Expression of multiple forms of ferredoxin NADP⁺ oxidoreductase in wheat leaves

J. O. Gummadova¹, G. J. Fletcher¹, A. Moolna¹, G. T. Hanke², T. Hase² and C. G. Bowsher¹,*

¹ Faculty of Life Sciences, The University of Manchester, 3. 614 Stopford Building, Oxford Road, Manchester M13 9PT, UK
² Division of Protein Chemistry, Institute for Protein Research, 3-2 Yamadaoka, Suita, Osaka 565-0879, Japan

Received 13 August 2007; Revised 7 September 2007; Accepted 11 September 2007

Abstract

In higher plants there are two forms of ferredoxin NADP⁺ oxidoreductase (FNR), a photosynthetic pFNR primarily required for the photoreduction of NADP⁺, and a heterotrophic hFNR which generates reduced ferredoxin by utilizing electrons from NADPH produced during carbohydrate oxidation. The aim of this study was to investigate the presence of multiple forms of FNR in wheat leaves and the capacity of FNR isoforms to respond to changes in reductant demand through varied expression and N-terminal processing. Two forms of pFNR mRNA (pFNRI and pFNRII) were expressed in a similar pattern along the 12 cm developing primary wheat leaf, with the highest levels observed in plants grown continuously in the dark in the presence (pFNRII) or absence (pFNRI) of nitrate respectively. pFNR protein increased from the leaf base to tip. hFNR mRNA and protein was in the basal part of the leaf in plants grown in the presence of nitrate. FNR activity in plants grown in a light/dark cycle without nitrate was mainly due to pFNR, whilst hFNR contributed significantly in nitrate-fed plants. The potential role of distinct forms of FNR in meeting the changing metabolic capacity and reductant demands along the linear gradient of developing cells of the leaf are discussed. Furthermore, evidence for alternative N-terminal cleavage sites of pFNR acting as a means of discriminating between ferredoxins and the implications of this in providing a more effective flow of electrons through a particular pathway in vivo is considered.

Key words: Ferredoxin NADP⁺ oxidoreductase (FNR), light, nitrogen assimilation, Triticum aestivum.

Introduction

Ferredoxin NADP⁺ oxidoreductase (FNR, EC 1.18.1.2) is a ubiquitous enzyme belonging to a large family of flavoproteins found in higher plants, eukaryotic algae, photosynthetic bacteria, and animals (Arakaki et al., 1997). In plants, this enzyme exists in two distinct forms as a photosynthetic (pFNR) and as a heterotrophic (hFNR) form. These two isoforms are encoded by different genes, linked to different metabolic pathways and are regulated by different factors. Six genes have been identified as producing FNR-like proteins in the Arabidopsis genome (Wang et al., 2003). Of these, four have been further confirmed as producing FNR proteins, that is two pFNR and two hFNR proteins (Hanke et al., 2005). Two pFNR isoforms have been identified in Oryza sativa and three in Zea mays (maize) (Okutani et al., 2005; Ohyanagi et al., 2006). Traditionally, pFNR has been considered to catalyse the final step in the photosynthetic electron transport chain. Based on its association with the cytochrome b₅f complex (Zhang and Cramer, 2004) and NADP(H) dehydrogenase complex (Quiles and Cuello, 1998) and on FNR-dependent quinone reduction measurements (Bojko et al., 2003), pFNR has also been implicated to have a role in cyclic electron transport. By contrast, a role for hFNR has been characterized for supplying reducing power in non-photosynthetic tissue to a variety of metabolic processes including nitrate assimilation.
(Bowsher et al., 1989, 1992). Sharing similar kinetic properties, pFNR and hFNR drive the same redox reaction but in opposite directions, with pFNR primarily reducing NADPH using electrons supplied by the photosynthetically reduced ferredoxin (Fd), and hFNR generating reduced Fd by accepting electrons from NADPH. pFNR has long been known to be regulated by light at both the transcriptional level (Neuhaus et al., 1993, 1997; Oelmüller et al., 1993; Bowler et al., 1994a, b) and the level of enzyme activity (Carrillo et al., 1981; Nikolaeva and Osipova, 1982; Pschorn et al., 1988a, b; Rühle et al., 1987). However, recent work has also shown that two Arabidopsis pFNR isoforms are differentially regulated by nitrate (Hanke et al., 2005). hFNR has been shown to be regulated by nitrate at both the transcriptional and enzyme levels (Bowsher et al., 1993b; Ritchie et al., 1994; Matsumura et al., 1997; Mattana et al., 1997; Wang et al., 2000, 2003; Hanke et al., 2005).

Despite the presence of multiple FNR isoforms and their importance in supporting a number of distinct processes which may operate simultaneously, few studies have examined whether the two differently regulated pFNR and hFNR exist in the same tissue or whether they have the potential to regulate different metabolic processes simultaneously. Both pFNR and hFNR have been detected in tomato fruit tissue (Green et al., 1991). In immature mung bean (Vigna radiata L.) immunoincubation and immunoblottings studies showed that with development and greening of young leaves hFNR was gradually replaced by pFNR (Jin et al., 1994; Wada et al., 1996). Such changes in abundance may be regulated, responding to the demand for specific metabolic processes by the plant during development and by the environment. Indeed two isoforms of hFNR have recently been identified in Arabidopsis leaves and found to be at the highest levels when grown in the presence of ammonium (Hanke et al., 2005). These hFNR isoforms or proteins have been proposed to meet the specific needs of non-photosynthetic cells in the leaf by supplying reduced Fd to support ammonium assimilation (Hanke et al., 2005). The 12 cm long developing primary wheat leaf has been well characterized as providing a profile of developing cells with a linear gradient of cell types. Using this developmental model of heterotrophic cells from the leaf meristem at the base of the leaf to photosynthetic cells at the tip of the leaf (Leech, 1985) provides important information as to whether, with increasing cell age, different FNR isoforms respond to changing metabolic demands during the development of photosynthesis.

This paper focuses on the identification and purification of multiple forms of FNR in wheat leaves. We first purified FNR and obtained N-terminal amino acid sequence data to investigate the FNR forms present in mature wheat leaves. Using an available wheat leaf cDNA library, FNR clones were then isolated. The effects on pFNR and hFNR transcription, protein, and activity in wheat grown in different light and nitrate environments were examined. pFNR with alternative N-terminal cleavage sites was overexpressed in Escherichia coli and, following purification, used to determine whether truncation led to potential differences in the enzymes affinity for photosynthetic and heterotrophic Fd. This complete study of the identified FNR isoforms allows us to speculate on the capacity of FNR isoforms to respond to changes in reductant demand through varied expression and N-terminal processing.

Materials and methods

Plant materials and growth conditions

For protein purification, spring wheat (Triticum aestivum var. Axona) was grown in John Innes No. 1 compost in a greenhouse between 15 °C and 25 °C, supplemented by sodium lights on a 16 h light (240 μmol m⁻² s⁻¹/8 h dark cycle. All green leaves were harvested between 20 and 30 d post-anthesis and stored at −20 °C until required. To examine the developmental gradient, spring wheat, Triticum aestivum L. cv. Maris Huntsman, was grown in sterile Magenta vessels on 50–100 ml 0.2% (w/v) phytagel nutrient media as described by Paul and Stüt (1993). Plants were grown in a controlled environment cabinet at 20 °C in either a 16 h light (240 μmol m⁻² s⁻¹/8 h dark cycle or continuous dark in the presence or absence of 10 mM KNO₃. Plants were harvested 4–5 h into the photoperiod after 6–7 d when leaf height had reached approximately 12 cm. Seedlings were removed from the phytagel and the roots separated from the shoots with a razor blade. Primary leaves were dissected from the seedling and then cut into six 2 cm sections, numbered section 1 at the base to section 6 at the tip. All samples were either used immediately or frozen in liquid nitrogen and stored at −80 °C until required. When using dark-grown tissue, all harvesting and extraction steps were performed in a dark room with a green safe light.

Purification of FNR from wheat leaf tissue

One kilogram of frozen wheat leaves were ground to a fine powder in a mortar and pestle in 100 g batches. All subsequent steps were carried out at 4 °C unless otherwise stated. The frozen powder was added to 2.0 l of extraction buffer [100 mM Tris-borate buffer, pH 7.7, 0.1% (v/v) β-mercaptoethanol, 1.5 % (w/v) polyvinylpyrrolidone, 2 mM benzamidine, 2 mM amino-n-caproic acid, 0.2 mM phenylmethlysulphonyl fluoride, 20 mM diethyliothiocarbamate] and homogenized with a polytron into a green slurry. Following acetone precipitation (33–77%, v/v) the resulting pellet was resuspended in buffer A (10 mM TRIS-HCl, pH 7.5) and dialysed overnight against buffer A before applying to a DEAE cellulose column (2.6 cm×100 cm). The FNR was eluted with buffer B containing 100 mM NaCl and following ammonium sulphate precipitation (40–60%, w/v) the precipitate was resuspended in buffer A and dialysed against the same buffer. The sample was loaded onto a DEAE Sephacel column (1.8 cm×50 cm) and after washing with buffer A containing 100 mM NaCl, the FNR was eluted with buffer A containing 200 mM NaCl and dialysed against buffer A. The FNR sample was further purified using a ferredoxin affinity column and eluted by buffer A containing 50 mM NaCl. Following dialysis against buffer A the FNR was concentrated using a microconcentrating column (10 KDa cut-off, Amicon).

Protein extraction

Leaf soluble proteins were extracted by grinding 8–10 frozen leaf sections in 0.8 ml of extraction buffer (100 mM phosphate buffer,
pH 7.8) in a Retsch vibration mixer for 2 min. The extracts were then centrifuged at 13 000 g for 5 min at 4 °C. The supernatant was transferred to a fresh tube and centrifuged again. The resulting supernatant was immediately used for protein estimation, enzyme activity assays, and immunotitration. Any remaining sample was stored at –80 °C for SDS PAGE analysis.

**Determination of protein concentration**

Protein concentration was determined using the method of Bradford (1976), or, where limited volume was available, the method of Schaffner and Weissmann (1973).

**Enzyme assays**

To monitor FNR activity in the purified proteins and in tissue sections, a cytochrome c reduction assay system in 0.6 ml containing 50 mM TRIS-HCl, pH 7.5, 200 μM cytochrome c, 100 mM NaCl, and 50 μM NADPH in the presence of an NADPH regenerating system was used (Onda et al., 2000). The reaction was carried out at room temperature and initiated by the addition of maize FdI or FdIII at final concentrations ranging from 0–40 mM. Cytochrome c reduction was measured by monitoring the increase in absorbance at 550 nm. Where appropriate the kinetic constants $K_m$ and $V_{max}$ were calculated with non-linear regression analysis using the Solver function in Microsoft Excel 97 (Microsoft, Redmond, WA).

To monitor FNR activity eluted from columns, diaphorase activity was measured in an NADPH-dependent reduction of nitroblue tetrazolium (Swegle and Mattoo, 1996), using a microtitre plate assay. 100 mM TRIS-HCl, pH 7.5, 1 mM nitro-blue tetrazolium (Swegle and Mattoo, 1996), using a microtitre plate assay was measured in an NADPH-dependent reduction of nitro-blue tetrazolium (Swegle and Mattoo, 1996), using a microtitre plate assay. Where appropriate the kinetic constants $K_m$ and $V_{max}$ were calculated with non-linear regression analysis using the Solver function in Microsoft Excel 97 (Microsoft, Redmond, WA).

**SDS polyacrylamide gel electrophoresis (PAGE)**

The SDS-PAGE of proteins was performed on 12.5% acrylamide gels according to Hames and Rickwood (1990). The gels were stained for protein with Coomassie Brilliant Blue (0.1% w/v Coomassie Brilliant Blue, 25% v/v methanol, 8% v/v acetic acid, 67% v/v water) or silver (Wray et al., 1981).

**Preparation of protein samples for N-terminal-amino-acid determination**

SDS-PAGE of FNR, blotting and amino acid sequence determination was performed as described previously (Bowsher et al., 1993a).

**Isoelectric point determination**

The isoelectric point of mature proteins was calculated using the available cDNA sequence information and the predictive program at EXPASY (http://www.expasy.org/tools/pi_tool.html). To determine the isoelectric points experimentally 0.1–1 μg overexpressed pFNR proteins were loaded onto 18 cm polyacrylamide gel strips with an immobilized linear gradient of pH 4–7 (Immobiline DryStrip, Amersham) and focused in a Protein isoelectric focusing cell (Bio-Rad) according to the manufacturers instructions. The second dimension was run with 15% SDS-PAGE using the Protein II xi vertical gel electrophoresis system (Bio-Rad). The location of the pFNR spots was visualized using either blue-silver Coomassie G-250 total protein staining (Candiano et al., 2004) or by immunoblotting with the PEPpFNR antibody. All gels and blots were scanned using Quantity One gel imaging software and the GS-710 calibrated imaging densitometer (Bio-Rad). The positions of each of the pFNR spots were measured according to the linear pH gradient and were consistent in four independent replicates.

**Antibody production**

Synthetic peptides were designed to produce PEPpFNR antisera based on information obtained from the Swissprot and Trembl databases of protein sequences (http://www.expasy.ch). In the absence of a complete amino acid sequence for wheat, FNR peptides were selected based on the pFNR sequence of rice, a related species (Aoki et al., 1994). hFNR and pFNR sequences were aligned using the Clustal W multiple sequence alignment program. Disparate regions of between 12–15 amino acids were selected from the rice pFNR amino acid sequence (Aoki et al., 1994) on the basis of low amino acid identity with rice root hFNR (Aoki and Ida, 1994). Regions were identified with less than 34% identity to hFNR and then mapped onto a three-dimensional model of pFNR based on the Karplus crystallographic structure of spinach pFNR (Karplus, 1991). Two surface-orientated peptides, peptide I, ISKKH-DEGVVPNK, and peptide IV, ADGDKNKGKPHKL, located in continuous domains of the Karplus crystallographic structure were used to synthesize multiple antigen peptides. One mg of each peptide was dissolved in 1 ml of 25 mM TRIS-HCl, pH 8.0, divided into three aliquots and each aliquot mixed with an equal volume of Freund’s adjuvant prior to injecting a sheep at monthly intervals (Diagnoscis Scotland). Three individual bleeds were taken from the sheep and dilutions of the final bleed used for all subsequent experiments without further purification.

**Immunoblotting**

For immunoblotting, following separation by SDS-PAGE proteins were electroblotted to a nitrocellulose or PVDF membrane and incubated with antibody raised against pFNR or hFNR. The antigen–antibody complex was visualized by reaction with alkaline phosphatase or horseradish peroxidase conjugated secondary antibody. Equal loading of protein samples was confirmed by total protein staining of the blots with Ponceau S solution prior to incubation with the specific antibody.

**Immunotitration using pea hFNR antiserum**

Pea hFNR antisemur (Bowsher et al., 1993a) was diluted with 100 mM TRIS-HCl pH 7.6 to 50% (v/v). One volume of leaf section extract was mixed with 1 vol. of the diluted antiserum and incubated with gentle shaking (150 rpm) at 4 °C for 90 min. The antibody–antigen complex was removed from the solution by centrifugation at 13 000 g for 30 min at 4 °C. The supernatant was carefully transferred to a fresh tube and assayed for cytochrome c reduction activity. Control serum had no effect on the FNR activity. Immunoblots of samples confirmed that immunodepletion of samples had occurred.

**RNA preparation**

RNA was prepared using LiCl precipitation based on the method of Knight and Gray (1994). mRNA was isolated from total RNA based on Poly (A) Quick mRNA isolation kit (Stratagene).

**RNA hybridization**

For RNA blot hybridization 10–20 μg of the isolated total RNA from each section was denatured with formaldehyde and subjected to electrophoresis and blotted onto a Hybond N+ membrane as described previously (Fonseca et al., 1997). The probe used for the hybridization was either a 1 400 bp Pisum sativum pFNR cDNA (Newman and Gray, 1988), a 252 bp partial clone for wheat hFNR, or a 1 000 bp Triticum aestivum rRNA cDNA clone. In all cases, DNA probes were radiolabelled with [α-32P]-dCTP using a random priming kit (Stratagene), prehybridized, hybridized, and exposed to film as described previously (Fonseca et al., 1997). Autoradiograms
were scanned and relative amounts of mRNAs determined as described previously (Fonseca et al., 1997).

**RNA loading control and relative mRNA levels**

All the RNA blots were incubated with specific probes under the same conditions and subsequently exposed to film for the same period of time. To equalize the total amount of RNA run on the gels for loading error and to allow comparisons of gene expression using the wheat leaf under different environmental conditions to be analysed statistically, all the membranes were reprobed with a 1 000 bp wheat tRNA cDNA as a housekeeping gene. Again, as with the specific probes, all the membranes were tested under the same conditions. The reprobed membranes were exposed for identical periods of time. The detected signals were scanned and quantitated by the densitometry analysis program AIDA200. The relative mRNA level of the gene was estimated as a ratio of its band density to that of the respective tRNA band. At least three replicates were performed for each treatment.

**Library screening**

A high-titre complementary DNA library was initially prepared with mRNA isolated from wheat (*Triticum aestivum* cv. Maris Huntsman) leaves grown for 14 d in the presence of 10 mM nitrate, using the ZAP Express cDNA synthesis kit and ZAP Express cDNA Gigapack III gold cloning kit and according to the manufacturer’s instructions (Stratagene Ltd). More than 300 000 phage were used to manufacture cDNA using the In-promII Gigapack III gold cloning kit and according to the manufacturer’s instructions (Stratagene) as described previously (Davis et al., 1996), or a 1 200 bp *Oryza sativa* hFNR cDNA (Aoki and Ida, 1994) fragment as a probe, radiolabelled with [α-32P]dCTP (3 000 Ci mmol−1) using Prime-it II Random Primer Labelling kit (Stratagene) as described previously (Davis et al., 2003). Following overnight hybridization at 50–65 °C and depending on the stringency required, filters were washed sequentially using 2× SSC and 0.1% SDS, 1× SSC and 0.1% SDS, and 0.1× SSC and 0.1% SDS at 50–65 °C. Positive clones were confirmed by subsequent secondary and tertiary screens.

**Reverse transcription-polymerase chain reaction**

Total RNA (0.05–0.5 μg) from wheat roots or basal leaf 2 cm sections was used to manufacture cDNA using the In-promII reverse transcription kit (Promega) according to the manufacturer’s instructions. Degenerate primers for polymerase chain reaction (PCR) were as follows: hFNR 5′-AGAGGAAGATCCGCAATGGCT-3′ and 5′-GCAATGCATCTGGCAGTCA-3′. Each PCR was performed with 1/10, 1/100, and 1/1000 dilutions of cDNA. PCR was hot started at 94 °C and carried out for 30 cycles of 94 °C for 1 min, 50 °C for 1 min, and 72 °C for 2 min. Following agarose gel electrophoresis, products were visualized by ethidium bromide staining.

**PCR amplification of the 5′ ends of pFNRI and pFNRII**

Primers for PCR amplification of the 5′ end of pFNRI cDNA coding region were as follows: 5′-GGCCAGCGCGCTCCTCACC-3′ and 5′-GACTTGGGACCCAGCTGACA-3′. Primers for PCR amplification of 5′ end of pFNRII cDNA coding region were as follows: 5′-ATGGCGCGCCACGCTGACA-3′ and 5′-GATGTCTGACC-GGCTTG-3′. PCR was hot started at 94 °C and carried out for 30 cycles of 94 °C for 45 s, 55 °C for 45 s, and 72 °C for 2 min, followed by a final extension at 72 °C for 7 min.

**Sequence alignment**

The FNR sequences were aligned using the Kalign multiple sequence alignment algorithm at http://msa.cgb.ki.se/cgi-bin/msa.
data obtained with that previously published for rice pFNR. Band IV showed no identity with either pFNR or hFNR.

**Isolation of wheat leaf FNR clones**

Following the final screen of the wheat leaf library, eight cDNA clones of varying length (1 350–3 050 bp) were sequenced. Based on their nucleotide similarity, the clones represented two main groups pFNRI and pFNRII. The nucleotide sequences corresponding to the coding regions of both clones and their deduced protein sequences were submitted to the GenBank database and given the accession numbers AJ457979 (CAD30025) and AJ457980 (CAD30024), respectively. pFNRI is composed of 1 047 bp, whereas pFNRII consists of 1 092 bp (Fig. 3). pFNRI and pFNRII share 80% overall similarity at the amino acid level. However, the N-terminal 45–55 amino acids, corresponding to the transit peptide region, share less than 25% sequence identity. Using the available N-terminal sequence data, alternative cleavage points of the chloroplast transit peptide for pFNRI and pFNRII could be identified in each cDNA clone. The mature protein regions of both pFNRI and pFNRII showed good sequence identity to other leaf FNRs (Fig. 3). The deduced amino acid sequence of wheat pFNRII had 84% identical residues with maize pFNRI (Onda et al., 2000), whilst the wheat pFNRII had 76% sequence identity to maize pFNRII and 88% sequence identity to maize pFNRII. These data, in conjunction with the N-terminal cDNA data, were used to predict the isoelectric point of the alternative mature proteins purified from wheat leaves (Table 1). pFNRII is notably more acidic than pFNRI.

The alternative N-terminal cleavage sites for pFNRI, pFNRIKKVS, and pFNRISKKQ appear to affect the predicted pI with the shorter pFNRISKKQ having a markedly more acidic isoelectric point. By contrast, the alternative N-terminal cleavage sites for pFNRII, pFNRIIISKK, and pFNRIIKKQD appear to have little effect on the predicted isoelectric point of the proteins.

When the wheat leaf library was screened using available rice (Aoki et al., 1994) or pea (Bowsher and Knight, 1996) root hFNR cDNA fragments as probes, no corresponding hFNR cDNA clones were identified. In an alternative RT-PCR approach, multiple alignment of available plant hFNR sequences was used to identify consensus sequences which could be used to design degenerate primers. A 252 bp product was amplified using total RNA from either nitrate-fed wheat roots or the leaf base. Alignment with known hFNR cDNAs confirmed that the isolated PCR product belonged to the hFNR class of proteins (Fig. 4). The wheat hFNR fragment shared 90–94% identity with rice (Aoki and Ida, 1994) and maize hFNR (Ritchie et al., 1994) and 57% with both the wheat pFNRI and pFNRII reported here.
Fig. 3. ClustalW multiple alignment of the deduced amino acid sequences of wheat pFNRI, pFNRII with some known pFNRI amino acid sequences. Sequence alignments were carried out as described in the Materials and methods. The alignment was performed using the deduced amino acid sequences of maize (Okutani et al., 2005; Onda et al., 2000), rice (Ohyanagi et al., 2006) and Arabidopsis (Hanke et al., 2005). Similar residues are shown in dark grey, strongly similar residues in grey, weakly similar in light grey.
Comparison of overexpressed wheat proteins with alternative N-terminal cleavage points

To investigate further whether there was any functional significance related to alternative N-terminal start sites, pFNRIKKVS, pFNRISKKQ, pFNRIIISKK, and pFNRIIKKQD were overexpressed in E. coli and purified using a ferredoxin affinity column as described previously (Onda et al., 2000). The pFNR proteins purified from recombinant bacterial cells showed absorption spectra typical of flavin containing enzymes (data not shown).

Table 1. Summary of the effect of N-terminal truncation on the isoelectric point (pI) of pFNRI and pFNRII

<table>
<thead>
<tr>
<th>pFNRI isoform</th>
<th>N-terminal sequence</th>
<th>pI (Predicted)</th>
<th>pI (Actual)</th>
<th>Molecular weight (kDa)</th>
</tr>
</thead>
<tbody>
<tr>
<td>pFNRII</td>
<td>KKVSQKE</td>
<td>8.12</td>
<td>6.73</td>
<td>33.52</td>
</tr>
<tr>
<td>pFNRI</td>
<td>SKQK</td>
<td>6.82</td>
<td>6.40</td>
<td>33.17</td>
</tr>
<tr>
<td>pFNRII</td>
<td>ISQQQD</td>
<td>5.50</td>
<td>5.42</td>
<td>33.49</td>
</tr>
<tr>
<td>pFNRII</td>
<td>KQD</td>
<td>5.50</td>
<td>5.44</td>
<td>33.29</td>
</tr>
</tbody>
</table>

Fig. 4. ClustalW multiple alignment of the deduced amino acid sequence of the wheat 252 bp hFNR cDNA fragment with other plant hFNRs. The alignment was performed using the deduced amino acid sequences of rice (Aoki and Ida, 1994), maize (Ritchie et al., 1995), and Arabidopsis (Hanke et al., 2005). Similar residues are shown in dark grey, strongly similar residues in grey, weakly similar in light grey.
isolectric point of each overexpressed protein was determined by isoelectric focusing and compared with the predicted values (Table 1). As expected, the pFNRII proteins had more acidic pIs than those of the pFNRI proteins. Furthermore, N-terminal truncation of pFNRII did lead to a more acidic pI, although the shift was not as marked as that predicted (Table 1). The kinetic properties of purified pFNRI proteins were determined using either photosynthetic type ferredoxin (FdI) or heterotrophic ferredoxin (FdIII) from maize as described for maize FNR (Onda et al., 2000; Table 2). Irrespective of ferredoxin type, recombinant pFNRII proteins had a 2-fold higher activity than pFNRI isoforms, but considerably lower affinity. Differences in the N-terminal of recombinant pFNRI, did not greatly affect $K_m$ and $V_{max}$ values with either FdI or FdIII, although the shorter pFNRI$^{KVS}$ had consistently lower activity than the longer pFNRI$^{KVS}$. By contrast, the longer version of pFNRII, pFNRII$^{KKQ}$ discriminated between photosynthetic and heterotrophic ferredoxins (having an approximate 3-fold lower $K_m$ for the heterotrophic FdIII) while the shorter pFNRII$^{KQD}$ did not. The removal of two N-terminal residues from pFNRII appeared to increase the affinity for FdIII to approximately the same value as FdI.

**pFNRI and hFNR expression in different light and nitrogen conditions**

FNR is involved not just in photosynthesis but also in providing reductant for a range of ferredoxin-dependent reactions. Previous experiments using microarray studies and the transfer of plants between different nitrogen regimes have confirmed isoform specific involvement of FNR in Arabidopsis (Wang et al., 2000; Hanke et al., 2005) and maize (Okutani et al., 2005). To monitor changes in pFNRI or hFNR relative steady-state transcript abundance along the primary wheat leaf, plants were grown continuously in the presence or absence of 10 mM nitrate and in either a 16/8 h light/dark cycle or continuous dark (Fig. 5). RNA was extracted from 2 cm sections along the 12 cm leaf from the base to the tip. To distinguish between pFNRI and pFNRII genes in northern blotting analysis, the areas of least identity in the pFNRI and pFNRII cDNAs, corresponding to the region coding for the transit peptide and the N-termini, were chosen for subsequent amplification of a probe. The 252 bp partial clone for wheat hFNR was used to study the gene expression of the hFNR isoform. The specificity of each probe was confirmed even under conditions of low stringency. The expression of each clone significantly varied between the base, middle, and tip of the wheat leaf.

Both pFNRI and pFNRII were expressed in a similar pattern along the wheat leaf. However, the highest levels of pFNRI and pFNRII mRNA were observed in the dark in the presence or absence of nitrate respectively (Fig. 5). hFNR mRNA expression was highest in the base and showed a marked decline along the wheat leaf under all growth conditions. The addition of nitrate in the light led to a significant up-regulation of hFNR mRNA levels. This effect of nitrate on gene expression was only significant in the light, although the effect of light alone on hFNR mRNA levels did not appear to be important. It was also shown that, in the dark, there was a highly significant dependence of hFNR mRNA production on the position along the leaf developmental gradient.

**Changes in the pFNRI and hFNR protein levels in variable light and nitrogen conditions**

Specific antibodies, PEPpFNR and pea hFNR, were used to distinguish between pFNRI and hFNR (Fig. 6). The amount of pFNRI protein detected by the PEPpFNR antibody gradually increased from the leaf base to tip in all sets of growth conditions (Fig. 6). By contrast, hFNR protein was detected in the basal part of the leaf under all growth conditions (Fig. 6). In leaves grown in the light and nitrate set of conditions, hFNR was detected along the entire wheat leaf (Fig. 6).

**Changes in pFNRI and hFNR enzyme activity in variable light and nitrogen conditions**

The contribution of each FNR isoform to measured enzyme activity in the wheat leaf was determined by using an immunoprecipitation approach (Fig. 7). Leaf extracts from different sections along the wheat leaf were incubated in the presence or absence of a constant amount of hFNR antibody and the total FNR activity measured in a cytochrome c reduction assay. The total FNR activity measured in each section in the absence of antibody was taken as 100% (comprising both pFNRI and hFNR). In the presence of hFNR antibody, hFNR activity was precipitated and therefore the activity measured in the assay represented that of pFNRI only. Figure 7 shows the proportion of pFNRI and hFNR activity as a percentage of the total FNR activity in the crude extract. In the light and nitrate treatment, pFNRI and hFNR activity represents 4%
and 96%, respectively, of the total FNR activity at the base (section 1) of the leaf. In the light without nitrate, pFNR and hFNR activity represented 27% and 73%, respectively, of the total FNR activity at the leaf base (section 1). The proportion of pFNR activity in the sections towards the leaf tip of plants grown in light without nitrate increased to reach 100% in sections 5 and 6. By contrast, there was no measurable enzyme activity detected in leaf tissue in the dark in either the presence or absence of nitrate.

Discussion

The aim of this study was to investigate the presence of multiple forms of FNR in wheat leaves and the capacity of FNR isoenzymes to respond to changes in reductant demand through varied expression and N-terminal processing. The 12 cm primary wheat leaf provides an ideal model system to investigate changes in response to development. To our knowledge, this is the first report of such a study on FNR. The purification of FNR from wheat leaf tissue led to the identification of four polypeptides. Based on N-terminal sequencing these proteins corresponded to two alternative N-termini versions of each pFNRI and pFNRII. Subsequent screening of a wheat leaf cDNA library confirmed the presence of pFNRI and pFNRII. Furthermore, the use of an RT-PCR approach led to the identification of a hFNRII clone in wheat leaves.

Changes in pFNR levels along the developmental gradient of a wheat leaf

12 cm long primary wheat leaves represent a linear developmental gradient of increasing cell age, with immature dividing heterotrophic cells at the base and mature photosynthetic cells at the tip. There is a gradient of cellular and plastid development along the wheat leaf.
confirmed in experiments with Arabidopsis where the steady-state transcript levels of AtLFNR1 were highest and AtLFNR2 transcripts were less abundant in plants grown in a light/dark cycle in the presence of high nitrate (Hanke et al., 2005). These data in wheat, together with that observed in Arabidopsis, support the suggestion that the two pFNR genes have distinct and separate functional roles. Recently, studies of Arabidopsis FNR knockout lines showed that AtLFNR1 and AtLFNR2 could not fully replace each others activity. The homozygous knockout line AtLFNR1 had reduced capacity for C fixation and a decreased biomass accumulation (Lintala et al., 2007).

Fig. 6. Changes in (A) pFNR and (B) hFNR protein levels along the length of a 12 cm primary wheat leaf grown for 7 d under variable light and nitrogen conditions as described in the Materials and methods. The 12 cm long primary wheat leaves were harvested, cut into six sections from the base (section 1) to the tip (section 6) and 50 μg of protein extract separated by SDS-gel electrophoresis, transferred onto a Protran membrane and treated with either (A) PEPpFNR or (B) pea hFNR primary antibody. Experiments were repeated at least three times and representative blots for each treatment are shown.

The physiological importance of N-terminal variants of pFNR

Since the original work of Keirns and Wang (1972) the presence of multiple forms of FNR have been confirmed by several laboratories (e.g. Karplus et al., 1984). The enzyme from Spinacia oleracea has been extensively studied and there have been discrepancies concerning the number of forms and their physical and chemical properties. In spinach, two closely migrating protein bands on SDS-polyacrylamide gels were suggested to arise from degradation of the larger form at the N-terminus (Shin et al., 1990). Although the alternative N-terminal cleavage sites identified in the work reported here for wheat may have arisen as a result of proteolytic degradation, this is unlikely since proteolytic protectants were included in all buffers during the initial stages of the purification. Furthermore where protein samples were prepared directly from fresh tissue of frozen extracts (stored at –20 °C or –80 °C) or directly from isolated chloroplasts, a similar pattern of multiple FNR proteins was identified using SDS-PAGE (data not shown). Recently, an NMR comparison of the effects on the N-terminal and C-terminal end of the FNR polypeptide chain of substrate interactions indicated that there are profound

(Dean and Leach, 1982a, b). By preparing sections along the leaf it is possible to investigate homogeneous populations of leaf cells and plastids of different developmental ages within the same leaves. Using this system allowed changes to be studied in FNR expression, protein, and enzyme activity with development, nitrate supply and light regime. The pattern of pFNR mRNA expression, protein and enzyme activity was similar in all experimental conditions, with lowest levels at the leaf base. Conversely, hFNR mRNA, protein, and enzyme activity tended to be highest at the leaf base and declined towards the tip of the leaf (Fig. 8).

Much of the available literature has identified the short-term response of pFNR at the transcriptional level via light-responsive cis elements and at the level of the enzyme (Neuhaus et al., 1993, 1997; Oelmüller et al., 1993; Bolle et al., 1994a, b; Bowler and Chua, 1996; Lübberstedt et al., 1994a, b). By contrast our study examined the impact of more long-term exposure to environmental factors, confirming that high levels of pFNR and pFNRII mRNA and protein were present in either set of light conditions. Interestingly, the highest mRNA levels for both pFNR and pFNRII were observed in the dark suggesting light does not play a significant long-term role in triggering and maintaining high levels of transcription of the pFNR encoding genes. In the dark-grown plants the highest expression was in the mid-leaf region and this may reflect an anatomical or developmental cue overridden and masked by light. In plants grown in the absence of light but in the presence of nitrate, there was a marked differential response with pFNRI expression increased and pFNRRII expression decreased. Although the level of pFNR protein increased towards the wheat leaf tip under all conditions, no other more subtle differences in the protein levels in response to nitrate or dark growth conditions were identified. This probably reflects the inability of the antibody used (PEPpFNR) to distinguish between the different pFNR forms. This is further
This is in agreement with work published on maize the predicted and actual isoelectric points, for example, its interaction with ferredoxin. Based on both important implications for the functionality of FNR with, focusing of the overexpressed latter difference was further confirmed by isoelectric measured using a cytochrome h FNR activity was measured using a cytochrome c six sections from the base (section 1) to the tip (section 6) and total methods. The 12 cm long primary wheat leaves were harvested, cut into nitrate and (B) light with nitrate as described in the Materials and effects on the N-terminus of the FNR (Maeda et al., 2005). In studies by Gadda et al. (1990) N-terminal truncated forms of FNR had full diaphorase capacity, but lost the ability to catalyse the ferredoxin-dependent reaction. As such, the alternative cleavage sites identified for both pFNRI and pFNRII in this study may have important implications for the functionality of FNR with, for example, its interaction with ferredoxin. Based on both the predicted and actual isoelectric points, pFNRI and pFNRII may be assigned as basic and acidic, respectively. This is in agreement with work published on maize (Okutani et al., 2005) and Arabidopsis (Hanke et al., 2005; Lintala et al., 2007) isoforms. Interestingly, although the alternative N-terminal forms of pFNRII had very similar isoelectric points, the alternative cleavage site for pFNRI, pFNRIKVS, and pFNRIKKK, resulted in a marked change in the predicted isoelectric point. This latter difference was further confirmed by isoelectric focusing of the overexpressed pFNRI, although the impact of the N-terminal truncation on the overall pI was not as marked as the predicted value. Overexpression of these alternative forms of pFNRI allowed us to investigate whether this N-terminal truncation has any potential effect on the kinetic properties of either pFNRI or pFNRII. Of all the overexpressed pFNRI forms assayed, only the longer form of pFNRII, pFNRIIKKK, showed a clear discrimination between the different forms of Fd with a notably higher Km for the photosynthetic FdII when compared to the heterotrophic FdIII. The two amino acid N-terminal truncation to pFNRIKDD eliminated this discrimination, bringing the Km for FdI to a level equivalent to that for non-photosynthetic FdIII. In previous work, maize FNR was also unable to discriminate between these photosynthetic and heterotrophic forms of Fd (Onda et al., 2000).

N-terminal truncation might be a form of post-transcriptional regulation operating alone or in combination with other regulatory mechanisms. For example, a suggested role for protein phosphorylation has been proposed. However, although possible sites have been identified in silico, these have still not been confirmed in vivo (Hodges and Miginiac-Maslow, 1993; Lintala et al., 2007). Multiple putative subforms of FNR exhibiting different enzymatic activities may reflect a specific metabolic role, for example, cyclic or non-cyclic electron flow. Truncated N-terminal forms of pFNRI with differential abilities to discriminate between different ferredoxins may enable the more effective flow of electrons through particular pathways.

The differences in the N-termini of the FNR may also result in different modes of association with thylakoids. This assumption was originally proposed by Wada et al. (1996) after detecting free-state hFNR and partly thylakoid-bound pFNRI in the same tissues. It was further supported when more than one leaf FNR were characterized for Arabidopsis (Hanke et al., 2005) and maize (Okutani et al., 2005). Multiple pFNRI subforms could represent free and thylakoid-bound populations within the chloroplast also differing in function. In previous work by Hanke et al. (2005), studying pFNRI, the more acidic ArFNRI was the most abundant form in the thylakoids and the less acidic ArFNRII was mostly present in the stroma. This might suggest the less acidic wheat pFNRIKVS is in the stroma and the more acidic truncated pFNRIKKK is in the thylakoids. Although the N-terminal truncation of pFNRII has no apparent impact on the isoelectric point, the observed effect of this truncation on the Km and Vmax of pFNRII and its ability to discriminate between different Fd isoproteins needs to be investigated further to understand the possible functional implications and whether this is of importance in vivo. This supports the idea that the plant system is similar to cyanobacteria where distribution of FNR between thylakoids and the phycobilosome fraction provides a mechanism for sharing electrons between linear and cyclic electron flow (Van Thor et al., 1998, 2000). Interestingly, recent work using Arabidopsis
Mutants has suggested that the absence of the acidic $AtLFNRI$ prevented association of the basic $AtLFNRII$ with the thylakoid membrane (Lintala et al., 2007). Dimer formation may, depending on environmental cues, be a way to regulate the distribution of electrons between the cyclic and linear electron transfer pathway (Lintala et al., 2007).

Although it is not known whether the specific isoforms of FNR identified in this study are catalytically more likely to support specific metabolic processes, we can propose a model where, for example, depending on the photosynthetic status of the cell, multiple $pFNR$ forms might be distributed between the cyclic and non-cyclic electron flow (Shahak et al., 1981). In cyclic electron flow, electrons from PSI are returned to the quinone pool via ferredoxin (and an unknown Fd:quinone reductase) or NADPH (and the Ndh complex), generating ATP with no net reduction of NADPH (Takabayashi et al., 2005). A close association of FNR with the chloroplastic Ndh complex was described in potato and barley (Guedeney et al., 1996; Quiles and Cuello, 1998; Funk et al., 1999). Teicher and Scheller (1998) demonstrated that Ndh in barley thylakoids was activated by light and accounted for 40% of ferredoxin-dependent cyclic electron-transport rates. Depending on the location of bound $pFNR$, it could create a NADPH-rich mini-environment, stimulating cyclic electron transfer through the Ndh complex. The over-reduced ferredoxin, resulting from an increased NADPH/NADP ratio, can enhance the diversion of the electrons into the cytochrome $b/f$ complex, thus triggering the cyclic flow (Kimata-Ariga et al., 2000). This has important implications for plant metabolism providing firstly, an additional supply of ATP and, secondly, a defensive mechanism against photo-oxidative damage and salt stress (Canaani, 1990; Heber and Walker, 1992). The presence of multiple FNR forms which are regulated independently and can interact differently along the wheat leaf provides added flexibility to allow the plant to adapt to a changing metabolic environment.

$hFNR$ in the developing wheat leaf

$hFNR$ mRNA was undetectable in leaf tissue grown in the light in the absence of nitrate, although there was protein and enzyme activity detectable in the basal section. The highest levels of $hFNR$ mRNA, protein, and enzyme activity were demonstrated in the leaf base section and decreased towards the leaf tip in the presence of nitrate. Since the basal cells of the cereal leaf are non-photosynthetic (Thompson et al., 1998), one can envisage a role for $hFNR$ in these cells in providing reductant non-photosynthetically via the oxidative pentose phosphate pathway for a range of metabolic processes (Bowsher et al., 2007). In agreement with previously published results on the short-term nitrate induction of $hFNR$-encoding genes in roots (Ritchie et al., 1994; Mattana et al., 1997), the continuous presence of nitrate has a significant effect on $hFNR$ expression, protein, and enzyme activity in leaf tissue.
transient induction of hFNR in Arabidopsis (Wang et al., 2003) and maize (Sakakibara, 2003) shoots transferred to nitrate has been reported. However, more recently, studies on Arabidopsis plants growing on high nitrate reported that only low steady-state levels of hFNR levels were detected in leaf tissue (Hanke et al., 2005). Interestingly when Arabidopsis plants were grown continuously in the presence of ammonium, the levels of hFNR increased, again confirming the potential importance of this isoform in green tissue during nitrogen metabolism (Hanke et al., 2005). Such a discrepancy in hFNR induction between wheat and Arabidopsis leaves may reflect a difference between photosynthetic and non-photosynthetic cells distribution in monocotyledonous and dicotyledonous plants.

A role for FNR in linking carbon and nitrogen metabolism

The use of the hFNR antibody in immunoprecipitation experiments meant the contribution of pFNR and hFNR to the total FNR enzyme activity measured could be distinguished. The apparent switch from one protein form to another occurs between the base and middle or between the middle and apical parts of the leaf. The presence of multiple pFNR subforms and hFNR in wheat leaf tissue appears to be dependent on the presence or absence of light and/or nitrate. Different isoforms of FNR may have similar functions within the cell, but operate in either distinct cell types or developmental periods. However, it is also possible that different isoforms may have distinct non-overlapping functions within the same cell. The heterogeneity of cell types along a wheat leaf and the apparent influence of the development gradient along the wheat leaf on the FNR forms present, most likely reflects the involvement of the different isoforms and indeed subforms specializing in specific types of electron transfer in response to changing metabolic demands along the leaf.

In leaves in the light, ferredoxin transfers photon-generated reducing power to nitrite reductase, to glutamate synthase, and to NADP+ via FNR. Therefore, enzymatic activities fundamental to carbohydrate synthesis from CO2 and the reduction of nitrite to ammonia and its incorporation into glutamate, all depend on reduced ferredoxin. Interestingly, the supply of reduced ferredoxin via hFNR seems to be a limiting factor when optimal nitrite reductase and glutamate synthase activity are operating in non-photosynthetic root plastids (Bowsher et al., 2007). The contribution of pFNR and hFNR in meeting the needs for reductant supply to support nitrogen assimilation in the leaves under non-photosynthetic conditions is not known. However, leaves contain a large proportion of cells with functions other than photosynthesis and of the total cell population in 12 cm illuminated wheat leaves, only 45% in the leaf base and 55% in the leaf tip are chloroplast containing mesophyll cells (Dean and Leach, 1982a, b). Furthermore while chloroplasts are located in mesophyll cells, plastids are also found in, for example, vascular tissue, which can contribute as much as 5% of the total leaf cells, and in epidermal cells, that contribute another 10–20% (Bowsher and Tobin, 2001). Such a distribution of pFNR and hFNR between mesophyll and heterotrophic cells, respectively, and the changing metabolic needs of different cells may account for the variation seen in gene expression, protein amount, and enzyme activity values observed in this current study. hFNR activity in a heterotrophic direction and therefore its presence along the wheat leaf most likely reflects an auxiliary role under conditions of high ferredoxin demand in the leaf, such as high nitrate growth conditions.

Using the model of the 12 cm primary wheat leaf has provided us with a system to examine how multiple FNR isoforms change in response to development and different light and nitrate environments along the leaf. The switch from hFNR at the base of the leaf to multiple pFNR forms reflects a change in heterotrophic metabolism in the leaf meristem, to increasing photosynthetic capacity as leaf cells move towards the leaf tip. Although the two forms of pFNR show a similar pattern of expression along the leaf, clearly different forms respond in a different manner to nitrate and the dark. Furthermore the forms may also associate differently based on the metabolic demands. This study has identified the presence of multiple forms of FNR in wheat leaves with varied expression and N-terminal processing. This provides the leaf with the potential flexibility to respond to changes in reductant demand in a number of ways. For example, depending on the photosynthetic status of the cell, the presence of multiple pFNR forms may reflect their distribution between the cyclic and non-cyclic electron flow. This situation may be further regulated by the role of isoform-dependent N-terminal truncations in functionality. Clearly N-terminal truncation leads to some discrimination betweenFd isoforms in vitro. We are currently investigating the physiological significance of this in vivo.

Acknowledgements

This work was supported in part by funding from The Royal Society, and the BBSRC (Grant JPA 886). GJF and AM were in receipt of BBSRC studentships. JGJ was funded by grants from the Open Society Institute (New York, USA) and the Central European University (Budapest, Hungary). Wheat was grown by Mr Thurston Heaton at the Firs Experimental Grounds, The University of Manchester. pFNR antibody was kindly supplied by Professor J Gray (Cambridge).

References


Bowsher CG, Dunbar B, Emes MJ. 1993a. The purification and properties of ferredoxin-NADP\(^+\) oxidoreductase from roots of *Pisum sativum* L. Protein Purification and Expression 4, 512–518.


Bowsher CG, Knight JS. 1996. The isolation of a pea root ferredoxin NADP\(^+\) oxidoreductase (FNR) cDNA. Plant Physiology 112, 861.


Canaani O. 1990. The role of cyclic electron flow around photosystem I and excitation energy distribution between the photosystems upon accumulation to high ionic strength in Dunaliella. Phytochemistry and Photobiology 52, 591–599.


Lintala L, Allahverdiyeval Y, Kidron H, Pippio M, Battchikoval N, Suorsa M, Rintamäki E, Salminen TA,


