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## High resolution studies of hydride transfer in the ferredoxin:NADP<sup>+</sup> reductase superfamily

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9 **High resolution studies of hydride transfer in the ferredoxin:NADP<sup>+</sup>**  
10 **reductase superfamily**

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36 **Running Title:** Hydride Transfer in Ferredoxin:NADP<sup>+</sup> Reductase

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38 **Database:** Structural data are available in the PDB database under the accession numbers **3LO8**  
39 (wild type), **5VW4** (Y316S:nicotinamide (P3<sub>2</sub>21)), **5VW9** (Y316S:nicotinamide (P3<sub>2</sub>21)),  
40 **5VW3** (Y316S:NADP<sup>+</sup> (P3<sub>2</sub>21)), **5VW8** (Y316S:NADP<sup>+</sup> (P3<sub>1</sub>21)), **5VW2** (Y316S:NADPH  
41 (P3<sub>2</sub>21)), **5VW5** (Y316A:nicotinamide (P3<sub>2</sub>21)), **5VW6** (Y316A:NADP<sup>+</sup> (P3<sub>2</sub>21)), **5VW7**  
42 (Y316A:NADPH (P3<sub>2</sub>21)), **5VWA** (Y316F (P3<sub>2</sub>21)), and **5VWB** (Y316F:NADP<sup>+</sup> (P3<sub>1</sub>21)).

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44 **Abbreviations:** FNR – ferredoxin:NADP<sup>+</sup> reductase; NOX – NADPH oxidase; Fd – ferredoxin;  
45 CTC-1 – charge-transfer complex 1; CTC-2 – charge-transfer complex 2

46 **Enzyme Commission number:** ferredoxin:NADP<sup>+</sup> reductase – EC 1.18.1.2

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48 **Keywords:** flavoenzyme, hydride transfer, enzyme mechanism, protein crystallography,  
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**ABSTRACT**

Ferredoxin-NADP<sup>+</sup> reductase (FNR) is an FAD-containing enzyme best known for catalyzing the transfer of electrons from ferredoxin (Fd) to NADP<sup>+</sup> to make NADPH during photosynthesis. It is also the prototype for a broad enzyme superfamily, including the NADPH oxidases (NOXs) that all catalyze similar FAD-enabled electron transfers between NAD(P)H and one-electron carriers. Here we define further mechanistic details of the NAD(P)H  $\rightleftharpoons$  FAD hydride-transfer step of the reaction based on spectroscopic studies and high resolution ( $\sim 1.5$  Å) crystallographic views of the nicotinamide-flavin interaction in crystals of corn root FNR Tyr316Ser and Tyr316Ala variants soaked with either nicotinamide, NADP<sup>+</sup>, or NADPH. The spectra obtained from FNR crystal complexes match those seen in solution and the complexes reveal active site packing interactions and patterns of covalent distortion of the FAD that imply significant active site compression that would favor catalysis. Furthermore, anisotropic B-factors show that the mobility of the C4 atom of the nicotinamide in the FNR:NADP<sup>+</sup> complex has a directionality matching that expected for boat-like excursions of the nicotinamide ring thought to enhance hydride transfer. Arguments are made for the relevance of this binding mode to catalysis, and specific consideration is given to how the results extrapolate to provide insight to structure-function relations for the membrane-bound NOX enzymes for which little structural information [has been](#) available.

## INTRODUCTION

Plant ferredoxin NADP<sup>+</sup> oxidoreductases (FNR) are flavoenzymes that catalyze the reversible electron transfer between NADP(H) and the iron-sulfur protein ferredoxin (Fd). In photosynthesis, plastidic FNRs serve to catalyze electron transfer from Fd to NADP<sup>+</sup> via the FAD prosthetic group, and in non-photosynthetic tissues distinct FNRs use NADPH as an electron source for the reverse reaction [1-3][1-3]. Plant-type FNRs – as seen in three-dimensional structures of FNRs from cyanobacteria [4][4], spinach (leaf [5][5]), pea (leaf [6][6]), paprika (leaf [7][7]), and corn (leaf [8][8] and root [9][9]) – are highly similar and contain two distinct domains: one for binding FAD and one for binding NADP<sup>+</sup> [10][10]. Importantly, this two-domain FNR-fold is also the prototype for an enzyme superfamily that uses a bound FAD or FMN to transfer redox equivalents between the hydride carrying NAD(P)<sup>+</sup> cofactors and diverse one-electron carriers [10-12][10-12]. Included among the FNR superfamily are the NADPH oxidases (NOXs), a biomedically important group of membrane bound enzymes that produce superoxide or hydrogen peroxide as part of many signaling pathways as well as during the oxidative burst of macrophages that is a key part of our immune defenses (recently reviewed in [13-16][13-16]).

Structures of plant type FNRs and other superfamily members, such as nitric oxide synthase [17][17], cytochrome P450 reductase [18, 19][18, 19], and phthalate dioxygenase reductase [20][20] showed that NADP(H) appears to bind non-productively to the wild-type enzymes because an aromatic side chain (a C-terminal Tyr residue in the case of plant type FNRs) sits in the site the nicotinamide ring must occupy for hydride transfer (Figure 1A). Deng *et al.* [6][6] resolved this mystery by showing for pea leaf FNR at 1.8 Å resolution that a mutant missing the aromatic placeholder residue (Tyr308Ser or Y308S) could bind NADP(H) tightly and in an

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9 apparently productive manner with the nicotinamide ring C4 atom adjacent to the FAD N5 atom  
10 and with a geometry reasonable for hydride transfer. The Y308S mutant bound its cofactor so  
11 tightly that NADP<sup>+</sup> co-purified with the enzyme. This mutant also showed an unexpected 500-  
12 fold change in cofactor specificity as the stronger binding of NAD(H) *increased* its steady-state  
13 turnover [21][24], but the stronger binding of NADP(H), *decreased* its steady state turnover, not  
14 because of impaired hydride transfer but because the NADP binding was so tight that  $k_{\text{off}}$  became  
15 rate-limiting [4, 21][4, 24].  
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22 With the support of spectroscopic studies, it was concluded that the wild-type enzyme binds  
23 NADP(H) in a bipartite fashion in which the 2'-P-AMP binds strongly to anchor the cofactor to  
24 the enzyme and the thermodynamics of nicotinamide displacing the aromatic placeholder is such  
25 that the nicotinamide ring remains largely in a solvent exposed conformation (“nicotinamide-  
26 out”) and is in rapid equilibrium with a smaller population of molecules in which the aromatic  
27 placeholder swings out and the nicotinamide ring takes its place (“nicotinamide-in”  
28 conformation) to allow for hydride transfer. The Tyr to Ser mutant changes the binding  
29 thermodynamics so that the nicotinamide is ~100% “in”. Since that report, studies of aromatic  
30 blocker variants of other superfamily members have shown a consistent geometry for the  
31 “nicotinamide-in” nicotinamide-flavin interaction [4, 19][4, 19] (Figure 1B) as well as changes  
32 from 50 to 1000-fold in NADP<sup>+</sup> vs NAD<sup>+</sup> specificity [4, 22-25][4, 22-25].  
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43 Although there is considerable structural information about FNRs and FNR-like modules in  
44 related enzymes, the mechanism of hydride transfer is still not fully understood. For instance, the  
45 previous NADP(H) complexes of the pea FNR mutants gave indications that the C4 in the  
46 nicotinamide ring of NADP<sup>+</sup> was mobile in a way that could favor hydride transfer [6][6], but  
47 higher resolution data are needed to better define the details of this mobility. Also, it has been  
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9 proposed that the geometry of the complex seen in the structures that have the aromatic  
10 placeholder mutated are artifacts that are not relevant to catalysis ~~[26-28],[10,11]~~. Here, we  
11 obtain further insight into the hydride-transfer step in FNR-like enzymes by combining  
12 spectroscopic studies and a series of higher resolution structures of FNR crystals soaked with  
13 nicotinamide, NADP<sup>+</sup>, and NADPH. To accomplish this, we chose corn root FNR, for which  
14 crystals of the wild-type enzyme diffract to near 1 Å resolution ~~[29]. We also[26]. We further~~  
15 argue for the general relevance of these insights for broader members of the FNR superfamily,  
16 including the NOX enzymes, which is further supported by the very recent structures of the core  
17 catalytic subunit domains of NOX5, published while this work was under review [30].  
18 ~~membrane proteins that have not yet been structurally defined [15].~~  
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## 30 RESULTS AND DISCUSSION

### 31 Strategy

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33 An inspection of the molecular packing in the well-diffracting crystals of corn root FNR that  
34 we reported earlier ~~[9, 29][9, 26]~~ revealed that the NADP(H) binding site was not involved in  
35 crystal packing interactions. We hypothesized that we could obtain high resolution structures of  
36 NADP(H) complexes of this enzyme by making mutants of Tyr316, the aromatic blocking  
37 residue. Not knowing which mutant would be more informative, we mutated Tyr316 to Ala  
38 (Y316A), Ser (Y316S), and Phe (Y316F). Crystal structures of NADP<sup>+</sup> soaks of Y316F FNR  
39 showed no nicotinamide binding, so this variant was not subjected to further study. For Y316S  
40 and Y316A, however, we present extensive characterization of the in-solution spectroscopic and  
41 catalytic properties, as well as crystal structures and in some cases the in-crystal spectroscopic  
42 properties of their complexes with NADP(H).  
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### Solution Properties of Mutants

Wild-type and mutant enzyme forms were produced in *Escherichia coli* and purified essentially as reported elsewhere [9][8]. The two protein variants were expressed at slightly lower levels than that the wild-type enzyme, and their purification required additional hydrophobic interaction chromatography on butyl Sepharose. This resulted in a substantial drop in overall yields, which in the case of Y316A was low enough to preclude its full functional characterization. As shown in Figure 2A, the two replacements induced very similar perturbations in the visible absorption spectrum of the flavoprotein, strongly reminiscent of those observed in the equivalent Tyr to Ser replacement in both pea and *Anabaena* FNRs [9, 31][9, 27]. Phenol – a mimic of the Tyr side-chain – was found to induce a spectral change in Y316S that qualitatively matched the difference between the absorption spectra of the wild-type and the variant protein (Figure 2B), implying that it can stack against the isoalloxazine ring in the active-site pocket. However, the affinity of Y316S FNR for phenol was too low to allow the accurate determination of the  $K_d$  of the complex.

*Titration with NADP(H)*. Anaerobic titration of oxidized Y316S FNR with  $\text{NADP}^+$  induced perturbations in the visible absorption spectrum of the FAD prosthetic group (Figure 3) very similar to those observed for the Y308S mutant of pea FNR [6][6]. The  $K_d$  of the complex was estimated to be  $\leq 0.02 \mu\text{M}$ , well below that of the wild-type complex ( $0.3 \mu\text{M}$ ) [9][9]. Whereas no interaction between  $\text{NAD}^+$  and wild-type FNR was detectable spectrophotometrically,  $\text{NAD}^+$

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9 induced in Y316S FNR a spectral shift virtually identical to that produced by NADP<sup>+</sup>, displaying  
10 a  $K_d$  of  $145 \pm 5 \mu\text{M}$  (Figure 3).  
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14 Next, using an EDTA-deazariboflavin system, stepwise anaerobic photoreductions of the  
15 FAD group of wild-type and mutant FNR forms were done in the absence and in the presence of  
16 roughly equimolar amounts NADP<sup>+</sup> or NAD<sup>+</sup> (Figure 4). These showed that the Y316S  
17 replacement decreased the amount of FAD semiquinone accumulated during the process and  
18 greatly increased the intensity of the broad long wavelength (~800 nm) absorption band  
19 attributable to charge-transfer electronic transitions of the interacting oxidized nicotinamide and  
20 2-electron reduced flavin [9],[8]. Upon complete reduction of the system, the charge-transfer  
21 band disappeared implying that the limited amount of NADP<sup>+</sup> present became reduced and no  
22 charge transfer interaction occurs between the reduced flavin and NADPH. Furthermore, the  
23 presence of NADP<sup>+</sup> favored the protonation of the dihydroquinone form of the prosthetic group  
24 of Y316S FNR, as indicated by the shift from 360 to 420 nm of the local maximum of the spectra  
25 of the fully reduced species (Figure 4). The results with NAD<sup>+</sup> (not shown) are qualitatively  
26 similar to those obtained with NADP<sup>+</sup>, although a significantly less intense charge-transfer band  
27 was observed, consistent with the lower affinity of the enzyme for NAD<sup>+</sup> as compared to  
28 NADP<sup>+</sup>.  
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41 Anaerobic titrations of wild-type FNR and its variants with NADPH were performed to  
42 further analyze the interactions between FAD and NADP(H). In such titrations, at one equivalent  
43 of NADPH added, for the “nicotinamide-in” species an equilibrium should exist between two  
44 charge transfer complexes depending on whether the nicotinamide or the flavin is in the reduced  
45 state. Charge-transfer complex 1 (CTC-1) has the nicotinamide ring of NADPH interacting with  
46 oxidized flavin (*i.e.* FNR<sub>ox</sub>:NADPH) and an absorption band with a maximum near 600 nm, and  
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9 charge transfer complex-2 (CTC-2) has the nicotinamide ring of  $\text{NADP}^+$  interacting with reduced  
10 flavin (*i.e.*  $\text{FNR}_{\text{red}}\cdot\text{NADP}^+$ ) and a broad absorption band with a maximum at ca. 800 nm [32,  
11 33][28, 29]. The spectra at one equivalent of NADPH show that for wild-type the CTC-1  
12 absorption band dominates (Figure 5A), whereas for both mutants the CTC-2 band dominates  
13 (Figure 5B,C), apparently accounting for ~90% of the population, based on the level of reduced  
14 flavin reduction as indicated by  $A_{460}$ . For the two mutants, the CTC-2 band is maximal at  
15 roughly one equivalent NADPH and decreases to about half of that height as the concentration of  
16 NADPH is increased to an ~135-fold excess (Figure 5B,C). The further additions of NADPH  
17 presumably result in the progressive decrease in the amount of CTC-2 due to the partial  
18 displacement of  $\text{NADP}^+$  by NADPH. For the wild-type enzyme, the amount of CTC-1 similarly  
19 decreases as NADPH concentration increases, but it is progressively replaced by a species  
20 attributable to a complex between NADPH and FNR carrying FAD semiquinone. We do not  
21 understand how the semiquinone is formed under these conditions, but wonder if it may reflect  
22 some flavin disproportionation between the CTC-2 and CTC-1 complexes or is the result of a  
23 small amount of residual oxygen or other one-electron acceptor in the titrating NADPH solution.  
24 Prolonged incubation of NO synthase with NADPH was also seen to promote semiquinone  
25 buildup [34][30].

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41 *Spectral properties during turnover as an NADPH oxidase.* To monitor the spectral changes  
42 that occur during steady-state NADPH oxidase activity of FNR, air was admitted into the  
43 anaerobic cell after the titration was complete and NADPH was >100-fold in excess. In each  
44 case, the enzyme underwent turnover until the NADPH was all converted to  $\text{NADP}^+$ . For wild-  
45 type FNR (Figure 5D), as has been seen before [9][9], there was substantial stabilization of the  
46 blue semiquinone form of FAD in the early parts of the reaction and CTC bands were hardly  
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9 detectable. For Y316S FNR (Figure 5E), less FAD semiquinone built up and the reaction had  
10 two distinct phases. During the first phase, the CTC-2 concentration progressively increased until  
11 at ~15 min it reached ~65% of the maximal value observed during the anaerobic titration (thin  
12 red line), then it decreased to near zero. The final spectrum (at ~1.7 h) had the 456 nm band  
13 shifted as is characteristic of an oxidized enzyme in complex with NADP<sup>+</sup> (Figure 5E). The  
14 behavior of Y316A FNR (Figure 5F) was qualitatively similar to that of the Y316S form, but  
15 with the reaction slowed by about 10-fold as the steady-state CTC-2 formation was reached after  
16 about 2.5 h (*vs* ~15 min) and the reaction was complete in 17 rather than 1.7 h. The Y316A  
17 variant also had still less stabilization of the FAD semiquinone, and its maximal CTC-2  
18 accumulation during turnover was ~85% of the highest amount observed in the anaerobic  
19 titration (thin red line). The slower reaction and higher CTC-2 buildup are consistent with the  $k_{\text{off}}$   
20 for NADP<sup>+</sup> being slower for the Y316A mutant. Also, the destabilization of the semiquinone  
21 seen in these variants mirrors what was seen for the corresponding mutants of *Anabaena* FNR  
22 [31][27] and NO synthase [22, 34][22, 30].

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*Ferricyanide reductase activity.* For the wild-type enzyme, ferricyanide is the most effective  
electron acceptor from reduced FNR, with  $k_{\text{cat}} = 520 \text{ s}^{-1}$  [9][9]. The Y316S and Y316A  
replacements impaired the steady-state diaphorase activity of FNR to a similar extent, and  $k_{\text{cat}}$   
and  $K_{\text{m}}^{\text{NADPH}}$  values determined for the Y316S variant were ca. 170 and 9-fold lower than those  
of the wild-type enzyme (Table 1) [9][8], respectively. Such large catalytic impairment parallels  
that reported for the Tyr to Ser variants of pea and *Anabaena* FNRs [21, 31][21, 27], with the  
slower turnover shown to be limited by the rate of NADP<sup>+</sup> (*i.e.* product) release. Consistent with  
this, we found that nicotinamide partially rescued the diaphorase activity of Y316S FNR in a  
concentration-dependent fashion, increasing its  $k_{\text{cat}}$  to 280 s<sup>-1</sup> at 800 mM nicotinamide (Table 1).

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9 As has been noted previously [2, 6][2, 6], by competing with the corresponding moiety of  
10 NADP<sup>+</sup> for stacking onto the flavin, nicotinamide will favor product dissociation and speed  
11 enzyme turnover if the  $k_{\text{off}}$  of NADP<sup>+</sup> is rate limiting.  
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14 *Interactions with small-molecule ligands.* The findings that added nicotinamide increased  
15 the catalytic activity of Y316S FNR, prompted us to quantify the affinities of nicotinamide and  
16 N-methyl-nicotinamide. As shown in Figure 3A, both were found able to interact with Y316S  
17 FNR, although only the former had an affinity high enough for its  $K_d$  value to be determined ( $3 \pm$   
18  $0.1$  mM). The difference spectrum of the Y316S-nicotinamide complex was blue-shifted by ca. 7  
19 nm with respect to that of the NAD(P)<sup>+</sup> complexes, but the spectral change induced by N-  
20 methyl-nicotinamide was almost identical to that induced by NAD(P)<sup>+</sup> (Figure 3B). This  
21 indicates that for Y316S FNR, N-methyl-nicotinamide but not nicotinamide is an excellent  
22 mimic of the redox-active moiety of the dinucleotides.  
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### 33 **Reanalysis of Stopped-Flow Results Reported for *Anabaena* FNR.**

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35 As noted in the introduction, in earlier work Lans *et al.* [26][34] called into question the  
36 relevance of the NADP(H) complexes that form in FNR when the aromatic placeholder is  
37 mutated. The origin of this view came from stopped-flow results for the Tyr303Ser (Y303S)  
38 mutant of *Anabaena variabilis* FNR. Per their interpretation, these results showed that Y303S  
39 promoted the hydride transfer from NADPH to FNR<sub>ox</sub> “slightly slower than the wild-type  
40 enzyme” but that “the reverse process is undetectable” (*e.g.* hydride transfer from FNR<sub>red</sub> to  
41 NADP<sup>+</sup> does not occur). This interpretation was based on the observation that when NADP<sup>+</sup> was  
42 mixed with Y303S FNR<sub>red</sub> the CTC-2 (~800 nm) charge transfer band that formed upon mixing  
43 did not substantially decrease over time (see Figure 3A,B of Lans *et al.* [26][34] reproduced  
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~~here in Figure S1~~). While the conclusion that no hydride transfer occurs is consistent with the unchanging CTC-2 band, there could be other explanations. We were skeptical of the inference both based on the thermodynamic principle that if the Y303S mutant catalyzed hydride transfer from NADPH to FNR<sub>ox</sub> it must equally well catalyze the reverse hydride transfer from FNR<sub>red</sub> to NADP<sup>+</sup>, and based on our photoreduction studies (Figure 4 above) implying that Y316S-bound NADP<sup>+</sup> is converted to NADPH. To clarify this question, we reanalyzed the Lans *et al.* [26][34] stopped-flow data for NADPH mixed with Y303S FNR<sub>ox</sub> and for NADP<sup>+</sup> mixed with Y303S FNR<sub>red</sub>.

Importantly, in the original work, when NADP<sup>+</sup> is mixed with Y303S FNR<sub>red</sub>, even though the 800 nm band changes little over time, the ~460 nm peak systematically increases indicating that oxidation of FAD is occurring. Lans *et al.* attributed this to a “side effect” rather than enzymatic activity; however, when comparing these spectra with those for NADPH mixed with Y303S FNR<sub>ox</sub>, it is striking that both reactions appear to reach the same endpoint, as would be expected if Y303S FNR were reaching the same equilibrium state independent of the direction of approach. Given this observation, we reanalyzed the reported stopped-flow results for both reactions using the changes in A<sub>460</sub> as an indicator of FAD oxidation/reduction (Figure S1). This reanalysis yielded excellent fits of  $k_{\text{obs}}$  of  $164 \pm 16 \text{ s}^{-1}$  for NADP<sup>+</sup> mixed with Y303S FNR<sub>red</sub> and  $146 \pm 21 \text{ s}^{-1}$  for NADPH mixed with Y303S FNR<sub>ox</sub> (Figure 6). These two  $k_{\text{obs}}$  values should both equal the sum of the elementary forward and back reaction rate constants ( $k_{\text{obs}} = k_1 + k_{-1}$ ) and are equal within the error of the analysis, leading us to conclude that the forward and back reactions both function as would be expected. We suggest the best number to use for  $k_{\text{obs}}$  for the *Anabaena* Y303S FNR is  $190 \pm 15 \text{ s}^{-1}$  which is the value originally reported by Lans *et al.* [26][34] for the forward reaction using the much more extensive original dataset.

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9 This revised interpretation is consistent with all the data and with basic thermodynamic  
10 principles, and implies that the nicotinamide-flavin interaction consistently formed in the  
11 aromatic placeholder mutants is in fact productive and relevant for understanding catalysis.  
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### 14 15 16 **Crystal structures of Y316S and Y316A NADP(H) complexes** 17

18 Co-crystallization of both FNR variants with nicotinamide yielded crystals that grew readily  
19 and could be soaked with NADP<sup>+</sup> or NADPH to obtain those complexes. Interestingly, the  
20 variants crystallized in two space groups, both the P<sub>3</sub><sub>2</sub>2<sub>1</sub> form seen for wild-type protein [9][9]  
21 and a new P<sub>3</sub><sub>1</sub>2<sub>1</sub> crystal form that had remarkably similar unit cell dimensions and related crystal  
22 packing. Here, we report a set of eight refined crystal structures of complexes that represent the  
23 Y316S and Y316A variants in complex with three ligands (NADP<sup>+</sup>, NADPH, and nicotinamide)  
24 with most structures refined at between 1.35 Å and 1.6 Å resolution (Table 2) and having very  
25 well defined active site electron density (FigureFigures 7-and-S2). As the corresponding  
26 complexes of Y316S and Y316A largely show equivalent features, ~~in the main text~~ we focus  
27 mainly on the highest resolution set of structures that provide unique information, ~~as well as~~  
28 ~~additional structures that reinforce the conclusions are in the supplementary material.~~  
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39 To check whether the complexes in the crystal represent those that form in solution, we  
40 obtained single-crystal absorption spectra [35] during data collection for a pair of Y316A  
41 crystals soaked with NADP<sup>+</sup> and NADPH.~~[32] during data collection for a pair of Y316A~~  
42 ~~crystals soaked with NADP<sup>+</sup> and NADPH.~~ These spectra qualitatively match those obtained in  
43 solution and indicate that even though the X-ray beam does cause reduction of the FAD, this  
44 does not undermine the relevance of the structures (Figure 7). Also, since Y316S and Y316A  
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both predominantly form CTC-2 after an NADPH soak (*e.g.* Figure 5B,C), we have modeled the NADPH soak structures as an  $\text{NADP}^+:\text{FADH}_2$  complex rather than NADPH:FAD.

In the following sections, we first provide a basic description of the complexes and then focus on three detailed aspects of complexes that provide evidence of compression in the active site for enhancing catalysis. In our structural comparisons, the unliganded wild-type structure we use is a 1.05 Å resolution structure refined much earlier (PDB entry 3LO8; released in 2010) and used in tests of new refinement strategies [36],[33]. As the statistics for this structure have not yet been described in the literature, they are included in Table 2.

*Overview of the NADP(H) complexes.* Globally, all Y316S and Y316A complexes, independent of space group, are very similar to the wild-type structure with rms  $C\alpha$  deviations between 0.1 Å and 0.4 Å. The broad features of NADP(H) binding match those that have been elaborated elsewhere [4, 6, 10, 12][4, 6, 10, 12], interacting mostly with the NADP-binding domain. Briefly, the 2'-phosphate anchor hydrogen bonds with the side chains of Ser237, Arg238, Lys247, and Tyr249, and the adenine is sandwiched between Tyr249 and Leu276. The 5'-phosphoryl group is largely exposed and the nicotinamide-side phosphoryl group hydrogen bonds with the Thr175 N and the Arg114 guanidinium group. The nicotinamide is surrounded by a triad of key side-chains, with Ser94 and Glu314 hydrogen bonding with the nicotinamide carboxamide nitrogen and Cys274 and Ser94 close to the C4 atom (Figure 7A). In the higher resolution structures of Y316S, an alternate conformation for the nicotinamide ribose is also observed (Figure S2). In both  $\text{Y316S}_{\text{NADP}^+}$  and  $\text{Y316S}_{\text{NADPH}}$  soaks in both space groups, the nicotinamide ribose is modeled as 60% 2'-endo and 40% 3'-endo. In each Y316A structure, the nicotinamide ribose adopts a single conformation, but in  $\text{Y316A}_{\text{NADP}^+}$  it is 2'-endo and in  $\text{Y316A}_{\text{NADPH}}$  it is 3'-endo. The reason for these conformational differences is not readily

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9 apparent but their presence suggests that they are the result of real but subtle differences in  
10 nicotinamide binding.

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12 *The complexes with free nicotinamide.* Surprisingly, the binding mode seen in nicotinamide  
13 soaks differ for Y316A and Y316S (Figure 3C). In Y316A<sub>nic</sub>, the nicotinamide orientation and  
14 hydrogen bonding roughly match how the NADP(H) nicotinamide binds. However, in Y316S<sub>nic</sub>,  
15 the nicotinamide binds in a flipped orientation that is stabilized by a hydrogen bond between  
16 Ser316-O $\gamma$  and nicotinamide-N1 as well as hydrogen bonds of the nicotinamide hydroxyamide  
17 with Ser94-O $\gamma$  and Thr 175-O $\gamma$  and O atoms (Figure 3C). Since this binding mode involves  
18 Ser316 as a hydrogen-bond partner, we conclude it is an artifact of the Y316S mutation. Also,  
19 the existence of the nicotinamide N1 to Ser316-O $\gamma$  hydrogen bond in this complex means that  
20 neither NADP(H) nor even N1-methyl-nicotinamide could adopt it. This provides a satisfying  
21 explanation for why nicotinamide binding to Y316S yields different spectral changes than  
22 NAD(P)<sup>+</sup>, whereas methyl-nicotinamide binds less tightly, but with spectral changes that match  
23 those of NAD(P)<sup>+</sup> (Figure 3B). It also serves as a reminder of the need to be cautious in inferring  
24 the binding mode of a molecule of interest based on that of an analog.

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37 *The nicotinamide-flavin interactions in the NADP<sup>+</sup>/NADPH complexes.* While the overall  
38 structures of the FNR variants are virtually superimposable, subtle differences exist in the details  
39 of nicotinamide binding seen in the NADP<sup>+</sup> and NADPH soaks, and having the structures for the  
40 two mutants allows us to gain confidence about which variations are reliably due to the  
41 difference between the active site redox state rather than the specific mutation or to experimental  
42 uncertainty. In terms of the active site, the difference between the structures is a single H-atom:  
43 the NADP<sup>+</sup> soaks with oxidized flavin and nicotinamide have one H-atom on the nicotinamide  
44 C4 and no H-atom on the flavin N5, and the NADPH soaks to produce CTC-2 have one H-atom  
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9 on the nicotinamide C4 as well as one H-atom on the flavin N5. (The CTC-1 complex would  
10 have the same number of H-atoms but distributed with two on the nicotinamide C4-atom and  
11 none on the flavin N5-atom.) As described in the following paragraphs, the high resolution  
12 structures provide evidence that the single additional H-atom increases active site crowding and  
13 a directional compression that we propose is a key promotor of catalysis.  
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18 A first line of evidence is provided by the temperature factors of the C4-atom of NADP<sup>+</sup>  
19 compared to NADPH. In our earlier work on pea FNR [6][6], a higher conformational freedom  
20 of the NADP<sup>+</sup> C4-atom was indicated by weaker electron density (and higher B-factors) for this  
21 atom in NADP<sup>+</sup> compared to NADPH; but at 1.8 Å resolution, this was not conclusive. Our  
22 higher resolution analyses here similarly show the additional mobility of the C4 atom of NADP<sup>+</sup>  
23 but also provide refined  $\sigma^2$ -anisotropic B-factors that give a more complete description of  
24 dynamics than do isotropic B-factors. The anisotropic B-factors very clearly show that the  
25 increased movement for the C4 atom of NADP<sup>+</sup> is perpendicular to the plane of the ring and  
26 towards N5 of the FAD (Figure 8A). This type of motion exactly corresponds to motions that  
27 would allow the nicotinamide ring to forms a boat-like conformation that calculations show will  
28 promote efficient hydride transfer [37][34].  
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39 A second observation of crowding derives from a visible distortion of the FAD  
40 isoalloxazine. In wild-type FNR, the isoalloxazine group appears to be pushed slightly by the  
41 Tyr316 side chain so that it is not coplanar with the N10-C1 ribose bond, but is about 3.5° non-  
42 planar (Figure 8B). In all FNR variant structures soaked with NADP<sup>+</sup>, this non-planarity  
43 increases to ~10°, and in the structures soaked with NADPH it increases further to ~14° (Figure  
44 8B). These deviations from planarity imply that the nicotinamide displacing the Tyr316 side  
45 chain leads to a more tightly packed active site and that the additional H-atom in the active site  
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9 present in the NADPH soaks increases crowding even more, with the isoalloxazine bending to  
10 relieve apparent pressure. It should be noted that the highly crowded complex seen in the  
11 NADPH soak is the one that is present during normal catalysis.  
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14 A third observation that completes the view of the tightly packed active site of FNR are the  
15 interaction distances of the nicotinamide C4 and H4 atoms that are involved in hydride transfer  
16 (Figure 8C). In all structures, these atoms are sandwiched between the flavin N5 atom in front  
17 and the Cys274-S $\gamma$  sulfhydryl behind, where it can act as a backstop. In comparing the  
18 Y316S<sub>NADP+</sub> complex to the Y316S<sub>NADPH</sub> complex which has an additional H-atom on the flavin  
19 N5-atom, the C4...N5 distance increases by  $\sim 0.3$  Å while the C4H... S $\gamma$  distance decreases by  
20  $\sim 0.1$  Å (Figure 8C). Taking into account the Y316A complexes (Figure 8C), the consistent  
21 increase of the C4...N5 distance in the NADPH complex leads us to conclude that Cys274 acts  
22 as a firm backstop while the isoalloxazine bends away from the nicotinamide by an additional  $\sim 4^\circ$   
23 in order to increase the distance between the flavin and nicotinamide and relieve the pressure due  
24 to the presence of the extra H-atom.  
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35 These observations lead us to conclude that the active site is experiencing compression,  
36 distorting the FAD and pushing the substrates together to enhance catalysis by aligning the  
37 substrates in an optimal position and, as has been noted by others, potentially promoting a  
38 quantum tunneling mechanism of hydride transfer [26][31].  
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#### 45 **Insights into factors promoting hydride transfer in FNRs**

46 Taken along with ~~our~~the reanalysis of previous stopped-flow kinetics work ~~that~~  
47 ~~called calling into question~~ the relevance of aromatic placeholder FNR mutants ~~into question~~,  
48 these in-solution spectroscopy, single-crystal spectroscopy, and high-resolution structures  
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9 provide insights into hydride transfer in FNRs using variants which have near wild-type hydride-  
10 transfer kinetics, but importantly, allow us to capture the productive binding mode of NADP(H)  
11 in the active site. ~~Such productive complexes have not been possible to capture in, something~~  
12 ~~which has proved impossible using~~ any wild-type member of the FNR superfamily ~~with (e.g. [5,~~  
13 ~~17-20]). This includes superfamily enzymes such as those already mentioned that have an~~  
14 aromatic ~~placeholder~~ residue ~~(e.g. [5, 17-20]) or that instead blocking the nicotinamide binding~~  
15 ~~site, as well as those found to have a C-terminal peptide extension as an alternate way to block~~  
16 ~~nicotinamide binding [3, 38][3, 35]. They have also been challenging to obtain for superfamily~~  
17 ~~enzymes that have or a shifted domain-domain interaction [39, 40][36, 37] instead of an aromatic~~  
18 ~~blocking residue, but for one of these enzymes, cytochrome b5 reductase, anaerobic co-~~  
19 ~~crystallization has yielded a productive, wild type complex with NADH in which the~~  
20 ~~nicotinamide-flavin interaction geometry is much like that seen here [41]. that appear to be~~  
21 ~~alternate ways to block nicotinamide binding.~~

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33 The high-resolution structures reported here illustrate the role of specific anisotropic  
34 motions as well as active site compression as catalytic strategies that promote hydride transfer.  
35 Key evidences are the anisotropic motion of the C4 atom of NADP<sup>+</sup>, that indicates boat-like  
36 perturbations in conformation are a preferred mode of vibrational freedom, as well as a strong  
37 decrease in the amplitude of that motion in the NADPH complex. Additionally, there is an  
38 incrementally increasing deviation from planarity at the flavin N10 atom in wild type,  
39 Y316<sub>NADP<sup>+</sup></sub>, and Y316<sub>NADPH</sub> indicating increased crowding and pressure associated with  
40 oxidized nicotinamide binding and even more so with reduced nicotinamide binding that is being  
41 relieved to some extent by bending of the flavin. Interestingly, a very similar flavin non-planarity  
42 was seen in the cytochrome b5 reductase-NADH complex (see Figure 3a of [41]). Finally,  
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9 comparison of Y316S<sub>NADP+</sub> and Y316S<sub>NADPH</sub> reveal close sub van der Waals contact distances  
10 consistent with a tightly packed active site. The central role of Cys274 in these interactions  
11 provides a rationale for its conservation across the whole superfamily. While steric compression  
12 has occasionally been noted as a key factor promoting catalysis for other enzymes [42-45][38-  
13 41], there are a limited number of studies which provide direct structural evidence for this [46-  
14 50][42-46], as this typically requires atomic or near-atomic resolution crystal structures. Also,  
15 that the Y316S and Y316A variants of corn root FNR, like similar mutants of other FNR  
16 superfamily members [22, 31, 34][22, 27, 30], show much less stabilization of the FAD  
17 semiquinone, is consistent with an earlier proposal [31][27] that the aromatic side chain is not  
18 solely a passive placeholder for the nicotinamide group, but, through its stacking interaction with  
19 the flavin, is an active agent that stabilizes the semiquinone form of the flavin to enhance the  
20 one-electron transfers required for these dehydrogenase-electron transferases.  
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### 33 Extrapolation of the results to the FNR superfamily members such as NOX enzymes

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35 Clarifying the relevance of the NADP<sup>+</sup> binding mode seen in the Tyr316 mutants is not just  
36 of interest for understanding catalysis for the whole superfamily, but can guide further studies of  
37 these enzymes as well as the generation of superfamily member variants that can be valuable  
38 tools. For instance, for the NOX enzymes that play crucial roles in the production of superoxide  
39 and hydrogen peroxide for diverse biological processes, the FNR-like module is at the C-  
40 terminus of a membrane-bound flavocytochrome catalytic subunit that has been relatively  
41 difficult to study [14][14]. Although a NOX structure has long eluded The only structural  
42 biologists studies of it are a crystal structure of the FNR-like module of NOX5 from  
43 Cylindrospermum stagnale (referred to as the NOX2-NADPH-dehydrogenase-binding domain)  
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~~was solved while this work was under review [30] as part of the Protein Structure Initiative (Figure 9). Sequence alignments of FNR with human NOXs aided by the *C. stagnale* NOX5 that structure (Figure 9A9) show that the NOX isozymes conserve many NADP<sup>±</sup> binding residues including the nicotinamide-interacting residues equivalent to corn root FNR Ser94, Cys274, Glu314 and Tyr316 (Phe693 in *C. stagnale* NOX5).~~

~~The NOX5 FNR-like module itself was unstable and crystals were only obtained for a construct with a “hyperstabilizing” C-terminal extension that included a Trp two residues after Phe693. Notably, in this structure, the expected natural C-terminal aromatic placeholder (Phe693) does not stack against the isoalloxazine, and, because the anticipated fifth  $\beta$ -strand of the NADP<sup>+</sup>-binding domain is perturbed and moved away from the protein core, the Trp residue is able to stack against the isoalloxazine ring instead (Figure 9B). However, if the NOX5 chain followed the path of the  $\beta$ -strand seen in other FNR superfamily members, Phe693 could stack against the isoalloxazine and we propose that this is what occurs in the native enzyme. Magnani et al [30] noted that the isolated dehydrogenase domain is deregulated and predicted that the normal role of the strictly conserved C-terminal aromatic residue as a Phe570 in NOX2).~~

~~We would “emerge only in the context of a full-length protein.” We agree and further suggest that the instability of the isolated dehydrogenase domain and the loose association of the C-terminal segment (including Phe693) is related to the missing cytochrome transmembrane domain and associated lipid bilayer that interact with this surface of the dehydrogenase domain in a full-length complex (see Fig 5A of [30]).~~

~~Consistent with the weak association observed for Phe693 with the protein core is in the isolated dehydrogenase domain, a Phe693Ser mutant had higher than wild-type activity and the stabilizing C-terminal extension mutant (adding the Trp695) dropped activity by 5-fold [30].~~

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9 Nevertheless, we predict that in the context of a full-length membrane-bound NOX, Phe693 will  
10 act as an aromatic placeholder residue and that Ser or Ala Phe570Ser and Phe570Ala (F570A)  
11 mutants ~~will of the aromatic placeholder residue should~~ bind both NADPH and NADH more  
12 tightly. ~~Such and also be more active with NADH, and such engineered~~ mutants could provide  
13 ~~be very~~ useful ~~for providing a~~ handle for purification (through the tight binding of NADP<sup>+</sup>) or  
14 ~~be a useful tool~~ for probing/controlling ~~the their~~ physiological roles of NOXs. A, potentially by  
15 ~~making a~~ solely NADH responsive version could potentially be made through additional  
16 mutations, such as the equivalent of FNR Ser237 to Asp (Ser237Asp (see Figure 9), as this  
17 position has been shown that would be predicted to discriminate against binding the 2'-  
18 phosphoryl group in multiple superfamily members [51-54],[47-50]. Interestingly, a full-length  
19 human NOX2 C-terminal PheF570A mutant (F570A) was characterized in 1998 and found to  
20 retain ~50% of the wild-type activity ~~[55][54]~~ rather than dropping 300-800 fold in activity as  
21 had been seen for the equivalent pea FNR mutant, and so it was not studied further. However,  
22 this study predated the knowledge that ~~activity loss of~~ the equivalent FNR mutant lost  
23 activity was due to the slow dissociation of NADP<sup>+</sup> and ~~also that the mutant~~ gained activity with  
24 NADH ~~[9, 21][9]~~, so those qualities of the variant were never characterized. In retrospect, given  
25 that the NOX2 K<sub>d</sub> for NADP<sup>+</sup> is ~40 μM (over ~~10400~~ fold higher than is typical for FNRs that of  
26 ~~corn root FNR~~) and the turnover number is slower, it would not be surprising if the expected  
27 enhanced binding of NADP<sup>+</sup> in the F570A variant would not be enough to make its dissociation  
28 highly rate limiting. Indeed, for NO synthase the equivalent mutation only showed about a 3-fold  
29 decrease in steady-state turnover, yet still showed a 50-fold change in specificity ~~[22, 34][22,~~  
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## MATERIALS AND METHODS

### Production of Recombinant Corn Root FNR and its Variants

Plasmids for the bacterial expression of the Y316A and Y316S proteins were generated from the pETrFNR2 [9][8] using the QuikChange II Site-Directed Mutagenesis kit (Agilent) and two appropriate oligonucleotide couples following the manufacturer directions. The wild-type and variant proteins were produced in *E. coli* HMS174(DE3) by induction with 0.1 mM IPTG for 4 h at 30 °C, and purified through a procedure similar to that previously reported for the wild-type protein [9] using an ÄKTA FPLC (GE Healthcare) apparatus.[8] using an ÄKTA FPLC (GE Healthcare) apparatus. Briefly, the crude cell lysate was brought to 40%  $\text{NH}_4\text{SO}_4$  and centrifuged. The supernatant was applied to a Sepharose 4B column (GE Healthcare), eluted with appropriate buffer and precipitated with 75%  $(\text{NH}_4)_2\text{SO}_4$ . In the case of the protein variants, the pellet was resuspended in 40%  $(\text{NH}_4)_2\text{SO}_4$  and chromatographed on a butyl Sepharose column (GE Healthcare) with a descending salt concentration gradient. The omission of this step prevented the adsorption of the FNR forms on the next cation exchange resin. After desalting, the sample was loaded on an SP-Sepharose HP column (GE Healthcare) and eluted through a NaCl concentration gradient. Notably, the visible spectra of both mutant forms, but not the wild-type enzyme, underwent a blue shift during the ion exchange step, suggesting release of some unidentified bound ligand at that stage. The resulting proteins were homogenous as judged by SDS-PAGE. The FNR forms were concentrated to ~25 mg/mL and stored at -20 °C in 10 mM HEPES, pH 7.0.

### Spectral Analyses and Ligand Binding

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9 All spectrophotometric measurements and steady-state enzyme kinetics were performed  
10 using an 8453 diode-array spectrophotometer (Agilent). Titration with NADP<sup>+</sup>, NAD<sup>+</sup>, phenol,  
11 nicotinamide and N-methyl-nicotinamide of the enzyme forms in their oxidized state were  
12 performed in 10 mM Tris-HCl, pH 7.7, at 15 °C.  
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### 17 18 **Photoreductions, Anaerobic Titrations and Activity Assays** 19

20 Flavin photoreduction experiments were carried out both in the absence and in the presence  
21 of NADP<sup>+</sup> or NAD<sup>+</sup> in anaerobic cuvettes in 10 mM HEPES-NaOH, pH 7.0, at 15 °C, according  
22 to the procedure described elsewhere [9].  
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24 ~~Flavin photoreduction experiments were carried out both in the absence and in the presence~~  
25 ~~of NADP<sup>+</sup> or NAD<sup>+</sup> in anaerobic cuvettes in 10 mM HEPES-NaOH, pH 7.0, at 15 °C, according~~  
26 ~~to the procedure described elsewhere [8].~~  
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31 Anaerobic titrations with NADPH of 14-18 μM wild-type, Y316A or Y316S FNR variants  
32 were performed on 1.2 mL samples in 50 mM HEPES-NaOH buffer, pH 7.0, at 15 °C, in a  
33 sealed cuvette. After recording the spectrum of the oxidized enzyme, five successive additions of  
34 anaerobic NADPH were made leading to ligand concentrations of 29 μM, 39 μM, 76 μM, 182  
35 μM, and 1.9 mM for wild-type titrations, 23 μM, 45 μM, 90 μM, 210 μM, and 2.2 mM for FNR  
36 Y316A titrations, and 24 μM, 48 μM, 95 μM, 225 μM, and 2.3 mM for FNR Y316S titrations,  
37 respectively. In each case, the volume after the final addition of NADPH was 1.4 mL. For each  
38 FNR form, the cuvette was then opened to air, and the spectrum of the mixture (kept at 15 °C)  
39 was monitored at over time until all the NADPH was oxidized. For technical reasons, the spectra  
40 for wild-type FNR during O<sub>2</sub> turnover was produced from a separate sample also prepared with  
41 2.4 mM NADPH.  
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NADPH—K<sub>3</sub>Fe(CN)<sub>6</sub> reductase activity assays were performed in 100 mM Tris-HCl, pH 8, at 25 °C, at different concentrations of both substrates, as reported elsewhere [56],[52]. As needed, nicotinamide was included in the assays at concentrations ranging from 50 to 800 mM.

### Reanalysis of Stopped-Flow Results Reported for *Anabaena* FNR

Spectra (Figure 3 in [26]) were enlarged by ~345% and printed.[31] were enlarged by ~345% and printed. The height of each peak at ~460 nm corresponding to the flavin oxidation state was measured in cm from the baseline of the spectra (-0.025 AU). The peak height at time 0 for FNR<sub>red</sub> with NADP<sup>+</sup> (corresponding to fully reduced flavin) was taken as a baseline and used to normalize the measurements by subtracting this height from the other peak heights of the corresponding variant. Whereas the final spectra (0.2547 s) for wild-type FNR for both FNR<sub>red</sub> with NADP<sup>+</sup> and FNR<sub>ox</sub> with NADPH matched in both peak height and shape, the final spectra (0.2547 s) for Y303S FNR varied slightly in peak height but both correspond to achieved equilibrium so were normalized to each other by subtracting the difference between the final peak heights from the measured peak heights for Y303S FNR<sub>red</sub> with NADP<sup>+</sup>.

Taking the peak height at time 0 for FNR<sub>ox</sub> with NADPH to correspond to fully oxidized flavin, the percentage of oxidized and reduced flavin and corresponding concentration of product (using the provided concentrations of 25 μM FNR and 125 μM NADP(H)) were calculated. A plot of concentration of product (μM) versus time (s) were fitted using GraFit 5 (Erithacus Software Limited) with a single-exponential decay equation to estimate the apparent rate constants of the hydride-transfer reactions.



## Crystallization and Structure Determinations

Initial co-crystallization trials of the corn root FNR variants with NADP<sup>+</sup> used similar conditions to those used for the wild-type enzyme [9][9] but yielded only ill-formed small crystals that grew very slowly. Solving their structure revealed that they contained a bound nicotinamide moiety rather than NADP<sup>+</sup>, so co-crystallization with nicotinamide was tried, and crystals grew readily. The best crystallizations were at room temperature in hanging drops formed by mixing equal volumes of the protein stock (Y316F: 20.2 mg/mL; Y316A: 7.3 mg/mL; Y316S: 10.8 mg/mL; all in 50 mM Tris-HCl, pH 7.4) with a reservoir solution containing 22-24% PEG 8000, 0.1 M sodium cacodylate (pH 6-7), 0.18-0.22 M magnesium acetate, and 100 mM nicotinamide. Typical crystals were ~0.3 mm on each side and grew within 1 week. Both FNR variants formed many prism-shaped crystals belonging to the space group P3<sub>2</sub>21 (same as wild type; [9][9]) and fewer hexagon-shaped crystals belonging to space group P3<sub>1</sub>21. Soaks of the nicotinamide-bound crystals were done aerobically at room temperature with either 10 mM NADP<sup>+</sup> or 10 mM NADPH for 1 h to obtain desired complexes before being flash frozen.

*Data collections.* For data collection, crystals were pulled through oil and flash-frozen in liquid nitrogen. In-house data were collected at 140 K on a Rigaku RU300 Cu-K $\alpha$  rotating anode X-ray source running at 50 kV and 100 mA equipped with a Raxis IV image plate detector. Synchrotron data were collected at beamline 5.0.1 or 5.0.3 at the Advanced Light Source (Lawrence Berkeley National Laboratory, Berkeley, CA) at 100 K. Oscillation images were collected with  $\Delta\phi = 1^\circ$  and were processed using Denzo and Scalepack [57][53] or iMosflm [58][54] and SCALA [59][55]. Unit cell parameters and data reduction statistics for each FNR variant are given in Table 2. In addition, data sets with  $\Delta\phi = 0.5^\circ$  and simultaneous single crystal

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visible absorption spectra were collected by Dr. Allen Orville at a National Synchrotron Light Source beamline as described in [35][32].

*Structure solutions and refinements.* All P3<sub>2</sub>21 FNR variant crystals were isomorphous with the published wild-type corn root FNR structure (PDB code 3LO8 [36][33]), and this was used as the starting model for the first refinement, with additional refinements built on partially refined models of the most similar structure. The P3<sub>1</sub>21 crystal form was trivial to solve by molecular replacement and these structures were then similarly refined. The same 10% of data were set aside for cross-validation [60][56] for each structure as had been set aside for the ~1 Å resolution refinement of the wild-type structure. Refinements were carried out by a number of researchers over many years using various versions of REFMAC [61][57] with Coot [62] used for manual fitting and Molprobity [63][58] used for manual fitting and Molprobity [59] used to help identify model problems. During iterative manual rebuilding, water molecules were added in Coot using standard criteria (>1  $\rho_{\text{rms}}$  intensity in the 2F<sub>o</sub>-F<sub>c</sub> map, >2.4 Å distance from nearest contact, no *B*-factors >80 Å<sup>2</sup>). For making a consistent set of structures for publication, a set of final refinements of each structure were done by the lead author using Phenix [64][60]. During this stage, we decided to model all NADPH soaks as having an oxidized nicotinamide (*i.e.* NADP<sup>+</sup>) and a reduced flavin since the spectra show that CTC-2 is present at higher amounts than is CTC-1. For structures refined with anisotropic *B*-factors (the P3<sub>2</sub>21 Y316S:NADPH, Y316S:NADP<sup>+</sup>, and Y316S:nicotinamide structures), no anisotropic restraints were used for the ligand in the final round of refinement to ensure the anisotropic *B*-factors of the nicotinamide atoms were based as much as possible on the diffraction data. Refinement statistics for all models are shown in Tables 2 and S1. Structural overlays were performed using the protein structure visualization software PyMOL [65][64].

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9 Also, in order to obtain a reliable representation of active site distances for Y316S<sub>NADPH</sub>,  
10 Y316S<sub>NADP+</sub> (P3<sub>2</sub>21 and P3<sub>1</sub>21), Y316A<sub>NADPH</sub>, and Y316A<sub>NADP+</sub> structures, ten different starting  
11 models for each were generated using the “shake” algorithm of Phenix with the setting  
12 “modify.sites.shake = 0.5”. This level of coordination disruption resulted in starting R/R<sub>free</sub>  
13 values of ~40%. Each of these models was re-refined and the distances between Cys274 Sγ and  
14 NADP<sup>+</sup> C4H and between NADP<sup>+</sup> C4 and FAD N5 were measured. The average distances and  
15 standard deviations for each set were calculated and reported.  
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22 *Accession Numbers.* Coordinates and structure factors for the wild-type corn root FNR,  
23 Y316S:nicotinamide (P3<sub>2</sub>21), Y316S:nicotinamide (P3<sub>1</sub>21), Y316S:NADP<sup>+</sup> (P3<sub>2</sub>21),  
24 Y316S:NADP<sup>+</sup> (P3<sub>1</sub>21), Y316S:NADPH (P3<sub>2</sub>21), Y316A:nicotinamide (P3<sub>2</sub>21),  
25 Y316A:NADP<sup>+</sup> (P3<sub>2</sub>21), Y316A:NADPH (P3<sub>2</sub>21), Y316F (P3<sub>2</sub>21), and Y316F:NADP<sup>+</sup> (P3<sub>1</sub>21)  
26 models have been deposited in the Protein Data Bank with accession numbers 3LO8, 5VW4,  
27 5VW9, 5VW3, 5VW8, 5VW2, 5VW5, 5VW6, 5VW7, 5VWA, and 5VWB, respectively.  
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**Author Contributions.** KMK analyzed data and wrote the paper; RAC planned and performed experiments and analyzed data; VP, ARH, and RF planned and performed experiments; GZ planned experiments; AA planned and performed experiments, analyzed data, and wrote the paper; PAK planned experiments, analyzed data and wrote the paper.

For Review Only

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## TABLES

**Table 1.** Effect of the Y316S replacement on the kinetic parameters of the NADPH—K<sub>3</sub>Fe(CN)<sub>6</sub> reductase reaction catalyzed by FNR.<sup>a</sup>

FNR form	$k_{\text{cat}}$ s <sup>-1</sup>	$K_{\text{m}}^{\text{NADPH}}$ μM	$k_{\text{cat}}/K_{\text{m}}^{\text{NADPH}}$ s <sup>-1</sup> μM <sup>-1</sup>
Wild type <sup>b</sup>	520 ± 10	12 ± 1	43 ± 4
Y316S	3 ± 0.3	1.4 ± 0.3	2.1 ± 0.5
Y316S, 0.8 M nicotinamide	280 ± 18	45 ± 10	6 ± 1

<sup>a</sup>Initial rate data were measured as moles of ferricyanide reduced per second.<sup>b</sup>Data on wild type FNR, taken from [9][9], are reported here for comparison.

**Table 2.** Data and refinement statistics for FNR variant structures<sup>a</sup>

Ligand soaked Space group	wt		Y316S		Y316A		Y316A		Y316A	
	None P3 <sub>2</sub> 21	Nicotinamide P3 <sub>2</sub> 21	Nicotinamide P3 <sub>1</sub> 21	NADP+ P3 <sub>2</sub> 21	NADP+ P3 <sub>1</sub> 21	NADPH P3 <sub>2</sub> 21	Nicotinamide P3 <sub>2</sub> 21	NADP+ P3 <sub>2</sub> 21	NADPH P3 <sub>2</sub> 21	NADPH P3 <sub>2</sub> 21
<i>Data Statistics</i>										
Wavelength (Å)	1.00	1.00	1.54	1.00	1.54	1.00	1.54	1.00	1.00	1.00
Unit cell <i>a, c</i> axes (Å)	59.1, 186.7	58.9, 184.8	58.9, 184.4	58.8, 187.5	58.9, 185.0	58.9, 187.3	59.4, 187.5	58.7, 186.7	59.2, 188.0	59.2, 188.0
Resolution (Å)	50-1.05 (1.07-1.05)	51-1.35 (1.42-1.35)	12-1.90 (1.97-1.90)	51-1.45 (1.53-1.45)	40-1.80 (1.90-1.80)	51-1.45 (1.53-1.45)	26-1.95 (2.06-1.95)	49-1.50 (1.55-1.50)	49-1.60 (1.66-1.60)	49-1.60 (1.66-1.60)
Unique reflections	177114 (8768)	75008 (9295)	29752 (2603)	65831 (8739)	35032 (5039)	66408 (8907)	24769 (1325)	44421 (1371)	50821 (4336)	50821 (4336)
Multiplicity	5.2 (2.9)	5.8 (2.6)	8.2 (3.8)	9.6 (2.5)	7.3 (3.9)	8.4 (2.5)	9.1 (2.3)	1.7 (1.1)	4.0 (3.2)	4.0 (3.2)
Average <i>I</i> / $\sigma$	57.0 (3.4)	6.2 (1.7)	19.4 (6.9)	14.7 (3.0)	6.5 (2.2)	12.7 (2.7)	34.3 (9.6)	10.4 (0.82)	24.5 (6.0)	24.5 (6.0)
R <sub>meas</sub> (%)	9.3 (50.6) <sup>b</sup>	30.2 (46.9)	4.0 (15.3) <sup>b</sup>	9.5 (29.0)	33.5 (53.2)	13.0 (39.5)	4.4 (6.4)	5.4 (71.1) <sup>b</sup>	2.8 (16.5) <sup>b</sup>	2.8 (16.5) <sup>b</sup>
Completeness (%)	99.8 (99.5)	91.0 (78.4)	97.6 (86.6)	97.9 (91.7)	99.9 (99.9)	98.1 (92.1)	85.8 (32.5)	72.9 (23.0)	98.6 (86.4)	98.6 (86.4)
<i>Refinement Statistics</i>										
Refinement method	Aniso	Aniso	TLS	Aniso	TLS	Aniso	TLS	TLS	TLS	TLS
Amino acid residues	302	309	309	309	309	309	309	309	309	309
Solvent atoms	646	<del>484</del> 486	<del>413</del> 414	491	<del>297</del> 300	500	556	451	<del>530</del> 531	<del>530</del> 531
Non-H atoms	3252	<del>3117</del> 3114	<del>3035</del> 3036	3160	<del>2923</del> 2926	3217	3165	2999	<del>3155</del> 3156	<del>3155</del> 3156
RMS bonds (Å)	0.017	0.009	0.009	0.008	0.010	0.008	0.010	0.009	<del>0.009</del> 0.008	0.009
RMS angles (°)	2.3	1.0	1.1	1.2	1.3	1.2	1.1	1.2	1.2	1.2
<B <sub>protein</sub> > (Å <sup>2</sup> )	21.7	14.68	<del>28.9</del> 20.8	20.9	<del>34.5</del> 35.4	21.3	17.8	26.7	24.0	24.0
<B <sub>FAD</sub> > (Å <sup>2</sup> )	15.1	10.43	<del>18.3</del> 17.9	16.6	<del>22.0</del> 21.7	16.7	13.1	21.1	18.53	18.53
<B <sub>NADP(H)</sub> > (Å <sup>2</sup> )	–	<del>13.8</del> 14.3	<del>22.7</del> 3	24.1	<del>28.3</del> 2	25.5	23.7	27.0	24.7	24.7
R <sub>work</sub> (%)	12.5	13.5	13.1	10.8	17.2	11.0	13.0	14.9	13.10	13.10
R <sub>free</sub> (%)	15.5	17.3	17.36	14.4	21.1	14.7	18.0	18.5	15.65	15.65
PDB code	3LO8	5VW4	5VW9	5VW3	5VW8	5VW2	5VW5	5VW6	5VW7	5VW7

<sup>a</sup> Numbers in parentheses are in the highest resolution shell<sup>b</sup> R<sub>merge</sub> reported in place of R<sub>meas</sub>

**FIGURE LEGENDS**

**Figure 1. Aromatic placeholder and nicotinamide binding in three FNR superfamily members.** (A) Stacking of aromatic placeholder side chain onto the flavin in wild-type forms of spinach FNR (PDB 1FNC; Tyr314 in forest green, FAD in orange), cytochrome P450 reductase (PDB 1AMO; Trp677 in green, FAD in dandelion), and NO synthase (PDB 1F20; Phe1395 in pale green, FAD in sand). (B) Mode of nicotinamide binding in the NADP<sup>±</sup> complexes of Y303S *Anabaena* FNR (PDB 2BSA; NADP<sup>±</sup> in blue), W677X cytochrome P450 reductase (PDB 1JA0; NADP<sup>±</sup> in cerulean), and Y303S pea FNR (1QFY; NADP<sup>±</sup> in seafoam); FAD colors as in panel A.

**Figure 2. Effect of the replacements of the Tyr316 residue on the visible absorption spectrum of FNR and rescue of the wild-type spectral features by phenol binding to the Y316S variant.** All spectra were determined in 50 mM Tris-HCl, pH 7.4 at 25 °C. (A) Extinction coefficients in the visible region of Y316S (solid line) and Y316A (dotted line) FNRs in comparison to that of wild-type enzyme (thin line). (B) Perturbation of the visible absorption spectrum of the Y316S variant (ca. 9 μM) induced by the presence of 150 mM phenol. The difference spectrum was rescaled (solid line) to match that expected for a theoretical enzyme concentration of 1 mM in order to be compared to the difference between the extinction coefficients of wild-type and Y316S FNR forms (dashed lines).

**Figure 3. Interaction of Y316S FNR with nicotinamide-containing ligands.** (A) Progress of the titrations of 10 μM Y316S FNR with NADP<sup>+</sup> (hollow circles), NAD<sup>+</sup> (filled circles), nicotinamide (hollow squares), and methyl-nicotinamide (filled squared). All titrations were

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9 performed in Tris-HCl, pH 7.7, at 15 °C. Data points and fitting curves were corrected to account  
10 for protein dilution. The logarithmic concentration scale was chosen to allow comparison of  
11 ligands displaying huge differences in affinity. **(B)** Computed difference spectra in the visible  
12 region of protein-ligand complexes. The difference between extinction coefficients of complexed  
13 and uncomplexed protein forms is shown for NADP<sup>+</sup> (solid line), NAD<sup>+</sup> (dashed line),  
14 nicotinamide (dash-dotted line), and methyl-nicotinamide (dotted line) are shown. **(C)** Overlay  
15 of Y316A<sub>nic</sub> (slate) and Y316S<sub>nic</sub> (hot pink) with nicotinamide bound, and as a reference, bound  
16 NADP<sup>+</sup> (ghosted grey). Hydrogen bonds to bound nicotinamide are shown (slate and hot pink  
17 dashed lines in the respective structures).  
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**Figure 4. Effect of the presence of NADP<sup>+</sup> on photoreduction of Y316S FNR.** Significant  
28 spectra recorded during the stepwise photoreduction of the FAD prosthetic group in the absence  
29 (black lines) and in the presence of a slight excess of NADP<sup>+</sup> (red lines). About 16 μM protein  
30 solutions in 10 mM HEPES, pH 7.0, were photoreduced at 15 °C in the presence of an EDTA-  
31 deazariboflavin system. Spectra corresponding to the maximal accumulation of semiquinone  
32 (dotted line) and charge-transfer complex (solid line) species are shown, and compared to those  
33 of the oxidized (thin lines) and fully reduced (dashed lines) respective enzyme mixtures.  
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**Figure 5. Spectra of wild-type FNR and its variants recorded during anaerobic titrations**  
43 **with NADPH and during NADPH-O<sub>2</sub> turnover. (Upper panels)** titrations of wild-type **(A)**,  
44 Y316S **(B)** and Y316A **(C)** FNRs showing the spectra of the free oxidized enzymes (thin black  
45 traces) and those recorded after the addition of ca. 1.2, 2.5, 5, 12, and 120-fold excess of  
46 NADPH (red, green, blue, magenta, and black bold traces, respectively). **(Lower panels)** Spectra  
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9 | ~~recorded~~ ~~reorder~~ at different reaction times after air was admitted into the cuvettes containing  
10 | the previously anaerobic solutions of the FNR forms in the presence of 120-fold excess NADPH.  
11 | The spectra of the free oxidized FNR forms (thin black traces) and those of the respective CT  
12 | complexes recorded in the presence of 1.2-fold NADPH excess (thin red traces) are reported for  
13 | comparison. **(D)** Spectra recorded after 15 s, 25 min, 35 min, 40 min, 45 min, and 60 min (red,  
14 | green, blue, magenta, lime, and black bold traces, respectively) turnover catalyzed by wild-type  
15 | FNR. **(E)** Spectra recorded after 2.5 min, 6 min, 11 min, 19 min, 46 min, and 100 min (red,  
16 | green, blue, magenta, lime, and black bold traces, respectively) turnover catalyzed by Y316S  
17 | FNR. **(F)** Spectra recorded after 1 min, 4 min, 53 min, 2.5 h, 3.5 h, and 17 h (red, green, blue,  
18 | magenta, lime, and black bold traces, respectively) turnover catalyzed by Y316A FNR.  
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30 | **Figure 6. Reanalysis of stopped-flow results reported for *Anabaena* FNR.** Data extrapolated  
31 | from [26] ~~Data extrapolated from [31] (as shown in Figure S1)~~ were fit to a single-exponential  
32 | decay. **(Upper panels)** Time-course of the approach to equilibrium upon mixing **(A)** wild-type  
33 | FNR<sub>red</sub> and NADP<sup>+</sup>, and **(B)** wild-type FNR<sub>ox</sub> and NADPH. Curve fitting yielded the  $k_{\text{obs}}$  values  
34 | of  $235 \pm 47 \text{ s}^{-1}$  and  $230 \pm 40 \text{ s}^{-1}$ , respectively, equivalent to each other within the error of the  
35 | analysis. Lans *et al.* [26] reported the rate constants for these reactions to be  $285 \text{ s}^{-1}$  and  $270 \text{ s}^{-1}$ ,  
36 | respectively. [31] ~~reported the rate constants for these reactions to be  $285 \text{ s}^{-1}$  and  $270 \text{ s}^{-1}$ ,~~  
37 | ~~respectively.~~ **(Lower panels)** Time-course of the approach to equilibrium upon mixing **(C)**  
38 | Y303S FNR<sub>red</sub> and NADP<sup>+</sup>, and **(D)** Y303S FNR<sub>ox</sub> and NADPH. Curve fitting yielded the  $k_{\text{obs}}$   
39 | values of  $165 \pm 16 \text{ s}^{-1}$  and  $145 \pm 20 \text{ s}^{-1}$ , respectively, equivalent to each other within the error of  
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9 **Figure 7. Active site environment of FNR (A)** A stereo view of the active site environment of  
10 FNR represented by the structure of Y316S<sub>NADP+</sub> with the 2F<sub>o</sub>-F<sub>c</sub> map shown at 3.0\* $\rho_{\text{rms}}$ . Key  
11 residues within the active site, Ser94, Cys274, and Glu314, which primarily interact with  
12 NADP(H), NADP(H), and FAD are shown along with hydrogen bonds (black dashed lines) and  
13 interaction distances (gray dashed lines). **(B)** Single crystal visible absorption spectra of Y316A  
14 FNR soaks with NADPH (right-hand panel) and NADP<sup>+</sup> (left-hand panel) and their change  
15 during data collection. Spectra collected corresponding to oscillation images 0, 10, 20, 50, 100  
16 and 360 go from light to dark shades of pink. These spectra qualitatively match those seen in  
17 solution in that the CTC-2 absorption band is seen for the crystals soaked in NADPH but not in  
18 those soaked in NADP<sup>+</sup>. The decrease in A<sub>460</sub> over time indicates reduction of the flavin in the  
19 X-ray beam during data collection. Over time the CTC-2 band does not disappear in the NADPH  
20 soaked crystal and does not appear in the NADP<sup>+</sup> soaked crystal so, we infer that at the cryo-  
21 temperatures of data collection, the flavin reduction is not leading to the same structural changes  
22 it would lead to in solution.  
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37 **Figure 8. Active site compression in corn root FNR as seen by mobility, covalent distortion,**  
38 **and interaction distances. (A)** Mobility of atoms in the active site of Y316S FNR. Atoms in the  
39 active sites of Y316S<sub>NADP+</sub> (orange) (i) and overlaid with Y316S<sub>NADPH</sub> (semi-transparent green)  
40 (ii) shown with ellipsoids, denoting the use of anisotropic B-factors. The nitrogen and oxygen  
41 atoms are shown in blue and red, respectively, in the complexes. The mobility of the C4 of  
42 NADP<sup>+</sup> is in the direction of the N5 of the FAD, forming a boat-like conformation which is  
43 favored for hydride transfer. **(B)** Active site compression in corn root FNR as seen by covalent  
44 distortion. The angle of distortion of the FAD moiety is shown in: (i) FAD in wild-type corn root  
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9 FNR (mauve), (ii) FAD in Y316S<sub>NADP+</sub> (orange) and the corresponding angles from Y316S<sub>NADP+</sub>  
10 (P3<sub>121</sub>, olive) and Y316A<sub>NADP+</sub> (salmon), (iii) FAD in Y316S<sub>NADPH</sub> (green) and the  
11 corresponding angle from Y316A<sub>NADPH</sub> (violet). (C) Active site interactions in FNR. Structural  
12 overlay of Y316S<sub>NADP+</sub> (orange) and Y316S<sub>NADPH</sub> (green) with relevant average distances shown  
13 in corresponding colors. The average distances from Y316S<sub>NADP+</sub> (P3<sub>121</sub>, olive), Y316A<sub>NADP+</sub>  
14 (salmon), and Y316A<sub>NADPH</sub> (violet) are also shown. The distances are averages from ten  
15 independent refinements of each complex (see Materials and Methods), and all standard  
16 deviations were < 0.02 Å. The C4H and N5H distance in Y316S<sub>NADPH</sub> is 3.0 Å (not labeled). If  
17 modeled as FADH<sub>2</sub> instead of FAD, the theoretical distance between C4H and N5H in  
18 Y316S<sub>NADP+</sub> would be 2.5 Å. Nitrogen, oxygen, and sulfur atoms colored blue, red, and yellow,  
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32 **Figure 9. Sequence alignment of corn root FNR with ~~human~~-NADPH oxidases. (A)**

33 Sequence alignment of segments of corn root FNR with human NOX1, NOX2, NOX3, NOX4,  
34 and NOX5 and C. stagnale NOX5. Select ~~with~~-FNR functional residues and similar aligned  
35 residues (in bold) have their-The function of FNR-conserved-residues denoted as are: aromatic  
36 placeholder (a), other catalytic center (c), 2'-phosphate2-phosphate (p) or other NADP  
37 interactions (n). Residue numbers are given for the structurally known ~~The numbering (first line)~~  
38 ~~corresponds to that of~~ corn root FNR and C. stagnale NOX5. ~~-(B)~~ Overlay of the C. stagnale  
39 NOX5 dehydrogenase domain (PDB 5O0X; pink with bound FAD and the natural C-terminal  
40 Phe693 and C-terminal extension Trp side chains in purple) and the Y316S<sub>NADP+</sub> structure  
41 (orange with bound FAD and NADPH in yellow). The anticipated fifth β-strand of the NADP<sup>+</sup>-  
42 binding domain of NOX5 which is perturbed and allows the Trp residue to stack against the  
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9 isoalloxazine ring is highlighted (purple trace) and an arrow indicates the direction we expect it  
10 to move in the full-length NOX5 context so that it will align with the rest of the  $\beta$ -sheet as does  
11 the fifth  $\beta$ -strand in FNR. The deposited structure of the NADPH binding domain of NOX2  
12 gp91(phox), determined through structural genomics efforts) (PDB 3A1F) but not yet described  
13 in the literature, also aligns well in a ; purple) and corn root FNR (FAD-binding domain in red,  
14 NADP-binding domain in green) with FAD (orange) and NADPH (blue) bound. In the structural  
15 overlay, FNR Ser237 which binds the 2' phosphate of NADP(H) aligns well with a conserved  
16 Thr in NOXs that could carry out the same role, and FNR Trp249 and Leu276 that stack against  
17 either side of the NADP adenine have roles that could reasonably be carried out by the conserved  
18 NOX Arg and Pro that align with them.  
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## High resolution studies of hydride transfer in the ferredoxin:NADP<sup>+</sup> reductase superfamily

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**Running Title:** Hydride Transfer in Ferredoxin:NADP<sup>+</sup> Reductase

**Database:** Structural data are available in the PDB database under the accession numbers **3LO8** (wild type), **5VW4** (Y316S:nicotinamide (P3<sub>2</sub>21)), **5VW9** (Y316S:nicotinamide (P3<sub>2</sub>21)), **5VW3** (Y316S:NADP<sup>+</sup> (P3<sub>2</sub>21)), **5VW8** (Y316S:NADP<sup>+</sup> (P3<sub>1</sub>21)), **5VW2** (Y316S:NADPH (P3<sub>2</sub>21)), **5VW5** (Y316A:nicotinamide (P3<sub>2</sub>21)), **5VW6** (Y316A:NADP<sup>+</sup> (P3<sub>2</sub>21)), **5VW7** (Y316A:NADPH (P3<sub>2</sub>21)), **5VWA** (Y316F (P3<sub>2</sub>21)), and **5VWB** (Y316F:NADP<sup>+</sup> (P3<sub>1</sub>21)).

**Abbreviations:** FNR – ferredoxin:NADP<sup>+</sup> reductase; NOX – NADPH oxidase; Fd – ferredoxin; CTC-1 – charge-transfer complex 1; CTC-2 – charge-transfer complex 2

**Enzyme Commission number:** ferredoxin:NADP<sup>+</sup> reductase – EC 1.18.1.2

**Keywords:** flavoenzyme, hydride transfer, enzyme mechanism, protein crystallography, NADPH oxidase

**ABSTRACT**

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5 Ferredoxin-NADP<sup>+</sup> reductase (FNR) is an FAD-containing enzyme best known for catalyzing the  
6 transfer of electrons from ferredoxin (Fd) to NADP<sup>+</sup> to make NADPH during photosynthesis. It is  
7 also the prototype for a broad enzyme superfamily, including the NADPH oxidases (NOXs) that  
8 all catalyze similar FAD-enabled electron transfers between NAD(P)H and one-electron carriers.  
9 Here we define further mechanistic details of the NAD(P)H ⇌ FAD hydride-transfer step of the  
10 reaction based on spectroscopic studies and high resolution (~1.5 Å) crystallographic views of the  
11 nicotinamide-flavin interaction in crystals of corn root FNR Tyr316Ser and Tyr316Ala variants  
12 soaked with either nicotinamide, NADP<sup>+</sup>, or NADPH. The spectra obtained from FNR crystal  
13 complexes match those seen in solution and the complexes reveal active site packing interactions  
14 and patterns of covalent distortion of the FAD that imply significant active site compression that  
15 would favor catalysis. Furthermore, anisotropic B-factors show that the mobility of the C4 atom  
16 of the nicotinamide in the FNR:NADP<sup>+</sup> complex has a directionality matching that expected for  
17 boat-like excursions of the nicotinamide ring thought to enhance hydride transfer. Arguments are  
18 made for the relevance of this binding mode to catalysis, and specific consideration is given to  
19 how the results extrapolate to provide insight to structure-function relations for the membrane-  
20 bound NOX enzymes for which little structural information has been available.  
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## INTRODUCTION

Plant ferredoxin NADP<sup>+</sup> oxidoreductases (FNR) are flavoenzymes that catalyze the reversible electron transfer between NADP(H) and the iron-sulfur protein ferredoxin (Fd). In photosynthesis, plastidic FNRs serve to catalyze electron transfer from Fd to NADP<sup>+</sup> via the FAD prosthetic group, and in non-photosynthetic tissues distinct FNRs use NADPH as an electron source for the reverse reaction [1-3]. Plant-type FNRs – as seen in three-dimensional structures of FNRs from cyanobacteria [4], spinach (leaf [5]), pea (leaf [6]), paprika (leaf [7]), and corn (leaf [8] and root [9]) – are highly similar and contain two distinct domains: one for binding FAD and one for binding NADP<sup>+</sup> [10]. Importantly, this two-domain FNR-fold is also the prototype for an enzyme superfamily that uses a bound FAD or FMN to transfer redox equivalents between the hydride carrying NAD(P)<sup>+</sup> cofactors and diverse one-electron carriers [10-12]. Included among the FNR superfamily are the NADPH oxidases (NOXs), a biomedically important group of membrane bound enzymes that produce superoxide or hydrogen peroxide as part of many signaling pathways as well as during the oxidative burst of macrophages that is a key part of our immune defenses (recently reviewed in [13-16]).

Structures of plant type FNRs and other superfamily members, such as nitric oxide synthase [17], cytochrome P450 reductase [18, 19], and phthalate dioxygenase reductase [20] showed that NADP(H) appears to bind non-productively to the wild-type enzymes because an aromatic side chain (a C-terminal Tyr residue in the case of plant type FNRs) sits in the site the nicotinamide ring must occupy for hydride transfer (Figure 1A). Deng *et al.* [6] resolved this mystery by showing for pea leaf FNR at 1.8 Å resolution that a mutant missing the aromatic placeholder residue (Tyr308Ser or Y308S) could bind NADP(H) tightly and in an apparently productive manner with the nicotinamide ring C4 atom adjacent to the FAD N5 atom and with a geometry reasonable for

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3 hydride transfer. The Y308S mutant bound its cofactor so tightly that NADP<sup>+</sup> co-purified with the  
4 enzyme. This mutant also showed an unexpected 500-fold change in cofactor specificity as the  
5 stronger binding of NAD(H) *increased* its steady-state turnover [21], but the stronger binding of  
6 NADP(H), *decreased* its steady state turnover, not because of impaired hydride transfer but  
7 because the NADP binding was so tight that  $k_{\text{off}}$  became rate-limiting [4, 21].  
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15 With the support of spectroscopic studies, it was concluded that the wild-type enzyme binds  
16 NADP(H) in a bipartite fashion in which the 2'-P-AMP binds strongly to anchor the cofactor to  
17 the enzyme and the thermodynamics of nicotinamide displacing the aromatic placeholder is such  
18 that the nicotinamide ring remains largely in a solvent exposed conformation (“nicotinamide-out”)  
19 and is in rapid equilibrium with a smaller population of molecules in which the aromatic  
20 placeholder swings out and the nicotinamide ring takes its place (“nicotinamide-in” conformation)  
21 to allow for hydride transfer. The Tyr to Ser mutant changes the binding thermodynamics so that  
22 the nicotinamide is ~100% “in”. Since that report, studies of aromatic blocker variants of other  
23 superfamily members have shown a consistent geometry for the “nicotinamide-in” nicotinamide-  
24 flavin interaction [4, 19] (Figure 1B) as well as changes from 50 to 1000-fold in NADP<sup>+</sup> vs NAD<sup>+</sup>  
25 specificity [4, 22-25].  
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41 Although there is considerable structural information about FNRs and FNR-like modules in  
42 related enzymes, the mechanism of hydride transfer is still not fully understood. For instance, the  
43 previous NADP(H) complexes of the pea FNR mutants gave indications that the C4 in the  
44 nicotinamide ring of NADP<sup>+</sup> was mobile in a way that could favor hydride transfer [6], but higher  
45 resolution data are needed to better define the details of this mobility. Also, it has been proposed  
46 that the geometry of the complex seen in the structures that have the aromatic placeholder mutated  
47 are artifacts that are not relevant to catalysis [26-28]. Here, we obtain further insight into the  
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3 hydride-transfer step in FNR-like enzymes by combining spectroscopic studies and a series of  
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5 higher resolution structures of FNR crystals soaked with nicotinamide, NADP<sup>+</sup>, and NADPH. To  
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7 accomplish this, we chose corn root FNR, for which crystals of the wild-type enzyme diffract to  
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9 near 1 Å resolution [29]. We also argue for the general relevance of these insights for broader  
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11 members of the FNR superfamily, including the NOX enzymes, which is further supported by the  
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13 very recent structures of the core catalytic subunit domains of NOX5, published while this work  
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15 was under review [30].  
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## 23 **RESULTS AND DISCUSSION**

### 24 **Strategy**

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27 An inspection of the molecular packing in the well-diffracting crystals of corn root FNR that  
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29 we reported earlier [9, 29] revealed that the NADP(H) binding site was not involved in crystal  
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31 packing interactions. We hypothesized that we could obtain high resolution structures of NADP(H)  
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33 complexes of this enzyme by making mutants of Tyr316, the aromatic blocking residue. Not  
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35 knowing which mutant would be more informative, we mutated Tyr316 to Ala (Y316A), Ser  
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37 (Y316S), and Phe (Y316F). Crystal structures of NADP<sup>+</sup> soaks of Y316F FNR showed no  
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39 nicotinamide binding, so this variant was not subjected to further study. For Y316S and Y316A,  
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41 however, we present extensive characterization of the in-solution spectroscopic and catalytic  
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43 properties, as well as crystal structures and in some cases the in-crystal spectroscopic properties  
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45 of their complexes with NADP(H).  
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## Solution Properties of Mutants

Wild-type and mutant enzyme forms were produced in *Escherichia coli* and purified essentially as reported elsewhere [9]. The two protein variants were expressed at slightly lower levels than that the wild-type enzyme, and their purification required additional hydrophobic interaction chromatography on butyl Sepharose. This resulted in a substantial drop in overall yields, which in the case of Y316A was low enough to preclude its full functional characterization. As shown in Figure 2A, the two replacements induced very similar perturbations in the visible absorption spectrum of the flavoprotein, strongly reminiscent of those observed in the equivalent Tyr to Ser replacement in both pea and *Anabaena* FNRs [9, 31]. Phenol – a mimic of the Tyr side-chain – was found to induce a spectral change in Y316S that qualitatively matched the difference between the absorption spectra of the wild-type and the variant protein (Figure 2B), implying that it can stack against the isoalloxazine ring in the active-site pocket. However, the affinity of Y316S FNR for phenol was too low to allow the accurate determination of the  $K_d$  of the complex.

*Titration with NADP(H).* Anaerobic titration of oxidized Y316S FNR with  $\text{NADP}^+$  induced perturbations in the visible absorption spectrum of the FAD prosthetic group (Figure 3) very similar to those observed for the Y308S mutant of pea FNR [6]. The  $K_d$  of the complex was estimated to be  $\leq 0.02 \mu\text{M}$ , well below that of the wild-type complex ( $0.3 \mu\text{M}$ ) [9]. Whereas no interaction between  $\text{NAD}^+$  and wild-type FNR was detectable spectrophotometrically,  $\text{NAD}^+$  induced in Y316S FNR a spectral shift virtually identical to that produced by  $\text{NADP}^+$ , displaying a  $K_d$  of  $145 \pm 5 \mu\text{M}$  (Figure 3).

Next, using an EDTA-deazariboflavin system, stepwise anaerobic photoreductions of the FAD group of wild-type and mutant FNR forms were done in the absence and in the presence of

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3 roughly equimolar amounts NADP<sup>+</sup> or NAD<sup>+</sup> (Figure 4). These showed that the Y316S  
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5 replacement decreased the amount of FAD semiquinone accumulated during the process and  
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7 greatly increased the intensity of the broad long wavelength (~800 nm) absorption band  
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9 attributable to charge-transfer electronic transitions of the interacting oxidized nicotinamide and  
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11 2-electron reduced flavin [9]. Upon complete reduction of the system, the charge-transfer band  
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13 disappeared implying that the limited amount of NADP<sup>+</sup> present became reduced and no charge  
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15 transfer interaction occurs between the reduced flavin and NADPH. Furthermore, the presence of  
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17 NADP<sup>+</sup> favored the protonation of the dihydroquinone form of the prosthetic group of Y316S  
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19 FNR, as indicated by the shift from 360 to 420 nm of the local maximum of the spectra of the fully  
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21 reduced species (Figure 4). The results with NAD<sup>+</sup> (not shown) are qualitatively similar to those  
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23 obtained with NADP<sup>+</sup>, although a significantly less intense charge-transfer band was observed,  
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25 consistent with the lower affinity of the enzyme for NAD<sup>+</sup> as compared to NADP<sup>+</sup>.  
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32 Anaerobic titrations of wild-type FNR and its variants with NADPH were performed to  
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34 further analyze the interactions between FAD and NADP(H). In such titrations, at one equivalent  
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36 of NADPH added, for the “nicotinamide-in” species an equilibrium should exist between two  
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38 charge transfer complexes depending on whether the nicotinamide or the flavin is in the reduced  
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40 state. Charge-transfer complex 1 (CTC-1) has the nicotinamide ring of NADPH interacting with  
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42 oxidized flavin (*i.e.* FNR<sub>ox</sub>:NADPH) and an absorption band with a maximum near 600 nm, and  
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44 charge transfer complex-2 (CTC-2) has the nicotinamide ring of NADP<sup>+</sup> interacting with reduced  
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46 flavin (*i.e.* FNR<sub>red</sub>:NADP<sup>+</sup>) and a broad absorption band with a maximum at ca. 800 nm [32, 33].  
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48 The spectra at one equivalent of NADPH show that for wild-type the CTC-1 absorption band  
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50 dominates (Figure 5A), whereas for both mutants the CTC-2 band dominates (Figure 5B,C),  
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52 apparently accounting for ~90% of the population, based on the level of reduced flavin reduction  
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3 as indicated by  $A_{460}$ . For the two mutants, the CTC-2 band is maximal at roughly one equivalent  
4 NADPH and decreases to about half of that height as the concentration of NADPH is increased to  
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6 an  $\sim 135$ -fold excess (Figure 5B,C). The further additions of NADPH presumably result in the  
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8 progressive decrease in the amount of CTC-2 due to the partial displacement of  $\text{NADP}^+$  by  
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10 NADPH. For the wild-type enzyme, the amount of CTC-1 similarly decreases as NADPH  
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12 concentration increases, but it is progressively replaced by a species attributable to a complex  
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14 between NADPH and FNR carrying FAD semiquinone. We do not understand how the  
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16 semiquinone is formed under these conditions, but wonder if it may reflect some flavin  
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18 disproportionation between the CTC-2 and CTC-1 complexes or is the result of a small amount of  
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20 residual oxygen or other one-electron acceptor in the titrating NADPH solution. Prolonged  
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22 incubation of NO synthase with NADPH was also seen to promote semiquinone buildup [34].  
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29 *Spectral properties during turnover as an NADPH oxidase.* To monitor the spectral changes  
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31 that occur during steady-state NADPH oxidase activity of FNR, air was admitted into the anaerobic  
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33 cell after the titration was complete and NADPH was  $>100$ -fold in excess. In each case, the enzyme  
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35 underwent turnover until the NADPH was all converted to  $\text{NADP}^+$ . For wild-type FNR (Figure  
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37 5D), as has been seen before [9], there was substantial stabilization of the blue semiquinone form  
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39 of FAD in the early parts of the reaction and CTC bands were hardly detectable. For Y316S FNR  
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41 (Figure 5E), less FAD semiquinone built up and the reaction had two distinct phases. During the  
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43 first phase, the CTC-2 concentration progressively increased until at  $\sim 15$  min it reached  $\sim 65\%$  of  
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45 the maximal value observed during the anaerobic titration (thin red line), then it decreased to near  
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47 zero. The final spectrum (at  $\sim 1.7$  h) had the 456 nm band shifted as is characteristic of an oxidized  
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49 enzyme in complex with  $\text{NADP}^+$  (Figure 5E). The behavior of Y316A FNR (Figure 5F) was  
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51 qualitatively similar to that of the Y316S form, but with the reaction slowed by about 10-fold as  
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3 the steady-state CTC-2 formation was reached after about 2.5 h (vs ~15 min) and the reaction was  
4 complete in 17 rather than 1.7 h. The Y316A variant also had still less stabilization of the FAD  
5 semiquinone, and its maximal CTC-2 accumulation during turnover was ~85% of the highest  
6 amount observed in the anaerobic titration (thin red line). The slower reaction and higher CTC-2  
7 buildup are consistent with the  $k_{\text{off}}$  for  $\text{NADP}^+$  being slower for the Y316A mutant. Also, the  
8 destabilization of the semiquinone seen in these variants mirrors what was seen for the  
9 corresponding mutants of *Anabaena* FNR [31] and NO synthase [22, 34].

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20 *Ferricyanide reductase activity.* For the wild-type enzyme, ferricyanide is the most effective  
21 electron acceptor from reduced FNR, with  $k_{\text{cat}} = 520 \text{ s}^{-1}$  [9]. The Y316S and Y316A replacements  
22 impaired the steady-state diaphorase activity of FNR to a similar extent, and  $k_{\text{cat}}$  and  $K_{\text{m}}^{\text{NADPH}}$   
23 values determined for the Y316S variant were ca. 170 and 9-fold lower than those of the wild-type  
24 enzyme (Table 1) [9], respectively. Such large catalytic impairment parallels that reported for the  
25 Tyr to Ser variants of pea and *Anabaena* FNRs [21, 31], with the slower turnover shown to be  
26 limited by the rate of  $\text{NADP}^+$  (*i.e.* product) release. Consistent with this, we found that  
27 nicotinamide partially rescued the diaphorase activity of Y316S FNR in a concentration-dependent  
28 fashion, increasing its  $k_{\text{cat}}$  to  $280 \text{ s}^{-1}$  at 800 mM nicotinamide (Table 1). As has been noted  
29 previously [2, 6], by competing with the corresponding moiety of  $\text{NADP}^+$  for stacking onto the  
30 flavin, nicotinamide will favor product dissociation and speed enzyme turnover if the  $k_{\text{off}}$  of  
31  $\text{NADP}^+$  is rate limiting.

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48 *Interactions with small-molecule ligands.* The findings that added nicotinamide increased the  
49 catalytic activity of Y316S FNR, prompted us to quantify the affinities of nicotinamide and N-  
50 methyl-nicotinamide. As shown in Figure 3A, both were found able to interact with Y316S FNR,  
51 although only the former had an affinity high enough for its  $K_{\text{d}}$  value to be determined ( $3 \pm 0.1$   
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3 mM). The difference spectrum of the Y316S-nicotinamide complex was blue-shifted by ca. 7 nm  
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5 with respect to that of the NAD(P)<sup>+</sup> complexes, but the spectral change induced by N-methyl-  
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7 nicotinamide was almost identical to that induced by NAD(P)<sup>+</sup> (Figure 3B). This indicates that for  
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9 Y316S FNR, N-methyl-nicotinamide but not nicotinamide is an excellent mimic of the redox-  
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11 active moiety of the dinucleotides.  
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### 17 **Reanalysis of Stopped-Flow Results Reported for *Anabaena* FNR.**

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20 As noted in the introduction, in earlier work Lans *et al.* [26] called into question the relevance  
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22 of the NADP(H) complexes that form in FNR when the aromatic placeholder is mutated. The  
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24 origin of this view came from stopped-flow results for the Tyr303Ser (Y303S) mutant of *Anabaena*  
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26 *variabilis* FNR. Per their interpretation, these results showed that Y303S promoted the hydride  
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28 transfer from NADPH to FNR<sub>ox</sub> “slightly slower than the wild-type enzyme” but that “the reverse  
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30 process is undetectable” (*e.g.* hydride transfer from FNR<sub>red</sub> to NADP<sup>+</sup> does not occur). This  
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32 interpretation was based on the observation that when NADP<sup>+</sup> was mixed with Y303S FNR<sub>red</sub> the  
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34 CTC-2 (~800 nm) charge transfer band that formed upon mixing did not substantially decrease  
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36 over time (see Figure 3A,B of Lans *et al.* [26]). While the conclusion that no hydride transfer  
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38 occurs is consistent with the unchanging CTC-2 band, there could be other explanations. We were  
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40 skeptical of the inference both based on the thermodynamic principle that if the Y303S mutant  
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42 catalyzed hydride transfer from NADPH to FNR<sub>ox</sub> it must equally well catalyze the reverse hydride  
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44 transfer from FNR<sub>red</sub> to NADP<sup>+</sup>, and based on our photoreduction studies (Figure 4 above)  
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46 implying that Y316S-bound NADP<sup>+</sup> is converted to NADPH. To clarify this question, we  
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48 reanalyzed the Lans *et al.* [26] stopped-flow data for NADPH mixed with Y303S FNR<sub>ox</sub> and for  
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50 NADP<sup>+</sup> mixed with Y303S FNR<sub>red</sub>.  
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Importantly, in the original work, when NADP<sup>+</sup> is mixed with Y303S FNR<sub>red</sub>, even though the 800 nm band changes little over time, the ~460 nm peak systematically increases indicating that oxidation of FAD is occurring. Lans *et al.* attributed this to a “side effect” rather than enzymatic activity; however, when comparing these spectra with those for NADPH mixed with Y303S FNR<sub>ox</sub>, it is striking that both reactions appear to reach the same endpoint, as would be expected if Y303S FNR were reaching the same equilibrium state independent of the direction of approach. Given this observation, we reanalyzed the reported stopped-flow results for both reactions using the changes in A<sub>460</sub> as an indicator of FAD oxidation/reduction. This reanalysis yielded excellent fits of  $k_{\text{obs}}$  of  $164 \pm 16 \text{ s}^{-1}$  for NADP<sup>+</sup> mixed with Y303S FNR<sub>red</sub> and  $146 \pm 21 \text{ s}^{-1}$  for NADPH mixed with Y303S FNR<sub>ox</sub> (Figure 6). These two  $k_{\text{obs}}$  values should both equal the sum of the elementary forward and back reaction rate constants ( $k_{\text{obs}} = k_1 + k_{-1}$ ) and are equal within the error of the analysis, leading us to conclude that the forward and back reactions both function as would be expected. We suggest the best number to use for  $k_{\text{obs}}$  for the *Anabaena* Y303S FNR is  $190 \pm 15 \text{ s}^{-1}$  which is the value originally reported by Lans *et al.* [26] for the forward reaction using the much more extensive original dataset.

This revised interpretation is consistent with all the data and with basic thermodynamic principles, and implies that the nicotinamide-flavin interaction consistently formed in the aromatic placeholder mutants is in fact productive and relevant for understanding catalysis.

### Crystal structures of Y316S and Y316A NADP(H) complexes

Co-crystallization of both FNR variants with nicotinamide yielded crystals that grew readily and could be soaked with NADP<sup>+</sup> or NADPH to obtain those complexes. Interestingly, the variants crystallized in two space groups, both the P<sub>3</sub><sub>2</sub>2<sub>1</sub> form seen for wild-type protein [9] and a new

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3 P3<sub>1</sub>21 crystal form that had remarkably similar unit cell dimensions and related crystal packing.  
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5 Here, we report a set of eight refined crystal structures of complexes that represent the Y316S and  
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7 Y316A variants in complex with three ligands (NADP<sup>+</sup>, NADPH, and nicotinamide) with most  
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9 structures refined at between 1.35 Å and 1.6 Å resolution (Table 2) and having very well defined  
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11 active site electron density (Figure 7). As the corresponding complexes of Y316S and Y316A  
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13 largely show equivalent features, we focus mainly on the highest resolution set of structures that  
14  
15 provide unique information.  
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19 To check whether the complexes in the crystal represent those that form in solution, we  
20  
21 obtained single-crystal absorption spectra [35] during data collection for a pair of Y316A crystals  
22  
23 soaked with NADP<sup>+</sup> and NADPH. These spectra qualitatively match those obtained in solution  
24  
25 and indicate that even though the X-ray beam does cause reduction of the FAD, this does not  
26  
27 undermine the relevance of the structures (Figure 7). Also, since Y316S and Y316A both  
28  
29 predominantly form CTC-2 after an NADPH soak (*e.g.* Figure 5B,C), we have modeled the  
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31 NADPH soak structures as an NADP<sup>+</sup>:FADH<sub>2</sub> complex rather than NADPH:FAD.  
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37 In the following sections, we first provide a basic description of the complexes and then focus  
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39 on three detailed aspects of complexes that provide evidence of compression in the active site for  
40  
41 enhancing catalysis. In our structural comparisons, the unliganded wild-type structure we use is a  
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43 1.05 Å resolution structure refined much earlier (PDB entry 3LO8; released in 2010) and used in  
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45 tests of new refinement strategies [36]. As the statistics for this structure have not yet been  
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47 described in the literature, they are included in Table 2.  
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51 *Overview of the NADP(H) complexes.* Globally, all Y316S and Y316A complexes,  
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53 independent of space group, are very similar to the wild-type structure with rms C $\alpha$  deviations  
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55 between 0.1 Å and 0.4 Å. The broad features of NADP(H) binding match those that have been  
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3 elaborated elsewhere [4, 6, 10, 12], interacting mostly with the NADP-binding domain. Briefly,  
4 the 2'-phosphate anchor hydrogen bonds with the side chains of Ser237, Arg238, Lys247, and  
5 Tyr249, and the adenine is sandwiched between Tyr249 and Leu276. The 5'-phosphoryl group is  
6 largely exposed and the nicotinamide-side phosphoryl group hydrogen bonds with the Thr175 N  
7 and the Arg114 guanidinium group. The nicotinamide is surrounded by a triad of key side-chains,  
8 with Ser94 and Glu314 hydrogen bonding with the nicotinamide carboxamide nitrogen and  
9 Cys274 and Ser94 close to the C4 atom (Figure 7A). In the higher resolution structures of Y316S,  
10 an alternate conformation for the nicotinamide ribose is also observed. In both Y316S<sub>NADP+</sub> and  
11 Y316S<sub>NADPH</sub> soaks in both space groups, the nicotinamide ribose is modeled as 60% 2'-endo and  
12 40% 3'-endo. In each Y316A structure, the nicotinamide ribose adopts a single conformation, but  
13 in Y316A<sub>NADP+</sub> it is 2'-endo and in Y316A<sub>NADPH</sub> it is 3'-endo. The reason for these conformational  
14 differences is not readily apparent but their presence suggests that they are the result of real but  
15 subtle differences in nicotinamide binding.  
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34 *The complexes with free nicotinamide.* Surprisingly, the binding mode seen in nicotinamide  
35 soaks differ for Y316A and Y316S (Figure 3C). In Y316A<sub>nic</sub>, the nicotinamide orientation and  
36 hydrogen bonding roughly match how the NADP(H) nicotinamide binds. However, in Y316S<sub>nic</sub>,  
37 the nicotinamide binds in a flipped orientation that is stabilized by a hydrogen bond between  
38 Ser316-O $\gamma$  and nicotinamide-N1 as well as hydrogen bonds of the nicotinamide hydroxyamide  
39 with Ser94-O $\gamma$  and Thr 175-O $\gamma$  and O atoms (Figure 3C). Since this binding mode involves Ser316  
40 as a hydrogen-bond partner, we conclude it is an artifact of the Y316S mutation. Also, the  
41 existence of the nicotinamide N1 to Ser316-O $\gamma$  hydrogen bond in this complex means that neither  
42 NADP(H) nor even N1-methyl-nicotinamide could adopt it. This provides a satisfying explanation  
43 for why nicotinamide binding to Y316S yields different spectral changes than NAD(P)<sup>+</sup>, whereas  
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3 methyl-nicotinamide binds less tightly, but with spectral changes that match those of NAD(P)<sup>+</sup>  
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5 (Figure 3B). It also serves as a reminder of the need to be cautious in inferring the binding mode  
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7 of a molecule of interest based on that of an analog.  
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10 *The nicotinamide-flavin interactions in the NADP<sup>+</sup>/NADPH complexes.* While the overall  
11 structures of the FNR variants are virtually superimposable, subtle differences exist in the details  
12 of nicotinamide binding seen in the NADP<sup>+</sup> and NADPH soaks, and having the structures for the  
13 two mutants allows us to gain confidence about which variations are reliably due to the difference  
14 between the active site redox state rather than the specific mutation or to experimental uncertainty.  
15 In terms of the active site, the difference between the structures is a single H-atom: the NADP<sup>+</sup>  
16 soaks with oxidized flavin and nicotinamide have one H-atom on the nicotinamide C4 and no H-  
17 atom on the flavin N5, and the NADPH soaks to produce CTC-2 have one H-atom on the  
18 nicotinamide C4 as well as one H-atom on the flavin N5. (The CTC-1 complex would have the  
19 same number of H-atoms but distributed with two on the nicotinamide C4-atom and none on the  
20 flavin N5-atom.) As described in the following paragraphs, the high resolution structures provide  
21 evidence that the single additional H-atom increases active site crowding and a directional  
22 compression that we propose is a key promotor of catalysis.  
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41 A first line of evidence is provided by the temperature factors of the C4-atom of NADP<sup>+</sup>  
42 compared to NADPH. In our earlier work on pea FNR [6], a higher conformational freedom of the  
43 NADP<sup>+</sup> C4-atom was indicated by weaker electron density (and higher B-factors) for this atom in  
44 NADP<sup>+</sup> compared to NADPH; but at 1.8 Å resolution, this was not conclusive. Our higher  
45 resolution analyses here similarly show the additional mobility of the C4 atom of NADP<sup>+</sup> but also  
46 provide refined anisotropic B-factors that give a more complete description of dynamics than do  
47 isotropic B-factors. The anisotropic B-factors very clearly show that the increased movement for  
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3 the C4 atom of NADP<sup>+</sup> is perpendicular to the plane of the ring and towards N5 of the FAD (Figure  
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5 8A). This type of motion exactly corresponds to motions that would allow the nicotinamide ring  
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7 to form a boat-like conformation that calculations show will promote efficient hydride transfer  
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9 [37].  
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12 A second observation of crowding derives from a visible distortion of the FAD isoalloxazine.  
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14 In wild-type FNR, the isoalloxazine group appears to be pushed slightly by the Tyr316 side chain  
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16 so that it is not coplanar with the N10-C1 ribose bond, but is about 3.5° non-planar (Figure 8B).  
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18 In all FNR variant structures soaked with NADP<sup>+</sup>, this non-planarity increases to ~10°, and in the  
19  
20 structures soaked with NADPH it increases further to ~14° (Figure 8B). These deviations from  
21  
22 planarity imply that the nicotinamide displacing the Tyr316 side chain leads to a more tightly  
23  
24 packed active site and that the additional H-atom in the active site present in the NADPH soaks  
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26 increases crowding even more, with the isoalloxazine bending to relieve apparent pressure. It  
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28 should be noted that the highly crowded complex seen in the NADPH soak is the one that is present  
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30 during normal catalysis.  
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36 A third observation that completes the view of the tightly packed active site of FNR are the  
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38 interaction distances of the nicotinamide C4 and H4 atoms that are involved in hydride transfer  
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40 (Figure 8C). In all structures, these atoms are sandwiched between the flavin N5 atom in front and  
41  
42 the Cys274-S<sub>γ</sub> sulfhydryl behind, where it can act as a backstop. In comparing the Y316S<sub>NADP<sup>+</sup></sub>  
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44 complex to the Y316S<sub>NADPH</sub> complex which has an additional H-atom on the flavin N5-atom, the  
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46 C4...N5 distance increases by ~0.3 Å while the C4H... S<sub>γ</sub> distance decreases by ~0.1 Å (Figure  
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48 8C). Taking into account the Y316A complexes (Figure 8C), the consistent increase of the C4...N5  
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50 distance in the NADPH complex leads us to conclude that Cys274 acts as a firm backstop while  
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52 the isoalloxazine bends away from the nicotinamide by an additional ~4° in order to increase the  
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3 distance between the flavin and nicotinamide and relieve the pressure due to the presence of the  
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5 extra H-atom.  
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8 These observations lead us to conclude that the active site is experiencing compression,  
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10 distorting the FAD and pushing the substrates together to enhance catalysis by aligning the  
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12 substrates in an optimal position and, as has been noted by others, potentially promoting a quantum  
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14 tunneling mechanism of hydride transfer [26].  
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### 17 18 19 **Insights into factors promoting hydride transfer in FNRs**

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21 Taken along with our reanalysis of previous stopped-flow kinetics work that called the  
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23 relevance of aromatic placeholder FNR mutants into question, these in-solution spectroscopy,  
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25 single-crystal spectroscopy, and high-resolution structures provide insights into hydride transfer  
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27 in FNRs using variants which have near wild-type hydride-transfer kinetics, but importantly, allow  
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29 us to capture the productive binding mode of NADP(H) in the active site. Such productive  
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31 complexes have not been possible to capture in any wild-type member of the FNR superfamily  
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33 with an aromatic placeholder residue (*e.g.* [5, 17-20]) or that instead have a C-terminal peptide  
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35 extension as an alternate way to block nicotinamide binding [3, 38]. They have also been  
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37 challenging to obtain for superfamily enzymes that have a shifted domain-domain interaction [39,  
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39 40] instead of an aromatic blocking residue, but for one of these enzymes, cytochrome b5  
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41 reductase, anaerobic co-crystallization has yielded a productive, wild type complex with NADH  
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43 in which the nicotinamide-flavin interaction geometry is much like that seen here [41].  
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51 The high-resolution structures reported here illustrate the role of specific anisotropic motions  
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53 as well as active site compression as catalytic strategies that promote hydride transfer. Key  
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55 evidences are the anisotropic motion of the C4 atom of NADP<sup>+</sup>, that indicates boat-like  
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3 perturbations in conformation are a preferred mode of vibrational freedom, as well as a strong  
4 decrease in the amplitude of that motion in the NADPH complex. Additionally, there is an  
5 incrementally increasing deviation from planarity at the flavin N10 atom in wild type,  
6 Y316S<sub>NADP+</sub>, and Y316S<sub>NADPH</sub> indicating increased crowding and pressure associated with  
7 oxidized nicotinamide binding and even more so with reduced nicotinamide binding that is being  
8 relieved to some extent by bending of the flavin. Interestingly, a very similar flavin non-planarity  
9 was seen in the cytochrome b5 reductase-NADH complex (see Figure 3a of [41]). Finally,  
10 comparison of Y316S<sub>NADP+</sub> and Y316S<sub>NADPH</sub> reveal close sub van der Waals contact distances  
11 consistent with a tightly packed active site. The central role of Cys274 in these interactions  
12 provides a rationale for its conservation across the whole superfamily. While steric compression  
13 has occasionally been noted as a key factor promoting catalysis for other enzymes [42-45], there  
14 are a limited number of studies which provide direct structural evidence for this [46-50], as this  
15 typically requires atomic or near-atomic resolution crystal structures. Also, that the Y316S and  
16 Y316A variants of corn root FNR, like similar mutants of other FNR superfamily members [22,  
17 31, 34], show much less stabilization of the FAD semiquinone, is consistent with an earlier  
18 proposal [31] that the aromatic side chain is not solely a passive placeholder for the nicotinamide  
19 group, but, through its stacking interaction with the flavin, is an active agent that stabilizes the  
20 semiquinone form of the flavin to enhance the one-electron transfers required for these  
21 dehydrogenase-electron transferases.  
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### 51 **Extrapolation of the results to the FNR superfamily members such as NOX enzymes**

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53 Clarifying the relevance of the NADP<sup>+</sup> binding mode seen in the Tyr316 mutants is not just  
54 of interest for understanding catalysis for the whole superfamily, but can guide further studies of  
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3 these enzymes as well as the generation of superfamily member variants that can be valuable tools.  
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5 For instance, for the NOX enzymes that play crucial roles in the production of superoxide and  
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7 hydrogen peroxide for diverse biological processes, the FNR-like module is at the C-terminus of  
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9 a membrane-bound flavocytochrome catalytic subunit that has been relatively difficult to study  
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11 [14]. Although a NOX structure has long eluded structural biologists, a crystal structure of the  
12  
13 FNR-like module of NOX5 from *Cylindrospermum stagnale* (referred to as the NADPH-  
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15 dehydrogenase domain) was solved while this work was under review [30]. Sequence alignments  
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17 of FNR with human NOXs aided by the *C. stagnale* NOX5 structure (Figure 9A) show that the  
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19 NOX isozymes conserve many NADP<sup>+</sup> binding residues including the nicotinamide-interacting  
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21 residues equivalent to corn root FNR Ser94, Cys274, Glu314 and Tyr316 (Phe693 in *C. stagnale*  
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23 NOX5).  
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29 The NOX5 FNR-like module itself was unstable and crystals were only obtained for a  
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31 construct with a “hyperstabilizing” C-terminal extension that included a Trp two residues after  
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33 Phe693. Notably, in this structure, the expected natural C-terminal aromatic placeholder (Phe693)  
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35 does not stack against the isoalloxazine, and, because the anticipated fifth  $\beta$ -strand of the NADP<sup>+</sup>-  
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37 binding domain is perturbed and moved away from the protein core, the Trp residue is able to stack  
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39 against the isoalloxazine ring instead (Figure 9B). However, if the NOX5 chain followed the path  
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41 of the  $\beta$ -strand seen in other FNR superfamily members, Phe693 could stack against the  
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43 isoalloxazine and we propose that this is what occurs in the native enzyme. Magnani et al [30]  
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45 noted that the isolated dehydrogenase domain is deregulated and predicted that the normal role of  
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47 the strictly conserved C-terminal aromatic residue would “emerge only in the context of a full-  
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49 length protein.” We agree and further suggest that the instability of the isolated dehydrogenase  
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51 domain and the loose association of the C-terminal segment (including Phe693) is related to the  
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3 missing cytochrome transmembrane domain and associated lipid bilayer that interact with this  
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5 surface of the dehydrogenase domain in a full-length complex (see Fig 5A of [30]).  
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8 Consistent with the weak association observed for Phe693 with the protein core is in the  
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10 isolated dehydrogenase domain, a Phe693Ser mutant had higher than wild-type activity and the  
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12 stabilizing C-terminal extension mutant (adding the Trp695) dropped activity by 5-fold [30].  
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14 Nevertheless, we predict that in the context of a full-length membrane-bound NOX, Phe693 will  
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16 act as an aromatic placeholder residue and that Ser or Ala mutants will bind both NADPH and  
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18 NADH more tightly. Such mutants could provide a useful handle for purification (through the tight  
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20 binding of NADP<sup>+</sup>) or be a useful tool for probing/controlling the physiological roles of NOXs. A  
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22 solely NADH responsive version could potentially be made through additional mutations such as  
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24 the equivalent of FNR Ser237 to Asp (Figure 9), as this position has been shown to discriminate  
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26 against binding the 2'-phosphoryl group in multiple superfamily members [51-54]. Interestingly,  
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28 a full-length human NOX2 C-terminal Phe mutant (F570A) was characterized in 1998 and found  
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30 to retain ~50% of the wild-type activity [55] rather than dropping 300-800 fold in activity as had  
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32 been seen for the equivalent pea FNR mutant, and so it was not studied further. However, this  
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34 study predated the knowledge that the equivalent FNR mutant lost activity due to the slow  
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36 dissociation of NADP<sup>+</sup> and also gained activity with NADH [9, 21], so those qualities of the variant  
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38 were never characterized. In retrospect, given that the NOX2 K<sub>d</sub> for NADP<sup>+</sup> is ~40 μM (over 10-  
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40 fold higher than is typical for FNRs) and the turnover number is slower, it would not be surprising  
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42 if the expected enhanced binding of NADP<sup>+</sup> in the F570A variant would not be enough to make  
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44 its dissociation highly rate limiting. Indeed, for NO synthase the equivalent mutation only showed  
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46 about a 3-fold decrease in steady-state turnover, yet still showed a 50-fold change in specificity  
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48 [22, 34].  
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## MATERIALS AND METHODS

### Production of Recombinant Corn Root FNR and its Variants

Plasmids for the bacterial expression of the Y316A and Y316S proteins were generated from the pETrFNR2 [9] using the QuikChange II Site-Directed Mutagenesis kit (Agilent) and two appropriate oligonucleotide couples following the manufacturer directions. The wild-type and variant proteins were produced in *E. coli* HMS174(DE3) by induction with 0.1 mM IPTG for 4 h at 30 °C, and purified through a procedure similar to that previously reported for the wild-type protein [9] using an ÄKTA FPLC (GE Healthcare) apparatus. Briefly, the crude cell lysate was brought to 40% NH<sub>4</sub>SO<sub>4</sub> and centrifuged. The supernatant was applied to a Sepharose 4B column (GE Healthcare), eluted with appropriate buffer and precipitated with 75% (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>. In the case of the protein variants, the pellet was resuspended in 40% (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub> and chromatographed on a butyl Sepharose column (GE Healthcare) with a descending salt concentration gradient. The omission of this step prevented the adsorption of the FNR forms on the next cation exchange resin. After desalting, the sample was loaded on an SP-Sepharose HP column (GE Healthcare) and eluted through a NaCl concentration gradient. Notably, the visible spectra of both mutant forms, but not the wild-type enzyme, underwent a blue shift during the ion exchange step, suggesting release of some unidentified bound ligand at that stage. The resulting proteins were homogenous as judged by SDS-PAGE. The FNR forms were concentrated to ~25 mg/mL and stored at -20 °C in 10 mM HEPES, pH 7.0.

### Spectral Analyses and Ligand Binding

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3 All spectrophotometric measurements and steady-state enzyme kinetics were performed using  
4 an 8453 diode-array spectrophotometer (Agilent). Titration with NADP<sup>+</sup>, NAD<sup>+</sup>, phenol,  
5 nicotinamide and N-methyl-nicotinamide of the enzyme forms in their oxidized state were  
6 performed in 10 mM Tris-HCl, pH 7.7, at 15 °C.  
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### 15 **Photoreductions, Anaerobic Titrations and Activity Assays**

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17 Flavin photoreduction experiments were carried out both in the absence and in the presence  
18 of NADP<sup>+</sup> or NAD<sup>+</sup> in anaerobic cuvettes in 10 mM HEPES-NaOH, pH 7.0, at 15 °C, according  
19 to the procedure described elsewhere [9].  
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24 Anaerobic titrations with NADPH of 14-18 μM wild-type, Y316A or Y316S FNR variants  
25 were performed on 1.2 mL samples in 50 mM HEPES-NaOH buffer, pH 7.0, at 15 °C, in a sealed  
26 cuvette. After recording the spectrum of the oxidized enzyme, five successive additions of  
27 anaerobic NADPH were made leading to ligand concentrations of 29 μM, 39 μM, 76 μM, 182  
28 μM, and 1.9 mM for wild-type titrations, 23 μM, 45 μM, 90 μM, 210 μM, and 2.2 mM for FNR  
29 Y316A titrations, and 24 μM, 48 μM, 95 μM, 225 μM, and 2.3 mM for FNR Y316S titrations,  
30 respectively. In each case, the volume after the final addition of NADPH was 1.4 mL. For each  
31 FNR form, the cuvette was then opened to air, and the spectrum of the mixture (kept at 15 °C) was  
32 monitored at over time until all the NADPH was oxidized. For technical reasons, the spectra for  
33 wild-type FNR during O<sub>2</sub> turnover was produced from a separate sample also prepared with 2.4  
34 mM NADPH.  
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50 NADPH—K<sub>3</sub>Fe(CN)<sub>6</sub> reductase activity assays were performed in 100 mM Tris-HCl, pH 8,  
51 at 25 °C, at different concentrations of both substrates, as reported elsewhere [56]. As needed,  
52 nicotinamide was included in the assays at concentrations ranging from 50 to 800 mM.  
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### Reanalysis of Stopped-Flow Results Reported for *Anabaena* FNR

Spectra (Figure 3 in [26]) were enlarged by ~345% and printed. The height of each peak at ~460 nm corresponding to the flavin oxidation state was measured in cm from the baseline of the spectra (-0.025 AU). The peak height at time 0 for FNR<sub>red</sub> with NADP<sup>+</sup> (corresponding to fully reduced flavin) was taken as a baseline and used to normalize the measurements by subtracting this height from the other peak heights of the corresponding variant. Whereas the final spectra (0.2547 s) for wild-type FNR for both FNR<sub>red</sub> with NADP<sup>+</sup> and FNR<sub>ox</sub> with NADPH matched in both peak height and shape, the final spectra (0.2547 s) for Y303S FNR varied slightly in peak height but both correspond to achieved equilibrium so were normalized to each other by subtracting the difference between the final peak heights from the measured peak heights for Y303S FNR<sub>red</sub> with NADP<sup>+</sup>.

Taking the peak height at time 0 for FNR<sub>ox</sub> with NADPH to correspond to fully oxidized flavin, the percentage of oxidized and reduced flavin and corresponding concentration of product (using the provided concentrations of 25 μM FNR and 125 μM NADP(H)) were calculated. A plot of concentration of product (μM) versus time (s) were fitted using GraFit 5 (Erithacus Software Limited) with a single-exponential decay equation to estimate the apparent rate constants of the hydride-transfer reactions.

### Crystallization and Structure Determinations

Initial co-crystallization trials of the corn root FNR variants with NADP<sup>+</sup> used similar conditions to those used for the wild-type enzyme [9] but yielded only ill-formed small crystals

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3 that grew very slowly. Solving their structure revealed that they contained a bound nicotinamide  
4 moiety rather than NADP<sup>+</sup>, so co-crystallization with nicotinamide was tried, and crystals grew  
5 readily. The best crystallizations were at room temperature in hanging drops formed by mixing  
6 equal volumes of the protein stock (Y316F: 20.2 mg/mL; Y316A: 7.3 mg/mL; Y316S: 10.8  
7 mg/mL; all in 50 mM Tris-HCl, pH 7.4) with a reservoir solution containing 22-24% PEG 8000,  
8 0.1 M sodium cacodylate (pH 6-7), 0.18-0.22 M magnesium acetate, and 100 mM nicotinamide.  
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10 Typical crystals were ~0.3 mm on each side and grew within 1 week. Both FNR variants formed  
11 many prism-shaped crystals belonging to the space group P3<sub>2</sub>21 (same as wild type; [9]) and fewer  
12 hexagon-shaped crystals belonging to space group P3<sub>1</sub>21. Soaks of the nicotinamide-bound  
13 crystals were done aerobically at room temperature with either 10 mM NADP<sup>+</sup> or 10 mM NADPH  
14 for 1 h to obtain desired complexes before being flash frozen.  
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29 *Data collections.* For data collection, crystals were pulled through oil and flash-frozen in  
30 liquid nitrogen. In-house data were collected at 140 K on a Rigaku RU300 Cu-K $\alpha$  rotating anode  
31 X-ray source running at 50 kV and 100 mA equipped with a Raxis IV image plate detector.  
32 Synchrotron data were collected at beamline 5.0.1 or 5.0.3 at the Advanced Light Source  
33 (Lawrence Berkeley National Laboratory, Berkeley, CA) at 100 K. Oscillation images were  
34 collected with  $\Delta\phi = 1^\circ$  and were processed using Denzo and Scalepack [57] or iMosflm [58] and  
35 SCALA [59]. Unit cell parameters and data reduction statistics for each FNR variant are given in  
36 Table 2. In addition, data sets with  $\Delta\phi = 0.5^\circ$  and simultaneous single crystal visible absorption  
37 spectra were collected by Dr. Allen Orville at a National Synchrotron Light Source beamline as  
38 described in [35].  
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53 *Structure solutions and refinements.* All P3<sub>2</sub>21 FNR variant crystals were isomorphous with  
54 the published wild-type corn root FNR structure (PDB code 3LO8 [36]), and this was used as the  
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3 starting model for the first refinement, with additional refinements built on partially refined models  
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5 of the most similar structure. The P3<sub>1</sub>21 crystal form was trivial to solve by molecular replacement  
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7 and these structures were then similarly refined. The same 10% of data were set aside for cross-  
8  
9 validation [60] for each structure as had been set aside for the ~1 Å resolution refinement of the  
10  
11 wild-type structure. Refinements were carried out by a number of researchers over many years  
12  
13 using various versions of REFMAC [61] with Coot [62] used for manual fitting and Molprobit  
14  
15 [63] used to help identify model problems. During iterative manual rebuilding, water molecules  
16  
17 were added in Coot using standard criteria (>1  $\rho_{\text{rms}}$  intensity in the 2F<sub>o</sub>-F<sub>c</sub> map, >2.4 Å distance  
18  
19 from nearest contact, no B-factors >80 Å<sup>2</sup>). For making a consistent set of structures for  
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21 publication, a set of final refinements of each structure were done by the lead author using Phenix  
22  
23 [64]. During this stage, we decided to model all NADPH soaks as having an oxidized nicotinamide  
24  
25 (*i.e.* NADP<sup>+</sup>) and a reduced flavin since the spectra show that CTC-2 is present at higher amounts  
26  
27 than is CTC-1. For structures refined with anisotropic B-factors (the P3<sub>2</sub>21 Y316S:NADPH,  
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29 Y316S:NADP<sup>+</sup>, and Y316S:nicotinamide structures), no anisotropic restraints were used for the  
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31 ligand in the final round of refinement to ensure the anisotropic B-factors of the nicotinamide  
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33 atoms were based as much as possible on the diffraction data. Refinement statistics for all models  
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35 are shown in Tables 2 and S1. Structural overlays were performed using the protein structure  
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37 visualization software PyMOL [65].  
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46 Also, in order to obtain a reliable representation of active site distances for Y316S<sub>NADPH</sub>,  
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48 Y316S<sub>NADP<sup>+</sup></sub> (P3<sub>2</sub>21 and P3<sub>1</sub>21), Y316A<sub>NADPH</sub>, and Y316A<sub>NADP<sup>+</sup></sub> structures, ten different starting  
49  
50 models for each were generated using the “shake” algorithm of Phenix with the setting  
51  
52 “modify.sites.shake = 0.5”. This level of coordination disruption resulted in starting R/R<sub>free</sub> values  
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54 of ~40%. Each of these models was re-refined and the distances between Cys274 S $\gamma$  and NADP<sup>+</sup>  
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3 C4H and between NADP<sup>+</sup> C4 and FAD N5 were measured. The average distances and standard  
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6 deviations for each set were calculated and reported.

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8 *Accession Numbers.* Coordinates and structure factors for the wild-type corn root FNR,  
9  
10 Y316S:nicotinamide (P3<sub>2</sub>21), Y316S:nicotinamide (P3<sub>1</sub>21), Y316S:NADP<sup>+</sup> (P3<sub>2</sub>21),  
11  
12 Y316S:NADP<sup>+</sup> (P3<sub>1</sub>21), Y316S:NADPH (P3<sub>2</sub>21), Y316A:nicotinamide (P3<sub>2</sub>21), Y316A:NADP<sup>+</sup>  
13  
14 (P3<sub>2</sub>21), Y316A:NADPH (P3<sub>2</sub>21), Y316F (P3<sub>2</sub>21), and Y316F:NADP<sup>+</sup> (P3<sub>1</sub>21) models have been  
15  
16 deposited in the Protein Data Bank with accession numbers 3LO8, 5VW4, 5VW9, 5VW3, 5VW8,  
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18 5VW2, 5VW5, 5VW6, 5VW7, 5VWA, and 5VWB, respectively.  
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44  
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46  
47 experiments and analyzed data; VP, ARH, and RF planned and performed experiments; GZ  
48  
49 planned experiments; AA planned and performed experiments, analyzed data, and wrote the paper;  
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51 PAK planned experiments, analyzed data and wrote the paper.  
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For Review Only

## TABLES

**Table 1.** Effect of the Y316S replacement on the kinetic parameters of the NADPH—K<sub>3</sub>Fe(CN)<sub>6</sub> reductase reaction catalyzed by FNR.<sup>a</sup>

FNR form	$k_{\text{cat}}$ s <sup>-1</sup>	$K_{\text{m}}^{\text{NADPH}}$ μM	$k_{\text{cat}}/K_{\text{m}}^{\text{NADPH}}$ s <sup>-1</sup> μM <sup>-1</sup>
Wild type <sup>b</sup>	520 ± 10	12 ± 1	43 ± 4
Y316S	3 ± 0.3	1.4 ± 0.3	2.1 ± 0.5
Y316S, 0.8 M nicotinamide	280 ± 18	45 ± 10	6 ± 1

<sup>a</sup>Initial rate data were measured as moles of ferricyanide reduced per second.

<sup>b</sup>Data on wild type FNR, taken from [9], are reported here for comparison.

For Review Only



**Table 2.** Data and refinement statistics for FNR variant structures<sup>a</sup>

	wt	Nicotinamide	Nicotinamide	Y316S	NADP+	NADPH	Nicotinamide	Y316A	NADPH
Ligand soaked	None	Nicotinamide	Nicotinamide	NADP+	NADP+	NADPH	Nicotinamide	NADP+	NADPH
Space group	P3 <sub>2</sub> 21	P3 <sub>2</sub> 21	P3 <sub>1</sub> 21	P3 <sub>2</sub> 21	P3 <sub>1</sub> 21	P3 <sub>2</sub> 21	P3 <sub>2</sub> 21	P3 <sub>2</sub> 21	P3 <sub>2</sub> 21
<i>Data Statistics</i>									
Wavelength (Å)	1.00	1.00	1.54	1.00	1.54	1.00	1.54	1.00	1.00
Unit cell <i>a</i> , <i>c</i> axes (Å)	59.1, 186.7	58.9, 184.8	58.9, 184.4	58.8, 187.5	58.9, 185.0	58.9, 187.3	59.4, 187.5	58.7, 186.7	59.2, 188.0
Resolution (Å)	50-1.05	51-1.35	12 – 1.90	51-1.45	40-1.80	51-1.45	26-1.95	49-1.50	49-1.60
	(1.07-1.05)	(1.42-1.35)	(1.97-1.90)	(1.53-1.45)	(1.90-1.80)	(1.53-1.45)	(2.06-1.95)	(1.55-1.50)	(1.66-1.60)
Unique reflections	177114 (8768)	75008 (9295)	29752 (2603)	65831 (8739)	35032(5039)	66408 (8907)	24769 (1325)	44421 (1371)	50821 (4336)
Multiplicity	5.2 (2.9)	5.8(2.6)	8.2 (3.8)	9.6(2.5)	7.3(3.9)	8.4(2.5)	9.1(2.3)	1.7 (1.1)	4.0 (3.2)
Average I/σ	57.0(3.4)	6.2 (1.7)	19.4 (6.9)	14.7(3.0)	6.5(2.2)	12.7(2.7)	34.3(9.6)	10.4 (0.82)	24.5 (6.0)
R <sub>meas</sub> (%)	9.3 (50.6) <sup>b</sup>	30.2 (46.9)	4.0 (15.3) <sup>b</sup>	9.5(29.0)	33.5(53.2)	13.0(39.5)	4.4(6.4)	5.4 (71.1) <sup>b</sup>	2.8 (16.5) <sup>b</sup>
Completeness (%)	99.8 (99.5)	91.0 (78.4)	97.6 (86.6)	97.9 (91.7)	99.9 (99.9)	98.1 (92.1)	85.8 (32.5)	72.9 (23.0)	98.6 (86.4)
<i>Refinement Statistics</i>									
Refinement method	Aniso	Aniso	TLS	Aniso	TLS	Aniso	TLS	TLS	TLS
Amino acid residues	302	309	309	309	309	309	309	309	309
Solvent atoms	646	484	413	491	297	500	556	451	530
Non-H atoms	3252	3117	3035	3160	2923	3217	3165	2999	3155
RMS bonds (Å)	0.017	0.009	0.009	0.008	0.010	0.008	0.010	0.009	0.009
RMS angles (°)	2.3	1.0	1.1	1.2	1.3	1.2	1.1	1.2	1.2
<B <sub>protein</sub> > (Å <sup>2</sup> )	21.7	14.6	28.9	20.9	34.5	21.3	17.8	26.7	24.0
<B <sub>FAD</sub> > (Å <sup>2</sup> )	15.1	10.4	18.3	16.6	22.0	16.7	13.1	21.1	18.5
<B <sub>NADP(H)</sub> > (Å <sup>2</sup> )	–	13.8	22.7	24.1	28.3	25.5	23.7	27.0	24.7
R <sub>work</sub> (%)	12.5	13.5	13.1	10.8	17.2	11.0	13.0	14.9	13.1
R <sub>free</sub> (%)	15.5	17.3	17.3	14.4	21.1	14.7	18.0	18.5	15.6
PDB code	3LO8	5VW4	5VW9	5VW3	5VW8	5VW2	5VW5	5VW6	5VW7

<sup>a</sup> Numbers in parentheses are in the highest resolution shell<sup>b</sup> R<sub>merge</sub> reported in place of R<sub>meas</sub>

**FIGURE LEGENDS**

**Figure 1. Aromatic placeholder and nicotinamide binding in three FNR superfamily members.** (A) Stacking of aromatic placeholder side chain onto the flavin in wild-type forms of spinach FNR (PDB 1FNC; Tyr314 in forest green, FAD in orange), cytochrome P450 reductase (PDB 1AMO; Trp677 in green, FAD in dandelion), and NO synthase (PDB 1F20; Phe1395 in pale green, FAD in sand). (B) Mode of nicotinamide binding in the NADP<sup>+</sup> complexes of Y303S *Anabaena* FNR (PDB 2BSA; NADP<sup>+</sup> in blue), W677X cytochrome P450 reductase (PDB 1JA0; NADP<sup>+</sup> in cerulean), and Y303S pea FNR (1QFY; NADP<sup>+</sup> in seafoam); FAD colors as in panel A.

**Figure 2. Effect of the replacements of the Tyr316 residue on the visible absorption spectrum of FNR and rescue of the wild-type spectral features by phenol binding to the Y316S variant.**

All spectra were determined in 50 mM Tris-HCl, pH 7.4 at 25 °C. (A) Extinction coefficients in the visible region of Y316S (solid line) and Y316A (dotted line) FNRs in comparison to that of wild-type enzyme (thin line). (B) Perturbation of the visible absorption spectrum of the Y316S variant (ca. 9 μM) induced by the presence of 150 mM phenol. The difference spectrum was rescaled (solid line) to match that expected for a theoretical enzyme concentration of 1 mM in order to be compared to the difference between the extinction coefficients of wild-type and Y316S FNR forms (dashed lines).

**Figure 3. Interaction of Y316S FNR with nicotinamide-containing ligands.** (A) Progress of the titrations of 10 μM Y316S FNR with NADP<sup>+</sup> (hollow circles), NAD<sup>+</sup> (filled circles), nicotinamide (hollow squares), and methyl-nicotinamide (filled squared). All titrations were performed in Tris-

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3 HCl, pH 7.7, at 15 °C. Data points and fitting curves were corrected to account for protein dilution.  
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5 The logarithmic concentration scale was chosen to allow comparison of ligands displaying huge  
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7 differences in affinity. (B) Computed difference spectra in the visible region of protein-ligand  
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9 complexes. The difference between extinction coefficients of complexed and uncomplexed protein  
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11 forms is shown for NADP<sup>+</sup> (solid line), NAD<sup>+</sup> (dashed line), nicotinamide (dash-dotted line), and  
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13 methyl-nicotinamide (dotted line) are shown. (C) Overlay of Y316A<sub>nic</sub> (slate) and Y316S<sub>nic</sub> (hot  
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15 pink) with nicotinamide bound, and as a reference, bound NADP<sup>+</sup> (ghosted grey). Hydrogen bonds  
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17 to bound nicotinamide are shown (slate and hot pink dashed lines in the respective structures).  
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24 **Figure 4. Effect of the presence of NADP<sup>+</sup> on photoreduction of Y316S FNR.** Significant  
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26 spectra recorded during the stepwise photoreduction of the FAD prosthetic group in the absence  
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28 (black lines) and in the presence of a slight excess of NADP<sup>+</sup> (red lines). About 16 μM protein  
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30 solutions in 10 mM HEPES, pH 7.0, were photoreduced at 15 °C in the presence of an EDTA-  
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32 deazariboflavin system. Spectra corresponding to the maximal accumulation of semiquinone  
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34 (dotted line) and charge-transfer complex (solid line) species are shown, and compared to those of  
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36 the oxidized (thin lines) and fully reduced (dashed lines) respective enzyme mixtures.  
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43 **Figure 5. Spectra of wild-type FNR and its variants recorded during anaerobic titrations**  
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45 **with NADPH and during NADPH-O<sub>2</sub> turnover. (Upper panels)** titrations of wild-type (A),  
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47 Y316S (B) and Y316A (C) FNRs showing the spectra of the free oxidized enzymes (thin black  
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49 traces) and those recorded after the addition of ca. 1.2, 2.5, 5, 12, and 120-fold excess of NADPH  
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51 (red, green, blue, magenta, and black bold traces, respectively). (Lower panels) Spectra recorded  
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53 at different reaction times after air was admitted into the cuvettes containing the previously  
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3 anaerobic solutions of the FNR forms in the presence of 120-fold excess NADPH. The spectra of  
4 the free oxidized FNR forms (thin black traces) and those of the respective CT complexes recorded  
5 in the presence of 1.2-fold NADPH excess (thin red traces) are reported for comparison. **(D)**  
6 Spectra recorded after 15 s, 25 min, 35 min, 40 min, 45 min, and 60 min (red, green, blue, magenta,  
7 lime, and black bold traces, respectively) turnover catalyzed by wild-type FNR. **(E)** Spectra  
8 recorded after 2.5 min, 6 min, 11 min, 19 min, 46 min, and 100 min (red, green, blue, magenta,  
9 lime, and black bold traces, respectively) turnover catalyzed by Y316S FNR. **(F)** Spectra recorded  
10 after 1 min, 4 min, 53 min, 2.5 h, 3.5 h, and 17 h (red, green, blue, magenta, lime, and black bold  
11 traces, respectively) turnover catalyzed by Y316A FNR.

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27 **Figure 6. Reanalysis of stopped-flow results reported for *Anabaena* FNR.** Data extrapolated  
28 from [26] were fit to a single-exponential decay. **(Upper panels)** Time-course of the approach to  
29 equilibrium upon mixing **(A)** wild-type FNR<sub>red</sub> and NADP<sup>+</sup>, and **(B)** wild-type FNR<sub>ox</sub> and  
30 NADPH. Curve fitting yielded the  $k_{\text{obs}}$  values of  $235 \pm 47 \text{ s}^{-1}$  and  $230 \pm 40 \text{ s}^{-1}$ , respectively,  
31 equivalent to each other within the error of the analysis. Lans *et al.* [26] reported the rate  
32 constants for these reactions to be  $285 \text{ s}^{-1}$  and  $270 \text{ s}^{-1}$ , respectively. **(Lower panels)** Time-course  
33 of the approach to equilibrium upon mixing **(C)** Y303S FNR<sub>red</sub> and NADP<sup>+</sup>, and **(D)** Y303S  
34 FNR<sub>ox</sub> and NADPH. Curve fitting yielded the  $k_{\text{obs}}$  values of  $165 \pm 16 \text{ s}^{-1}$  and  $145 \pm 20 \text{ s}^{-1}$ ,  
35 respectively, equivalent to each other within the error of the analysis.

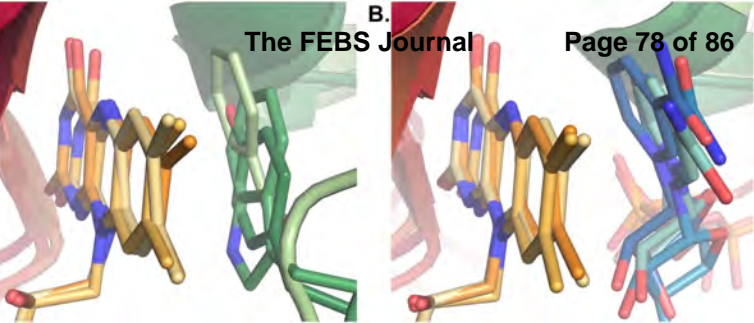
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51 **Figure 7. Active site environment of FNR (A)** A stereo view of the active site environment of  
52 FNR represented by the structure of Y316S<sub>NADP+</sub> with the 2F<sub>o</sub>-F<sub>c</sub> map shown at  $3.0 \cdot \rho_{\text{rms}}$ . Key  
53 residues within the active site, Ser94, Cys274, and Glu314, which primarily interact with  
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3 NADP(H), NADP(H), and FAD are shown along with hydrogen bonds (black dashed lines) and  
4 interaction distances (gray dashed lines). **(B)** Single crystal visible absorption spectra of Y316A  
5 FNR soaks with NADPH (right-hand panel) and NADP<sup>+</sup> (left-hand panel) and their change  
6 during data collection. Spectra collected corresponding to oscillation images 0, 10, 20, 50, 100  
7 and 360 go from light to dark shades of pink. These spectra qualitatively match those seen in  
8 solution in that the CTC-2 absorption band is seen for the crystals soaked in NADPH but not in  
9 those soaked in NADP<sup>+</sup>. The decrease in A<sub>460</sub> over time indicates reduction of the flavin in the  
10 X-ray beam during data collection. Over time the CTC-2 band does not disappear in the NADPH  
11 soaked crystal and does not appear in the NADP<sup>+</sup> soaked crystal so, we infer that at the cryo-  
12 temperatures of data collection, the flavin reduction is not leading to the same structural changes  
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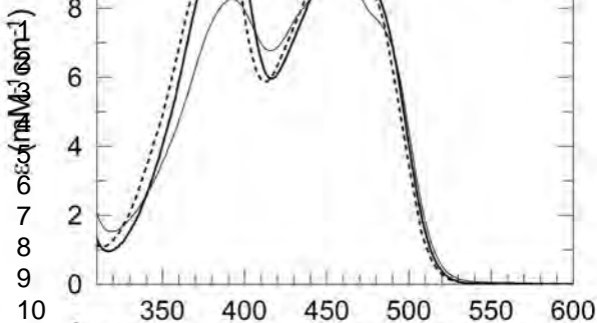
32 **Figure 8. Active site compression in corn root FNR as seen by mobility, covalent distortion,**  
33 **and interaction distances. (A)** Mobility of atoms in the active site of Y316S FNR. Atoms in the  
34 active sites of Y316S<sub>NADP<sup>+</sup></sub> (orange) (i) and overlaid with Y316S<sub>NADPH</sub> (semi-transparent green)  
35 (ii) shown with ellipsoids, denoting the use of anisotropic B-factors. The nitrogen and oxygen  
36 atoms are shown in blue and red, respectively, in the complexes. The mobility of the C4 of NADP<sup>+</sup>  
37 is in the direction of the N5 of the FAD, forming a boat-like conformation which is favored for  
38 hydride transfer. **(B)** Active site compression in corn root FNR as seen by covalent distortion. The  
39 angle of distortion of the FAD moiety is shown in: (i) FAD in wild-type corn root FNR (mauve),  
40 (ii) FAD in Y316S<sub>NADP<sup>+</sup></sub> (orange) and the corresponding angles from Y316S<sub>NADP<sup>+</sup></sub> (P3<sub>121</sub>, olive)  
41 and Y316A<sub>NADP<sup>+</sup></sub> (salmon), (iii) FAD in Y316S<sub>NADPH</sub> (green) and the corresponding angle from  
42 Y316A<sub>NADPH</sub> (violet). **(C)** Active site interactions in FNR. Structural overlay of Y316S<sub>NADP<sup>+</sup></sub>  
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3 (orange) and Y316S<sub>NADPH</sub> (green) with relevant average distances shown in corresponding colors.  
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5 The average distances from Y316S<sub>NADP+</sub> (P3<sub>121</sub>, olive), Y316A<sub>NADP+</sub> (salmon), and Y316A<sub>NADPH</sub>  
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7 (violet) are also shown. The distances are averages from ten independent refinements of each  
8  
9 complex (see Materials and Methods), and all standard deviations were < 0.02 Å. The C4H and  
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11 N5H distance in Y316S<sub>NADPH</sub> is 3.0 Å (not labeled). If modeled as FADH<sub>2</sub> instead of FAD, the  
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13 theoretical distance between C4H and N5H in Y316S<sub>NADP+</sub> would be 2.5 Å. Nitrogen, oxygen, and  
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15 sulfur atoms colored blue, red, and yellow, respectively.  
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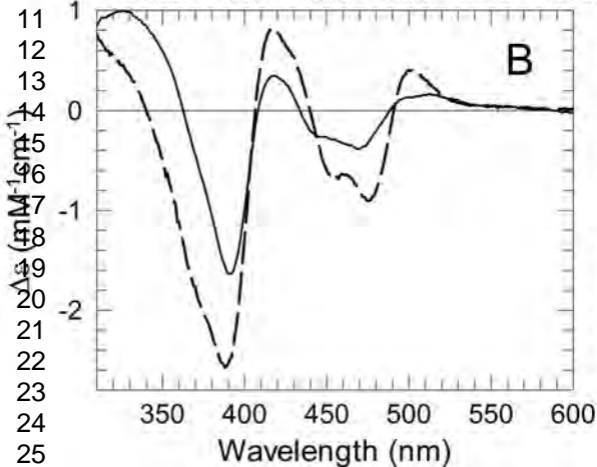
22 **Figure 9. Sequence alignment of corn root FNR with NADPH oxidases. (A)** Sequence  
23 alignment of segments of corn root FNR with human NOX1, NOX2, NOX3, NOX4, and NOX5  
24 and *C. stagnale* NOX5. Select FNR functional residues and similar aligned residues (in bold) have  
25 their function denoted as: aromatic placeholder (a), other catalytic center (c), 2'-phosphate (p) or  
26 other NADP interactions (n). Residue numbers are given for the structurally known corn root FNR  
27 and *C. stagnale* NOX5. **(B)** Overlay of the *C. stagnale* NOX5 dehydrogenase domain (PDB 5O0X;  
28 pink with bound FAD and the natural C-terminal Phe693 and C-terminal extension Trp side chains  
29 in purple) and the Y316S<sub>NADP+</sub> structure (orange with bound FAD and NADPH in yellow). The  
30 anticipated fifth β-strand of the NADP<sup>+</sup>-binding domain of NOX5 which is perturbed and allows  
31 the Trp residue to stack against the isoalloxazine ring is highlighted (purple trace) and an arrow  
32 indicates the direction we expect it to move in the full-length NOX5 context so that it will align  
33 with the rest of the β-sheet as does the fifth β-strand in FNR. The deposited structure of the  
34 NADPH binding domain of NOX2 gp91(phox), determined through structural genomics efforts  
35 (PDB 3A1F) but not yet described in the literature, also aligns well in a structural overlay.  
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7**The FEBS Journal****B.****Page 78 of 86**

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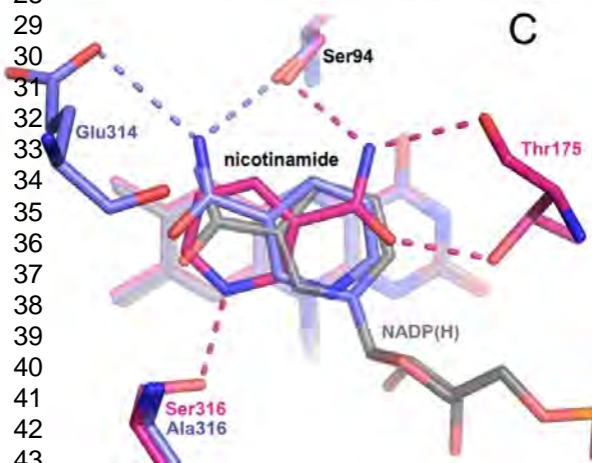
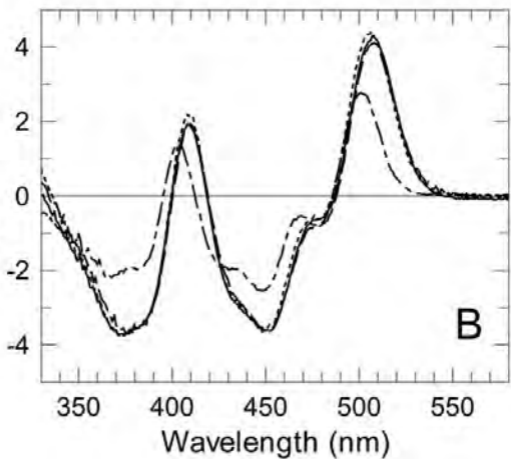
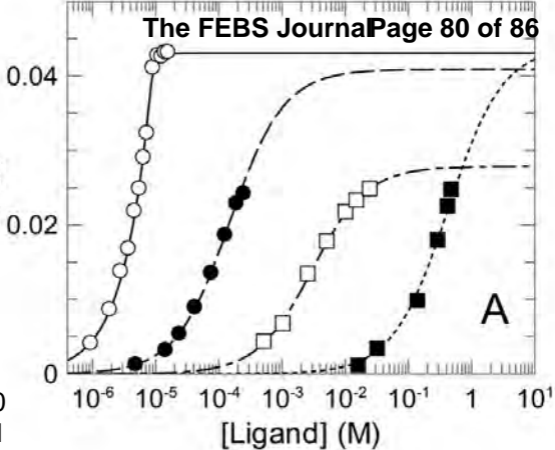


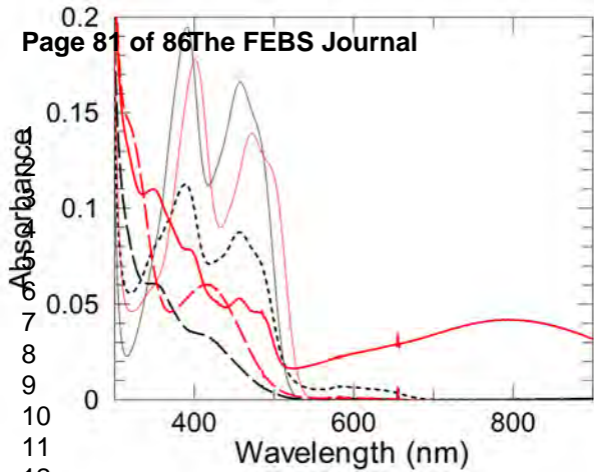
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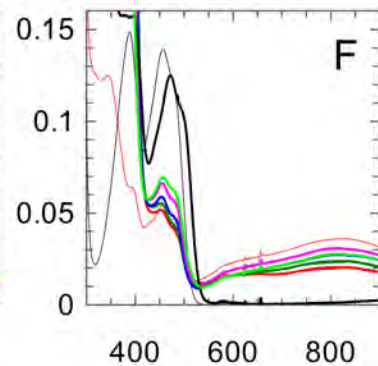
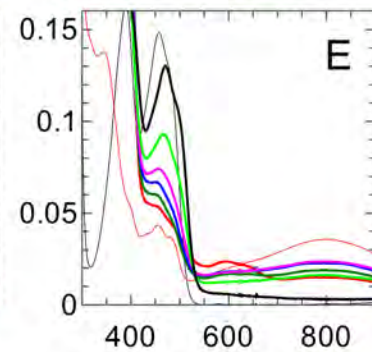
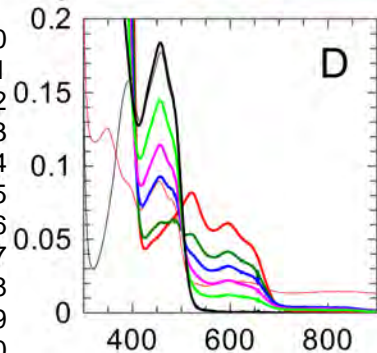
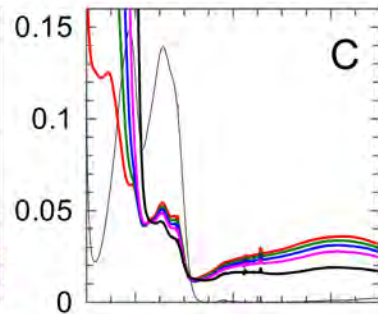
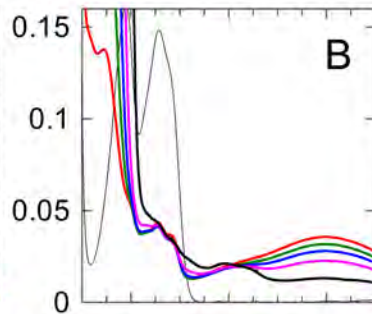
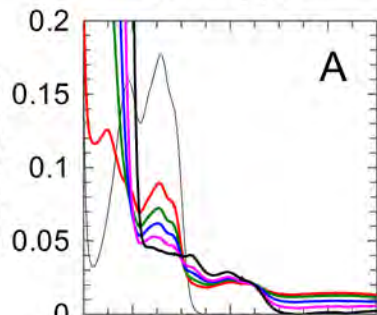


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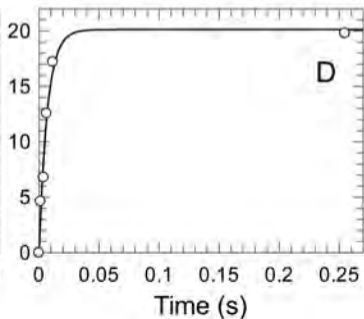
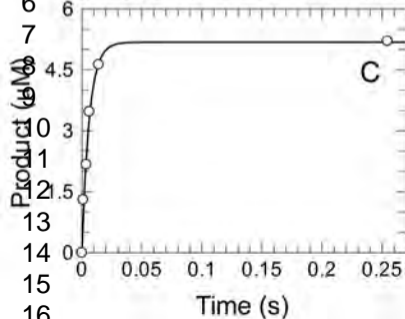
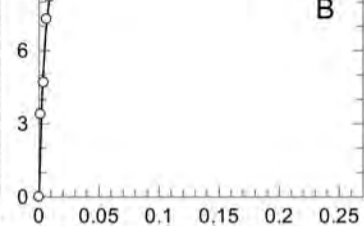
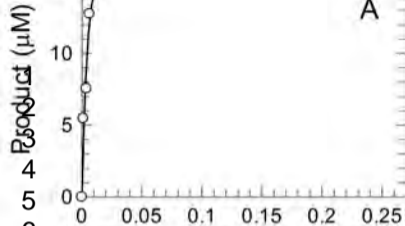
The FEBS Journal  
Y316S

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Page 82 of 86

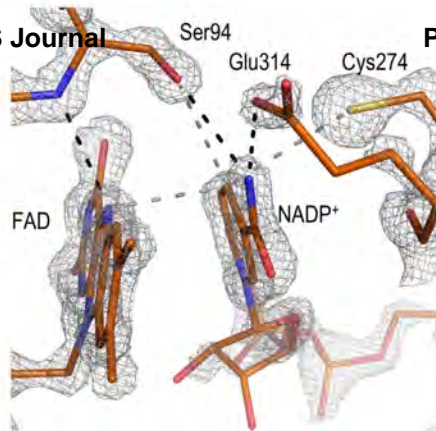
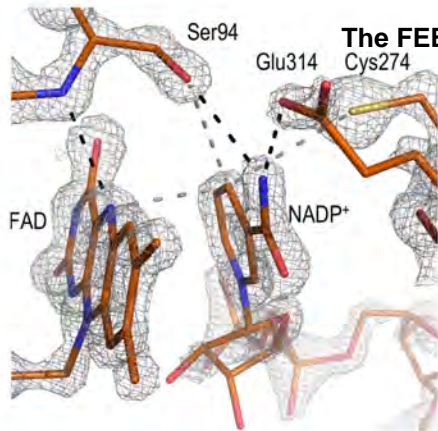
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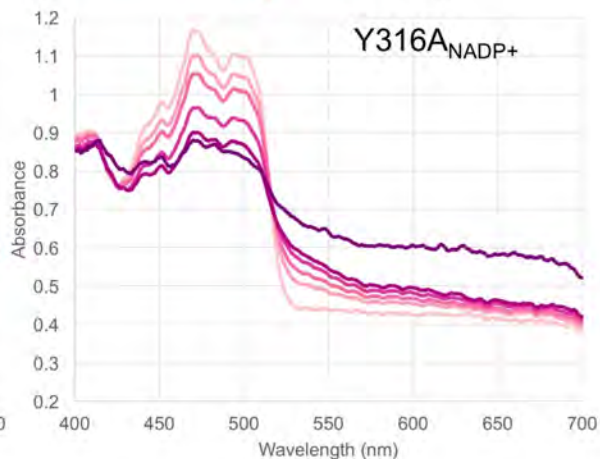
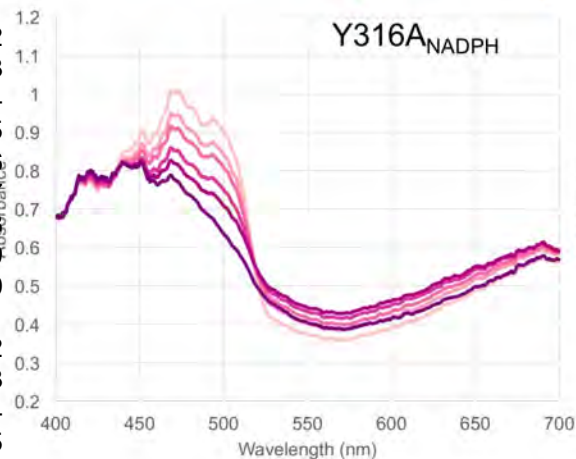


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Page 84 of 86

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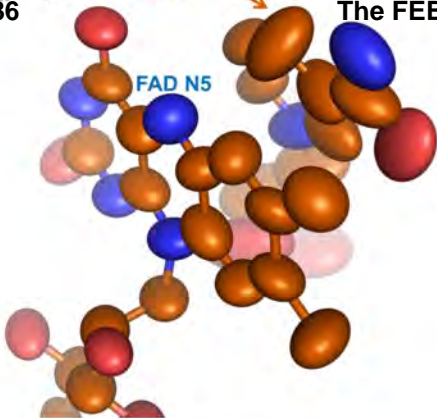


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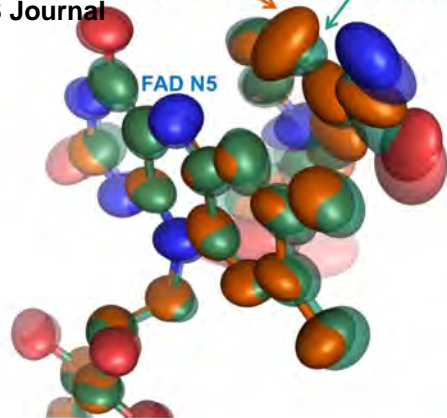
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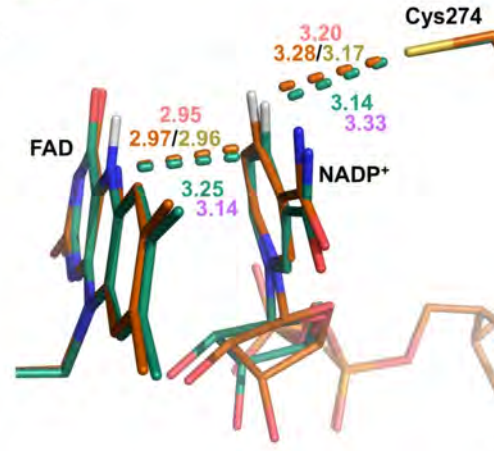
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**A.** corn root FNR ...<sup>31</sup>RLYSIAS...<sup>113</sup>VRRA...<sup>172</sup>ATGTG...<sup>237</sup>SRE...<sup>246</sup>GKMYVD...<sup>273</sup>FCGLK...<sup>312</sup>HVEVY  
 human NOX1 ...HPFTLTS...<sup>312</sup>VRRA...<sup>312</sup>FCGLK...<sup>312</sup>HVEVY  
 human NOX2 ...HPFTLTS...<sup>1</sup>IRRS...<sup>1</sup>GAGIG...<sup>1</sup>TGW...<sup>1</sup>LYGRPNW...<sup>1</sup>LCGPE...<sup>1</sup>NKENF  
 human NOX3 ...HPFTLTS...<sup>1</sup>IRQA...<sup>1</sup>AAGIG...<sup>1</sup>TGW...<sup>1</sup>FYGRPNW...<sup>1</sup>FCGPK...<sup>1</sup>NKESF  
 human NOX4 ...HPFTLTM...<sup>1</sup>IRVS...<sup>1</sup>AGGIG...<sup>1</sup>SQT...<sup>1</sup>FIGRPNW...<sup>1</sup>CCGPN...<sup>1</sup>NKESFS  
 human NOX5 ...HPFTISS...<sup>1</sup>IRRS...<sup>1</sup>GAGIG...<sup>1</sup>TSA...<sup>1</sup>QPGRPRW...<sup>1</sup>FCGSP...<sup>1</sup>FQENF  
 1 *C. stagnale* NOX5 ...<sup>459</sup>HPFTISS...<sup>477</sup>IRAV...<sup>535</sup>CAGIG...<sup>604</sup>TGA...<sup>641</sup>KTGRPDW...<sup>667</sup>FCGPT...<sup>689</sup>RKENF  
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