A conserved mechanism for Bni1- and mDia1-induced actin assembly and dual regulation of Bni1 by Bud6 and profilin

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Abstract

Formins have conserved roles in cell polarity and cytokinesis and directly nucleate actin filament assembly through their FH2 domain. Here, we define the active region of the yeast formin Bni1 FH2 domain and show that it dimerizes. Mutations that disrupt dimerization abolish actin assembly activity, suggesting that dimers are the active state of the FH2 domains. The Bni1 FH2 domain protects growing barbed ends of actin filaments from vast excesses of capping protein, suggesting that the dimer maintains a persistent association during elongation. This is not a species-specific mechanism, as the activities of purified mammalian formin mDia1 are identical to those of Bni1. Further, mDia1 partially complements BNI1 function in vivo, and expression of a dominant active mDia1 construct in yeast causes similar phenotypes to dominant active Bni1 constructs. In addition, we purified the Bni1-interacting half of the cell polarity factor Bud6 and found that it binds specifically to actin monomers and, like profilin, promotes rapid nucleotide exchange on actin. Bud6 and profilin show additive stimulatory effects on Bni1 activity and have a synthetic lethal genetic interaction in vivo. From these results, we propose a model in which Bni1 FH2 dimers nucleate and processively cap the elongating barbed end of the actin filament, while Bud6 and profilin generate a local flux of ATP-actin monomers to promote actin assembly.
Introduction

Formins, a conserved family of proteins defined by formin homology (FH) domains, are required for the establishment and maintenance of cell polarity and cytokinesis (reviewed in Frazier and Field, 1997; Evangelista et al., 2003; Wallar and Alberts, 2003). In the budding yeast *Saccharomyces cerevisiae*, two formin proteins (Bni1 and Bnr1) drive the assembly of actin cables and the cytokinetic ring (Evangelista et al., 2002; Sagot et al., 2002a; Tolliday et al., 2002). In fission yeast, the formin For3p is required for the formation of interphase actin cables (Feierbach and Chang, 2001), and together with the formin Cdc12p is required for the formation and proper constriction of the cytokinetic ring (Chang et al., 1997; Pelham and Chang, 2002). In animal cells, formins are similarly involved in the assembly of specialized F-actin containing structures, such as stress fibers and the cytokinetic ring (Wasserman, 1998; Watanabe et al., 1999; Tominaga et al., 2000; Severson et al., 2002).

Several recent studies have shown that purified carboxyl terminal fragments of the budding yeast formin Bni1 and the fission yeast Cdc12p stimulate actin filament assembly in vitro (Pruyne et al., 2002; Sagot et al., 2002b; Kovar et al., 2003). Both of these formins associate with the barbed ends of actin filaments (Pruyne et al., 2002; Kovar et al., 2003; Pring et al., 2003); however, the mechanism of filament growth may differ. Studies on Bni1 concluded that it forms a specialized barbed end association, acting as a processive cap to allow filament elongation at the barbed end. Studies on Cdc12p concluded that it acts as a standard barbed end capping protein, limiting filament growth to the pointed end, until it is ‘gated’ by profilin to allow growth at the barbed end. Many open questions about formin activities remain. What is the minimal active region of the FH2 domain of formins? What are its biophysical properties? How do formins maintain an association with a barbed end that is growing rapidly? How do formin-associated proteins affect formin activities? What are the activities of different mammalian formins, and are they more similar in function to Bni1 or Cdc12p?

The inherent actin nucleation function of Bni1 is mediated by its FH2 domain in vitro (Pruyne et al., 2002; Sagot et al., 2002b), but regions flanking the FH2 domain contribute to the assembly of polarized actin cables in vivo (Evangelista et al., 2002; Sagot et al., 2002a). The actin monomer binding protein profilin is required in vivo for the assembly of actin cables (Haarer et al., 1990; Wolven et al., 2000; Evangelista et al., 2002). Profilin stimulates formin-induced actin assembly in vitro, which requires its interactions with the FH1 domain and with
actin monomers (Sagot et al., 2002b; Kovar et al., 2003; Pring et al., 2003).

Another potential regulator of the yeast formin Bni1 is Bud6/Aip3, first identified in a two-hybrid screen as an actin interacting protein (Amberg et al., 1995). Bud6 interacts by two-hybrid assay with a carboxyl terminal region of Bni1 (Evangelista et al., 1997; also see Figure 1) and has a similar in vivo localization to Bni1 (Amberg et al., 1997; Ozaki-Kuroda et al., 2001; Segal et al., 2002). Genetic evidence suggests that Bud6 has an important role in cell polarity and actin cable organization (Amberg et al., 1997; Glynn et al., 2001; Evangelista et al., 2002; Sagot et al., 2002a), but the biochemical activities of Bud6 have remained elusive, leaving its exact function unclear.

Here, we show that the Bni1 FH2 domain dimerizes, which has important implications for how formins maintain association with the growing end of an actin filament. Further, the FH2 dimers protect elongating barbed ends from the inhibitory effects of excess capping protein, demonstrating their persistent association with the growing filament ends. Activities similar to Bni1 were observed for mouse formin mDia1 in vitro and in vivo, demonstrating the strong evolutionary conservation of this mechanism. We also report a striking and unexpected parallel between the cellular functions of two Bni1-interacting proteins, Bud6 and profilin. The purified carboxyl terminal half of Bud6 (C-Bud6) binds to actin monomers and dramatically accelerates nucleotide exchange on actin. C-Bud6 and profilin additively stimulate Bni1-mediated actin assembly in vitro, and pfy1-4 and bud6Δ mutations are synthetic lethal in vivo. Thus, central to the formin mechanism in vivo is that two actin monomer binding proteins/actin nucleotide exchange factors interact on either side of the FH2 domain and stimulate formin activity.
Materials and Methods

Yeast strains, media, and DNA manipulations

Standard methods were used for DNA manipulations and for yeast growth, sporulation, and transformation (Rose et al., 1989). For testing mDia1 function in yeast cells (Figure 2), strains previously described were used: \textit{bni1-FH2#1 bnr1Δ} (PY3744) and \textit{bni1Δ bnr1Δ} carrying the plasmid 2\(\mu\) URA3 \textit{BNII} (PY4315) (Sagot et al., 2002b). These and the \textit{bud6Δ} strain are isogenic to the wild type \textit{BY4741} strain from Research Genetics (Invitrogen, Carlsbad, CA). The \textit{pfy1-4} yeast strain (DDY2008) has been described (Wolven et al., 2000).

For mDia1 expression in \textit{E. coli} and yeast, the coding sequence of mDia1, residues 739-1255 (FH2-COOH) or residues 553-1255 (FH1-COOH), were PCR amplified from pEGFP-mDia1 (a gift from T. Fujiwara). The PCR products were subcloned into either pET-GST-TEV (pET derivative for expression in \textit{E. coli}, which has a TEV protease site inserted after GST) or a pRS415 derivative (a \textit{LEU2}, CEN vector with an insertion of the yeast actin (\textit{ACT1}) promoter (Christianson et al., 1992)). For expression of Bni1 fragments in \textit{E. coli}, the appropriate coding sequences were PCR amplified and subcloned into pET-GST-TEV. To generate the FH2(core) ww mutant, site directed mutagenesis was performed using a PCR-based method to generate Trp1363(TGG) to Ala(GCG) and Trp1374(TGG) to Ala(GCC) substitutions. FH2(core) ts contains two mutations (R1528A and R1530A) previously described in Sagot et al. (2002a). For complementation analysis, we used pB1005 (\textit{URA3}, 2\(\mu\)) and pB1006 (\textit{LEU2}, CEN). The latter, a generous gift from J. Pringle and M. Longtine, allows expression of full length \textit{BNII} under the control of its own promoter. \textit{pBNII-mini} (\textit{LEU2}, CEN) encodes a fusion protein between protein A and amino acid residues 1215-1750 of Bni1 under the control of the \textit{ACT1} promoter. For expression of C-Bud6 in \textit{E. coli}, the sequence encoding amino acids 489-788 of Bud6 was PCR amplified from genomic DNA and subcloned into pET-GST-TEV. All DNA constructs were verified by sequencing. Details of the constructs are available upon request.

Imaging of actin organization in yeast cells

Alexa-568-phalloidin staining and imaging of yeast cells was performed as described previously (Sagot et al., 2002a).
**Protein purification**

Rabbit skeletal muscle actin, pyrene-labeled rabbit skeletal muscle actin, and human profilin were purchased from Cytoskeleton (Denver, CO). Yeast capping protein, coflin, and profilin were purified by methods described (Amatruda and Cooper, 1992; Lappalainen and Drubin, 1997; Wolven et al., 2000). Mouse mDia1 and yeast Bni1 proteins were expressed as GST-fusion proteins in the *E. coli* strain BL21-Codon Plus (DE3)-RP (Stratagene), which were grown at 37°C to log phase, then induced by addition of 0.2mM IPTG and grown for 16 hours at 25°C. Cells were pelleted and resuspended in PBS buffer supplemented with a protease inhibitor mixture (final concentration of 1mM PMSF and 0.5µM each of antipain, leupeptin, pepstatin A, chymostatin, aprotinin), then lysed by sonication. The lysate was cleared by centrifugation at 17,000 g, and GST-fusion proteins were purified using glutathione-conjugated agarose beads (Sigma, St. Louis, MO). After washing beads into HEK buffer (20mM Hapes, pH 7.5, 1mM EDTA, 50mM KCl), Bni1 or mDia1 fragments were released by digestion with recombinant TEV protease (Invitrogen, Carlsbad, CA), then further purified by size exclusion chromatography. Peak fractions were concentrated in a Microcon-10 device (Millipore, Billerica, MA). Cleavage of GST from the Bni1 construct FH1-COOH was inefficient, so we used the GST-fusion protein for biochemical assays. The presence of GST on this construct has been shown previously to have no effect on its activity (Pruyne et al., 2002).

The carboxyl terminal fragment of Bud6 (C-Bud6) was purified as a GST fusion protein by methods similar to above. After release from glutathione agarose by digestion with TEV protease, C-Bud6 was purified further on a Mono Q (5/5) anion exchange column (Amersham Pharmacia Biotech, Piscataway, NJ). Proteins were eluted from the column in HEK buffer and a linear gradient of KCl (0.1-0.5M); C-Bud6 eluted at ~250mM KCl. Peak fractions were concentrated and exchanged into HEK buffer by multiple rounds of concentration and dilution in a Microcon-10 device.

**Actin filament assembly and elongation assays**

Actin filament assembly assays were performed essentially as described (Humphries et al., 2002). Briefly, 42µl of monomeric actin (final concentration in reactions 2µM, 1% pyrene-labeled) was mixed with 15µl of HEK buffer or proteins in HEK buffer (Bni1, mDia1, profilin, and/or C-Bud6), then mixed with 3µl of 20x initiation mix (40mM MgCl2, 10mM ATP, 1M
KCl). Pyrene fluorescence was monitored at excitation 365nm and emission 407nm in a fluorescence spectrophotometer (Photon Technology International, Lawrenceville, NJ). In actin filament elongation assays, 35µl preformed actin filaments (1µM final) were mixed with 15µl HEK buffer alone or premixed proteins in HEK buffer. The actin filaments were sheared by five passages through a 27-gauge needle and syringe. After 10 seconds, 20µl of sheared filaments (333nM final) were added to 40µl actin monomers (0.5µM final, 1% pyrene labeled) in F-buffer to initiate filament elongation (10mM Tris, pH7.5, 0.7mM ATP, 0.2mM CaCl₂, 2mM MgCl₂, 50mM KCl, 0.2mM DTT). Pyrene fluorescence was monitored as above. The relative rates of actin assembly and elongation were determined from the slopes of the reaction curves during the early phase (first 200 seconds) of actin polymerization.

**Analytical ultracentrifugation and gel filtration analyses**

Sedimentation equilibration was performed in a Beckman XLA analytical ultracentrifuge using an An-Ti60 rotor and 6-channel cells. 110µl samples (0.2, 0.4, 0.7 mg/ml) were centrifuged at 10,000 and 13,000 rpm for 24 hours at 4°C in HND buffer (20mM Hepes (pH 7.5), 200mM NaCl, 2mM DTT). Absorbance scans were obtained at 280nm. The calculated partial specific volume for FH2(core) was 0.7299 ml/g and buffer density was 1.007. Concentration profiles were analyzed by SEDFIT85 (Lebowitz et al., 2002).

Gel filtration analysis was performed using a Superdex 200 (10/30) column (AP Biotech) equilibrated in HND buffer. Elution peaks were monitored by absorbance at 280nm and verified by SDS-PAGE of column fractions. The size standards used were Aldolase (158kDa), Albumin (67kDa), Focal Adhesion kinase FERM fragment (46kDa), and Chymotrypsinogen A (25kDa).

**Actin monomer binding and nucleotide exchange assays**

To measure rates of nucleotide exchange on actin monomers, 57.2µl G-buffer (10mM Tris pH 7.5, 0.2mM CaCl₂, 0.2mM DTT, no ATP) containing actin (2.5µM final) was mixed with 10µl HEK buffer or proteins in HEK buffer (profilin, coflin, or C-Bud6) and added to 2.8µl etheno-ATP (50µM final). The reaction was monitored at 25°C in a fluorescence spectrophotometer as above. Nucleotide exchange rates were determined from the initial, linear slopes of the curves. For native gel electrophoresis, C-Bud6 and actin (each 10µM final) were mixed for 15 minutes, then fractionated on 7.5% native gels as described (Safer, 1989).
Results

Defining the active region of the Bni1 FH2 domain

Based on the sequence alignment of multiple formins, the boundaries of the Bni1 FH2 domain have been defined previously as amino acid residues 1348-1824, and this region alone induces actin assembly in vitro (Pruyne et al., 2002). To better define the active portion of the FH2 domain, we expressed a large series of constructs spanning residues 1327 to 1953, some of which are shown in Figure 1A. The shortest soluble construct with actin assembly-promoting activity was Bni1 FH2(core) (residues 1348-1750). This fragment had equivalent actin nucleation activity to that of a longer 1344-1824 FH2 domain (Figure 1D, curves 2 and 3). As previously shown for the 1348-1824 FH2 domain (Pruyne et al., 2002), FH2(core) displayed a slightly lower actin nucleation activity than FH1-COOH (compare Figures 1B and 1C; also see Figure 1D). The mutations R1528A and R1530A, which confer thermosensitivity to Bni1 for actin cable formation in vivo (Sagot et al., 2002a), abolished actin nucleation activity of the FH2(core) (Figure 1D, curve 6). This demonstrates that the biochemical activities of the FH2 domain correlate with its in vivo functions.

Analysis of FH2(core) and other Bni1 constructs using a panel of proteases revealed a chymotrypsin cleavage site at Leu1407 (our unpublished results), which led us to generate the FH2Δ fragment (1408-1766). This truncation abolished actin nucleation activity (Figure 1D, curve 4). Sequence alignments across species showed that this 60 amino acid region (1348-1408) contains two highly conserved tryptophan residues located on the carboxyl-terminal side of the FH1 domain (residues 1363 and 1374 in Bni1). Tryptophan to alanine substitutions at these two sites (W1363A/W1374A) to generate the construct FH2(core) ww abolished actin nucleation activity (Figure 1D, curve 5). These data define residues 1348-1750 as a minimal active region of the FH2 domain and indicate that the sequence 1348-1408, and the two conserved tryptophan residues, play crucial roles in actin assembly.

Active Bni1 FH2 domain polypeptides dimerize

In the course of purifying Bni1 FH2(core), we noted that it eluted from a gel filtration column as a single symmetrical peak at approximately the volume expected for a dimer. We therefore measured its molar mass by analytical ultra-centrifugation. Sedimentation equilibrium
analysis of FH2(core) yielded a mass of 93,000g/mol, very close to the mass expected for an
FH2(core) dimer (95,400g/mol) based on its primary sequence (Figure 2A). Using analytical gel
filtration, we compared the elution profiles of FH2(core), FH2Δ, and FH2(core) ww. In contrast
to the active FH2(core) dimers, the inactive FH2Δ construct behaved as a monomer by gel
filtration (Figure 2B). Furthermore, the inactive FH2(core) ww polypeptide was impaired for
dimerization; its migration as a mixed population suggests that this mutant forms an unstable
dimer. These data demonstrate that residues 1348-1407 are critical for dimerization of the FH2
domain and that W1363 and W1374 are required for stable dimerization. There is also a
correlation between ability to form stable dimers and ability to promote actin assembly. Taken
together, these data suggest that the dimer is the functional state of formins. This conclusion is
supported by our observations that a Bni1 fragment similar to the FH2(core) polypeptide

**Bni1 dimers processively cap the growing barbed ends of actin filaments and block the
inhibitory effects of excess capping protein**

Bni1 associates with the barbed ends of actin filaments with an apparent Kd of 20 nM and
reduces only modestly the rate of barbed end elongation (Pruyne et al., 2002; Pring et al., 2003).
The addition of higher concentrations of Bni1 (600nM) fails to reduce further the rate of
elongation, suggesting Bni1 forms a processive cap that maintains association with the barbed
end during filament elongation. As a further test of this mechanism, we assayed the ability of
Bni1 to associate with growing barbed ends of filaments in the presence of capping protein,
which terminates barbed end elongation. We reasoned that if Bni1 maintains a strong and
persistent association with barbed ends during filament growth, it should be able to protect ends
from the inhibitory effects of excess capping protein.

First, we measured the rate of actin filament elongation at barbed ends upon addition of
0.5μM monomeric actin to preformed actin filament seeds (free barbed end concentration
~0.3nM). As expected, yeast capping protein inhibited elongation in a dose-dependent manner
(Figure 3A). Half maximal inhibition occurred at 20nM capping protein, defining this as the
apparent Kd for association with the barbed ends of filaments, and 500nM capping protein
abolished elongation (Figure 3B). The affinity of yeast capping protein for actin filament barbed
ends has not previously been determined, and our data show that it is over an order of magnitude
weaker than mammalian CapZ isoforms (Caldwell et al., 1989; Schafer et al., 1996). These differences in affinity may reflect varying requirements for barbed end dynamic behavior in different organisms and cell types.

This assay monitors the rate of actin filament growth specifically at the barbed ends, as indicated by three lines of evidence. First, capping protein abolishes filament elongation in the assay. Because capping protein terminates elongation specifically at filament barbed ends, we conclude that under these conditions polymerization is limited to barbed ends. Second, addition of 5µM profilin had no effect on the rate of filament elongation (our unpublished results). Since profilin binds to actin monomers and prohibits their addition to the pointed ends of filaments (Pollard and Cooper, 1984), this means that pointed end elongation does not make a detectable contribution in the assay. Third, the total actin concentration in the elongation reactions is 0.8µM, and the critical concentration for growth of actin filament pointed ends is 0.5-0.8µM (Pollard and Cooper, 1986; Rickard and Sheterline, 1986; Pollard et al., 2000). These three points clearly demonstrate that pointed end growth is not being measured.

Next, we tested the ability of Bni1 to protect filament ends from capping protein by pre-mixing filament seeds with a combination of 500nM capping protein and varying concentrations of Bni1 FH2(core), then adding actin monomers to initiate elongation. As shown in Figures 3C and 3D, Bni1 FH2(core) rescued barbed end elongation in a dose-dependent manner, demonstrating that the formin protects filament ends from capping protein while allowing rapid elongation. The rate of elongation of FH2(core)-capped filaments was only slightly reduced compared to uncapped filaments in agreement with previous reports (Pruyne et al., 2002; Pring et al., 2003). In a control experiment, 500nM FH2(core) did not promote filament assembly in the absence of actin filament seeds (Figure 3C), indicating that in this assay, the effects of FH2(core) are specific to elongation and cannot be attributed to nucleation of new filaments.

The half maximal effect for protecting filament ends from 500nM capping protein was 7nM Bni1 FH2(core) (Figure 3D). This indicates a very strong association of the FH2 domain with the barbed end, significantly greater than capping protein, which may provide a key part of the mechanism in cells that allows elongation of particular actin structures (e.g. yeast cables) in the presence of capping protein. These data provide complementary support for the proposed mechanism of Pruyne et al. (2002), that Bni1 maintains a persistent association with the barbed end during filament growth. As shown in Figure 3E, all of the Bni1 polypeptides that were
active for actin nucleation also were active in protecting barbed ends of filaments from capping protein, and polypeptides that were defective in actin nucleation did not show barbed end protection activity. Thus, there is a close correlation between the activities in Bni1 required for actin assembly and the ability of Bni1 to associate with the growing barbed end.

**Bud6 binds to actin monomers and accelerates nucleotide exchange on actin**

In previous studies, we and others have shown that profilin binds to the FH1 domain of Bni1 and stimulates Bni1-induced actin assembly in vitro (Sagot et al., 2002b; Pring et al., 2003). Profilin is also required for formation and maintenance of formin-dependent actin cables in vivo (Haarer et al., 1990; Wolven et al., 2000; Evangelista et al., 2002), suggesting that the cellular mechanism of formin-induced actin assembly involves regulation by formin-associated proteins. Here, we addressed the role of another Bni1-interacting protein, Bud6. Full-length Bud6 expressed in *E. coli* was insoluble and did not re-fold following urea denaturation and dialysis into aqueous buffers. However, we were able to purify a soluble carboxyl terminal fragment of Bud6 (C-Bud6; residues 489-788) that encompasses the domains interacting with actin and Bni1 by two-hybrid assay (Figure 4A) (Amberg et al., 1995; Evangelista et al., 1997).

Bud6 was identified as an actin interacting protein by the two hybrid assay, but it has not previously been shown whether it has biochemical affinity for actin, and if so, whether it prefers actin monomers or filaments. C-Bud6 showed no affinity for actin filaments in co-sedimentation assays (our unpublished results), but interacted with actin monomers in three independent assays. First, C-Bud6 retarded the migration of monomeric actin in native gels (Figure 4B), causing a modest but reproducible shift (see lane 2, Figure 4B). This result is consistent with a dynamic association between C-Bud6 and actin monomers under these conditions, similar to the reported profilin-actin interaction (Mockrin and Korn, 1980). Second, 1.5µM C-Bud6 inhibited the maximal rate of 2µM actin assembly in spontaneous actin assembly reactions, and did so more efficiently than 1.5µM yeast profilin (Figure 4C). This activity is common to many actin monomer-binding proteins and reflects their ability to suppress spontaneous actin nucleation by interfering with G-actin associations required to form polymerization intermediates. Third, and perhaps most significantly, C-Bud6 affected the rate of nucleotide exchange on actin monomers (Figure 4D). Profilin and CAP (cyclase associated protein) are the only two other actin monomer binding proteins previously shown to promote nucleotide exchange on actin (Eads et
al., 1998; Wolven et al., 2000; Moriyama and Yahara, 2002). Many other actin monomer-binding proteins inhibit nucleotide exchange on actin (e.g. cofilin and twinfilin; Lappalainen et al., 1997; Goode et al., 1998). As expected, cofilin inhibited nucleotide exchange and profilin promoted nucleotide exchange on actin. C-Bud6 increased the rate of nucleotide exchange approximately three times more efficiently than profilin. The observation that profilin and Bud6 each have this unique activity and each associate with Bni1 suggests that their cellular functions may be highly related.

_Bud6 and profilin additively stimulate Bni1-mediated actin assembly and share a genetically overlapping function_

Next, we tested the ability of C-Bud6 to affect Bni1-induced actin assembly. Because Bud6 interacts by two-hybrid assay with the carboxyl terminus of Bni1, residues 1647-1954 (Evangelista et al., 1997), we tested the effects of C-Bud6 on Bni1 FH1-COOH, which contains the Bud6 binding domain (see Figure 1A). C-Bud6 reproducibly stimulated the actin assembly activity of 25nM FH1-COOH, with a maximal effect at 50nM C-Bud6 (Figure 5A). At higher concentrations of C-Bud6, the inhibitory effect from actin monomer sequestering reduced the overall rate of actin assembly in a formin-independent manner. Yeast profilin (200nM) also stimulated actin assembly mediated by Bni1 FH1-COOH, consistent with previous reports (Sagot et al., 2002b; Pring et al., 2003). As shown in curve A (Figure 5A), 200nM profilin and 50nM C-Bud6 together reproducibly stimulated the activity of Bni1 FH1-COOH in an additive manner, and importantly, at these concentrations they did not affect the rate of assembly of actin alone (no Bni1 present). Further, C-Bud6 and profilin had no activity on Bni1 FH2(core), which lacks the FH1 domain (profilin binding site) and a portion of the Bud6-binding domain (Figure 5B). Therefore, the ability of C-Bud6 and profilin to stimulate Bni1 activity requires the presence of their known binding sites on Bni1.

Given the similarity in biochemical activities between Bud6 and profilin, we tested their genetic interactions in vivo. A _bud6Δ_ strain was crossed to a strain bearing a thermosensitive allele of the yeast profilin gene (_pfy1-4_) with defects in actin monomer binding and nucleotide exchange (Wolven et al., 2000). The _bud6Δ pfy1-4_ double mutant cells displayed a severe additive growth defect at 25°C (Figure 5C); 22 tetrads were dissected, and 27 out of 27 of the
resulting double mutants showed this phenotype. These genetic data support our biochemical data suggesting that profilin and Bud6 act together to promote formin-dependent actin assembly.

**The mammalian formin mDia1 promotes actin assembly by a similar mechanism to Bni1 and partially complements loss of BNI1 function in vivo**

Recent studies have shown that the budding yeast formin Bni1 (Pruyne et al., 2002; Sagot et al., 2002b) and the *S. pombe* formin Cdc12p (Kovar et al., 2003) directly nucleate actin assembly, but by apparently different mechanisms. To address whether mammalian formins have a similar activity to either Bni1 or Cdc12p, we purified two fragments of the *Diaphanous*-related mouse formin mDia1 (Figure 6A). The fragment FH1-COOH (amino acids 553 to 1255) includes the FH1, FH2, and carboxyl-terminal domains. The fragment FH2-COOH (amino acids 739 to 1255) includes the FH2 and carboxyl-terminal domains. As shown in Figures 6B and 6C, both fragments of mDia1 stimulated the polymerization of mammalian actin in a dose-responsive manner. As previously demonstrated for yeast profilin with Bni1 (Sagot et al., 2002b; Pring et al., 2003), mammalian profilin enhanced the rate of actin polymerization induced by mDia1, and this acceleration required the mDia1 FH1 domain (Figure 6D). Further, mDia1 displayed the same ability to protect growing barbed ends of filaments from termination by excess capping protein (Figure 3E). These data suggest that formin-mediated actin assembly is a highly conserved function.

To investigate conservation of formin activity in vivo, we tested whether mDia1 could promote actin filament assembly in yeast. Previously, it has been shown that yeast cells expressing dominant active Bni1 fragments accumulate excess actin cables, resulting in enlarged, depolarized cells (Evangelista et al., 2002; Sagot et al., 2002a). Similarly, the expression of carboxyl terminal constructs of mDia1 in mammalian cells promotes the assembly of excessive filamentous actin (Copeland and Treisman, 2002). To test mDia1 activity in yeast, we expressed the two mDia1 fragments under the control of a constitutive promoter (the yeast *ACT1* promoter) in a formin temperature-sensitive mutant yeast strain (*bni1-FH2#1 bnr1Δ*) (Sagot et al., 2002a). When grown at the permissive temperature, expression of mDia1 FH1-COOH promoted the assembly of numerous filamentous actin structures (Figure 7A, lower panel). Similar to previous in vivo domain analysis on Bni1 (Evangelista et al., 2002; Sagot et al., 2002a), we found that the FH1 domain of mDia1 was required for these effects. In three independent experiments (n > 250
cells), 40 ± 1 % of the cells expressing mDia1 FH1-COOH displayed this phenotype, compared to less than 1% of cells transformed with a control vector (middle panel). These data show that mDia1 can promote the assembly of supernumerary F-actin structures in vivo, consistent with its biochemical activities described above. Further, expression of the FH1-COOH, but not the FH2-COOH, fragment of mDia1 partially rescued the lethality of a strain deleted for the two yeast formin genes, *BNJ1* and *BNRI* (Figure 7B).
Discussion

*FH2 dimers nucleate actin assembly and processively cap growing barbed ends of filaments*

Bni1 nucleates the initial formation of an actin filament by a mechanism that appears to involve stabilization of otherwise short-lived actin dimers, intermediates in actin polymerization (Pring et al., 2003). After nucleation, formins have been proposed to remain associated with the barbed ends of filaments as they elongate (Pruyne et al., 2002; Pring et al., 2003). A similar mechanism of ‘riding’ the fast-growing ends of actin filaments also has been suggested for Ena/VASP proteins (Bear et al., 2002). However, this model has presented a paradox, because it is difficult to imagine how a protein remains bound to a filament end that is rapidly elongating (insertion of 4-5 actin subunits per second in our assays in Figure 3). Our data shed new light on the mechanism underlying this activity in two important ways. First, we demonstrate that formins protect the barbed ends of growing actin filaments from the inhibitory effects of excess capping protein, which provides new experimental support for the model that Bni1 maintains a persistent (or processive) association with the barbed end during growth. Bni1 and mDia1 both have this activity, which means that this is a conserved formin function. In fact, formins (or possibly Ena/VASP proteins) may underlie a previously reported activity in neutrophil extracts that protects growing barbed ends of filaments from capping protein (Huang et al., 1999). This ability to associate with and protect the growing barbed end of an actin filament from capping protein may be crucial in vivo for generating specific actin structures, such as yeast actin cables that appear to be free of capping protein (Amatruda and Cooper, 1992). Second, we show that Bni1 dimerizes and that ability to dimerize correlates with ability to nucleate actin filament assembly and to protect growing barbed ends from capping protein. Dimerization of the FH2 domain may help to explain how persistent barbed end association is maintained. Each half of the dimer (each FH2 domain) may be capable of making independent contacts with actin subunits exposed at the barbed end of the filament. This would permit new actin subunit insertion while allowing the formin dimer to alternate contacts and thus, walk up the growing filament end (see model below).

The activities of Bni1 and mDia1 differ significantly from those recently reported for the *S. pombe* formin Cdc12p (Kovar et al., 2003). These authors concluded that formins cap actin filament barbed ends in the absence of profilin, preventing all barbed end growth until gated by
profilin to allow barbed end growth. In contrast, we have demonstrated that both Bni1 and mDia1 direct barbed end elongation in the absence of profilin. Further, the addition of profilin does not affect the rate of elongation in the presence of these formins (our unpublished results). These activities are supported by another recent study, which shows that mDia1 activities are similar to those of Bni1 (Li and Higgs, 2003). We do not believe that the differences in activities between Cdc12p and other formins (Bni1 and mDia1) are due to construct variation, because we have purified Cdc12p constructs and observed activities similar to those reported by Kovar and colleagues (J.M., B.G., A. Yonetani, and F. Chang, unpublished results). Thus, Bni1 and mDia1 behave distinctly from Cdc12p in the absence of profilin in vitro. What remains uncertain is the physiological relevance of the differences (profilin-gated versus non-gated). The simplest prediction is that all of the formins discussed above should allow barbed end elongation in vivo, because profilin-bound actin monomers are abundant in most eukaryotic cells. Therefore, unless profilin accessibility is limited under specific cellular conditions or in particular sub-cellular locations, all of these formins may have similar activities in vivo (i.e. promote barbed end assembly of actin filaments).

**Regulation of Bni1 activities by profilin and Bud6**

A central aspect of formin function in vivo is the association of multiple interacting proteins to build large protein complexes that regulate cell polarity and cytokinesis (reviewed in Pruynne and Bretscher, 2000). The known Bni1 associated proteins include profilin, Bud6, eF1α, Spa2, and the Rho-family GTPases Rho1 and Cdc42 (Kohno et al., 1996; Evangelista et al., 1997; Imamura et al., 1997; Fujiwara et al., 1998; Umikawa et al., 1998), and the in vivo mechanism of formin-induced actin assembly likely involves the coordinated activities of these factors. *Diaphanous*-related formins, such as Bni1 and mDia1, are proposed to exist in an autoinhibited conformation until released by binding of Rho GTPases to the amino terminal Rho binding domain (RBD) (see Figure 1) (Watanabe et al., 1999; Alberts, 2001; Palazzo et al., 2001; Li and Higgs, 2003). In budding yeast, this regulation is controlled by distinct Rho signaling pathways acting on Bni1 and Bnr1 at different stages of bud morphogenesis and in response to cell stress (Dong et al., 2003). Rho GTPase activation is thought to release the carboxyl terminus, containing the FH1, FH2, and COOH domains, which stimulate actin assembly. The functions of the carboxyl terminus (FH1-COOH) are likely subject to coordinated regulation by
multiple associated proteins. We and others have shown that profilin, which binds the FH1 domain, stimulates the actin assembly-promoting activity of yeast formins (Sagot et al., 2002b; Kovar et al., 2003; Pring et al., 2003). Here we have extended this analysis to mammalian formins, showing that human profilin stimulates mDia1-induced actin assembly.

In addition, we defined the regulatory role of the formin-associated protein Bud6, which has an established role in cell polarity. Although Bud6 bears no sequence identity to profilin, the carboxyl terminal half of Bud6 (C-Bud6) has similar biochemical activities to profilin, dramatically accelerating nucleotide exchange on actin and enhancing Bni1-induced actin filament assembly. This similar activity suggests that Bud6, like profilin (Schutt et al., 1993), may bind to actin monomers at their barbed ends, promoting nucleotide exchange on actin and leaving free the pointed ends of the monomers for addition to filament barbed ends. Further, the stimulatory effects of Bud6 on Bni1 are additive with those of profilin, which binds to sequences located on the opposite side of the FH2 domain. These activities are consistent with the synthetic lethal genetic interaction we observed between \textit{bud6}Δ and \textit{pfy1-4}. Taken together, these data suggest that Bud6 and profilin have related functions in vivo in stimulating formin-induced actin assembly. This is supported by reports that \textit{bud6} and \textit{pfy1} mutants have similar defects in the formation and organization of formin-nucleated actin cables (Haarer et al., 1990; Amberg et al., 1997; Wolven et al., 2000; Glynn et al., 2001; Evangelista et al., 2002; Sagot et al., 2002a).

\textit{A model for processive capping by formins during actin filament elongation}

Based on our data showing 1) dimerization of the Bni1 FH2 domain, 2) persistent and protective association at the growing barbed ends of actin filaments, and 3) additive stimulation of Bni1 activity by profilin and Bud6, we propose a mechanism for processive capping by formins during filament elongation (Figure 8). At any given time, there are two actin subunits exposed at the barbed end of an uncapped actin filament, one protruding and one recessed. In our model, one of the formins in a dimer contacts the protruding actin subunit, and new actin subunit insertion occurs at the recessed site (left frame in model). This leads to binding of the other formin in the dimer to the newly inserted actin subunit, which is now the protruding end. This event may in turn trigger dissociation of the first formin, exposing the next free site for actin subunit insertion (right frame in model). In this manner, we hypothesize that the formin dimer
walks or teeter-totters up the barbed end of the actin filament, maintaining a stable association while the filament end while allowing actin subunit insertion and filament growth. Key to this mechanism is that the FH2 domain forms a stable dimer, which allows it to maintain steady association with the filament end via alternating contacts of two linked FH2 domains. Bud6 and profilin bind to sequences flanking the FH2 domain and to actin monomers, promoting rapid nucleotide exchange on actin. They facilitate formin-mediated actin filament assembly by generating a local flux of ATP-actin and/or by hand feeding ATP-actin monomers to the open sites at the filament end.

Our data emphasize a functional distinction between the CapZ family of capping proteins and the formins Bni1 and mDia1. Kovar et al. (2003) hypothesized that formins may fold like capping protein to terminate barbed end growth, based on the activities of Cdc12p and suggested sequence homology between capping protein and formin FH2 domains. However, our alignments show no significant sequence homology between formin FH2 domains and capping protein. Further, our recently solved crystal structure of the Bni1 FH2 domain reveals a unique structure that bears no resemblance to that of capping protein (Yamashita et al., 2003), but instead is a flexibly tethered dimer, consistent with our model (Y.Z., I.S., J.M., B.G., D.P., and M.E., submitted).

One key future challenge is to define the activities and regulation of formins from diverse species in the context of larger regulatory complexes. We have demonstrated that an FH2 domain-containing fragment of the mammalian formin mDia1, like Bni1, nucleates actin filament assembly and processively caps the barbed ends of filaments during elongation. Further, mDia1 partially complements yeast formin function in vivo. We speculate that many of the mechanisms for regulating formin-dependent actin assembly also may be conserved. Most formin homologues contain a profilin-binding FH1 domain, located on the amino terminal side of the FH2 domain. Many formins also contain a domain on the carboxyl terminal side of the FH2 domain, but the function(s) of these sequences are less well understood. Here we defined a regulatory role for Bud6, which binds to the carboxyl terminal extension of Bni1. Bud6 homologues are described in other fungi but no clear functional homologue has been identified in animals. Interestingly, database searches here and in a previous study (Glynn et al., 2001) identified a region of homology (22% identity, 46% similarity over 236 amino acids) between residues 517-753 of Bud6 and residues 501-734 of human p160 ROCK serine/threonine kinase
(see supplementary material). Moreover, ROCK kinase has been linked to formin activity in vivo (Nakano et al., 1999; Watanabe et al., 1999). The vertebrate formin Daam1, which has an important role in Wnt signaling, interacts through its carboxyl terminal extension with Disheveled, which binds to actin (Torres and Nelson, 2000; Habas et al., 2001; Capelluto et al., 2002). Thus, it will be important to determine if these and other formin binding partners have functions related to Bud6.
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**Figure legends**

**Figure 1:** Mapping the core activity of the Bni1 FH2 domain. (A) Schematic diagram of Bni1 domain organization and Bni1 polypeptides tested. FH: Formin Homology; GBD: Rho GTPase Binding Domain; BD: Binding Domain; DAD: Diaphanous Autoregulatory Domain; NA: Not Applicable. The bars in constructs 5 and 6 mark the residues mutated (W1363A and W1374A for construct 5; R1528A and R1530A for construct 6). (B-C) 2µM Actin (1% pyrene-labeled) was assembled in the presence of the indicated concentration (nM) of (B) Bni1 GST-FH1-COOH or (C) Bni1 FH2(core). (D) 2µM Actin (1% pyrene-labeled) was assembled in the presence of 100nM of each Bni1 construct: FH1-COOH (curve 1: closed diamonds); FH2 (curve 2: open squares); FH2(core) (curve 3: closed triangles); FH2(core) ts mutant (curve 6: open circles); FH2(core) ww mutant (curve 5: open triangles); FH2Δ (curve 4: crosses). (E) Coomassie stained SDS-PAGE gel showing purified Bni1 fragments (100-200ng each). Lanes correspond to construct numbers in (A).

**Figure 2:** Dimerization of Bni1 FH2 domain polypeptides. (A) Sedimentation equilibrium analysis of the Bni1 FH2(core) polypeptide (0.4 mg/ml) by analytical ultracentrifugation. The data points define a concentration distribution expected for a single species of 93,000g/mol (solid black line), confirming the dimeric nature of the Bni1 FH2 domain. The lower panel shows the distribution of the residuals for the single exponential fit. (B) Analytical gel filtration analysis of Bni1 polypeptides. Bni1 FH2(core) elutes at 13.3 ml, suggesting a molecular weight of 100kDa (consistent with dimerization). Bni1 FH2Δ elutes at 14.0 ml, corresponding to a molecular weight of 50kDa (consistent with a monomer). Bni1 FH2(core) ww elutes at two different peaks, one at a dimer position (13.3 ml) and one at a monomer position (14.0 ml).

**Figure 3:** Bni1 protects the barbed ends of actin filaments from capping protein. (A) 0.5µM monomeric actin (1% pyrene labeled) was polymerized at the barbed ends of mechanically sheared actin filament seeds (333nM) in the presence of the indicated concentrations (nM) of yeast capping protein. Complete inhibition of actin filament elongation was reached at 500nM capping protein. (B) The data from (A) were graphed as rate of filament elongation versus concentration of capping protein. Half-maximal inhibition of elongation occurred at 20nM
capping protein. (C) Bni1 FH2(core) protects filament barbed ends from 500nM capping protein in a dose-responsive manner. 0.5μM actin monomers were assembled as above on actin filament seeds sheared in the presence of 500nM capping protein and the indicated concentrations (nM) of Bni1 FH2(core). The unlabeled curve at the bottom, paralleling the “no seed” curve, demonstrates that 500nM FH2(core) has no effect on the rate of assembly of 0.5μM actin in the absence of filament seeds. (D) The rates of filament elongation from (C) were plotted versus the concentration of Bni1 FH2(core). Half-maximal protection occurred at 7nM Bni1 FH2(core). (E) Rates of filament elongation, calculated as in C, for different Bni1 and mDia1 constructs in the presence of 500nM capping protein. Shaded labels indicate reactions that contain 500nM capping protein.

**Figure 4:** The carboxyl terminus of Bud6/Aip3 binds to actin monomers and promotes nucleotide exchange. (A) SDS-PAGE gel stained with Coomassie blue showing purified proteins used for these experiments: Lane 1, 100ng C-Bud6 (residues 489-788); Lane 2, 100ng yeast profilin; Lane 3, 200ng yeast cofolin. (B) Coomassie stained native gel showing the interaction of C-Bud6 with actin monomers: Lane 1, 10µM C-Bud6; Lane 2, 10µM C-Bud6 and 10µM actin monomers; Lane 3, 10µM actin monomers. (C) Actin monomer binding proteins profilin and C-Bud6 reduce the rate of actin assembly. The rates of actin assembly (2µM actin, 1% pyrene labeled) alone and in the presence of 1.5µM profilin or 1.5µM C-Bud6 were compared during the early linear phase of polymerization. (D) Rate of nucleotide exchange on 2.5µM actin monomers in the presence of varying concentrations of C-Bud6, profilin, or cofolin. The rate of nucleotide exchange was determined from the change in etheno-ATP fluorescence upon binding to actin (see Methods).

**Figure 5:** C-Bud6 and profilin enhance Bni1-induced actin assembly in an additive manner. (A) Actin (2µM, 1% pyrene labeled) was assembled in the presence of 25nM FH1-COOH (curve D), 25nM FH1-COOH + 200nM profilin (curve C), 25nM FH1-COOH + 50nM C-Bud6 (curve B), or 25nM FH1-COOH + 200nM profilin + 50nM C-Bud6 (curve A). Control curves (E-H) show assembly of 2µM actin alone, 2µM actin + 200nM profilin, 2µM actin + 50nM C-Bud6, and 2µM actin + 200nM profilin + 50nM C-Bud6, respectively. (B) Actin was assembled as in (A), in the presence of 100nM FH2(core) (curve A), 100nM FH2(core) + 200nM profilin (curve B),
100nM FH2(core) + 50nM C-Bud6 (curve C), 100nM FH2(core) + 200nM profilin + 50nM C-Bud6 (curve D), 200nM profilin (curve E), 50nM C-Bud6 (curve F), 200nM profilin + 50nM C-Bud6 (curve G), and actin alone (curve H). (C) Genetic dissection of parental diploid strains resulting from two crosses: PFY1 x bud6Δ and pfy1-4 x bud6Δ. Parental strains were sporulated and tetrads were dissected. Growth of the resulting haploid colonies after three days of growth at 25°C on rich medium is shown for representative tetrads. Double mutant pfy1-4 bud6Δ cells showed little or no colony growth (arrows).

Figure 6: mDia1 promotes actin assembly and is activated by profilin. (A) Coomassie stained SDS-PAGE gel showing purified mDia1 fragments. Lane 1: mDia1 FH1-COOH (200ng). Lane 2: mDia1 FH2-COOH (200ng). (B) 2µM actin (1% pyrene-labeled) was assembled in the presence of the indicated concentrations (nM) of mDia1 FH2-COOH. (C) Rate of actin assembly plotted versus concentration of mDia1 FH1-COOH or mDia1 FH2-COOH. Rates of actin assembly were calculated from the slopes of the assembly curves during the initial rapid phase of growth and standardized against rate of assembly of actin alone. (D) Rate of actin assembly mediated by 25nM mDia1 FH1-COOH or mDia1 FH2-COOH in the presence and absence of 300nM human profilin. Rates were normalized as above to the rate of assembly of actin alone.

Figure 7: Effects of expressing mouse formin mDia1 in yeast cells. (A) mDia1 FH1-COOH promotes actin cable formation in yeast. Panels show Alexa-phalloidin staining for wild type cells (upper panel), bni1-FH2#1 bnr1Δ cells carrying a control vector (middle panel: note that actin cables are fainter and less numerous than in wild type cells), and bni1-FH2#1 bnr1Δ cells expressing mDia1 FH1-COOH (lower panel). All cells were grown at the permissive temperature of 24°C. Bar is 2µm. (B) Expression of mDia1 FH1-COOH partially rescues the lethality of a bnr1Δ bni1Δ strain. Left: serial dilutions of bni1Δ bnr1Δ cells spotted on selective medium, where both the 2µ BNI1 URA3 plasmid and the LEU2 plasmids designated on the left are maintained. Right: the same cells spotted onto medium containing 5-fluoroorotic acid to select against the 2µ BNI1 URA3 plasmid. From top to bottom, cells are transformed with a LEU2 control vector, a plasmid expressing full length BNI1, a plasmid expressing Bni1-mini (amino acid residues 1215-1750; (Sagot et al., 2002b)), a plasmid expressing mDia1 FH2-
COOH, and a plasmid expressing mDia1 FH1-COOH.

**Figure 8:** A model for how Bni1 and possibly other formins processively cap the elongating barbed end of an actin filament (see Discussion for details). FH2 domains are depicted as shaded ovals and are labeled ‘Bni1’. FH1 domains are shown as black lines that bind to profilin (labeled ‘P’). The carboxyl terminal regions of Bni1 are shown as black lines that bind to Bud6 (darkly shaded hexagons). ADP-actin subunits are shaded, and ATP-actin subunits are white.

**Figure S1 (Supplemental):** (A) Schematic diagram of domain structures for ROCK and Bud6. CC = predicted coiled coil domain. PH = Pleckstrin Homology domain. The region of sequence homology (22% amino acid identity and 46% similarity) is located between the two dashed lines. (B) Alignment of the primary sequences in the region of homology between budding yeast (*S. cerevisiae*) Bud6 and human ROCK. ClustalW analysis of amino acid sequences was performed, introducing gaps (-) to optimize the alignment. Residues that are similar are boxed, while residues that are identical are boxed and shaded.
A

1. FH1-COOH
2. FH2
3. FH2(core)
4. FH2∆
5. FH2(core) ww
6. FH2(core) ts

Bni1 FH1-COOH

Bni1 FH2(core)

Actin Nucleation
Activation by Profilin

+++ +
++ -
++ -
- NA
- NA
- NA

Actin Nucleation
Activation by Profilin

+++ +
++ -
++ -
- NA
- NA
- NA

Bni1 FH1-COOH

Bni1 FH2(core)
A

Residuals Absorbance (A.U.)

Absorbance (A.U.)

Residuals

Radius (cm)

B

Absorbance (280nm)

Absorbance (280nm)

Elution volume (ml)

FH2(core), 47.7kDa

FH2², 41.5kDa

FH2(core)ww, 47.7kDa

158 67 46 25 kDa
Figure A shows the fluorescence over time for different concentrations of capping proteins. Figure B displays the elongation rate against the concentration of capping protein. Figure C illustrates similar data with an added condition of seeds. Figure D presents the elongation rate against the concentration of Bni1 FH2(core). Figure E provides a bar chart comparing the elongation rates under various conditions, including +500nM capping protein.
### Figure A

A gel electrophoresis showing bands at different positions, labeled as C-Bud6 and actin.

### Figure B

A densitometric scan showing shifted actin and actin bands.

### Figure C

A bar graph showing the rate of actin assembly in different conditions:
- **Actin**
- **Actin + profilin**
- **Actin + C-Bud6**

### Figure D

A plot showing the nucleotide exchange rate with concentration of actin binding protein (μM):
- **C-Bud6**
- **profilin**
- **cofilin**
A

wild type

bni1-FH2#1 bnr1Δ

bni1-FH2#1 bnr1Δ

p mDia1 FH1-COOH

B

+p BNI1 URA3 - p BNI1 URA3

vector

p BNI1

p BNI1-mini

p mDia1 FH2-COOH

p mDia1 FH1-COOH

bni1Δ bnr1Δ