

Article

Bacterial NLR-related proteins protect against phage

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SUMMARY

Bacteria use a wide range of immune pathways to counter phage infection. A subset of these genes shares homology with components of eukaryotic immune systems, suggesting that eukaryotes horizontally acquired certain innate immune genes from bacteria. Here, we show that proteins containing a NACHT module, the central feature of the animal nucleotide-binding domain and leucine-rich repeat containing gene family (NLRs), are found in bacteria and defend against phages. NACHT proteins are widespread in bacteria, provide immunity against both DNA and RNA phages, and display the characteristic C-terminal sensor, central NACHT, and N-terminal effector modules. Some bacterial NACHT proteins have domain architectures similar to the human NLRs that are critical components of inflammasomes. Human disease-associated NLR mutations that cause stimulus-independent activation of the inflammasome also activate bacterial NACHT proteins, supporting a shared signaling mechanism. This work establishes that NACHT module-containing proteins are ancient mediators of innate immunity across the tree of life.

INTRODUCTION

Bacteria are in constant conflict with viruses called bacteriophages (phages) and have evolved elaborate antiphage signaling systems to halt infections. These phage defense systems are typically multi-gene operons encoding proteins that cooperate to sense infection and inhibit virion production through diverse mechanisms.¹ The best-understood antiphage systems are restriction-modification and CRISPR-Cas; however, there are many additional antiphage genes/systems that we are only beginning to understand.^{2–7} Most antiphage systems are a form of innate immunity, meaning they protect against a wide variety of phages and do not require previous exposure, unlike CRISPR-Cas systems, which are a form of adaptive immunity. Bacteria typically encode multiple antiphage systems, often on mobile genetic elements, which are shared across the pangenome. This arsenal of antiphage systems creates a “pan-immune system,”¹ which depends on the ability of antiphage genes to function well in diverse host cells, protect against disparate phages, possess potential addiction modules, and encode most of their essential components in one gene/operon.

The endeavor to catalog antiphage signaling systems from the bacterial pan-immune system has led to an unexpected finding: some bacterial antiphage proteins are homologous to core components of the human immune system. One clear example is bacterial cyclic-oligonucleotide-based antiphage signaling sys-

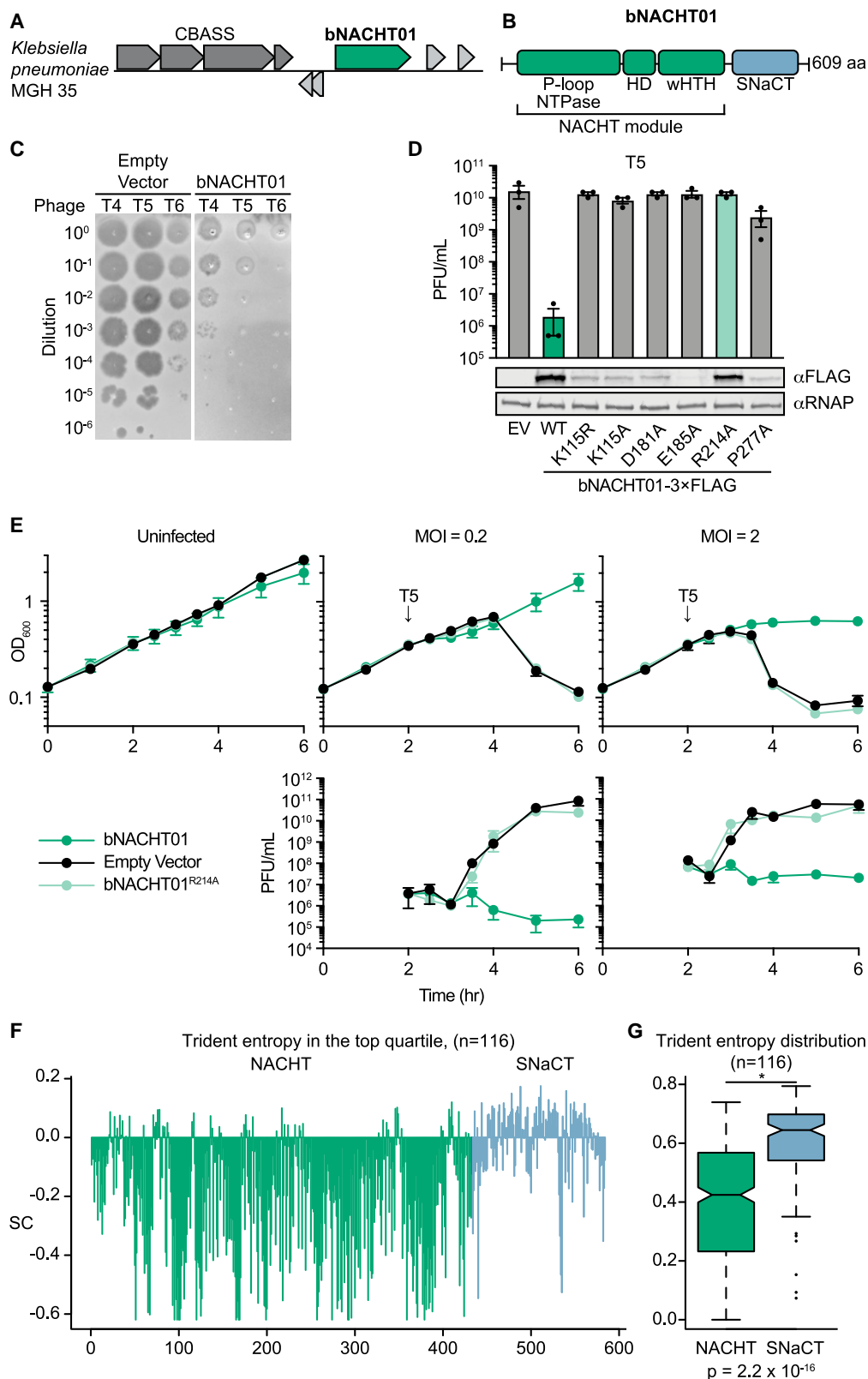
tems (CBASSs^{6,8–11}), which encode proteins homologous to the human cGAS and STING proteins. Other examples are bacterial Viperins¹² and bacterial Gasdermins,¹³ which are homologous to human Viperin and Gasdermin D, respectively. These genes are antiviral in both humans and bacteria, and bioinformatic evidence supports that these genes share a common ancestor.^{6,11–13} It appears that the pervasive horizontal gene transfer of antiphage systems between bacteria may have also resulted in metazoans horizontally acquiring antiphage genes from bacteria, which were then adapted to fight viruses in eukaryotic cells. We therefore hypothesized that additional components of the metazoan innate immune system originated from antiphage signaling systems and searched for those genes in bacteria.

RESULTS

A bacterial NACHT protein is antiphage

Antiphage systems frequently cluster together into “defense islands” throughout bacterial genomes.¹⁴ This phenomenon has been used to identify novel antiphage systems by interrogating genes of unknown function that are co-located with known antiphage systems.^{2,3,7} We investigated genes of unknown function located near CBASS systems and identified a gene encoding a NACHT protein module in *Klebsiella pneumoniae* MGH 35 (Figures 1A and 1B, Genbank: WP_015632533.





(legend on next page)

1). We named this gene bacterial NACHT module-containing protein 1 (bNACHT01), and it did not appear to be in an operon with any other genes. We hypothesized that bNACHT01 was antiphage and tested that hypothesis by expressing bNACHT01 from its endogenous promoter in *Escherichia coli* (*E. coli*), then challenging those bacteria with diverse phages. bNACHT01 conferred over a 100-fold increase in protection against phage T4 and over a 1,000-fold increase in protection against phages T5 and T6 (Figures 1C, 1D, and S1C–S1E). We further confirmed that bNACHT01 is constitutively expressed and that expression is not impacted by phage infection (Figure S1B).

Within 3 h post-infection at a multiplicity of infection (MOI) of 0.2, phage replicated and lysed cultures of bacteria expressing an empty vector (EV). However, bacteria expressing bNACHT01 restricted phage virion production and continued growing (Figure 1E). When cultures were infected at an MOI of 2, bacteria expressing bNACHT01 continued to restrict virion production, but the OD₆₀₀ was stagnant and did not show continued growth or overt bacterial lysis (Figure 1E).

The NACHT module of bNACHT01 shares core features with NACHT modules in eukaryotes, including a Walker A motif that binds NTPs in those proteins^{15–17} (Figures 1B and S1). Mutation of the conserved lysine residue within the bNACHT01 Walker A motif (K115) to arginine or alanine not only abrogated phage defense but also decreased protein expression (Figure 1D). We therefore tested a range of previously published NACHT-inactivating mutations to find an inactivating mutation that did not impact bNACHT01 expression.¹⁸ Mutation of R214 to alanine maintained expression of the protein but abrogated phage defense (Figures 1D, 1E, and S1C–S1E). Based on alignments to nucleotide-binding and oligomerization domain (NOD)1/2, this residue is expected to sense the γ -phosphate of ATP, indicating that NTP binding may be required for antiviral function (Figure S1A).¹⁸

We next interrogated a multiple sequence alignment of the clade of bacterial NACHT proteins defined by bNACHT01 to better understand the mechanism of phage defense. This analysis revealed that the NACHT module is relatively stable in its sequence conservation; however, the region to the C terminus of the NACHT module appears to be rapidly diversifying (Figures 1F and 1G). We named this region the short NACHT-

associated C-terminal domain (SNaCT). Three SNaCT domain models illustrating the observed diversity are provided in Figure S1K. Deletion of the SNaCT domain or mutation to a conserved aspartate at the N terminus of the domain abrogated bNACHT01 phage defense (Figures S1G–S1I). We used trident entropy scores to compare the degree of amino acid conservation for the NACHT and SNaCT domains and confirmed that the mean entropy score is higher for the SNaCT domain (Figures 1F and 1G; see STAR Methods for a description of entropy calculation).¹⁹ The rapid diversification of the SNaCT domain is a hallmark of a host-pathogen “arms race,” where evolutionary pressure from interactions between an immune sensor and pathogen selects for amino acid substitutions that change protein-protein interactions. In eukaryotic NACHT proteins, the C terminus is often the “sensor” or “receptor” region of the protein that responds to infection stimuli.²⁰ The C-terminal leucine-rich repeat (LRR) region of a subset of these maintains the protein in an auto-inhibited state that is alleviated by stimulant-induced conformational changes. The predicted structure of bNACHT01 (Figure S1J) shows the C-terminal SNaCT domain occludes the NTP-binding region, suggesting that the C terminus serves a similar function for bNACHT01 as for some animal NACHT proteins. It is therefore paradoxical that bNACHT01 is both capable of detecting a variety of unrelated phages (T4, T6, and T5) and is rapidly diversifying in the sensor region. These data may suggest that the bNACHT01 SNaCT domain is evolutionarily diverging under pressure from constant antagonism by phage-encoded proteins that enable immune evasion (e.g., phage-encoded bNACHT inhibitors).

Diversity and ubiquity of NACHT module-containing proteins in bacteria

The NACHT module was first discovered in eukaryotes where it is often found in proteins that mediate immunity and inflammation. The best-understood metazoan NACHT proteins belong to the nucleotide-binding domain and LRR-containing gene family (NLRs),²¹ which have a core NOD whose function is fulfilled by a NACHT module. Mammalian NLRs are immune components that play an important role in the formation of inflammasomes (such as NAIP/NLRC4, NLRP1, and NLRP3), transcriptional regulation (CIITA), and other inflammatory responses.¹⁷

Figure 1. A bacterial NACHT domain-containing protein is antiphage

(A) Genome context of bNACHT01, which is located near a CBASS system in *Klebsiella pneumoniae* MGH 35.

(B) Schematic of bNACHT01 (Genbank: WP_015632533.1) protein domains, annotated by alignment to the NACHT module of NLRC4. The P loop NTPase domain is also known as a nucleotide-binding domain (NBD), the helical domain (HD), and the winged helix-turn-helix (wHTH, also called WHD for winged helical domain) are indicated. See Figure S1 for a protein alignment of bNACHT01 with eukaryotic NACHT modules.

(C) Efficiency of plating of indicated phages infecting *E. coli* expressing bNACHT01 or an empty vector (EV). Data are representative images of $n = 3$ biological replicates.

(D) Above: efficiency of plating of phage T5 infecting *E. coli* expressing the indicated genotype. Data represent the mean \pm standard error of the mean (SEM) of $n = 3$ biological replicates, shown as individual points. See Figure S1 for the efficiency of plating of phage T4 and T6. Below: western blot analysis of *E. coli* expressing empty vector or FLAG-tagged bNACHT01 of the indicated genotype. Representative image of $n = 2$ biological replicates.

(E) Above: growth curve of *E. coli* expressing the indicated plasmid. Arrows indicate the time each culture was infected with phage T5 at the indicated multiplicity of infection (MOI). Below: efficiency of plating of the phage present in each sample at the indicated time points. Data represent the mean \pm SEM of $n = 3$ biological replicates.

(F) Scaled trident entropy (SC) values (see STAR Methods) for individual residues of bNACHT01-like proteins. The trident entropy for each column of the alignment, including both the NACHT module and SNaCT domain, is scaled with respect to the top quartile. Positions with values greater than 0 are those with diversity in the top quartile.

(G) Distribution of trident entropy (S) values across NACHT and SNaCT modules in bNACHT01-like proteins showing significantly different mean trident entropy. Values were compared using a two-sample t test.

NACHT modules are also found in fungi where HetE/D proteins can mediate self/non-self discrimination after two hyphal cells have fused their cytosols, so-called heterokaryon incompatibility.^{22–24}

NACHT proteins have been rigorously investigated in eukaryotes, but little is known about their roles in bacteria. The NACHT module belongs to the large family of STAND NTPases, which describes many divergent proteins. Both active and inactive (with disrupted Walker A/B motifs) STAND NTPase domains were previously identified computationally in predicted antiviral conflict systems that are enriched in multicellular bacteria.^{25,26} STAND NTPases were also observed in the antiviral ATPase/NTPase of the STAND superfamily (AVAST) antiphage signaling system,^{3,27} as well as in prophage-encoded antiphage systems,⁴ but NACHT domains were not specifically recognized in these studies.

We undertook an exhaustive bioinformatic analysis of NACHT module-containing proteins across publicly available genomes of both eukaryotes and prokaryotes. We started with identifying NACHT modules based on the amino acid sequence and predicted structural features. The NACHT module belongs to the STAND-Cdc6-Orc family of AAA+ NTPases, which in turn belong to the ASCE division of P loop NTPases. STAND NTPase modules are unified structurally by a characteristic C-terminal extension to the core AAA+ domain in the form of a winged helix-turn-helix (wHTH) domain, also referred to as a winged helix domain (WHD), which they share with Cdc6-Orc AAA+ and transposase ATPases.^{28–30} Within the STAND clade, the NACHT subclade is unambiguously separated from other STANDs based on characteristic motifs, including the D[GAS]hDE signature (a small amino acid directly C-terminal to the Mg²⁺-coordinating aspartate within the Walker B motif, followed by two acidic residues one position away), as well as signatures in the region N-terminal to the NTPase domain and the above-mentioned C-terminal wHTH extension (Figure S1).^{28,29}

Our analysis identified approximately 15,000 unique bacterial NACHT proteins (Table S1). They are encoded by about 9%–10% of complete, published bacterial genomes (Figure S2; Table S2). NACHT proteins are found throughout the bacterial superkingdom, including in the genomes of pathogenic bacteria, members of the human gut microbiome, and other important bacteria from environmental niches. Some bacterial phyla show a much higher tendency than average to encode NACHT proteins (Figure S2): we found that 58% of the cyanobacteria, 25% of the actinobacteria, and 24% of the deltaproteobacteria encode NACHT proteins (Figure S2D). Cyanobacteria also tend to display a large number of paralogous versions per genome—for instance, a record number of 23 paralogous NACHT proteins are seen in *Rivularia* sp. PCC 7116 (Figure S2A). Moreover, organisms with 3 or more NACHT proteins have a significant tendency to be multicellular bacteria (Figure S2B, *p* value = 1.0089e–12). Notably, the multiple copies of the NACHT proteins in these organisms tend to possess distinct effector and sensor domains, suggesting that they are not merely duplications representing iterations of the same theme, but a diversified biochemical repertoire potentially optimized to deal with the unique immune challenges related to multicellularity. This situation mirrors the previously described class of immunity

and apoptosis mechanisms shared by a range of multicellular bacteria.^{25,26}

The NACHT modules from a representative subset of these proteins were aligned and related proteins were grouped into clades (Figures 2 and S3; Table S2). By aligning proteins based on the NACHT module, this analysis was independent of fused protein domains on each polypeptide. Our analysis also included NACHT module-containing proteins from Archaea and eukaryotes, which allowed us to group related proteins from different domains of life into a total of 25 major clades and establish evolutionary relationships (Figures 2 and S3; Tables S2 and S3). Phylogenetic analysis suggests that proteins with more closely related NACHT modules tend to have similar C-terminal sensor domains, possibly due to the need for the C termini to coevolve with the NACHT domain to maintain autoinhibitory function. Conversely, the effector domains to which the NACHT module is fused can vary more dramatically between species of the same genus or bacterial lineage, suggesting that the effector domains may be in an arms race with viral inhibitors that target them.

The C-terminal regions of bacterial NACHT proteins can be placed into different, broad categories: the antigen receptor or infection signal recognition type, those that have transmembrane (TM) domains, those with short C-terminal extensions, or those with a combination of these features (Tables S2 and S4). There are two types of antigen receptor-type domains: formylglycine-generating enzyme sulfatase (FGS) domains and supersecondary structure-forming tandem repeats (e.g., LRR, tetratricopeptide repeat [TPR], and Huntington, elongation factor 3, PR65/A, TOR [HEAT]). Antigen receptor-type and TM domains are found across many different clades; however, bacterial NACHT proteins with short C-terminal extensions that lack supersecondary structure-forming elements, such as bNACHT01, are predominantly found in the monophyletic clade 14 (Figures 2 and S3). These and other characteristics that predominate each NACHT clade are annotated (Figure S3) and a quantification of the frequency of NACHT protein architectures is found in Table S4.

The N-terminal regions of bacterial NACHT proteins encode many enzymatic domains that have previously been associated with biological conflict, including nucleases (RNases and DNases), peptidases, nucleotide signal-generating or degrading domains, and NAD⁺-targeting enzymes (toll/interleukin receptor [TIR] and Sirtuin).^{31–34} Other domains also include predicted “adaptors” that lack a predicted enzymatic function but may mediate interactions with other factors such as effector-associated domains, death-like domains, RNA-binding domains, transcription regulatory domains, and cyclic nucleotide sensors (Tables S1, S2, and S4).

The tripartite domain architecture we observe in bacterial NACHT proteins is consistent with the domain architectures previously observed in eukaryotic NACHT proteins.^{28,29} The central NACHT in mammalian NLRs is flanked by an N-terminal “effector” domain that coordinates signaling and a C-terminal sensor domain that often consists of supersecondary structure-forming tandem repeats, such as LRRs. The similarity of the tripartite domain architecture in bacterial and mammalian NACHT proteins suggests that the sensor and effector domains

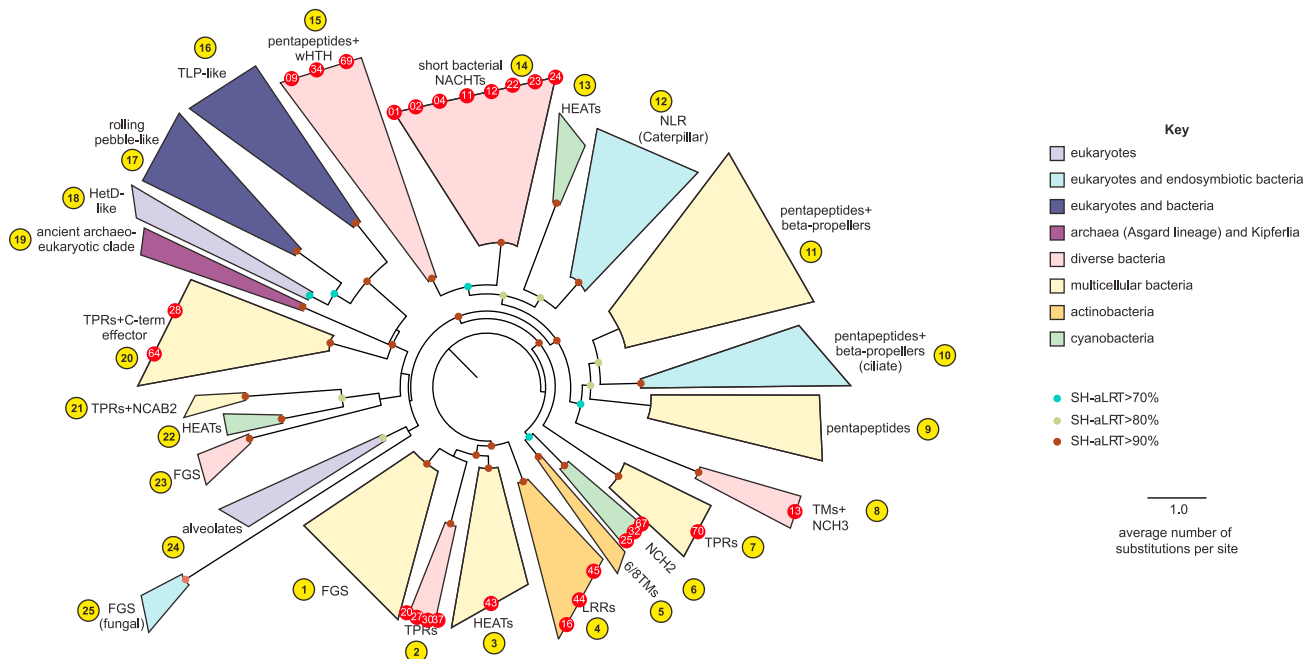


Figure 2. NACHT module-containing proteins in bacteria are widespread and diverse

A sequence-based phylogenetic tree of NACHT modules was generated using NACHT module-containing proteins from eukaryotes and prokaryotes. The NACHT module, not accessory domains, was used for tree building. Clades are color coded based on the indicated key and numbered arbitrarily in yellow circles. Red dots indicate the bacterial NACHT proteins from each clade that were selected for analysis in this study. Bootstrap values are provided where applicable. See [Figure S2](#) for bNACHT gene distribution, [Figure S3](#) for representative domain architectures from each clade, and [Table S4](#) for the most common domain architectures found in each clade. Additional details on genes used to construct the phylogenetic tree can be found in [Table S2](#), and [Table S1](#) contains a full list of all NACHT module-containing proteins identified. See [Table S3](#) for tree topology tests used to validate proposed evolutionary relationships.

of canonical eukaryotic NLRs represent a broader organizational strategy across the tree of life.²⁸ Therefore, we suggest that these bacterial NACHT proteins are NLR-related, even though many of them lack the LRRs required to be classified as true NLRs. These observations further imply that the role of the NACHT module is to act as a signaling hub that transduces the detection of an invader signal into diverse biochemical outputs, enabling the host to respond to a threat.

Although most of the effector domains on bacterial NACHTs are found at the N terminus of the protein, before the core NACHT module, some occur at the extreme C terminus of the protein, after tandem repeat or FGS domains. Published structures of metazoan NLRs suggest that, in these cases, the C-terminal effectors are likely in a similar position as the N-terminal effectors in the folded polypeptide. We predict that the toroidal or helical elements of the C terminus may allow the effector domain to maintain its normal spatial location, despite being located at the opposite terminus of the protein.

Evolutionary history of NACHT proteins

Phylogenetic relationships between prokaryotic and eukaryotic NACHT proteins suggest these genes have horizontally transferred from prokaryotes to eukaryotes on multiple occasions. The clades in [Figure 2](#) are categorized by the organisms that are most represented in that clade. This includes eukaryotes, a mixture of eukaryotes and bacteria, as well as various groups

of prokaryotic organisms. Eukaryotic NACHT modules are found in multiple clades ([Figures 2](#) and [S3](#)), suggesting that NACHT modules were acquired on several distinct occasions and subsequently experienced lineage-specific expansions.²⁸ Fungi have acquired NACHT proteins from multiple horizontal gene transfer events and one of these resulted in the expansion of the heterokaryon incompatibility NACHT proteins (clade 18, HetE/D-like). In the mammalian lineage, there are three distinct clades of NACHT modules. The first of these is telomerase associated protein 1 (TEP1), named for the TEP1 (previously named TP-1), found in clade 16, and was acquired early in eukaryotic evolution from bacteria. The second is the Rolling pebbles clade, a sister group of the fungal HetD/E (clade 17, e.g., human neural development protein TANC2). The third, typified by the mammalian NLR/Caterpillar NACHT proteins, (clade 12) represents a separate transfer from bacteria.

Despite these postulated horizontal gene transfer events being ancient, clade 12 includes extant prokaryotic NACHT proteins encoded by *Rickettsiales*. The *Rickettsiales* are an order of obligate intracellular bacteria that have coevolved extensively with animals. Similarly, clade 10 includes NACHTs expanded in fungi that were horizontally acquired from their bacterial endosymbionts. Thus, our phylogenetic analysis suggests that metazoan and fungal hosts acquired their NACHT genes involved in immune mechanisms from obligate intracellular symbionts/pathogens. This likely origin of metazoan and fungal NLRs

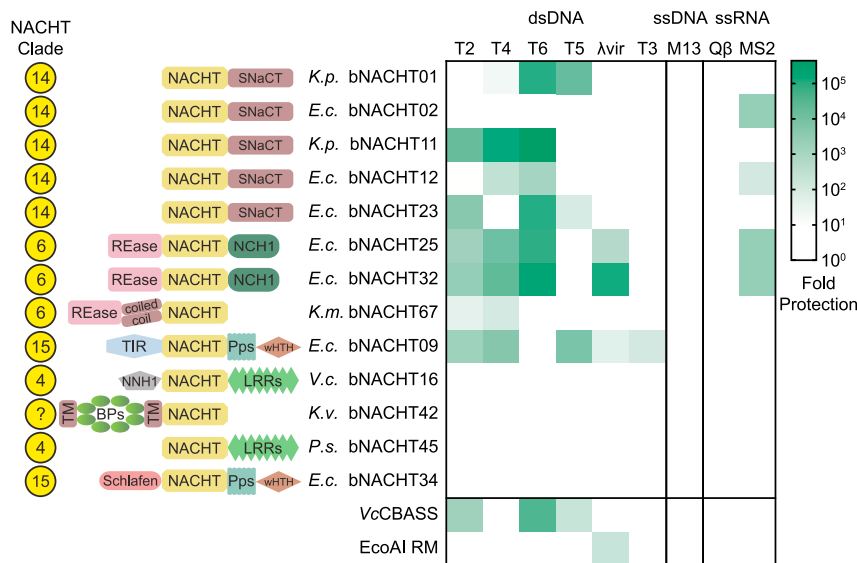


Figure 3. Bacterial NACHT proteins are anti-phage

Heatmap of fold defense provided by the indicated bNACHT gene for a panel of diverse phages. *E. coli* expressing the indicated defense system were challenged with phages and fold defense was calculated for each defense system-phage pair by dividing the efficiency of plating (in PFU/mL) on empty vector by the efficiency of plating on defense system-expressing bacteria. The NACHT clade, domain architecture, and species of origin for each bNACHT are shown. bNACHT genes displayed in this figure are a subset of the 27 candidates interrogated, selected based on their robust antiphage activity or the diversity of domain architectures sampled. *Vibrio cholerae* CBASS (VcCBASS) and *E. coli* UPEC-36 restriction-modification system (EcoAI RM) were included as positive controls. Data represent the mean of $n = 3$ biological replicates. *Escherichia coli* (*E.c.*), *Klebsiella michiganensis* (*K.m.*), *Klebsiella pneumoniae* (*K.p.*), *Klebsiella varicola* (*K.v.*), *Pseudomonas* sp. LAIL14HWK12:16 (*P.s.*), and *Vibrio campbellii* (*V.c.*). Domain abbreviations as described in Figure S3. See Table S5 and Figures S4 and S5 for details on all 27 bNACHT genes analyzed. See Figure S6 for the raw efficiency of plating data.

and NLR-related proteins, from intracellular bacteria, stands in contrast to the potential horizontal transfer of STING from an extracellular bacterial symbiont.^{6,10}

Multiple bacterial NACHT proteins provide a broad spectrum of antiphage immunity

The bNACHT01 protein was potently antiphage (Figures 1 and S1); however, our bioinformatic analysis demonstrated that there are many additional clades of NACHT proteins in bacteria (Figures 2 and S3). To measure the breadth of antiphage activity of NACHT proteins, we expressed 27 representative NACHT module-encoding genes in our *E. coli*-based phage resistance assay (Figures 3, S5, and S6; Table S4). Representatives were selected based on protein domain, the similarity of domain architecture to eukaryotic NACHT proteins, and the phylogenetic distance of the source genome to *E. coli* (to recapitulate native host cell conditions). Specifically, we focused on proteins primarily from the family Enterobacteriaceae to increase the likelihood that proteins would be functional in our *E. coli* heterologous system and/or recognize *E. coli*-specific phages. Bacterial NACHT proteins were expressed from promoters in their native genomic context and were only rarely within poorly conserved operons (Table S5). Bacteria were challenged with a diverse panel of double-stranded DNA (T2–T6, λ_{vir}), single-stranded DNA (M13), and positive-sense single-stranded RNA (MS2, Qβ) phages. We also included a previously characterized CBASS system from *Vibrio cholerae* and a restriction-modification system from *E. coli* as positive controls in these experiments. Diverse bacterial NACHT proteins from different clades exhibited robust antiphage activity across a wide range of phages (Figures 3, S5, and S6). Intriguingly, some bacterial NACHT proteins defended against both DNA and RNA phages.

Our interrogation of a wide range of bacterial NACHT proteins demonstrates that bacterial NACHT proteins are related

to metazoan NLRs because they share several features: (1) a broad role in antipathogen immune activity, (2) conserved sequence and structural characteristics of their NACHT modules, and (3) a characteristic tripartite protein architecture (C-terminal supersecondary structure-forming tandem repeats, central NACHT, and N-terminal effector). These findings extend the scope of STAND NTPases involved in viral defense beyond the previously identified AVAST systems^{3,27} to include NACHT proteins.

NACHT proteins are activated in response to phage infection

We next sought to understand how bNACHT proteins restrict virion production and selected bNACHT09 and bNACHT25 for in-depth investigation because their N-terminal effector domains could be readily identified as TIR and restriction endonuclease (REase) domains, respectively. We first confirmed that both proteins functioned in single-copy when expressed under their native promoters from the chromosome of *E. coli* and conferred similar magnitudes of protection to phages T4–T6 when compared with low-copy plasmid-based expression (Figures 4A, 4C, S5B, and S5C). Next, we investigated TIR domain activation upon phage infection for bNACHT09. TIR domains in plant resistosomes and other bacterial antiphage systems degrade NAD⁺^{32,33,35} upon activation. We therefore measured NAD(H) in bacteria expressing bNACHT09 and found a dramatic reduction in NAD(H) levels upon phage infection. The decrease in NAD(H) was dependent on phage and the catalytic glutamate of the TIR domain (Figure 4B). We then performed a similar analysis of effector activation for bNACHT25, which encodes a predicted N-terminal REase domain. Infection of bNACHT25-expressing bacteria with MS2 phage resulted in the rapid destruction of plasmid DNA that was dependent on the predicted catalytic aspartate (Figure 4D). Taken together,

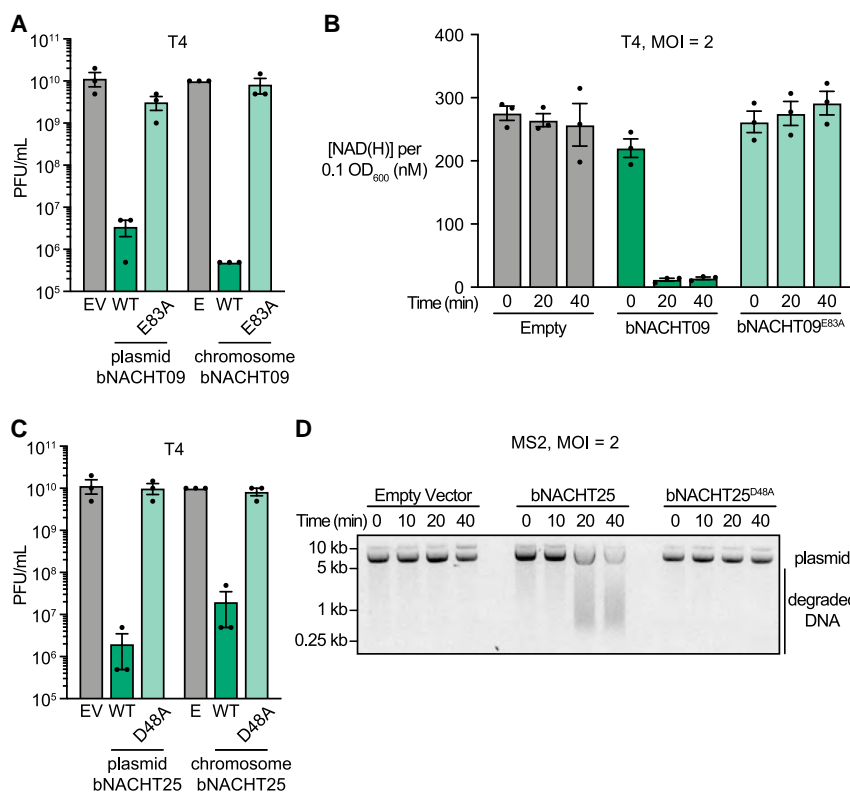


Figure 4. Bacterial NACHT effector modules are activated by phage

(A) Efficiency of plating of phage T4 infecting *E. coli* expressing the indicated genotype from a low-copy plasmid or from the chromosome. See Figure S5 for the efficiency of plating data for phages T5 and T6. (B) Measurement of [NAD(H)] in each sample when normalized to an OD₆₀₀ of 0.1 from *E. coli* expressing the indicated chromosomal genotype at the indicated time points after infection with phage T4. (C) Efficiency of plating of phage T4 infecting *E. coli* expressing the indicated genotype from a low-copy plasmid or from the chromosome. See Figure S5 for the efficiency of plating data for phages T5 and T6. For (A)–(C), empty (E) indicates *E. coli* with the chromosomal expression of a kanamycin resistance cassette and *gfpmut3*. Data represent the mean ± SEM of *n* = 3 biological replicates, shown as individual points.

(D) Visualization of plasmid integrity in *E. coli* expressing the indicated plasmid at the indicated time points after infection with MS2. Data are a representative image from *n* = 3 biological replicates.

these data demonstrate that phage infection results in the activation of the N-terminal effector domains of bNACHT proteins.

Phage proteins alter bacterial NACHT phage defense

We sought to understand how phages alter bacterial NACHT protein signaling by generating phage mutants that evade defense (suppressor mutant phage). Phage T5 was selected for analysis. Wild-type T5 plaque formation is robustly inhibited by bNACHT01 (Figures 1D and 1E); however, when bacteria were infected with a high number of T5 PFU, suppressor mutants capable of escaping bNACHT01 and forming a plaque were isolated (Figure 5A). These mutants were extremely rare, appearing at an average rate of one suppressor for every 5×10^7 PFU of wild-type phage (Table S6). Accordingly, genome sequencing revealed that every suppressor phage encoded at least two mutations that affected the same ORFs: one mutation that altered *orf008* (Genbank: AAX11945.1), an SH3-like fold β -barrel protein, and the other that altered *orf015* (Genbank: AAX11952.1, Table S6), a 5-stranded β -meander protein. The majority of the suppressor mutations identified were missense mutations, although some included frameshifts, nonsense, and promoter mutations (Table S6). Genes *orf008* and *orf015* are encoded in the “first-step transfer” region of the T5 genome, a 10 kb section that is injected first into host bacteria and coordinates injection pausing for approximately 5 min before the remainder of the genome is injected.³⁶ During those first minutes of infection, other pre-early genes of the first-step transfer region remodel core processes and shut down signaling within the host cell.³⁷ We were unable to

identify homologs of *orf008* or *orf015* outside of the *Tequintavirus* genus (T5-like phages).

We hypothesized that the low frequency of isolating suppressor phages reflects that T5 must encode mutations in both *orf008*

and *orf015* to evade bNACHT01-mediated protection. To measure the impact of these genes on bNACHT01 antiphage activity, we constructed an assay where bNACHT01 was co-expressed with either *orf008*, *orf015*, or both phage genes, then challenged with phages T4 and T6 (Figures 5B and S7A). bNACHT01 provided 1,000-fold protection against phage T4 in this assay, and the expression of either *orf008* or *orf015* individually had a modest impact on the efficiency of plating. However, the expression of both genes together resulted in a 100-fold recovery of phage T4 virulence (Figure 5B). These data suggest that *orf008* and *orf015* act together to allow phage to evade bNACHT01.

Relatively few bacterial NACHT proteins protected *E. coli* against phage T5 (Figures 3 and S5), and we hypothesized that *orf008* and *orf015* might be broadly responsible for T5 immune evasion. To test this, we selected bNACHT genes that defended against phage T4, but not phage T5, then repeated our assay for measuring the impact of these phage genes. Expression of *orf015* significantly decreased the protection by bNACHT11, 12, 25, and 32 against phage T4 (Figure 5C). Similar results could be obtained for a subset of these genes when phage T2 and T6 were used (Figures S7B and S7C). We next analyzed the effect of *orf008* on bNACHT activity. Interestingly, we found that bacterial growth was inhibited when *orf008* was co-expressed with bNACHT genes but not when co-expressed with an EV (Figure 5D). The growth inhibition was specific to *orf008*; *orf015* did not alter bacterial growth when expressed with bNACHT genes (Figure S7D). Because *orf008* resulted in growth inhibition, we did not measure its impact on phage defense beyond bNACHT01 (Figure 5B). These data demonstrate that two phage

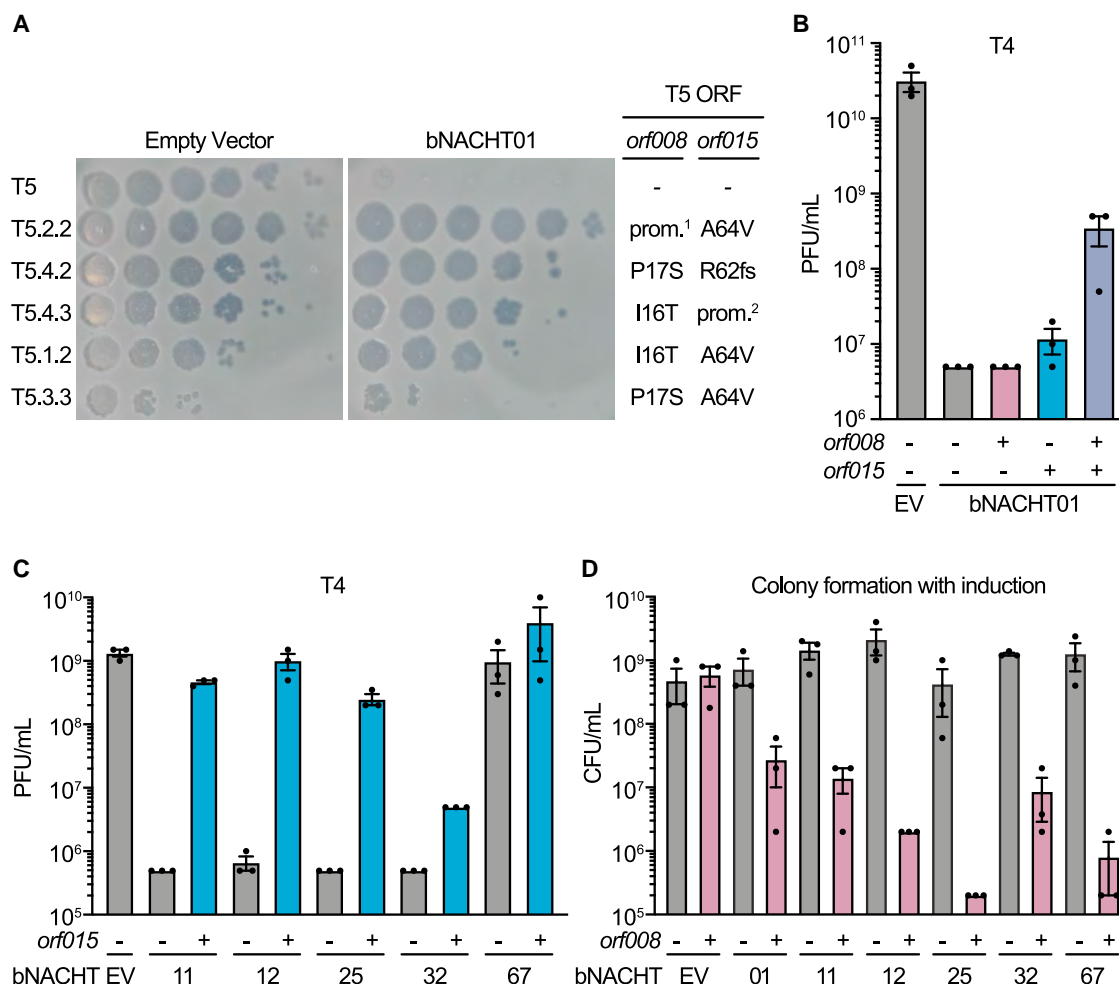


Figure 5. Phage proteins modulate host immune responses

(A) Efficiency of plating of wild-type or suppressor T5 phage when infecting *E. coli* expressing the indicated plasmid. The impact of *orf008* and *orf015* suppressor mutations is indicated. Data are representative images of $n = 3$ biological replicates. Wild-type alleles (-); mutations in the promoter region of *orf008* (prom.¹); frameshift mutations at the indicated position (fs); and mutations deleting *orf009-012* predicted to disrupt the promoter of *orf015* (prom.²) are indicated. See Table S6 for rates of suppressor phage isolation, suppressor mutations identified in *orf008* and *orf015*, and a complete list of mutations identified.

(B) Efficiency of plating of phage T4 when infecting *E. coli* expressing bNACHT01 or an empty vector (EV) on one plasmid and the indicated phage T5 gene(s) on a second plasmid. See Figure S5 for the efficiency of plating of phage T6.

(C) Efficiency of plating of phage T4 when infecting *E. coli* expressing the indicated bNACHT gene on one plasmid and phage T5 *orf015* on a second plasmid. See Figure S7 for the efficiency of plating of phages T2 and T6.

(D) Quantification of colony formation of *E. coli* expressing the indicated bNACHT system on one plasmid and *orf008* on a second plasmid. See Figure S7 for colony formation in the presence of *orf015*. For (B)–(D), the expression of *orf008*, *orf015*, and *mCherry* is IPTG-inducible. (–) symbols denote the induction of an *mCherry* negative control. (+) symbols denote induction of the indicated phage gene. Data represent the mean \pm SEM of $n = 3$ biological replicates, shown as individual points.

genes alter the activity of a wide variety of bNACHT genes and provide evidence for a complicated relationship between phage genes and bNACHT-based host defense systems, where *orf008* activates and *orf015* inhibits bNACHT proteins.

Human disease mutations activate bacterial NLRs

Human NLR protein activation has potent signaling consequences and rare, monoallelic mutations in patients cause serious diseases that include bare lymphocyte syndrome, Crohn's/inflammatory bowel disease, and autoimmune conditions.^{16,38} A subset of these diseases are inflammasomopa-

thies, which are point mutations in NLRs that result in stimulus-independent hyperactivation of inflammasome signaling. Patients encoding H443P mutations in NLRC4 display familial cold autoinflammatory syndrome (FCAS)³⁹ and H443L mutations also result in NLRC4 activation in cells.¹⁵ Histidine 443 is a highly conserved and defining residue located within the wTHH (WHD) domain of the NACHT module (Figure S7E). In NLRC4, H443 is thought to interact with ADP to stabilize an inactive conformation.¹⁵

Given the high degree of conservation between human and bacterial NACHT modules, we hypothesized that mutations that

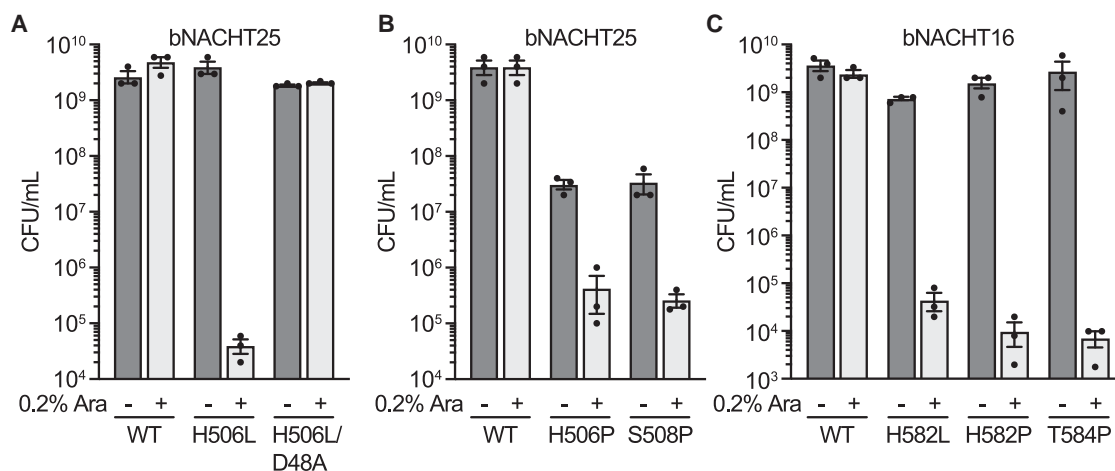


Figure 6. Human disease-associated mutations hyperactivate bacterial NACHT proteins

(A and B) Quantification of colony formation of *E. coli* expressing wild-type (WT) bNACHT25 or alleles with the indicated mutations.

(C) Quantification of colony formation of *E. coli* expressing bNACHT16 with the indicated mutations. See Figure S7 for an alignment of NLRC4, bNACHT16, and bNACHT25. For (A)–(C), gene expression was induced with arabinose. Symbols denote induction (+) or lack of induction (–). Data represent the mean \pm SEM of $n = 3$ biological replicates, shown as individual points.

hyperactivate human NLRs might also hyperactivate bacterial NLR-related proteins. Structure-guided alignments between NLRC4 and bNACHT25 identified the analogous residue to H443 and mutations at this location were constructed in bNACHT25 (Figure S7). The effector domain of bNACHT25 is a predicted Mrr-like REase. When activated, bNACHT25 cleaves DNA, resulting in toxicity to the host cell and/or destruction of the DNA phage chromosome (Figures 3, 4, and S4; Table S5). For this reason, we expressed wild-type bNACHT25 and mutant alleles using an inducible system. Expression of the histidine mutant, but not wild-type bNACHT25, resulted in potent bacterial growth inhibition (Figure 6A). An additional mutation predicted to disrupt the nuclease activity of the effector domain (D48A) rescued the growth inhibition of hyperactive bNACHT25 (Figure 6A). We next interrogated bNACHT16 because this gene is similar to human NLR proteins, encoding LRRs at the C terminus (Figures 3 and S4; Table S5). The introduction of mutations at the H443 equivalent residue of bNACHT16 also resulted in bacterial growth inhibition, consistent with NLR hyperactivation (Figure 6C). Histidine to leucine mutations, shown to synthetically activate NLRC4,¹⁵ or histidine to proline mutations, found in patients with inflammasomopathies,³⁹ both inhibited growth equivalently (Figure 6). Our findings for bNACHT16 suggest that even though we did not observe a phage protection phenotype, the protein is expressed and capable of effector activation in *E. coli*, despite originating in *Vibrio campbellii*. bNACHT16 may therefore be unable to respond to the phages tested due to a lack of the appropriate stimulus or host components required for phage sensing.

The disease-associated mutation at S445 also results in hyperactivation of NLRC4.^{40,41} To test whether we could recapitulate the effects of mutation to this residue in bacteria, we mutated the corresponding residue in bNACHT25 (S508P) and bNACHT16 (T584P). Overexpression of both the bNACHT25 and the bNACHT16 mutants resulted in the inhibition of bacterial growth (Figures 6B and 6C). These data demonstrate that

NACHT modules in humans and bacteria can be hyperactivated by similar mutations, suggesting these proteins have a similar mechanism of effector domain activation.

Our analysis of *orf008* and *orf015* from phage T5 demonstrated that these proteins alter bNACHT-dependent phage resistance and growth (Figure 5). To further characterize the effect of these two phage genes on bNACHT activity, we co-expressed *orf008* and *orf015* with a hyperactive allele of bNACHT25, then measured bacterial growth. As predicted from our previous experiments demonstrating *orf008*-mediated activation of bNACHT proteins (Figure 5), *orf008* did not appreciably alter colony formation as hyperactive bNACHT25 already leads to growth inhibition (Figures 6A and 7). However, the *orf015* gene was sufficient to rescue the growth inhibition of the bNACHT25 hyperactive allele (Figure 7A). These data are consistent with the impact of *orf015* on bNACHT25-mediated resistance during infection (Figure 5C). Similar results were also obtained for bNACHT01 when this protein was hyperactivated by overexpression, a method used to activate other anti-phage systems (Figure 7B).⁴²

To analyze the impact of *orf008* and *orf015* on bNACHT25 effector activation, we returned to our assay for plasmid integrity and measured the REase activity of bNACHT25 during co-expression. These experiments showed that the induced expression of hyperactive alleles of bNACHT25 results in plasmid destruction, which is inhibited by *orf015*. Conversely, expression of *orf008* with wild-type bNACHT25 results in REase activation (Figure 7C). These data demonstrate that in the absence of phage, *orf015* interrupts, and *orf008* activates, bNACHT signaling.

DISCUSSION

Here we identify that NACHT module-containing proteins are abundant and widespread in the genomes of bacteria where

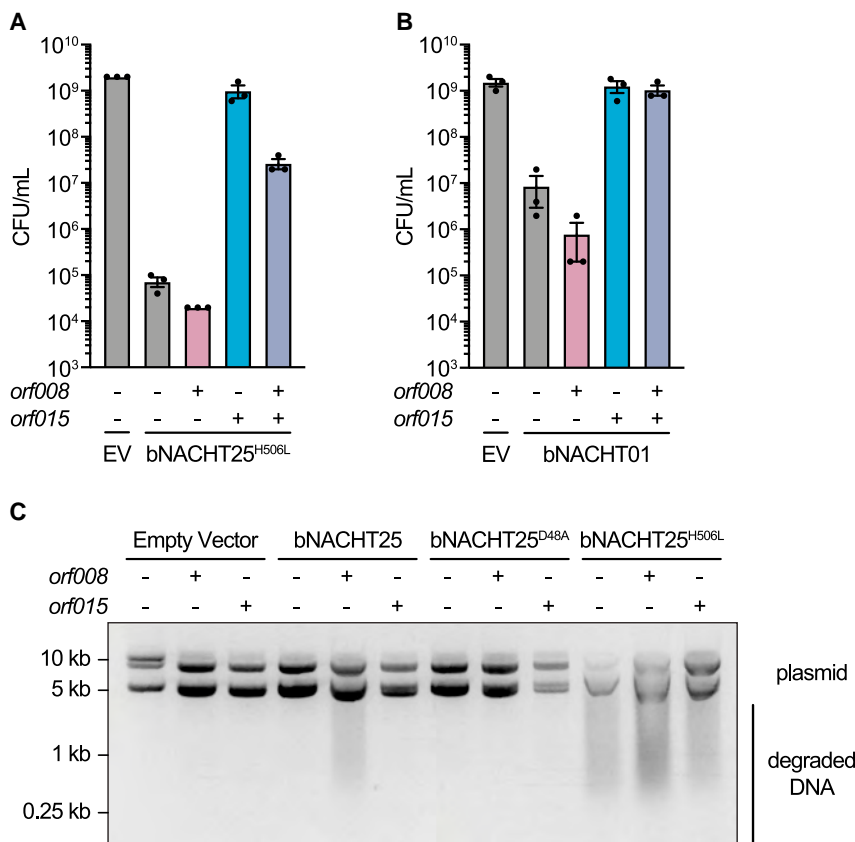


Figure 7. Phage proteins alter the activity of hyperactive bacterial NACHT proteins

(A and B) Quantification of colony formation of *E. coli* expressing a bNACHT gene with the indicated genotype on one plasmid and the indicated phage T5 gene(s) on a second plasmid. Data represent the mean \pm SEM of $n = 3$ biological replicates, shown as individual points.

(C) Visualization of plasmid integrity in *E. coli* expressing a bNACHT gene with the indicated genotype on one plasmid and the indicated phage T5 gene on a second plasmid. For (A)–(C), the expression of *or008*, *or015*, and *sfGFP* was induced with IPTG. Symbols denote the induction of an *sfGFP* negative control gene (–) or induction of the indicated phage gene (+). Expression of the indicated bNACHT gene or empty vector (EV) was arabinose-inducible. Data are an image representative of $n = 3$ biological replicates.

they are potent phage defense systems. Bacterial and animal NACHT proteins are highly similar in their overall domain architecture, the predicted structure of their NACHT module, and their role in immune signaling. These data establish bacterial NACHT proteins are related to eukaryotic NLRs. In support of a shared molecular mechanism of NACHT module activation, point mutations that hyperactivate NACHT modules in human cells also hyperactivate NACHT modules in bacteria.^{15,39,40} Hyperactivated alleles of bacterial NACHT proteins inhibited the growth of bacteria. Further, phage infection also appeared to inhibit the growth of NACHT protein-expressing bacteria, suggesting that these systems may inhibit phage replication via abortive infection. Abortive infection is a form of programmed cell death that interrupts the viral life cycle by prematurely destroying a host component essential to virion production.⁴³ In this way, the antimicrobial signaling outcome of bacterial NLR-related proteins may also be similar to mammalian inflammasomes, which initiate a caspase-dependent programmed cell death called pyroptosis when activated.⁴⁴ We anticipate that further understanding of the molecular mechanisms of bacterial NACHT protein signaling will provide valuable insights into human NLRs.

Our expansive bioinformatic analysis found that bacteria encode the largest diversity of NACHT module sequences compared with other superkingdoms, which suggests that this protein module first evolved in bacteria before being acquired into the genomes of eukaryotes. However, not all eukaryotic NACHT module sequences are monophyletic and each often

clusters with distinct groups of bacterial NACHT proteins, implying that horizontal gene transfer of NACHT modules from prokaryotes to eukaryotes has occurred on multiple occasions. Evidence for one transfer event is found in NACHT module clade 12, which groups mammalian NLRs (aka Caterpillar genes) with bacterial NACHT proteins from *Rickettsiales*, an order of intracellular bacteria. This observation suggests that metazoans acquired their NLRs from *Rickettsiales*. A similar horizontal gene transfer event has been suggested for the innate immune gene STING; however, the most probable bacterial source for that event is the Bacteroidetes.⁶ Both Bacteroidetes (living extracellularly as a symbiont) and *Rickettsia* (living intracellularly) have intimate interactions with eukaryotes yet distinctly different lifestyles. The shared evolutionary history of NACHT genes may enable future investigators to take advantage of studying bacterium-phage interactions to learn about cryptic aspects of human NLR signaling.

Fungi also encode NACHT proteins that are uniquely suited to their lifestyle. The HET-D and HET-E proteins from the filamentous fungus *Podospora anserina* are NACHT proteins that mediate kin recognition after two cells have fused their cytoplasms. When kin cells expressing these proteins fuse, the subsequent heterokaryon survives; however, when non-kin cells expressing HET-D or HET-E fuse, the NACHT protein initiates programmed cell death.^{45,46} HET-E/D recognizes allelic differences in the HET-C protein to distinguish kin, i.e., self from non-self.⁴⁷ This phenomenon is known as heterokaryon incompatibility. In related systems, heterokaryon incompatibility has been shown to restrict the spread of endogenous viruses between non-kin fungi.^{48,49} Thus, in fungi, as in animals and bacteria, NLR-related proteins are part of the innate immune system.

NLRs within the mammalian inflammasome require additional factors to induce cell death. NLRC4 requires the pore-forming protein Gasdermin D to execute cell death (pyroptosis).^{44,50} Gasdermin D homologs can also be found in fungi, where they

mediate heterokaryon incompatibility, and in bacteria, where they mediate antiphage signaling.^{13,51,52} The bacterial NLR-related proteins interrogated here do not require Gasdermin D homologs for signaling. However, they are encoded in operons also coding for NACHT proteins in a few bacteria (Table S2). Heterokaryon incompatibility loci are highly polymorphic across fungi and there are many more than *het-d/e* (NLRs) and *rcd-1* (Gasdermin D).^{22,53} These observations suggest that heterokaryon incompatibility loci, like bacterial antiphage systems, may be an important repository for identifying mammalian innate immune genes.

Bacterial NACHT proteins are the first example of an innate immune antiphage system in bacteria capable of defending against RNA phages. Although adaptive immune systems like CRISPR can be programmed to defeat RNA phages,⁵⁴ this may not represent their natural function. Bacterial NLRs capable of recognizing RNA phages also recognize DNA phages, suggesting that the stimulus recognized is highly conserved between disparate viruses. We do not yet know what the stimulus might be, or if the stimulus is the same for all bacterial NACHT proteins. However, we are able to synthetically activate these proteins using mutations that hyperactivate mammalian NLRs. Many NACHT-associated effector domains are highly conserved and found across multiple known and predicted antiphage systems but remain as yet biochemically uncharacterized. Given that they cannot be readily activated in the absence of a phage (which might be unknown), synthetic activation might prove highly useful to study the large array of effector domains fused to the N terminus of bacterial NACHTs. Some noteworthy examples include: (1) the Schlafen RNase domain found at the N terminus of bNACHT34 that is related to human Schlafen proteins involved in HIV1 restriction⁵⁵; (2) the polyribonucleotide nucleotidyl transferase (PNPase) domain that is predicted to degrade nucleotides or NAD⁺ by removal of the base¹¹; and (3) bacterial domains related to the death-superfamily domains found in metazoan apoptosis.^{25,26}

Our data support a unifying role for proteins encoding NACHT modules and related STAND NTPases as mediators of innate immunity across the tree of life. NACHT module-encoding NLRs in mammals initiate inflammation and are potentially antimicrobial. Fungal NACHT proteins mediate heterokaryon incompatibility, which can stop viral transmission. Here, we demonstrate that bacterial NACHT proteins are antiphage. Land plants also show an expansion of the antibacterial and antiviral R (NB-ARC) proteins that contain another clade of STAND NTPase modules, i.e., the AP-ATPase, which is a sister group of the NACHT clade. Further, a contemporary analysis of bacterial STAND NTPases outside the NACHT clade by Gao *et al.* shows that those proteins use a similar tripartite domain architecture to recognize structural motifs of specific phage proteins.²⁷ Thus, it appears that the NACHT and related STAND modules define an architectural theme that is especially suited for immune signaling and apoptosis. One potential explanation is that these modules can serve as switches that combine sensing of infection signals (either pathogen or endogenous molecules), activation-threshold setting, signal transduction, and effector deployment, all in a single protein.^{56,57}

Understanding the unique qualities of the NACHT module is an exciting area for future investigation.

Limitations of the study

There are important limitations to our study. First, NLRs are defined as nucleotide-binding domain and LRR-containing proteins,²¹ not by the presence of a NACHT domain. As our analysis focused solely on NACHT domain-containing proteins, we are unable to draw conclusions about NLRs that encode other STAND NTPases such as the AP-ATPase domains found in plants. Second, we have measured phage resistance using a heterologous system in *E. coli*, which limits our ability to test bNACHT proteins from outside the Enterobacteriaceae family as these are often poorly expressed or unable to recognize *E. coli* phages. This limits our ability to interrogate the *Rickettsiales* NACHT proteins found in clade 12. Despite this limitation, we were able to test many of the most common domain architectures found in Enterobacteriaceae (Table S4). We found these genes are antiphage when expressed from the chromosome but have been unsuccessful in investigating *E. coli* strains natively expressing NACHT proteins because these strains are refractory to genetic manipulation and encode many redundant antiphage systems that obscure interpretation. Third, we do not understand the effector mechanism of bNACHT01 and other clade 14 proteins that lack N-terminal effector domains. These proteins inhibit growth upon activation; however, without a predicted effector signaling outcome we cannot conclusively show that growth inhibition is cell death, and thus abortive infection. Nevertheless, other bNACHT proteins with catalytic effector domains are likely to limit infection through abortive infection by destroying NAD(H) (TIR domain, bNACHT09) and degrading the genome (REase domain, bNACHT25).

STAR★METHODS

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SUPPLEMENTAL INFORMATION

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AUTHOR CONTRIBUTIONS

Conceptualization, E.M.K., A.N.C., L.A., and A.T.W.; methodology, E.M.K., A.N.C., and A.T.W.; software, A.M.B. and L.A.; formal analysis, A.M.B. and L.A.; investigation, all authors; writing – original draft, E.M.K., A.M.B., L.A., and A.T.W.; writing – review & editing, all authors; visualization, E.M.K. and A.M.B.; supervision, L.A. and A.T.W.; funding acquisition, L.A. and A.T.W.

DECLARATION OF INTERESTS

The authors declare no competing interests.

INCLUSION AND DIVERSITY

We support inclusive, diverse, and equitable conduct of research. One or more of the authors of this paper self-identifies as an underrepresented ethnic minority in their field of research or within their geographical location. One or more of the authors of this paper self-identifies as a gender minority in their field of research. One or more of the authors of this paper self-identifies as a member of the LGBTQIA+ community. One or more of the authors of this paper received support from a program designed to increase minority representation in their field of research. While citing references scientifically relevant for this work, we also actively worked to promote gender balance in our reference list.

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STAR★METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Antibodies		
Rabbit polyclonal anti-FLAG	Sigma-Aldrich	Cat#F7425; RRID: AB_439687
Mouse monoclonal anti- <i>E. coli</i> RNA polymerase B	Biolegend	Cat#663006; RRID: AB_2565555
IRDye® Goat 680RD anti-Mouse	Li-Cor	Cat#926-68070; RRID: AB_10956588
IRDye® Goat 800CW anti-Rabbit	Li-Cor	Cat#926-32211; RRID: AB_621843
Bacterial and virus strains		
See Table S7 for a complete list of bacterial strains	N/A	N/A
See Table S7 for a complete list of virus strains	N/A	N/A
Chemicals, peptides, and recombinant proteins		
NAD (β-Nicotinamide Adenine Dinucleotide)	Gold Biotechnology	Cat#N-030-1
Carbenicillin	Gold Biotechnology	Cat#C-103-50
Chloramphenicol	Gold Biotechnology	Cat#C-105-25
Tetracycline Hydrochloride	Gold Biotechnology	Cat#T-101-25
Kanamycin Monosulfate	Gold Biotechnology	Cat#K-120-10
IPTG (Isopropyl-beta-D-thiogalactoside)	Gold Biotechnology	Cat#I2481C
X-Gal (5-Bromo-4-chloro-3-indolyl β-D-galactopyranoside)	Gold Biotechnology	Cat#X4281C
PBS (Phosphate Buffered Saline)	Corning	Cat#21-040-CMX12
DNase I (RNase-free)	New England BioLabs	Cat#M0303S
RNase A (Bovine ribonuclease A from pancreas)	VWR Chemicals	Cat#E866-5ML
EDTA (Ethylenediaminetetraacetic acid) Disodium, dihydrate	Gold Biotechnology	Cat#E-210
Critical commercial assays		
NAD/NADH-Glo Assay	Promega	Cat#G9071
DNeasy Cleanup Kit	Qiagen	Cat#69506
PureLink RNA Minikit	Invitrogen	Cat#12183018A
SuperScript III First-Strand Synthesis System	Invitrogen	Cat#18080051
Tagment DNA Enzyme and Buffer Small Kit	Illumina	Cat#20034197
NEBNext® Multiplex Oligos for Illumina®	New England BioLabs	Cat#E7335S
NEBNext® dsDNA Fragmentase	New England BioLabs	Cat#M0348S
Oligonucleotides		
oAC0025: gccaaaacagccaagctttgggtgtaactagccaagcag	This Study	N/A
Recombinant DNA		
See Table S7 for a complete list of plasmids	N/A	N/A
Software and algorithms		
Geneious Prime® 2022.2.2	Biomatters Ltd.	RRID: SCR_010519
PSI-BLAST	Altschul et al. ⁵⁸	RRID: SCR_001010
JACKHMMER	Potter et al. ⁵⁹	RRID: SCR_005305
BLASTClust	NCBI	RRID: SCR_016641
HHpred	Zimmermann et al. ⁶⁰	RRID: SCR_010276
PFAM	Mistry et al. ⁶¹	RRID: SCR_004726
PDB	Berman et al. ⁶²	RRID: SCR_012820
Kalign	Lassmann et al. ⁶³	RRID: SCR_011810
Muscle	Edgar et al. ⁶⁴	RRID: SCR_011812

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REAGENT or RESOURCE	SOURCE	IDENTIFIER
JPred	Drozdetskiy et al. ⁶⁵	RRID: SCR_016504
RoseTTa Fold	Baek et al. ⁶⁶	N/A
IQ-TREE	Minh et al. ⁶⁷	RRID: SCR_017254
FigTree	tree.bio.ed.ac.uk	RRID: SCR_008515
MAFFT	Katoh et al. ⁶⁸	RRID: SCR_011811
Adobe Illustrator 2021	Adobe	RRID: SCR_010279
GraphPad Prism 9.2.0	GraphPad Software	RRID: SCR_002798

RESOURCE AVAILABILITY**Lead contact**

Correspondence and requests for materials should be addressed to and will be fulfilled by the lead contact, Aaron Whiteley (aaron.whiteley@colorado.edu).

Materials availability

Strains, plasmids, and phages used in this study are available upon request.

Data and code availability

- All data reported in this paper will be shared by the [lead contact](#) upon request.
- This paper does not report original code.
- Any additional information required to reanalyze the data reported in this paper is available from the [lead contact](#) upon request.

EXPERIMENTAL MODEL AND SUBJECT DETAILS**Bacterial strains and culture conditions**

E. coli strains used in this study are listed in [Table S7](#). *E. coli* were cultured in LB medium (1% tryptone, 0.5% yeast extract, and 0.5% NaCl) shaking at 37 °C and 220 rpm in 1–3 mL of media in 14 mL culture tubes, unless otherwise indicated. Where applicable, carbenicillin (100 µg/mL), chloramphenicol (20 µg/mL), and tetracycline (15 µg/mL) were added. We defined “overnight” bacterial cultures as 16–20 hours post-inoculation from a glycerol stock or single colony. All strains were frozen for storage in LB plus 30% glycerol at –70 °C. *E. coli* OmniPir was used for construction and propagation of all plasmids. *E. coli* MG1655 (CGSC6300) was used to collect all experimental data.

E. coli OmniPir was constructed from OmniMAX 2 T1^R *E. coli* (Thermo Fisher Scientific) and pGRG36*pir-116* as previously described.⁶⁹ Briefly, the *pir116* gene was integrated at the Tn7 attachment site by conjugating pGRG36*pir-116* into OmniMAX *E. coli*, cultivating bacteria at the permissive temperature with arabinose induction, then curing the plasmid at 42 °C. Integration of *pir116* was confirmed by PCR and retention of the F' plasmid was confirmed by tetracycline resistance. *E. coli* MG1655 F+ strain was constructed by isolating the F' plasmid from OmniPir following a previously described protocol.⁷⁰ Briefly, 3 mL of an overnight culture was pelleted and resuspended in 200 µL resuspension buffer (50 mM glucose, 10 mM EDTA, 10 mM Tris-Cl, pH 8.0). 400 µL Buffer P2 (0.2 M NaOH, 1% sodium dodecyl sulfate) was then added and the sample was incubated for 5 minutes at 25 °C. 300 µL of ammonium acetate (7.5 M) and 300 µL chloroform were then added. The sample was incubated at 4 °C for 10 minutes, and pelleted spinning at 21,000 × g for 10 minutes at 25 °C. The supernatant was transferred to a new tube containing 400 µL precipitation solution (30% polyethylene glycol 8000, 1.5 M NaCl) and incubated for 15 minutes at 4 °C. After incubation, DNA was pelleted by centrifugation for 5 minutes at 15,000 × g at 25 °C. The supernatant was discarded, the DNA pellet was resuspended in 100 µL UltraPure water and allowed to dissolve at 4 °C for 2 hours. Purified plasmid was then electroporated into electrocompetent MG1655, followed by selection with tetracycline.

MMCG medium (47.8 mM Na₂HPO₄, 22 mM KH₂PO₄, 18.7 mM NH₄Cl, 8.6 mM NaCl, 22.2 mM Glucose, 2 mM MgSO₄, 100 µM CaCl₂, 3 µM Thiamine, Trace Metals at 0.1× (Trace Metals Mixture T1001, Teknova, final concentration: 5 mM Ferric chloride, 2 mM Calcium chloride, 1 mM Manganese chloride, 1 mM Zinc Sulfate, 0.2 mM Cobalt chloride, 0.2 mM Cupric chloride, 0.2 mM Nickel chloride, 0.2 mM Sodium molybdate, 0.2 mM Sodium selenite, 0.2 mM Boric acid)) with appropriate antibiotics was used to collect all experimental data. When experiments required bacteria expressing two plasmids, strains were grown using reduced antibiotic concentrations to enhance growth rate (MMCG with 20 µg/mL carbenicillin and 4 µg/mL chloramphenicol).

When growing strains that required induction, 100 µM IPTG or 0.2% arabinose was used to induce, as appropriate.

Phage amplification and storage

The phages used in this study are listed in Table S7. Phages were amplified via either liquid or plate amplification using a modified double agar overlay.⁷¹ For liquid amplification, 5 mL mid-log cultures of *E. coli* MG1655 in LB plus 10 mM MgCl₂, 10 mM CaCl₂, and 100 μM MnCl₂ were infected with phage at an MOI of 0.1 and grown, shaking, for 2–16 hours. The supernatant was harvested and filtered through a 0.2 μm spin filter to remove bacterial contamination. For plate amplification, 400 μL of mid-log MG1655 were mixed with 3.5 mL LB soft agar mix (LB with 0.35% agar and 10 mM MgCl₂, 10 mM CaCl₂ and 100 μM MnCl₂) and 100–1,000 PFU. Plates were then incubated for 16 hours at 37 °C. 5 mL of SM buffer (100 mM NaCl, 8 mM MgSO₄, 50 mM Tris-HCl pH 7.5, 0.01% gelatin) was added to the plate and allowed to soak out the phages for 1 hour before SM buffer was collected and passed through a 0.2 μm filter or treated with 1–3 drops of chloroform to remove viable bacteria. All phages were stored at 4 °C in SM buffer or LB.

Validation of phages used in this study

All phages were first tested for F plasmid-dependent infection, which confirmed that only M13, MS2, and Qβ required the F plasmid for successful infection of MG1655, as previously reported.^{72,73}

Genomes of dsDNA phage were purified as previously described.⁷⁴ Briefly, 450 μL of phage lysate (>10⁷ PFU/mL) was treated with DNase I (final concentration 3 × 10⁻³ U/μL) and RNase A (final concentration 3 × 10⁻² μg/μL) and incubated for 1.5 hours at 37 °C to remove extracellular nucleic acids. EDTA was added (final concentration 20 mM) to stop the reaction. Phage genomes were subsequently isolated and purified using the Qiagen DNeasy cleanup kit, starting at the proteinase K digestion step.⁷⁴ Purified phage genomes were sequenced using 200Mbp Illumina sequencing (SeqCenter). Reads were mapped to the following NCBI Genome accessions using Geneious software's *Map to Reference* feature: AP018813.1 (T2), NC_047864.1 (T3), NC_000866.4 (T4), AY587007 (T5), NC_054907.1 (T6), and NC_001416.1 (λvir).

To purify the RNA genomes of MS2 and Qβ, 172.8 μL of phage lysate (>10⁸ PFU/mL) was treated with DNase I (final concentration 3 × 10⁻³ U/μL) in DNase I buffer (10mM Tris-HCl pH 7.6, 2.5 mM MgCl₂, 0.5 mM CaCl₂) and incubated for 1 hour at 37 °C. EDTA was added to a final concentration 20 mM to stop the reaction. RNA was extracted following the PureLink™ RNA Minikit (Invitrogen) protocol for RNA clean-up and purification from liquid samples with omission of the on-column DNase treatment. RNA was eluted in 30 μL nuclease-free water.

Qβ RNA was sequenced directly using RNA sequencing, 12M reads with rRNA depletion and omitted DNase treatment (SeqCenter). Reads were mapped to NCBI Genome accession AB971354.1 using Geneious software's *Map to Reference* feature, default settings.

MS2 cDNA was synthesized using the Invitrogen SuperScript III First-Strand Synthesis System. Briefly, 4 μL of phage RNA was combined with dNTP mix (final concentration 1 mM), random hexamers (final concentration 5 ng/μL), a primer that anneals to the 3' end of the genome (final concentration 0.2 μM, oAC0025: gccaaaacagccaagcttgggtgtaactagccaagcag), and 3 μL of nuclease-free water. The RNA/primer mix was incubated at 65 °C for 5 minutes, and from this step the rest of the protocol was followed as described in the manufacturer instructions. The MS2 genome was amplified in 3 overlapping fragments from the First-Strand cDNA using OneTaq PCR using previously reported primers.⁷⁵ Amplified MS2 genome was prepared for Illumina sequencing using a modification of the Nextera kit protocol as previously described.⁷⁶ Illumina sequencing was performed using a MiSeq V2 Micro 300-cycle kit (CU Anschutz Genomics and Microarray Core). Reads were mapped to NCBI Genome accession NC_001417 using Geneious software's *Map to Reference* feature, default settings.

METHOD DETAILS

Plasmid construction

The plasmids used in this study are listed in Table S7. DNA manipulations and cloning were performed as previously described.⁸ Briefly, genes of interest were amplified from phage or bacterial genomic DNA using Q5 Hot Start High Fidelity Master Mix (NEB, M0494L), or synthesized as GeneFragments (Genewiz) flanked by ≥ 18 base pairs of homology to the vector backbone. Ligation of genes into restriction-digested, linearized vectors was accomplished using modified Gibson Assembly.⁷⁷ Gibson reactions were transformed via heat shock or electroporation into competent OmniPir and plated onto appropriate antibiotic selection. Where possible, bNACHT coding sequences and endogenous regulatory regions were amplified from the genomic DNA of *E. coli* strains from the ECOR collection.⁷⁸ All other bNACHT gene inserts were ordered as GeneFragments (Genewiz). bNACHT point mutations were generated by amplifying out the gene of interest in two parts from a plasmid template, with the desired mutation occurring in the overlapping region between the two amplicons. Inserts for expression of all *orf008* and *orf015* alleles were amplified from appropriate phage genomic DNA. Unless otherwise indicated, all enzymes were purchased from New England Biolabs.

For all vectors using the pLOCO2 backbone, pAW1382 was amplified and purified from OmniPir. Purified plasmid was then linearized using SbfI-HF and NotI-HF or FseI-HF. Gibson ligation was used to circularize the plasmid with a new insert.

For all vectors using the pTACxc backbone, pAW1608 was amplified and purified from OmniPir. Purified plasmid was then linearized using BamHI-HF and NotI-HF. Gibson ligation was used to circularize the plasmid with the new insert.

For all vectors using the pBAD30x backbone, pAW1367 was amplified and purified from OmniPir. Purified plasmid was then linearized using EcoRI-HF and HindIII-HF. Gibson ligation was used to circularize the plasmid with the new insert.

Sanger sequencing (Genewiz) was used to validate the correct sequence within the multiple cloning site. Additionally, all plasmids expressing bNACHT genes were sequence verified by Illumina sequencing (CU Boulder Sequencing Facility). A NextSeq V2 Mid Output 150-cycle kit was used to sequence the plasmids. Reads were mapped to the predicted plasmid sequence using the *Map to Reference* feature of Geneious Prime (default settings).

Construction of *E. coli* expressing bNACHT genes on the chromosome

MG1655 strains expressing bNACHT alleles at the chromosomal *lacZ* locus were constructed by Lambda red methodology, as previously described.⁷⁹ Sequences were inserted by replacing the *lacZ* coding sequence, eg. replacing ATG...TAA. Synthesis by overlap extension (SOE) PCR was used to generate dsDNA products that contained in order: homology to 50 bp immediately upstream of the MG1655 *lacZ* gene, a kanamycin resistance cassette amplified from pKD4,⁷⁹ GFPmut3 or the indicated bNACHT allele with its endogenous regulatory regions, and homology to 50bp immediately downstream of the MG1655 *lacZ* gene. Purified PCR products were transformed into electrocompetent MG1655 expressing pKD46 and Lambda red was induced with 0.2% arabinose for 2 hours at 30 °C. Cultures were then plated on LB plus kanamycin (25 µg/mL) and grown overnight at 37 °C. Resulting colonies were patched onto LB plus kanamycin (50 µg/mL), and LB plus IPTG (500 µM) and X-Gal (40 µg/mL) to screen for integration of the kanamycin resistance cassette and deletion of the *lacZ* gene, respectively. PCR was used to confirm insertion of the bNACHT genes at the *lacZ* locus.

Efficiency of plating/phage resistance analysis

A modified double agar overlay was used to measure the efficiency of plating (EOP) of phages.^{71,80} Briefly, overnight cultures of *E. coli* MG1655 expressing the indicated plasmids cultured in MMCG plus appropriate antibiotics were diluted 1:10 into the same media and cultivated for an additional two hours to reach mid-log phase (OD₆₀₀ 0.1–0.8). 400 µL of the mid-log culture was mixed with 3.5 mL MMCG (0.35% agar), plus an additional 5 mM MgCl₂ and 100 µM MnCl₂. The mixture was poured onto an MMCG (1.6% agar) plate and cooled for ~15 minutes. 2 µL of a phage dilution series in SM buffer was spotted onto the overlay and allowed to adsorb for 10 minutes before incubating the plate overnight at 37 °C.

Plaque formation was enumerated the following day. For instances with a hazy zone of clearance rather than individual plaque formation, the lowest phage concentration at which clearance was observed was counted as ten plaques. In instances where no clearance or plaque formation was visible, 0.9 plaques at the least dilute spot were used as the limit of detection.

Fold protection was calculated using the inverse of EOP. The PFU of a given phage lysate was measured on sensitive host bacteria, expressing an empty vector, then divided by the PFU for the same phage lysate measure on test bacterial strains. In this way, a 10-fold decrease in EOP is a 10-fold increase in phage protection.

bNACHT22 is included in Figure S4 but not selected for inclusion in Figure 3. Although we did observe a decrease in T3 PFU for this system, we did not observe an expected decrease in T3 plaque size, which undermined our confidence in this result.

Time course of phage infection

Overnight cultures of the appropriate strains were inoculated in 30 mL MMCG plus 5 mM MgCl₂, and 100 µM MnCl₂ to an OD₆₀₀ of 0.1. Cultures were then cultivated shaking at 37 °C for two hours and infected with phage at the indicated MOI. Culture OD₆₀₀ was measured at indicated times.

To enumerate PFU, 250 µL of culture was harvested at each time point and centrifuged at 20,000 × g for 5 minutes at 4 °C. The supernatant was transferred to a new Eppendorf tube and 20–50 µL of chloroform was added to kill any remaining bacteria. Phage lysates were titered using the Efficiency of Plating assay described above.

Validation of bNACHT01 expression

For bNACHT01 mutant expression analysis, 5 mL of the indicated strains were grown to mid-logarithmic phase in MMCG and 5 × 10⁸ CFU were pelleted. For analysis of bNACHT01 expression in response to phage infection, overnight cultures of the appropriate strains were inoculated in 10 mL MMCG plus 5 mM MgCl₂ and 100 µM MnCl₂ to an OD₆₀₀ of 0.1. Cultures were then cultivated shaking at 37 °C for two hours and infected with phage at an MOI of 2. One milliliter of sample was collected at indicated time points and pelleted. Bacterial pellets were washed with water and resuspended in 50 µL of 1 × LDS buffer (106 mM Tris-HCl pH7.4, 141 mM Tris Base, 2% w/v Lithium dodecyl sulfate, 10% v/v Glycerol, 0.51 mM EDTA, 0.05% Orange G). Samples were then incubated at 95 °C for 10 minutes followed by a 5-minute centrifugation at 20,000 × g to remove debris. Samples in LDS were loaded at equal volumes and resolved using SDS-PAGE, then transferred to PVDF membranes charged in methanol. Membranes were blocked in Licor Intercept Buffer for one hour at 24 °C, followed by incubation with primary antibodies diluted in Intercept buffer overnight at 4 °C with rocking. αFLAG antibody (Sigma) was used at 1:10,000 to detect bNACHT01-3×FLAG and α*E. coli* RNA polymerase B antibody (Biolegend) was used at 1:5,000 as a loading control. Blots were then incubated with Licor infrared (800CW/680RD) αRabbit/Mouse secondary antibodies at 1:30,000 dilution in TBS-T (0.1% Triton-X) for one hour at 24 °C and visualized using the Licor Odyssey CLx. Representative images were assembled using Adobe Illustrator CC 2021.

Identification of bacterial NACHTs

We started with an initial sequence library of known NACHT modules from prior studies.^{26,28,81} Upon identification of additional homologs these were then integrated into the initial library for further large-scale sequence analysis as described below. We iterated this

procedure for several rounds, and eventually generated an exhaustive collection of NACHT module homologs. To detect distant relationships, iterative sequence profile searches were conducted using the PSI-BLAST (RRID:SCR_001010)⁵⁸ and JACKHMMER (RRID:SCR_005305)⁵⁹ programs with a profile-inclusion threshold of expect (e)-value at 0.005 against the non-redundant database of the National Center for Biotechnology Information (NCBI) clustered down to 50%. Clustering of proteins based on bit score density and length of aligned sequence was performed using the BLASTCLUST program (<http://ftp.ncbi.nih.gov/blast/documents/blastclust.html>). Remote homology searches were performed using profile-profile comparisons with the HHpred program (RRID:SCR_010276)⁶⁰ against profile libraries comprised of the PFAM (RRID:SCR_004726)⁶¹ and PDB (RRID:SCR_012820)⁶² databases as well as an in-house library of profiles of conserved domains. Multiple sequence alignments were built using the Kalign (RRID:SCR_011810)⁶³ and Muscle (RRID:SCR_011812)⁶⁴ programs followed by manual adjustments based on profile-profile alignment, secondary structure prediction, and structural alignment. Secondary structures were predicted using the JPred (RRID:SCR_016504)⁶⁵ and RoseTTa Fold⁶⁶ programs.

Searches for establishing taxonomic counts of NACHT domains from lineages across the tree of life and viruses was performed using a custom database of 14785 completely sequenced genomes (6847 bacteria) using known NACHT domains as queries for PSI-BLAST searches run for 3 iterations with an inclusion threshold of 0.0001. The detected candidates were then run through a confirmation step with the RPS-BLAST program to obtain the final count of NACHT proteins.

Phylogenetic analysis

The input multiple alignment for this analysis contained 437 proteins and 1112 aligned columns, spanning NACHT domains from across the Tree of Life. Phylogenetic analysis was conducted using the maximum likelihood method implemented in the IQtree program (RRID:SCR_017254)⁶⁷ under multiple parameter regimes using: 1) the Q.pfam substitution matrix derived from alignments in the Pfam database and 1 invariant site category with 8 gamma distributed sites; 2) the LG substitution matrix with 1 invariant site category with 8 gamma distributed sites; 3) with a 20-profile mixture model. Bootstrap values were calculated using the Shimodaira-Hasegawa-like approximate likelihood ratio (SH-aLRT) and the bootstrap proportion-RELL approximation tests^{82,83}. The trees were rendered using the FigTree program (RRID:SCR_008515) (<http://tree.bio.ed.ac.uk/software/figtree/>). Clades with tree topologies evidencing HGT events (Figures 2 and S3) were further tested with a range of tree topology tests. Briefly, this involved construction of the complete set of possible tree topologies within a clade. A subset of these were randomly selected for testing by a range of tree topology tests including approximately unbiased (AU), Kishino-Hasegawa, Shimodaira-Hasegawa, and expected likelihood weights with the IQtree program^{67,84–86} (see Table S3 for a complete list of tests). Trees passing all tests were visually inspected for adherence to the proposed HGT events, and all screened trees retained the proposed HGT topology (Table S3).

Tests for association with multicellularity

Tests for significance of the bNACHT proteins with bacterial multicellularity used the hypergeometric distribution implemented in the phyper command of the R language as previously described, using the available curated database of multicellularity.²⁵ Bacterial multicellularity is defined as reported in the literature (for review, see Lyons and Kolter, 2015).⁸⁷ These include presence of obligate colonial aggregates; namely, the rosettes of planctomycetes, cooperating bacteroid aggregates with branching structures, aggregating cells forming fruiting bodies like the Myxobacteria in the deltaproteobacteria, filaments with differentiated cells (cyanobacteria) and hyphal filamentous aggregates (actinobacteria).

Domain detection

To establish the domain architectures of the NACHT proteins, they were first searched for previously known domains using the RPS-BLAST program with the Pfam database and a custom database including all of domains detected by the Aravind group and augmentations of the Pfam profiles to improve detection. Unknown regions were then investigated. Profile-profile searches were performed with the HHpred program against libraries of profiles based on non-redundant PDB structures, the Pfam database, and a custom collection of profiles of domains not detected by Pfam. Kalign with default parameters and Mafft with maxiterate= 3000, globalpair, op= 1.9 and ep= 0.5 were used to generate input multiple sequence alignments (MSA), followed by refinements using HHpred profile-profile matches or HMM-align. For specific cases structural modeling was performed using the RoseTTAFold program, which uses a “three-track” neural network, utilizing patterns of sequence conservation, distance inferred from coevolutionary changes in MSAs, and coordinate information.⁶⁶ MSAs of related sequences (>30% similarity) were used to initiate HHpred searches for the initial step of correlated position and contact identification to be used by the neural networks.

Analysis of differential diversity of the NACHT module and the SNaCT domain

The analysis of the Shannon entropy (H) for a given multiple sequence alignment was performed using the equation:

$$H = - \sum_{i=1}^M P_i (\log_2(P_i))$$

P is the fraction of residues of amino acid type i and M is the number of amino acid types. The Shannon entropy for the ith position in the alignment ranges from 0 (only one residue at that position) to 4.32 (all 20 residues equally represented at that position).

Trident entropy was used as the metric to analyze the differential divergence of the NACHT module and the SNaCT domain in clade 14. This measure simultaneously unites three distinct elements (hence trident) of positional variability¹⁹ namely: 1) residue diversity; 2) Biochemical diversity among residues; 3) Gapiness of an alignment column. The first $t(x)$ is measured using normalized Shannon entropy (see above); the second $r(x)$ is measured using dissimilarity between two amino acids based on Karlin's formula using a substitution matrix computed from the alignment; the third $g(x)$ measures the number of gaps in the column. The three united as a product ($S=t(x)^a \cdot r(x)^b \cdot g(x)^c$), with each factor scaled with an exponent. The respective exponents used here are: $a=1$, $b=1/2$ and $c=3$. The analysis of the entropy values which were thus derived were performed in the R language.

bNACHT gene selection

bNACHT proteins were selected for screening by considering relatedness of the source genome to *E. coli* and protein domain diversity. For each gene tested, we included the coding sequence of the bNACHT gene, as well as any other genes in the operon. We also included the endogenous regulatory elements of each system, using bPROM⁸⁸ to predict bacterial promoters and Arnold to predict terminators.⁸⁹ We included at least 100 nucleotides to the 3' and 5' region of the gene of interest, to ensure that even unidentified regulatory elements would be included.

DNA degradation measurements

Overnight cultures of the appropriate strains were diluted in 20 mL MMCG plus carbenicillin (100 μ g/mL), 5 mM $MgCl_2$, and 100 μ M $MnCl_2$ to an OD_{600} of 0.1. Cultures were then grown shaking at 37 °C to an OD_{600} of 0.6–0.8 and infected with MS2 at an MOI of 2. 5×10^9 CFU of each sample were harvested at the indicated time points, pelleted by centrifugation at 4,000 \times g for 10 minutes at 4 °C, and plasmid DNA extracted using a standard plasmid miniprep protocol (Qiagen). 10 μ L DNA sample was combined with 2 μ L of 6 \times DNA loading dye (final concentration 3.3 mM Tris-HCl pH 8, 2.5% Ficoll-400, 10 mM EDTA, 0.05% Orange G) and run for 30 min at 130 V on a 1% agarose gel (1% agarose, 40 mM Tris, 20 mM acetic acid, 1 mM EDTA, SYBR Safe DNA stain). Gels were imaged using an Azure Biosystems Azure 200 Bioanalytical Imaging System.

To measure impact of *orf008* and *orf015* on DNA degradation, overnight cultures of the appropriate strains were inoculated in 20 mL MMCG plus carbenicillin (50 μ g/mL) and chloramphenicol (10 μ g/mL) to an OD_{600} of 0.1. Cultures were then cultivated shaking at 37 °C to an OD_{600} of 0.6–0.8 before the addition of 0.2% arabinose and 500 μ M IPTG to induce expression of bNACHT25 and phage *orfs*, respectively. Cultures were then harvested after an additional two hours of growth and analyzed as described above.

NAD(H) degradation measurements

Concentrations of NAD(H) were measured using the Promega NAD/NADH-Glo Assay following the manufacturer instructions. Briefly, overnight cultures of the appropriate strain were diluted in 25 mL MMCG to an OD_{600} of 0.1 and grown for 2 hours at 37 °C. Cultures were then split into 1.5 mL aliquots and infected with phage T4 at an MOI of 2 in 14 mL culture tubes. At the indicated time points, 500 μ L culture was harvested by centrifugation at 21,000 \times g for 5 minutes at 4 °C. Bacterial pellets were resuspended in 50 μ L PBS (Corning) and incubated with 50 μ L 0.2 M NaOH with 0.1% dodecyltrimethylammonium bromide at 24 °C for 8 minutes to lyse. 100 μ L HCl/Tris solution was added to neutralize the sample and incubated at 24 °C for 5 minutes. 50 μ L sample was transferred to a white opaque 96-well plate (Pierce) and mixed with 50 μ L NAD/NADH-Glo™ Detection Reagent. Luminescence was measured after 30 min incubation at 25 °C using a Tecan Spark multimode microplate reader.

The amount of NAD(H) present in each sample was calculated based on the NAD(H) standard curve for each experiment and normalized to the amount of NAD(H) present in an equivalent volume of sample with an OD_{600} of 0.1 to allow for accurate comparisons between samples. Each biological sample was analyzed in technical triplicate.

Growth inhibition measurements

The impact of bNACHT expression with and without *orf008* and *orf015* alleles on bacterial growth was quantified using a colony formation assay. *E. coli* was cultivated overnight in MMCG with appropriate antibiotics. Cultures were diluted in a 10-fold series into MMCG and 5 μ L of each dilution was spotted onto a MMCG agar plate containing the appropriate antibiotics, as well as IPTG and/or arabinose as appropriate. Spotted bacteria were allowed to dry for ~10 minutes before the plates were incubated overnight at 37 °C.

Growth inhibition was measured the following day by enumerating the colony forming units of each strain, reported as CFU/mL for the starting culture. For instances where bacteria were growing but no individual colonies could be counted, the lowest bacterial concentration at which growth was observed was counted as ten CFU. In instances where no growth was visible, 0.9 CFU at the least dilute spot was used as the limit of detection.

Phage suppressor generation and amplification

T5 phages able to evade bNACHT01-mediated protection were generated by mixing 400 μ L of mid-log bacteria expressing bNACHT01 in MMCG plus 100 μ g/mL carbenicillin with wild-type T5 at an MOI ~10 and pouring the mixture onto a MMCG agar plate. Individual plaques were isolated and spot-plated onto *E. coli* MG1655 expressing bNACHT01 to confirm that phages were able to

replicate in the presence of bNACHT01 and to plaque-purify each clone. Phage bNACHT01 suppressors were generated using three separate wild-type T5 stocks amplified from individual plaque purifications. Phage T5 suppressors were subsequently plate amplified on *E. coli* MG1655 expressing bNACHT01 in MMCG.

Genome sequencing and analysis of phage suppressors

Suppressor phage genomes were extracted as described above. Extracted phage genomes were prepared for Illumina sequencing using a modification of the Nextera kit protocol as previously described.⁷⁶ Illumina sequencing was performed using a MiSeq V2 Micro 300-cycle kit (CU Boulder Sequencing Facility). Reads were mapped to Genome accession AY587007 (empirically determined to be most similar to the T5 phage used in this study) using Geneious software's *Map to Reference* feature. Reads were trimmed to remove the Nextera adapter sequences before mapping (sequence trimmed: AGATGTGTATAAGAGACAG) using the "Trim primers" option, with otherwise default settings. Sequences were mapped using default settings, selecting "map multiple best matches to all locations" to accommodate repetitive T5 sequences.

Geneious was also used for variant detection from the reference T5 genome. Variants that were present in ≥ 75 percent of reads from the suppressor phage genome but not the parent phage genome were identified as potential suppressor mutations.

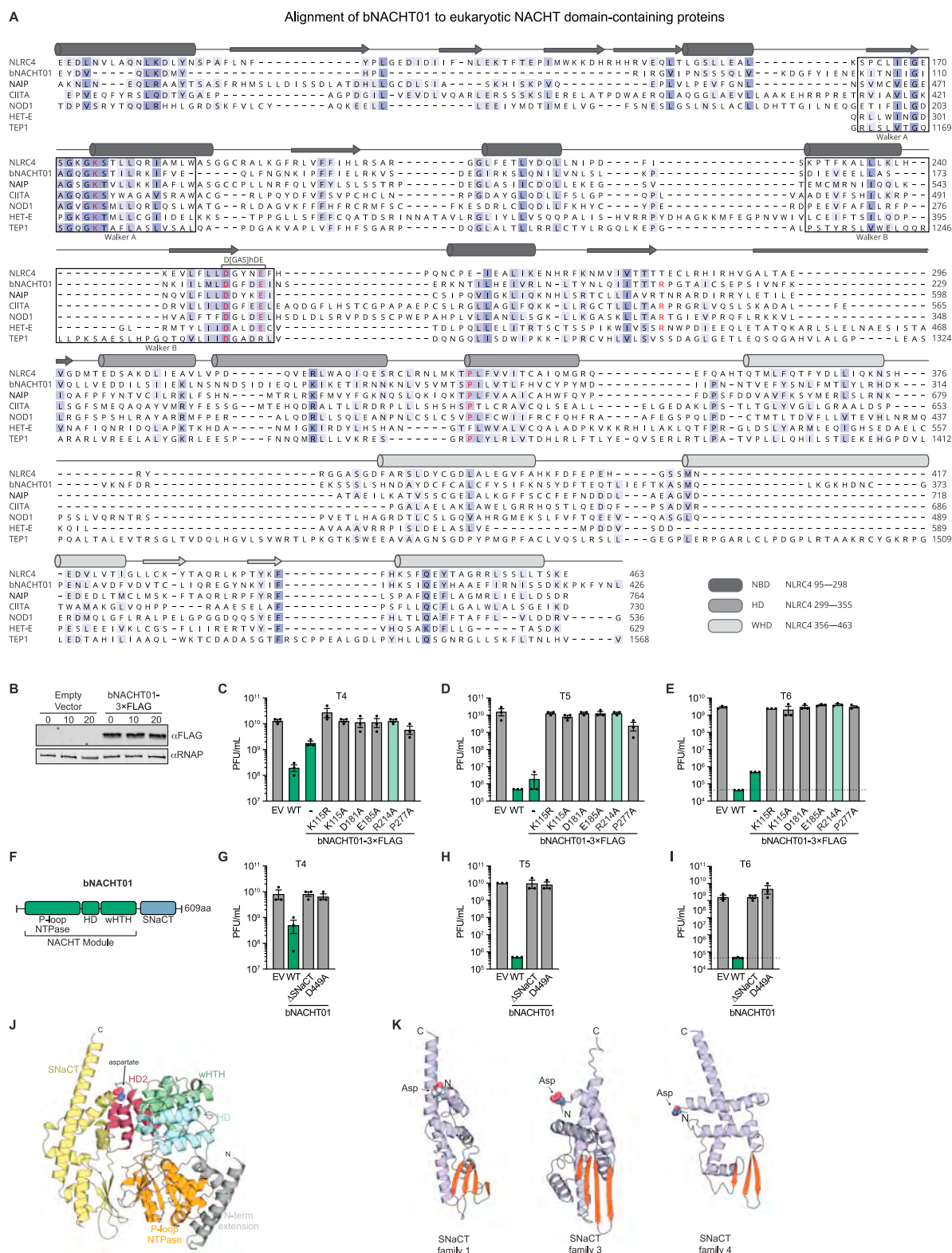
Effect of phage genes on bNACHT protection against phage

Bacterial strains were cultured overnight in MMCG plus 20 $\mu\text{g/mL}$ carbenicillin and 4 $\mu\text{g/mL}$ chloramphenicol. Cultures were then diluted 1:10 into the same media with or without 100 μM IPTG and grown for 4 more hours to reach mid-log phase. Phage resistance was measured as described above, with the addition of IPTG to the MMCG top agar (0.35%) to continue inducing conditions.

QUANTIFICATION AND STATISTICAL ANALYSES

All experiments were performed in biological triplicate using cultures grown on three separate days. Data was plotted using Graphpad Prism 9 at an n of 3 with error bars indicating standard error of the mean. Illumina sequencing results were analyzed using Geneious Prime Software. Geneious Prime was also used to generate alignments, using MAFFT alignment⁶⁸ and default settings. Figures were created using Adobe Illustrator CC. For statistical analysis of trident entropy, see [STAR Methods](#) section titled "[analysis of differential diversity of the NACHT module and the SNaCT domain](#)".

Supplemental figures



(legend on next page)

Figure S1. Protein alignment of the NACT modules from bNACT01 and other NACT proteins, related to Figure 1

(A) Protein alignment of the NACT modules of NLRC4 (*Mus musculus*; GenBank: NP_001028539), bNACT01 (*Klebsiella pneumoniae* MGH 35; GenBank: WP_015632533.1), NAIP (*Homo sapiens*; GenBank: NP_001333799.1), CIITA (*Homo sapiens*; GenBank: NP_000237.2), NOD1 (*Homo sapiens*; GenBank: NP_006083.1), HET-E (*Podospira anserina*; Uniprot: Q00808), and TEP1 (*Homo sapiens*; GenBank: NP_009041.2). The secondary structure of NLRC4 as determined using structure PDB: 4KXF,¹⁵ is indicated above with alpha helices depicted as cylinders and beta sheets depicted as arrows. Secondary structure elements are color coded to represent the NBD, HD, and WHD of NLRC4. Amino acid residues are color coded based on conservation such that darker colors indicate a higher degree of conservation in this alignment. Black boxes indicate the Walker A and Walker B motifs. Residues mutated and analyzed for expression in Figure 1D are highlighted in red. The D[GAS]hDE motif within the Walker B region that distinguishes NACT modules from other STAND NTPases is indicated.

(B) Western blot analysis of *E. coli* expressing empty vector or FLAG-tagged bNACT01 at 0, 10, and 20 min post-infection with phage T5. Representative image of n = 2 biological replicates.

(C–E) Efficiency of plating of the indicated phage infecting *E. coli* expressing the indicated genotype. Data represent the mean \pm SEM of n = 3 biological replicates, shown as individual points.

(F) Schematic of bNACT01 (GenBank: WP_015632533.1) protein domains, annotated by alignment to the NACT module of NLRC4. The P loop NTPase domain is also known as a nucleotide-binding domain (NBD), the helical domain (HD), and the winged helix-turn-helix (wHTH, also called WHD for winged helical domain) are indicated.

(G–I) Efficiency of plating of the indicated phage infecting *E. coli* expressing the indicated genotype. Data represent the mean \pm SEM of n = 3 biological replicates, shown as individual points.

(J) Predicted structure of bNACT01. Aspartate represents the conserved residue D449.

(K) Predicted structures of 3 different SNaCT domain families. N represents the N terminus of the domain.

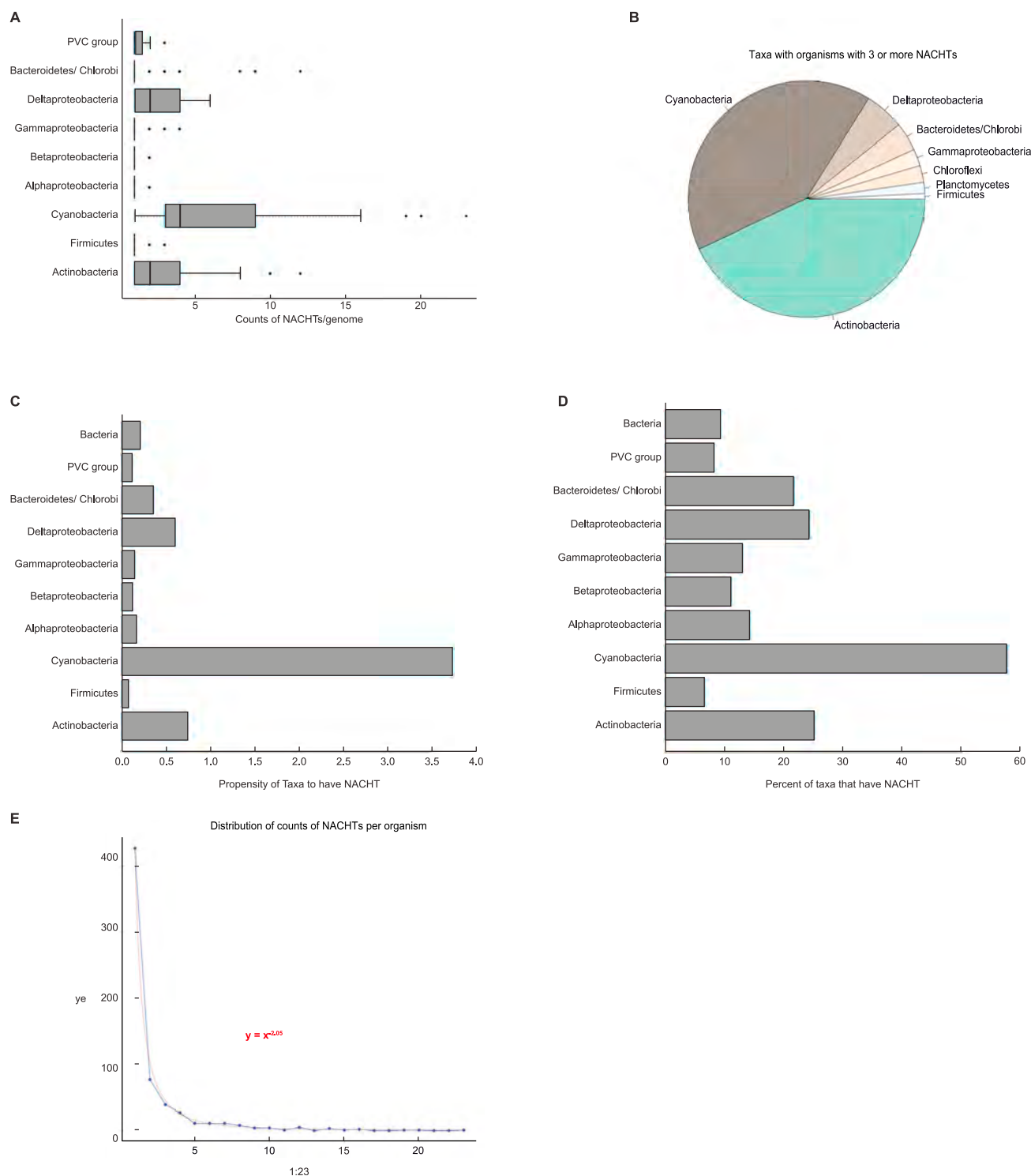


Figure S2. Distribution of NACHT proteins in bacterial taxa, related to Figure 2

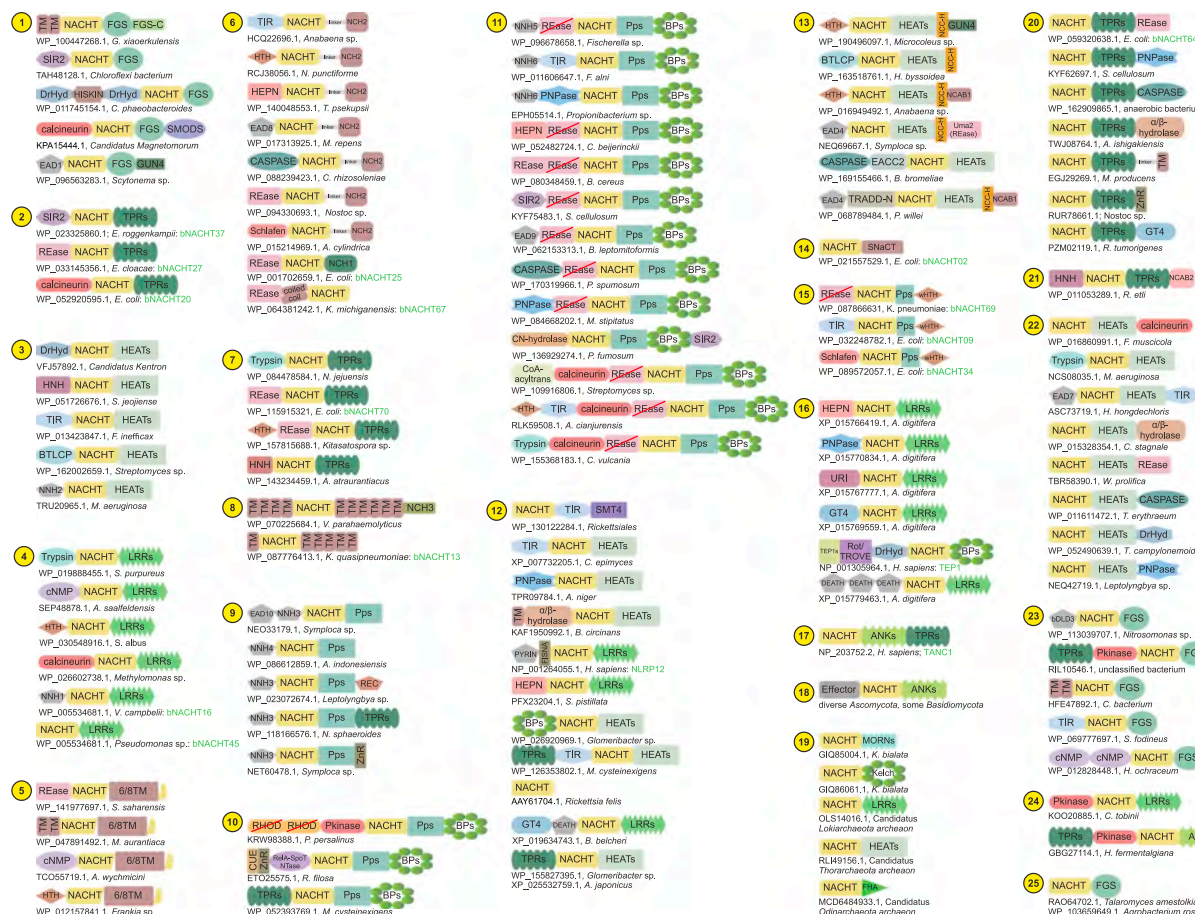
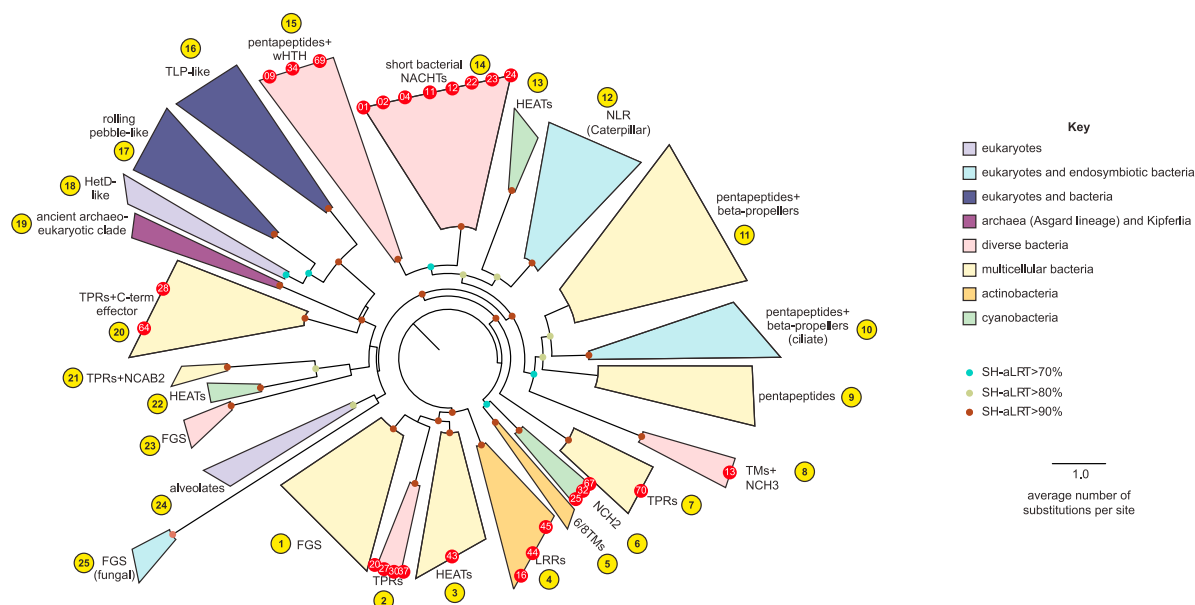
(A) Quantification of the number of individual NACHTs found within a single genome across different bacterial taxa. The maximum for the x axis is 23. Planctomycetota, Verrucomicrobiota, Chlamydiota (PVC).

(B) Relative distribution of taxa that have organisms with 3 or more NACHTs in a single genome.

(C) Propensity of organisms within the indicated taxa to encode a NACHT module-containing protein.

(legend continued on next page)

(D) Percent of organisms in the indicated taxa to encode at least one NACHT module-containing protein.
 (E) Distribution of the number of NACHT proteins per organism, which can be fitted to the equation $y = x^{-2.05}$. For (A)–(E), a custom database of complete bacterial genomes was used to analyze the distribution of NACHT proteins in diverse bacterial taxa.



(legend on next page)

Figure S3. NACHT module-containing proteins in bacteria are widespread and diverse, related to Figure 2

A sequence-based phylogenetic tree of NACHT modules was generated using NACHT module-containing proteins from eukaryotes and prokaryotes. The NACHT module, not accessory domains, was used for tree building. Clades are color coded based on the indicated key and numbered arbitrarily in yellow circles. Red dots indicate the bacterial NACHT proteins from each clade that were selected for analysis in this study. Bootstrap values are provided where applicable. The scale bar indicates the average number of substitutions per site. See Table S4 for the most common domain architectures found in each clade. The genes used to construct this phylogenetic tree can be found in Tables S1 and S2 contains a full list of all NACHT domain-containing proteins identified. See Table S3 for tree topology tests used to validate proposed evolutionary relationships. Representative domain architectures for each clade are provided, including NCBI Protein accession number, species of origin, and gene name in green where appropriate. 3',5' cyclic nucleotide-generating cyclase (cNMP), Ankyrin repeats (ANKs), bacterial death-like domain-3 (bDLD3), beta-propeller repeats (BPs), bacterial transglutaminase-like cysteine protease (BTCLP), carbon-nitrogen hydrolase (CN-hydrolase), CoA-dependent acyltransferase (CoA-acyltrans), deoxyribohydrolase (DrHyd), effector-associated constant component (EACC), effector-associated domain (EAD), formylglycine-generating enzyme sulfatase (FGS), FGS C-terminal domain (FGS-C), forkhead-associated domain (FHA), fish-specific NACHT-associated domain (FISNA), glycosyltransferase 4 (GT4), Huntington, elongation factor 3, PR65/A, TOR (HEAT), higher eukaryotes and prokaryotes nucleotide-binding domain (HEPN), homing endonuclease (HNN), helix-turn-helix (HTH), leucine-rich repeat (LRR), membrane occupation and recognition nexus (MORN), NACHT N-terminal helical domain (NNH), NACHT C-terminal helical domain (NCH), NACHT C-terminal α/β domain (NCAB), NACHT C-terminal cysteine and histidine-containing domain (NCC-H), protein kinase domain (Pkinase), polyribonucleotide nucleotidyl transferase (PNPase), Pentapeptide repeat (Pp), restriction endonuclease (REase), receiver domain (Rec), rhodanese domain (RHOD), Sirtuin (SIR2), second messenger oligonucleotide or dinucleotide synthetase domain (SMODS), telomerase associated protein 1 (TEP1), toll/interleukin receptor (TIR), transmembrane (TM) number indicates how many copies of this motif, tetratricopeptide repeat (TPR), N-terminal domain of tumor necrosis factor receptor type 1 associated death domain protein (TRADD-N), UvRC and intron-encoded endonuclease domain (URI), zinc ribbon (ZnR). The red slash indicates a catalytically inactive domain.

bNACHT Number	Domain Architecture	NACHT Clade	Source genome	N-term (effector)	C-term (sensor)	Length (aa)
bNACHT01		14	<i>Klebsiella pneumoniae</i> MGH 35	-	SNaCT	609
bNACHT02		14	<i>Escherichia coli</i> ECOR34	-	SNaCT	634
bNACHT04		14	<i>Klebsiella pneumoniae</i> UCICRE 8	-	SNaCT	652
bNACHT09		15	<i>Escherichia coli</i> 3-105-05S4 C1	TIR	Pentapeptide Repeat + Winged Helix-Turn-Helix	957
bNACHT11		14	<i>Klebsiella pneumoniae</i> 4300STDY6470423	-	SNaCT	639
bNACHT12		14	<i>Escherichia coli</i> HVH 41	-	SNaCT	612
bNACHT13		8	<i>Klebsiella quasipneumoniae</i> TUM14048	TM	TM	1021
bNACHT16		4	<i>Vibrio campbellii</i> CAIM 519	α-h1	LRR	950
bNACHT20		2	<i>Escherichia coli</i> 2014C-3338	Metallophosphoesterase	Tetratricopeptide Repeat	1050
bNACHT22		14	<i>Escherichia coli</i> ECOR36	-	SNaCT	652
bNACHT23		14	<i>Escherichia coli</i> ECOR08	-	SNaCT	645
bNACHT24		14	<i>Escherichia coli</i> ECOR35	-	SNaCT	652
bNACHT25		6	<i>Escherichia coli</i> ECOR11	REase	HEAT	752
bNACHT27		2	<i>Enterobacter asburiae</i> RHBSTW-00831	REase	Tetratricopeptide Repeat	1473
bNACHT28		20	<i>Escherichia coli</i> TzEc064	-	Tetratricopeptide Repeat	615
bNACHT30		2	<i>Klebsiella pneumoniae</i> 4300STDY6470402	-	-	572
bNACHT32		6	<i>Escherichia coli</i> RU1 MA4	REase	HEAT	753
bNACHT34		15	<i>Escherichia coli</i> NCTC9088	Schlafen	Pentapeptide Repeat + Winged Helix-Turn-Helix	979
bNACHT37		2	<i>Enterobacter roggenkampii</i> MGH 34	Sirtuin	Tetratricopeptide Repeat	1118
bNACHT42		7	<i>Klebsiella varicola</i> 5012STDY7312728	Beta propeller	-	1042
bNACHT43		3	<i>Klebsiella varicola</i> WUSM_KV_03	Peptidase	-	749
bNACHT44		4	<i>Pseudomonas viridiflava</i> CDRTc14	-	LRR	630
bNACHT45		4	<i>Pseudomonas</i> sp. LAIL14HWK12:16	-	LRR	954
bNACHT64		20	<i>Escherichia coli</i> JML132	-	Tetratricopeptide Repeat + REase	1468
bNACHT67		6	<i>Klebsiella michiganensis</i> 2880STDY5682685	REase	-	921
bNACHT69		15	<i>Klebsiella pneumoniae</i> CRK0035	REase	Pentapeptide Repeat + Winged Helix-Turn-Helix	743
bNACHT70		7	<i>Escherichia coli</i> LMR3158	REase	Tetratricopeptide Repeat	1330

Figure S4. Domain architectures and details of experimentally tested NACHT proteins, related to Figure 3

Diagram of the N-terminal domains, central NACHT modules, and C-terminal domains of each experimentally tested bNACHT protein. The clade, species of origin, and length of each protein in amino acids (aa) are also indicated. Leucine-rich repeat (LRR), Huntington, elongation factor 3, PR65/A, TOR (HEAT), restriction endonuclease (REase), toll/interleukin receptor (TIR), transmembrane (TM), NACHT N-terminal helical domain 1 (NNH1), NACHT C-terminal helical domain 1 (NCH1).

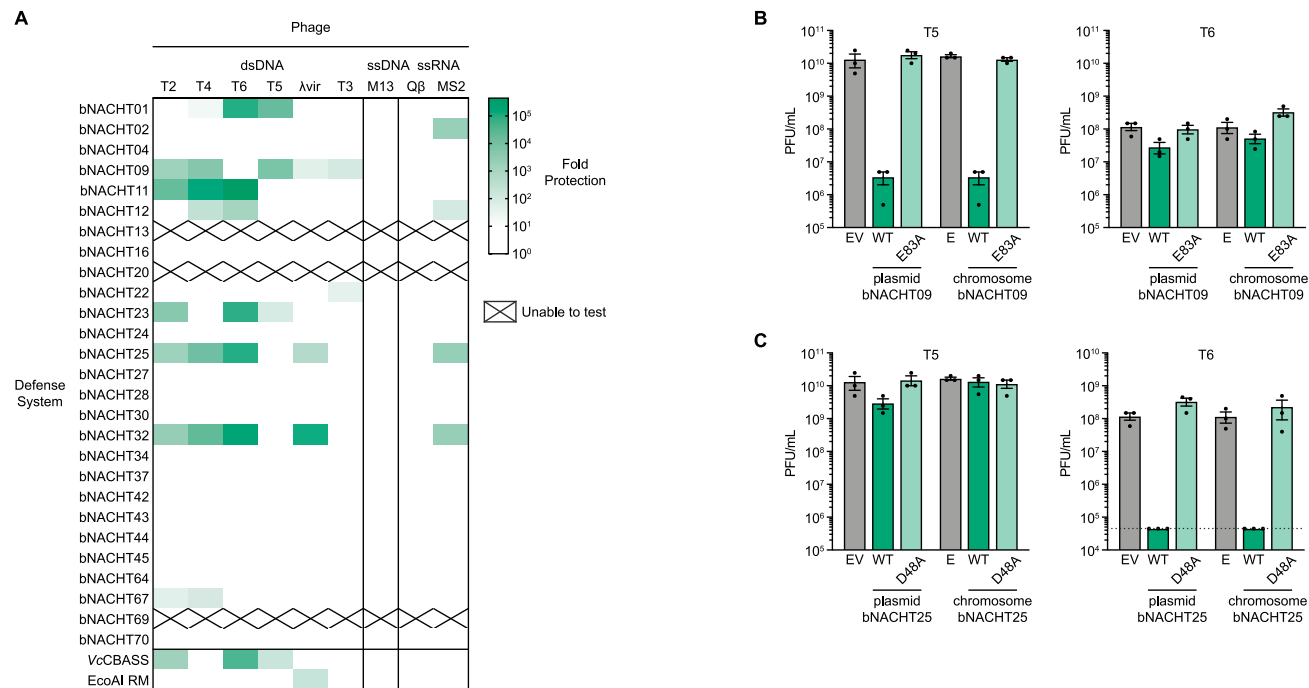
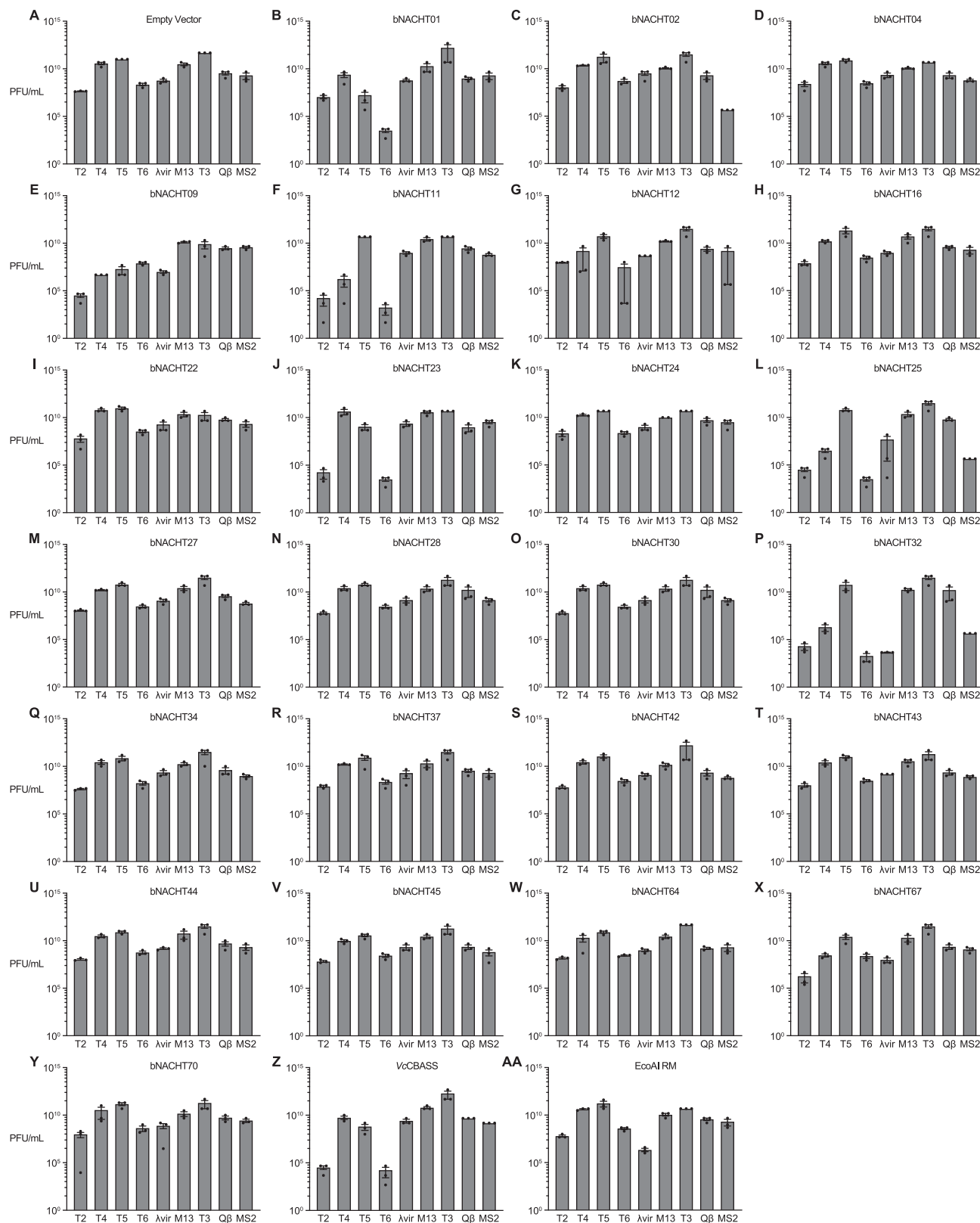


Figure S5. Expanded screen for bacterial NACHT protein phage defense and comparison of defense conferred by plasmid- or chromosomally expressed bacterial NACHTs, related to Figure 3

(A) Heatmap of fold defense provided by the indicated bNACHT gene for a panel of diverse phages. Data are as described in Figure 3 and show all experimentally interrogated bNACHT genes. Data were not collected for bacterial NACHT proteins that exhibited insufficient growth under these experimental conditions, indicated as "X." Data represent the mean of $n = 3$ biological replicates. See Table S5 and Figure S4 for details on all 27 bNACHT genes analyzed. See Figure S6 for the raw efficiency of plating data.

(B and C) Efficiency of plating of the indicated phage infecting *E. coli* expressing the indicated genotype. Empty (E) indicates *E. coli* with the chromosomal expression of a kanamycin resistance cassette and *gfpmut3*. Data represent the mean \pm SEM of $n = 3$ biological replicates, indicated as individual points.



(legend on next page)

Figure S6. Efficiency of plating of phages infecting bacteria expressing NACHT proteins, related to [Figure 3](#)

Efficiency of plating of a phage panel infecting *E. coli* expressing the indicated bNACHT gene. The negative control is an empty vector, which expresses an inactive *gfp* gene. Positive controls are *V. cholerae* CBASS (VcCBASS) and *E. coli* UPEC-36 (EcoAI RM) restriction-modification system (Z and AA). Data represent the mean \pm SEM of $n = 3$ biological replicates, indicated as individual points.

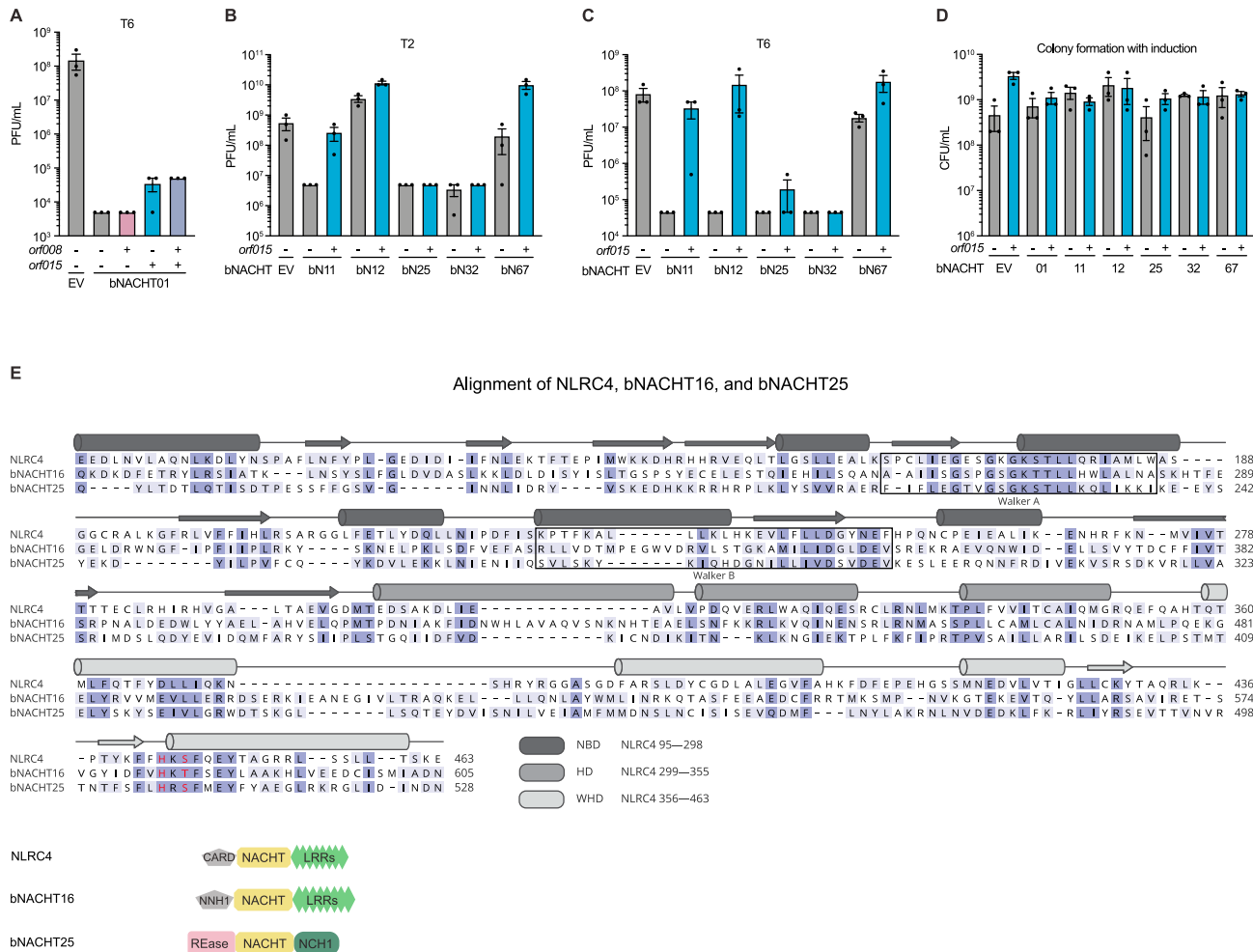


Figure S7. Effect of *orf008* and *orf015* expression on bNACHT01 activity and comparison of bNACHT modules to the NACHT domain of NLRC4, related to Figures 5 and 6

(A) Efficiency of plating of phage T6 when infecting *E. coli* expressing bNACHT01 or empty vector (EV) on one plasmid and the indicated phage T5 gene(s) on a second plasmid.

(B) Efficiency of plating of phage T2 when infecting *E. coli* expressing the indicated bNACHT gene on one plasmid and phage T5 *orf015* on a second plasmid.

(C) Efficiency of plating of phage T6 when infecting *E. coli* expressing the indicated bNACHT gene on one plasmid and phage T5 *orf015* on a second plasmid.

(D) Quantification of colony formation of *E. coli* expressing the indicated bNACHT system on one plasmid and phage T5 *orf015* on a second plasmid. For (A)–(D), the expression of *orf008*, *orf015*, or *sfGFP* is IPTG-inducible. (–) symbols denote the induction of an *sfGFP* negative control. (+) symbols denote induction of *orf015*. Data represent the mean \pm SEM of $n = 3$ biological replicates, shown as individual points.

(E) Protein alignment of the NACHT modules of NLRC4 (*Mus musculus*; GenBank: NP_001028539), bNACHT16 (*Vibrio campbellii* CAIM 519; GenBank: WP_005534681.1), and bNACHT25 (*E. coli* ECOR11; GenBank: WP_001702659.1). The secondary structure of NLRC4 as determined by structure PDB: 4KXF,¹⁵ is indicated above with alpha helices depicted as cylinders and beta sheets depicted as arrows. Secondary structure elements are color coded and labeled as in Figure S1. Amino acid residues are color coded based on conservation in the sequence alignment. Black boxes indicate Walker A and B motifs. The conserved histidine and serine/threonine in the WHD are highlighted in red.