The assembly of iron–sulfur (Fe–S) clusters requires dedicated protein factors inside the living cell. Striking similarities between prokaryotic and eukaryotic assembly proteins suggest that plant cells inherited two different pathways through endosymbiosis: the ISC pathway in mitochondria and the SUF pathway in plastids. Fe–S proteins are also found in the cytosol and nucleus, but little is known about how they are assembled in plant cells. Here, we show that neither plastid assembly proteins nor the cytosolic cysteine desulfurase ABA3 are required for the activity of cytosolic aconitase, which depends on a [4Fe–4S] cluster. In contrast, cytosolic aconitase activity depended on the mitochondrial cysteine desulfurase NFS1 and the mitochondrial transporter ATM3. In addition, we were able to complement a yeast mutant in the cytosolic Fe–S cluster assembly pathway, dre2, with the Arabidopsis homologue AtDRE2, but only when expressed together with the diflavin reductase AtTAH18. Spectroscopic characterization showed that purified AtDRE2 could bind up to two Fe–S clusters. Purified AtTAH18 bound one flavin per molecule and was able to accept electrons from NAD(P)H. These results suggest that the proteins involved in cytosolic Fe–S cluster assembly are highly conserved, and that dependence on the mitochondria arose before the second endosymbiosis event leading to plastids.

1. Introduction

Iron–sulfur (Fe–S) clusters are cofactors found in a large number of essential proteins [1,2]. Their versatile chemical properties enable enzymes to carry out electron transfer, Lewis acid catalysis, radical chemistry or to provide stability to protein folds. Fe–S-rich sediment may have functioned as a catalytic surface to form the first building blocks of life [3], but when primordial life became dependent on Fe–S chemistry, the catalyst needed to be assembled inside the cell. Whereas simple rhombic and cubane Fe–S clusters can form spontaneously under anaerobic conditions in a test tube, a significant number of dedicated assembly proteins have been uncovered over the past 15 years (reviewed in [4–6]). These are the nif pathway in nitrogen fixing bacteria, which bears similarity to the isc (iron–sulfur cluster) pathway in bacteria and mitochondria. The suf (sulfur mobilization) pathway operates in (cyano)bacteria under oxidative stress and in chloroplasts, while suf BC genes are also commonly found in Archaea. In eukaryotes, the CIA pathway (cytosolic iron–sulfur protein assembly) is involved in the maturation of cytosolic and nuclear Fe–S proteins.

Based on extensive biochemical and cell biological studies, the assembly proteins are needed for controlled delivery of sulfur in the form of persulfide (S0), reduction to sulfide (S2−), pre-assembly together with Fe on a scaffold protein and ATP-dependent transfer of the Fe–S cluster to a target protein. In most bacteria and in organelles, the source of sulfur for Fe–S clusters is cysteine, which is converted to alanine and S0 by a pyridoxal phosphate-dependent
cysteine desulfurase. Phylogenetically, the cysteine desulfurases fall into two groups, type I and type II [7]. The mitochondrial cysteine desulfurase NFS1 is type I, whereas the chloroplast CpNifS (also known as SUP5 or NFS2) is a type-II enzyme. A third cysteine desulfurase found in the cytosol, named ABA3 for its role in the synthesis of the plant hormone abscisic acid, also belongs to type II. The two types differ in the level of surface exposure of the active site, as well as in the requirement of helper proteins.

While it is clear that the mitochondrial NFS1 and the plastid CpNifS provide sulfur for the assembly of clusters in the mitochondria and plastids, respectively, the source of sulfur for cytosolic/nuclear Fe–S proteins is more enigmatic. In yeast, a small pool of the single NFS1 gene product is localized in the nucleus [8], but does not play a role in Fe–S protein assembly [9]. In human cell lines, a mitochondrial and a cytosolic/nuclear isoform of the NFS1 homologue (called ISCS) have been observed [10,11], but only the mitochondrial form contributes functionally to the assembly of cellular Fe–S proteins [11]. In Arabidopsis, two independent studies found that NFS1::GFP fusion proteins are exclusively targeted to the mitochondria [12,13], although the existence of a minor non-mitochondrial pool can as yet not be ruled out. If the mitochondrial pool of NFS1 provides a sulfur-containing precursor for the CIA pathway, then this needs to be exported to the cytosol via a transporter. A probable candidate is the ABC transporter of the mitochondria, Atm1 in yeast [14,15]. The orthologues in mammals (ABC87) and plants (ATM3) are also required for the activity of cytosolic Fe–S enzymes [16–18]. However, disruption of Arabidopsis ATM3 is not lethal, although growth is severely compromised [18]. Therefore, in plants the possibility exists that either the plastid CpNifS or the cytosolic ABA3 can provide persulfide to cytosolic Fe–S proteins.

The reduction of $S^0$ to $S^{2−}$, and possibly the reductive coupling of two rhombic [2Fe–2S] clusters to [4Fe–4S], require electrons. NAD(P)H, ferredoxin reductase and ferredoxin form an electron transfer chain that provides electrons to the mitochondrial ISC pathway in all eukaryotes, including plants [19–21]. In plastids, electrons for the assembly of Fe–S clusters are derived from light or NADP(H) [22], probably via ferredoxins, but the latter remains to be shown. In the yeast cytosol, the diflavin reductase Tah18 and its partner Dre2 form an electron input module for the CIA pathway [23]. The Tah18 and Dre2 proteins were recently described in Arabidopsis, as ATR3, a homologue of ATR1 and ATR2 (Arabidopsis thaliana P450 reductases), and AtCIAPIN1 (Arabidopsis thaliana homologue of the human cytokine-induced apoptosis inhibitor 1) [24]. In this paper, we use the yeast gene names for the Arabidopsis homologues, with the prefix At for Arabidopsis thaliana. Varadarajan et al. [24] found that AtTAH18 interacted specifically with AtDRE2 identified in a yeast two-hybrid screen. Moreover, cell fractions of yeast or Nicotiana benthamiana overexpressing AtTAH18::GFP were shown to have NADPH-dependent cytochrome c reductase activity. However, the biological role of AtTAH18 and AtDRE2, particularly regarding Fe–S protein biogenesis, was not established, partly owing to lack of viable loss-of-function mutants and incomplete characterization of the yeast proteins at the time.

To investigate the assembly of Fe–S proteins in the plant cytosol, we have addressed the issues of sulfur source and electron input. Arabidopsis mutants in the three cysteine desulfurase genes, encoding the plastid CpNifS, cytosolic ABA3 and mitochondrial NFS1, were analysed for cytosolic Fe–S protein activities. Our results show that CpNifS is required for plastid Fe–S enzymes and that AtBA3 function is restricted to Molybdenum cofactor (Moco) maturation. For activity of the [4Fe–4S]-dependent enzyme aconitase in the cytosol, NFS1 was required. We also show that AtTAH18 and AtDRE2 could functionally complement the yeast dre2 mutant only when expressed together, indicating that AtDRE2–AtTAH18 complex formation is of functional importance.

## 2. Methods

### (a) Plant materials and growth

T-DNA insertion mutants in DRE2 (SAIL_1222_B12) and NFS1 (SALK_083681 and WiscDsLox445D06) were obtained from the Nottingham Arabidopsis Stock Centre. nfs1-1/nfs1-2 hemizygous knock-down individuals were obtained by crossing the homozygous SALK_083681 line (nfs1-1/nfs1-1) with heterozygous individuals (nfs1-2/NFS1) of the WiscDsLox445D06 line, followed by PCR genotyping of the F1. The other mutant lines were as previously described: CpNifS [25]; nfs2-1 (SALK_039254) [26,27]; the str3-3 allele of ABA3 [28]; atm3-1 and atm3-4 [18]. Plants were grown on compost in a controlled growth cabinet at 21 °C, 16 h light:8 h dark, humidity 65 per cent. For 3-day-old seedlings, seeds were germinated in ½ Murashige and Skoog medium.

### (b) Yeast strains, growth and plasmids

Saccharomyces cerevisiae W303-1A (MATa, ura3-1, ade2-1, trp1-1, his3-11,15 and leu2-3,112) was used as wild-type (WT) strain. The Gal-DRE2 and Gal-TAH18 strains (carrying the Gal1-10 promoter) in this study were constructed using the plasmid pYM-N23 [29] as the template for PCR amplification of linear DNA for homologous recombination of the WT strain using nourseothricin selection. A strain having both DRE2 and TAH18 genes under the control of the Gal1-10 promoter was constructed from Gal-DRE2 in a similar fashion, but using the plasmid pYM-N23 in which the NAF gene was replaced by the Schizosaccharomyces pombe HIS5 gene. Correct promoter insertion was confirmed by PCR of chromosomal DNA with primers hybridizing to flanking regions and the inserted cassette. The AITA18 coding region was previously isolated [24] and the AITA18 coding region was PCR amplified from Arabidopsis thaliana cDNA with gene-specific primers. AITA18 and AITA2 were cloned into p414 and p416 yeast vectors, respectively [30]. For constructs with natural promoters (NPs) of yeast, 450 bp or 518 bp 5′ of the start codons of DRE2 or TAH18, respectively, were used. All cloned regions in the plasmids were confirmed by sequencing. Yeast methods were according to [23].

### (c) Enzyme assays and spectroscopic analysis

In-gel assays for aldehyde oxidases, xanthine dehydrogenase and aconitases were as previously described [18]; isopropylmalate isomerase (Leu1) activity in yeast cell extracts was determined according to [14]. Spectroscopy and kinetic analysis of horse heart cytochrome c reduction by AITA18 was performed as described in [23].

### (d) Protein expression and purification

AtDRE2 or AITA18 were cloned into the SacI and NcoI sites of the pASK-IBA43plus vector, generating a N-terminal 6xHis- and C-terminal Streptag in fusion with the encoded protein. Expression of AtDRE2 or AITA18 in E. coli HMS174 was
started by inoculation of Terrific Broth medium with 1 per cent (v/v) of an overnight culture. For AtDRE2, the culture at 37 °C was shifted to 25 °C at an OD_{600nm} of 0.5, followed by the addition of 0.2 mg l^{-1} anhydrotetracyclin. Cells were harvested after 3 h. In the case of AtTAH18, growth was at 20 °C, induction was at OD_{600nm} = 0.3 and the harvest was after 16 h. Extracts were prepared from defrosted cells with an Avestin liquidizer and clarified by centrifugation for 2 h at 100 000 × g (4 °C). Purification according to the manufacturers’ protocols was with IBA Streptactin Agarose superflow (AtDRE2) or with Ni-NTA HP Sepharose with 20 mM imidazol in the lysis/wash buffer (AtTAH18). AtTAH18 was further purified by Superdex 200pg gel filtration in 25 mM Tris–HCl pH 8.0, 300 mM NaCl buffer using an Äkta system.

3. Results
(a) Plant mitochondria, but not plastids, are required for cytosolic Fe–S enzyme activities

Because plant cells have two functional pathways for Fe–S cluster assembly, ISC in mitochondria and SUF in plastids, each could potentially contribute to the assembly of Fe–S clusters in the cytosol. We have previously shown that the mitochondrial cysteine desulfurase NSF1 is required for activity of cytosolic aldehyde oxidases (AO), which bind two [2Fe–2S] clusters, FAD and Moco [13]. In addition, we have demonstrated that the mitochondrial ABC transporter ATM3 is required for the activities of at least three cytosolic Fe–S enzymes, AO, the structurally related xanthine dehydrogenase (XDH) and the [4Fe–4S] enzyme aconitase (ACO) [18]. To investigate whether the plastid Fe–S assembly pathway also contributes to cytosolic Fe–S cluster assembly, we analysed the activities of cytosolic Fe–S enzymes in two mutants of plastid assembly proteins. Knockout of the cysteine desulfurase CpNiFS is lethal, whereas silencing of its expression leads to decreased activity and/or protein stability of all resident plastid Fe–S proteins tested [25]. In contrast, plants lacking the Fe–S scaffold protein NFU2 are viable, but have severely decreased levels of Photosystem I and ferredoxin, while plastid Fe–S proteins such as sulfite reductase (Sir) and nitrite reductase were less affected, and glutamate synthase and Rieske protein not at all [26,27]. Plants expressing an ethanol-inducible RNAi fragment complementary to CpNiFS were grown for two weeks and treated with 2 per cent (v/v) ethanol to downregulate CpNiFS expression, as previously described [25]. After two weeks, the plants were daisy and chlorotic compared with normal growth of WT plants that had undergone the same treatment (figure 1a). In CpNiFS silenced leaves, the levels of Sir and the Fe–S subunit of cytochrome b$_{5}$ complex (CpRieske) were decreased, but activities of cytosolic AO, XDH and ACO were normal (figure 1b,c). nfs2-1 mutant plants also had normal activities of cytosolic Fe–S enzymes. For comparison, AO, XDH and cytosolic ACO activities were very low or undetectable in plants homozygous for either a weak or strong allele of atm3. Despite the chlorotic appearance and severe growth phenotype of atm3-1 mutant plants, the Sir protein was stable. In addition to the mitochondrial and plastid cysteine desulfurases, plants also contain a cysteine desulfurase in the cytosol, ABA3. The ABA3 protein consists of two domains: an N-terminal cysteine desulfurase domain which transfers a sulfur to Moco bound to the C-terminal domain, prior to transfer and insertion of sulfurated Moco into AO and XDH [31]. Using the same in-gel activity assays, we found that the sir3-3 allele of ABA3 has strongly decreased activities of AO and XDH, but that the cytosolic ACO activity is unaffected (figure 1b,c). These results are in agreement with previous in vitro data suggesting that activity of ABA3 is specific for Moco sulfuration [31,32]. Moreover, the phenotypic features of abac mutants, which include a darker green leaf colour and wilty leaves, are very different from atm3 mutants [18,33], also suggesting that ABA3 is not involved in cytosolic Fe–S cluster assembly.

To confirm that the mitochondrial NSF1 is not only required for Fe–S/Moco enzymes like AO [13], but also for cytosolic Fe–S enzymes, we analysed ACO activity in an nfs1 mutant. A previously reported line in which NSF1 was downregulated by transgenic expression of the coding sequence in antisense direction was genetically unstable, whereas T-DNA insertions in the NSF1 coding sequence cause embryo lethality [13]. Therefore, to obtain a viable mutant with consistently decreased NSF1 expression levels, we isolated a mutant with a T-DNA insertion in the 5’ untranslated region of NSF1 (nfs1-1, SALK_083681). Although transcript levels of NSF1 were decreased by approximately 50 per cent in homozygous plants, they had no obvious visual or biochemical phenotypes (data not shown). However, in combination with a knock-out allele (nfs1-2), the hemizygous knock-down seedlings showed delayed germination. During germination of WT Arabidopsis, high activity of cytosolic aconitase (see the electronic supplementary material, figure S1) supports a modified glyoxylate cycle involved in mobilization of storage lipids (Hooks & Balk, manuscript in preparation). In 3-day-old nfs1-1/nfs1-2 seedlings, however, aconitase activity is strongly decreased compared with WT, while the level of aconitase protein is unaffected (figure 1d). These results indicate a deficiency in Fe–S cofactor assembly on cytosolic aconitase and suggest that the mitochondrial cysteine desulfurase is the source of sulfur for cytosolic Fe–S proteins in plants. Moreover, our data show that neither the plastid Fe–S cluster assembly machinery, nor the cytosolic cysteine desulfurase ABA3 play a role in the maturation of Fe–S proteins in the plant cytosol.

(b) The Arabidopsis homologue of DRE2 is essential for embryo development

To investigate proteins downstream of NFS1 and ATM3 that could be involved in the assembly of Fe–S clusters in the plant cytosol, we have previously annotated Arabidopsis and Chlamydomonas homologues of the yeast CIA genes [3,34]. All CIA genes were easily identified, except for CFD1, which is absent in the green lineage [35]. AtDRE2, AtTAH18, AtNBP35, AtNAR1, AtCIA1 and AtMET18 occur as single genes, whereas three genes with homology to CIA2 are found in Arabidopsis (table 1). T-DNA insertion mutants are available for most genes. It has been shown that AtTAH18, AtNBP35, AtNAR1, AtCIA1 and one of the AtCIA2 genes (named AE7) are each essential for embryo development (see table 1 and references). In contrast, Arabidopsis met18 knock-out mutants have no obvious phenotype [37].
Since the phenotype of atdre2 mutants has not been described, we screened the stock centres for T-DNA insertion mutants. Several lines were obtained, but only one line in the SAIL collection [40] displayed a phenotype. The T-DNA is inserted in the fifth exon, which is likely to cause complete disruption of expression. No homozygous mutant plants were found among the offspring of heterozygous parents. Further inspection of the seed pods revealed inviable seeds among healthy-looking seeds (see the electronic supplementary material, figure S2), at a ratio of 75 : 190 \((n = 6\) siliques), or approximately 1 : 3 \((\chi^2 = 1.634; \chi^2_{0.05} = 3.841)\). Based on the small size of aborted seeds, embryo development is arrested at an early stage, similar to mutants in AtTAH18, AtNBP35, AE7, AtCIA1 and AtNAR1 [37]. Evidently, the CIA genes perform a fundamental function in plants.

(c) AtDRE2 and AtTAH18 fully complement a yeast dre2 mutant when expressed together

To further investigate the function of AtDRE2, the Arabidopsis gene was cloned and expressed from a plasmid in the yeast Gal1-10-DRE2 regulatable mutant (abbreviated Gal-DRE2). Upon downregulation of DRE2 by changing the carbon source to glucose, the Gal-DRE2 cells stopped growing (figure 2a). While constitutive expression of ScDRE2 was able to rescue growth, expression of AtDRE2 from either the natural yeast DRE2 promoter (NP) or the strong TDH promoter did not rescue.

We also investigated whether AtTAH18 could complement a yeast Gal1-10-TAH18 mutant (abbreviated Gal-TAH18). Upon downregulation of the yeast TAH18 gene, constitutive
Table 1. Arabidopsis genes of the CIA pathway and their mutant phenotypes.

<table>
<thead>
<tr>
<th>locus</th>
<th>gene name</th>
<th>yeast respectively human homologues</th>
<th>mutant allele(s)</th>
<th>phenotype</th>
<th>references</th>
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<td>At5g18400</td>
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<td>DRE2, CIAPIN1/ anamorin</td>
<td>T-DNA insertion</td>
<td>embryo lethal</td>
<td>this study</td>
</tr>
<tr>
<td>At3g02280</td>
<td>TAH18 , ATR3</td>
<td>TAH18, NDOR1</td>
<td>T-DNA insertion</td>
<td>embryo lethal</td>
<td>[24]</td>
</tr>
<tr>
<td>At5g0960</td>
<td>NBP35</td>
<td>NBP35, NUBP2</td>
<td>T-DNA insertion</td>
<td>embryo lethal</td>
<td>[35, 36]</td>
</tr>
<tr>
<td>At4g16440</td>
<td>NAR1</td>
<td>NAR1, IOP1/NARF</td>
<td>T-DNA insertion</td>
<td>ovule abortion or early</td>
<td>[37, 38]</td>
</tr>
<tr>
<td>At2g26060</td>
<td>CIA1</td>
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<td>T-DNA insertion</td>
<td>embryo lethal</td>
<td>[37]</td>
</tr>
<tr>
<td>At1g68310</td>
<td>AE7</td>
<td>CIA2, MIP18</td>
<td>T-DNA insertion</td>
<td>embryo lethal</td>
<td>[37]</td>
</tr>
<tr>
<td></td>
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<td></td>
</tr>
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<td>At3g50845</td>
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<td>MET18, MMS19</td>
<td>T-DNA insertion</td>
<td>no phenotype</td>
<td>[37]</td>
</tr>
</tbody>
</table>

expression of the Arabidopsis homologue could only partially rescue growth when the expression level was increased using the MET25 promoter (figure 2a). Because Dre2 and Tah18 proteins in yeast, human and Arabidopsis are known to form a protein–protein interaction [23, 24, 41], we tested whether co-expression of AtTAH18 and AtDRE2 could complement dre2 yeast. When expressed from either the NP or MET25 promoter for AtTAH18, the combined expression of the plant genes almost fully complemented growth of a Gal-DRE2 single- and Gal-TAH18/Gal-DRE2 double mutant.

To confirm that yeast growth in the complemented strains was owing to restoration of Fe–S protein biogenesis, the activity of the cytosolic Fe–S enzyme isopropylmalate isomerase (Leu1 protein) was measured. Co-expression of Arabidopsis TAH18 and DRE2 did restore approximately 30 per cent of the Leu1 activity measured in control cells (figure 2b). These data indicate that the AtTAH18 and AtDRE2 proteins form a functional unit that can serve the same function as the respective yeast homologues.

(d) AtTAH18 is a flavoprotein and AtDRE2 binds Fe–S clusters

To characterize the cofactors and activities of AtTAH18 and AtDRE2, we expressed these proteins in E. coli. Upon expression at 30 °C the majority of AtTAH18 was found in inclusion bodies. Induction at 20 °C resulted in soluble full-length AtTAH18, which could be purified by Ni-NTA affinity chromatography and gel filtration (0.1 mg g⁻¹ E. coli cells). The yellow monomeric protein contained approximately 1 flavin per molecule, as calculated from the characteristic absorbance maxima at 365 and 450 nm (figure 3a). Electron transfer from NADPH and NADH (0.074 and 0.046 s⁻¹, respectively) to the commonly used electron acceptor cytochrome c (see figure 3b and electronic supplementary material, figure S3) occurred at a rate typical for other diflavin reductases from the same family [23, 42]. Kₘ values for cytochrome c, NADPH and NADH were 10, 50 and 28 μM, respectively. The specificity constants (Vₘₐₓ/Kₘ) are thus almost identical, indicating that in vivo both reduced coenzymes can act as an electron donor.

AtDRE2 could be obtained in moderate yield following purification on a Strept-tactin column (0.3 mg g⁻¹ E. coli cells). The red-brownish protein had a UV-visible absorbance spectrum with maxima at 315 and 420 nm, suggesting it binds a flavin within the [2Fe–2S] centre (figure 3c). Reduction of the protein sample revealed an electron paramagnetic resonance (EPR) spectrum characteristic of a rhombic [2Fe–2S] cluster, which was detectable up to 84 K (figure 3d). The g values as determined by simulation of the EPR spectrum (1.998, 1.959 and 1.916) are similar to those reported for the purified yeast Dre2 protein [43, 44]. Since chemical analysis detected approximately 1.3 Fe and 1.8 per polypeptide, we anaerobically reconstituted AtDRE2. As seen in figure 3c, the amount of visible Fe–S chromophores bound to AtDRE2 increased approximately sevenfold. The intensity of the contribution of the [2Fe–2S]³⁺ cluster increased and was accompanied by a general increase of less-structured absorbance in the 400–700 nm region. These results indicate that a [4Fe–4S] cluster is present in the holo-form in addition to the [2Fe–2S] cluster.

4. Discussion

The cells of plants and green algae have both mitochondria and plastids, each with an evolutionarily distinct pathway for the assembly of Fe–S clusters, termed ISC and SUF, respectively [6]. Previous reports showed that the pathways act independently: downregulation of the plastid cysteine desulphurase CpsNifS did not impact on mitochondrial respiratory function [25]. Mutants in the mitochondrial ABC transporter ATM3 affected cytosolic Fe–S enzymes, but not plastid Fe–S proteins such as Photosystem I, nitrite reductase...
and chlorophyllide a oxygenase [18]. Here, we extend these observations to include evidence that the cytosolic Fe–S enzymes depend on the mitochondrial assembly pathway, but not the plastid assembly machinery. The findings suggest that the dependence of the cytosolic Fe–S clusters on the mitochondria was established before the second endosymbiosis leading to plastids, and that this situation was maintained after the introduction of the SUF Fe–S cluster assembly pathway into the cell.

Interestingly, from yeast to human cells to plants, the assembly of cytosolic/nuclear Fe–S proteins depends on a cysteine desulfurase that is localized in the mitochondria. Are there constraints to re-localize certain steps of the cluster assembly process, or is there a specific, biochemical reason for NFS1 to be in the mitochondria? Examples of assembly proteins in unexpected locations have been reported. The Arabidopsis SUFE protein fused to GFP localized to both plastids and mitochondria [45]. The unicellular eukaryote Blastocystis, which has rudimentary mitochondria and no plastids, appears to have acquired a functional SufC–SufB fusion protein by lateral gene transfer. The SufCB protein is localized in the cytosol, alongside putative CIA proteins [46]. Also, as mentioned earlier, the mitochondrial cysteine desulfurase is found in the cytosol/nucleus of yeast and human cells, although this pool does not serve extra-mitochondrial Fe–S protein biogenesis. So given that targeting of a protein is subject to evolutionary change, there must be another reason why the source of sulfide for cytosolic

Figure 2. Arabidopsis TAH18 and DRE2 complement yeast dre2 when expressed together. (a) The indicated yeast strains were transformed with the listed plasmids. After growth for 40 h on liquid glucose-containing minimal medium, successive 10-fold dilutions of a cell suspension with OD600 nm = 0.5 were spotted on agar plates. Photographs were taken after 40 h of growth at 30 °C. (b) Isopropylmalate isomerase (Leu1) specific activities of cell extracts of yeast strains as in (a). Extracts were prepared from cells grown for 40 h on liquid glucose-containing minimal medium. —, no plasmid; Ø, empty plasmid.
Fe–S clusters is restricted to the mitochondrial matrix. It has been suggested that the ISC pathway is oxygen sensitive, and can only function in the matrix because of lower oxygen partial pressure in this compartment.

The biochemical characterization of Arabidopsis TAH18 and DRE2 reported here, together with other recent studies of plant homologues of the yeast CIA proteins (table 1), allow a first draft of the cytosolic Fe–S cluster assembly pathway in plants (figure 4) as well as comparison with other eukaryotes. Successful complementation of the yeast dre2 mutant with the Arabidopsis TAH18-DRE2 module and characterization of their redox-active cofactors suggest that these proteins also act as an electron transfer chain in the plant CIA machinery. Based on 55Fe incorporation studies in yeast, TAH18-DRE2 act early in the pathway before NBP35, whereas the CIA targeting complex acts late. In plants, NBP35 forms a dimer carrying a stable N-terminal cluster, and a labile C-terminal cluster that spans the two protomers [35,36]. Because of its labile nature, the C-terminal cluster is probably the cluster that is transferred to target apo-proteins. The presence of a nucleotide binding domain indicates that NBP35 function needs ATP or GTP, most probably for loading of the Fe–S cluster on the scaffold, as shown in yeast [47]. In fungi and metazoa NBP35 forms a heterotetramer with Cfd1/NUBP1. Cfd1 has sequence similarity to the C-terminus of Nbp35, including the nucleotide binding domain and the CxxC motif binding a labile Fe–S cluster. As mentioned before, Cfd1 is absent in the green lineage [35].

The CIA targeting complex consists of four proteins: NAR1 has intriguing similarity to [Fe–Fe] hydrogenases; the CIA1 protein is thought to act as a protein binding platform of the complex; the AE7 protein has an hyper-reactive thiol; and the MET18 protein belongs to the HEAT repeat proteins (figure 4). Although the precise molecular functions of these proteins remain to be determined, the formation of the complex and its biological function is conserved from yeast to humans and plants [37,48,49]. A universal phenotype of cia mutants is the loss of nuclear genome integrity. Interestingly, early observations in Arabidopsis already pointed to altered nuclear morphology and increased expression of DNA repair proteins in an atm3 mutant [12]. Also, AtTAH18 was suggested to have a role in DNA

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**Figure 3.** Cofactors and activity of AtTAH18 and AtDRE2. (a) UV-visible absorbance spectrum of purified AtTAH18 (3 µM) in 25 mM Tris–HCl, pH 8.0, 150 mM NaCl. Buffer as in (a). (c) UV-visible absorbance spectra of 30 µM purified AtDRE2 before (as isolated) and after chemical reconstitution and desalting (reconstituted). Buffer as in (a). (d) X band EPR spectra of purified (as isolated) AtDRE2 (100 µM) reduced with 2 mM sodium dithionite in buffer. EPR conditions: microwave frequency, 9.458 GHz; microwave power, 1.3 mW (10 K) or 20 mW (84 K); modulation amplitude, 1.25 mT; modulation frequency, 100 kHz. Simulation parameters for the EPR signal: $g_x, g_y$ and $g_z$: 1.998, 1.959 and 1.916 with line widths of 2.3, 4.1 and 3.4 mT, respectively.

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The CIA targeting complex consists of four proteins: NAR1 has intriguing similarity to [Fe–Fe] hydrogenases; the CIA1 protein is thought to act as a protein binding platform of the complex; the AE7 protein has an hyper-reactive thiol; and the MET18 protein belongs to the HEAT repeat proteins (figure 4). Although the precise molecular functions of these proteins remain to be determined, the formation of the complex and its biological function is conserved from yeast to humans and plants [37,48,49]. A universal phenotype of cia mutants is the loss of nuclear genome integrity. Interestingly, early observations in Arabidopsis already pointed to altered nuclear morphology and increased expression of DNA repair proteins in an atm3 mutant [12]. Also, AtTAH18 was suggested to have a role in DNA
replication [24]. These observations are now understood because replicative DNA polymerases have been shown to require an Fe–S cluster [50]. However, differences between eukaryote species have also been observed. For example, the Arabidopsis met18 mutant has no obvious phenotype, although in combination with a single allele of a partial ac7 mutant, the phenotype is strongly enhanced compared with the ac7 homozygote. In contrast, met18 mutants in yeast do have a slow-growth phenotype and methionine auxotrophy, whereas mouse embryos lacking the MET18 homologue MMS19 are embryo lethal [36,47].

Key questions that remain with respect to the assembly of cytosolic/nuclear Fe–S clusters are (i) what is the sulfur-containing substrate of the mitochondrial ABC transporters? (ii) What is the source of Fe and how is it delivered to NBP35? While fast progress has been made in yeast, studies in other eukaryotes are also providing new information and different perspectives.

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ABA3 cysteine desulfurase enzyme encoded by the ABA3 gene (ABSCISIC ACID3). The enzyme is required for sulfuration of Moco in aldehyde oxidases and xanthine dehydrogenases. Aldehyde oxidase 3 carries out the final step in the biosynthesis of abscisic acid, an important plant hormone.

ABC ATP-binding cassette

ABCDEFG ATP-binding cassette transporter, B group, number 7

ACO aconitase

AO aldehyde oxidase

At as prefix to gene and protein names, meaning Arabidopsis thaliana

Atm1 ATP-binding cassette (ABC) transporter of the mitochondria in Saccharomyces cerevisiae

ATM3 ATP-binding cassette (ABC) transporter of the mitochondria in Arabidopsis thaliana

ATR1, ATR2 or ATR3

CFD1 cytosolic Fe–S cluster deficient

CIA genes genes identified based on their requirement for the assembly of Fe–S clusters on cytosolic and nuclear proteins, but not on mitochondrial Fe–S proteins

CIA pathway cytosolic iron–sulfur protein assembly pathway

CIAPIN1 cytokine-induced apoptosis inhibitor 1; human homologue of DRE2

CpNifS chloroplast NifS-like (nitrogenase fixation S-like) protein, the cysteine desulfurase enzyme localized in chloroplasts

CxxC cysteine-x-x-cysteine amino acid sequence, with x denoting any amino acid

DRE2 Fe–S binding protein encoded by the DRE2 gene, derepressed for ribosomal protein S14 expression

EPR electron paramagnetic resonance

Fe–S iron–sulfur

Gal galactose; in this text used as an abbreviation for galactose-inducible promoter

GFP green fluorescent protein

HEAT tandemly repeated, 37–47 amino acid long-module occurring in a number of cytoplasmic proteins, including the four name-giving proteins huntingtin, elongation factor 3, alpha regulatory subunit of protein phosphatase 2A and TOR1

isc gene cluster iron–sulfur cluster assembly

ISC pathway iron–sulfur cluster (assembly) pathway

ISCS human homologue of the bacterial IscS protein, a type-I cysteine desulfurase

MET25 yeast gene involved in methionine synthesis. The promoter is used in this study to drive constitutive expression of transgenes

Moco molybdenum cofactor

NFS1 NifS-like gene encoding a cysteine desulfurase

NFU2 NifU-like protein 2

NP natural promoter

Sc as prefix to gene and protein names, meaning Saccharomyces cerevisiae

Sir sirtinol resistant allele 3–3 (not to be confused with sulfite reductase!)

suf operon bacterial operon of six genes involved in sulfur mobilization for the biosynthesis of Fe–S clusters under oxidative stress and iron limitation.

SUF pathway sulfur mobilization pathway

TAH18 gene that was first identified in a mutant screen for Top1T722A Hypersensitivity.

TDH yeast gene encoding a Triose-phosphate DeHydrogenase. The promoter is used in this study to drive the expression of transgenes at very high levels.

T-DNA transfer DNA; large segment of DNA transferred by Agrobacterium into the plant genome

WT wild-type

XDH xanthine dehydrogenase