

FORUM REVIEW ARTICLE

# Cytochromes *b561*: Ascorbate-Mediated Trans-Membrane Electron Transport

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## Abstract

**Significance:** Cytochromes *b561* (CYB561s) constitute a family of trans-membrane (TM), di-heme proteins, occurring in a variety of organs and cell types, in plants and animals, and using ascorbate (ASC) as an electron donor. CYB561s function as monodehydroascorbate reductase, regenerating ASC, and as Fe<sup>3+</sup>-reductases, providing reduced iron for TM transport. A CYB561-core domain is also associated with dopamine  $\beta$ -monooxygenase redox domains (DOMON) in ubiquitous CYBDOM proteins. In plants, CYBDOMs form large protein families. Physiological functions supported by CYB561s and CYBDOMs include stress defense, cell wall modifications, iron metabolism, tumor suppression, and various neurological processes, including memory retention. CYB561s, therefore, significantly broaden our view on the physiological roles of ASC. **Recent Advances:** The ubiquitous nature of CYB561s is only recently being recognized. Significant advances have been made through the study of recombinant CYB561s, revealing structural and functional properties of a unique “two-heme four-helix” protein configuration. In addition, the DOMON domains of CYBDOMs are suggested to contain another heme *b*. **Critical Issues:** New CYB561 proteins are still being identified, and there is a need to provide an insight and overview on the various roles of these proteins and their structural properties. **Future Directions:** Mutant studies will reveal in greater detail the mechanisms by which CYB561s and CYBDOMs participate in cell metabolism in plants and animals. Moreover, the availability of efficient heterologous expression systems should allow protein crystallization, more detailed (atomic-level) structural information, and insights into the intra-molecular mechanism of electron transport. *Antioxid. Redox Signal.* 19, 1026–1035.

## Introduction

THE IMPACT OF oxidation-reduction (redox) reactions on many fundamental cellular processes is increasingly recognized. An intricate network of redox proteins and low-molecular-weight electron donors and acceptors control redox homeostasis in many cell types. Ascorbate (ASC) is an essential component of this redox network in plants and animals, mainly as an electron donor in enzymatic reactions, including H<sub>2</sub>O<sub>2</sub> scavenging by ASC peroxidase (APX) (19). However, ASC is also a principal electron donor to a family of di-heme, trans-membrane (TM) proteins, cytochromes *b561* (CYB561s), catalyzing TM electron transport (TMET). CYB561s occur in a wide range of animal and plant phyla, and

are generally encoded by small gene families. Their name is derived from the characteristic  $\alpha$ -band absorption maximum (reduced minus oxidized).

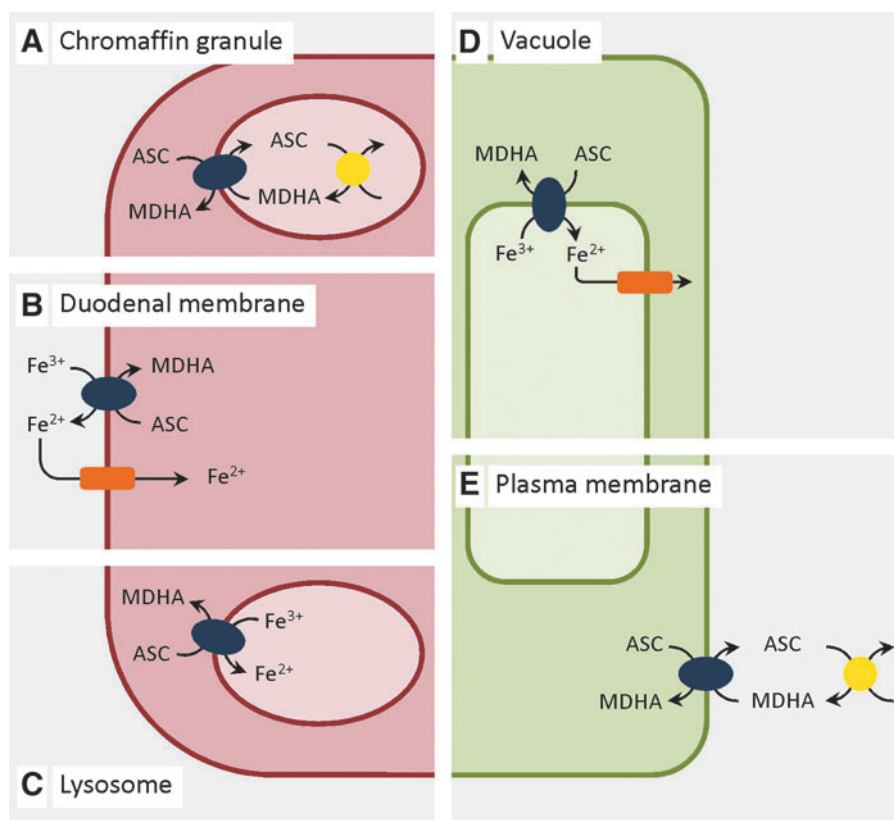
CYB561 members are implicated in physiological processes, including plant ‘front-line’ (apoplastic) defense; iron uptake; ASC regeneration in neuroendocrine tissues; and tumor growth. CYB561s are also of interest because of their particular biochemical (structure-function) properties. They are composed of six TM  $\alpha$ -helices, the central four of which make up the ‘CYB561-core,’ coordinating two heme *b* molecules with four conserved His. The core domain harbors the intra-molecular electron transfer, and, intriguingly, is a building block in other proteins, often in association with other redox domains such as the N-terminal domain of

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**FIG. 1. Subcellular localization and orientation of CYB561 family members in plant and animal cells.** CYB561s (blue ovals) catalyze trans-membrane electron transport (TMET) through membranes, separating cytoplasm and acid compartments (secretory vesicles, vacuole, and extracellular matrix). Localization, orientation, and electron donor/acceptor couples are based on experimental evidence (on native and recombinant proteins), and ‘most plausible’ interpretations (see text). Ascorbate (ASC) appears a principal electron donor for all CYB561s, whereas both monodehydroascorbate (MDHA) and  $\text{Fe}^{3+}$  may act as electron acceptors. Animal CYB561s: **(A)** Chromaffin granules: CGCytb/CYB561A1 in chromaffin granule vesicle membranes, involved in catecholamine synthesis operating in concert with ASC-dependent oxidases (yellow circles); **(B)** duodenal DCytb/CYB561A2 in the duodenal cell membrane, reducing intestinal  $\text{Fe}^{3+}$ -chelates before uptake through  $\text{Fe}^{2+}$ -transporters (orange rectangles); **(C)** lysosomal membrane LCytb/CYB561A3. Plant CYB561: **(D)** plant tonoplast TCytb/CYB561B1 may reduce vacuolar  $\text{Fe}^{3+}$  for transport to the cytoplasm; **(E)** (putative) plant plasma membrane CYB561B acting in concert with extracellular oxidases (e.g., ASC oxidase). To see this illustration in color, the reader is referred to the web version of this article at [www.liebertpub.com/ars](http://www.liebertpub.com/ars)

dopamine  $\beta$ -monoxygenase (DOMON). Together, the CYB561 protein family expands our horizon on the role of ASC in cell metabolism.

### Physiological Functions and Biological Relevance

#### ASC recycling

Some CYB561s support ASC regeneration, through one-electron reduction of partially oxidized ASC, monodehydroascorbate (MDHA) to ASC, using electrons provided by ASC, on ‘the other’ side of the membrane. In animals, such TMET activity is catalyzed by the chromaffin granule CYB561 (CGCytb/CYB561A1, for systematic nomenclature, see Supplementary Table S1; Supplementary Data are available online at [www.liebertpub.com/ars](http://www.liebertpub.com/ars)), the first CYB561 identified (18). Chromaffin granules harbor dopamine synthesis, involving ASC-dependent dopamine  $\beta$ -hydroxylation, generating MDHA. A similar ASC-regenerating function is required in neuropeptide storage vesicles to sustain ASC-dependent amidated peptide biosynthesis. The essential role of CYB561s in this process, and its impact on neurological functions, was

recently illustrated in the *Drosophila* mutant *nemy*, where decreased expression of a CYB561 homologue resulted in reduced memory retention (24). The ‘electron shuttle’ activity of CGCytb/CYB561A1, between ASC and membrane-impermeable ferricyanide or MDHA, was demonstrated in native chromaffin granule ‘ghost’ vesicles and reconstituted membrane systems (11, 44) (Fig. 1A). Further support on the role of CYB561s in ASC regeneration is found in erythrocytes, where intracellular ASC reduces extracellular MDHA (54, 60).

Work on plant membranes revealed the presence of ASC-reducible, di-heme-like *b*-type cytochromes, with similar biophysical properties as bovine CGCytb/CYB561A1, in several species (4, 5, 22). A *Phaseolus vulgaris* CYB561, presumably plasma membrane (PM)-located, mediated TMET in purified, ASC-loaded, vesicles (2). Extracellular ASC provides antioxidant defense against gaseous oxidants (ozone, NO<sub>x</sub>), and CYB561 could be involved in apoplasmic ASC regeneration (Fig. 1E). The *Arabidopsis* genome reveals four genes encoding six-TM CYB561s with an average of around 30% sequence identity to CGCytb/CYB561A1 (3), but it is not yet clear which of the gene products is involved in PM TMET.

### *Fe<sup>3+</sup>-reductase activity and iron homeostasis*

A new CYB561 member was identified in the cell membrane of mouse duodenum epithelial cells, in response to hypoxia and iron deficiency (39). Mouse duodenal DCytb/CYB561A2 reduces intestinal Fe<sup>3+</sup> before uptake (Fig. 1B). ASC is an electron donor to mouse DCytb/CYB561A2, at least *in vitro* (54). The elevated expression of DCytb/CYB561A2 in mice with systemic iron overload (hemochromatosis) provides additional support for a role in iron uptake (23).

ASC-mediated TM Fe<sup>3+</sup>-reductase activity was confirmed by complementation of *Δfre1Δfre2* Fe<sup>3+</sup>-reductase-deficient yeast lines. Reduction of extracellular Fe<sup>3+</sup>-chelates in transformed yeast cells, expressing genes for three mouse and one Arabidopsis CYB561 protein (CGCytb/CYB561A1, DCytb/CYB561A2; LCytb/CYB561A3; TCytb/CYB561B1, respectively), was demonstrated (10, 54). For CGCytb/CYB561A1, the TM Fe<sup>3+</sup>-reductase activity was also demonstrated in ASC-loaded proteo-liposomes (11). These experiments suggest that, at least *in vitro*, all CYB561s may have Fe<sup>3+</sup>-reductase activity. Therefore, the actual *in vivo* activity of CYB561s, that is, Fe<sup>3+</sup> reduction versus MDHA reduction, is probably dependent on substrate availability and biological context.

Free Fe<sup>3+</sup> would readily be reduced by ASC, if occurring in the same compartment, and Fe<sup>2+</sup> catalyzes free radical generation through Fenton-type reactions. Separation of Fe<sup>3+</sup> and ASC, along with CYB561 catalyzing TMET, thus provides efficient control over ASC-mediated Fe<sup>3+</sup> reduction.

Arabidopsis TCytb/CYB561B1, used in the yeast Fe<sup>3+</sup>-reduction assays, is localized in the tonoplast (20, 50) (Fig. 1D). Given that the Arabidopsis TCytb/CYB561B1 ASC-binding (electron acceptor) site is predicted on the cytoplasmic protein side, this suggests that this CYB561 could reduce Fe<sup>3+</sup> in the vacuolar lumen, supporting transport to the cytoplasm. Vacuolar Fe<sup>2+</sup>-transporters (NRAMP3, NRAMP4), which may act in conjunction with TCytb/CYB561B1, have been identified (27). Possibly plant vacuoles store iron for remobilization when metabolic needs exceed supply. ASC-dependent CYB561-mediated Fe<sup>3+</sup>-reductase activity in plants may complement the well-characterized NADH-dependent activity of the FRO proteins (27).

### *Other physiological functions*

In addition to a role in ASC and Fe metabolism, CYB561 members may be involved in other physiological processes. A watermelon PM CYB561B is up-regulated under drought and high-light stress and was proposed to constitute a route for thermal dissipation of excess light energy by functional interaction with apoplastic ASC oxidase (43) (Fig. 1E). In humans, a CYB561 (TSCytb/CYB561D2/101F6) is encoded in chromosomal region 3p21.3 and is related to suppression of tumor cell growth. Treating lung cancer cells with TSCytb/CYB561D2 plasmid-DNA nanoparticles and ASC, synergistically suppressed lung cancer cell development (46). For yet other CYB561s, limited functional information is still available. For example, mouse LCytb/CYB561A3 is located in lysosomal membranes in various tissues and is possibly associated specifically with macrophages (64) (Fig. 1C). However, its physiological impact is not yet clear. Lysosomal iron pools play a role in oxidative stress control (30), which could involve LCytb/CYB561A3-Fe<sup>3+</sup>-reductase activity. Finally, one should consider that other electron

donors and acceptors (than ASC and MDHA/Fe<sup>3+</sup>) may still be identified, linking CYB561s to yet other physiological processes.

### *Tissue- and subcellular-distribution*

CGCytb/CYB561A1 was first identified in catecholamine-synthesizing (chromaffin) vesicles, and was, consistently, demonstrated in various neuroendocrine tissues (53). It is now clear that plant and animal CYB561s occur in a wide range of organs, tissues, and cell types. Remarkably specific cell-type localizations are sometimes observed, but not yet understood, such as testicular Sertoli cells for mouse LCytb/CYB561A3 (64). Some animal organs (*e.g.*, lung, spleen, brain, and duodenum) contain multiple CYB561 isoforms. However, in these cases, the proteins are apparently not located in identical cell types or membrane types. In Arabidopsis, CYB561 isoforms show some, but not complete, overlapping organ distribution patterns, and CYB561s occur in different membrane types (52, 62). Therefore, CYB561 isoform expression does not appear to be redundant. In a recently isolated Arabidopsis homozygous KO of TCytb/CYB561B1, transcription of at least three other CYB561 isoforms was not enhanced, supporting this notion (unpublished).

The subcellular membranes containing CYB561 consistently are membranes separating the cytoplasm and an acidic compartment, including acidic extracellular environments in plants and animals (Fig. 1). This pH gradient is most likely maintained by the activity of H<sup>+</sup>-transporting ATPases, and is probably important in CYB561 TMET function.

### **CYB561 Structure and Structure-Function Relationship**

The CYB561 protein family, in plants and animals, was identified on the basis of sequence comparison to the amino acid sequence of the bovine chromaffin granule CGCytb/CYB561A1 (59) (Fig. 2). Until today, this is also the only native CYB561 protein for which biochemical and biophysical measurements are available. Biochemical and structural information of other plant and animal CYB561s is derived from recombinant proteins, expressed in yeast (*Saccharomyces cerevisiae*, *Pichia pastoris*), insect (Sf9) cells, and *Escherichia coli* (8, 11, 15, 20, 33, 35–37, 41, 45, 54). CYB561s are membrane-spanning proteins that are generally accepted to contain six TM  $\alpha$ -helix domains and two pairs of His residues, arranged on four consecutive TM helices, coordinating two *b*-type hemes, located on each side of the membrane and accepting electrons from ASC (Figs. 2 and 3). Crystal-structure-level 3D atomic information is not (yet) available for any CYB561. However, 3D atomic resolution models were calculated for human CGCytb/CYB561A1, and Arabidopsis TCytb/CYB561B1 to support data analysis (6) (Fig. 4).

### *Midpoint redox potentials*

Redox titration results consistently reveal the presence of two one-electron redox centers (*b*-type hemes), with midpoint redox potential ( $E_0'$ ) values ranging from +80 to +190 mV for the 'high potential' (HP) heme, and –20 to +60 mV for the 'low potential' (LP) heme, (15, 35, 56). Among CYB561 isoforms, there is a relatively consistent difference of about 100 mV between the  $E_0'$  of the heme centers, supporting intra-molecular electron transfer mechanisms. Recently, it was demonstrated

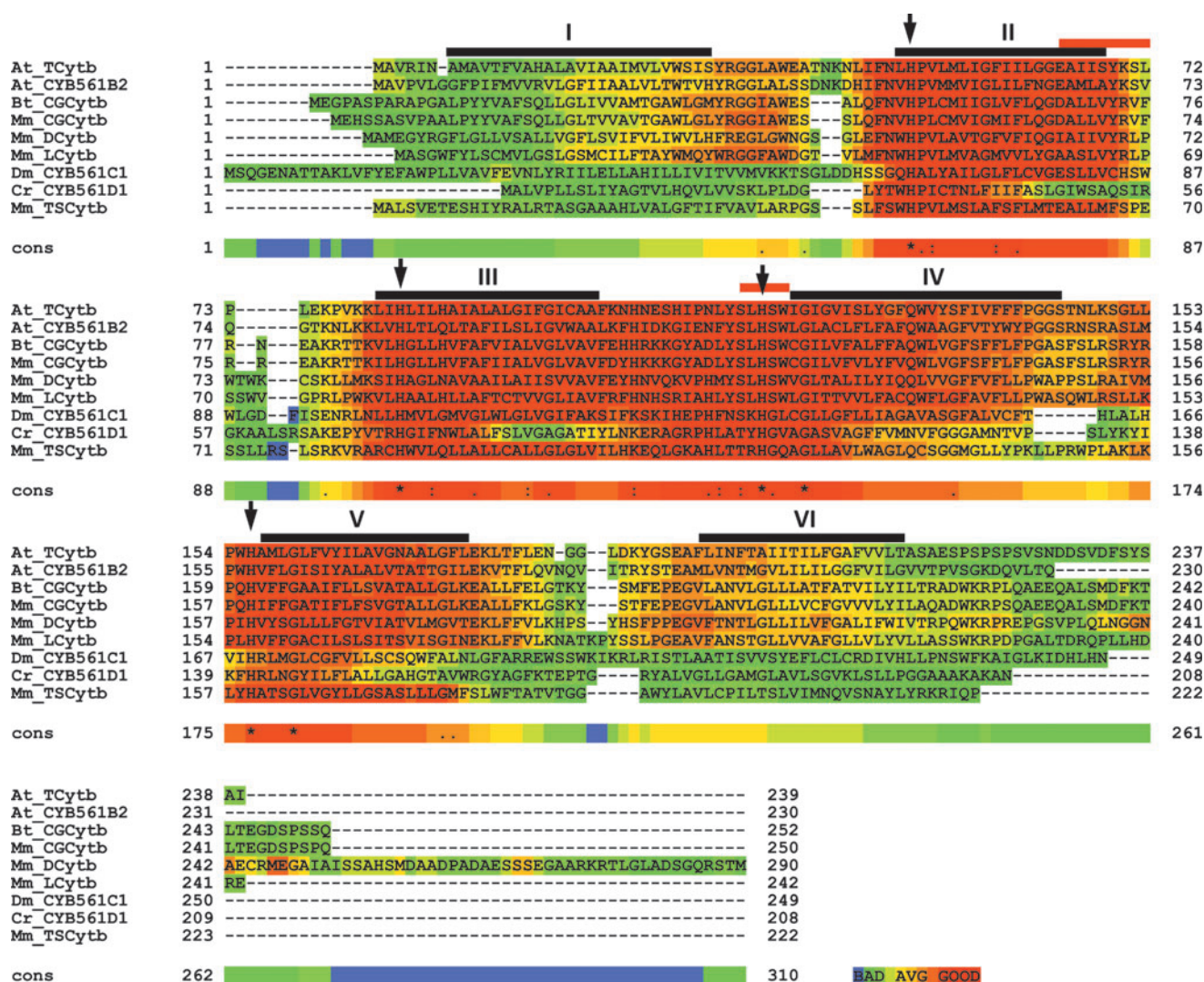


FIG. 2. Multiple alignment of CYB561s, showing TM domains, conserved residues, and substrate-binding domains. ClustalW2.1 alignment of selected CYB561 sequences, representing isoforms from most evolutionary clusters (59) (see Fig. 5), showing TM domains (I–VI, black bars), conserved His residues (arrows), and putative substrate interacting sites (red bars). Sequences are as follows: (i) *Arabidopsis thaliana* tonoplast CYB561 (At\_TCytb/CYB561B1; Q8L856); (ii) At\_CYB561B2 (Q9SW51); (iii) *Bostaurus* chromaffin granule CYB561 (Bt\_CGCytb/CYB561A1; P10897); (iv) *Mus musculus* chromaffin granule CYB561 (Mm\_CGCytb/CYB561A1; Q60720); (v) *M. musculus* duodenal CYB561 (Mm\_DCytb/CYB561A2; Q925G2); (vi) *M. musculus* lysosomal CYB561 (Mm\_LCytb/CYB561A3; Q6P1H1); (vii) *Drosophila melanogaster* CYTB561 (Dm\_CYB561C1; Q9W4U9); (viii) *Chlamydomonas reinhardtii* CYB561 (Cr\_CYB561E1; XP\_001702111.1); and (ix) *M. musculus* tumor suppressor CYB561 (Mm\_TSCytb/CYB561D2; Q9WUE3). Color coding of sequence similarity is performed using the T-Coffee alignment evaluation (17). To see this illustration in color, the reader is referred to the web version of this article at [www.liebertpub.com/ars](http://www.liebertpub.com/ars)

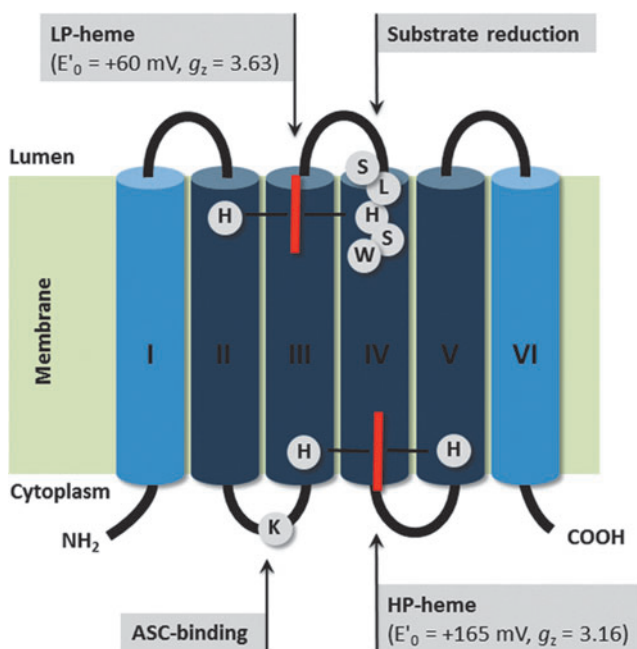
by electron paramagnetic resonance (EPR) and site-directed mutagenesis that the HP-heme is located on the cytoplasmic side of the TM proteins (16).

#### ASC reducibility

Reducibility by ASC is recognized as a particular CYB561 feature, generally reaching reduction levels of 50% to 90% (relative to Na-dithionite [DTH]). Why 100% reduction levels are not reached, even at high ASC concentrations, is not yet fully understood, in part because the ASC and/or DTH reduction mechanism is not yet fully unraveled. The midpoint redox potentials of the electron transfer reactions involving ASC are around +320 mV (MDHA/ASC) and –200 mV

[dehydroascorbate (DHA)/MDHA], and +60 mV for the DHA/ASC redox couple (25). Since the MDHA concentration in the cell is orders of magnitude lower than that of ASC (due to disproportionation), the electron transfer from ASC to CYB561 is favored under physiological conditions. Thermodynamic evidence supports the idea that CYB561-mediated TMET may be coupled to a local protonation/de-protonation cycle (42, 44).

Direct molecular-binding interactions between ASC and CYB561 have not been measured, but steady-state and fast kinetic measurements on plant and animal CYB561s suggest the existence of two distinct ASC-affinity sites, with approximate values of 0.01 mM and 1 mM, for CGCytb/CYB561A1 (7, 11, 28). These parameters are not real binding constants but rather characterize the interaction between ASC



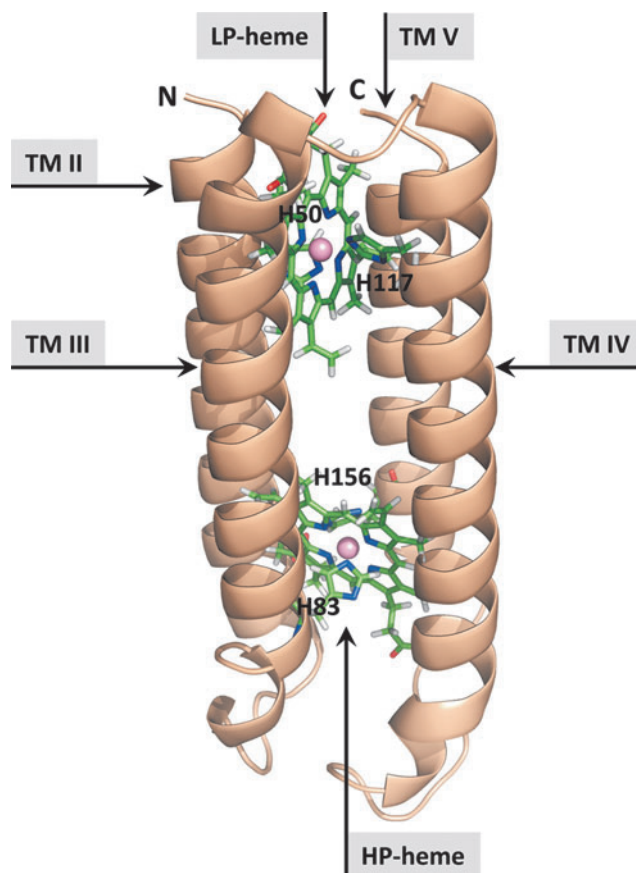
**FIG. 3. Schematic 2D structure of the Arabidopsis TCytb/CYB561B1 TM organization, including hemes, heme coordination, and putative substrate binding sites.** CYB561s are composed of six TM domains (I-VI), with two heme-*b* molecules (red bars), coordinated by four conserved His on TM helices II-V. The high potential (HP)-heme is located on the cytoplasmic protein side, accepting electrons from ASC (or other electron donors). The heme coordination and orientation is confirmed through electron paramagnetic resonance studies on site-directed-mutant proteins. Also shown is a Lys residue, possibly involved in substrate (ASC) binding, and a well-conserved putative substrate-binding site (SLHSW). Dark blue: CYB561 four helix core domain. To see this illustration in color, the reader is referred to the web version of this article at [www.liebertpub.com/ars](http://www.liebertpub.com/ars)

and CYB561, resulting in the reduction of the HP and LP-hemes. The effect of diethylpyrocarbonate on ASC reduction kinetics also supports this hypothesis (58). A comparison of conserved sequence regions in the CYB561 core led to the tentative assignment of two substrate binding sites located on the cytoplasmic and luminal side of the protein (47, 59) (Fig. 2).

Spectrum analysis of mouse and bovine CGCytb/CYB561A1 and of TSCytb/CYB561D2 revealed the presence of two distinct, split  $\alpha$ -bands in the spectrum of ASC-reduced proteins, which is consistent with the presence of two heme pockets (9, 12, 28). In Arabidopsis TCytb/CYB561B1 and mouse TSCytb/CYB561D2, singular value decomposition analysis identified two distinct *b*-type heme spectra that could be assigned to the two CYB561 hemes (12, 16). The property of split  $\alpha$ -bands points to the fact that each heme is located in an anisotropic electrostatic field, originating from the distribution of charged amino acid residues around the heme pockets. Given the limited primary structure similarity between CYB561 isoforms, it is not surprising that the fine structure of the split  $\alpha$ -band slightly differs, not only for the two hemes but also between isoforms (9, 12, 15, 16).

#### Other reducing agents

Common reductants such as NAD(P)H and reduced glutathione (GSH) are poor electron donors to CYB561s.



**FIG. 4. Calculated 3D atomic structure model of the Arabidopsis TCytb/CYB561B1-core domain.** Based on extensive CYB561s sequence analysis, structural constraints were identified to predict 3D structural models, using molecular modeling tools. Counterclockwise helix topologies are shown, including the HP- and low-potential (LP) hemes (ball-and-stick). Balls represent the Fe-atoms [modified after (6)]. To see this illustration in color, the reader is referred to the web version of this article at [www.liebertpub.com/ars](http://www.liebertpub.com/ars)

Reduction by NAD(P)H of various CYB561 isoforms never exceeded 5% (relative to DTH) (12, 57). GSH does not reduce CYB561, but other thiol reagents, in particular dihydrolipoic acid (DHLA), are almost as efficient as ASC in reducing mouse CGCytb/CYB561A1 and TSCytb/CYB561D2; however, with only one apparent, high affinity, binding site (12). In addition, the Arabidopsis TCytb/CYB561B1 was reduced by DHLA, although with much lower sensitivity and biphasic concentration dependence (12). Thus, it is possible that DHLA plays a different role in CYB561 reduction in plants and animals. In cells, DHLA is present mostly in its lipoyl-accepting form, tethered to conserved Lys residues on lipoyl-accepting protein domains (13). It may, therefore, be worth to investigate lipoyl-domain-containing proteins as potential CYB561 redox partners. In addition, dithioerythritol, and its stereoisomer, dithioerythritol, reduce Arabidopsis TCytb/CYB561B and mouse TSCytb/CYB561E (12, and unpublished).

#### Site-directed mutagenesis

Important structure-function information is derived from targeted mutations in the highly conserved heme-coordinating

His residues of the CYB561-core (Fig. 2 and 3). For example, detailed EPR studies of Arabidopsis TCytb/CYB561B1 mutant proteins revealed that the HP-heme center is located on the cytosolic side of the protein (16), opposite to that previously suggested (56, 58). Western blot analysis and spectroscopy demonstrated that mutation of His residues coordinating the LP-heme (intra-vesicular-side) of Arabidopsis TCytb/CYB561B1, and mouse CGCytb/CYB561A1, resulted in nearly undetectable protein levels. In contrast, mutations in the HP-heme-coordinating residues hardly affected CYB561 expression, but resulted in altered ASC-reduction kinetics and reduced heme content (11, 28, 34). Somewhat contradicting results were obtained with His mutations in human DCytb/CYB561A2, and need further investigation (37, 45).

Positively charged residues near the hemes may facilitate the interaction with the ASC and MDHA anions, as is, for example, the case in the ASC-mediated reduction of soybean APX (14, 31). Lys and Arg residues are frequently found in the loop regions between the TM helices (Fig. 2), and an Arg residue (R72 in mouse CGCytb/CYB561A1), located near the HP-heme, is fully conserved among CYB561A members (59). Mutagenesis of R72 resulted in the loss of the 'high affinity' ASC reduction, supporting its role in substrate binding and suggesting that the high-affinity ASC-binding site was close to the HP heme (11). Arabidopsis TCytb/CYB561B1 contains no Arg in the vicinity of the heme, but still displays two distinct affinity sites (7). Its role is possibly replaced by a conserved Lys (K80) residue. This idea is supported by mutagenesis of K83 in maize CYB561B1, which resulted in altered midpoint redox potentials and ASC-reduction kinetics (41).

Mutations in other conserved and partially conserved residues of mouse CYB561s had variable effects on TMET activity and ASC reduction (34, 41, 45, 54). Consistent with the putative location of the electron-accepting and -donating sides, mutations in loop regions on the intra-vesicular side severely decreased LCytb/CYB561A3 TM  $\text{Fe}^{3+}$ -reductase activity, but mutations on the cytoplasmic side had comparatively less effect (54).

A key function of CYB561s is TMET, through sequential reduction and oxidation of the HP and LP-hemes. Conserved residues presumably located in TM domains (e.g., Q131 in mouse LCytb/CYB561A3/) have been identified, and mutation resulted in strongly decreased  $\text{Fe}^{3+}$ -reductase activity (54). This suggests that electron transfer may involve oxidation/reduction of intramolecular amino acids. On the other hand, electron transfer along chains of prosthetic groups is well established, as long as the "edge-to-edge" distance between groups is smaller than 14 Å (15, 48).

### CYB561-Domain-Containing Proteins

#### CYB561-DOMAIN

Identification of a CYB561-core domain at the C-terminus of a protein named stromal cell-derived receptor 2 (SDR2) provided evidence that CYB561s could be part of multi-domain proteins (49) (Fig. 5). Bio-informatic analyses demonstrated that most commonly a single CYB561-core follows a DOMON-domain (CYB561-DOMAIN), but some proteins contain up to four DOMONs in a row (e.g., C13B4.1 of *C. elegans*; Fig. 5) (1, 26, 49, 51).

The DOMON-domain refers to the N-terminal domain of dopamine  $\beta$ -monoxygenase (or dopamine[ $\beta$ ]-hydroxylase, DBH); (1, 49), catalyzing the reduction of dopamine to

norepinephrine in chromaffin granules. In the process, ASC is oxidized to MDHA, which may be regenerated by CYB561 (Fig. 1A). The DOMON-domain is widespread in nature, often in association with redox domains such as the  $\text{Cu}^{2+}$ -monoxygenase domain in DBH or CYB561 in CYB561-DOMAINs, and/or with adhesion modules (e.g., reelin-domain in SDR2) (Fig. 5) (26). In DBH, this module is probably involved in tetramerisation (29), but not in catalysis or catecholamine binding (61).

CYB561-DOMAIN sequences have been identified in the genome of animals, fungi, plants, and prokaryotes (26). Eukaryotic CYB561-DOMAINs were classified in two major groups. The 'F'-group, including plant proteins with single CYB561- and DOMON-domains, and the 'G'-group, including SDR2-related sequences (from animals or plants), sometimes with multiple DOMON domains (59) (Fig. 5). G-group CYB561-DOMAINs often also contain the reelin domain (animals) or DM13 domain (plants) (26, 51).

It is now clear that plants contain larger CYB561-DOMAIN gene families than do other organisms (51, 59). For example, Arabidopsis contains at least ten CYB561-DOMAIN-containing genes: eight belonging to the F-group, one to the G-group, and a divergent gene that could not easily be classified (51). In addition, Arabidopsis and other plant genomes contain CYB561-DOMAIN-related genes with either no DOMON or no CYB561-domain (e.g., see section on heme-b binding). On the contrary, SDR2 is the only CYB561-DOMAIN of the human genome, and single SDR2 orthologous genes are found in mouse and fly genomes, while *C. elegans* contains four G-group CYB561-DOMAINs (59).

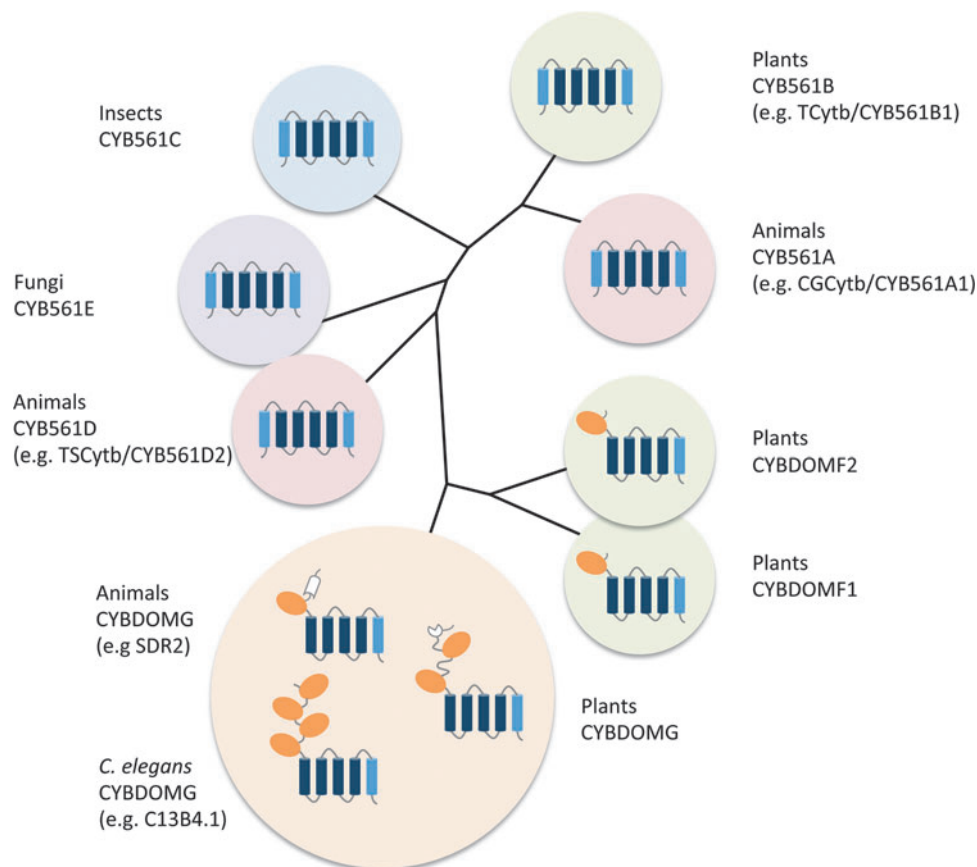
All of these CYB561-DOMAINs contain a canonical CYB561-core, with four fully conserved His, at the N-terminus of four consecutive TM  $\alpha$ -helices, plus an additional C-terminal  $\alpha$ -helix, corresponding to the sixth  $\alpha$ -helix of CYB561s (Fig. 5). In spite of these strictly conserved features, CYB561s and CYB561-DOMAINs have highly divergent sequences (59).

A common feature of CYB561-DOMAIN sequences, at least in eukaryotes, is the presence of a signal sequence directing these proteins to the secretory pathway. Consistently, proteomic studies in plants have identified CYB561-DOMAIN proteins on the PM (55), in some cases in association with lipid rafts (32). In tobacco roots expressing a chimeric CYB561-DOMAIN-GFP from soybean, fluorescence was clearly associated with the PM (A. Costa and P. Trost, unpublished). In mammals, SDR2 is also believed to reside in the PM (61).

Topology predictions show that the C-terminus of CYB561-DOMAINs (corresponding to the C-terminus of CYB561s) is cytosolic and, consequently, the N-terminus of the five- $\alpha$ -helices CYB561-domain, connecting to the DOMON domain(s), is extracellular. Indeed, DOMON domains are commonly extracellular and either heme- or sugar-binding domains (26). The heme-binding DOMON domain of cellobiose dehydrogenase (CDH) of ligninolytic fungi plays a role in the attack to the lignocellulosic extracellular matrix of plants (63). Sugar-binding DOMON domains are also found in bacterial extracellular cellulases (26), overall suggesting a role of DOMON-containing proteins in modifying extracellular polymers (see section on Why CYB561-DOMAINs?).

#### $\text{Fe}^{3+}$ -reductase activity

The CYB561-core in CYB561-DOMAINs contains all necessary elements to catalyze TMET. Recombinant fly and mouse SDR2 demonstrate  $\text{Fe}^{3+}$ -reductase activity in *Xenopus* oocytes (61).



**FIG. 5. Evolutionary relationship between CYB561s and CYB561-core-containing CYBDOM proteins.** CYB561 and CYBDOM proteins, cluster in various subgroups. The CYB561-core domain (four TM helices containing four conserved His and two hemes in bis-His coordination) is dark blue (see also Fig. 3). In the CYBDOM protein family, the CYB561 domain constituted by a CYB561-core plus the C-terminal  $\alpha$ -helix of CYB561s is combined with one or more dopamine  $\beta$ -mono-oxygenase N-terminal domain (DOMON) domains (brown ovals). Cluster identifications are based on (59), except that clusters D and E were exchanged, for nomenclature reasons (see Supplementary Table S1). Each CYB561 cluster contains sequences from only animals or plants or insects or fungi, with the notable exception of *Chlamydomonas reinhardtii* CYB561 (XP\_001702111.1, see also Supplementary Table S1) that clusters with the 'fungal' cluster CYB561E. CYBDOMF is divided into F1 and F2 clusters with diverging DOMON sequences. CYBDOMG includes sequences with one DOMON plus a reelin domain (white), sequences with four DOMONS, and plant sequences with two DOMONS and a DM13 domain (white). To see this illustration in color, the reader is referred to the web version of this article at [www.liebertpub.com/ars](http://www.liebertpub.com/ars)

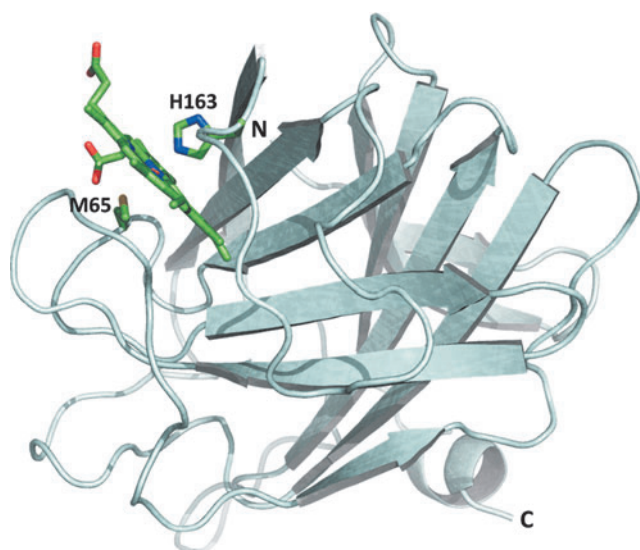
SDR2 is, therefore, considered a TM reductase and was renamed ferric chelate reductase 1 (e.g., in UniProt). Whether  $\text{Fe}^{3+}$ -reductase activity can be extended to other CYBDOMs is unknown. Since the CYB561-domain can catalyze by itself the TM reduction of  $\text{Fe}^{3+}$ -chelates, the role of the DOMON domain in SDR2 redox activity is uncertain. However, DOMON domains are most likely *b*-type cytochromes and may form a redox chain with CYB561 domains.

#### Heme-*b* binding in the DOMON domain

The DOMON-containing CDH of wood-degrading and phytopathogenic fungi is an extracellular flavo-cytochrome that participates in early events of lignocellulose degradation (38). The 3D structure of the DOMON-domain of CDH shows heme binding through an uncommon His-Met coordination (21) (Fig. 6). The heme is tightly fitted into a pocket of the concave inner  $\beta$ -sheet of the overall  $\beta$ -sandwich structure that is typical of the DOMON domain (26). Interestingly, both heme-axial ligands of *Phanerochaete chrysosporium* CDH (M65,

H163) are conserved in CYBDOM sequences, strongly suggesting that DOMON-domains of CYBDOMs may also bind a heme *b*. These residues are conserved neither in the DOMON of DBH (that bind no hemes) nor in DOMON proteins that bind a sugar in place of the heme (26).

A close relative to CYBDOMs is the plant protein AIR12, which is composed of a single DOMON domain, including the conserved Met-His pair that in *P. chrysosporium* is involved in heme ligation (21, 51). AIR12 is attached to the external leaflet of plant PMs via a GPI anchor and was experimentally demonstrated to bind a single heme *b* (51). The peculiar electron paramagnetic resonance spectrum of AIR12 is consistent with the Met-His heme-coordination displayed by CDH. AIR12 is the only CYBDOM-like DOMON-domain for which heme binding was experimentally demonstrated. Arabidopsis contains a single AIR12 gene, and AIR12-knock out mutants are resistant to the necrotrophic pathogen *Botrytis cinerea* (R. Barbaro, A. Costa, F. Sicilia, P. Trost, unpublished). Whether the function of AIR12 depends on the interaction with a CYB561 isoform (reminiscent of CYBDOM proteins) remains to be demonstrated.



**FIG. 6.** DOMON domain of *P. chrysosporium* cellobiose dehydrogenase (CDH), showing the uncommon *b*-type heme coordination by a Met-His pair. The DOMON domains of CYBDOM proteins are predicted to have the same 3D-fold based on two antiparallel  $\beta$ -sheets forming a  $\beta$ -sandwich structure as the DOMON domain of CDH (26). Although the sequence similarity between CYBDOM and CDH-DOMONS is limited, the pair of heme-coordinating residues Met65 and His163 (*P. chrysosporium* numbering) is strictly conserved. The structure was drawn from pdb file 1d7b (21). Heme and side chains of heme-coordinating residues are represented in sticks with the following color codes C (green), N (blue), O (red), and S (yellow). To see this illustration in color, the reader is referred to the web version of this article at [www.liebertpub.com/ars](http://www.liebertpub.com/ars)

#### Why CYBDOMs?

The architecture of CYBDOM proteins, with a heme *b*-binding DOMON-domain linked to a five- $\alpha$ -helix CYB561-domain, suggests an 'electron wire,' extending well beyond the PM surface. Asking how the function of CYBDOMs may be related to their redox activity and architecture is particularly challenging in plants, where CYBDOMs are often coded by ten genes or more.

*P. chrysosporium* lines, defective in the DOMON-enzyme CDH, fail to colonize wood. It has been proposed that CDH may participate in lignin degradation (38, 63). In this view, CDH, extending into the cell wall, reduces an  $\text{Fe}^{3+}$ -chelate (e.g.,  $\text{Fe}^{3+}$ -oxalate) with cellobiose being the reductant. In the presence of hydrogen peroxide (possibly produced by the flavo-domain of CDH itself), the reduced iron could generate hydroxyl radicals through Fenton chemistry, which may attack cell wall polymers, including lignin. CYBDOMs may well be involved in a similar reaction. The CYB561-core could provide cytoplasmic reducing equivalents to the extracellular heme(s), supporting hydroxyl radical-based modifications of cell wall polymers at a safe distance from the PM. As cell wall-loosening agents in plants, hydroxyl radicals are involved in fundamental processes such as for example, seed germination and cell elongation (40). In this context, CYBDOMs may constitute a redox bridge between cytosolic ASC and cell wall sites where hydroxyl radicals may be locally produced

by DOMON-catalyzed Fenton chemistry, similar to the DOMON-containing CDH of basidiomycetes.

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#### Abbreviations Used

APX = ascorbate peroxidase
ASC = ascorbate
CDH = cellobiose dehydrogenase
DHA = dehydroascorbate
DHLA = dihydrolipoic acid
DOMON = dopamine $\beta$ -monoxygenase N-terminal domain
HP = high potential
LP = low potential
MDHA = monodehydroascorbate
PM = plasma membrane
TM = trans-membrane
TMET = trans-membrane electron transport
SDR2 = stromal cell-derived receptor 2