Localization, Regulation, and Substrate Transport Properties of Bpt1p, a *Saccharomyces cerevisiae* MRP-Type ABC Transporter

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*Saccharomyces cerevisiae* Bpt1p is an ATP-binding cassette (ABC) protein that belongs to the MRP subfamily and is a close homologue of the glutathione conjugate (GS conjugate) transporter Ycf1p. The function of Bpt1p has previously been evaluated only in vitro, by using nonphysiological substrates. In the present study we examined the localization, regulation, and transport properties of Bpt1p in vivo, as well as its capacity to transport a set of prototypical MRP substrates in vitro. Our results show that Bpt1p, like Ycf1p, localizes to the yeast vacuolar membrane, plays a role in cadmium detoxification and ade2 pigmentation in vivo, and can participate in the transport of GS conjugates and glucuronate conjugates, as well as free glutathione, in vitro. However, in all of these cases the contribution of Bpt1p is substantially less than that of Ycf1p. In addition, the expression patterns of *YCF1* and *BPT1* differ significantly. Whereas *YCF1* expression is markedly increased by cadmium, adenine limitation in an ade2 strain, or overexpression of the stress-responsive transcription factor *Yap1p*, *BPT1* expression is only modestly affected under these conditions. Thus, although the functional capabilities of Bpt1p and Ycf1p overlap, their differences in regulation and substrate preference imply that they contribute to cellular detoxification processes in different ways.

With the complete genome of numerous organisms now in hand, the ATP-binding cassette (ABC) transporter superfamily has emerged as the largest membrane protein superfamily in both prokaryotes and eukaryotes, including microbes, plants, and animals (10–12, 17, 40, 45). Members of this superfamily catalyze the MgATP-energized transport of a broad range of substrates across biological membranes. Mutational loss of function of ABC proteins has been implicated in an increasing number of inherited diseases (11), and overexpression of certain ABC transporters has been shown to enhance multidrug resistance and the elimination of xenobiotics (1, 2). Consequently, elucidation of the biochemical activity, substrate specificity, and physiological regulation of ABC transporters is of both clinical and general biological significance.

Phylogenetic analysis has provided a valuable road map for formulating hypotheses about the function and substrate(s) of a particular transporter. ABC proteins can be divided into seven subfamilies based on sequence relatedness, designated ABCA through ABCG (11, 12, 45; http://nutrigene.4t.com/humanabc.htm). The emergent view is that members of a paragon have been rendered anionic by conjugation to GSH or glucuronate, processes by excreting potentially harmful endogenously and exogenously derived compounds, after these compounds have been rendered anionic by conjugation to GSH or glucuronate, or through complexing with GSH.

To gain further insight into multidrug resistance, it is important to understand how an organism may benefit by having multiple, highly homologous MRPs. Comparisons of the properties of the human MRP1, MRP2, and MRP3 transporters suggest several possibilities (3, 18, 21). First, these three MRPs appear to have overlapping, but distinct, substrate and kinetic profiles. Second, their intracellular sites of localization differ. Third, their regulatory properties differ, resulting in distinct patterns of tissue expression and/or response to extracellular stress (i.e., oxidative stress) (7). The combined effect of differences in these three parameters (spectrum of substrate specificity, location, and regulation) could impart a sufficiently high degree of specificity and flexibility to explain the distinctive
physiological roles of particular MRPs and to account for their large numbers in all organisms.

The yeast Saccharomyces cerevisiae presents a highly manipulable system for the genetic and biochemical analysis of both endogenous and heterologous ABC proteins (12, 45). Yeast cells can transport organic GSH conjugates and endogenous and heterologous ABC proteins (12, 45). Yeast SEY6210 mutants can transport and cadmium resistance in a significant manner, human MRP1 restores GSH conjugate trans- port and cadmium resistance in vivo (23, 25). Ycf1p can transport organic GSH conjugates and Cd- glutathione (Cd-GS) complexes into the vacuole (23, 25, 44). Deletion of YCF1 results in hypersensitivity to cadmium, while overexpression confers cadmium resistance. Ycf1p shares many biochemical properties with mammalian MRPs, and significantly, human MRP1 restores GS conjugate transport and cadmium resistance in ycf1-defective yeast strains (46). In addition, Ycf1p has been found to confer resistance to arsenite and diamide, and it is required for the vacuolar accumulation of a pigment precursor in ade2 mutants (5, 14, 49).

The closest homologe of YCF1 in yeast is YLL015. Since YLL015 was discovered through the yeast genome sequencing project and not from mutant or overexpression screens, its role in vivo as a transporter has been only hypothetical. Recently, a biochemical study indicated that YLL015, like YCF1, can mediate the in vitro transport of unconjugated bilirubin into yeast vacuole-derived vesicles (37). Accordingly, the gene has been designated BPT1 (named BPT for bile pigment transporter), even though this particular substrate is unlikely to have direct physiological relevance for yeast. Bpt1p and Ycf1p have also been shown to have overlapping activity for the in vitro transport of other nonphysiological substrates, including several Gd-based magnetic resonance imaging (MRI) contrast agents (34).

In the present study we have compared the localization, regulation, and transport properties of Bpt1p and Ycf1p both in vivo and in vitro. Our results show that Bpt1p, like Ycf1p, localizes to the vacuolar membrane, participates in cadmium detoxification and ade2 pigmentation in vivo, and has the capacity to catalyze the transport of GS and glucuronate conjugates, as well as that of free GSH in vitro. Although the transport properties of Bpt1p and Ycf1p overlap, their cellular impacts seem to differ considerably: the overall phenotypic and biochemical contributions of YCF1 consistently exceed those of BPT1. These differences in contribution may be due in part to differences in substrate selectivity and in part to the relative levels of each transporter, as determined by differences in their patterns of regulation. An interesting feature observed for some of the substrates was an unexpected dependence of Bpt1p on Ycf1p for maximal transport activity in vitro. Of the several potential explanations for this phenomenon, one intriguing possibility is that Bpt1p and Ycf1p form functional heterodimers.

MATERIALS AND METHODS

Yeast strains. The strains used in this study are listed in Table 1. A chromosomal disruption of BPT1 was generated in the ade2 haploid strains YPH499 (here designated ABC470 and ABC500) by using PCR-mediated gene disruption with the KanMX module (47) to replace codons 428 to 736 of Bpt1p, yielding ABC791 and ABC790, respectively. A double mutant, ABC794 (ycf1::KanMX2 bpt1::KanMX2), was constructed by dissecting tetrads derived from a diploid generated by a cross between ABC470 and ABC790. Double-disruption segregants were found to be viable. The presence of both disruptions was confirmed by PCR.

Single and double disruptions of YCF1 and BPT1 in the ade2 diploid strain SM1060 (30) were also generated. Chromosomal YCF1 was disrupted from codon 1 to 1439 by the standard two-step integration-excision method using plasmid pJAW53, as previously described (50). The diploid was sporulated and the tetrads were dissected to obtain both disruptions was confirmed by PCR.

A strain containing chromosomal BPT1-GFP was generated by transforming Smal-digested pSM1489 into strain SM3851 (bpt1::KanMX4-HIS4-GFP) and selecting for Ura" transformants. One of the transformants, SM4787, which contains BPT1-GFP, was generated by the standard two-step integration-excision method using ArH2-digested pSM1817 into strain SM3851 (bpt1::KanMX4-HIS4-GFP) and selecting for Ura" transformants. The resulting strain, SM4643, contains YCF1-GFP in the YCF1 locus. Correct integration of YCF1-GFP was confirmed by Southern blot analysis, and the presence of YCF1-GFP was confirmed by Western blot analysis.

Plasmids and their construction. Plasmids used in this study are listed in Table 2. Plasmid pSM1487 (CEN Ira3 BPT1) contains the complete BPT1 ORF flanked by its native 5' upstream and 3' downstream regions (515 and 500 bp, respectively) and was constructed in two steps. First, the 5' upstream and 3' downstream regions of BPT1 were amplified from the genome by PCR and subcloned by three-way ligation into pRS316 (CEN URA3) (42). Next, the BPT1 ORF was cloned into this vector by ligating it from the genome (32). The BPT1 coding sequence in the resulting plasmid, pSM1487 (CEN Ira3 BPT1), was confirmed by DNA sequence analysis. The multicopy plasmid pSM1488 (CEN Ira3 BPT1) contains a 5.7-kb KpmI-NolI fragment from pSM1487 subcloned into the same sites of pSM217 (2μm Ira3) (6). A GFP-tagged version of BPT1 was constructed by fusing the GFP coding sequence (F64L, S65T) immediately before the stop codon of BPT1 by PCR-mediated recombinational cloning (32).
to create pSM1489 (CEN URA3 BPT1-GFP). To generate a multicopy version of this construct, a 6.5-kb KpnI-Nol fragment from pSM1489 was subcloned into the same sites of pSM217, generating pSM1490 (2 μm URA3 BPT1-GFP). A GFP-tagged version of YCFL was generated by subcloning a 6.0-kb KpnI-SacI fragment from pJAW53 (50) into KpnI-SacI-digested pSM217. Next, the coding sequence for GFP (pQBI25; Quantum Biotechnologies, Inc., Montreal, Quebec, Canada) was fused immediately before the stop codon of YCFL by PCR-mediated recombinational cloning (32) to yield pSM1753 (2 μm URA3 YCFL-GFP). To generate an integrating plasmid expressing BPT1-GFP, we constructed pSM1849 by subcloning an 8.2-kb PvuI fragment from pSM1490 into the same sites of pRS306 (42). The YCFL ORF was amplified by PCR product containing 448 bases upstream of the YCFL lacZ gene at the BamHI site in pLG699-Z. The sequences of the XhoI-HindIII-digested pSM217. Next, the coding sequence from pSM1753 and 1,092 bases of the region downstream of the YCFL ORF amplified from pJAW53 (50) subcloned into pRS306 (42).

To construct the BPT1 promoter-reporter fusion plasmid pBPT1 (2 μm URA3 BPT1-lacZ), a purified PCR product containing 448 bases upstream of the BPT1 ORF was subcloned into the Smal site of pHBluescript (Stratagene, La Jolla, Calif.). Next, an XhoI-BamHI fragment was excised and cloned into the BamHI-XhoI sites of pLG699-Z (2 μm URA3 lacZ) (16), yielding pBPT1. The YCFL reporter-fusion plasmid pBPT1 was constructed by amplifying 418 bases upstream of the YCFL ORF and subcloning this purified PCR product into the XhoI-BamHI sites of pLG699-Z. The YCFL promoter differed from the published sequence at position –302, containing an insertion of C at this position. The same insertion was observed in clones from completely independent and separate PCRs and is therefore not an artifact of PCR. The primers for both plasmids were designed to enable in-frame fusions with the LacZ gene at the BamHI site in pLG699-Z. The sequences of the oligonucleotides used for the strain and plasmid constructions will be furnished upon request.

Media and growth conditions. Cells were routinely grown at 30°C in complete (yeast extract-peptone-dextrose [YPD]) medium or minimal drop-out medium (i.e., synthetic complete medium without Ura [SC-Ura]) as previously described (yeast extract-peptone-dextrose [YPD]) medium or minimal drop-out medium). Yeast cells were grown to log phase in selective medium, harvested, washed, resuspended in Z buffer, and assayed for fluorescence (HQ Endow GFP) and Nomarski optics (Zeiss, Thornwood, N.Y.). Images were determined by liquid scintillation counting. Nonenergized (MgATP-independent) uptake was estimated by the same procedure, except that ATP was omitted from the uptake medium.

Transport substrates. Radiolabeled 2,4-dinitrophenyl-glutathione ([3H]DNP-GS) (55 mCi/mmol) was purchased from DuPont, NEN.

Transport assays. For in vitro transport measurements, vacuoles were isolated by the ficoll flotation technique from cells grown in YPD medium and were vesiculated at a 1:50 dilution into the same medium, and grown overnight at 30°C to an OD600 of approximately 1.0. This overnight culture was diluted to an OD600 of 0.1, which in turn was diluted in 10-fold increments. Five microliters of each 10-fold dilution was spotted onto SC-Ura plates containing different concentrations of CdSO4 and incubated for 3 days at 30°C.

Transport substrates. Radiolabeled 2,4-dinitrophenyl-glutathione ([H3]DNP-GS) (418.7 mCi/mmol) was synthesized and purified as described previously (24).

To test the transport activities of Ycf1p in vivo and in vitro are well established, cells were grown to an OD600 ranging between 2.9 and 3.2 in YPD medium containing 0.5% yeast extract. An equal number of cells from each strain (as determined by OD600) was harvested and washed with sterile water. Cells were lysed in 5% sulfosalicylic acid by using glass beads, and cell debris was removed by centrifugation at 15 K for 5 min. The amount of the pigment was estimated by recording the absorbance at 530 nm (kmax). Control extracts were prepared in a similar manner by using strain ABC591 (gsh1Δ) (41) grown in the presence of excess adenine (5).

Growth inhibition by cadmium. To examine growth in liquid medium, cells grown to saturation in YPD medium were subcultured at a starting OD600 of ~0.1 into YPD medium containing different concentrations of CdSO4, and the extent of growth after 30 h was determined by measuring the OD600. The growth rate of each culture was also determined by taking OD600 readings at multiple intervals throughout the entire 30-h period. In all cases, an alteration in the extent of growth after 30 h is reflected as a corresponding alteration in the rate of growth (data not shown).

To examine growth on plates, cells were grown overnight to saturation in minimal medium, and subcultured at a 1:50 dilution into the same medium, and grown overnight at 30°C to an OD600 of approximately 1.0. This overnight culture was diluted to an OD600 of 0.1, which in turn was diluted in 10-fold increments. Five microliters of each 10-fold dilution was spotted onto SC-Ura plates containing different concentrations of CdSO4 and incubated for 3 days at 30°C.

Media and growth conditions. Cells were routinely grown at 30°C in complete (yeast extract-peptone-dextrose [YPD]) medium or minimal drop-out medium (i.e., synthetic complete medium without Ura [SC-Ura]) as previously described (30). A PD medium containing different concentrations of CdCl2, was prepared by addition of the required amount of CdCl2 immediately prior to the growth experiment. For pigment quantitation, the liquid YPD medium was limiting for adenine and contained 0.5% yeast extract rather than 1% yeast extract. The minimal drop-out plates used for the pigmentation test in Fig. 2 contained a limited amount of adenine (7.5 mg per liter). The plates used for the cadmium spot test were prepared by adding the indicated concentration of CdSO4 to the minimal plate medium.

Fluorescence microscopy. Log-phase cells expressing GFP-tagged Bpt1p and GFP-tagged Ycf1p were examined at ×100 magnification on a polystyrene-coated slide by using a Zeiss Axioplan 2 microscope equipped with fluorescence (HQ Endow GFP) and Nomarski optics (Zeiss, Thornwood, N.Y.). Images were captured with a Cooke charge-coupled device camera and IP Lab Spectrum software (Biowision Technologies, Inc., Exton, Pa.).

β-Galactosidase reporter assays. Saccharomyces cerevisiae strain 512100 (wild type [WT]) or SM13 ( yap1Δ-Δ2) cells were singly transformed with the β-galactosidase reporter plasmids YEp55 and YEp55-Δ2 or doubly transformed with one of these reporter plasmids and Yep55-Δ2 (50), which overproduces Yap1p. The WT strain ABC154 (ade2Δ) was also transformed with the reporter plasmids to compare the transcriptional regulation of YCF1 and BPT1 in the presence of CdCl2, or a limited amount of adenine. Transformants were grown to log phase in selective medium, harvested, washed, resuspended in Z buffer, and assayed for β-galactosidase activity (38). Measurements were carried out in triplicate, and activity was expressed as β-galactosidase units per OD600 (optical density at 600 nm) unit of cells.

Quantitation of the red pigment that accumulates in an ade2 strain. To quantitate the extent of pigment formation during growth of ade2 strains on adenine-limiting medium, cells were grown to an OD600 ranging between 2.9 and 3.2 in YPD medium containing 0.5% yeast extract. An equal number of cells from each strain (as determined by OD600) was harvested and washed with sterile water. Cells were lysed in 5% sulfosalicylic acid by using glass beads, and cell debris was removed by centrifugation at 15 K for 5 min. The amount of the pigment was estimated by recording the absorbance at 530 nm (kmax). Control extracts were prepared in a similar manner by using strain ABC591 (gsh1Δ) (41) grown in the presence of excess adenine (5).

Growth inhibition by cadmium. To examine growth in liquid medium, cells grown to saturation in YPD medium were subcultured at a starting OD600 of ~0.1 into YPD medium containing different concentrations of CdSO4, and the extent of growth after 30 h was determined by measuring the OD600. The growth rate of each culture was also determined by taking OD600 readings at multiple intervals throughout the entire 30-h period. In all cases, an alteration in the extent of growth after 30 h is reflected as a corresponding alteration in the rate of growth (data not shown).

To examine growth on plates, cells were grown overnight to saturation in minimal medium, and subcultured at a 1:50 dilution into the same medium, and grown overnight at 30°C to an OD600 of approximately 1.0. This overnight culture was diluted to an OD600 of 0.1, which in turn was diluted in 10-fold increments. Five microliters of each 10-fold dilution was spotted onto SC-Ura plates containing different concentrations of CdSO4 and incubated for 3 days at 30°C.

TABLE 2. Plasmids used in this study

<table>
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<th>Plasmid</th>
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<tr>
<td>pABB51</td>
<td>2μm URA3 P_{BPT1-lacZ}</td>
<td>This study</td>
</tr>
<tr>
<td>pABB52</td>
<td>2μm URA3 P_{YCF1-lacZ}</td>
<td>This study</td>
</tr>
<tr>
<td>pJAW53</td>
<td>YEp 5'-1.018 nt-YCF1-hisG-URA3-hisG-3'1.356 nt-YCF1</td>
<td>50</td>
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<tr>
<td>pSM217</td>
<td>2μm URA3</td>
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<tr>
<td>pSM1753</td>
<td>2μm URA3 YCF1-GFP</td>
<td>This study</td>
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<tr>
<td>pSM1817</td>
<td>YEp URA3 YCF1-GFP</td>
<td>This study</td>
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<tr>
<td>pSM217</td>
<td>YEp URA3 BPT1-GFP</td>
<td>This study</td>
</tr>
<tr>
<td>YEp55-Δ2</td>
<td>2μm LEU2 YAPI</td>
<td>50</td>
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* Yip denotes a yeast integrating plasmid; nt, nucleotides.
A. Bpt1p-GFP

<table>
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<td>GFP</td>
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B. Ycf1p-GFP

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FIG. 1. Bpt1p, like Ycf1p, localizes to the vacuolar membrane. The GFP fluorescence pattern of cells expressing Bpt1p-GFP (A) or Ycf1p-GFP (B) from the chromosome (panels 1 and 3) or from a multicopy plasmid (panels 5 and 7) is shown. In the corresponding Nomarski (DIC) images of these cells (A and B, panels 2, 4, 6, and 8), the vacuole appears as an indentation. Strains used are SM4787 (A, panels 1 to 4), SM3794/pSM1490 (A, panels 5 to 8), SM4643 (B, panels 1 to 4), and SM3851/pSM1753 (B, panels 5 to 8).

established (23, 25). In this study we analyzed Bpt1p, the closest yeast homologue of Ycf1p. Bpt1p shares strong overall homology with Ycf1p (42% identity and 59% similarity) as well as with human MRP1 (38% identity and 57% similarity) (9). It possesses a domain organization typical of most members of the MRP subclass of ABC transporters, consisting of two membrane-spanning domains (MSDs), each with six spans, and two nucleotide binding domains (NBDs). Notably, Bpt1p and Ycf1p also contain the N-terminal extension that is the hallmark of a subset of the MRP subfamily members and consists of an additional membrane-spanning domain (MSD0) with five spans, followed by a linker (L0) region (3).

Ycf1p localizes to the vacuolar membrane, as shown by immunofluorescence and subcellular fractionation (23, 49). To localize Bpt1p, we generated a Bpt1p-GFP fusion construct, as well as its Ycf1p-GFP counterpart. The latter is fully functional, as measured by its ability to confer cadmium resistance on a ycf1Δ strain (data not shown). Fluorescence and Nomarski microscopy were carried out for Bpt1p-GFP and Ycf1p-GFP expressed from either the chromosome or a multicopy plasmid. Bpt1p-GFP (Fig. 1A), like Ycf1p-GFP (Fig. 1B), exhibits a clear-cut vacuolar membrane localization, as can be seen by the correspondence of the vacuole indentation observed by differential interference contrast (DIC) and GFP fluorescence patterns. The brighter areas of fluorescence appear to correspond to regions where the multiple vacuolar lobes intersect. The grainy fluorescence pattern in cells expressing single-copy versus multicopy Bpt1p-GFP and Ycf1p-GFP is due to the long exposure required to enhance the dim signal for the former. Based on the results above, Bpt1p, like Ycf1p, would therefore be expected to be an ABC transporter that delivers its cargo into the vacuole lumen.

Bpt1p contributes to the transport of the GS-conjugated compound responsible for the red pigmentation of an ade2 strain in vivo. Since Bpt1p and Ycf1p reside in the same membrane and share significant sequence identity, we were interested to determine if they have overlapping transport functions. As a starting point for these analyses, we generated a bpt1 null mutant by homologous gene replacement using the KanMX2 cassette to disrupt the BPT1 ORF in both MATa and MATα haploid strains. Growth of the bpt1 null mutant is indistinguishable from that of the isogenic WT strain (data not shown), indicating that BPT1 is not an essential gene under normal growth conditions. Likewise, the growth of a double null mutant (ycf1Δ bpt1Δ) generated by a cross between the single mutants is similar to that of the WT, although occasionally a slight growth impairment could be observed with this strain (for example, see Table 3, growth in 0 μM CdCl2). YCF1 has been shown to play an important role in the red pigmentation of adenine biosynthetic mutants in S. cerevisiae (5). Specifically, strains carrying ade1 or ade2 mutations accumulate an intense red pigment in their vacuoles when they are grown under conditions of adenine limitation. The formation of this pigment involves a series of steps, including conjugation of GSH to the adenine biosynthetic intermediate phosphoribosylaminomimidazole (AIR), transport of AIR-GS conjugates into the vacuole in a Ycf1p-dependent manner, and further metabolism of the pigment precursor after its vacuolar sequestration. In a ycf1 null mutant, pigmentation of an ade2 mutant is diminished but not completely eliminated (5). Presumably, residual pigmentation occurs because another cellular transporter(s) can mediate the transport of AIR-GS conjugates.

To examine whether the residual pigmentation of the ade2 ycf1Δ strain involves Bpt1p, we quantitated the extent of pigment formation in strains that had single and double disruptions of BPT1 and YCF1. Quantitation was performed as described in Materials and Methods, and data are averages of four experiments ± standard deviations. Pigment accumulation, expressed as a percentage of that in the WT strain ABC154, was 96% ± 8.4% for strain ABC791 (bpt1Δ), 31% ± 4.3% for strain ABC470 (ycf1Δ), and 15% ± 6.8% for strain ABC794 (bpt1Δ ycf1Δ). Thus, as expected, pigment accumulation was substantially decreased in the ycf1Δ mutant (to 31% of the WT level). Notably, the deletion of bpt1 in a ycf1Δ strain background resulted in a further decrease in pigmentation (to 15% of the WT level in the ycf1Δ bpt1Δ double mutant), clearly implicating Bpt1p in vacuolar pigment sequestration. However, the accumulation of pigment in the bpt1 null mutant was not significantly different from that in the WT strain (96 versus 100%, respectively), presumably because the contribution of Ycf1p overwhelmed that of Bpt1p. Thus, while Bpt1p is not a major player in ade2 pigmentation when Ycf1p is present, it does indeed appear to contribute to this process, as evidenced by the finding that when the Ycf1p transporter is absent, a role for Bpt1p is clearly revealed. From a genetic standpoint, the
ycf1Δ and bpt1Δ mutants exhibit “synthetic enhancement” of the ade2 phenotype (15); the phenotype of the double mutant is markedly more severe than that of either of the single mutants.

The results of plate tests of the capacity of the BPT1 gene expressed from a multicopy plasmid to restore ade2 pigmentation in the ycf1Δ bpt1Δ double mutant are consistent with these conclusions. Whereas the ycf1Δ bpt1Δ strain containing an empty vector formed white to pale-pink colonies (Fig. 2A), the ycf1Δ bpt1Δ strain harboring a BPT1 plasmid formed dark-pink colonies (Fig. 2C), with a pigmentation intensity comparable to that of the double-mutant strain harboring a YCF1 plasmid (Fig. 2B). Importantly, although this result is not quantitative, it suggests that Bpt1p on its own, like Ycf1p, has the capacity to function as a transporter for the GS-conjugated pigment precursor.

Bpt1p contributes to cadmium detoxification, as evidenced by synthetic enhancement of cadmium sensitivity in a ycf1Δ bpt1Δ double mutant. Ycf1p contributes to cadmium detoxification in yeast by mediating the transport of Cd-GS complexes into the vacuole (25). Compared to an isogenic WT (YCF1) strain, a ycf1Δ mutant is hypersensitive to cadmium salts (44). To determine whether Bpt1p also plays a role in cadmium detoxification, we compared the growth of a WT strain with that of strains that had single and double disruptions of BPT1 and YCF1, both in liquid media and on plates containing different concentrations of cadmium.

For liquid growth experiments, saturated overnight cultures of each strain were subcultured into YPD medium containing 0, 50, or 100 μM CdCl₂ and were propagated for 30 h. The extent of growth was determined by measuring the OD₆₀₀ after 30 h (Table 3). All of the strains, including the WT, are subject to growth retardation by 50 or 100 μM CdCl₂. As reported previously, growth of the ycf1Δ strain was markedly retarded, here attaining only 55 and 36% of the OD₆₀₀ of the WT at 50 and 100 μM CdCl₂, respectively, confirming that Ycf1p is important for cadmium resistance. In contrast, growth of the bpt1Δ strain was unaffected; this mutant strain attained 100% of the OD₆₀₀ of the WT at 50 and 100 μM CdCl₂, indicating that Bpt1p is dispensable for cadmium detoxification when Ycf1p is functional. However, a discernible contribution of Bpt1p to cadmium detoxification was revealed in the double (ycf1Δ bpt1Δ) mutant, whose growth was significantly diminished in CdCl₂ relative to that of the ycf1Δ single mutant (36% versus 55% in 50 μM CdCl₂ and 21% versus 36% in 100 μM CdCl₂). Augmentation of the ycf1Δ cadmium-sensitive phenotype through the abolition of Bpt1p function is consistent with a synthetic enhancement, similar to that observed for the Ade2 phenotype. Bpt1p can contribute to cadmium detoxification in yeast, presumably by mediating the transport of Cd-GS complexes into the vacuole, albeit at modest levels compared to that of Ycf1p.

The results of plate tests yielded the same basic conclusion, regardless of the background strain in which the single and double mutants were generated. Results with strains derived from YPH499 (42) (here designated ABC154) and EG123 (43)
membranes from wild-type, single-mutant, and double-mutant strains, respectively. A culture of each strain was diluted to an OD600 of 0.1, and serial dilutions were spotted onto plates without (0 mM CdSO4), with (10 and 30 mM CdSO4), and with (10 and 30 mM CdSO4) cadmium. In the absence of cadmium, the growth of all strains is essentially equivalent (Fig. 3, A and B). However, a role for Bpt1p is clearly revealed by the double-mutant strain at 10 mM cadmium, the growth of all strains is essentially equivalent (Fig. 3, A and B). The strains used for the preparation of vacuolar vesicles are ABC154 (wild type), ABC470 (ycf1Δ), ABC791 (bpt1Δ), and ABC794 (ycf1Δ bpt1Δ).

**FIG. 4.** MgATP-dependent uptake of [3H]DNP-GS, [3H]E217βG, and [3H]GSH by vacuolar membrane vesicles from WT and mutant strains. For the large graphs in panels A, B, and C, MgATP-dependent uptake of the indicated transport substrates was measured for 10 min at the concentrations of substrate indicated, in the presence or absence of 3 mM ATP. The level of uptake at each substrate concentration was calculated by subtracting the amount of radioactivity detected in the absence of ATP from that detected when ATP was present. The data were fitted to the Michaelis-Menten equation by nonlinear least-squares analysis. The values obtained for each substrate and each strain are as follows: (A) For the WT, Km_{DNP-GS} = 5.8 ± 2.3 μM and V_{max(DNP-GS)} = 13.5 ± 0.2 nmol/mg/10 min; for the bpt1Δ strain, Km_{DNP-GS} = 23.2 ± 3.8 μM and V_{max(DNP-GS)} = 2.3 ± 0.4 nmol/mg/10 min. (B) For the WT, Km_{E217βG} = 268.4 ± 157 μM and V_{max(E217βG)} = 76.3 ± 6.0 nmol/mg/10 min; for the ycf1Δ strain, Km_{E217βG} = 35.3 ± 3.3 μM and V_{max(E217βG)} = 76.3 ± 6.0 nmol/mg/10 min; for the ycf1Δ bpt1Δ double mutant, Km_{E217βG} = 3.8 ± 0.4 mM and V_{max(E217βG)} = 149.4 ± 76.3 nmol/mg/10 min; for the ycf1Δ strain, Km_{GSH} = 3.8 ± 0.4 mM and V_{max(GSH)} = 87.5 ± 43.3 nmol/mg/10 min. Kinetic parameters were not calculated for DNP-GS uptake by the ycf1Δ bpt1Δ strain or for E217βG and GSH uptake by the ycf1Δ and ycf1Δ bpt1Δ mutants, because the data did not approximate the Michaelis-Menten equation. Individual data points are means ± standard errors (n = 3 to 6). The insets in panels A and B show the time courses for DNP-GS and E217βG uptake, respectively. The strains used for the preparation of vacuolar vesicles are ABC154 (wild type), ABC470 (ycf1Δ), ABC791 (bpt1Δ), and ABC794 (ycf1Δ bpt1Δ).

**Bpt1p plays a role in the in vitro transport of GS conjugates, glucuronides, and GSH into vacuole-derived vesicles.** The in vivo tests described above indicate that Bpt1p function partially overlaps that of Ycf1p. To compare Bpt1p- and Ycf1p-dependent transport biochemically, we isolated yeast vacuolar membranes from wild-type, single-mutant, and double-mutant strains and examined their capacities to mediate the MgATP-energized transport of three canonical MRP substrates: the GSH conjugate DNP-GS, the glucuronate conjugate E217βG, and reduced GSH. For all of these substrates, it was found that the transport capabilities of Bpt1p overlapped with those of Ycf1p to varying degrees; for two of the three substrates, Bpt1p appeared to require Ycf1p for maximal activity.

Vesicular DNP-GS uptake was the first transport function to be assayed, because this model GS conjugate is firmly established as a substrate for Ycf1p. When assayed for time dependence at a concentration of 10 μM, DNP-GS uptake by WT vesicles was 18.8 nmol of substrate in 5 min (Fig. 4A, inset). Uptake by vesicles from a bpt1Δ single mutant was 13.5 nmol of substrate in 5 min, representing a 28% decrease from the transport capacity of the WT and indicating a modest but observable contribution of Bpt1p to MgATP-dependent transport of this conjugate. By contrast, the amount of DNP-GS uptake by vesicles from a ycf1Δ single mutant and from a ycf1Δ bpt1Δ double mutant was <0.2 nmol of substrate in 5 min,
representing a $>95\%$ decrease in transport capacity compared to that of the WT.

These effects were similarly evident in the concentration dependence of DNP-GS uptake (Fig. 4A). In a WT strain, the bulk of the transport measured is attributable to Ycf1p, since vesicles from a ycf1Δ strain are markedly decreased in their transport efficacy compared to vesicles from a WT strain. The $K_m$ for the WT versus ycf1Δ vesicles is $5.8 \pm 2.3$ versus 268.4 ± 157.0 μM; the corresponding $V_{max}$ values are 30.7 ± 2.3 versus 20.3 ± 7.8 nmol/mg/10 min, respectively (Fig. 4A). In contrast to YCF1, the effects of disrupting the BPT1 gene were modest: $K_m$ was negligibly affected relative to that of the WT (6.3 ± 3.8 versus 5.8 ± 2.3 μM) and $V_{max}$ was only slightly attenuated (25.3 ± 3.0 versus 30.7 ± 2.3 nmol/mg/10 min) (Fig. 4A). As expected if Ycf1p mediates the bulk of DNP-GS transport, the difference between the concentration-dependent uptake by ycf1Δ single-mutant membranes and that by ycf1Δ bpt1Δ double-mutant membranes was small (Fig. 4A).

By comparison with that of DNP-GS, transport of the other substrates, the glucuronate conjugate E17βG (Fig. 4B) and reduced GSH (Fig. 4C), followed a different pattern. In both cases, the relative contribution made by Bpt1p was greater, implying that that this transporter has a preference for E17βG and GSH over DNP-GS. Vesicles from the bpt1Δ strain show a 59% decrease in the capacity for uptake of 100 μM E17βG (1.8 nmol of substrate in 5 min versus a WT value of 4.4 nmol of substrate in 5 min) (Fig. 4B, inset). However, unexpectedly, the contributions of Bpt1p and Ycf1p are not additive, as a decrease in the amount of E17βG uptake in the ycf1Δ single mutant was $>95\%$, not 41%. This low level of uptake was similar to that of the ycf1Δ bpt1Δ double mutant (Fig. 4B).

A pattern comparable to that seen for E17βG was also observed for GSH. Disruption of the BPT1 gene diminished the total capacity of the membranes for transport by more than 40%, whereas uptake of GSH was negligible throughout the concentration range examined not only in the membrane vesicles from ycf1Δ bpt1Δ double-mutant cells but also in those from ycf1Δ single-mutant cells (Fig. 4C).

**BPT1 and YCF1 are differentially regulated transcriptionally.** Yap1p is a transcription factor that modulates the expression of a large number of yeast genes implicated in the response to oxidative stress, for instance, stress resultant from exposure to cadmium (31, 50). Since YCF1 is under the regulatory control of Yap1p, it was of interest to determine whether BPT1 expression shares this characteristic. To compare the influence of Yap1p on the expression patterns of YCF1 and BPT1, we generated constructs in which the 5′ upstream regions of YCF1 and BPT1 were fused to the coding region of lacZ. These reporter plasmids were transformed into WT, yap1Δ-defective, or YAP1-overexpressing strains that were subsequently assayed for β-galactosidase activity (Table 4).

Two features are evident from these expression studies. First, the basal levels of YCF1 and BPT1 expression in a WT strain are essentially the same (Table 4, rows 1 and 6), suggesting that in the absence of inducer these transporters are present at similar levels. Second, while the expression of YCF1 is strongly modulated by Yap1p, BPT1 expression is only marginally affected if at all. Mutational loss of Yap1p activity results in an approximately 3-fold decrease in YCF1 expression. Conversely, Yap1p overexpression leads to an approximately 10-fold increase in YCF1 expression (Table 4, compare rows 2 and 3 to row 1). This is in complete agreement with the results of previous studies (50). In strict contrast, BPT1 expression was not affected by a lack of Yap1p and was only modestly increased (∼1.5-fold) by an excess of Yap1p.

Expression of the fusion constructs was also examined under conditions of adenine deficiency and excess in order to assess the effects of endogenous AIR-GS conjugates on YCF1 and BPT1 expression. Limiting levels of adenine (0 μg/ml) elicited an approximately 5-fold increase in $P_{YCF1-lacZ}$ expression over that in cells subjected to excess adenine (60 μg/ml), but an increase of only 2.5-fold in $P_{BPT1-lacZ}$ expression (Table 4, compare rows 5 and 4). Analogous experiments employing cadmium at 0 and 50 μM as an inducer increased the expression of $P_{YCF1-lacZ}$ and $P_{BPT1-lacZ}$ by approximately 2.5- and 1.5-fold, respectively (Table 4, compare rows 6 and 7). The strong induction of YCF1 observed under conditions of high cadmium and limiting adenine may be partly or wholly mediated through Yap1p. The modest transcriptional effects seen for Bpt1p may also be mediated by Yap1p, or instead by another, as yet undetermined transcription factor.

**DISCUSSION**

Bpt1p and Ycf1p are localized to the vacuolar membrane and show overlapping transport properties in vivo. In this study we compared the localization, expression, and substrate transport properties of Bpt1p and its close homologue Ycf1p in order to gain more insight into the MRP subfamily of ABC transporters in yeast. The sequence and structural similarities between Bpt1p and Ycf1p suggest that they might have similar functions in vivo. To initiate studies of Bpt1p, we compared the localization of C-terminally GFP-tagged Ycf1p and Bpt1p in vivo. Fluorescence microscopy revealed that the localization of Bpt1p-GFP is indeed vacuolar, like that of Ycf1p-GFP (Fig. 1). These findings corroborate and extend the results of previous in vitro experiments showing that vacuole-enriched membrane preparations, which are at best only partially pure, mediate Bpt1p-dependent transport activity (34, 37).

The vacuolar localization of Bpt1p is consistent with the notion that it, like Ycf1p, contributes to the vacuolar sequestration of otherwise toxic compounds. Several possibilities can be envisioned for how these transporters might function in the

<table>
<thead>
<tr>
<th>Trial no.</th>
<th>Straina or substrate concentration</th>
<th>$P_{YCF1-lacZ}$</th>
<th>$P_{BPT1-lacZ}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>SEY6210 (WT)</td>
<td>2.53 ± 0.12</td>
<td>2.37 ± 0.12</td>
</tr>
<tr>
<td>2</td>
<td>SM13 (yap1Δ)</td>
<td>0.85 ± 0.07</td>
<td>2.17 ± 0.08</td>
</tr>
<tr>
<td>3</td>
<td>SM13 (yap1Δ YEp351-YAP1)</td>
<td>22.93 ± 0.85</td>
<td>3.26 ± 0.07</td>
</tr>
<tr>
<td>4</td>
<td>60 mg of adenine/liter</td>
<td>3.14 ± 0.15</td>
<td>2.22 ± 0.11</td>
</tr>
<tr>
<td>5</td>
<td>0 mg of adenine/liter</td>
<td>15.76 ± 0.7</td>
<td>5.40 ± 0.36</td>
</tr>
<tr>
<td>6</td>
<td>0 μM CdCl₂</td>
<td>2.72 ± 0.45</td>
<td>2.34 ± 0.34</td>
</tr>
<tr>
<td>7</td>
<td>50 μM CdCl₂</td>
<td>6.72 ± 0.84</td>
<td>3.70 ± 0.47</td>
</tr>
</tbody>
</table>

a Strain ABC154 (ade2) was used for each trial unless otherwise specified.

b Measured as described in Materials and Methods and expressed as β-galactosidase units per OD₆₀₀ unit of cells.
same intracellular environment. For example, Bpt1p and Ycf1p could have identical substrate transport and regulatory properties, and so function redundantly. Alternatively, these transporters could have a distinct pattern of substrate selectivity and regulation, and function complementarily. Finally, a combination of both of these schemes could apply, such that Bpt1p and Ycf1p might exhibit functional and regulatory overlap under some circumstances and unique properties under others. As detailed below, the latter combined view seems to best explain the properties of Bpt1p and Ycf1p both in vivo and in vitro.

To directly compare the contributions of Bpt1p and Ycf1p in vivo, we generated bpt1Δ and ycf1Δ (single) and ycf1Δ bpt1Δ (double) mutants. Neither YCF1 nor BPT1 is essential for viability. We used these mutants to examine two well-characterized phenotypes known to involve Ycf1p: ade2 pigmentation and cadmium resistance. These substrates were considered to be particularly appropriate because they share the property of being conjugated to, or complexed with, GSH yet are either of endogenous (the biosynthetic intermediate AIR) or exogenous (cadmium) origin. We show here that Bpt1p, like Ycf1p, plays a role in ade2 pigmentation and cadmium detoxification (Table 3; Fig. 2 and 3). While Ycf1p accounted for most of the transport measured for both substrates, Bpt1p made a minor but demonstrable contribution. Specifically, in the ycf1Δ single mutant, ade2 pigmentation was strongly reduced (by 70%) compared to that in the WT, and growth in the presence of Cd²⁺ was strongly inhibited (by 64%) at 100 μM CdCl₂ (Table 3). Although the bpt1Δ mutation alone did not result in a defect in ade2 pigmentation or cadmium resistance, a function for Bpt1p was clearly evident in the ycf1Δ bpt1Δ double mutant, whose phenotype was significantly more severe than that of the ycf1Δ single mutant. Both ade2 pigmentation accumulation and growth in 100 μM CdCl₂ (Table 3) were reduced by a further 15% in the ycf1Δ bpt1Δ double mutant versus the ycf1Δ single mutant.

The phenomenon observed here for Bpt1p, in which the single bpt1Δ and ycf1Δ mutants show no phenotype, or only a modest one, but the double mutant is severely impaired, can be referred to as “synthetic enhancement” (15). Generally, genes which exhibit synthetic enhancement are intimately connected functionally, either as interacting components or, in the case of deletion mutations, more likely as functionally redundant proteins. With Bpt1p and Ycf1p, the latter mode of synthetic enhancement is the more likely. Thus, Ycf1p alone (i.e., in a bpt1Δ mutant) is sufficient for full transport of the substrates tested in vivo here. In the absence of Ycf1p (i.e., in a ycf1Δ mutant), there is still residual function. This must be due to Bpt1p, since in the ycf1Δ bpt1Δ double mutant, transport is decreased even further. Independent corroboration of the ability of Bpt1p to function as a transporter is provided by the finding that its overexpression in the ycf1Δ bpt1Δ double mutant rescues the ade2 pigmentation defect (Fig. 2).

**BPT1 and YCF1 are differentially regulated.** The smaller contribution of Bpt1p, compared to Ycf1p, to the transport of in vivo substrates could be due solely to the distinct substrate preferences of these transporters. Alternatively, or additionally, the lower contribution of Bpt1p could reflect a lower relative level of expression than that of Ycf1p. The possibility of differentially regulated expression of YCF1 versus BPT1 is supported by promoter fusion studies (Table 4). YCF1 expression is exquisitely sensitive to the dosage of Yap1p, a transcription factor involved in responses to oxidative stress. P_{YCF1-lacZ} expression is decreased 3-fold in the absence of Yap1p and is increased 10-fold when Yap1p is overexpressed. In contrast, P_{BPT1-lacZ} expression exhibits no, or at most very little, dependence on Yap1p; it is insensitive to the absence of Yap1p and only modestly elevated (~1.5-fold) by Yap1p overexpression. Since cadmium resistance involves regulatory effects exerted through Yap1p, it is not surprising that P_{YCF1-lacZ} expression is increased to a greater extent than that of P_{BPT1-lacZ} in 50 μM Cd (~2.5–versus 1.5-fold, respectively [Table 4]). It is established that the strong Cd²⁺-induced increase in YCF1 expression is indeed dependent on Yap1p (50), but it is not known if the modest increase in BPT1 expression elicited by Cd²⁺ is mediated by Yap1p or by another transcription factor.

By analogy with the results obtained with Cd²⁺, we find that P_{YCF1-lacZ} expression is substantially induced (fivefold) under conditions of adenine limitation, in which the ade2 gene expression mediated through Yap1p, it is not surprising that P_{BPT1-lacZ} expression is increased to a greater extent than that of P_{BPT1-lacZ} in 50 μM Cd (~2.5–versus 1.5-fold, respectively [Table 4]). It is established that the strong Cd²⁺-induced increase in YCF1 expression is indeed dependent on Yap1p (50), but it is not known if the modest increase in BPT1 expression elicited by Cd²⁺ is mediated by Yap1p or by another transcription factor.

Extrapolating from our gene expression data in Table 4, it is likely that at the level of protein expression Ycf1p and Bpt1p are roughly equivalent under normal conditions. However, the relative amounts of Ycf1p may exceed those of Bpt1p when cells are treated with Cd²⁺. Therefore, Ycf1p should make proportionately the greatest contribution to Cd²⁺ resistance in WT cells, masking the contribution of Bpt1p. Thus, differences in BPT1 and YCF1 gene expression mediated through Yap1p and potentially other transcription factors may be largely responsible for the distinct phenotypic contributions of Bpt1p and Ycf1p in vivo. Clearly, the ability of these transporters to be differentially regulated expands the range of environmental or endogenous insults which yeast cells can tolerate.

**Bpt1p and Ycf1p exhibit distinct substrate preferences in in vitro transport assays.** We were interested in comparing the transport properties of Bpt1p and Ycf1p for canonical MRP substrates in vitro. To do so, we isolated a vacuolar membrane-enriched fraction from WT, single-mutant, and double-mutant cells and determined the kinetics of transport of a model GS transporter (Fig. 4) reflects the basal levels of Bpt1p and Ycf1p, since the cells used for preparation of the vacuole-enriched fraction were grown in the absence of inducer. From the results of these in vitro measurements, it is clear that Bpt1p and Ycf1p overlap in their transport capabilities but exhibit distinct substrate preference profiles. On the one hand, Bpt1p appears to favor glucuronate conjugates and free GSH over GS conjugates (Fig. 4). On the other hand, vesicles from a ycf1Δ mutant are severely (>95%) impaired for the transport of all three substrates, suggesting that Ycf1p plays a major and essential role in the transport of all three substrates (see below).

Although for the substrates tested here, ycf1Δ vesicles were less effective in transport than bpt1Δ vesicles, this pattern ap-
pears not to be universal for all substrates. Two recent studies examined the substrate transport properties for unconjugated bile pigment (UCB) (37) and several MRI contrast reagents (Gd-EOB-DTPA, Gd-BOPTA, Gd-B 20790, and Gd-B 21690) (34), comparing vesicles from WT and mutant strains. Taken together, the results from our study and those previous studies show that Ycf1p is the major (>50%) contributor to the transport of UCB, Gd-BOPTA, DNP-GS, E178G, and GSH. In contrast, Bpt1p and Ycf1p contribute approximately equally to Gd-B 20790 transport, and Bpt1p is the major contributor to Gd-EOB-DTPA transport. This pattern of overlapping but nonidentical substrate preference for yeast MRPs is reminiscent of what has been documented for the mammalian MRPs to MRPs of different proteins (20, 22) and the vascular GS conjugate pumps AMR1 and AMR2 from plants (26–28), suggesting that this is a recurrent theme within the MRP subfamily in all organisms.

A notable difference between the in vitro transport of some substrates (i.e., Bpt1p) and that of others (i.e., UCB) (37) is the lack of additivity of Bpt1p and Ycf1p activities. For example, since we observed a 59% diminution of E178G transport in bpt1Δ vacuolar membrane vesicles, we expected that the equivalent fraction from ycf1Δ cells would show a reduction of ≤41%. However, unexpectedly, what we saw instead was a diminution of >95% in the ycf1Δ single mutant strain and no further reduction of transport activity in the ycf1Δ bpt1Δ double-mutant strain. Thus, Bpt1p appears to contribute to the transport of E178G (Fig. 4B), and similarly GSH (Fig. 4C), only in the presence of Ycf1p. How might this surprising dependence be explained? One model to account for these findings is that Bpt1p and Ycf1p may interact combinatorially to form a heterodimer. For the three substrates tested here, such a Bpt1p-Ycf1p heterodimer would have optimal transport activity, while Bpt1p alone or as a homodimer is inactive and Ycf1p alone or as a homodimer has intermediate activity. This model is appealing not only because it is consistent with the present data but also in light of a recent structural study of MRPs where Ycf1p has a predominant and dimeric structure (39). An alternative model is that another component which is lacking or down-regulated in both the ycf1Δ and ycf1Δ bpt1Δ strains is necessary for maximal Bpt1p activity.

MRPs in yeast. Comparison of the yeast MRPs Bpt1p and Ycf1p highlights the point that despite their sequence similarity, MRPs can exhibit biologically important differences, including differential substrate specificity and regulatory properties. There are four additional genes that encode MRPs in yeast: YBT1 (first published as BAT1), YOR1, YHL035, and YKR103/YKR104 (12, 45). Sequence alignments suggest that all of these contain a full or partial N-terminal extension. Ybt1p is a vacuolar membrane protein that can transport bile acid, but apparently not GS-conjugated compounds (33), and Yor1p is a plasma membrane protein that plays a role in oligomycin resistance (13, 19). The products of the remaining two genes, YHL035 and YKR103/YKR104, remain uncharacterized. It is noteworthy that the latter actually encodes two adjacent reading frames separated by a nonsense codon, likely representing a sequencing error or a mutation in the strain used to sequence the yeast genome. It is clear from the present study that developing a complete picture of detoxification and related processes in yeast will require an understanding of the distinct localization patterns, expression profiles, and substrate transport properties of all of the yeast MRPs.

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K. G. Sharma and D. L. Mason contributed equally to this work.

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