Kinetics of Endophilin N-BAR Domain Dimerization and Membrane Interactions*

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The recruitment to plasma membrane invaginations of the protein endophilin is a temporally regulated step in clathrin-mediated endocytosis. Endophilin is believed to sense or stabilize membrane curvature, which in turn likely depends on the dimeric structure of the protein. The dynamic nature of the membrane association and dimerization of endophilin is thus functionally important and is illuminated herein. Using subunit exchange Förster resonance energy transfer (FRET), we determine dimer dissociation kinetics and find a dimerization equilibrium constant orders of magnitude lower than previously published values. We characterize N-BAR domain membrane association kinetics under conditions where the dimeric species predominates, by stopped flow, observing prominent electrostatic sensitivity of membrane interaction kinetics. Relative to membrane binding, we find that protein monomer/dimer species equilibrate with far slower kinetics. Complementary optical microscopy studies reveal strikingly slow membrane dissociation and an increase of dissociation rate constant for a construct lacking the amphipathic segment helix 0 (H0). We attribute the slow dissociation kinetics to higher-order protein oligomerization on the membrane. We incorporate our findings into a kinetic scheme for endophilin N-BAR membrane binding and find a significant separation of time scales for endophilin membrane binding and subsequent oligomerization. This separation may facilitate the regulation of membrane trafficking events.

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§The abbreviations used are: N-BAR, Bin/amphiphysin/Rvs domain with N-terminal amphipathic helix; AF-488, Alexa Fluor® 488; PB, Pacific Blue™; DOPC, 1,2-dioleoyl-sn-glycero-3-phosphocholine; DOPG, 1,2-dioleoyl-sn-glycero-3-phosphoglycerol; DPH-PC, 2-(3-(diphenylhexatrienyl)propargyl)-1-hexadecanoyl-sn-glycero-3-phosphocholine; FCS, fluorescence correlation spectroscopy; GUV, giant unilamellar vesicle; PI(4)P, L-α-phosphatidylinositol (4)-phosphate; PI(4,5)P, L-α-phosphatidylinositol (4,5)-bis-phosphate; DMSO, dimethyl sulfoxide; SEC, size exclusion chromatography.
ates with membranes as a dimer in an effective two-step process leading to oligomerization even in the absence of pronounced membrane deformation. We assess this mechanistic hypothesis as follows. The evaluation of N-BAR dimerization in solution serves as a prerequisite of our mechanistic analysis of membrane binding. We employ stopped-flow and microscopy-based techniques for kinetic studies. We find evidence for an important contribution of protein oligomerization on the membrane in the mechanism. We arrive at a kinetic model that is fitted to the experimental data. Finally, we discuss the physiological relevance of the separation of time scales for N-BAR membrane interactions observed here.

**EXPERIMENTAL PROCEDURES**

**Materials**—Lipids 1,2-dioleoyl-sn-glycero-3-phosphocholine (DOPC), 1,2-dioleoyl-sn-glycero-3-phospho-(1'-rac-glycerol) (DOPG), 1-α-phosphatidylglycerol (4)- and (4,5)-bis-phosphate (PI(4)P and PI(4,5)P₂; brain, ammonium salts) were obtained from Avanti Polar Lipids (Alabaster, AL). Folch Fraction I was from Sigma. 2-(3-(Diphenylhexatrienyl)propanoyl)-1-hexadecanoyl-sn-glycero-3-phosphocholine (DPH-PC), Pacific Blue™ (PB) C₅-maleimide, Alexa Fluor® 488 (AF-488) C₅-maleimide, and Texas Red 1,2-dihexadecanoyl-sn-glycero-3-phosphothanolamine (triethylammonium salt) were from Invitrogen/Life Technologies. Casein was from Fisher Scientific. Tris(2-carboxylethyl)phosphine was from Pierce/Thermo Fisher Scientific. All commercial reagents were used without further purification.

**Membrane Preparation**—Extruded vesicles were prepared by standard techniques (31). Chloroform solutions of lipid stock were gently evaporated, by using compressed air, from a round-bottom flask, which was subsequently evacuated. Vesicle compositions are represented as the molar fraction DOPG, with DOPC comprising the remaining portion. DPH-PC was included at 0.7 mol % for stopped-flow Förster resonance energy transfer (FRET) studies. Lipids were hydrated at 2 mg/ml with 20 mM HEPES buffer with indicated NaCl concentrations, pH 7.4, under bath sonication at 40 °C for 10–15 min. Lipid dispersions were extruded 13 times through single polycarbonate membranes (Whatman/GE Healthcare). Dynamic light scattering consistently revealed an average hydrodynamic radius of 75 ± 15–30 nm for all lipid compositions utilized in stopped-flow studies. For tubulation assays, 400-nm pore sizes were used. Phosphate concentrations of extruded preparations were determined by standard analysis (32). Giant unilamellar vesicles (GUVs) were prepared by electroformation (33) as described (34).

**Plasmids and Protein Preparation**—A plasmid encoding rat endophilin A1, kindly provided by P. De Camilli, served as the template for generation of a single Cys construct comprising residues 1–247 with C108A/E241C substitutions (N-BAR_C241), ΔNH (BAR) (residues 33–247 with and without the A66W substitution), and N-BAR F10W. We employed sequential QuikChange (Stratagene) reactions for mutagenesis. A plasmid encoding residues 1–247 of human endophilin A1 with the A66W mutation in the pGEX6p vector (GE Healthcare) was kindly provided by N. Mochizuki (24) and served as the template for N-BAR_C234 (with A66W/C108S/Q234C substitutions). Relative to rat, human N-BAR differs by an M133I substitution. A plasmid encoding residues 1–247 of rat endophilin A1 with the C108S/A247C substitutions (3) (N-BAR_C247) was kindly provided by R. Langen. All sequences were verified by DNA sequencing.

GST fusion proteins were purified from the soluble portion of bacterial lysates (BL21(DE3) RIL CodonPlus, Stratagene) using glutathione affinity. Fusion proteins were cleaved at the GST moiety using PreScission protease or thrombin. BAR domains were purified further by ion exchange with a linear NaCl gradient and size exclusion chromatography (SEC) (Superdex200 (GE Healthcare)), concentrated, aliquoted, and flash-frozen in liquid nitrogen. Purity was assessed by SDS-PAGE and analytical SEC. For measurements with thawed samples, we used ultracentrifugation supernatant portions. Concentrations were determined by Bradford analysis using bovine serum albumin (Thermo Fisher Scientific) as a standard (35) and absorption at 280 nm. Concentrations indicated refer to total N-BAR in terms of monomeric units. Labeling was conducted at introduced Cys residues for N-BAR_C241 and N-BAR_C247 or endogenous Cys108 for ΔNH (BAR) (both with and without the A66W substitution). 4-fold excess of maleimide dye reagent was used via a DMSO solution for reaction at 4 °C, in 150 mM NaCl, 20 mM HEPES, 1 mM tris(2-carboxyethyl)phosphine, pH 7.4 solution (HNTcep buffer). Commonly, glutamate/arginine/HEPES, pH 7.4, was added to promote protein solubility during labeling (36). Reactions were quenched with excess dithiothreitol (DTT), and excess dye reagent was removed via three 5-ml HiTrap desalting columns (GE Healthcare) connected in series.

**Liposome Tubulation Assay**—Endophilin N-BAR variants were incubated with 400-nm membrane-extruded vesicles at room temperature for 30 min. Samples were applied to carbon/Formvar-supported copper grids (Electron Microscopy Sciences, Hatfield, PA). After a 5-min incubation, excess material was removed by blotting on filter paper. Grids were then stained with 2% (w/v) uranyl acetate for 2 min, washed in buffer, and air-dried at room temperature. Grids were observed with a JEM 1011 transmission electron microscope (JEOL) with the accelerating voltage set to 100 kV.

**Protein Cross-linking**—Site-specific irreversible cross-linking by reaction of Cys at residue 241 (Cα-Cα distance, 38 Å) was conducted with Bis-MAL-dPEG®11 (Quanta BioDesign, Powell, OH; linker length, 30 Å). Reactions proceeded for 4 h at 4 °C with 20 μM protein in HNTcep buffer with equimolar cross-linker added via a DMSO solution. Reactions were quenched using 2-fold excess DTT, and unreacted reagent was removed using a HiTrap desalting (GE Healthcare) column achieving buffer exchange for stopped flow and SEC.

**Size Exclusion Chromatography**—N-BAR samples in HNTcep buffer were run at 4 °C over a Superdex200 (GE Healthcare) column pre-equilibrated with the same buffer. The retention volume at peak absorption value (V) was used to calculate the distribution coefficient according to:

\[ K_{d} = \frac{V - V_0}{V_I - V_0} \]
Dimerization and Membrane Binding Kinetics of Endophilin N-BAR

where the column void $V_0$ and total volume $V_t$ were determined using blue dextran (GE Healthcare) and acetone, respectively.

Fluorescence Correlation Spectroscopy (FCS)—Measurements were carried out at 20 °C in glass-bottomed chambers (Nunc, Rochester, NY) passivated with a polylysine-conjugated polyethylene glycol solution. The concentration of labeled protein (N-BAR_C247-AF-488) was held constant (between 7 and 17 nM) for each titration. The total protein concentration was varied from 40 nM to 40 μM via the addition of unlabeled protein. The concentration of unlabeled protein was determined by absorption, and the concentration of labeled protein was determined by FCS. For measurements following a 24-h incubation, DTT was refreshed.

Details of the instrument used for FCS measurements have been described (37). Laser power was adjusted to 5.3–5.6 microwatts (prior to entering the microscope).

For each concentration of protein, 30 traces of 10-s duration were measured. The curves were averaged and analyzed as representing a single diffusing species in three dimensions. The ratio of radial to axial dimensions of the observation volume was determined from measurements of a solution of free AF-488 and was fixed for the protein measurements.

Fluorescence Anisotropy Decay—Experiments were carried out at the University of Pennsylvania Ultrafast Optical Processes Laboratory using time-correlated single photon counting instrumentation described in Ref. 38, at ambient temperature with N-BAR_C241-PB in HNTcep buffer with 0.05 mM phenylmethylsulfonyl fluoride, using 405 nm excitation. Samples were prepared to desired concentrations at 4 °C 18 h before measurements. For 1 μM, a sample prepared 60 h prior to measurements yielded results indistinguishable from the 18-h sample (data not shown).

Fluorescence Spectroscopy—Measurements employed a Cary Eclipse fluorometer with a Peltier-controlled temperature block. Donor (PB) and acceptor (AF-488) N-BAR_C241 conjugates (D and A, respectively) were examined with a fluorescence ratio of 1:2, chosen to optimize FRET. For kinetic studies, a temperature of 27 °C was selected to accelerate subunit exchange relative to 22 °C. D and A samples in HNTcep buffer were adjusted to identical concentrations and separately preincubated at the experimental temperature for a duration equal to that of the measurement. Protein adsorption to cuvette walls was <5%, as confirmed by excitation spectra (not shown) in the acceptor emission peak region (519 nm), at the start and end of kinetic monitoring. To achieve this, quartz cuvettes were treated with casein solution for 1 h and washed once with sample buffer. The casein solution comprised the soluble portion of a 2.5 mg/ml suspension in sample buffer following sedimentation.

The dimer dissociation rate $k_{dissn}$ was determined from fluorescence emission time courses monitoring donor quenching relative to the initial time point (right after mixing D and A solutions) by single-exponential fitting (39). The reaction scheme for subunit exchange yields the rate of heterodimer (DA) formation

$$\frac{d[DA]}{dt} = -k_{dissn}[DA] + k_{assn}[D][A]$$  
(Eq. 2)

where $k_{dissn}$ is the dimerization rate constant for association of monomers. In this mixing experiment, the concentration of monomers is unchanged. Hence, the second term in Equation 2 is constant, and a single exponential with time constant $k_{dissn}$ results as the solution to Equation 2 (40).

For equilibrium measurements, the pre-equilibration period for acceptor-donor mixtures was selected as 12 h at 27 °C, as guided by the kinetics results. Emission measurements at the acceptor wavelength upon excitation of the donor (400 nm) were processed by the following previously reported routine (41, 42). Briefly, acceptor-only and donor-only protein conjugate reference standards enabled accounting for emission arising from donor bleedthrough and acceptor cross-excitation. All spectra were buffer-subtracted. Emission spectra were first corrected for acceptor cross-excitation contribution, as determined by comparing excitation spectra at the acceptor emission wavelength of the standard and sample. Subsequently, the contribution of the donor to measured emission in the acceptor peak region was subtracted, as determined from the donor-only standard equalized to the donor-acceptor mixture sample spectrum at the donor emission wavelength (455 nm). Finally, the values were referenced to that observed at the highest protein concentration used in the titration, yielding relative FRET, $F$, proportional to the dimer fraction of the sample, and fitting was conducted of

$$F = \frac{1}{4c_T}(4c_T + K_D - \sqrt{K_D^2 + 8c_TK_D})$$  
(Eq. 3)

where $c_T$ is the total protein concentration, and $K_D$ is the equilibrium dimerization dissociation constant (41).

Stopped Flow—Measurements were carried out with an Applied Photophysics (Surrey, UK) SX.18MV stopped-flow spectrometer (43) using excitation at 280 nm and collecting emission after passage through a 400-nm long-pass filter. Between two and six traces were averaged for each condition, and 50 μl each of protein and lipid solutions were mixed. For light scattering measurements, we used an incident wavelength of 430 nm, and lipid-only traces were subtracted from protein-lipid mixing traces prior to fitting. Data before the dead time of 2 ms were excluded from fitting. All measurements were conducted at 22.0 ± 0.05 °C, in HEPES, pH 7.4, with varying NaCl concentrations as indicated for particular experiments. For concentration variations analyzed in conjunction with microscopy data (see below), 33 mM NaCl was selected as opposed to physiological levels to improve signal-to-noise ratios.

For global analysis of progress curves incorporating off-rates from microscopy experiments (see “Results”), the reaction scheme (44) detailed below yielded differential equations (Equations 4–7) characterizing chemical species as dimeric N-BAR D, the membrane as L, the intermediate as $D^*L$ (where

\[ A, \text{ protein conjugated to acceptor fluorophore (AF-488); } D, \text{ protein conjugated to donor fluorophore (PB); } \]

\[ k_{dissn} \text{ and } k_{dissn} \text{ are protein dimer association and dissociation rates; } k_{dissn} \text{ is a protein dimerization equilibrium constant; } k_1 \text{ and } k_2 \text{ are rate constants for membrane binding and unbinding of non-oligomerized N-BAR; and } k_3 \text{ and } k_4 \text{ are oligomerization and disassembly rates for protein subunits on the membrane surface.} \]
the asterisk is meant to represent an inserted species), and oligomerized species as \( D_2^*L \), treated as a dimer of N-BAR dimers represented as \( D_2^*L \).

\[
\frac{d[D]}{dt} = -k_1[L][L] + k_{1-}[D^*L] \\
\frac{d[L]}{dt} = -k_1[D][L] + k_{1-}[D^*L] \\
\frac{d[D^*L]}{dt} = k_2[L][D^*L] - k_{1-}[D^*L] - 2k_2[L]_0[D^*L]^2 + 2k_{L^*}[D_2^*L] \\
\frac{d[D_2^*L]}{dt} = \frac{k_2}{[L]^0}D^*L]^2 - k_{L^*}[D_2^*L]
\]

Terms for which \( k_2 \) appears as a coefficient (related to the membrane-bound bimolecular reaction) are divided by \([L]_0\), the initial vesicle concentration in terms of total lipid used in the mixing experiment. This approach considers the concentration of membrane-bound species in terms of a dimensionless density. Equations 4–7 were numerically integrated and fitted to experimental data employing MATLAB (The MathWorks, Natick, MA).

Microscopy-monitored Protein Dissociation—Protein and GUVs containing 25% DOPG were incubated in ~180 \( \mu \)l of an equimolar solution of sucrose and glucose, pH 7.4, containing HEPES and varying NaCl concentration, with the same osmolarity as the prepared GUV dispersion (measured with an Advanced Instruments Inc. (Norwood, MA) micro-osmometer) in a polypropylene tube for 1 h.

Imaging chambers (2 mm thick) were formed from two coverslips (24 \( \times \) 40 mm) overhanging a glass microscope slide. The protein-GUV mixture was added to the chamber on a surface pretreated with 2 \( \mu \)l of 2.5 mg/ml casein, 20 mm Tris, and 2 mm EDTA.

Micropipettes were prepared as described (34) and treated with the casein solution applied to the chamber. Transfer capillaries (outer/inner diameters: 1.5 mm/1.12 mm, World Precision Instruments Inc., Sarasota, FL) were cut to desired lengths with a ceramic glass cutter and equipped with a rubber O-ring as a handle.

GUVs were aspirated in micropipettes at constant aspiration pressure (less than 60 pascals). The transfer capillary was manually positioned as to enclose the aspirated GUV. Protein dilution was achieved by transfer of the GUV using a motor-controlled micromanipulator from the protein chamber into an adjacent chamber, composed typically of ~400 \( \mu \)l of buffer solution, under the protection of the transfer capillary (see Fig. 4A, below). Finally, the transfer capillary was removed to expose the protein-covered GUV.

The protein dissociation process was monitored by using a confocal fluorescence microscope (34). The protein fluorescence intensity was determined by fitting a Gaussian ring to the GUV contour (excluding the aspirated region) using MATLAB.

RESULTS

Subnanomolar Dimerization Affinity—We first asked the question whether the membrane binding mechanism of endophilin involves a dimerization step. This hypothesis is supported by previous reports of low dimerization affinities for this protein (16, 35, 45, 46).

Because the measurement of affinities requires establishment of monomer/dimer equilibria, we first designed an experiment to accurately assess N-BAR dimerization kinetics using FRET (Fig. 1A). A single Cys N-BAR construct, N-BAR_C241, was separately labeled with donor (PB) and acceptor (AF-488) fluorophores. We showed that this mutant exhibits functionality in liposome tubulation in labeled and unlabeled form (Fig. 1, B–E).

To uncover the equilibration time for dimer dissociation at 27 °C, we monitored FRET (Fig. 2A), characterizing dimer dissociation kinetics for N-BAR_C241. We selected this optimized variant as N-BAR_C247 (see “Experimental Procedures”) exhibited low (although detectable) FRET, hindering quantitative analysis, whereas N-BAR_C234 showed appreciable FRET but poor tubulation functionality (data not shown). We determined a value for \( k_{dissn} \) of 1.4 \( \pm \) 0.3 \( \times \) 10⁻⁴ s⁻¹ at 27 °C (means \( \pm \) S.E. of five measurements) from fitting FRET time traces such as those shown in Fig. 2B with a model resulting from Equation 2.
We excluded putative “proximity” FRET due to random localization of fluorophores, based on measurements with free dyes in solution (Fig. 2C). Our interpretation of the FRET signal as reporting on protein dimerization (Fig. 1A) is further substantiated by the elimination of FRET upon SDS-induced dimer dissociation (Fig. 2D).

Knowledge of the kinetics determined above allowed us to equilibrate endophilin samples to study dimerization equilibria at 27 °C (Fig. 2, E and F). We determined a $K_D$ at 27 °C of $0.41 \pm 0.15 \text{nm}$ (mean ± S.E. of three measurements exemplified in Fig. 2E) by fitting FRET data with Equation 3. The determined $K_D$ value corresponds to a substantially higher affinity than a value resulting from published measurements at reduced temperature (10 μM) (16). Additional transitions at protein concentrations much higher than $K_D$ were not revealed by FRET measurements (Fig. 2F).

Protein cross-linking (Fig. 3, A and B), SEC (Fig. 3, B and C), time-resolved fluorescence anisotropy (Fig. 3D), and FCS measurements (Fig. 3E) are also inconsistent with 10 μM dimerization affinity. SEC of 2 μM N-BAR under conditions of the subunit exchange FRET studies (Fig. 3, B and C) reported a Stokes radius in agreement with that calculated for dimeric N-BAR, using HYDROPRO (47), as opposed to monomeric N-BAR. In support of this reasoning, we note that HYDROPRO analysis outputs a radius of gyration of 35 Å for dimeric N-BAR in agreement with the experimental value of 35.8 Å (48). Although a nearly 2-fold change in rotational correlation time is predicted comparing N-BAR monomer and dimer (using HYDROPRO (47)), indistinguishable values are observed for 1 and 16 μM samples (Fig. 3D). Similarly, the diffusion coefficients of monomeric and dimeric N-BAR are expected to differ, although indistinguishable diffusion times are reported by FCS over a range of concentrations (Fig. 3E). Hence, these data do not report a monomer-dimer transition occurring in the concentration regime investigated. We next aimed to elucidate the N-BAR membrane binding mechanism through measurements of endophilin N-BAR membrane binding and unbinding kinetics.

Dissociation from Single GUVs Reports Long Membrane Residence Time—We studied dissociation kinetics via a rapid dilution strategy through the transfer of single protein-bound GUVs to a solution devoid of protein (Fig. 4A). To reduce the complexity of kinetic processes, we limited our experiments to conditions where membrane morphology transitions, such as tubulation, are not observed (Fig. 4, B and C).

Considering the electrostatic component of N-BAR/membrane interactions (10, 16, 18), we selected a membrane composition containing 25 mol % of negatively charged lipids that was recently employed in N-BAR studies (50). A fluorescence decay time of $1010 \pm 245$ s (mean ± S.E. for nine GUVs) was determined (Fig. 4, D and E).

Early Membrane Association Assessed by Stopped Flow—We next extended the membrane binding investigation to shorter timescales by using stopped-flow rapid mixing and FRET between endophilin N-BAR A66-W and DPH-PC embedded in vesicle membranes, or alternatively, by monitoring light scattering changes induced by protein binding (Fig. 5A). The kinetic traces are well fitted by a single exponential (Fig. 5B) for a wide range of conditions investigated. We used this approach to compare membrane binding kinetics of cross-linked (irreversibly dimerized) and non-cross-linked endophilin N-BAR domains. Consistent with our finding of subnanomolar dimerization affinity, we determined indistinguishable membrane binding kinetics for cross-linked and non-cross-linked samples under the same experimental conditions (Fig. 5C).

Taken on its own, the observation via stopped flow of single-exponential membrane binding kinetics could be consistent with a one-step membrane binding process (51). However, the slow dissociation kinetics observed by microscopy (Fig. 4E)
suggest (as further discussed below) the presence of a second step in the membrane binding mechanism.

**Kinetic Model**—We formulated a kinetic two-step model (see Equations 4–7) as a minimal model consistent with our observations for the endophilin-membrane interaction. This model bridges the time scales characterized by stopped flow and microscopy (Fig. 6A). In this model, dimeric protein components in solution (D in Fig. 6A) represent the membrane-binding species, achieving both association with the lipid membrane (L) and insertion of amphipathic helices effectively in the same step. Our model thus does not include separate steps for association of the protein with the membrane and the insertion of amphipathic segments. This is due to the fact that the stopped-flow data are well described by a single exponential (see e.g. Ref. 52) and that the folding transition associated with membrane insertion is likely not rate-limiting (53). We designate the resultant N-BAR-membrane intermediate species \( D^*L \), where the asterisk is meant to represent an inserted species, and \(-L\) indicates membrane-bound species. Intermediate \( D^*L \) subsequently oligomersizes on the membrane, generating the species \( D'\text{L} \), which is proposed to govern membrane residence time.

To test this kinetic model, we identified a range of protein and vesicle concentrations in which N-BAR is dimeric but does not alter membrane morphology (Fig. 6, B and C) for collection of stopped-flow data (Fig. 6D). We note that this model corresponds to that recently proposed in analysis of the binding mechanism of an amphipathic membrane-binding peptide (44).

To determine the rate constants specified in the model, we employed global fitting of numerically integrated rate equations to the entire data set of Fig. 6D (results in Fig. 6A), treating membrane-bound species (\( D^*L \)) within a surface density perspective (see “Experimental Procedures”). The oligomeric species was approximated as a dimer of N-BAR dimers (i.e. \( n = 2 \) in Fig. 6A). We used the single-exponential decay constant observed in GUV dissociation experiments (see example trace in Fig. 6E; using the acidic lipid content and ionic strength of the stopped-flow series of Fig. 6D) as an estimate of \( k_{2-} \) (Fig. 6A). Global fitting with this two-step model yields rate constants for the first step of \( k_1 = 0.11 \pm 0.01 \text{ M}^{-1} \text{s}^{-1} \) and \( k_{1-} = 36.4 \pm 2.4 \text{ s}^{-1} \). These values can be compared with those resulting from fitting a single-step model to the data shown in Fig. 6D; we find \( k_1 = 0.122 \pm 0.001 \text{ M}^{-1} \text{s}^{-1} \) and \( k_{1-} = 35.1 \pm 2.4 \text{ s}^{-1} \), i.e. identical values within fit errors when compared with the two-step model. Consistent with this finding is a large uncertainty of the fit value for \( k_2 \) of the two-step model; our stopped-flow measurements report on the first membrane binding step (Fig. 6A, \( k_2 \) not reported) only. These findings are consistent with a large separation in time scales for membrane binding/unbinding of dimeric BAR domain proteins on the one hand and oligomerization/deoligomerization of membrane-bound BAR domain protein on the other hand.

**Substantiating the Role of Oligomerization in the Mechanism of N-BAR Membrane Binding**—Our hypothesis of oligomerization following membrane binding is supported by the direct observation (via cryo-electron microscopy imaging) of oligo-
merization of membrane-bound endophilin N-BAR (23). Importantly, endophilin N-BAR domain lattice formation has been observed even in the absence of morphological changes relative to the bare membrane, as is true of our conditions (Figs. 4, B and C, and 6, B and C).

Comparative studies with an endophilin variant lacking the H0 sequence (H9004 NH (BAR) (residues 33–247 (16))) further support our incorporation of membrane-mediated oligomerization in the model and exclusion of an explicit step for membrane insertion. Recently published cryo-electron microscopy reconstructions showing intermolecular interactions between helix 0 components (21, 23), site-specific cross-linking studies (21), and recent optical imaging studies (28) document the role of H0 in mediating oligomerization of membrane-bound endophilin N-BAR. Considering these studies, our model predicts that H9004 NH (BAR) would exhibit accelerated dissociation relative to N-BAR. Indeed, remarkably accelerated dissociation was observed for the helix deletion construct H9004 NH (BAR) relative to N-BAR under the same conditions (Fig. 7, A and B).

In further support of our view on oligomerization being a second step in the membrane binding mechanism, we found a dependence of the dissociation time on the protein density on the membrane (Fig. 7 B). Conversely, no density dependence of membrane dissociation rates is expected for a protein that does not oligomerize on the membrane because in that case, protein unbinding will follow a kinetic first-order process. The density

Dimerization and Membrane Binding Kinetics of Endophilin N-BAR

FIGURE 4. Dissociation kinetics under nontubulating conditions. A, diagram of dissociation kinetics experiments employing rapid dilution via micropipette-mediated GUV transfer. B and C, electron micrographs of liposomes (composed of 25% DOPG) in the absence of endophilin N-BAR (B) and in the presence of 300 nM N-BAR_C241-AF488 (4 μM lipids, 33 mM NaCl) (C). D, fluorescence micrograph of N-BAR_C241-AF-488 bound to a GUV (25% DOPG) aspirated in a micropipette, in 33 mM NaCl, 1 mM DTT following preincubation at 300 nM protein, 4 μM lipids, and identical ionic strength. Scale bar, 5 μm. E, fluorescence (Fl) intensity record and single-exponential fit (solid line) charting dissociation of N-BAR, as diagrammed in A, with conditions of D. norm., normalized.

FIGURE 5. Stopped-flow assessment of early timescale of membrane association. A, FRET between endophilin N-BAR A66W and DPH-PC (alternatively, excess light scattering using unlabeled proteins and membranes) used to follow binding upon rapid mixing of protein with vesicles. B, representative stopped-flow record of light scattering monitoring the process diagrammed in A, using 100% DOPG vesicles, 72 μM total lipid, and 0.4 μM N-BAR, C241 in 100 mM NaCl with 2 mM DTT, and single-exponential fit (gray). AU, arbitrary units. C, single-exponential decay constants from stopped-flow progress curves as in B, with and without cross-linking (Fig. 3, A and B).

FIGURE 6. Global experimental data analysis to rationalize two prominent timescales of endophilin-membrane interactions under nontubulating conditions. A, scheme as described under “Results.” Amphipathic helices are indicated by outlined circles. Rate constants represent global fitting results from panel D. B and C, electron micrographs of liposomes (55 μM total lipid) in presence of 0.65 μM (B) and 1.3 μM (C) N-BAR A66W. Liposomes in the absence of protein are imaged in Fig. 4B. Scale bars, 200 nm. D, stopped-flow DPH-PC FRET data (black) used for quantitating the proposed model below. Gray lines display global fitting to predictions of the model, as described under “Experimental Procedures,” yielding parameters in A. Each graph shows post-mixing concentrations of 25% DOPG vesicles of 55, 80, 115, and 160 μM total lipid (with increasing signal magnitude) for fixed N-BAR A66W concentrations of 0.65, 0.81, 1.0, and 1.3 μM as indicated in 33 mM NaCl, 1 mM DTT. Protein solutions were prepared to premixing concentrations 24 h prior to measurements. AU, arbitrary units.
Dimerization and Membrane Binding Kinetics of Endophilin N-BAR

FIGURE 7. N-BAR exhibits density-dependent membrane dissociation and ΔNH (BAR) exhibits accelerated membrane dissociation. A, protein dissociation records, as in Fig. 4, D and E, within a solution without added salt (to increase membrane binding) following preincubation at −10 mM NaCl, pH 7.4, and 400 mM protein. GUVs contained 25 mol % DOPG. Lines represent single-exponential fits. For both constructs, the decay constant is orders of magnitude different from the timescale of the unbinding rate constant yielded by analysis of stopped-flow data assuming a single-step binding/association mechanism (data not shown). FL, fluorescence; norm., normalized. B, dependence of membrane dissociation time of N-BAR and ΔNH (BAR) determined as in A on the initial protein fluorescence (corrected for differing labeling efficiencies of the two proteins) measured on the GUV, proportional to membrane-bound protein density. Protein concentration used for preincubation was varied from 100 to 400 nM for N-BAR and from 100 to 800 nM for ΔNH (BAR), to yield a range of initial membrane-bound protein densities. Other conditions are the same as in A. Lines represent linear fits. Error bars are the errors of single exponential fits to data shown in A. AU, arbitrary units.

dependence is significantly more prominent for N-BAR relative to ΔNH (BAR), consistent with changed N-BAR protein organization at elevated membrane-bound density, leading to slower dissociation.

In sharp contrast to the slow timescale for membrane dissociation of endophilin N-BAR (Figs. 4E and 6A), direct/competitive binding stopped-flow measurements of peripheral proteins not known to oligomerize on the membrane have revealed membrane dissociation rate constants typically between 0.1 and 100 s⁻¹ (see e.g. Refs. 54–57). We note that the single dimer unbinding rate constant (kₚ = 36 s⁻¹; Fig. 6A) determined for endophilin does indeed lie within this range. For HIV1-Nef and a myristoylated alanine-rich C kinase substrate (MARCKS) peptide, both known to insert into the bilayer, respective dissociation rate constants of 0.5 s⁻¹ (58) and 1.5 s⁻¹ (59) have been measured. Thus, the alternative hypothesis that a deinsertion process underlies the slow dissociation of N-BAR and the influence of the H0 segment on dissociation kinetics (Fig. 7A) is not consistent with previously measured rate constants. We note in passing that the on-rate for endophilin membrane binding (k⁺ = 0.1 μM⁻¹ s⁻¹) lies within the range of values typically determined for peripheral membrane proteins (43, 54, 57, 58, 60–63).

Role of Electrostatics in the Association Phase of the Mechanism of N-BAR Membrane Binding—We applied stopped-flow analysis to describe the determinants of the first step in the binding mechanism (Fig. 6A). Promotion of electrostatic interaction strength by varying vesicle acidic lipid (DOPG) content accelerated association and decelerated dissociation (Fig. 8A). This trend is inferred (via the relation kobs = kₚ + [L]kₚ) from increased slopes and decreased y-intercepts (Fig. 8A) extracted from plots exemplified in Fig. 5C. Increasing ionic strength from 33 to 750 mM NaCl significantly decelerated association, as indicated by similar analysis (Fig. 8B), albeit with a lower influence on dissociation (data not shown).

FIGURE 8. Prominent role of electrostatics in the first step of the membrane binding mechanism. A, parameters extracted from analysis of titrations as shown in Fig. 5C, for indicated membrane compositions, averaged over 3–7 vesicle preparations (error bars indicate S.E.), with 33 mM NaCl and 0.3 μM N-BAR A66W. B, averaged slope as in A for experiments varying solution ionic strength via the NaCl concentration, with 72% DOPG and 0.3 μM N-BAR A66W (error bars indicate S.D. and S.E. over ≥3 vesicle preparations). C, DPH-PC stopped-flow association analysis in the pseudo-first-order regime, using 0.3 μM N-BAR A66W, 33 mM NaCl, with 25% DOPG and phosphoinositides (PI(4)P and PI(4,5)P₂) incorporated to 3 mol %. D, pseudo-first-order parameters extracted from plots exemplified in C. Error bars represent S.D. and S.E. of datasets as in C from ≥3 vesicle preparations. A similar trend was observed using 150 mM NaCl and a background composition of 35% DOPG (data not shown).

The sensitivity of stopped-flow parameters to electrostatic interaction strength is consistent with the assumption that the association step (rather than helix insertion) governs kinetics of processes on this timescale. However, carefully delineating the kinetic contributions of membrane association and amphi-pathic membrane insertion remains an aim for future study. Electrostatic influences on kinetics are likely to have implications in the context of a model (64) accounting for the role of endophilin in clathrin-coated vesicle formation (13, 65). In particular, this model attributes a role in vesicle fission to a preference of BAR domain proteins for PI(4,5)P₂ over PI(4)P. This affinity difference has been supported by equilibrium lipid binding studies for endophilin (13, 28). We found a marked influence of phosphoinositides on kinetics, although revealing no significant difference comparing PI(4,5)P₂ and PI(4)P in association or dissociation (Fig. 8, C and D). Thus, localization of N-BAR proteins to necks of budding vesicles may rely more heavily on membrane curvature than on membrane lipid composition.

DISCUSSION

Our in vitro results document the separation of time scales for the following three principal interaction features of endophilin N-BAR at 22–27 °C. Assembled dimers remain associated for hours to days, bind the membrane in tens of milli-
onds, and following higher-order oligomerization, remain membrane-bound for tens of minutes.

Implications of Dimerization Affinity and Kinetics—An influence of monomeric species on the mechanism of endophilin membrane association has been suggested previously. Particularly, the dimerization dissociation constant for N-BAR was measured at 4 °C as 10 μM (16), a far lower affinity than that determined here at 27 °C.

The higher affinity for endophilin dimerization found here relative to previously reported results is, in fact, closer to a value that may be predicted by consideration of dimer-interface surface area. Notably, endophilin exhibits high dimer interface area, as do BAR domains in general (27, 66, 67), calculated using PISA (68) as 2743 Å² for the human N-BAR structure (24). This value is over two times larger than the maximum of a recent tabulation of homodimers classified as weak affinity, which have measured $K_D$ values down to 12 nM (69). Using the affinity curation of Nooren and Thornton (70) this interface area predicts a subnanomolar $K_D$ consistent with our experimentally determined value at 27 °C.

Our finding of subnanomolar affinity rationalizes previous studies that are consistent with the predominance of N-BAR dimers over monomers at sub/lowl micromolar concentrations: distance measurements via EPR methods in solution at low micromolar concentrations (17), biochemical analysis of extracts from rat brain (71) and from cultured cells exogenously expressing endophilin A2 (45), and fluorescence microscopy analysis of endophilin A2 and A1 fluorescent protein chimeras in cultured cells (35). An analogous notion has been advanced for the BAR protein sorting nexin 9 (72). We note in passing that $K_{assn}$ for endophilin dimerization may be calculated from our measured dissociation equilibrium and rate constants (Fig. 2), as $K_{assn} = K_{dissn}/K_D = (1.4 \times 10^{-9} \text{ s}^{-1})/(0.4 \times 10^{-9} \text{M}) = 3.5 \times 10^{6} \text{ M}^{-1} \text{s}^{-1}$ at 27 °C. We conclude that our results do not support a role for dimerization in the kinetic mechanism of membrane binding.

Deoligomerization Kinetics—Our value for $k_{2-}$ suggests far slower dissociation of membrane-bound oligomers for N-BAR relative to a peptide examined recently (44), likely reflecting intersubunit interactions of higher complexity for N-BAR. We note that disassembly of membrane lattice components in live cells is commonly not spontaneous, but depends on additional agents, as has been illuminated in recent in vitro studies (73–75).

Electrostatic Influence on Membrane Binding—Equilibria of electrostatic membrane binding are well predicted by Gouy-Chapman-Stern theory (76–78). However, predictions for electrostatic influences on kinetics (79–81) of membrane binding are less well developed. Trends in kinetics for varying vesicle acidic lipid content observed here resemble those previously reported for a C2 domain (54), a C1 domain (61), and the HIV-1 Nef protein (58). The increase in association rates with DOPG content for N-BAR (Fig. 8A) indicates that a stage in the N-BAR binding mechanism in which PG hydrogen-bonding networks are disassembled (82) is not rate-limiting.

Physiological Relevance of the Separation of Time Scales of Endophilin Interactions—Live-cell imaging reports slower apparent membrane recruitment times (5–15 s) (7, 9) relative to the present stopped-flow measurements (Figs. 5B and 6D). Numerous features do not present in our in vitro experiments likely contribute to this difference, with dynamin GTPase activity (83) and actin scaffolding (5) representing factors with documented influences on endophilin membrane recruitment in vivo.

It has been suggested that BAR domains oligomerize at clathrin-coated pits (84). Based on the marked extension of membrane residence time observed here and attributed to oligomerization for endophilin N-BAR, this process may underlie the detection of endophilin in internalized vesicle preparations (85, 86). However, recycling of endophilin between membranes and the cytoplasm would likely require mechanisms for the dissociation of endophilin oligomers to thereby regulate the membrane residence time of the protein. Indeed, a regulatory role for membrane-mediated oligomerization of synaptotagmin has been proposed (87). Recent work has shown the effects of phosphorylation within endophilin H1 on membrane binding and tubulation, and in turn, functional recycling of endophilin between membrane and cytoplasm. A direction for future studies is thus to pursue whether phosphorylation within H0 (88) may provide alternative regulation by impacting endophilin oligomerization on the membrane (89).

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Dimerization and Membrane Binding Kinetics of Endophilin N-BAR

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