<td>1.3 ± 0.6</td>
<td>0.31</td>
</tr>
<tr>
<td>FRDA4576</td>
<td>800</td>
<td>800</td>
<td>1.8 ± 0.6</td>
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</tr>
<tr>
<td>HV0001</td>
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<td>ND</td>
<td>5.2 ± 1.9</td>
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<tr>
<td>HV0002</td>
<td>ND</td>
<td>ND</td>
<td>1.7 ± 0.3</td>
<td>0.01</td>
</tr>
</tbody>
</table>

Patients and controls were dosed for 1 week with a 1% dyclonine oral rinse, twice daily for 30 s. Buccal cell samples were taken by clinician use swabs and blinded prior to analysis. GAA1 and GAA2 refer to the number of GAA repeats in both alleles of the patient’s FXN gene (ND, not determined). FXN was measured in blinded samples using dipstick immunoassay normalized to complex IV expression. Fold induction is a value post-treatment divided by pre-treatment FXN/CIV ± SD (n = 4 measurements per patient). P-value determined by the t-test.

Gorilla and Pongo abeli) in the region upstream of the FXN gene (29).

Chromatin immunoprecipitation

Patient lymphoblasts were seeded at 0.5 × 10⁶/ml in tissue culture flasks, and incubated with 0.1% DMSO, 5 μM dyclonine or 5 μM dimethyl-fumarate. About ~10 × 10⁶ cells were used for each condition. After 24 h, cells were treated with 18.5% paraformaldehyde to crosslink proteins to DNA. ChIP was performed using an EZ-Chip kit (EMD Millipore). After shearing by sonication, each cell lysate was split and incubated with either Anti-Nrf2 (#H-300) or anti-mouse IgG negative control (Santa Cruz Biotechnology, Inc.) and protein-G-conjugated agarose beads to immunoprecipitate cross-linked protein/DNA. Crosslinks were reversed by incubating at 65°C overnight, and DNA was purified with spin columns and buffers. DNA concentration was determined using a Nanodrop 2000c (Thermo Scientific Corp.), and then analyzed by quantitative real-time PCR.

Quantitative real-time PCR

Quantitative PCR was performed using the Superscript III One Step kit (Invitrogen Corp) in a Roche Lightcycler 480 (Roche Diagnostics). Standard curves were generated for each primer set, and samples fitted to the linear portion of the curve. Additionally, PCR products were run on DNA gels to verify single products. Primer sequences used were described previously and were as follows: FXNFWD 5′- ATCTTCTCCATCCAGTGGACCT-3′ and FXNREV 5′- GCTGGGCATCAAGCCTTCTT-3′; ARE 16223FWD 5′- CCTGCCGTACTCGATCCCTC-3′ and ARE 16223REV 5′- CCACCTCGGTACTGTCGTG-3′; NQO1FWD 5′- CCCTTTTGAAGCCTTGGCACGAAA-3′ and NQO1REV 5′- TGCACCAGGAAAGTTGTTGTAT-3′; HO1FWD 5′- CCCTGCTGATATACCTTTCCGAA-3′ and HO1REV 5′- ATGTCGCCACTCCAGACTCCA-3′ (67).

Table 2. Buccal FXN expression in FA patients after treatment with a 1% dyclonine oral rinse
with adherence to the NIH Guide for the Care and Use of Laboratory Animals. The age of mice used for all studies was between 6 and 14 months and was age-matched within experiments. Drug dosing was done i.p. formulated in a DMSO and PBS mixture, and p.o. formulated in a DMSO, water and peanut butter mixture. Behavior testing was performed at the University of California, Davis Mouse Behavioral Analysis Laboratory. A colony of YG8R B6.Cg-Fxntm1Mkn Tg(FXN)YG8Pook/J mice was established at UC Davis (Jackson Labs, # 012253) (68). For our testing, we used both hemizygous mice [FA-YG8 (hFXN\(^{+/+}\) with FXN (GAA)\(_{190}\) expansion; mFxn\(^{-/-}\)] containing one allele of the mutant FXN transgene and one KO (and thus the least amount of FXN) and homozygous mice [FA-YG8 (hFXN\(^{+/+}\) with FXN (GAA)\(_{190}\) expansion; mFxn\(^{-/-}\)], containing two alleles of the transgene (and thus higher levels of FXN), and wild-type (WT) mice. In addition, the B6.Cg-Fxn\(^{tm1Mkn}\) Fxn\(^{tm1Pand}\) J strain was also established and used (Jackson Labs, # 014162), and is referred to as FA-PandKIKO mice [mFxn\(^{+/+}\) with FXN (GAA)\(_{230}\) expansion; mFxn\(^{-/-}\)] (69,70). Both of these mice are bred on a C57BL/6J background.

Level beam performance was measured by placing mice on a 100 cm long level beam with a line marked 10 cm from each end as the start and end lines. A 60 W desk lamp was used as an aversive stimulus at the starting end of the beam, and an enclosed...
safety in this study. The Animal Care and Use Committee at the University of California, Los Angeles approved all procedures involving animals. All experimental protocols were designed in accordance with the Institutional animal Care and Use Committee at the University of California, Davis.

SUPPLEMENTARY MATERIAL

Supplementary Material is available at HMG online.

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Conflict of Interest statement. The authors have no competing financial interests. S.S. and G.A.C. are inventors on a pending patent on identification of agents useful for the treatment of Friedreich’s ataxia.

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