

# A novel motif in the yeast mitochondrial dynamin Dnm1 is essential for adaptor binding and membrane recruitment

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To initiate mitochondrial fission, dynamin-related proteins (DRPs) must bind specific adaptors on the outer mitochondrial membrane. The structural features underlying this interaction are poorly understood. Using yeast as a model, we show that the Insert B domain of the Dnm1 guanosine triphosphatase (a DRP) contains a novel motif required for association with the mitochondrial adaptor Mdv1. Mutation of this conserved motif specifically disrupted Dnm1–Mdv1 interactions, blocking Dnm1 recruitment and mitochondrial fission.

Suppressor mutations in Mdv1 that restored Dnm1–Mdv1 interactions and fission identified potential protein-binding interfaces on the Mdv1  $\beta$ -propeller domain. These results define the first known function for Insert B in DRP–adaptor interactions. Based on the variability of Insert B sequences and adaptor proteins, we propose that Insert B domains and mitochondrial adaptors have coevolved to meet the unique requirements for mitochondrial fission of different organisms.

## Introduction

Eukaryotic cells possess a family of dynamin-related proteins (DRPs), each of which is responsible for a specific cellular membrane-remodeling event. For example, the dynamin-related GTPase Drp1 (Dnm1 in yeast) mediates mitochondrial (Bleazard et al., 1999; Labrousse et al., 1999; Sesaki and Jensen, 1999) and peroxisomal fission (Koch et al., 2003; Li and Gould, 2003; Kuravi et al., 2006), Atlastin (Hu et al., 2009; Orso et al., 2009; Moss et al., 2011) and Mitofusins (Hales and Fuller, 1997; Hermann et al., 1998; Rapaport et al., 1998; Chen et al., 2003; Eura et al., 2003) play roles in ER and mitochondrial membrane fusion, respectively, and ARC5 (Gao et al., 2003) facilitates chloroplast membrane division. Like classical dynamin, DRPs self-assemble into highly ordered oligomers that use the energy of GTP hydrolysis to remodel lipid bilayers (Praefcke and McMahon, 2004).

Dynamin and DRPs have conserved GTPase, middle, and GTPase effector domains (see Fig. 1 A; van der Bliek, 1999). These domains mediate self-assembly and modulate GTPase activity. Dynamin and DRPs also contain nonconserved domains that, in some cases, have been shown to determine their cellular

distribution and heterotypic interactions. For example, a proline-rich domain at the C terminus of dynamin facilitates its binding to a variety of actin-binding proteins. Dynamin also harbors a pleckstrin homology (PH) domain between its GTPase and GTPase effector domain, which is essential for interactions with the plasma membrane. In place of the PH domain, DRPs contain a region called Insert B (InsB) whose length and sequence varies. Whether or not there is a conserved function for InsB is not clear.

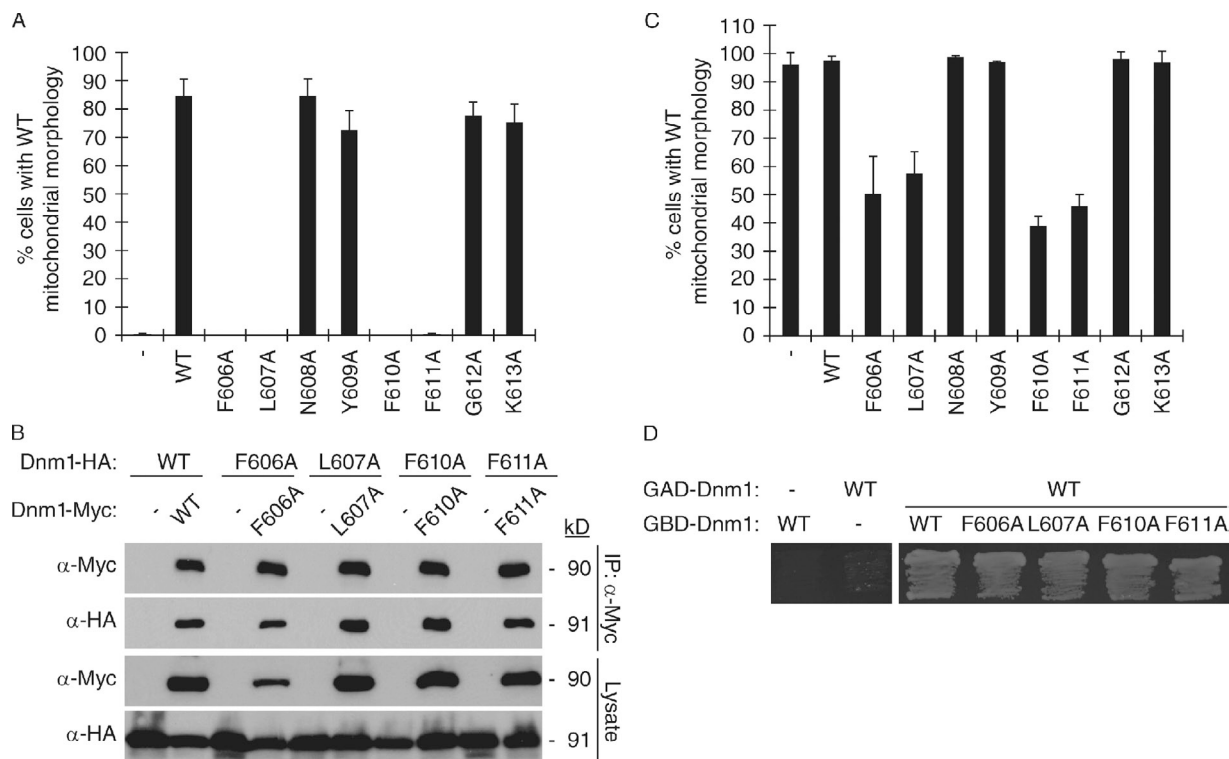
Although dynamin interacts directly with the lipid bilayer via its PH domain, DRPs do not initially interact with the lipid bilayers they remodel. Instead, they are recruited to specific cellular sites via interactions with membrane-associated adaptor proteins. In most cases, the DRP domains necessary for adaptor interactions have not been identified. However, structural studies of dynamin and several DRPs (Mears et al., 2007, 2011; Faelber et al., 2011; Ford et al., 2011) suggest that InsB is a likely candidate because it is predicted to reside at the base of the DRP oligomer, closest to the membrane.

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Abbreviations used in this paper: coIP, coimmunoprecipitation; DRP, dynamin-related protein; DSP, dithiobis(succinimidyl propionate); InsB, Insert B; mt-ffRFP, mitochondrial-targeted fast-folding RFP; PH, pleckstrin homology; WT, wild-type.

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**Figure 2. Dnm1 InsB conserved residues are important for Dnm1 function in mitochondrial fission.** (A) Quantification of mitochondrial morphology in *dnm1Δ* cells expressing indicated Dnm1 variants. (B) Lysates from cells expressing the indicated C-terminal HA- or Myc-tagged Dnm1 proteins were used for immunoprecipitation with anti-Myc agarose beads. Immunoprecipitated fractions (top) and lysates (bottom) were analyzed by SDS-PAGE and Western blotting with anti-Myc and anti-HA antibodies. (C) Quantification of mitochondrial morphology in WT cells expressing indicated Dnm1 variants. (A and C) Black columns and error bars represent the mean and standard deviation of at least three independent experiments ( $n = 100$ ). (D) pGBD and pGAD plasmids expressing the indicated fusion proteins were cotransformed into the Y187 yeast two-hybrid reporter strain and grown on S-Dextrose minus adenine plates at 30°C for 3 d.

We also observed that coexpression of WT and Dnm1<sup>InsBmut</sup> proteins caused dominant-negative fission defects (Fig. 2 C). Although expression of a second copy of WT *DNM1* in WT cells did not cause significant changes in mitochondrial morphology, expression of Dnm1<sup>InsBmut</sup> proteins in WT cells blocked fission in up to 60% of the population. One explanation for these dominant-negative fission phenotypes is that Dnm1<sup>InsBmut</sup> proteins are able to assemble with WT Dnm1. Consistent with this idea, we observed an interaction between WT Dnm1 and Dnm1<sup>InsBmut</sup> proteins in a two-hybrid assay (Fig. 2 D). The interaction of Dnm1<sup>InsBmut</sup> with WT Dnm1 could interfere with multiple steps in mitochondrial fission including fission complex formation and Dnm1–adaptor interactions.

### The InsB motif is essential for Dnm1 binding to the Mdv1 adaptor

After mitochondrial membrane recruitment and self-assembly, the majority of GFP-tagged Dnm1 can be visualized as puncta on mitochondrial tubules (Fig. 3 A; Otsuga et al., 1998). In contrast, all of the GFP-Dnm1<sup>InsBmut</sup> proteins failed to associate with mitochondria (Fig. 3 A, a representative cell is shown). Instead, GFP-Dnm1<sup>InsBmut</sup> proteins assembled into punctuate structures that moved rapidly through the cytoplasm and could not be captured by digital imaging. In a few cells, GFP-Dnm1<sup>InsBmut</sup> also formed larger, immobile aggregates in the cytoplasm (Fig. 3 A). Importantly, this localization of GFP-Dnm1<sup>InsBmut</sup>

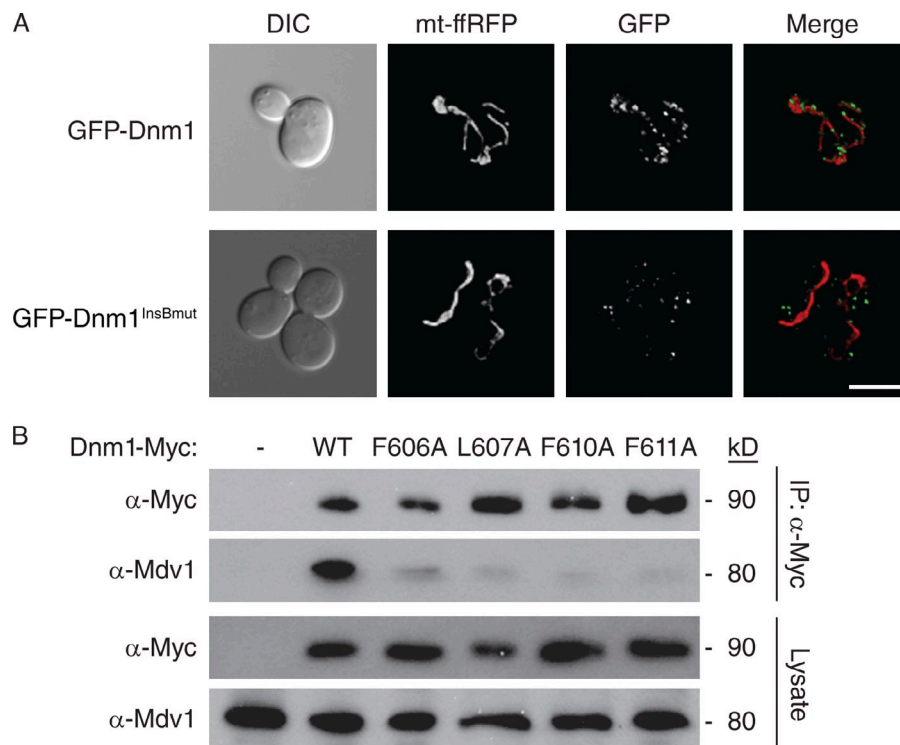
proteins is indistinguishable from that observed for GFP-Dnm1 in cells lacking Fis1 or fission adaptor proteins (Mozdy et al., 2000; Griffin et al., 2005). This observation raised the possibility that mutations in Dnm1<sup>InsBmut</sup> proteins were disrupting interactions with the Mdv1 adaptor.

In vitro pull-down assays using purified GST-Mdv1 β-propeller and His-tagged Dnm1 showed that the Mdv1 β-propeller domain binds directly to Dnm1 (Fig. S1 B). To determine whether InsB mutations affected the Dnm1–Mdv1 interaction, we performed coIP experiments from *dnm1Δ* cells expressing Myc-tagged Dnm1<sup>InsBmut</sup> proteins. Although full-length Mdv1 was efficiently coprecipitated by WT Dnm1-Myc, Mdv1 interaction with all of the Dnm1<sup>InsBmut</sup>-Myc proteins was dramatically reduced (Fig. 3 B). These combined results identify a novel motif in InsB essential for Dnm1–Mdv1 interaction and Dnm1 recruitment to mitochondria.

### Suppressors of a *dnm1*<sup>InsBmut</sup> mutation cluster in the Mdv1 β-propeller

Using an integrated form of the *dnm1*<sup>F610A</sup> mutation, we performed a suppressor screen to identify residues in Mdv1 important for Dnm1–Mdv1 interaction (see Materials and methods and Table S1). Although the screen covered 84% of the *MDV1* coding sequence, all but one suppressor mutation fell in the Mdv1 β-propeller domain (Fig. 4 A). Most of the affected residues localized to the top of the Mdv1 β-propeller model (Fig. 4 B, right),

**Figure 3. InsB conserved residues are critical for Dnm1–Mdv1 interaction.** (A) Representative images of GFP-Dnm1 and GFP-Dnm1<sup>InsBmut</sup> (F606A, L607A, F610A, or F611A) localization in *dnm1Δ* cells. Differential interference contrast microscopy (DIC), mitochondrial matrix-targeted dsRed (mt-ffRFP), GFP, and merged images are shown. Bar, 5 μm. (B) Lysates from cells expressing the indicated C-terminal Myc-tagged Dnm1 were used for immunoprecipitation with anti-Myc agarose beads. Immunoprecipitated fractions (top) and lysates (bottom) were analyzed by SDS-PAGE and Western blotting with anti-Myc and anti-Mdv1 antibodies.



suggesting that they are part of an interaction interface. Suppressors affecting residues on the bottom of the structure may define an additional binding interface. Two additional mutations, S557C and T558I, lay in a short sequence that was eliminated during homology modeling. These results are consistent with our finding that the Mdv1 β-propeller is sufficient for direct interaction with Dnm1 (Fig. S1 B). The final suppressor mutation altered residue Q288 in the Mdv1 coiled-coil domain. This suppressor was not analyzed further, as the structure and function of the coiled-coil domain has been extensively studied (Koirala et al., 2010; Zhang et al., 2012).

In addition to Mdv1, yeast encodes a second fission adaptor protein called Caf4 (Griffin et al., 2005). Although these two adaptors are paralogues and have similar domain structures, Caf4 is not essential for mitochondrial fission. Sequence alignment revealed that most of the suppressor mutations affected conserved or similar amino acids in the Mdv1 and Caf4 β-propeller domains (Fig. 4 C). This conservation is consistent with an important functional role for these amino acids in vivo.

All but one of the residues (S541) identified in our suppressor screen differ from those reported by others (Cervený and Jensen, 2003; Naylor et al., 2006). The mutations in these previous studies were selected based on homology modeling with known β-propeller structures, whereas the mutations we identified here were selected by the organism in a suppressor screen and are specifically relevant to defects caused by InsB mutations. The exception, S541, was reported not to have a phenotype when replaced by glutamine (Q) in the full-length Mdv1 protein (Naylor et al., 2006). In contrast, we found that an Mdv1 S541G mutant protein was unable to rescue mitochondrial fission defects in an *mdv1Δ* strain (Table 1). This difference may be because of the different yeast strain backgrounds used

in these two studies. It is also possible that mutation to glycine has a more significant effect on local protein structure and flexibility than mutation to glutamine.

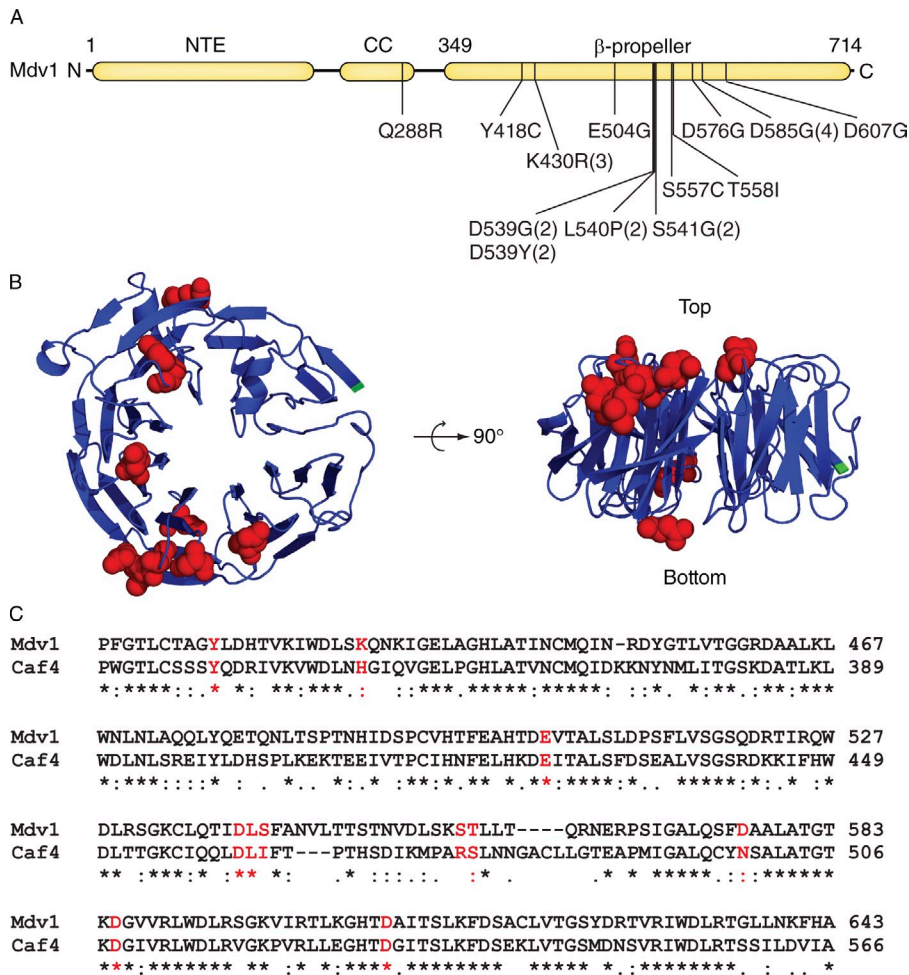
#### Suppressor mutations in the Mdv1 β-propeller rescue mitochondrial fission defects caused by Dnm1<sup>F610A</sup>

In control studies, the Mdv1 suppressor proteins were all stably expressed in vivo (Fig. S1 C). To verify that the Mdv1 suppressor proteins were capable of rescuing fission, we expressed them from a plasmid in cells lacking WT Mdv1 and expressing Dnm1<sup>F610A</sup> protein from the genome. All of the Mdv1 suppressors rescued mitochondrial fission defects in this strain (up to 70% rescue). Representative results are shown in Fig. 5 A for the five best rescuing Mdv1 suppressors (Y418C, D539G, D539Y, L540P, and D576G). The same Mdv1 suppressors also rescued mitochondrial morphology defects caused by other Dnm1<sup>InsBmut</sup> proteins (F606A, L607A, and F611A; Fig. 5 B).

To test whether Mdv1 suppressors could rescue the mitochondrial recruitment of Dnm1<sup>F610A</sup>, Mdv1 suppressors and GFP-Dnm1<sup>F610A</sup> were coexpressed in an *mdv1Δdnm1Δ* strain. GFP mitochondrial puncta were visible in up to 70% of these cells (Fig. 5 C). The mitochondrial recruitment and puncta formation by GFP-Dnm1<sup>F610A</sup> suggested that Dnm1 interaction with the Mdv1 suppressor proteins had been restored. However, this interaction could not be detected in coIP assays, even after chemical cross-linking (unpublished data). Thus, the restored interaction between Dnm1<sup>F610A</sup> and Mdv1 suppressors is less robust than WT.

As an alternative, we evaluated Dnm1<sup>F610A</sup>–Mdv1 suppressor interactions using the yeast two-hybrid growth assay. Previous studies established that WT Dnm1 and WT Mdv1 interact in





**Figure 4. Suppressors of a *dnm1<sup>InsB</sup>* mutation cluster in the Mdv1  $\beta$ -propeller.** (A) Domain structure of Mdv1 with indicated Dnm1<sup>F610A</sup> suppressor mutations (the number of mutations obtained for each allele is in parentheses). NTE, N-terminal extension; CC, coiled coil. (B) Top and side views of the Mdv1  $\beta$ -propeller model (blue). Red indicates residues changed by suppressor mutations. The N terminus is in green. (C) Alignment of the portions of the Mdv1 (aa 408–643) and Caf4 (aa 329–566)  $\beta$ -propeller sequences. Residues affected by suppressor mutations are shown in red. Symbols below the sequence alignment indicate identity (\*), strong similarity (:), and weak similarity (.) of amino acids.

this assay (Fig. 5 D; Tieu and Nunnari, 2000; Cervený and Jensen, 2003; Karren et al., 2005). Although the interaction was severely disrupted when Dnm1<sup>F610A</sup> was paired with WT Mdv1,

the interaction was partially (D539Y and L540P) or completely (Y418C and D539G) restored by substituting an Mdv1 suppressor protein for WT Mdv1 (Fig. 5 D). Surprisingly, the Mdv1<sup>D576G</sup>

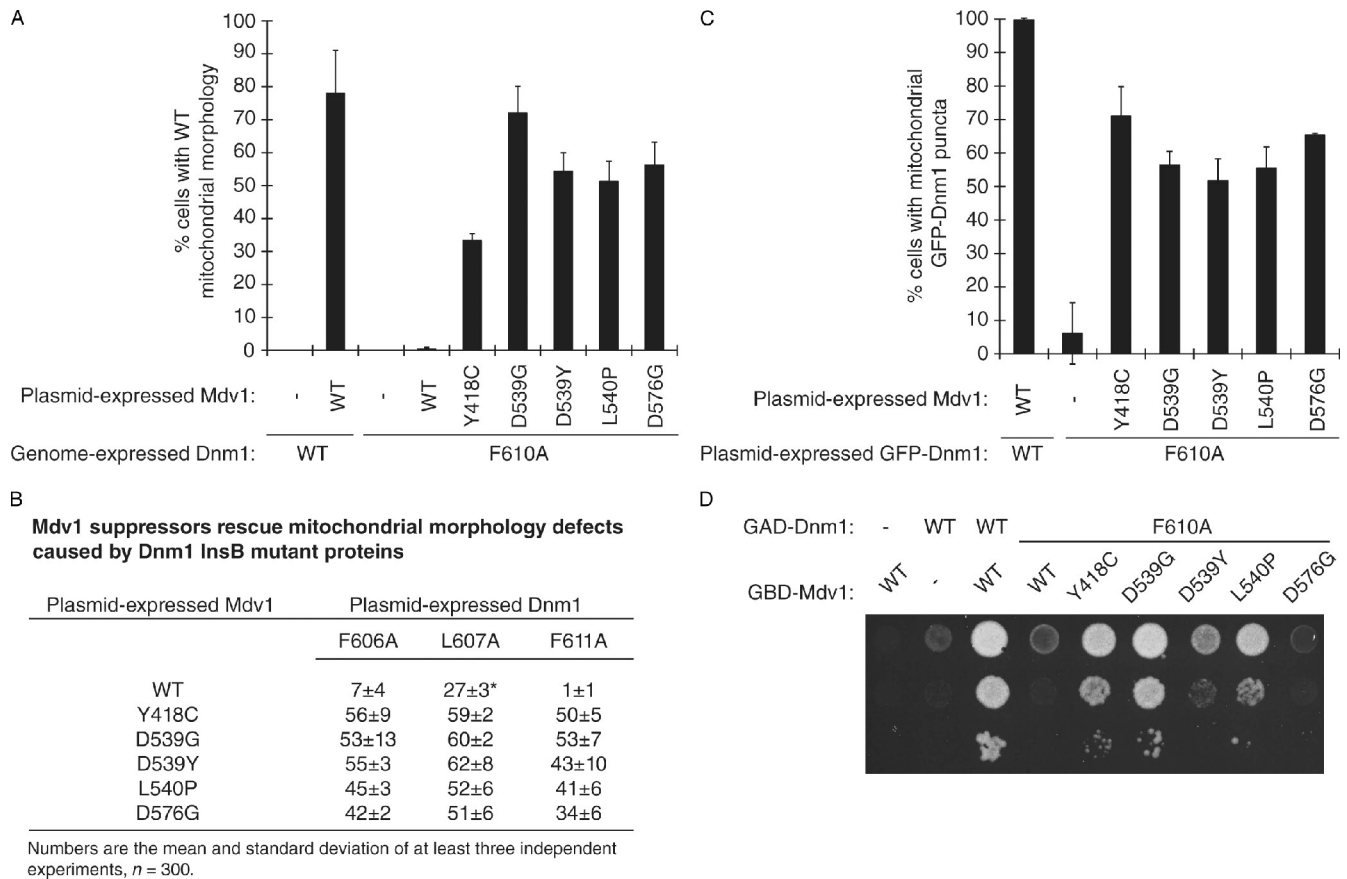
**Table 1. Characterization of Mdv1 suppressors of the *dnm1<sup>F610A</sup>* allele**

Mutations	WT mitochondrial morphology <sup>a</sup>		GFP mitochondrial puncta <sup>b</sup>		Interaction with Dnm1 <sup>c</sup>	
	<i>DNM1 mdv1Δ</i>	<i>dnm1<sup>F610A</sup> mdv1Δ</i>	Dnm1	Dnm1 <sup>F610A</sup>	Dnm1	Dnm1 <sup>F610A</sup>
	%	%				
Mdv1	67 ± 7	0 ± 0	+	+	+	–
Y418C	6 ± 3	41 ± 16	+	+	++	+
K430R	9 ± 8	28 ± 4	+	+	+	–
E504G	19 ± 8	32 ± 16	+	+	++	–
D539G	9 ± 2	68 ± 11	+	+	++	+
D539Y	11 ± 1	61 ± 14	+	+	++	+
L540P	12 ± 5	58 ± 14	+	+	++	+
S541G	8 ± 4	38 ± 12	+	+	++	+
S557C	9 ± 8	45 ± 8	+	+	++	+
T558I	10 ± 8	34 ± 6	+	+	++	+
D576G	68 ± 17	53 ± 9	+	+	–	–
D585G	45 ± 3	30 ± 8	+	+	–	–
D607G	15 ± 6	31 ± 12	+	+	++	–

<sup>a</sup>Numbers are the mean and standard deviation of at least three independent experiments, *n* = 300.

<sup>b</sup>+, GFP puncta localized to mitochondrial tubules.

<sup>c</sup>Yeast two-hybrid assays. –, No growth on His or Ade minus medium; +, growth on His but not Ade minus medium; ++, growth on both His and Ade minus media. Growth on Ade minus medium is the more stringent indicator of protein–protein interaction.



**Figure 5. Suppressor mutations in the Mdv1  $\beta$ -propeller rescue mitochondrial fission defects caused by Dnm1<sup>F610A</sup>.** (A) Quantification of mitochondrial morphology in *mdv1 $\Delta$  DNM1* and *mdv1 $\Delta$  dnm1::dnm1<sup>F610A</sup>* strains expressing the indicated Mdv1 suppressor proteins. (B) Quantification of mitochondrial morphology in *mdv1 $\Delta$  dnm1 $\Delta$*  strains expressing indicated Mdv1 and Dnm1 proteins from plasmids. The mitochondria were visualized with MitoFluor Red 589 (Molecular Probes). (\*) Plasmid-expressed WT Mdv1 rescued Dnm1<sup>L607A</sup> better than genome-expressed WT Mdv1 (Fig. 2 A). This is likely because of the higher steady-state abundance of the plasmid-expressed protein. (C) Quantification of GFP-Dnm1 localization in *mdv1 $\Delta$  dnm1 $\Delta$*  cells expressing the indicated Mdv1 and GFP-Dnm1 variants. (A and C) Black columns and error bars represent the mean and standard deviation of at least three independent experiments (A,  $n = 300$ ; C,  $n = 150$ ). (D) pGBD and pGAD plasmids expressing the indicated fusion proteins were cotransformed into the Y187 yeast two-hybrid reporter strain and grown on S-Dextrose minus histidine plates at 30°C for 3 d.

suppressor protein reproducibly rescued both mitochondrial morphology and GFP-Dnm1<sup>F610A</sup> localization (Fig. 5, A–C), but was unable to restore growth in the two-hybrid assay (Fig. 5 D). In the morphology rescue and localization studies, binding of Dnm1<sup>F610A</sup> to this Mdv1 suppressor may be sufficient to recruit Dnm1<sup>F610A</sup> to the membrane, after which co-oligomerization of both proteins into mitochondrial fission complexes could further stabilize the interaction. Such stabilizing forces may be compromised by the spheroplasting/lysis required for coIP studies or by the fusion of both proteins to nuclear targeting sequences used in the two-hybrid assay. This interpretation is supported by our finding that GFP-Mdv1<sup>D576G</sup> assembled into punctate fission complexes in the presence of Dnm1<sup>F610A</sup> (Fig. S1 D).

## Discussion

Although Dnm1 binding to Mdv1 and recruitment to the mitochondrial membrane is essential for fission, the Dnm1 domains required for this interaction were not known. Here we identify a novel motif in Dnm1 InsB that is specifically required for interaction with the Mdv1  $\beta$ -propeller. Second-site suppression studies

and cell-based assays confirm that the InsB– $\beta$ -propeller interaction is critical for Dnm1–Mdv1 binding and Dnm1 membrane recruitment. The Dnm1 InsB motif and Mdv1 adaptor sequences required for this interaction are conserved in fungi but not in mammals or plants (Fig. 6). Thus, different InsB domains and adaptors may have coevolved in different organisms to mediate membrane targeting of mitochondrial dynamin-related GTPases.

Our mutational analysis demonstrates that four hydrophobic residues in the Dnm1 InsB motif are necessary for interaction with Mdv1. However, the purified Dnm1 InsB domain (aa 535–639) is not sufficient to bind the Mdv1  $\beta$ -propeller in vitro (unpublished data). Because the conserved motif in InsB is hydrophobic, it may not adopt a functional conformation when InsB is expressed on its own. In full-length Dnm1, the InsB motif may normally be buried in the interior of the protein. In this case, a conformational change in Dnm1 would be required to expose critical residues in InsB for adaptor binding. A conformational change of this type could be stimulated by GTP binding to Dnm1. This scenario would be consistent with the previous finding that Mdv1 preferentially binds to the GTP-bound form of Dnm1 (Cervený and Jensen, 2003; Naylor et al., 2006; Lackner et al., 2009).





mitochondrial adaptor called Mff (Strack and Cribbs, 2012). The model that InsB domains mediate DRP interactions in other organisms can be directly tested in genetic and cellular studies, as well as structural studies of DRPs bound to their cognate adaptor proteins.

## Materials and methods

### Yeast strains and plasmids

Yeast strains used in this study include JSY5740 (*MATa leu2Δ1 his3Δ200 trp1Δ63 ura3-52 lys2Δ202*), JSY1361 (*MATα leu2Δ1 his3Δ200 trp1Δ63 ura3-52 lys2Δ202 dnm1::HIS3*), JSY9134 (*MATα leu2Δ1 his3Δ200 trp1Δ63 ura3-52 lys2Δ202 mdv1::HIS3 dnm1::HIS3*), JSY9744 (*MATα leu2Δ1 his3Δ200 trp1Δ63 ura3-52 lys2Δ202 mdv1::HIS3 dnm1::dnm1<sup>F610A</sup> fzo1-1*), JSY9983 (*MATα leu2Δ1 his3Δ200 trp1Δ63 ura3-52 lys2Δ202 mdv1::HIS3 dnm1::dnm1<sup>F610A</sup>*), JSY3903 (*MATα leu2Δ1 his3Δ200 trp1Δ63 ura3-52 lys2Δ202 mdv1::HIS3*), and JSY5148 (*MATa trp1-901 leu2-3, 112 ura3-52 his3-200 gal4Δ gal80Δ LYS2::GAL1-HIS3, GAL2-ADE2 met2::GAL7-lacZ*).

A list of plasmids used in this study is provided in Table S2, except for the His-Dnm1 expression plasmid, which was received from J. Nunnari (University of California, Davis, CA). For *pRS415-dnm1<sup>InsBmut</sup>*, site-directed mutagenesis of a *pRS415-DNM1* template was used to introduce sequences encoding single substitutions F606A, L607A, N608A, Y609A, F610A, F611A, G612A, and K613A. Similar methods were used to generate *pRS415MET25-GFP-dnm1<sup>InsBmut</sup>*, *pRS425-dnm1<sup>InsBmut</sup>-MYC*, and *pRS426-dnm1<sup>InsBmut</sup>-3xHA*. To create *pGAD-C1-DNM1* and *pGAD-C1-dnm1<sup>F610A</sup>*, BamHI-DNM1-Sall and BamHI-dnm1<sup>F610A</sup>-Sall fragments were cloned between the BamHI and Sall sites of the *pGAD-C1* vector. Similarly, BamHI-MDV1-Sall and BamHI-mdv1<sup>suppressor</sup>-Sall fragments were cloned between the BamHI and Sall sites of the *pGBD-C1* vector to generate *pGBD-C1-MDV1* and *pGBD-C1-mdv1<sup>suppressor</sup>*.

### Fluorescence microscopy

Mitochondrial morphologies were quantified in WT, *dnm1Δ*, and *dnm1Δmdv1Δ* strains expressing the indicated proteins. The WT morphology category includes unbudded or budded cells with more than two free tubule ends in the mother cell. The formation of GFP-Dnm1 mitochondrial puncta was quantified by analysis of deconvolved epifluorescence images of random fields of cells. Phenotypic quantification is reported as the mean and standard deviation of three independent experiments (total  $n \geq 300$  cells unless noted). Unless specified in the figure legend, the mitochondria were visualized by expressing mitochondrial-targeted fast-folding RFP (mt-fFRFP). Dnm1 InsB variants were expressed from the *DNM1* promoter in the *pRS415* vector. GFP-tagged Dnm1 variants were expressed from the *pRS415-MET25* vector. Mdv1 variants were expressed from the *pRS416-MET25* plasmid. Yeast cells were grown at 30°C in selective synthetic dextrose medium containing 0.1 mg/ml methionine. Overnight cultures were diluted to 0.2 OD<sub>600</sub> and grown for 3–5 h (OD<sub>600</sub>, 0.5–1.0). The methionine-repressible *MET25* promoter is leaky under these conditions and expresses approximately four-fold more protein at steady state than that expressed from the endogenous *DNM1* or *MDV1* promoters (Karren et al., 2005).

A microscope (Axioptan 2; Carl Zeiss) equipped with a 100× oil immersion objective was used to observe and image cells. For mitochondria and GFP-Dnm1 puncta, 0.275 μm optical sections encompassing the entire yeast cells were deconvolved and analyzed using Axiovision version 4.6 (Carl Zeiss). All slices were projected on the transparency setting and the three-dimensional projections were converted to a single image. Final images were assembled using Photoshop and Illustrator (Adobe). Linear brightness and contrast adjustments were applied to the entire image.

### ColP assays

For Dnm1–Dnm1 interaction experiments, functional HA- and Myc-tagged Dnm1 variants were expressed in *dnm1Δ* cells. ColPs with anti-c-Myc agarose-conjugated beads (Sigma-Aldrich) were performed as described previously (Koirala et al., 2010). 30 OD<sub>600</sub> cell equivalents were harvested and resuspended in 500 μl IP buffer (0.5% Triton X-100, 150 mM NaCl, 1 mM EDTA, 50 mM Tris, pH 7.4, and 1:500 protease inhibitor cocktail set III [EMD]). Cells were lysed with glass bead and cleared by centrifugation at 18,000 g for 10 min. 400 μl of supernatant was incubated with 40 μl of anti-c-Myc-conjugated agarose beads (Sigma-Aldrich) for 1 h at 4°C. Agarose beads were collected and washed in immunoprecipitation buffer. The bound proteins were released by incubating the beads in 60 μl SDS-PAGE

sample buffer lacking β-mercaptoethanol at 60°C for 8 min. 3.2 μl β-mercaptoethanol was added to the samples before boiling. Immunoprecipitated proteins were analyzed by SDS-PAGE and Western blotting with anti-Myc (Santa Cruz Biotechnology, Inc.) and anti-HA (University of Utah Core Facility) antibodies.

Mdv1–Dnm1 interaction was analyzed by colP after dithiobis(succinimidyl propionate) (DSP) cross-linking in *dnm1Δ* cells expressing endogenous Mdv1 and plasmid-borne Myc-tagged Dnm1 variants (Koirala et al., 2010). 50 OD<sub>600</sub> cell equivalents were spheroplasted by treating with 0.2 mg/ml zymolase for 60 min at 30°C followed by treatment with 2.5 mM DSP (Thermo Fisher Scientific) at 30°C for 30 min. 50 mM glycine was added to the cell suspensions and all subsequent buffers to quench DSP. Spheroplasts disrupted using a dounce homogenizer (Wheaton) were spun at 18,000 g for 10 min. Pellets were solubilized for 10 min at 4°C in 500 μl of immunoprecipitation buffer (1% Triton X-100, 150 mM NaCl, 30 mM Hepes-KOH, pH 7.4, and 1:500 protease inhibitor cocktail set III) and centrifuged at 18,000 g for an additional 10 min. 400 μl of supernatant was incubated for 1 h at 4°C with anti-HA-conjugated agarose beads (Sigma-Aldrich). Proteins released from agarose beads were separated by SDS-PAGE and analyzed by ECL Western blotting with anti-Myc and anti-Mdv1 antibodies.

### Screen for mdv1 suppressors of the dnm1<sup>F610A</sup> allele

The growth phenotypes of strains used for this screen are summarized in Table S1. PCR amplification with Taq DNA polymerase was used to introduce random mutations into the *MDV1* coding region. The PCR products were introduced into linearized *pRS416-MET25* using gap repair (Orr-Weaver et al., 1983) in *mdv1Δ dnm1::dnm1<sup>F610A</sup> fzo1-1* cells. In temperature-sensitive *fzo1-1* cells, ongoing mitochondrial fission causes fragmentation, mitochondrial genome loss, and inability to grow on glycerol medium at 37°C (Hermann et al., 1998). Disrupting fission in this strain by introducing an *mdv1Δ* mutation and expressing *dnm1<sup>F610A</sup>* from the endogenous *DNM1* locus (*dnm1::dnm1<sup>F610A</sup>*) prevents mitochondrial fragmentation and genome loss, allowing *mdv1Δ dnm1::dnm1<sup>F610A</sup> fzo1-1* strains to grow on glycerol at the elevated temperature. Expression of WT Mdv1 from a plasmid does not restore the temperature-sensitive glycerol growth defect in this strain. In contrast, *mdv1Δ dnm1::dnm1<sup>F610A</sup> fzo1-1* cells expressing Mdv1<sup>suppressor</sup> from a plasmid fail to grow on glycerol at 37°C, indicating that Mdv1<sup>suppressor</sup> restores mitochondrial fission. Cells containing Mdv1<sup>suppressor</sup>-expressing plasmids were identified by their ability to grow on glycerol at 25°C, but not at 37°C. Candidate clones with verified phenotypes were sequenced to identify *MDV1* mutations. In alleles with multiple amino acid changes, mutations were separated by site-directed mutagenesis. Mutations contributing to growth phenotypes were analyzed for mitochondrial morphology and GFP-Dnm1 localization.

### Mdv1 β-propeller modeling

The β-propeller model shown in Fig. 4 was generated from the crystal structure of the Cdc4 WD40 repeat (PDB accession no. 1NEX) using the PHYRE Protein Fold Recognition server (Kelley and Sternberg, 2009). Residues 349–713 of Mdv1 are variably modeled as a seven- or an eight-bladed β-propeller, depending on the structures most recently deposited in the PDB. The eight-bladed β-propeller model shown in Fig. 4 includes the majority of residues identified in the second-site suppressor analysis described here.

### Yeast two-hybrid analysis

Yeast two-hybrid studies to analyze Dnm1–Mdv1 and Dnm1 self-interactions were performed in the Y187 *S. cerevisiae* strain background (Takara Bio Inc.) via a growth assay as described previously (Guthrie and Fink, 2002). *pGAD* and *pGBD* plasmid expressing the indicated fusion proteins were cotransformed into the Y187 reporter strain. Interaction between two fusion proteins leads to expression of one of several reporter genes in this strain, allowing the yeast cells to grow on S-dextrose minus histidine or minus adenine. WT Dnm1–Dnm1<sup>InsBmut</sup> interactions were performed in cells co-expressing *GAD-Dnm1* WT and *GBD-Dnm1<sup>InsBmut</sup>*. The Dnm1–Mdv1 interaction was tested in both directions. However, the interaction was only detected when Dnm1 and Mdv1 were fused with the *GAD* and *GBD* domains, respectively.

### Online supplemental material

Table S1 shows a screen for *mdv1* suppressors of *dnm1<sup>F610A</sup>*. Table S2 shows the plasmids used in this study. Fig. S1 shows expression, interaction, and assembly properties of Dnm1 and Mdv1 variants. Online supplemental material is available at <http://www.jcb.org/cgi/content/full/jcb.201207079/DC1>.



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