Mutations Define Cross-talk between the N-terminal Nucleotide-binding Domain and Transmembrane Helix-2 of the Yeast Multidrug Transporter Pdr5

**POSSIBLE CONSERVATION OF A SIGNALING INTERFACE FOR COUPLING ATP HYDROLYSIS TO DRUG TRANSPORT**

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The yeast Pdr5 multidrug transporter is an important member of the ATP-binding cassette superfamily of proteins. We describe a novel mutation (S558Y) in transmembrane helix 2 of Pdr5 identified in a screen for suppressors that eliminated Pdr5-mediated cycloheximide hyper-resistance. Nucleotides as well as transport substrates bind to the mutant Pdr5 with an affinity comparable with that for wild-type Pdr5. Wild-type and mutant Pdr5s show ATPase activity with comparable $K_m$(ATP) values. Nonetheless, drug sensitivity is equivalent in the mutant pdr5 and the pdr5 deletion. Finally, the transport substrate clotrimazole, which is a non-competitive inhibitor of Pdr5 ATPase activity, has a minimal effect on ATP hydrolysis by the S558Y mutant. These results suggest that the drug sensitivity of the mutant Pdr5 is attributable to the uncoupling of NTPase activity and transport. We screened for amino acid alterations in the nucleotide-binding domains that would reverse the phenotypic effect of the S558Y mutation. A second-site mutation, N242K, located between the Walker A and signature motifs of the N-terminal nucleotide-binding domain, restores significant function. This region of the nucleotide-binding domain interacts with the transmembrane domains via the intracellular loop-1 (which connects transmembrane helices 2 and 3) in the crystal structure of Sav1866, a bacterial ATP-binding cassette drug transporter. These structural studies were supported by biochemical and genetic evidence presented here that interactions between transmembrane helix 2 and the nucleotide-binding domain, via the intracellular loop-1, may define at least part of the translocation pathway for coupling ATP hydrolysis to drug transport.

Multidrug transporters, including members of the ATP-binding cassette (ABC) family, show unusual flexibility toward their substrate cargo. They efflux structurally diverse xenobiotic compounds and confer broad-spectrum hyperresistance when they are overexpressed, a property that impedes chemotherapeutic treatment of pathogens and cancer.

Several fungi, including the clinically relevant human pathogenic species *Candida albicans* and *Cryptococcus neoformans*, contain major multidrug transporters that are close homologues to the *Saccharomyces cerevisiae* transporter Pdr5 (1). It is well established that many clinical isolates of *C. albicans* overproduce the Pdr5 homologue Cdr1 (2). Multidrug-resistant fungi are an increasing problem in the treatment of immunocompromised patients with AIDS and cancer (3).

The Pdr5 subfamily of multidrug transporters shows significant differences in molecular architecture from their mammalian counterparts such as P-glycoprotein (P-gp) or Mrp1. The nucleotide-binding domains (NBDs) of these important fungal transporters precede the transmembrane domains (TMDs) and thus their orientation is the reverse of P-gps and Mrp1s. Furthermore, the Walker A, B, and signature motifs are degenerate when compared with nonfungal ABC counterparts. For example, a cysteine replaces a lysine in Walker A (Cys-199) of the N-terminal NBD (NBD1) that is essential in other ABC transporters (1).

Not surprisingly, the ATPase activities of Pdr5 and *C. albicans* Cdr1p appear to be similar (4–7). For example, both have high basal (unstimulated) levels. Recently, we demonstrated that Pdr5 also hydrolyzes GTP at physiological concentrations and that this activity is considerably more resistant to some substrates, such as clotrimazole (clo), than is its ATPase activity. Furthermore, GTP fuels Pdr5-mediated chloramphenicol efflux in purified vesicles (5). These results suggest that Pdr5 has great flexibility not only in the drug cargo that it transports but also in its efflux energy source.

Considerable progress has been made in recent decades in understanding the domain organization of ABC transporters and the mechanism of the transport cycle (8). A typical ABC protein consists of four domains, two TMDs and two NBDs. The TMDs contain the drug-binding sites, which show little differences in molecular architecture from their mammalian counterparts such as P-glycoprotein (P-gp) or Mrp1. The nucleotide-binding domains (NBDs) of these important fungal transporters precede the transmembrane domains (TMDs) and thus their orientation is the reverse of P-gps and Mrp1s. Furthermore, the Walker A, B, and signature motifs are degenerate when compared with nonfungal ABC counterparts. For example, a cysteine replaces a lysine in Walker A (Cys-199) of the N-terminal NBD (NBD1) that is essential in other ABC transporters (1).
sequence similarity in different ABC transporters. By contrast, the NBĐs exhibit very high sequence similarity.

The activity of an ABC transporter hinges on the effective coupling of the ATP catalytic cycle, which occurs at the NBĐs, and the drug-transport cycle, which occurs at the TMDs. Mutational analyses, labeling with the photoaffinity drug or nucleotide analogues, and cross-linking with the thiol-reactive substrate derivatives have yielded a wealth of information about interactions at the drug-binding domains as well as about the role of conserved residues in the NBĐs. Biochemical and genetic studies suggested the existence of a transmission interface that couples the energy of ATP binding and/or hydrolysis to conformational changes at the TMDs (9). These studies have taken indirect approaches, however, such as the determining the effect of drugs on ATP hydrolysis.

The recent elucidation of the x-ray structure of Sav1866, a bacterial ABC transporter (10), has permitted the delineation of such a transmission interface in more precise detail. A novel feature of this structure is that both NBĐs contact the TMDs. These contacts occur via intracellular loops (ICLs) that link the transmembrane helices. The organization of the structure of this domain suggests a plausible explanation for the mechanical coupling between the NBĐs and TMDs of ABC transporters. In addition, recent studies with P-gp and Yor1 used site-directed mutagenesis and cross-linking agents to demonstrate the proximity of ICL2 and the Q-loop (11, 12). These studies show or imply a physical interface between the NBDs and TMDs of ABC transporters. In addition, the use of site-directed mutagenesis. This powerful approach can also be applied to any mammalian transporter that is expressed in yeast and has a readily observable phenotype.

**EXPERIMENTAL PROCEDURES**

*Strains and Genetic Transformation*—The strains used in this study are listed in Table 1. All are isogenic to each other and derived from AD1–7, which lacks all PM ABC transporters but contains a PDR1–3 allele that results in overexpression of PDR5 when a plasmid containing this gene is integrated into its chromosomal location. The construction of JG2000 and JG2001 was described previously (5).

We also constructed several new AD1–7 derivatives. The first series of strains carry two copies of PDR5 either as wild type (WT; JG2004) or mutant (JG2010). These strains produce significantly more Pdr5 than does the popular single-copy strain AD124567 and are thus invaluable for biochemical studies. JG2004 was created by transforming JG2001 with pSS607, the plasmid containing the WT gene PDR5.

The JG2009 strain contains a single copy of the S558Y allele and was used to create JG2010. A PDR5-bearing strain (JG2001) was transformed with S558Y to create JG2003. Treatment with 5-fluoroorotic acid (5-FOA) (14) yielded segregants (9/22). These studies show or imply a physical interaction, but none establish that these regions are involved in communication between the TMDs and NBĐs. Demonstrating the functional relevance of such interactions remains a challenge.

Here we describe the behavior of an S558Y mutation in transmembrane helix 2 (TMH2) that retains significant ATPase activity and drug-binding capability but nevertheless tests as phenotypically null for drug resistance and transport. Furthermore, an allosteric inhibitory signal mediated by the Pdr5-specific transport substrate clotrimazole is greatly diminished. These deficiencies are significantly alleviated by a suppressor mutation that lies near the Q-loop of NBĐ1. This is a particularly important observation because recent work suggests that the noncanonical NBĐ1 of Pdr5 may not play a catalytic role in ATP hydrolysis (13).

Our results are thus consistent with the x-ray crystallographic structure of Sav1866 and suggest that even evolutionarily distant ABC transporters have a similar coupling interface. Moreover, our strategy of using suppressor mutations to identify the interacting residues in the TMDs and NBĐs has obvious advantages over the use of site-directed mutagenesis. This powerful approach can also be applied to any mammalian transporter that is expressed in yeast and has a readily observable phenotype.

**TABLE 1**

**Yeast strains used in this study**

All the strains are isogenic derivatives of AD1–7, with the exception of JG365-3B.

| Strain       | Relevant genotype                                                                 | Source or Ref. |
|--------------|----------------------------------------------------------------------------------|----------------|
| JG365-3B     | Mat a PDR5 PDR1-3 ura3 leu2                                                        | 35             |
| AD1–7        | MATa PDR1-3 ura3 his3 yer1 pdr5 arg2 pdr10 ycf1 pdr11 pdr3                         | 5              |
| JG2000       | AD1–7 + pSS607                                                                    | This study     |
| JG2001       | ura3, PDR5 5-FOA derivative of JG2000                                             | 5              |
| JG2001       | JG2001 + pSS558Y                                                                  | This study     |
| JG2004       | JG2001 + pSS607                                                                   | This study     |
| JG2005       | AD1–7 + pSS558Y                                                                  | This study     |
| JG2009       | ura3, S558Y derivative of JG2003                                                   | This study     |
| JG2010       | JG2009 + pSS558Y                                                                  | This study     |
| R-1          | Δpdr5:KANMX4 replaces PDR5 in JG2001                                              | This study     |
| JG2011       | R-1 + pSS558Y                                                                    | This study     |
| JG2012       | R-1 + pSS558A                                                                    | This study     |
| JG2015       | R-1 + pSS607 (wild-type control)                                                  | This study     |
| JG2016       | R-1 + pN242K                                                                     | This study     |
| JG2020       | clo-3-0 + pSS558Y, N242K                                                         | This study     |
| JG2021       | R-1 + pSS558Y, N242K                                                             | This study     |

**Source or Ref.**

- 35: JG2001
- 36: JG2001
- 5: JG2001
- This study: JG2001

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We also constructed several new AD1–7 derivatives. The first series of strains carry two copies of PDR5 either as wild type (WT; JG2004) or mutant (JG2010). These strains produce significantly more Pdr5 than does the popular single-copy strain AD124567 and are thus invaluable for biochemical studies. JG2004 was created by transforming JG2001 with pSS607, the plasmid containing the WT gene PDR5.

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Interactions between TM Helix-2 and NBD-1 of Pdr5

All of these new isogenic strains are available from the corresponding author. We used a Gietz transformation kit (Medi-
corp, Montreal, Quebec, Canada) to introduce plasmids and PCR-generated products into yeast.

Plasmids and Site-directed Mutagenesis—We created all site-
directed mutants from the plasmid pSS607, which was described previously (5). To obtain pSS58Y, pSS58A, and 
Pn424K, we carried out PCR-based site-directed mutagenesis 
with the Quick Change kit (Stratagene, La Jolla, CA). PCR primers 
carrying the desired base pair substitution were prepared, 
and PAGE was purified by Operon Biotechnologies (Huntsville, 
AL). The resulting mutants were confirmed by DNA sequenc-
ing the entire PDR5 insert. Sequencing was performed by Ret-
rogen (San Diego, CA). We constructed the 14 sequencing 
primers so that the resulting sequencing data would contain 
substrant overlapping segments, thus ensuring accuracy.

Chemicals and Media—All chemicals were purchased from Sigma. We dissolved cycloheximide (cyc) in sterile water and all other compounds in dimethyl sulfoxide. We purchased 
8-azido-[α-32P]ATP (15–20 Ci/mmol) from Affinity Labeling 
Technologies, Inc. (Lexington, KY). We obtained [125I]iodoary-
lazidoprazosin ([125I]IAAP), 2,200 Ci/mmol, from PerkinElmer 
Life Sciences.

We cultured cells in yeast extract, peptone, glucose (YPD) 
medium. Cyh, tritylimidazole, and clo were added to YPD after 
autoclaving. Tetrabutyltin was added to yeast extract, peptone, 
glycerol (YPG) medium. After sterilization, 5-FOA (1 g/liter) 
avoclaved. Tetrabutyltin was added to yeast extract, peptone, 
glycerol (YPG) medium. After sterilization, 5-FOA (1 g/liter) 
was used to add synthetic dextrose complete medium.

Determining IC50 Values in Liquid Culture—We grew cul-
tures of yeast strains to be tested for hyporesponsiveness over-
night in 5 ml of YPD broth in 15-ml clinical centrifuge tubes at 
30 °C in a shaking water bath. Then cells were pelleted in 
a clinical centrifuge at high speed for 5 min. We decanted the 
growth medium and washed the pellets with 10 ml of sterile 
water before resuspension. We determined cellular concentra-
tion by spectroscopy (absorbance at 600 nm). We removed 
0.5 × 10⁶ cells, added this to 2 ml of YPD broth containing cyc 
(0.18–43.2 μM) or tritylimidazole (2–20 μM), and incubated 
the cultures at 30 °C with shaking. After 48 h, we monitored 
relative inhibition by determining the absorbance at 600 nm 
and comparing measurements to cells grown in YPD alone.

Qualitative Analysis of Relative Drug Resistance—We tested 
muteant and control strains qualitatively for drug resistance by 
spotting 2-fold dilutions of cells in 5 μl of water on plates with a 
fixed concentration of a xenobiotic compound. Cultures to be 
tested were grown overnight in 10 ml of YPD broth. The cultures 
were washed and concentrated by centrifugation and resuspended in sterile water at 1/10th the original culture vol-
ume. We diluted a small sample 10-fold and determined the 
absorbance at 600 nm to check cell concentration. We made 
2-fold dilutions in sterile water before plating in a 5-μl sample.

UV Mutagenesis of JG365-3B—The first S558Y allele was 
created by UV mutagenesis as described previously (17).

Estimating the Frequency of Reversion—To estimate the rate 
of reversion of S558Y, 10 1-ml YPD cultures were seeded 
with 100 cells and grown to saturation. We plated 10⁶ cells in 4-ml sterile agar on plates containing 7.5 μM clotrimazole and incu-
bated them for 5 days at 30 °C. We counted the suppressors and 
picked them for further testing.

Preparation of Purified Membrane Vesicles—We prepared plasma membrane (PM) vesicles as described previously (18), 
with further modification (5). We determined protein concen-
tration in the vesicles with a bicinchonic acid kit (Perbio, Rock-
land, IL).

Immunoblots of Pdr5 in PM Vesicles—Proteins in the purified 
PM vesicles were separated by gel electrophoresis and trans-
ferred to nitrocellulose membrane at constant current (400 
ma, 1 h). The nitrocellulose membranes were blocked for 30 
min with 20% nonfat milk in PBS containing Triton X-100 
(PBST); incubated with a 1:200 dilution of polyclonal goat, anti-
Pdr5 antibody, and yC-18 (Santa Cruz Biotechnology, Santa Cruz, 
CA); washed with PBST (three times for 15 min); incubated with a 
1:5,000 dilution of secondary antibody (donkey anti-goat IgG 
horseradish peroxidase in 5% nonfat milk) for 2 h; and washed with 
PBST (three times for 15 min). The chemiluminescence signal was 
developed by adding 10 ml of ECL reagent for 10 min and then 
exposing to x-ray film for 0.5–1 min.

Assay of ATPase Activity—ATPase activity of purified PM ves-
icles was measured by the end point P₆ assay as described previ-
ously for P-gp (19, 20), with minor modifications for Pdr5 (5).

One concern about ATPase measurements made with PM preparations is that these contain Pma1p [H⁺], which exhibits 
vanadate (Vi)-sensitive ATPase activity. We described our resolu-
tion of this issue in detail in a previous study (5). Briefly, the optimal 
ph for Pma1p [H⁺] is ~5.5–6.0, and the enzyme is only min-
imal active at pH 7.5. We therefore characterized ATPase activity in PMs from the Pdr5 at pH 7.5. In addition, measure-
ments were made with purified PM vesicles from both Pdr5+ 
strains and the Δpdr5 strains, and the small amount of residual 
ATPase activity (<5%) observed in Δpdr5 strains was subtracted.

Rhodamine 6G (R6G) Transport Assay—We measured R6G 
transport at 30 °C as described previously (21). The concentra-
tion of R6G used in this study was 2.5 μM.

Cross-linking of 8-Azido-[α-32P]ATP to Pdr5—we carried 
out binding studies in purified PM vesicles at 4 °C. Membrane 
proteins (18 μg/assay) were suspended in MOPS ATPase assay 
buffer containing 10 μM 8-azido-[α-32P]ATP (2.5 μCi/nmol). 
Samples were incubated in the dark for 5 min at 4 °C before 
cross-linking with 365 nm UV light on ice for 10 min. Reactions 
were stopped by adding 12.5 μl of 5× SDS-PAGE sample-loading 
buffer. After electrophoresis on a Tris acetate gel at constant 
voltage, gels were dried. We quantified the radioactivity 
corporated into the Pdr5 band with a STORM 860 Phosphor-
Imager system (GE Healthcare) and ImageQuant software.

Cross-linking of [125I]IAAP to Pdr5—Purified membranes 
prepared from yeast cells overexpressing Pdr5 were incubated 
at room temperature in the ATPase buffer with [125I]IAAP (7 nM) 
for 5 min under subdued light. The samples were photocross-linked 
for 10 min with 365 nm UV light at room temperature followed by 
electrophoresis with 7% NuPAGE gels and quantified as described 
previously (21). Modifications to this procedure in specific exper-
iments are described in the figure legends.

DNA Extraction and PCR Recovery of Suppressor Mutant 
Sequences—we amplified PDR5 from suppressor mutants with 
a PureLink Genomic DNA kit from Invitrogen. The DNA was
FIGURE 1. Drug sensitivity of S558Y. A, 2-fold serial dilutions of Pdr5 strains were plated on YPD media or on YPD media containing 15 μM clotrimazole, triethylimazol, or tetrabutyltin. The number of cells plated ranged from 1.0 × 10^7 to 0.5 × 10^7; scans of plates that were incubated for 96 h at 30 °C are shown. A representative figure from three independent experiments is shown. B, quantitative analysis of drug sensitivity was carried out by inoculating 500 cells bearing the WT and mutant Pdr5s in the presence of increasing concentrations of cyh (0.05–21.6 μM). The cells were incubated at 30 °C for 48 h and then quantified by measuring the absorbance at 600 nm. The strains used were R-1 (Δpdr5), R-1 + pS5607 (WT = ), and R-1 + pS558Y (Δ). Growth at each concentration of cycloheximide is shown as (% control), i.e. growth in the absence of cyh. C, same protocol was used to compare single- and double-copy mutant strains: single-copy WT (JG2000 = ), single-copy S558Y (R-1 + pS558Y = ); double-copy S558Y (JG2010 = A), and R-1 ( ). The figures (B and C) show the mean values of three independent experiments, and the error bars represent the S.D.

extracted as described in the accompanying handbook from Invitrogen, with several modifications. We lysed 4 × 10^7 cells for each extraction. We purchased zymolyase made up in buffer from Sigma. We used 1 ml (~60 units of activity) for each sample. We also doubled the volume of proteinase K and RNase A.

We carried out PCR amplification with Platinum PCR Supermix High Fidelity from Invitrogen, with the following protocol: 1 cycle at 95 °C for 5 min, 35 cycles of denaturation at 95 °C for 1 min, annealing at 55 °C for 1 min, and extension at 68 °C for 7 min. Reactions were then placed on hold at 4 °C until they were processed further. The primers used to amplify PDR5 were left, TAAGACTTCCGTTGAGTTG (Tm of 59 °C), and right, GCACGTTCTGTACTTCCA (Tm of 60 °C).

RESULTS

S558Y Mutation—The S558Y allele was identified in the course of a large mutant search designed to uncover suppressors that eliminated the cyh resistance of a strain overproducing Pdr5 (JG365-3B), the major yeast multidrug transporter. Following UV mutagenesis, we screened ~70,000 survivors and isolated 17 cyh-sensitive colonies that failed to complement a known pdr5 deletion. Of these, 16 were severely impaired and phenotypically similar to the deletion; a single very leaky allele was also recovered.

Immunoblotting of purified PM vesicles indicated that in 15 of these 17 mutants, Pdr5 was significantly reduced or missing (data not shown). One such mutant, pdr5-107 was the subject of recent study (17). Another mutant, although severely drug-sensitive, had unaltered levels of Pdr5 in its PM. Complete nucleotide sequencing of the PDR5 gene indicated that it contained an S558Y missense mutation. The reported TMHs of Pdr5 suggested that this residue lie just outside TMH2. The original TMH assignments, however, were slightly different from those that we have established from seven topology modeling programs that allow us to predict core consensus sequences. Results from all these programs place Ser-558 in TMH2.3

Drug Phenotypes of S558Y and S558A—The S558Y mutation was recreated in pSS607, a yeast integrative vector that we recently described (5). We also made an S558A mutation. The mutant plasmids were completely sequenced to verify that each had only the desired alteration. From these plasmids we created several strains used in this study; all are isogenic to each other. Fig. 1A shows the relative clo sensitivity of four strains as follows: R-1 (pdr5::KANMX4); R-1 + pSS607, referred to as WT; R-1 + pS558Y or S558Y; and R-1 + pS558A or S558A.

These data indicate that the S558Y mutation is drug hypersensitive as we predicted. By contrast, the S558A allele is phenotypically similar to the WT PDR5. This result suggests that it is the size of the tyrosine residue rather than the hydrophobicity of its aromatic ring that is primarily responsible for the observed mutant phenotype. A similar phenotype is observed with tritylimidazole and tetrabutyltin. These substrates were used to define different transport sites in Pdr5 that may or may not overlap (22, 23). In each case, S558Y is considerably more sensitive than the WT control to all of these xenobiotic agents and is qualitatively the same as the Δpdr5::KANMX4 isogenic negative control.

3 R. Rutledge and D. Xia, manuscript in preparation.
Several studies have indicated that the drug-binding sites of ABC transporters consist of a large pocket, and individual transport substrates show specific interactions with different amino acid side chains (24–26). The fact that the S558Y mutant is not substrate-specific (Fig. 1A) suggests that the reversal of drug resistance is a consequence of general steric hindrance and that the Ser-588 residue may not be involved in specific interactions with the drugs. This is consistent with the finding that the mutation S558A does not affect Pdr5 function.

We carried out a quantitative analysis of relative cyh resistance in liquid culture. The data in Fig. 1B show that the S558Y mutant has an IC_{50} value that is ~30-fold lower (0.5 μM) than the WT and is similar to the Δpdr5 control.

Fig. 1C illustrates the behavior of double- and single-copy strains at low cyh concentrations. Strains containing 1 (JG2009) or 2 copies of S558Y show similar drug hypersensitivity. Therefore, doubling the amount of mutant protein does not increase cyh resistance. This is significant because the increased levels of Pdr5 in the double-copy strains are useful in biochemical assays, where they provide more robust measurements.

**S558Y Mutant Strain Is Transport-deficient**—We analyzed the transport capability of S558Y with a well established, whole-cell R6G efflux assay that uses flow cytometry (21, 27). We compared the transport capability of the WT (JG2000), Δpdr5 (AD1–7), and S558Y (JG2009) isogenic strains. These data are found in Table 2, and representative histogram plots are shown in Fig. 2. These data clearly demonstrate that the S558Y mutation creates an R6G transport deficiency that is as severe as that observed in the Δpdr5 strain. Both of these strains retain ~60–70-fold more R6G than does the WT PDR5 strain, JG2000.

**S558Y Shows Normal Membrane Localization and Nucleotide Binding and Slightly Diminished ATP Hydrolysis**—We evaluated the effect of adding a second copy of WT or mutant protein to a haploid strain. The colloidal blue-stained gel and the immunoblots (Fig. 3A) clearly show a 2-fold increase in Pdr5 levels in WT (JG2004) and S558Y mutant (JG2010) compared with the isogenic, single-copy counterpart (JG2000). Thus, although the single-copy strain overproduces Pdr5, it is clear that more protein can be localized to the membrane by simply raising the copy number. We therefore used the PMs prepared from these strains for all biochemical studies.

**TABLE 2**

| Strain and pertinent genotype | Median fluorescence | S.D. |
|------------------------------|--------------------|------|
| AD1–7 (Δpdr5)                | 408.21             | 96.94|
| JG2005 (AD1–7 + pSS558Y)    | 389.47             | 52.94|
| JG2000 (AD1–7 + pSS607)     | 671.17             | 0.59 |

**FIGURE 2.** R6G transport is drastically diminished in S558Y. The transport activity of WT and mutant Pdr5s was estimated with a flow cytometry-based efflux assay with R6G as the fluorescent substrate. The null strain (Δpdr5 = AD1–7) and yeast strains containing the WT (JG2000) and mutant (JG2009) Pdr5 were used in these experiments. The assay was carried out as described previously (21), and the representative histograms from four independent experiments are depicted. The strains used are labeled on the figure.

**FIGURE 3.** Interaction of nucleotides with WT and S558Y. A, a colloidal blue staining and immunoblotting of purified PM vesicles prepared from various strains of yeast. Lanes 1–4 depict a colloidal blue-stained gel and lanes 5–8 an immunoblot with the anti-Pdr5 antibody yC-18. Lanes 1 and 5, Δpdr5; lanes 2 and 6, WT Pdr5, single copy; lanes 3 and 7, WT Pdr5, double copy; lanes 4 and 8, mutant Pdr5, S558Y, double copy. Arrow shows the position of the Pdr5 protein. B, photopaffinity labeling of Pdr5 with 0.5 mM 8-azido-[α-32P]ATP. Cross-linking was carried out in purified plasma membrane vesicles at 4 °C as described under “Experimental Procedures.” Upper panel, 8-azido-[α-32P]ATP incorporated into the Pdr5 band; lower panel, quantification of the [32P]P incorporated with a STORM 860 Phospholmager system. The cross-linking in the absence of ATP is shown as filled bars and the cross-linking the presence of ATP as empty bars. C, 8-azido-[α-32P]ATP binding as a function of ATP concentration (0–1 mM) was determined in WT Pdr5 (■) and the S558Y mutant (□). The 8-azido-[α-32P]ATP incorporated in the Pdr5-bands was quantified with a Phospholmager, and the curves were generated with the curve-fitting Prism GraphPad software (GraphPad Software, La Jolla, CA). The inset depicts the data with an expanded x axis in the range 0–0.125 mM ATP. The x and y axis legends for the inset are the same as those for the main figure. Figure shows representative data from two independent experiments.
Interactions between TM Helix-2 and NBD-1 of Pdr5

PMs prepared from WT and S558Y Pdr5-expressing strains show comparable levels of photolabeling with 8-azido-[α-32P]ATP (Fig. 3B). Furthermore, because the [32P] signal is completely eliminated by the addition of 2 mM ATP, the interaction is clearly taking place at the NBDs.

It is of course possible that although nucleotides bind to both WT and mutant Pdr5s, the affinity is altered. The concentration-dependent displacement of 8-azido-[α-32P]ATP by ATP provides a measure of the apparent affinity of ATP for the transporter. We found that the IC_{50} values for the WT and mutant Pdr5 are almost identical (Fig. 3C). As expected, the Δpdr5 strain shows no significant cross-linking of 8-azido-[α-32P]ATP even though the colloidal blue-stained gels clearly indicate that all three lanes were loaded with equal amounts of vesicle protein (data not shown). These results strongly suggest that the WT and S558Y strains have similar nucleotide-binding capabilities.

The kinetics of Pdr5-mediated ATP hydrolysis was the subject of several previous studies, and it is clear that this transporter has a high level of unstimulated (basal) activity (4, 5, 13). We compared the kinetics of ATP hydrolysis in WT Pdr5 and in S558Y. We found an ~2-fold decrease in the V_{max} of ATP hydrolysis. Several vesicle preparations from each strain were assayed. The V_{max} for the WT ranged from 180 to 279 nmol/mg/min; the V_{max} for the S558Y mutant ranged from 69 to 93 nmol/mg/min (representative data from one preparation is shown in Fig. 4A).

Two aspects of these data are notable. Although the WT and mutant Pdr5s differ in V_{max}, they show comparable K_{m} values (about 0.5 mM). This is consistent with the finding that the IC_{50} for displacement of 8-azido-[α-32P]ATP by ATP is identical in WT and S558Y (Fig. 3C). Furthermore, the ATPase activity of the double-copy S558Y mutant Pdr5 is comparable with that found in the single-copy WT protein (5). Nevertheless, the single-copy WT strain is ~30-fold more resistant to cyh than is the double-copy S558Y strain (see Fig. 1B above). Therefore, the decrease in ATPase activity observed in the double-copy mutant strain alone cannot explain why the mutant Pdr5 is profoundly hypersensitive to all of the tested drugs (Fig. 1A).

Pdr5-ATPase activity is sensitive to inhibition by its transport substrate clo, which acts as a noncompetitive inhibitor at an allosteric site (5). When WT Pdr5 was treated with 25 μM clo, the V_{max}(ATP) decreased by ~90% (Fig. 4B). Although clo also reduces the V_{max}(ATP) for Pdr5-mediated ATP hydrolysis in S558Y, the inhibition is only partial (40%; Fig. 4C). These results suggest two possibilities as follows: either the drug substrate site itself is modified, or the allosteric inhibition of ATPase activity cannot be efficiently communicated to the NBDs so that a significant amount of the ATPase activity is allowed to proceed.

8-IAAP Binding in the S558Y Mutant—To understand whether it is the interaction between drug substrate(s) and Pdr5 per se that is affected in the S558Y mutant, we exploited the photoaffinity labeling of IAAP. We previously demonstrated that IAAP, an [125I]-labeled photoaffinity analogue of prazosin, cross-links to the transport substrate sites of Pdr5 (21). We further demonstrated that IAAP cross-links to both WT and S558Y Pdr5 but not to PMs prepared from the Δpdr5 strain (AD1–7) (Fig. 5A). Moreover, in the presence of clo, cross-
linking of IAAP is significantly reduced in both WT and mutant Pdr5 (Fig. 5A).

To determine the apparent affinities of WT and mutant Pdr5, we measured the displacement of cross-linked IAAP in the presence of increasing concentrations of clortrimazole (Fig. 5B). We found that the IC_{50} values for the displacement of IAAP by clo are comparable in WT and mutant Pdr5 (13.8 and 12.6 \mu M, respectively).

Taken together, our results demonstrate that both the drug substrate and nucleotide bind to the S558Y mutant with affinities that are comparable with WT Pdr5. However, the drug efflux mediated by the S558Y mutant is severely impaired and as a consequence exhibits drug sensitivity.

Genetic Analysis of S558Y Suppressors—Our results suggest that the S558Y mutation in Pdr5 disrupts the coupling between the NBDs and the transport substrate site. We isolated clo-resistant (clo^r) suppressors of the S558Y mutation present in JG2011 (R-1 + pS558Y) by plating cultures on medium containing 15 \mu M clo. We analyzed 15 independent cultures and determined a frequency of 2.2 \times 10^{-5}. This is a relatively high value for a true back mutation (Tyr \rightarrow Ser) and thus suggested that at least some of the suppressors might represent mutations at other sites in PDR5 or conceivably in other genes.

Because suppressors generally have dominant phenotypes, complementation testing cannot be used to determine whether a mutation resides in PDR5. The R-1 strain was built to quickly plaque the mutant plasmid sequence containing URA3 and other PDR5 sequences. These events result in Ura^- colonies that are selected on 5-FOA medium (14) and are tested on clotrimazole and geneticin medium. If the mutation is attributable to a second mutation in the S558Y copy, clo^r, gen^r, and clo^r, gen^r colonies will be recovered. A mutation in another gene would yield only clo^r colonies.

To determine whether the clo resistance is in the PDR5 gene, we employed 5-FOA selection (14). This drug selects for Ura^- colonies. If the reversion is attributable to a second mutation in the S558Y copy, clo^r, gen^r, and clo^r, gen^r colonies would be recovered. A mutation in another gene would yield only clo^r colonies.

Ten-fold serial dilutions of cells were prepared and spotted as described under “Experimental Procedures.” Plates were scored at 96 h after incubation at 30 °C.

| Strain and pertinent genotype | Clo, 15.0 \mu M | Clo, 22.5 \mu M | Cyh, 0.78 \mu M |
|------------------------------|----------------|----------------|----------------|
| WT (R-1 + pS6607)            | 4              | 0              | 0              |
| S558Y (R-1 + pS558Y)         | 0              | 0              | 0              |
| Clo-1-0                      | 4              | 0              | 0              |
| Clo-3-0                      | 3              | 2              | 3              |

To determine whether the clo resistance is in the PDR5 gene, we employed 5-FOA selection (14). This drug selects for Ura^- colonies. If the reversion is attributable to a second mutation in the S558Y allele (possibly bearing a second mutation causing the clo^r phenotype) or the disrupted/deleted (KANMX4) copy. The loss of the latter leaves a geneticin-sensitive S558Y segregant that is presumably clo-resistant. Loss of the former, however, would make the colony geneticin-resistant but clo-sensitive if the clo resistance is caused by a second mutation in the S558Y copy of the gene. If the clortrimazole resistance was not caused by a genetic change in the S558Y copy, all of the 5-FOA-selected recombinants would remain clo-resistant.

Each of 12 independently derived suppressors was plated on 5-FOA, and ura3 recombinant survivors arose at a frequency of \sim 2 \times 10^{-3}. 5-FOA survivors were tested for each suppressor to determine whether the clo^r mutation was in PDR5. Of 31 5-FOA survivors recovered from clo-1-0, 13 were clo^r, Gen^r and 18 were clo^r, Gen^r. Similarly, of 25, 5-FOA survivors recovered from clo-3-0, 14 were clo^r, Gen^r and 11 were clo^r, Gen^r. These results confirm that each suppressor phenotype was attributable to a second mutation in the S558Y allele. Comparable results were obtained for the remaining 10 suppressors (data not shown).

The clo-1-0 and clo-3-0 suppressors were spot-tested on two different Pdr5 substrates, clo (15.0 and 22.5 \mu M) and cyh (0.78 \mu M). A summary of these results is given in Table 3. The numerical score indicates the lowest concentration of cells showing confluent growth on the particular drug plate. Thus, a score of 1 means the suppressor grew only at the highest cellular concentration, 0.5 \times 10^{-3}; a score of 4 means it grew at the lowest concentration, 0.5 \times 10^{-6}. The JG2011 strain used to isolate suppressors failed to grow at any cellular concentration and thus received a 0.

The data in Table 3 suggest that the clo-1-0 suppressor mutant is as drug-resistant as the WT control. The strong but not completely WT phenotype of the clo-3-0 suppressor suggested it might be particularly useful in further biochemical studies, especially in distinguishing various models of S558Y dysfunction, and was therefore analyzed further. The other suppressors are the subject of continuing study.

We compared the clo-3-0 and clo-1-0 suppressor mutants and controls for relative resistance to triethylamine, a substrate that defines a transport site that is at least somewhat different from the one used by clo and R6G (21). These results are shown in Fig. 7A. As with R6G transport, the clo-1-0 suppressor is indistinguishable from WT positive control, but the clo-3-0 suppressor increases resistance to a level of the drug.
between the levels of the S558Y and WT strains. Similarly, flow cytometry data demonstrated that R6G transport is restored to a level that is also between that of the S558Y and WT strains (Fig. 7B). In addition, neither the initial drug hypersensitivity of S558Y nor the restored resistance of clo-3-0 shows any indication of substrate specificity. The clo-1-0 suppressor exhibited transport properties that were indistinguishable from the WT (data not shown).

The PDR5-specific DNA was recovered from the clo-1-0 and clo-3-0 suppressors by PCR amplification, and the entire coding region was sequenced. The sequence recovered from clo-1-0 contains a true back mutation from Tyr back to Ser. The completely WT phenotype it exhibits confirms the pleiotropic nature of the original mutation (deficiencies in transport and drug resistance).

In the clo-3-0 suppressor, the original S558Y base pair substitution remains, but a second alteration, N242K was identified. The S558Y,N242K double mutation was confirmed in two separate sequencing reactions from overlapping primers. N242K is located in NBD1 near the Q-loop between the Walker A and the signature motifs. No other alterations were observed.

We constructed the double mutation in pSS607 to create pS558Y,N242K. We also made the single N242K mutation in this plasmid. We demonstrated that transformation of R-1 with a pS558Y,N242K plasmid recreates a clo-3-0 phenotype when it is used to transform the R-1 strain (pdr5::KANMX4; Fig. 7C).

**FIGURE 7.** The clo-3-0 (S558Y,N242K) suppressor reverses the drug-sensitive phenotype of S558Y. A, quantitative analysis of clo-3-0 growth in the presence of trityl was carried out as described under “Experimental Procedures.” Growth of WT (■), S558Y (●), clo-1-0 (○), and clo-3-0 (▲) in the presence of increasing concentrations of tritylimidazole was performed. Growth was estimated on the basis of absorbance at 600 nm and plotted as percentage growth, where the absorbance in the absence of tritylimidazole was taken to be 100%. The mean values (± S.D.) of three independent experiments are plotted. B, R6G (2.5 μM) accumulation in yeast strains bearing WT and the mutants S558Y and S558Y,N242K (clo-3-0 strain) was analyzed with flow cytometry (21). Representative histograms from four independent experiments are shown. The strains used are labeled on the figure. C, clo-3-0 phenotype was recreated by inserting pS558Y,N242K plasmid DNA into R-1 (pdr5::KANMX4) by lithium chloride transformation. Quantitative growth in liquid medium containing trityl was carried out as described in A with WT (R-1+pSS607; ■), S558Y (R-1+pS558Y; ●), transformants of clo-3-0 (S558Y,N242K;▲), and R-1+pS558Y,N242K;○). The mean values (± S.D.) of three independent experiments are plotted.
the transport-deficient S558Y strain and the S558Y,N242K suppressor exhibit comparable values for the $K_m$ and $V_{\text{max}}$ of ATP hydrolysis (Fig. 8B). Although both the single S558Y mutant and the suppressor show robust ATPase activity, there is an ~1.5-fold decrease in the $V_{\text{max}}$ compared with WT Pdr5. Thus, the behavior of the S558Y ATPase is qualitatively similar to the experiment shown in Fig. 4. Furthermore, it is clear that the second-site mutation does not affect the kinetics of ATP hydrolysis per se.

We demonstrated in earlier studies that clo has a strong inhibitory effect on the ATPase activity of WT Pdr5 (5). We also showed that clo-induced inhibition of ATP hydrolysis is significantly reduced in the S558Y mutant (Fig. 4C). We find that inhibition of ATPase activity approaches levels observed in the WT protein in the S558Y,N242K suppressor (Fig. 8C). These data are consistent with our earlier hypothesis that the S558Y mutation exhibits drug susceptibility because the two key activities of Pdr5, ATP hydrolysis and drug-substrate transport, are uncoupled. These results demonstrate that coupling is clearly restored in clo-3-0.

N242K Mutation Exhibits Drug Sensitivity in a Ser-558 Background—To determine whether the N242K substitution has a drug-sensitive phenotype when separated from S558Y, we transformed R-1 ($pdr5::\text{KANMX4}$) with pN242K. Two independent transformants were tested on 22.5 mM clotrimazole (Figs. 3 and 5). Finally, we showed that the S558Y mutant hydrolyzes ATP, although the $V_{\text{max}}$ is lower than that observed for WT protein.

We screened UV-treated survivors to identify suppressors that eliminated the cyh resistance mediated by Pdr5 and recovered S558Y. Several observations implicate this residue in interdomain communication. For example, yeast cells overexpressing the S558Y mutant Pdr5 and an isogenic $\Delta pdr5$ strain show similar drug hypersensitivity and transport deficiency (Figs. 1 and 2 and Table 2). Nonetheless, nucleotide (8-azido-[\(3^3\)P]ATP) and transport substrate (IAAP) analogues both show photocross-linking to S558Y mutant Pdr5 (Figs. 3 and 5). Furthermore, the WT and mutant Pdr5s have comparable affinities for ATP and clo (Figs. 3 and 5). Finally, we showed that the S558Y mutant hydrolyzes ATP, although the $V_{\text{max}}$ is lower than that observed for WT protein.

We inserted double copies of the $PDR5$ gene to obtain higher levels of the protein for biochemical studies (compare lanes 2 and 3 with 6 and 7 in Fig. 3A). Earlier studies were conducted with purified PMs prepared from single-copy strains (5). In these experiments, the $V_{\text{max}}$ (ATP hydrolysis) value for WT Pdr5 was comparable with that found in the double-copy S558Y mutant (Fig. 4A). Nevertheless, yeast strains with a single-copy WT Pdr5 show 30-fold greater cyh resistance than the double-copy S558Y mutant strain (Fig. 1B). Therefore, it appears very unlikely that the 1.5-fold decrease in the ATPase activity in the mutant is sufficient to account for the null-like phenotype of the S558Y mutant in transport and drug resistance assays (Figs. 1 and 2). Taken together, these data strongly suggest that the S558Y drug hypersensitivity is a consequence of impaired cou-
pling between ATP hydrolysis and the conformational changes in the TMDs that bring about transport.

The S558Y mutation thus provides a unique opportunity to probe the structural elements of the intramolecular domains that couple ATP hydrolysis and transport of drug substrate. Unlike with mammalian cells, it is possible in yeast to screen for second-site suppressors even if they occur at very low frequencies. The strategy is described in detail in Fig. 6. Previous studies involving large scale mutagenesis of several proteins provided evidence that the functional coupling of amino acids in interacting proteins and long range interactions in different domains of a protein are not always predictable from the atomic structure (see Ref. 28 and references therein). The molecular chaperone Hsp70, like Pdr5, exhibits two independent functions (ATP hydrolysis and polypeptide binding) that occur at distinct domains of the protein. Nonetheless, the coupling of these two activities is critical for the chaperone to function. A genetic screen that identified amino acid alterations in the ATPase domain that reverse defects caused by mutations in the peptide-binding domain was successfully used to locate the hot spots in these two domains (29). The yeast as a model system is well suited to such genetic screens. We found that a second mutation at position 242 (N242K) resulted in an almost complete reversal of the null phenotype of the S558Y phenotype (Fig. 7). Moreover, the S558Y,N242K double mutant also showed a $V_{\text{max}}$ for the ATPase comparable with the mutant, S558Y, and 1.5-fold lower than that observed for the WT Pdr5 (Fig. 8B). Thus, we demonstrate equivalent levels of ATP hydrolysis in the drug-sensitive and transport-deficient S558Y mutant Pdr5 and the drug-resistant suppressor S558Y,N242K (Fig. 8B). There is additional evidence to support our contention that the absence of coupling between the NBDs and transport domains rather than a very modest loss in ATPase function is the cause of the drug-sensitive phenotype of the S558Y mutant Pdr5. Clo is a potent noncompetitive inhibitor of ATPase activity. This inhibition occurs at a location that is distinct from the transport site of the drug and the NBDs (5). The loss of inhibition in S558Y and its nearly complete restoration in the suppressor strongly imply that the NBD1/TMH2 interface is a route for both productive and inhibitory signals. Fig. 10 proposes a model showing the structural changes in the WT, S558Y mutant, and suppressor (S558Y,N242K) strains. Fig. 10A shows the arrangement of the NBD and TMD domains in Pdr5. The organization of part of the N-terminal NBD (between the

FIGURE 9. Drug sensitivity of WT Pdr5 and S558Y and N242K. A, 10-fold serial dilutions (0.5 $\times$ 10$^{6}$ cells to 0.5 $\times$ 10$^{3}$ cells in 5 $\mu$L of sterile H$_2$O) were plated on YPD and on YPD containing 3.1 $\mu$M cycloheximide or 22.5 $\mu$M clotrimazole. Plates were scanned after 96 h of incubation at 30 °C. Row 1, R-1 (pdr5) cells; row 2, WT (R-1 + pS607) cells; row 3, R-1 + pS558Y; rows 4 and 5, two independent transformants of R-1 + pN242K. B, quantitative growth analysis of WT (○), S558Y (●), and N242K (▲) in the presence of increasing concentrations of cyc.h. Growth in the absence of cyc.h was taken to be 100% in each curve. C, growth of WT (○), S558Y (●), and N242K (▲) was compared as described in B, except that the experiment was carried out in the presence of increasing concentrations of clotrimazole.
Walker A domain and the signature sequence) is illustrated in Fig. 10B. Residue Ser-558 is located toward the extracellular end of TMH2. Hydrophobicity plots and sequence alignment predict that Ser-558 is lipid facing and lies toward the extracellular end of the TMH. Sequence alignments of the NBDs of Pdr5 show that the residue Asn-242 lies between the Walker A and Signature sequence of ABC transporters. The location of N242K, identified in a mutant screen in this study, is shown by a red star. Note that this residue as well as Glu-256 which co-varies with Ser-558 in correlated mutation analysis (see text) are located slightly upstream of the Q-loop. The x-ray crystallographic structure of the ABC transporter Sav1866 describes the “x-loop” in the NBD, which is implicated in the coupling of NBDs and TMDs via the ICL1 (10). The x-loop lies between the Walker A and signature regions of the NBD; the Q-loop also lies in this region and has been implicated in communication between the NBDs and transport sites (34). The central panel shows that the S558Y mutation (yellow circle) involves replacement with a bulkier residue that could result in the tilting of TM helix 2 (see structures in the black boxes for comparison). As the TM helix 2 is directly linked to the ICL1, the interface between the ICL1 and the NBD1 could be disrupted (purple arrows). This is consistent with our results that demonstrate that the S558Y has no effect on the affinity of Pdr5 for nucleotide or transport substrate but shows complete loss of transport activity because the NBD and transport-substrate sites are uncoupled. The right panel shows the S558Y,N242K double mutant. The structures show that the lysine that replaces the asparagine has more rotamers and a potentially longer reach. It is plausible that this change reestablishes the contacts between NBD1 and ICL1 (red arrow).
Pdr5 and Sav1866, including the organization of the NBDs and TMDs as well as key residues in the NBDs. It is nonetheless noteworthy that we find TMH2 and the region upstream of the signature sequence associated in the coupling of ATP hydrolysis to drug-substrate translocation. Additionally, correlated mutation analysis predicts that mutations at Ser-558 co-vary with those at Glu-256. This highly predictive statistical analysis is based on the concept that the functional coupling of two positions, even if distantly positioned in the structure, should be reflected in the mutually constrained evolution at the two positions (28, 31). In the case of Pdr5 our analysis shows that a mutation at position 256 is the most likely “response” to the mutation S558Y, demonstrating that the residue Ser-558 has been subjected to evolutionary pressure as a pair. It is interesting that the residue 256 also lies in the region between the Walker A and signature sequence of NBD1 of Pdr5 (Fig. 10B). A plausible explanation of how the S558Y mutation (in TMH2) uncouples the contacts between ICL1 and NBD1 is provided in Fig. 10C. We speculate that the additional bulk and/or conformational flexibility of the lysine residue compared with the asparagine permits the reestablishment of these contacts in the S558Y,N242K double mutant. The molecular models of Asn and Lys in Fig. 10C clearly show that Lys has a larger size as well as significantly more rotamers that could permit the reestablishment of contacts between the NBD and ICL1. The N242K mutant in the WT background is more sensitive to drugs compared with the WT. However, the N242K mutant exhibits greater resistance than the Δpdr5 or the S558Y mutant. An alignment of NBDs of fungal ABC transporters shows that residues at the position equivalent to Asn-242 in Pdr5 are not conserved. Besides asparagine, two other residues are found to occur at this location, threonine and serine. Thus a small polar amino acid that can participate in hydrogen bonding always occurs at this position. This study indicates that the asparagine residue at position 242 plays an important role in inter-domain communication between the NBDs and TMDs, plausibly via hydrogen bonds. It is thus likely that an Asn → Lys mutation at this location could disrupt these contacts. Nonetheless, as stated above, the increased conformational flexibility of the lysine residue could restore these contacts albeit not as efficiently as in the WT Pdr5.

It is important to note that this study suggests that conformational changes in TMH2 are coupled to those at the ATP site in the N-terminal half of Pdr5. The structures of NBDs of numerous ABC proteins support the notion that the ATP is sandwiched between the Walker A and B domains and H-loop of one NBD and the signature sequence and the D-loop of the other (for review see Ref. 8). Thus the interactions described here would involve Walker A and B domains and the H-loop of NBD1 and the signature sequence of NBD2. In Pdr5 several residues that are highly conserved among ABC proteins (and have also been shown to play key functional roles) are altered in the NBD1. These include the conserved lysine in the Walker A, the catalytic carboxylate in Walker B, and the conserved histidine that defines the H-loop. This has led to the proposal that the NBD1 either lacks function or plays a regulatory role (1, 13). The asymmetry in the two ATP sites of Pdr5 is similar to that observed in most members of the ABCC subfamily of mamma-

lial ABC proteins (32). Of these, the cystic fibrosis transmembrane conductance regulator (ABCC7) has been studied in considerable detail, and the consensus appears to be that the NBD1 is regulatory (33).

Our data show that residues in TMH2 and NBD1 are strongly coupled and clearly have a functional role and appear to define the transmission interface. This would suggest that the NBD1 of Pdr5 has, at the very minimum, a regulatory function in the transport cycle. In addition unlike many ABC proteins, the ATPase activity of Pdr5 is not stimulated by the transport substrate (5, 13). In bacterial transporters this coupling is dependent on the conserved histidine in the H-loop. Pdr5 lacks this residue in NBD1, and Schmitt and co-workers (13) have shown that the His → Ala mutation in Pdr5 has no effect on the ATPase activity. This has led them to posit that Pdr5 exhibits uncoupled ATPase activity. This may well be true; however, our data strongly suggest that structural domains involved in the transmission of conformational changes generated by ATP binding/hydrolysis to the TMDs are intact and may share features common to other ABC proteins.

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