A biosensor-based framework to measure latent proteostasis capacity

Rebecca J. Wood1, Angelique R. Ormsby1, Mona Radwan1, Dezerae Cox1, Abhishek Sharma2, Tobias Vöpel2, Simon Ebbinghaus2, Mikael Oliveberg3, Gavin E. Reid1,4, Alex Dickson5,6 & Danny M. Hatters1

The pool of quality control proteins (QC) that maintains protein-folding homeostasis (proteostasis) is dynamic but can become depleted in human disease. A challenge has been in quantitatively defining the depth of the QC pool. With a new biosensor, flow cytometry-based methods and mathematical modeling we measure the QC capacity to act as holdases and suppress biosensor aggregation. The biosensor system comprises a series of barnase kernels with differing folding stability that engage primarily with HSP70 and HSP90 family proteins. Conditions of proteostasis stimulation and stress alter QC holdase activity and aggregation rates. The method reveals the HSP70 chaperone cycle to be rate limited by HSP70 holdase activity under normal conditions, but this is overcome by increasing levels of the BAG1 nucleotide exchange factor to HSPA1A or activation of the heat shock gene cluster by HSF1 overexpression. This scheme opens new paths for biosensors of disease and proteostasis systems.

1 Department of Biochemistry and Molecular Biology, Bio21 Molecular Science and Biotechnology Institute, University of Melbourne, Parkville, VIC 3010, Australia. 2 Department of Physical Chemistry II, Ruhr-University Bochum, Universitätsstraße 150, 44780 Bochum, Germany. 3 Department of Biochemistry and Biophysics, Arhenius Laboratories of Natural Sciences, Stockholm University, 10691 Stockholm, Sweden. 4 School of Chemistry, University of Melbourne, Parkville, VIC 3010, Australia. 5 Department of Biochemistry & Molecular Biology Michigan State University, East Lansing, MI 48824, USA. 6 Department of Computational Mathematics, Science and Engineering Michigan State University, East Lansing, MI 48824, USA. Correspondence and requests for materials should be addressed to D.M.H. (email: dhatters@unimelb.edu.au)
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rotein-folding homeostasis (proteostasis) in humans is controlled by a quality control (QC) network of about 800 proteins. Key cogs of the QC network are the chaperones, such as HSP70 and HSP90 family members, which monitor the foldedness of proteins almost from the moment they emerge from the ribosome and thereon throughout their lifespan. While the QC network can dynamically respond to stresses to maintain proteostasis, it can also become depleted in protein misfolding diseases. A challenge has been to define the buffering depth of the QC network in managing proteostasis and to track how it changes when stimulated or challenged.

To understand baseline proteostasis buffering capacity requires the development of new quantitative approaches. Prior schemes have used aggregation of ectopically expressed conformationally destabilized (i.e., metastable) proteins as tags for when proteostasis had declined. These schemes operated on the principle that QC systems actively suppress the aggregation rates of metastable “bait” proteins and hence when proteostasis was depleted cells lost the capacity to suppress aggregation. Others have also examined the in-cell folding rates of a test protein through rapid temperature jumps to follow rates for reestablishment of equilibrium. However, these approaches lack a quantitative capacity to understanding proteostasis; namely through the inability to define the effectiveness of QC systems to engage with the bait proteins.

Our motivation was to develop a new biosensor system modeled on these prior schemes—but with a substantially improved quantitative capacity. Here, we describe a biosensor system based on a series of metastable bait proteins that report on foldedness and aggregation state by fluorescence resonance energy transfer (FRET). We show that we can measure the engagement of QC (in terms of the net holdase activity from many individual components) to the unfolded state of the biosensor and concomitant influences on the biosensor aggregation. We describe a new mathematical framework that can extract quantitative information from the holdase activity of QC as well as ability to suppress aggregation. These approaches provide insight to the depth of the pool of QC resources that regulate proteostasis.

Results

Barnase as a sensing kernel for a new tunable biosensor. The prior biosensor schemes using metastable proteins typically display complex folding mechanisms, which make good substrates for QC but pose great challenges in mechanistically quantifying the effects in terms of the thermodynamics of the system. We hence chose a bait protein through which we could more deeply examine the thermodynamics of protein folding and aggregation in the context of QC engagement. Barnase (in a catalytically inactive form (H102A mutant) defined hereon as wild-type based on established nomenclature), was chosen as our bait because it can be predictably tuned to different free energies of folding ($\Delta G_F$) by mutation. Knowledge of, and the capacity to predictably alter $\Delta G_F$, provides a strategy to alter the dynamic range of foldedness and hence provide a handle for more control over defining the extent to which proteostasis alters the folded state of barnase away from thermodynamic equilibrium and basal aggregation levels (Fig. 1).

We predicted that a FRET strategy—flanking the barnase moiety with two fluorescent proteins—could be used to monitor barnase conformations; including chaperone-unfolded client complex (Fig. 1). To make the biosensor, we first screened a mini-library of circularly permuted variants of monomeric teal fluorescent protein 1 (mTFP1) and Venus to identify the optimal combination for tracking barnase foldedness (Supplementary Fig. 1). Upon determining the best combination (mTFP1 cp175-barnase-Venus cp173), we made 15 mutants predicted to span a range of equilibrium constants of folding ($K_f$), corresponding to $\Delta G_F$ values between $-25$ kJ/mol (most stable) to $1$ kJ/mol (least stable). Denaturation curves of these constructs in purified form or directly in mammalian lysates yielded $\Delta G_F$ values that correlated with the previously recorded or predicted $\Delta G_F$ values, demonstrating that the FRET scheme authentically reports on barnase folding equilibrium (Fig. 2a, b; Supplementary Table 1). Furthermore, assessment of FRET in cells expressing the mutant biosensor revealed the cells containing extensively aggregated biosensor have higher FRET signal than cells lacking aggregates, suggesting that aggregation can be detected as a higher FRET state than the folded and unfolded states (Fig. 2c). Hence, these data confirm the capacity to follow the three FRET states as depicted in Fig. 1: high-FRET (aggregated), medium-FRET (folded), and low-FRET (unfolded).

Next we assessed whether barnase was indeed an appropriate client for sampling mammalian QC. Binding partners to the biosensor were measured by quantitative proteomics on immunoprecipitates of a barnase biosensor tuned to be folded (wild-type) vs. a variant tuned to be substantially unfolded (I25A, I96G). A variety of heat shock family proteins bound abundantly to both forms of the biosensor based on the Exponentially Modified Protein Abundance Index (emPAI) score. Four of the five most enriched binding partners to the more unfolded biosensor (with a $p < 0.01$, Student’s t-test) were HSP70 family members (3.9-fold more HSPA1B, 1.7-fold more HSPA8) and HSP90 family members (1.9-fold more HS90AB1 and HS90AA1) (Supplementary Table 2). Furthermore, these four chaperones were the most abundant proteins in the immunoprecipitates based on the emPAI score. Hence, we concluded that the basic barnase module is a suitable probe for some of the major chaperones of the QC network.

Two quantitative measures of QC engagement. To calculate the extent of QC engagement (which we hereon refer to in terms of holdase activity) we needed to devise a strategy to distinguish the conformers of barnase as depicted in Fig. 1a in intact live cells. To do this, we used flow cytometry to measure donor and FRET (i.e., sensitized emission) channels. For the wild-type barnase biosensor, we saw a strict linear relationship between donor and FRET emission (Fig. 3a). Because the slope of this dependence is
proportional to FRET efficiency (i.e., a higher FRET will result in a greater slope), its linearity indicated that all cells belong to a single FRET population with similar fraction of folded barnase, which we anticipated would be close to 100% based on the ΔG\textsubscript{F} value of ~25 kJ/mol. By contrast, cells expressing destabilized barnase mutants, such as the V45T, L89G double mutant, segregated into two populations with different FRET slopes (Fig. 3a). Sorted cells from the Lower-slope population were enriched with unaggregated barnase, whereas those in the Upper-slope were enriched with visibly aggregated barnase, accounting for the increased FRET (Fig. 3b). These data suggested that the gradient of the Lower-slope population provided a measure of the balance at thermodynamic equilibrium of folding (Fig. 4a). Barnase folding and unfolding rates are slower than this—synthesis occurs on the scale of milliseconds to seconds\textsuperscript{15}. Assuming that synthesis, degradation and aggregation rates are slower than this—synthesis occurs in mammals on the scale of at least 10 s of seconds per protein\textsuperscript{16} and degradation occurs on the rate of minutes to hours\textsuperscript{17}—the ratio between native and free unfolded barnase (K\textsubscript{D}) will remain the same at equilibrium, but both states will decrease in concentration at higher proteostasis capacity.

To test this model, we overexpressed human chaperones HSPA1A and DNAJB1 (family members of HSP70 and HSP40 respectively). Immunoprecipitation experiments of a biosensor variant tuned to be substantially unfolded (I25A, I96G) verified this treatment produced a greater interaction of HSPA1A with the biosensor as anticipated from elevated holdase activity (Supplementary Fig. 3a—c). And in accordance with the model, this treatment also decreased the Lower-slope values compared to a negative control biosensor lacking the barnase kernel, consistent with a greater holdase activity to the unfolded biosensor (Fig. 4b).

Next we sought to use the model to define the extent of the pool of QC resources capable of holdase activity, defined here as the latent chaperone concentration (C). Changes in C between a control condition and treatment (ΔC) can be determined with knowledge of a binding affinity constant for the average interactions of all chaperones in the cell with barnase (K\textsubscript{D}) and the fraction folded of a given barnase mutant under control (f\textsubscript{c}) and treatment (f\textsubscript{t}).
fraction folded (left axis, scaled to arbitrary, cannot be compared between different panels and can be measured as a single scalar parameter by the translational amount for all barnase mutants (Fig. 5d). These results when all the mutants were considered, it was apparent that the barnase concentration at which 50% of the cells contained aggregates at a particular time point of expression (A50%) and also to higher protein levels in each cell (Fig. 5b). Overexpression of DNAJB1 and HSPA1A increased the correlation with biosensors tuned to more positive values due to changes in instrument calibration settings.

Analysis of the Lower-slope data for the set of barnase mutants upon the DNAJB1 and HSPA1A over-expression enabled a calculation of ΔC (Fig. 4c) and hence provided a useful scalar measure of QC holdase activity. Details of the analysis are explained in Supplementary Note 2 (for derivation) and Supplementary Note 3 (for application of Equation 1 to our data).

To provide a quantitative measure of aggregation, we investigated the Upper-slope FRET population and the proportion of cells that fell into this population (Fig. 5a). As anticipated, the proportion of cells containing barnase aggregates increased in correlation with biosensors tuned to more positive ΔGF values and also to higher protein levels in each cell (Fig. 5b). Overexpression of DNAJB1 and HSPA1A increased the concentration-threshold of aggregation for two of the barnase variants (I51A and V54T, I88G), consistent with chaperone-mediated suppression of aggregation (Fig. 5c). This data indicated that changes in aggregation propensity could be measured by the barnase concentration at which 50% of the cells contained aggregates at a particular time point of expression (A50%) (Fig. 5c). When all the mutants were considered, it was apparent that the A50% correlated linearly to ΔGF (Fig. 5d). The DNAJB1 and HSPA1A overexpression treatment offset A50% by a comparable amount for all barnase mutants (Fig. 5d). These results collectively indicated that the change in proteostasis efficiency can be measured as a single scalar parameter by the translational offset due to the treatment of interest (ΔA50%) (Fig. 5e; further discussion of the A50% analysis is provided in Supplementary Note 1).

Using the biosensor to probe how QC manages proteostasis. Next we assessed whether we could use the biosensor to measure changes in QC engagement upon stress of proteostasis. First we inhibited HSP90 with novobiocin, which impairs HSP90 activity without activating the heat shock response.18,19 We predicted this treatment would lead to negative ΔC and ΔA50% values, which was the case (Figs 4d and 5e). Furthermore, the activity of novobiocin could be extracted with this method in terms of a dose response curve, providing an IC50 in reasonable accordance with its known value (1.8 mM (our data) c.f. 700 μM (literature)18; Supplementary Fig. 4). As a control, we overexpressed HSP90 family member HSP90AA1, which led to the anticipated reverse response with respect to suppression of aggregation although the change in holdase activity was not significant (Fig. 6a). As a second test for stress, we depleted ATP levels using a glycolysis inhibitor (2-deoxy-d-glucose) in combination with an inhibitor of oxidative phosphorylation (valinomycin)20,21 (Supplementary Fig. 5). Under such conditions, we predicted that chaperone networks would be impaired from functioning, and hence lead to a net loss of engagement with client. This appeared to be the case, with a negative ΔC (Fig. 4d) resulting as well as a negative ΔA50% (Fig. 5e).

Next we explored the capacity of the biosensor system to gain insight into the mechanics of QC systems that regulate proteostasis. First, we applied a selective inhibitor of HSP70 ATPase activity VER-155008, which binds competitively to the ATP binding pocket of HSP70 family proteins.22 While this inhibitor does not prevent client binding in vitro, it

Fig. 3 Barnase foldedness and aggregation can be readily assessed by flow cytometry. a Flow cytometry analysis to determine FRET. Data points indicate fluorescence signatures of individual cells for donor and FRET channels. b Representative confocal images of cells recovered by cell sorting (scale bar=10 μm; “Soluble” image scaled at 4× brightness vs. “Aggregated”). c Conceptual framework for how flow cytometry data reports on foldedness vs. aggregation. The Lower-slope gradients of barnase mutants measured by flow cytometry (right axis) follows the expected relationship between stability (ΔGF) and fraction folded (left axis, scaled to fit). Each data point reflects one barnase mutant (mean ± SD; three replicates). Note that the AFU or AU scales, while arbitrary, cannot be compared between different panels and figures due to changes in instrument calibration settings.

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\Delta C = \frac{-K_d K_f (f_i - f_f)}{f_d f_c} - B(f_i - f_f) \left(1 + \frac{1}{K_f}\right)
\]
did lead to an increase in the Lower-slope values and decrease in $A_{95\%}$ consistent with an overall reduction in the pool of chaperone supply with dynamic capacity to act as a holdase and suppress aggregation (Supplementary Fig. 6).

Next we probed HSP70-HSP40 holdase activity in greater context of nucleotide exchange factors (NEFs), which catalyze the disengagement of client bound to HSP40-HSP70 to complete the chaperone cycle\cite{24-26}. As described above (Fig. 4c, d), overexpression of HSPA1 and DNAJB1 increased the net unfolded pool of the biosensor through a gain in holdase activity. This suggested that overloading a cell with HSPA1 and DNAJB1 by their selective overexpression created a bottleneck in QC capacity to complete the chaperone cycle. To test this possibility, we co-expressed BAG1, the NEF co-factor to HSPA1, with HSPA1 and DNAJB1 (Fig. 6b). This treatment significantly increased foldedness of a biosensor tuned to be moderately unfolded (L89G) compared to when HSPA1 and DNAJB1 were overexpressed without BAG1. This is hence consistent with a decrease in HSP70 holdase activity as anticipated by a greater NEF activity to facilitate release of client bound to HSPA1 and DNAJB1. Assessment of aggregation by $\Delta A_{95\%}$ analysis revealed the NEF to not provide any further enhancement to proteostasis than HSPA1 and DNAJB1 overexpression without BAG1, suggesting that the holdase activity may be sufficiently effective to mitigate inappropriate aggregation (Fig. 6b).

An interesting result from this experiment was that the NEF-HSP40–HSP70 overexpression increased foldedness of the biosensor to a greater extent than what was observed under baseline conditions (Fig. 6b). This suggested that the HSP70 chaperone cycle did not operate at maximum potential under normal conditions whereby NEF activity was the rate limiting step. To further probe this possibility, we overexpressed HSFl, which induces expression of the heat shock response genes (Fig. 6a). This treatment significantly increased foldedness of the L89G variant biosensor (and also suppressed its aggregation) beyond baseline conditions (Fig. 6a). Hence, this supported the conclusion that HSP70 chaperone activity is rate-limited under normal conditions by NEF activity and that this can be overcome by a coordinated upswing in cellular QC resources driven through the heat shock response gene elements.

**Discussion**

We describe here a strategy for quantifying the holdase activity of the QC network as an indicator of proteostasis health. Using barnase as a bait protein to the QC we were able to define the extent to which the QC system altered the folding equilibrium and aggregation and also validate how manipulation of QC machinery alters the biosensor readout in a predictable way. The key objective of the biosensor was to provide a simple numerical metric of “holdase” activity and capacity to suppress aggregation. As such, this system provides a quantitative measure of the health of proteostasis. However, as indicated by the studies with selective chaperone overexpression and proteomics, the biosensor also has capacity for nuanced analysis of the specific changes in chaperones that interact with the biosensor under different conditions. One area of potential application is in the study of selectivity in how different HSP70 proteins respond to client under different stresses. Prior work has suggested HSP70 family members display selectivity in function: some are better at refolding misfolded proteins and others at preventing aggregation\cite{27}. Others seem to be involved in protein disaggregation\cite{28}. The biosensor platform described here provides a manner to explore the dual role of holdase activity and aggregation of individual HSP70 proteins (e.g., by their selective knock out) in an intact proteostasis system. An interesting finding was how the HSP70 chaperone cycle appeared to not be operating a maximal capacity under baseline conditions. This may provide a mechanism for HSP70 to accumulate client as part of HSP70-mediated client triage to other nodes of the QC system, such as degradation\cite{29}. It may also provide QC systems an instant buffering strategy after an acute stress during the time period for stress responses to synthesize additional QC resources.

Collectively, the new biosensor system and methods described here offer promise to probe mechanisms intersecting proteostasis and disease. In particular the biosensor system has great potential to be developed into a tool to measure early changes in neurodegenerative disease where early diagnostics are desperately needed. The development of treatments for neurodegenerative diseases including Huntington’s (HD), Alzheimer’s (AD) and Motor Neuron Disease (MND) remains one of the toughest scientific challenges of our times. Between 2002 and 2012 only one of 244 AD clinical trials yielded a treatment with therapeutic benefit\cite{30}. Similarly Motor Neuron disease remains largely untreatable with only two drugs approved by the US Food and Drugs Administration that mildly delay disease progression (edaravone and riluzole)\cite{31,32}. The failure of clinical trials in neurodegenerative diseases may arise from trials beginning too late in the disease course. In turn, beginning trials at pre-symptomatic stages of disease has been hampered by a lack of
Fig. 5 Quantifying proteostasis by changes in biosensor aggregation propensity ($A_{50\%}$). a Conceptual framework for how chaperone levels affect barnase-biosensor aggregation. b Baseline barnase aggregation “landscape” in cells as measured by the proportion of cells in the Upper-slope flow cytometry population. Data show mean of three replicates for each barnase mutant coexpressed with Y66L EGFP control. c Impact on chaperone overexpression on aggregate landscape. Data show mean of three replicates for each barnase mutant coexpressed with Y66L EGFP control. Data points are means ± SEM of three replicates. Lines show linear regressions with same slope (preferred model by extra sum of squares F-test, $p = 0.57$). Δ$A_{50\%}$ is the translational offset, calculated for each mutant (example indicated by arrow). d Impact of proteostasis on changes in $A_{50\%}$ (Δ$A_{50\%}$). Data show chaperone co-expression (compared to Y66L EGFP negative control), treatment with proteostasis-modulating drugs (compared to untreated control). Data are means ± SEM of 12 mutants of barnase. Results of Wilcoxon signed rank test are shown: Results coded as *** $p < 0.001$

 METHODS

Expression constructs. A toolkit for FRET biosensor comprising a combinatorial library of circularly permuted mTFP1 and Venus fluorescent proteins (cpFRET library in pTriEx4 expression vector) was used34. Barnase mutants were synthesized (Thermo Fisher Scientific) and inserted via PCR-mediated cloning with XmaI and NotI flanking restriction sites between BspEI (complementary to XmaI) and NotI sites of the cpFRET library such that barnase was fused to mTFP1 and Venus at the amino and carboxy termini, respectively. Additional single mutations were introduced and double-mutant constructs using BspEI and XbaI restriction sites. All constructs were verified by sequencing and the basic construct sequence is shown in Supplementary Table 3. Plasmids expressing HSPA1A, DNAJB1 and mCherry were prepared as described previously33,35: pCMV6-AC RAG1 (cat# SC319483), pCMV6-AC HSF1 (cat# SC321225) and pCMV6-Entry myc-HSP70/A1 (cat# RC212496) were purchased from Origene.

Cell culture. AD293 and HEK293 cell-lines (from lab cultures originally obtained from ATCC) were used in this study and tested and cleared for mycoplasma. Cells were not tested for cross-contamination of other cell lines or misidentification. AD293 cells were maintained in Dulbecco’s modified Eagles Medium (DMEM) supplemented with 2 mM L-glutamine, 200 U mL$^{-1}$ penicillin/streptomycin and 10% v/v fetal bovine serum (Thermo Fisher Scientific) in a humidified 37 °C incubator with 5% v/v atmospheric CO$_2$. HEK293T cell-lines were maintained as for AD293 cells except DMEM was supplemented with 2 mM L-alanyl-L-glutamine, 100 U mL$^{-1}$ penicillin/streptomycin and 10% v/v fetal bovine serum. For biosensor screening, 2 $\times$ 10$^5$ AD293 cells were seeded on poly-L-lysine coated 12-well plates and transfected using 1.6 µL P3000 reagent and 0.75 µL Lipofectamine 2000 and 4 µL Lipofectamine 2000 as per the manufacturer’s instructions. For urea denaturation, 5 $\times$ 10$^5$ HEK293T cells were seeded on poly-L-lysine coated 48-well plates (Corning) or 8-well µ-slides (for microscopy, Ibidi). Cells were transfected using 0.25 µg DNA, 0.5 µL P3000 reagent and 0.75 µL Lipofectamine 3000 according to the manufacturer’s instructions.

Drug treatments. To deplete ATP, cells were kept in glucose-free DMEM (Thermo Fisher Scientific) for 16 h then treated with 10 mM 2-deoxy-d-glucose.
Fig. 6 Probing changes in proteostasis through selective activation of different chaperone systems. a) Investigation of the effect of induction of the heat shock response by HSF1 overexpression or selective overexpression of HSP90. Shown on left are western blots of the HEK293T cells matched for total protein (a) and the L89G barnase biosensor (which also contains a myc tag). The right graphs show the Lower-slope and ns post-hoc test. Results coded as ****p < 0.0001, **p < 0.01, *p < 0.05, ns => 0.05. b) Same paradigm as panel a. In this case, BAG1 was cotransfected with the barnase biosensor, HSP40 (DNAJB1) and HSP70 (HSPA1A). The right panels show the proportional dosage of each construct (by mass of DNA) in the transfection. Bars indicate means ± SEM. The right panel shows the lower-slope graph was analysed via a two-way ANOVA subjected to Dunnett’s post hoc test and the ANOVA analyses of these treatments and one or three of the biosensor variants as indicated vs. the wild-type* variant. Bars indicate means ± SEM. The right panels were analysed via a two-way ANOVA subjected to a Dunnett’s post-hoc test, results coded as ****p < 0.0001, **p < 0.01, *p < 0.05, ns => 0.05. b) Same paradigm as panel a. In this case, BAG1 was cotransfected with the barnase biosensor, HSP40 (DNAJB1) and HSP70 (HSPA1A). The right panels show the proportional dosage of each construct (by mass of DNA) in the transfection. Bars indicate means ± SEM. The right panel shows the lower-slope graph was analysed via a two-way ANOVA subjected to Dunnett’s post hoc test and the ANOVA analyses of these treatments and one or three of the biosensor variants as indicated vs. the wild-type* variant. Bars indicate means ± SEM. The right panels were analysed via a two-way ANOVA subjected to a Dunnett’s post-hoc test, results coded as ****p < 0.0001, **p < 0.01, *p < 0.05, ns => 0.05. b) Same paradigm as panel a. In this case, BAG1 was cotransfected with the barnase biosensor, HSP40 (DNAJB1) and HSP70 (HSPA1A). The right panels show the proportional dosage of each construct (by mass of DNA) in the transfection. Bars indicate means ± SEM. The right panel shows the lower-slope graph was analysed via a two-way ANOVA subjected to Dunnett’s post hoc test and the ANOVA analyses of these treatments and one or three of the biosensor variants as indicated vs. the wild-type* variant. Bars indicate means ± SEM. The right panels were analysed via a two-way ANOVA subjected to a Dunnett’s post-hoc test, results coded as ****p < 0.0001, **p < 0.01, *p < 0.05, ns => 0.05. b) Same paradigm as panel a. In this case, BAG1 was cotransfected with the barnase biosensor, HSP40 (DNAJB1) and HSP70 (HSPA1A). The right panels show the proportional dosage of each construct (by mass of DNA) in the transfection. Bars indicate means ± SEM. The right panel shows the lower-slope graph was analysed via a two-way ANOVA subjected to Dunnett’s post hoc test and the ANOVA analyses of these treatments and one or three of the biosensor variants as indicated vs. the wild-type* variant. Bars indicate means ± SEM. The right panels were analysed via a two-way ANOVA subjected to a Dunnett’s post-hoc test, results coded as ****p < 0.0001, **p < 0.01, *p < 0.05, ns => 0.05. b) Same paradigm as panel a. In this case, BAG1 was cotransfected with the barnase biosensor, HSP40 (DNAJB1) and HSP70 (HSPA1A). The right panels show the proportional dosage of each construct (by mass of DNA) in the transfection. Bars indicate means ± SEM. The right panel shows the lower-slope graph was analysed via a two-way ANOVA subjected to Dunnett’s post hoc test and the ANOVA analyses of these treatments and one or three of the biosensor variants as indicated vs. the wild-type* variant. Bars indicate means ± SEM. The right panels were analysed via a two-way ANOVA subjected to a Dunnett’s post-hoc test, results coded as ****p < 0.0001, **p < 0.01, *p < 0.05, ns => 0.05.

Recombinant production of the Venus cp173 construct. The Venus cp173 construct in the pTrEx based vectors was expressed in T7 Express E. coli (NEB) using ampicillin as the selection antibiotic. 10 mL of an overnight starter culture (grown in 2 × YT at 37 °C in a shaking incubator), was inoculated in 1 L of 2 × YT grown to an OD600nm of 0.5 AU at 37 °C in a shaking incubator. The culture was cooled to 18 °C and then expressed was induced with 0.4 mM Isopropyl β-D-1-thiogalactopyranoside with cells grown overnight at 18 °C in a shaking incubator. Cells were pelleted (5000 × g; 4 °C) and resuspended in 100 mM Tris, pH 8.0 supplemented with Complete EDTA-free protease inhibitor cocktail (Roche) and 1 mM phenylmethylsulfonyl fluoride. Hen egg white lysozyme was added to a concentration of 1 mg mL⁻¹ and the lysate was frozen at −20 °C. The lysate was thawed, Benzonase nuclease was added according to the manufacturer’s instructions (EMD-Millipore). The debris was pelleted and removed by centrifugation (16,000 × g; 20 min; 4 °C). Imidazole was added to the supernatant to a concentration of 5 mM and the solution was applied to a 1 mL His-tag column presaturated in buffer B (5 mM imidazole). The column was washed with 20 mL buffer C and then with PBS and 25 mM imidazole until no proteins further eluted (as assessed by Bradford assay). The column was then washed with 10 mL PBS and 50 mM imidazole before elution with PBS and 200 mM imidazole. Eluted proteins from the most concentrated fractions were buffer exchanged into PBS using a PD-10 column (GE Healthcare). Proteins were immediately snap frozen in liquid nitrogen and stored at −80 °C until further use. (Sigma) and 0.5 μM valinomycin (Sigma) in glucose-free DMEM for 45 min. ATP levels in ATP-depleted and control (untreated) cells were measured using the ATPLite luminescence assay system (Perkin Elmer) on a ClarioStar microplate reader, as per the manufacturer’s protocol. HSP90 was inhibited with 800 μM novobiocin in maintenance media for 6 h. HSP70 was inhibited with 20 μM VER-155008 (Sigma cat # SML0271) in maintenance media for 18 h.

FRET biosensor screen. The cpFRET library containing wild type* (H102A) barnase was transfected into AD293 cells. 24 h post-transfection cells were washed with PBS then lysed by extrusion through a 27 gauge syringe in native lysis buffer (20 mM Tris pH 8.0, 2 mM MgCl2, 1% v/v Triton X-100, 1 × EDTA-free protease inhibitor (Roche), 150 mM NaCl, 20 U mL⁻¹ benzamid, 1 mM PMSF). Lysate was centrifuged at 100,000 × g for 30 min at 4 °C to remove cell debris. 80 μL supernatant was added to 200 μL native lysis buffer or native lysis buffer with 6 M urea in a 96-well plate. mTFP1 was excited at 462 nm and emission spectra were collected from 480 to 600 nm using a Varioskan Flash microplate reader (Thermo Fisher Scientific). Relative FRET efficiency was calculated as A(D)/A(A); that is, fluorescence intensity at the acceptor (A) maximum (532 nm) divided by the sum of intensities at the donor (D) (492 nm) and acceptor maxima. Because the readings were ratiometric, we deemed it unnecessary to normalize protein concentration in the lysates.
Recombinant production of the barnase biosensors. The barnase constructs in the pTriEx based vectors were expressed in T7 Express E. coli (NEB) using ampicillin as the selection antibiotic. 10 mL of an overnight starter culture (grown in 2 x YT at 37°C in a shaking incubator), was inoculated in 100 mL of 2 x YT and grown to an OD600 of 0.5 AU at 37°C in a shaking incubator. The culture was cooled to 18°C and expression was induced with 0.4 mM Isopropyl β-D-1-thiogalactopyranoside with cells grown overnight at 18°C in a shaking incubator. Cells were harvested by centrifugation at 120 × g for the duration of the experiment. Relative FRET efficiencies (430 nm excitation, 492 nm emission and 532 emission) were measured at 23 °C.

Urea denaturation stability measurements. 384-well plates were prepared with 80 µL of a concentration series of urea (0 M to – 6 M) in phosphate buffered saline (PBS). For the recombinant purified proteins, 5 µL of protein was used in the assay. For the curves performed with cell lysates, cells were lysed 24 h after transfection by pipetting in native lysis buffer. Aggregates and cell debris were pelleted by centrifugation at 16,000 × g for 10 min at 4 °C. 5 µL supernatant was added to each urea concentration. Samples were not matched for protein concentration; however, as the measurements were ratiometric and both fluorophores were on the same molecule, concentration was not expected to influence results. Fluorescence readings (490 nm excitation, 532 nm emission) were measured using a plate reader (Roche); 10 min at 37°C with a 15 min equilibration at 10 °C or less. The resultant lysate was supplemented to 500 µL with HENG buffer and pelleted (16,000 × g for 10 min; 4°C). The supernatant was removed and used for the FRET fluorescence screen. Fluorescence readings were taken at 23°C. The FRET signal of barnase in the native (αα) or denatured (αβ) conformation with [U] = 0 and β is the rate of change of the FRET signal with increasing [U]. As small error in the cooperativity value m has large impact on ΔGm, m was constrained as a shared parameter for all mutants. The data were fitted to the two-state unfolding model with terms for pre- and post-transition baselines (Equation 2).

\[
\text{FRET} = \frac{(m_0 + β_0 \cdot [U])}{(1 + \frac{β_0}{(m_0 - β_0) \cdot [U]} \cdot e^{-\frac{U}{m_0}} - \frac{U}{m_0}}
\]

Where [U] is the urea concentration, \(m_0\) is the urea concentration at which barnase is 50% denatured, m is the cooperativity value and \(e^{-\frac{U}{m_0}}\) is the urea concentration at which barnase is 50% denatured, and \(α_β\) and \(β_0\) describe the FRET baselines of barnase in the native (αα) or denatured (αβ) conformation when \([U]\) = 0 and \(β\) is the rate of change of the FRET signal with increasing [U]. As small error in the cooperativity value m has large impact on ΔGm, m was constrained as a shared parameter for all mutants. The data were fitted to the two-state unfolding model with terms for pre- and post-transition baselines were shared for all mutants. Reported ΔGm were means of three experiments fit independently.

Microscopy. To estimate cell volume, cells expressing Venus (pcp173) were imaged 24 h after transfection on a Leica TCS SPS Confocal microscope using a HCX APO CS 63 × 1.40 Oil objective and 1 Air pinhole as 2-stacks in 0.2 µm steps (514 nm excitation, 520–650 nm emission) and analyzed using ImageJ. To measure FRET in intact cells, cells were imaged with a Leica SP8 confocal microscope using a HCX APO CS 63 × 1.40 Oil objective and 1 Air pinhole. Emission spectral scans were acquired with a 405 nm excitation laser, from 480 to 576.6 nm in 3.45 nm steps (514 nm stop, 30 s on, 30 s off) cycles at 6800 rpm with temperature monitored and maintained at 10°C or less. The resultant lysate was supplemented to 500 µL with HENG buffer and pelleted (16,000 × g for 10 min; 4°C). The supernatant was removed and used for Venus fluorescence (ex = 514 nm, em = 527 nm) using a plate reader (ClarioStar; BMG).

Immunoprecipitation for western blotting. In total 1.25 × 10^7 HEK293T cells were seeded into 25 cm² flasks and transfected the following day with 6.25 µg DNA constructs, 12.5 µg P3000 and 18.75 µg Lipofectamine 3000 (Life Technologies) according to the manufacturer’s instructions (Life Technologies). Cells were harvested at 24 h post-transfection for drug treatments and 48 h for co-expressions. Cells were harvested by gentily pipetting with PBS. Cells were pelleted by centrifugation at 120 × g for 6 min and resuspended in lysis buffer (0.5% v/v IGEpal, 50 mM Tris, 5 mM MgCl₂, pH 7.4 supplemented with EDTA-free protease inhibitor (Roche). Lysate was incubated on ice for 10 min. Debris was removed by centrifugation (13,000 rpm; 10 min; 4°C) and the supernatant placed in a new microcentrifuge tube. For samples matched for barnase fluorescence, fluorescence was measured as a ClarioStar microplate reader (ex = 513 ± 10 nm, em = 530 ± 30 nm). 20 µL of GFP-Trap_MA (Chromotek) beads were washed in washing buffer (0.5% v/v IGEpal, 50 mM Tris, 150 mM NaCl, 5 mM EDTA, pH 7.4) was added to each sample and the mixture was incubated for 2 h at 4°C with constant rotation. The unbound fraction was removed from beads immobilized with a magnet, and beads were washed three times with washing buffer at room temperature. Proteins were eluted off the beads by boiling the beads in SDS-PAGE Laemmli sample buffer for 10 min.

Western blotting. Proteins were transferred to PVDF membrane, blocked in blocking buffer (5% w/v skim milk powder in PBS-T) for 1 h at room temperature. The barnase constructs in the pTriEx based vectors were expressed in T7 Express E. coli (NEB) using ampicillin as the selection antibiotic. 10 mL of an overnight starter culture (grown in 2 x YT at 37°C in a shaking incubator), was inoculated in 100 mL of 2 x YT and grown to an OD600 of 0.5 AU at 37°C in a shaking incubator. The culture was cooled to 18°C and expression was induced with 0.4 mM Isopropyl β-D-1-thiogalactopyranoside with cells grown overnight at 18°C in a shaking incubator. Cells were harvested by centrifugation at 120 × g for the duration of the experiment. Relative FRET efficiencies (430 nm excitation, 492 nm emission and 532 nm emission) were measured at 23 °C.

Sample preparation for proteomics. In total 3.5 × 10^6 HEK293T cells were seeded into 75 cm² flasks and transfected the following day with either WT or I25A, I96G barnase Antibody (18.75 µg DNA, 37.5 µL P3000 and 66.52 µL Lipofectamine 3000 (Life Technologies)) according to the manufacturer’s instructions (Life Technologies). The experiment was designed as 4 biological matched pair replicates. Media was refreshed 5 h after transfection. At 24 h post-transfection, cells were gently rinsed with PBS and harvested in PBS by gently pipetting. Cells were pelleted (120 × g, 6 min; room temperature) and resuspended in 1 mL PBS and pelleted again (4000 × g, 6 min; room temperature). The pellet was resuspended in ice-cold 200 µL HENG buffer (50 mM HEPES-KOH, pH 7.9, 150 mM NaCl, 20 mM Na₂MoO₄, 2 mM EDTA, 5% v/v glycerol, 1 mM PMSF and Complete EDTA-free protease inhibitor (Roche)). Cells were mechanically lysed using a cryomill (Prepsilyzer, Jun-24, 2015; 548872 entries). The search was conducted with 20 ppm MS tolerance against the Swissprot Homo sapiens (version 2015_07: Jun-24, 2015; 548872 entries). The search was conducted with 20 ppm MS tolerance, 0.8 Da MS/MS tolerance, 2 missed cleavages allowed. The following modifications were allowed: Oxidation (M), Acetylation (Protein N-term), Dimethylation (K), Dimethylation (N-term), 1H(3)D(13)C(2) Dimethylation (K), 2H(6)13C(2) Dimethylation (K), 2H(4) (N-term) Dimethylation (N-term) (Variable); Carbamidomethyl (C) (Fixed). The fast discovery rate (FDR) maximum was set to 0.1% at the peptide identification level.
actual was 0.06% for each replicate) and 1% at the protein identification level. Proteins were filtered for those containing at least one unique peptide in all four replicates. The common contaminant, Keratin, was excluded from the dataset. Peptide quantification was performed in Proteome Discoverer v.2.1.0.81 using the precursor ion quantifier node. Dimethyl labelled peptide pairs (between two comparison of light, medium or heavy) were published with a 2 ppm mass precision, a signal to noise threshold of 3, and filtered for a minimum Mascot ion score of 30. The retention time tolerance of isotope pattern multiplets was set to 0.8 min. Three single peak or missing channels were allowed for peptide identification. The protein abundance in each replicate was calculated by summation of the unique peptide abundances that were used for quantification (light, medium and-or heavy dimethyl derivatives). In the cases where quan values were missing or were apparent outliers to the other two replicates, raw data was manually checked and added or adjusted where relevant. The protein ratios (from two of the light, medium and heavy labels) were manually calculated from the data. Data were normalized to the Venus protein peak abundance in the dataset. These correction values were additive on the L-slope and Upper-slope population. Since many datasets contained high numbers of cells with a signal below background to accurately determine the Lower-slope gradient.

Equation 1 was derived as described in Supplementary Note 2 and applied to our data as described in Supplementary Note 3.

Flow cytometry. After 24 h (drug treatments) or 48 h (cotransfections) of post-transfection, cells were washed once in PBS then harvested by gentle pipetting in PBS and transferred to a 96-well U-bottom microplate. 100 µL cell suspension was analysed on 4 µl-1 using the high throughput sampler in an LS-R/Fortessa flow cytometer equipped with 405, 488, and 561 nm lasers (BD Biosciences). Forward scatter threshold was set to 5000. Acceptor (Venus) fluorescence was collected with the 488 nm laser and FITC (530/30) filter. Acceptor sensitization emission (FRET) and donor (mTFP1) fluorescence were collected with the 405 nm laser with PE (575/72) and V500 (525/50) filters, respectively. All flow cytometry data were processed with FlowJo (Tree Star Inc) to exclude cell debris, cell aggregates and untransfected cells. The Venus channel was compensated to remove bleedthrough from mTFP1 and FRET channels. mTFP1, FRET and Venus data were exported as csv files and analyzed in MATLAB (see section on data analysis). The gating strategy is explained in the associated Protocols Exchange manuscript.

The expression level at which there was 50% aggregation (A50%) was determined from the sigmoidal fits. For hypothesis testing (i.e., to determine whether a treatment resulted in a significant change in aggregation propensity) the difference between average A50% of control and treatment was calculated for each barnase mutant (excluding wild-type*, V45T, and 155G). A Wilcoxon signed rank test was used to calculate whether the difference was significantly different to zero, treating the twelve mutants as replicate values.

Data availability. Raw data not presented in the manuscript are available upon request.

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**Author contributions**

R.J.W. designed and performed the experiments and analytical approaches, and co-wrote the manuscript. A.R.O., M.R., D.C., G.E.R., A.S., T.V., S.E. and M.O. assisted in project design and/or performing experiments. A.D. developed the mathematical models. D.M. H. devised the original concept of the project, oversaw its implementation and co-wrote the manuscript.

**Additional information**

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