



# Functional Characterization of COG1713 (YqeK) as a Novel Diadenosine Tetraphosphate Hydrolase Family

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**ABSTRACT** Diadenosine tetraphosphate (Ap<sub>4</sub>A) is a dinucleotide found in both prokaryotes and eukaryotes. In bacteria, its cellular levels increase following exposure to various stress signals and stimuli, and its accumulation is generally correlated with increased sensitivity to a stressor(s), decreased pathogenicity, and enhanced antibiotic susceptibility. Ap<sub>4</sub>A is produced as a by-product of tRNA aminoacylation, and is cleaved to ADP molecules by hydrolases of the ApaH and Nudix families and/or by specific phosphorylases. Here, considering evidence that the recombinant protein YqeK from *Staphylococcus aureus* copurified with ADP, and aided by thermal shift and kinetic analyses, we identified the YqeK family of proteins (COG1713) as an unprecedented class of symmetrically cleaving Ap<sub>4</sub>A hydrolases. We validated the functional assignment by confirming the ability of YqeK to affect *in vivo* levels of Ap<sub>4</sub>A in *B. subtilis*. YqeK shows a catalytic efficiency toward Ap<sub>4</sub>A similar to that of the symmetrically cleaving Ap<sub>4</sub>A hydrolases of the known ApaH family, although it displays a distinct fold that is typical of proteins of the HD domain superfamily harboring a diiron cluster. Analysis of the available 3D structures of three members of the YqeK family provided hints to the mode of substrate binding. Phylogenetic analysis revealed the occurrence of YqeK proteins in a consistent group of Gram-positive bacteria that lack ApaH enzymes. Comparative genomics highlighted that *yqeK* and *apaH* genes share a similar genomic context, where they are frequently found in operons involved in integrated responses to stress signals.

**IMPORTANCE** Elevation of Ap<sub>4</sub>A level in bacteria is associated with increased sensitivity to heat and oxidative stress, reduced antibiotic tolerance, and decreased pathogenicity. ApaH is the major Ap<sub>4</sub>A hydrolase in gamma- and betaproteobacteria and has been recently proposed as a novel target to weaken the bacterial resistance to antibiotics. Here, we identified the orphan YqeK protein family (COG1713) as a highly efficient Ap<sub>4</sub>A hydrolase family, with members distributed in a consistent group of bacterial species that lack the ApaH enzyme. Among them are the pathogens *Staphylococcus aureus*, *Streptococcus pneumoniae*, and *Mycoplasma pneumoniae*. By identifying the player contributing to Ap<sub>4</sub>A homeostasis in these bacteria, we disclose a novel target to develop innovative antibacterial strategies.

**KEYWORDS** Ap<sub>4</sub>A hydrolase, adenosine tetraphosphate, dinucleoside polyphosphates, Gram-positive bacteria, nucleotides

Ap<sub>4</sub>A is a ubiquitous metabolite whose physiological function is still a matter of debate. Unknown is whether it represents simply a “damage metabolite” or is instead an “alarmone” with stress-signaling functions, or even a modulator of the stress response (1, 2). It is formed by a side reaction of aminoacyl-tRNA synthetase that, in the absence of tRNA, catalyzes the transfer of the AMP moiety of the bound aminoacyl

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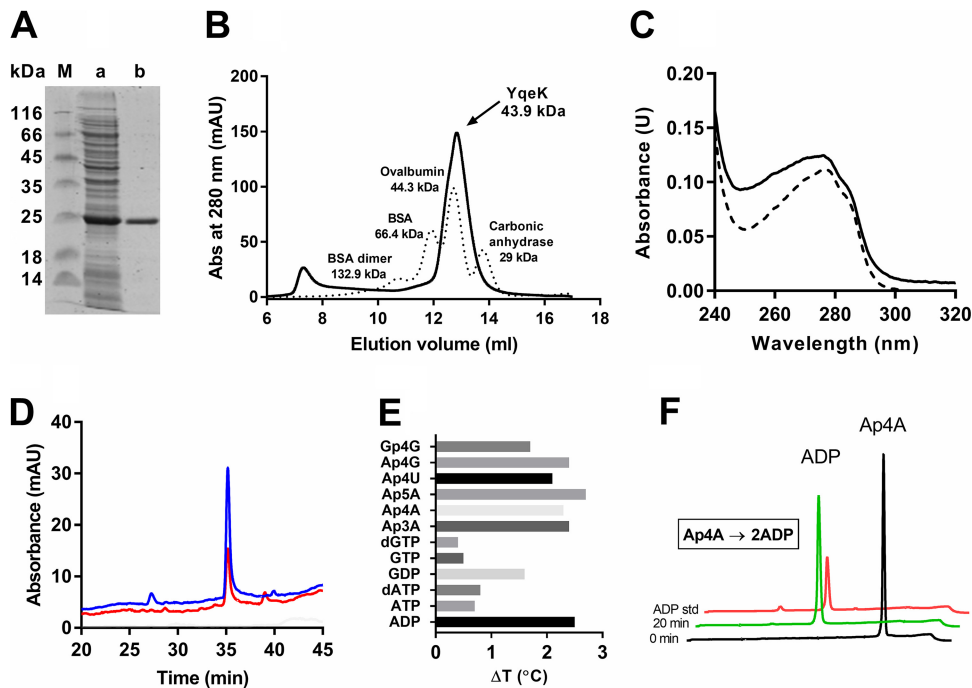
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adenylate to an ATP molecule (3). In mammalian and bacterial cells, Ap<sub>4</sub>A is present at a low micromolar concentration under normal conditions, but these levels rise significantly under various stress conditions, including heat, antibiotics, nutritional, and oxidative stresses (4–6). Accumulation of Ap<sub>4</sub>A is pleiotropic, resulting in loss of motility and defects in catabolite repression, increased sensitivity to heat, oxidative stress, and aminoglycoside antibiotics, along with impairment of biofilm formation and inhibition of sporulation under starvation conditions (4, 5, 7–9). The mechanism of action of Ap<sub>4</sub>A is still under investigation. Recent evidence showing that elevation of Np<sub>4</sub>A levels is associated with a novel Np<sub>4</sub> capping of *E. coli* transcripts, leading to longer lifetimes, suggests that Ap<sub>4</sub>A might be involved in gene expression regulation (2). Ap<sub>4</sub>A homeostasis is maintained by the activity of hydrolases and/or phosphorylases, depending on the organism (10). Ap<sub>4</sub>A hydrolases belong to the ApaH (EC 3.6.1.41) and Nudix (EC 3.6.1.17) families. Hydrolases of the ApaH family catalyze the symmetrical cleavage of Ap<sub>4</sub>A, yielding two ADP molecules (11). They are widely distributed among gamma- and betaproteobacteria. Loss of ApaH function results in a marked increase in Ap<sub>4</sub>A levels, with consequent enhanced sensitivity to stressful conditions and decreased pathogenicity (8, 9, 12). Interestingly, the recent finding that *apaH* deletion in *P. aeruginosa* caused a significant increase in the bacterium sensitivity to kanamycin identified ApaH as a novel target for enhancing the killing potency of aminoglycosides (4). Ap<sub>4</sub>A hydrolases of the Nudix family (EC 3.6.1.17) asymmetrically hydrolyze Ap<sub>4</sub>A to ATP and AMP and are present both in eukaryotes and bacteria (13, 14). Notably, in pathogenic bacteria, both ApaH and Nudix hydrolase activities are essential for the intracellular invasion of the host (12). Finally, an Ap<sub>4</sub>A phosphorylase (EC 2.7.7.53) that converts Ap<sub>4</sub>A into ADP and ATP in the presence of phosphate can be found in lower eukaryotes (15) and some bacteria (16), including *M. tuberculosis* (17), where its absence is shown to cause a constant stress response that impairs bacterial growth and proliferation (18). In this work, based on biochemical studies and bioinformatic analyses, we identified the *Staphylococcus aureus* YqeK protein as the member of a novel bacterial family of symmetrically cleaving Ap<sub>4</sub>A hydrolases, distinct from the ApaH family, and conserved in a consistent group of bacterial species lacking the ApaH enzyme.

## RESULTS

**Biochemical characterization of *S. aureus* YqeK.** YqeK from *S. aureus* (SaYqeK) was expressed in *E. coli* and purified to homogeneity. A molecular weight of about 23 kDa was estimated upon SDS-PAGE analysis (Fig. 1A). Gel filtration experiments showed a native molecular weight of about 44 kDa, which is consistent with a dimeric structure of the protein (Fig. 1B). During the purification of the protein, we noticed that the UV spectrum of the fraction eluted from the Ni-nitrilotriacetic acid (Ni-NTA) column had a shape and a  $A_{260}/A_{280}$  ratio that indicated a mixture of protein and nucleotides (Fig. 1C) (19). Conversely, the spectrum of the protein after gel filtration was typical of a pure protein, with a  $A_{260}/A_{280}$  ratio very close to 0.6 (20) (Fig. 1C). This suggested that an unknown nucleotide(s), likely bound to YqeK during its expression in *E. coli*, could have been released during the gel filtration chromatography. To identify such a ligand(s), we denatured the protein obtained from the affinity chromatography either by acidic treatment or by heating and performed a reversed-phase high-performance liquid chromatography (RP-HPLC) analysis on the deproteinized samples. In both cases, one single peak was observed, whose identity as genuine ADP was confirmed by coelution with an ADP standard (Fig. 1D).

The finding that an ADP molecule copurified with YqeK, together with the observation that molecules of GDP and dGDP have been found in the active sites of the protein's available 3D structures (PDB codes: 2O08 and 2OGI), prompted us to screen as possible ligands a series of metabolites comprising ADP and GDP in their moieties. To this end, we used a thermal shift assay as a preliminary screening test. Several compounds were tested, and those affecting the thermal stability of the protein are shown in Fig. 1E. Among them, ADP, Ap<sub>5</sub>A, Ap<sub>4</sub>A, Ap<sub>3</sub>A, Ap<sub>4</sub>U, and Ap<sub>4</sub>G were the most



**FIG. 1** Biochemical characterization of *SaYqeK*. (A) SDS-PAGE analysis of recombinant *SaYqeK* purification. M, molecular weight standards; a, crude extract; b, Ni-NTA pool. (B) Gel filtration chromatography of Ni-NTA pool (solid line) and standard proteins (dotted line). (C) UV spectral analysis of the Ni-NTA pool (solid line) and gel filtration pool (dashed line). (D) HPLC analysis of the acid-soluble fraction of the Ni-NTA pool (red) and ADP standard (blue). (E) *SaYqeK* ligand screening by thermal shift analysis. Ligands were tested at 5  $\mu$ M concentration. For ease of viewing, bars are colored with different shades of gray. (F) HPLC profile at 260 nm of the reaction mixture of *SaYqeK* (50  $\mu$ g/ml) at 0 and 20 min of incubation in the presence of 0.2 mM Ap<sub>4</sub>A. A standard ADP profile is also shown (std).

effective. Also, GDP and Gp<sub>4</sub>G increased the melting temperature of the protein, although to a lesser extent, whereas ATP, GTP, and the corresponding deoxyribonucleotides were the least effective. No effect was exerted by Ap<sub>2</sub>A, AMP or GMP, pyrimidine nucleotides, NAD or nicotinate adenine dinucleotide (NaAD), ADP- or GDP-linked sugars (like ADP-ribose, ADP-mannose, ADP-glucose, or GDP-glucose), cyclic AMP or GMP, or cyclic di-GMP or cyclic di-AMP. Likewise, no thermal shift was observed in the presence of phosphate, pyrophosphate, or phosphoribosyl pyrophosphate.

Prompted by these results, we investigated whether YqeK was able to use the candidate ligands as substrates. As a first screening, we incubated the enzyme at 0.05 mg/ml with 0.2 mM ligand and analyzed the reaction mixture by RP-HPLC after a 10 min incubation at 37°C. While GDP, ADP, GTP, ATP, and the corresponding deoxyribonucleotides were not consumed, Ap<sub>3</sub>A, Ap<sub>4</sub>A, and Ap<sub>5</sub>A entirely disappeared, being converted to ADP plus AMP, ADP, and ADP plus ATP, respectively. Likewise, efficient hydrolysis of Ap<sub>4</sub>U, Ap<sub>4</sub>G, and Gp<sub>4</sub>G was observed, always releasing ADP or GDP as one of the products. Hydrolysis of Ap<sub>4</sub>A is shown in Fig. 1F. By assaying YqeK activity under initial velocity conditions, we found that the activity with Ap<sub>3</sub>A was only 5% of that measured with Ap<sub>4</sub>A. Therefore, Ap<sub>3</sub>A was not considered further. The results of the kinetic analysis performed with the tested nucleotides are shown in Table 1. The

**TABLE 1** *SaYqeK* kinetic constants for preferred substrates

Substrate	$K_m$ ( $\mu$ M $\pm$ SD)	$k_{cat}$ ( $s^{-1}$ $\pm$ SD)	$K_{cat}/k_m$ ( $s^{-1}$ $\mu$ M <sup>-1</sup> )
Ap <sub>4</sub> A	3.2 $\pm$ 0.3	107 $\pm$ 2	34
Ap <sub>5</sub> A	21.5 $\pm$ 2.5	32 $\pm$ 1	1.5
Gp <sub>4</sub> G	68.0 $\pm$ 22	1,002 $\pm$ 186	15
Ap <sub>4</sub> G	19.7 $\pm$ 4.0	210 $\pm$ 16	11
Ap <sub>4</sub> U	17.6 $\pm$ 2.3	324 $\pm$ 12	18

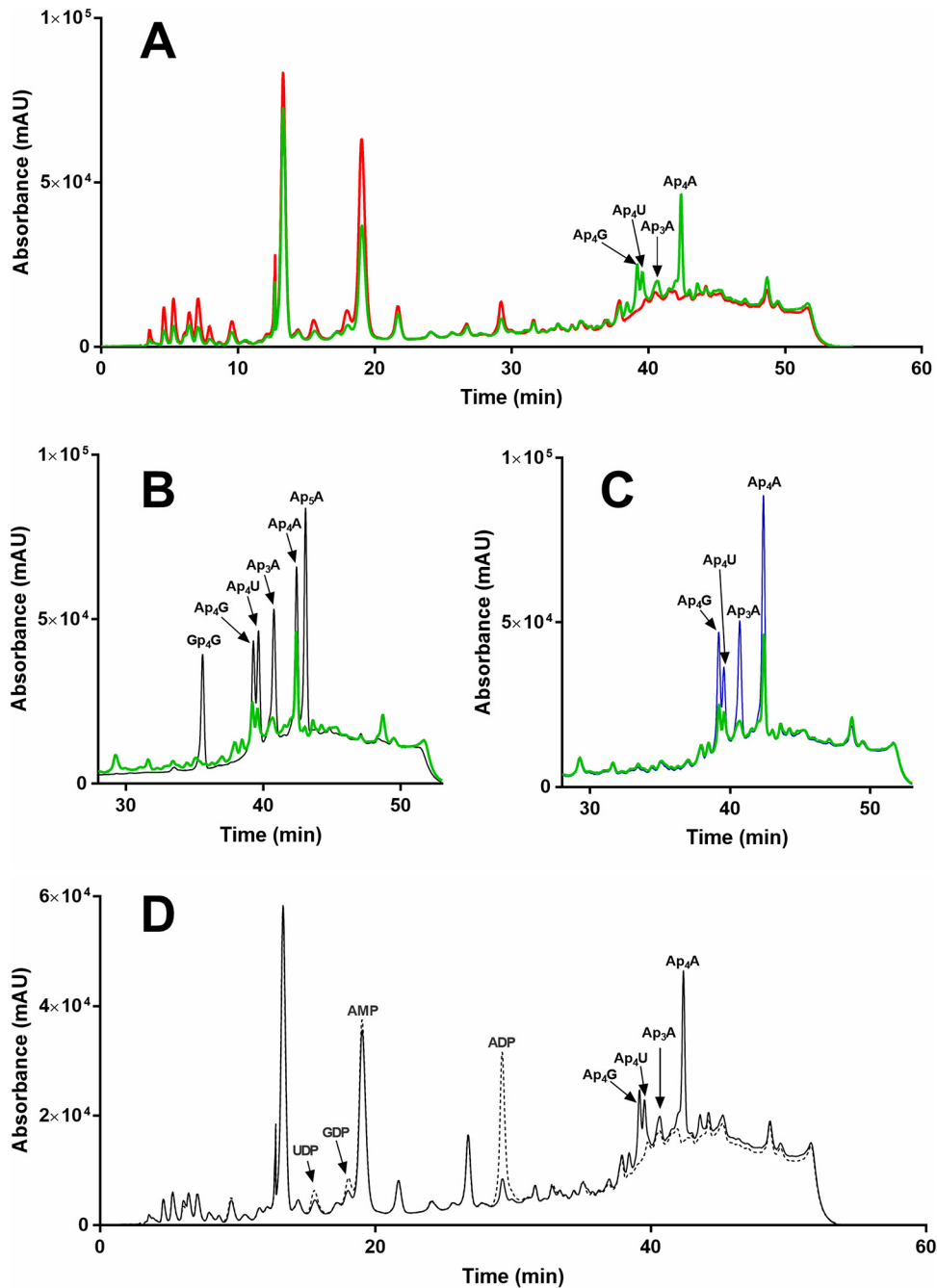
enzyme exhibits a marked preference for Ap<sub>4</sub>A, which is mainly exerted at the substrate affinity level. Indeed, the catalytic efficiency is at least twice that determined for the other dinucleoside tetraphosphates tested, despite a lower  $k_{\text{cat}}$  value. Analysis of product inhibition revealed that the enzyme is not inhibited by the ADP product. Based on all the above results, we proposed a functional assignment of SaYqeK as a symmetrically cleaving Ap<sub>4</sub>A hydrolase.

As outlined below, the available crystal structures of YqeK bacterial orthologs show the presence of a diiron cluster in the putative active site, indicating that the enzymatic activity might be iron dependent. Indeed, we found that an overnight preincubation of YqeK with 50 mM EDTA at room temperature fully inactivated the enzyme, provided that 50 mM EDTA was also present in the assay mixture, whereas a 25% inhibition was measured without the preincubation step. These results confirm that YqeK is metal dependent and suggest that the metal-binding site is poorly accessible to chelators. As we found that the recombinant enzyme was active in the absence of added metal ions in the reaction mixtures, we hypothesized that the metal was taken up from the host during protein synthesis.

**In vivo functional activity of yqeK gene.** To validate our functional assignment, we grew *B. subtilis* wild type and the strain lacking the *yqeK* gene (GenBank BSU25630, Uniprot ID P54456) to compare the *in vivo* levels of the dinucleotides that are hydrolyzed *in vitro* by recombinant YqeK. No difference in the growth rate of wild-type and mutant cells was observed. As shown in Fig. 2A, the HPLC analysis of the mutant revealed several peaks that were not detectable in the wild type. Among them, we identified Ap<sub>4</sub>G, Ap<sub>4</sub>U, Ap<sub>3</sub>A, and Ap<sub>4</sub>A based on the finding that the corresponding chromatographic peaks (i) coeluted with standard molecules (Fig. 2B), (ii) increased in spiked samples (Fig. 2C), and (iii) disappeared after incubation with an excess of SaYqeK (Fig. 2D). Notably, a concomitant increase of UDP, GDP, AMP, and ADP, i.e., the expected products of YqeK-catalyzed reactions, was observed (Fig. 2D). These results indicate that YqeK affects the intracellular levels of these dinucleotides.

**Phylogenetic distribution of yqeK and apaH.** As current literature suggests that Ap<sub>4</sub>A hydrolase symmetrical cleavage activity belongs to the ApaH family, we next explored the distribution of *yqeK* and *apaH* genes using comparative genomics. Using *SayqeK* as the query, we performed a homology search analysis against a selection of one hundred bacterial genomes that maximized the diversity within the eubacterial kingdom. As a result, we observed an overall presence of YqeK in Gram-positive bacteria. All analyzed *Firmicutes*, including the class *Mollicutes*, invariably possess the *yqeK* gene (Fig. 3A). Outside *Firmicutes*, *yqeK* is found in the *Thermotoga* and *Thermus-Deinococcus* groups, a few species of *Cyanobacteria* and *Spirochaetes*, and in an isolated case of *Actinobacteria* (*Rubrobacter xylanophilus*) (Fig. 3A). In contrast, *apaH* is consistently distributed in Gram-negative bacteria of the classes *Beta-* and *Gammaproteobacteria*, and no instances of *yqeK* and *apaH* genomic cooccurrence have been detected (Fig. 3A). This observation, along with our biochemical and *in vivo* functional characterization, supports the notion that YqeK represents the symmetrically cleaving Ap<sub>4</sub>A hydrolase in *Firmicutes*.

**Genomic neighborhood of yqeK.** Comparative genome analysis shows that in *Firmicutes* *yqeK* is always adjacent to *nadD*, a key gene in the NAD biosynthetic pathway, with the only exception being *Carboxydotherrmus hydrogenoformans* (Fig. 3B). Such genetic proximity is also found outside the *Firmicutes*, although less frequently. Nonetheless, *yqeK*, when present, invariably cooccurs with *nadD* in the genome. In addition, the two genes are fused in *Mycoplasma genitalium* and *Elusimicrobium minutum*, thus reinforcing their putative functional association (Fig. 3B). To further investigate such functional association, we asked whether NaAD, the product of the NadD-catalyzed reaction, and NAD, the ultimate product of the NadD-driven pathway, might be substrates or inhibitors of the Ap<sub>4</sub>A hydrolase activity of YqeK. However, YqeK proved to be insensitive to both dinucleotides. Likewise, NadD activity was not affected by Ap<sub>4</sub>A. In addition, NadD activity did not change in the presence of YqeK, and YqeK

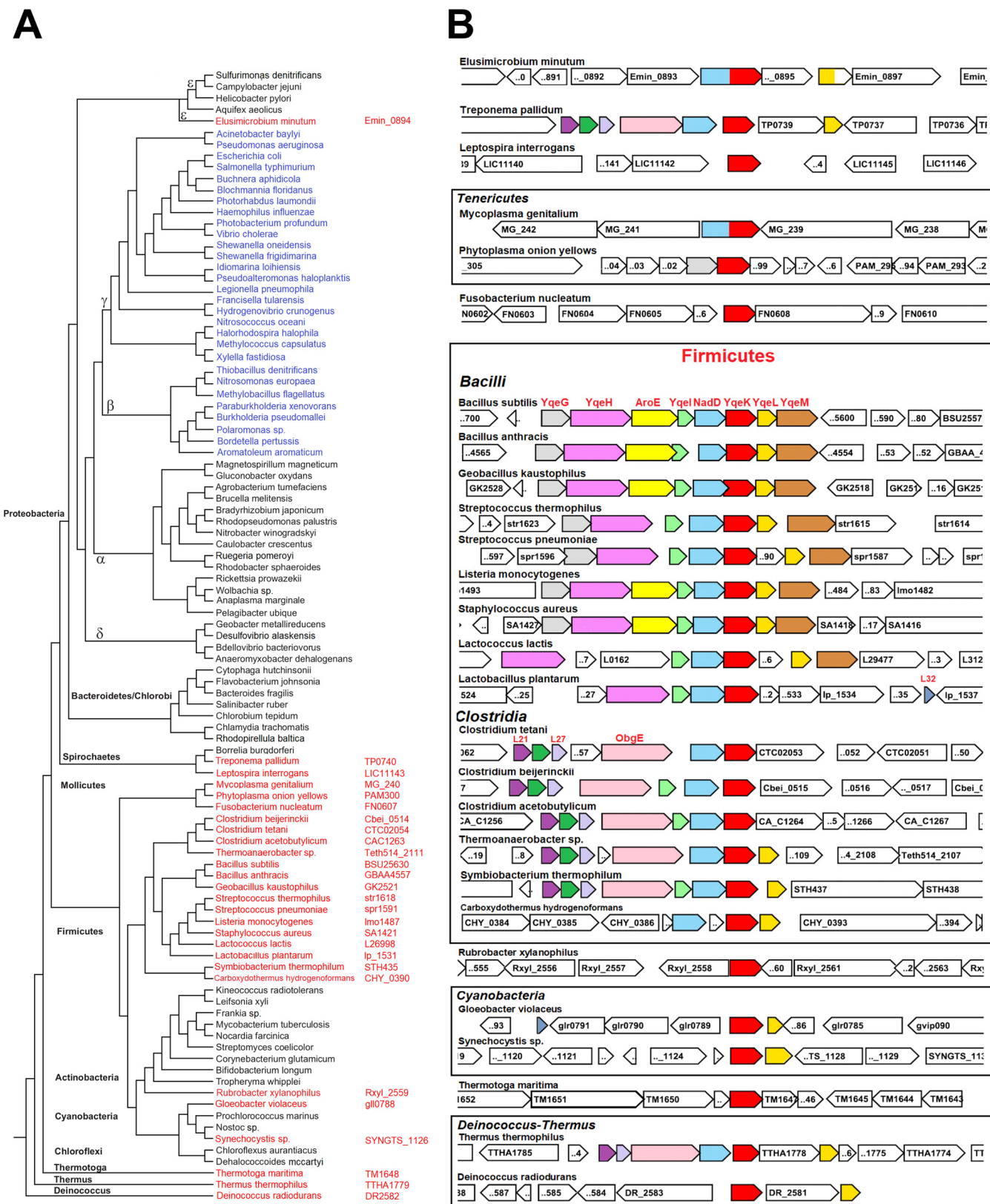


**FIG. 2** *In vivo* validation of YqeK function. HPLC absorbance profiles at 260 nm of ethanol-boiled extracts from wild type *B. subtilis* (red) and the  $\Delta yqeK$  mutant (green) (A), mutant extract (green) and nucleotide standards (black) (B), mutant extract (green) and nucleotide-spiked extract (blue) (C), or mutant extract incubated in the absence (solid line) and in the presence (dotted line) of SaYqeK enzyme (D).

activity was insensitive to the presence of NadD. These results appear to rule out direct cross talk between the two enzymes.

Another strong association is with *yqeL*, encoding the ribosomal silencing factor RsfS (formerly lojap), which is often next to *yqeK* and even fused to it in some cases (as in *Clostridium* sp. strain SY8519 and *Eubacterium nodatum* [not shown]). Other closely neighboring genes, often belonging to the same operon as in *Bacilli*, include *yqeG*, *yqeH*, *aroE*, *yqeI*, and *yqeM* (Fig. 3B). The *yqeG* gene encodes a 5'-nucleotidase of broad specificity with an as yet uncertain physiological role. In *B. subtilis*, YqeG is necessary for





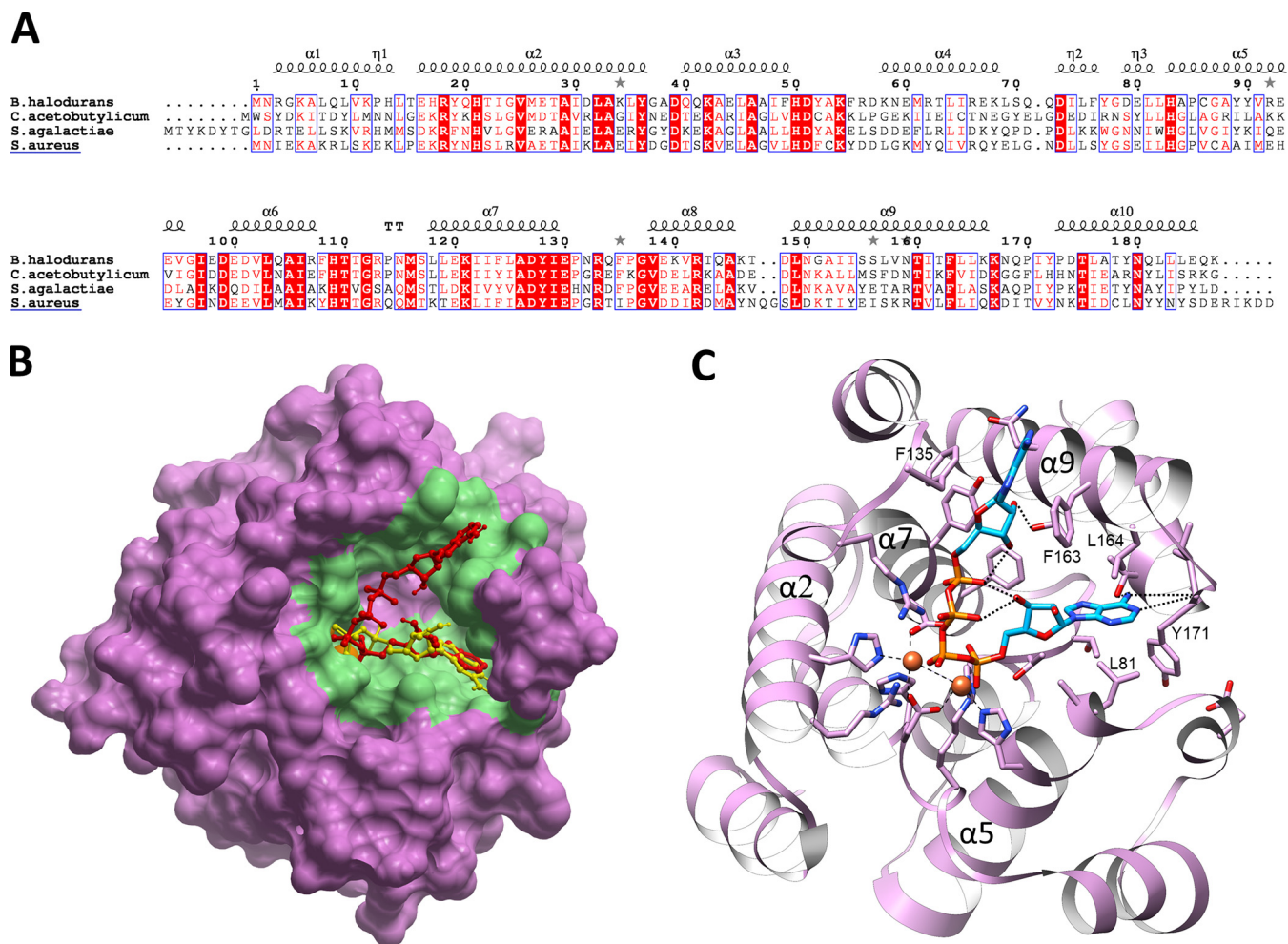
**FIG. 3** Phylogenomics of *yqkE*. (A) Phylogenetic distribution of YqkE (red) and ApaH (blue) in bacteria. Locus tags of YqkE orthologs are next to the species name. (B) Genomic context of the *yqkE* gene. Genes are colored according to the locus organization in *B. subtilis* and *Clostridium tetani*. The gene abbreviations are explained in the text.

normal colony formation on solid medium and is induced by oxidative stress (21). The *yqeH* gene encodes an essential GTPase of the Era/Obg family involved in ribosome 30S assembly (22). In *Clostridia*, *Thermus thermophilus*, and *Treponema pallidum*, the *obgE* gene replaces *yqeH* in the *yqeK*-containing operons. Notably, ObgE is another essential GTPase that has been recently proposed as a (p)ppGpp binding target with a role in the stringent response to amino acid starvation and in other downstream effects of (p)ppGpp metabolism (23). The *aroE* gene, found in the majority of *Bacilli*, encodes NAD(P)-dependent shikimate dehydrogenase, an indispensable enzyme committed to the biosynthesis of aromatic amino acids (24). YqeI, present in nearly all *Firmicutes*, is a putative rRNA-binding protein, required for the proper assembly of 50S and 30S ribosome subunits (25). The last member of the *yqeK*-containing operon is *yqeM* (*smtA* in *E. coli*), a gene typically found in *Bacilli* and shown to be a dispensable, putative S-adenosylmethionine-dependent methyltransferase overexpressed under biofilm conditions in *S. aureus* (26).

**YqeK structural analysis.** YqeK is a member of the largely uncharacterized HD domain superfamily of enzymes (27). This superfamily comprises phosphohydrolases with nucleotidase or phosphodiesterase activity and oxygenases that catalyze the O<sub>2</sub>-dependent cleavage of C-C or C-P bonds (28). HD-domain proteins are metal dependent, and enzymes with one, two, or three metal-binding sites have been characterized. YqeK belongs to a subfamily identified as COG1713 (a predicted HD-superfamily hydrolase involved in NAD metabolism) that features a diiron catalytic cluster, as seen in the three crystal structures of YqeK orthologs from *Streptococcus agalactiae* (PDB ID, 2OGI), *Bacillus halodurans* (PDB ID, 2O08), and *Clostridium acetobutylicum* (PDB ID, 3CCG), solved by the Joint Center for Structural Genomics. Of note, the *S. agalactiae* and *B. halodurans* YqeK structures have dGDP and GDP molecules bound to the dimetal centers, whereas a free phosphate is present in the *C. acetobutylicum* protein. By using the structure of *B. halodurans* YqeK, which shares the highest sequence identity (42%) with *SaYqek* (Fig. 4A), we employed ICM Pocket Finder to predict the shape of the putative active site. The prediction was for an extended, V-shaped cleft (with an area of 633 Å<sup>2</sup> and a volume of 651 Å<sup>3</sup>) half occupied by the GDP molecule, with the dinuclear cluster located at its vertex (Fig. 4B). Molecular docking of Ap<sub>4</sub>A revealed that one of the two ADP moieties of Ap<sub>4</sub>A ligand overlaps with the natural nucleotide found in the ligand-bound structure (Fig. 4B). The adenine ring of this ADP moiety might be stabilized by two H-bonds with the Tyr171 backbone amide and further coordinated by van der Waals contacts with Leu81 and Leu164 (Fig. 4C). Notably, this conformation also fits with the docking pose of the ADP stand-alone ligand (not shown), suggesting that this moiety of Ap<sub>4</sub>A leaves the active site last. Indeed, in our docking simulation, the other ADP moiety occupies the less-buried arm of the V-shaped binding cleft, with the adenine slightly protruding outward and only partially stabilized by Phe135 and Phe163 (Fig. 4C). In agreement with our gel filtration experiment, all the available YqeK structures share a dimeric assembly with an average interface area of ~650 Å<sup>2</sup>. Analysis by the PISA server classifies the dimeric structures in complex with the nucleotides as stable, in contrast with the unstable phosphate-bound complex, suggesting that the dimer formation is favored by the ligand binding.

## DISCUSSION

Despite great advances in the functional annotation of genes since the publication of the first fully sequenced genome, a significant fraction of bacterial genes still lack functional characterization (29). Among them is the gene *yqeK*, which is conserved in most Gram-positive species. In this work, we identified it as an Np<sub>n</sub>N hydrolase gene, highly specific for Ap<sub>4</sub>A. Among known Np<sub>n</sub>N hydrolases, YqeK mostly resembles the ApaH enzyme, which is the major Ap<sub>4</sub>A hydrolase in proteobacteria. Both enzymes are active toward several Np<sub>n</sub>N nucleotides but exhibit the highest specificity toward Ap<sub>4</sub>A, which is symmetrically cleaved into ADP molecules. The efficiency of YqeK in hydrolyzing Ap<sub>4</sub>A is very similar to that exhibited by ApaH (30), and YqeK significantly affects the *in vivo* intracellular level of the dinucleotide in *B. subtilis*, as reported for ApaH in



**FIG. 4** YqeK structural analysis. (A) Structure-based sequence alignment of SaYqeK and structurally characterized YqeK proteins.  $\alpha$  and  $3_{10}$  helices ( $\eta$  labels) are shown as spirals on top of the alignment according to the structure of *B. halodurans* YqeK (PDB 2008). TT represents sharp turns. Identical amino acids are indicated by a solid red background, while similar amino acids are boxed. UNIPROT IDs for the sequences used are as follows: Q9KD90, *Bacillus halodurans*; Q97JL1, *Clostridium acetobutylicum*; Q8DY32, *Streptococcus agalactiae*; A0A0Y9ZU75, *Staphylococcus aureus*. (B) Ap<sub>4</sub>A (red) docking pose in putative binding site (colored in green) of *B. halodurans* YqeK. Ap<sub>4</sub>A is superposed with cocrystallized GDP ligand (yellow). (C) Close-up view of *B. halodurans* YqeK active site with modeled Ap<sub>4</sub>A. Residues that are likely to interact with the dinucleoside tetraphosphate are displayed. Iron ions are shown as orange spheres, and the side chains of the coordinating residues are shown in stick format and colored by atom type.

gammaproteobacteria (7, 12). Deletion of the *yqeK* gene in *B. subtilis* also increased the content of Ap<sub>4</sub>G and Ap<sub>4</sub>U. The *in vivo* occurrence of these dinucleotides has been well documented in *E. coli* as the result of the activity of aminoacyl-tRNA transferases (31). To our knowledge, this is the first report showing their occurrence in *B. subtilis*. Despite the described similarities, ApaH and YqeK belong to different families, i.e., the serine/threonine phosphatase family (32) and the HD domain superfamily (27), respectively, which is suggestive of a different catalytic mechanism. Our phylogenetic analysis revealed that the two enzymes occur in a mutually exclusive manner, with YqeK present in all bacterial species that lack ApaH. Furthermore, both genes play a role in the stress response, as evidenced by gene deletion studies. In particular, *E. coli* *apaH* mutants show impaired growth and change in morphology under starvation conditions or heat stress (1), suggesting that the enzyme is essential for the cell to adapt to stress. Similarly, the deletion of *yqeK* in *B. subtilis* impairs the bacterium's capability to form biofilm, which is a typical adaptation to environmental cues, as cells living in biofilms are better adapted to survive periods of environmental stress (33). *apaH* and *yqeK* also share similarities in genomic context. In *E. coli*, *apaH* is part of an operon comprising the gene *ksgA*, encoding an rRNA modification enzyme involved in 16S rRNA maturation



and ribosome biogenesis, and is essential for the fidelity of translation during antibiotic stress (34, 35). Similarly, *yqeK* is often found associated with genes encoding proteins involved in ribosome assembly/maturation (*yqeH*, *yqeG*, and *yqeI*) and the control of protein translation under nutrient shortage (*yqeL*). Notably, in *B. subtilis* these genes together with *yqeK* form a large operon, also comprising genes encoding metabolic enzymes like shikimate dehydrogenase (*aroE*) and NadD (*nadD*) that catalyze key steps in the biosynthesis of aromatic amino acids and the coenzyme NAD, respectively. Expression of this operon is constitutive, but increases during germination and under conditions of oxidative stress (36), indicating that it might represent a complex and integrated response to stress signals. Consistent with this, and with the findings that shortage of aromatic amino acids is a hallmark of starvation (37) and that coping with stress-induced events requires the activity of several NAD-consuming enzymes (38), the upregulation of *aroE* and *nadD* might ensure amino acid and NAD replenishment, respectively. In general, the stress response requires cellular reprogramming at metabolic, transcriptional, and translational levels. Thus, a *yqeK*-containing operon might control the coordination of multiple responses at these various levels. Although other biological substrates could exist for YqeK, our finding that the enzyme efficiently hydrolyzes Ap<sub>4</sub>A both *in vitro* and *in vivo* indicates that maintenance of Ap<sub>4</sub>A levels should be coordinated with NAD and amino acids synthesis, as well as with the control of translation.

Evidence is accumulating that targeting of Ap<sub>4</sub>A hydrolase might present a promising approach to weakening the bacterial response to stress signals (including antibiotics) and to decreasing pathogenicity. In this view, the functional annotation of YqeK as a novel Ap<sub>4</sub>A hydrolase in a group of Gram-positive pathogenic bacteria discloses a novel target for antibacterial strategies.

## MATERIALS AND METHODS

**Preparation of recombinant protein.** Plasmid pOHypY.SA.4, carrying *Staphylococcus aureus yqeK* under the control of the T7 promoter and encoding a protein with an N-terminal His tag, was kindly provided by Andrei L. Osterman (Sanford Burnham Prebys Medical Discovery Institute, San Diego, CA). The construct was used to transform BL21(DE3) cells for protein expression. Cells were grown at 37°C in Luria Bertani medium supplemented with 0.1 mg/ml ampicillin. After reaching an optical density at 600 nm (OD<sub>600</sub>) of 0.6, expression was induced with 1 mM isopropyl-β-D-thiogalactoside. After 4 h of induction, cells were harvested by centrifugation at 5,000 × *g* for 10 min. A pellet from 250 ml of culture was resuspended in 13 ml of lysis buffer (50 mM HEPES [pH 7.5], 0.3 M NaCl) containing 1 mM phenylmethanesulfonyl fluoride and proteases inhibitors. The recombinant protein was purified to homogeneity by nickel affinity chromatography. Briefly, the cell suspension was passed twice through a French press (18,000 lb/in<sup>2</sup>) and centrifuged at 20,000 × *g* for 25 min at 4°C. The supernatant was loaded onto a 1 ml Ni-NTA resin (GE Healthcare) equilibrated with lysis buffer. After a washing with 30 mM imidazole in lysis buffer, elution was performed with a linear gradient from 30 to 350 mM imidazole. Fractions containing the recombinant protein were pooled and applied to a Superose 12 HR 10/30 column (GE Healthcare) eluted with lysis buffer. About 6 mg of pure protein was obtained from a starting volume of 250 ml of culture.

**Thermal shift assays.** The assay was performed according to reference 39. The mixtures contained 1 μM SaYqeK, 5 μM ligand, 10× Sypro Orange (Invitrogen), 40 mM HEPES (pH 7.5), and 0.25 M NaCl in a final volume of 25 μl. They were heated from 40 to 99°C with a heating rate of 0.5°C every 5 s in a real-time PCR device (Rotor-Gene 3000, Corbett Life Science). The fluorescence intensity was measured with excitation and emission wavelengths of 470 and 585 nm, respectively.

**Enzyme activity assays.** For YqeK activity, reaction mixtures contained 40 mM HEPES (pH 7.5), 0.25 M NaCl, and appropriate amounts of substrate and YqeK protein in 100 μl volume. For kinetic analyses, 1 to 2 ng protein was used, and the substrates ranged from 1 μM to 160 μM. Incubation was carried out at 37°C for an appropriate incubation time ensuring both a linear change of product formation with time and a substrate consumption of less than 5%. Reactions were stopped with 0.6 M HClO<sub>4</sub>, and after 10 min on ice, the samples were centrifuged for 1 min at 12,000 × *g*. The supernatants were neutralized with 0.8 M K<sub>2</sub>CO<sub>3</sub> and injected into an HPLC system. Nucleotide separation was performed as described in reference 40. Control mixtures in the absence of YqeK were always processed in parallel. *K<sub>m</sub>* and *k<sub>cat</sub>* values were calculated by fitting initial rates to a standard Michaelis-Menten model using the software Prism 6 (GraphPad). The effect of NAD, NaAD, and ADP on the Ap<sub>4</sub>A hydrolase activity of YqeK was assayed in the presence of 80 μM Ap<sub>4</sub>A and 0.25 mM effector.

Pure recombinant NadD from *B. anthracis* was prepared as described in (41). NadD reaction mixtures contained 50 mM HEPES (pH 7.5), 10 mM MgCl<sub>2</sub>, 0.1 mM nicotinate mononucleotide (NaMN), 0.1 mM ATP, and 0.05 mU of enzyme in a final volume of 50 μl, in the presence and absence of 80 μM Ap<sub>4</sub>A. Incubation was carried out at 37°C for 20 min, and product formation was analyzed by HPLC as described

above. The effect of the NadD protein on the Ap<sub>4</sub>A hydrolase activity and vice versa was tested by assaying YqeK or NadD in the presence of the other enzyme at molar ratios ranging from 1:1 to 1:10.

**Deletion mutant analysis.** *Bacillus subtilis* subsp. *subtilis* 168 (BGSCID: 1A1) and *ΔyqeK* knockout mutant (BGSCID: BKE25630) (42) were obtained from the Bacillus Genetic Stock Center (OH). Cells were grown to mid-log phase in Luria-Bertani medium and harvested by centrifugation for 10 min at 5,000 × *g*. Pellets were washed twice with 0.9% NaCl and stored at –80°C. For the extraction of the nucleotides, pellets deriving from 15 ml of culture were resuspended in 3 ml of boiling buffered ethanol (75% ethanol in 10 mM HEPES, pH 7.1) and suspensions were incubated at 84°C for 3 min (43). After cooling on ice for 3 min, samples were centrifuged for 5 min at 20,000 × *g* and supernatants were dried in a Speed-vac concentrator before resuspension in 0.75 ml double-distilled water. Samples were acidified to pH 5.0 by addition of 1 μl of 0.4 M HClO<sub>4</sub> and nucleotides were analyzed by HPLC as reported previously (40). The column was maintained at 25°C in order to separate Ap<sub>4</sub>G from Ap<sub>4</sub>U. Spiked samples were prepared by adding standard nucleotides in the samples before HPLC injection.

To evaluate the effect of YqeK activity on the nucleotide extract prepared from the mutant, 100 μl of extract was incubated with 0.5 μg recombinant SaYqeK in 50 mM HEPES, pH 7.5, and 0.3 M NaCl, in a final volume of 100 μl. A control mixture in the absence of YqeK was processed in parallel. After a 15-min incubation at 37°C, samples were analyzed by HPLC as described above.

**Comparative genome analysis and bioinformatics tools.** Orthologs of YqeK and ApaH were identified by PSI-BLAST (44) searches (E value cutoff, e<sup>–20</sup>). We used a representative set of 100 bacterial genomes as the organismal search set (Genomic Encyclopedia of Bacteria and Archaea project) (45). Physical clustering and gene fusions were analyzed with the microbial genomic context viewer (46) and STRING 11.0 (47). Structure-based multiple alignment was constructed using PROMALS3D (48) and rendered with Esript 3.0 (49). Visualization, comparison of protein structures, and molecular docking were performed with Molsoft ICM 3.8. We used Pocket Finder (50) in ICM for automatic detection of ligand-binding sites in the high resolution (1.9 Å) crystal structure of *B. halodurans* YqeK (PDB entry 2O08). One of the identified pockets, encompassing the cocrystal ligand and the metal dinuclear center, has been used for docking simulations. Ligands were fully flexible, and the receptor was set as rigid during all docking studies.

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