The Tyrosine-Autokinase UbK Is Required for Proper Cell Growth and Cell Morphology of Streptococcus pneumoniae

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Protein phosphorylation is a key post-translational modification required for many cellular functions of the bacterial cell. Recently, we identified a new protein-kinase, named UbK, in Bacillus subtilis that belongs to a new family of protein-kinases widespread in bacteria. In this study, we analyze the function of UbK in Streptococcus pneumoniae. We show that UbK displays a tyrosine-kinase activity and autophosphorylates on a unique tyrosine in vivo. To get insights into its cellular role, we constructed a set of pneumococcal ubk mutants. Using conventional and electron microscopy, we show that the ubk deficient strain, as well as an ubk catalytic dead mutant, display both severe cell-growth and cell-morphology defects. The same defects are observed with a mutant mimicking permanent phosphorylation of UbK whereas they are not detected for a mutant mimicking defective autophosphorylation of UbK. Moreover, we find that UbK phosphorylation promotes its ability to hydrolyze ATP. These observations show that the hydrolysis of ATP by UbK serves not only for its autophosphorylation but also for a distinct purpose essential for the optimal cell growth and cell-morphogenesis of the pneumococcus. We thus propose a model in which the autophosphorylation/dephosphorylation of UbK regulates its cellular function through a negative feedback loop.

Keywords: Streptococcus pneumoniae, protein phosphorylation, tyrosine-kinase, ATP hydrolysis, cell-morphogenesis

INTRODUCTION

Streptococcus pneumoniae (the pneumococcus) is a Gram-positive bacterium, living as a commensal in healthy adults and children. In immature and/or immunocompromised people, the pneumococcus can, however, become pathogenic and causes diseases that range from otitis, pneumonia to meningitis with sepsis (Kadioglu et al., 2008; Henriques-Normark and Normark, 2010). Despite the availability of antibiotic treatments and vaccines, S. pneumoniae still kills more than 1.2 million persons each year and is in the WHO list of priority pathogens for research and development of new antibiotics (Tacconelli, 2017).
Evidences have accumulated that protein phosphorylation on hydroxylated residues (i.e., serine, threonine and tyrosine) catalyzed by serine/threonine-kinases and tyrosine-kinases is crucial for the biology of bacteria (Manuse et al., 2016; Mijakovic et al., 2016). eSTKs (for eukaryotic-like Serine and Threonine kinases), that possess a catalytic domain structurally homologous to that of eukaryotic protein-kinases, have been shown to regulate different physiological processes like the cell cycle, virulence and central and secondary metabolisms (Burnside and Rajagopal, 2012; Mijakovic and Macek, 2012; Canova and Molle, 2014; Fleurie et al., 2014b; Wright and Ulijasz, 2014; Dworkin, 2015; Manuse et al., 2016). eSTKs are widespread in bacteria, but with a highly variable distribution (Dworkin, 2015). On the other hand, phosphorylation on tyrosine is mainly achieved by the bacterial idiosyncratic protein-tyrosine kinase family BY-kinases (Bacterial tyrosine kinases) (Grangeasse et al., 2007; Jadeau et al., 2008; Mijakovic et al., 2016). Like eSTKs, BY-kinases regulate several biological processes and their best studied function concerns their role in the biosynthesis and export of extracellular polysaccharides (Standish et al., 2014; Nourikyan et al., 2015; Mijakovic et al., 2016). They are also widely conserved in bacterial genomes and most of bacterial species encode for at least one BY-kinase (Jadeau et al., 2012).

In the pneumococcus, only one eSTK and one BY-kinase, StkP and D respectively, are produced. Recent studies have demonstrated the critical role of D in the polysaccharide capsule synthesis and export as well as the coordination of this process with the cell cycle (Henriques et al., 2011; Nourikyan et al., 2015; Mercy et al., 2019). The capsule is the main virulence factor of the pneumococcus and its composition is highly variable (more than 90 serotypes known to date). On the other hand, StkP is the central regulator of cell division and morphogenesis (Beilharz et al., 2012; Fleurie et al., 2012, 2014a,b; Grangeasse, 2016; Zucchini et al., 2018).

We have recently identified an unprecedented type of protein-kinase in Bacillus subtilis (Nguyen et al., 2017). This protein was named UbK for Ubiquitous bacterial Kinase as it is present in most bacterial genomes. Strikingly, ubk genes are found neither in Archea nor in eukaryotic genomes (Tepljakov et al., 2002). UbK proteins possess the canonical Walker A-motif G/AX4GKT/S found in the large family of the P-loop proteins, including BY-kinases (Leipe et al., 2002; Grangeasse et al., 2007). However, besides this, the crystal structures of UbK from Haemophilus influenzae and B. subtilis showed that the structure of UbK proteins share little similarities with that of BY-kinases and other ATP-binding proteins with a Walker A motif (Reinstein et al., 1990; Nguyen et al., 2017). Interestingly, UbKs of H. influenzae, E. coli or B. subtilis show a weak ATPase activity (Campbell et al., 2007; Karst et al., 2009) but their ability to autophosphorylate and to phosphorylate in vitro the surrogate substrate Myelin-Basic-Protein has been reported only for UbK from B. subtilis (Nguyen et al., 2017). In B. subtilis, a recent investigation suggests a possible role in resistance to oxidative stress (Nguyen et al., 2017). ubks were described as being essential genes in S. pneumoniae (Zalacain et al., 2003; Havarstein et al., 2006), B. subtilis (Kobayashi et al., 2003), Mycoplasma pulmonis (French et al., 2008) and E. coli (Freiberg et al., 2001; Campbell et al., 2007). On the other hand, other studies have concluded that ubk would be dispensable in S. pneumoniae (Molzen et al., 2011), B. subtilis (Hunt et al., 2006; Karst et al., 2009; Nguyen et al., 2017), and Anabaena sp. strain (Yoon et al., 2003). One can therefore not exclude that these opposing observations either convey the presence of suppressive mutations appearing when deleting ubk genes or that the dispensability/essentiality of ubk is strain-specific.

Here, we analyze the cellular function of the pneumococcal UbK (spr1761 in the R6 strain) together with its kinase and ATP hydrolysis activities. We notably demonstrate that UbK ATP hydrolysis activity, but not autophosphorylation, is crucial for S. pneumoniae growth. However, we also show that UbK autophosphorylation increases ATP hydrolysis and is detrimental for the cell growth. Last, we observe that cell growth defects correlate with an abnormal cell morphology. Therefore, we propose a model in which autophosphorylation/dephosphorylation of UbK finely tunes and balances its ATP hydrolysis, an activity that is required for the optimal growth and proper morphogenesis of the pneumococcus cell growth. Altogether, this work highlights the importance of this new atypical tyrosine-kinase in the bacterial cell physiology.

RESULTS

ubk Is Essential for Streptococcus pneumoniae

To delete ubk, we used the Janus markerless and non-polar strategy (Sung et al., 2001). While transformants of the R6 derivative strains were usually obtained after a 16-h culture with transformation levels ranging from 1 to 10%, small colonies of kanamycin resistant mutants were obtained with very low frequency (0.001%) after 30 h of growth. We confirmed that ubk was properly deleted and substituted for the Janus cassette (Δubk:kan-rpsL) in these mutant colonies by PCR amplification and sequencing of the targeted locus. This observation suggested that ubk was either crucial for the pneumococcus biology or that deletion of ubk generates a polar effect affecting the expression of downstream genes. To differentiate between the two possibilities, we first expressed ubk ectopically using the pCEN maltose-inducible platform (Guiral et al., 2006) in WT cells (ubk+:ubkCEP). Then, we substituted the chromosomal copy of ubk for the Janus cassette (Δubk:kan-rpsL-ubkCEP). Colonies were readily obtained with normal growth and transformation frequency. This excluded the second hypothesis and suggested that the deletion of ubk requires suppressive mutations(s). To check this, we statistically analyzed the number of clones obtained by transformation of the WT and the ubk+:ubkCEP strains either with a whole genomic DNA of a Δubk:spc-rpsL mutant or with a Δubk:spc-rpsL. PCR amplified DNA fragment (Figure 1A). The ubk+:ubkCEP strain transformed efficiently the two donor DNAs. By contrast, the WT strain transformed the genomic DNA with a 2 log decreased efficiency compared to the ubk+:ubkCEP strain and transformed the PCR DNA with a 4 log decreased efficiency compared to the ubk+:ubkCEP strain.
strain. This suggested the presence of a single suppressive mutation co-selected with ubk deletion in the genome of original mutants. To identify this suppressive mutation, we performed whole genome sequencing of two independent Δubk mutants together with that of the WT parental strain. A single mutation differentiating the WT and the Δubk mutant genomes was unambiguously identified. This mutation consisted in an A to C transversion in position 362 relative to the ATG of the spr1397 (asnS) coding sequence. The mutation led to an aspartate to alanine amino-acid substitution in position 121 of the AsnS protein. spr1397 was shown already to be an essential gene (Thanassi et al., 2002). We nevertheless transferred the D121A mutation in the spr1397 gene of WT strain (asnS-D121A) and the resulting mutant, called suppressor-WT, was transformed again by the Janus cassette to inactivate chromosomal ubk. Colonies were readily obtained with normal transformation level and growth. Confirming that the single A362 to C362 transversion in the asnS gene is required to get cells devoid of ubk (Figure 1B).

UbK Autophosphorylates on a Single Tyrosine Residue in vivo

UbK of S. pneumoniae was previously shown in vitro either able to autophosphorylate on tyrosine and to phosphorylate the surrogate substrate myelin basic protein (MBP) (Nguyen et al., 2017). We first tested the relevance of UbK autophosphorylation in vivo. For that, we purified UbK fused to the GFP (GFP-UbK) from pneumococcal cells (strain Δubk-gfp-ubkCEP). As control, we also constructed a strain expressing a UbK kinase-dead form in which the conserved lysine 36 of the Walker A motif is mutated for an arginine (strain Δubk-gfp-ubkK36RCEP). Indeed, mutation of this catalytic lysine prevents ATP hydrolysis of Walker A ATP-binding proteins (Reinstein et al., 1990; Soulat et al., 2007; Karst et al., 2009). In addition, we checked that Ubk-K36R overexpressed and purified from E. coli cells is unable to autophosphorylate on tyrosine in vitro (Figure 2). The same observation was made previously with UbK (i.e., YdiB) from B. subtilis mutated on lysine 41 of the Walker A (Nguyen et al., 2017). We then prepared crude extracts of the Δubk-gfp-ubkCEP and the Δubk-gfp-ubkK36RCEP pneumococcal strains and performed GFP trap using GFP antibodies immobilized on agarose. Immunopurified proteins were then analyzed by SDS-PAGE and UbK phosphorylation was analyzed by Western blot using the 4G10 antiphosphotyrosine antibody. As shown in Figure 2, a phosphorylation signal was detected for GFP-UbK but not for GFP-UbK-K36R confirming that UbK is autophosphorylated on tyrosine in vivo. Then, we analyzed GFP-UbK by LC-MS/MS mass spectrometry. It showed unambiguously that GFP-UbK was phosphorylated on the tyrosine 58 (Supplementary Figure 1A). To confirm this, and check if Y58 represented the single autophosphorylation site of UbK, we constructed a strain expressing GFP-UbK which is Y58 was substituted for a phenylalanine (strain Δubk-gfp-ubkK58FCEP). Using the same procedure, GFP-UbK-Y58F was GFP-trapped and analyzed by SDS-PAGE and Western blot with the 4G10 antiphosphotyrosine antibody. No phosphorylation signal was detected (Figure 2). Then, we performed Western blots and immunodetections with the antiphosphotyrosine antibodies on whole-cells extracts of WT and Δubk cells to determine if UbK is responsible for the phosphorylation of pneumococcal endogenous proteins in vivo (Supplementary Figure 1B). We did not detect phosphorylation signals in WT cells that are not found in Δubk cells, suggesting that UbK is unable to phosphorylate a substrate, or only at very low levels undetectable by immunolabelling. The signal

![FIGURE 1](image-url)
detected around 40 kDa in both strains suggested that an unknown protein might be phosphorylated on tyrosine but independently of UbK. In addition, to determine whether UbK autophosphorylates in cis or trans, we incubated inactive UbK-K36R and UbK with radioactive [α-32P]-ATP. The difference in size of the tags fused to UbK-K36R and UbK (Supplementary Table 1) that were used for their purification allowed to separate the two proteins by SDS-PAGE. Autoradiography of the gel showed that UbK displayed a radioactive signal whereas UbK-K36R did not (Supplementary Figure 1C). This showed that UbK is unable to trans-phosphorylate UbK-K36R and that UbK only cis-autophosphorylated. UbK is therefore a tyrosine-autokinase cis-autophosphorylating on a single tyrosine, namely tyrosine 58.

ATP Hydrolysis by UbK but Not UbK Autophosphorylation Is Required for Cell Growth

To assess the function of UbK autophosphorylation, we first analyzed the growth of different ubk mutants. As control, we first checked that the suppressive mutation D121A has not effect on the pneumococcal growth (Supplementary Figure 2). Then, we generated strains expressing the different forms of UbK (mutated on the catalytic lysine and Y58) from the ubk chromosomal locus under the control of the native promoter. Two strains expressing UbK carrying either a mutation substituting Y58 for a phenylalanine or for a glutamic acid (strains ubk-Y58F and ubk-Y58E respectively) were constructed. Indeed, mutation of a phosphorylatable tyrosine for a phenylalanine (phospho-ablative) or a glutamic acid (phospho-mimetic) is classically used as the best proxy to mimic a deficient or permanent phosphorylation, respectively (Bidnenko et al., 2013; Lee et al., 2014; Vilcheze et al., 2014). The mutated chromosomal copy represented therefore the only source of UbK protein in these mutant strains (ubk-K36R, ubk-Y58F, and ubk-Y58E).

The analysis of the cell growth of these mutants revealed an unexpected behavior. While cells expressing UbK-K36R grew slowly and similarly to Δubk cells, reflecting that UbK activity is crucial for the pneumococcal growth, ubk-Y58E cells also displayed the same slow and delayed growth (Figure 3). These observations implied that the ability of UbK to bind and hydrolyze ATP is not only dedicated to its autophosphorylation but likely serves another purpose required for the growth of the pneumococcus. On the other hand, it also suggested that permanent phosphorylation of UbK is detrimental for the cell. Supporting this, ubk-Y58F cell grew properly (Figure 3). Moreover, we constructed two other mutants in which we mutated both the catalytic lysine and the autophosphorylation site, generating thus the ubk-K36R-Y58F and the ubk-K36R-Y58E strains. We observed that these two strains behaved like the ubk-K36R and Δubk mutants with a slow and delayed growth (Supplementary Figure 3 and Figure 3). The ubk-K36R mutation is therefore epistatic over ubk phosphorylation and ATP hydrolysis is predominant for the cell growth. This indicates that the ability to bind and/or hydrolyze ATP is crucial for the pneumococcus and further suggests that UbK autophosphorylation could modulate ATP hydrolysis.

The ATP Hydrolysis Activity of UbK Is Modulated by Autophosphorylation

To further check our hypothesis, we analyzed the ability of UbK mutant proteins to hydrolyze ATP. For that, we...
overproduced in *E. coli* and purified UbK mutants fused to a C-terminal 6His-tag (UbK-6His, UbK-K36R-6His, UbK-Y58F-6His and UbK-Y58E-6His) (Supplementary Figure 4). Then, we used the pyruvate-kinase/lactate-deshydrogenase assay to measure the ATP hydrolysis activity of UbK (Karst et al., 2009; Typas et al., 2011). As expected, the inactive UbK-K36R-6His protein was unable to hydrolyze ATP (Figure 4A). By contrast, and as previously reported for Ubk of *B. subtilis*, UbK-6His harbored a low ATP hydrolysis activity (31.5 ± 2.8 nanomoles of ATP/mg of UbK/min) (Figure 4B). We further observed a significant difference regarding the ability to hydrolyze ATP between UbK-Y58F-6His and UbK-Y58E-6His. Indeed, UbK-Y58F-6His displayed a reduced activity (23.4 ± 2.7 nanomoles of ATP/mg of UbK/min) compared to UbK-6His whereas UbK-Y58E-6His hydrolyzed ATP more efficiently than UbK-6His (38.3 ± 2.7 nanomoles of ATP/mg of UbK/min). UbK autophosphorylation therefore influences its ability to hydrolyze ATP. To strengthen this observation, we incubated purified UbK-6His with radiolabeled [α-32P]-ATP and varying concentrations of non-radioactive ATP for different times. We observed increasing radioactive phosphorylation signals for UbK as a function of the incubation time (Supplementary Figure 5). The same experiment performed in the presence of increasing concentrations of non-radioactive ATP, showed a decrease of the UbK phosphorylation signal (Supplementary Figure 5). This reflected that non-radioactive ATP competes with the incorporation of radiolabeled [α-32P]-ATP. In light of the immunodetection of tyrosine autophosphorylation made with UbK purified directly from *E. coli* cells (Figure 2), it shows that purified UbK-6His contained a mix of phosphorylated and non-phosphorylated forms of UbK. ATP hydrolysis activity of UbK-6His reflects therefore the activity of both phosphorylated and dephosphorylated forms. The difference between ATP hydrolysis activities of UbK-Y58F-6His and UbK-Y58E-6His confirms that UbK dephosphorylation hampers its ability to hydrolyze ATP. Consistent with the growth defect observed for Δubk, *ubk*-K36R and *ubk*-Y58E pneumococcal mutants compared to WT and *ubk*-Y58F (Figure 3), these findings show that ATP hydrolysis by UbK is modulated by its autophosphorylation and suggest that a balanced ratio between the phosphorylated form and the non-phosphorylated form is required for the pneumococcal growth.

**ubk Mutants With Growth Defects All Show an Abnormal Morphology**

The growth defects of *ubk* mutants prompted us to analyze the morphology of cells by contrast phase microscopy. As shown in
**Figure 5**, all mutants with growth defects (Δubk, ubk-K36R and ubk-Y58E) showed a phenotype characterized by cells with an aberrant morphology. Indeed, cells appeared swelled and bigger. This visual impression was confirmed statistically by analyzing a large number of cells (**Figure 5**). More precisely, the analysis of cell size-parameters showed that these cells are wider than for the WT cells. Consistent with the normal growth measured for ubk-Y58F and ubk complemented (ubk<sup>CEP</sup>-Δubk) cells, these cells did not display any defects and are similar to WT cells. This indicates that the slow and delayed growth of UbK mutants is likely due to a defect in cell morphogenesis and that UbK is required in this process.

The bacterial cell wall, whose main component is the peptidoglycan, is responsible of the bacterial cell shape (Typas et al., 2011). We therefore observed the cells by transmission and scanning electron microscopy to evaluate the presence of potential defects in the cell wall ultrastructure of ubk mutants. As shown in **Figure 6**, transmission electron microscopy (TEM) images showed that the integrity of the cellular envelope was compromised. Indeed, we detected small cell wall extrusion at the surface of cells, reminiscent of cell wall peeling, in Δubk, ubk-K36R and ubk-Y58E mutants. In addition, these defects were associated with an abnormal positioning and/or orientation of the septum plane and shape (**Figure 6**). Such defects were not observed in WT and ubk-Y58F cells. On the other hand, scanning electronic microscopy (SEM) further revealed the presence of gashes in the cell walls of Δubk, ubk-K36R and ubk-Y58E mutants with cells even not properly covered by the cell wall envelop at the poles (**Figure 6**).

Considering the aberrant cell shape of Δubk, ubk-K36R and ubk-Y58E mutants, we questioned the localization of UbK. We therefore analyzed the cellular localization of the diverse UbK mutant forms using fluorescence microscopy. For that, we generated strains expressing the different forms of GFP-UbK (mutated on the catalytic lysine and Y58) from the ubk chromosomal locus. Expression of the GFP-UbK fusions in gfp-ubk, gfp-ubk K36R, gfp-ubk Y58F and gfp-ubk Y58E strains was analyzed by Western blot using an anti-GFP antibody and showed that fusions are produced at comparable levels (**Supplementary Figure 6A**). In addition, these mutant strains grew and presented a similar morphology as their non-fused gfp counterparts (**Supplementary Figures 6B,C and Figure 3**). We observed that UbK localized in the cytoplasm and that phosphorylation or dephosphorylation of UbK had no impact on its localization (**Figure 7**). Indeed, GFP-UbK, GFP-UbK K36R, GFP-UbK Y58F and GFP-UbK Y58E all localize in the cytoplasm. Altogether, these observations indicate that the assembly of the cell wall is compromised in absence of active UbK suggesting...
a role in cell morphogenesis that does not rely on particular localization of UbK.

DISCUSSION

In this work, we have shown that UbK of *S. pneumoniae*, which belongs to a newly identified class of ubiquitous bacterial protein kinases (Nguyen et al., 2017), is crucial for the pneumococcal growth and its morphology. Opposing observations were previously reported concerning the essentiality of UbK (Freiberg et al., 2001; Kobayashi et al., 2003; Zalacain et al., 2003; Havarstein et al., 2006; Hunt et al., 2006; French et al., 2008; Karst et al., 2009; Molzen et al., 2011). We have now clarified the question for *S. pneumoniae* by showing that *ubk* deletion is actually co-selected with a single mutation (Figure 1) leading to the substitution of aspartic acid 151 for an alanine in the AsnS protein. The latter is described as a potential Asn-tRNA synthetase. AsnS enzymes are known to catalyze specific aminoacylation of tRNA<sub>Asn</sub> with asparagine (Asn-tRNA<sub>Asn</sub>) (Hausmann and Ibba, 2008). Another indirect route to Asn-tRNA<sub>Asn</sub> formation is often found in bacteria and involves the GatA/B/C subunit-proteins that transamidate Asp-tRNA<sub>Asn</sub> (Akochy et al., 2004; Mladenova et al., 2014). In the R6 *S. pneumoniae* strain, both the gatA/B/C and the *asnS* genes are found in the genome, suggesting that *asnS* could be dispensable in order to obtain Asn-tRNA<sub>Asn</sub>. Nevertheless, we confirmed the essentiality of *asnS* but not that of the aspartate 151 of AsnS that can be substituted for an alanine and be suppressive for the deletion of *ubk* (Figure 1). A direct relationship between the incorporation of Asn in proteins and the morphogenesis of *S. pneumoniae* has not been reported yet. Studying the potential connection between AsnS and UbK represents therefore a promising venue to better understand the physiology of the pneumococcus in the future. It is, however, tempting to make some assumptions. Indeed, the function of AsnS might be altered or enhanced when aspartate 151 is substituted for an alanine and sufficient to render UbK dispensable. Interestingly, substitution of an aspartic acid for an alanine could mimic a phospho-ablative mutation. A recent comparative genomic study indicates that nature uses this trick by evolving serine, threonine, and tyrosine phosphorylation sites from Asp/Glu residues (Pearlman et al., 2011). An interesting speculation could be that AsnS activity could have been regulated by UbK and that this regulatory switch would have been lost in the course of evolution by the substitution of an Alanine for an Aspartic acid.

Despite *ubk* is largely but exclusively conserved in the bacterial kingdom, it is also intriguing that its genetic environment in *S. pneumoniae* is so different from that of *B. subtilis* and other bacteria like *E. coli* in which the genetic environment of *yjeE*, the *ubk* homolog, is still different (Mangat and Brown, 2008). This suggests that the cellular function of UbK could operate differently and even differ from one bacterium to another. In *S. pneumoniae*, *ubk* is described as the second gene of an operon of 4 genes, the first being *comM*. *comM* codes for a membrane immunity factor that protects competent cells against their own lytic enzymes (Havarstein et al., 2006). ComM was also shown to delay pneumococcal division during competence (Berge et al., 2017). The third *orf*, *spr1760*, codes for a protein of unknown function while the fourth codes for the essential LytR protein, a regulator of the LytR-Cps2A-Psr (LCP) protein family. LytR is proposed to belong to a type of enzymes required for the attachment of anionic cell wall polymers, like teichoic acids and capsular polysaccharides, to peptidoglycan (Kawai et al., 2011) and is also proposed to be involved in cell division and morphogenesis of the pneumococcus (Johnsborg and Havarstein, 2009). As described by Johnsborg and Havarstein (2009; Johnsborg and Havarstein, 2009), a *ΔlytR* mutant was obtained with a low frequency close to that of the *Δubk* mutant, confirming that *lytR* is also essential in absence of suppressive mutation(s). We constructed a *ΔlytR* strain and observed that the deletion of *lytR* did not select the same suppressive mutation as that found with *ubk* deletion (i.e., in the *asnS* gene) (data not shown). Supported by the ectopic complementation of *Δubk*, this substantiates distinct roles of UbK and LytR in morphogenesis and/or morphology.
FIGURE 7 | Cytoplasmic localization of the different GFP-UbK forms. Phase-contrast images (top row) and GFP images (bottom row) of exponentially growing cells expressing GFP-UbK or GFP-UbK-K36R or GFP-UbK-Y58F or GFP-UbK-Y58E. Bar scale 1 µm. Heat maps represent the localization patterns of GFP-UbK or GFP-UbK-K36R or GFP-UbK-Y58F or GFP-UbK-Y58E. The n values represent the number of cells analyzed.

Based on the absence of *ubk* only in few *Mycoplasma* or *Ureaplasma*, that are devoid of cell-wall, and because the *ubk* homolog *yjeE* is potentially co-transcribed with an amidase gene in *H. influenzae*, it had been proposed that UbK would be implicated in the cell-wall assembly (Teplyakov et al., 2002). Nevertheless, if these arguments constitute clues, they are not direct proofs to assign a role to UbK in cell-wall homeostasis. Our findings that mutations or deletion of *ubk* affect the morphology of *S. pneumoniae* (*Figures 5, 6*) however support this claim and strongly suggest that UbK is required for the cell wall assembly in the pneumococcus.

An important feature is also that Y58 of UbK in not conserved in YdiB of *B. subtilis* (Nguyen et al., 2017). Indeed, the tyrosine 58 is replaced by a phenylalanine (F63). Considering that YdiB autophosphorylates mainly on Ser and Thr (Nguyen et al., 2017), and in view of all the observations discussed above, one can suggest that the mode of regulation of UbKs as well as their cellular functions have evolved independently to operate differently and to potentially serve distinct purposes. To strengthen this point of view, we tried to functionally complement the *ubk* pneumococcal mutant with the WT or F63Y alleles of the *ydiB* gene of *B. subtilis* but none of the two alleles allowed complementing the growth defects due to *ubk* deletion in *S. pneumoniae* (data not shown).

We have also shown that the ability of UbK to hydrolyze ATP prevails over its autophosphorylation since a phosphoablative mutant (*ubk*-Y58F) grows like a WT strain and does not display the morphological defects of the catalytic *ubk*-K36R mutant or the *ubk*-Y58E mutant. In addition, the K36R mutation is epistatic toward the Y58F or Y58E mutations since double mutants grow like the *ubk*-K36R mutant. This allows us to propose a model in which the level of initial phosphorylation enhances and determines the level of ATP hydrolysis activity and subsequently the ability of the cells to grow properly. This model also implies that only a moderate range of ATP hydrolysis by UbK is required by the cell (*Figure 8*). In this model, one can speculate that a phosphatase, not yet identified, could consistently adjust the level of UbK phosphorylation and consequently the enhancement of UbK ATP hydrolysis. To date, the only well-known phosphatase of *S. pneumoniae* susceptible to play such a role is CpsB, the cognate phosphotyrosine-phosphatase of the BY-kinase CpsD (Standish et al., 2013). Nevertheless, as for protein-kinases, one cannot exclude that other types of atypical phosphatase idiosyncratic to bacteria may exist and that a specific phosphatase works together with UbK. Similarly, one cannot exclude the existence of potential phosphorylation substrate(s) of UbK that still remain unknown. However, we never obtained either *in vivo* evidence of UbK-mediated phosphorylation of an endogenous substrate (by probing whole cell lysates with anti-phosphotyrosine antibodies, *Supplementary Figure 1B*), nor *in vitro* UbK-mediated phosphorylation signals for proteins found to be phosphorylated on Tyr in the phosphoproteome (Sun et al., 2010) (data not shown). Altogether, these data suggest that UbK of *S. pneumoniae* could either act as a substrateless-kinase whose autophosphorylation level is crucial for the interaction with its partners and/or as a kinase phosphorylating with low efficiency some endogenous substrates. Whatever the case, UbK autophosphorylation and/or phosphorylation of its potential targets should be finely controlled for normal growth and morphogenesis of *S. pneumoniae*. Further phosphoproteomic studies are needed to gain a better understanding on the level of occupancy of the phosphosite of UbK and its potential substrates.

Until now, *S. pneumoniae* was thought to possess only one Y-kinase (CpsD) and one S/T-kinase (StkP). Our characterization of UbK supports the concept that bacteria could have evolved
other types of protein-kinases that will be first identifiable only biochemically. In the long term, it would therefore come as no surprise if all the phosphoproteins identified by high-throughput phosphoproteomic studies (84 in the phosphoproteome of S. pneumoniae, Sun et al., 2010) are phosphorylated by different types of protein-kinases, potentially forming a complex and interconnected regulatory network indispensable for the biology of the pneumococcal cell.

**MATERIALS AND METHODS**

**Bacterial Strains, Plasmids and Oligonucleotides**

Strains and plasmids, and primers used in this study are listed in Supplementary Material as Supplementary Tables 1, 2 respectively.

**Growth Conditions, Media and Bacterial Transformation**

*Streptococcus pneumoniae* strains were cultivated at 37°C in Todd-Hewitt Yeast (THY) broth (Difco) or in C + Y medium (Martin et al., 1995). For growth curves, cells were previously grown in THY to an OD of 0.5 at 550 nm and aliquoted. From a frozen aliquot, cells were counted and numbered by plating onto THY-agar plates. Cells were then diluted so that 2.10^5 to 10^8 bacteria/mL were inoculated in a fresh THY or C + Y liquid medium at 37°C. *S. pneumoniae* mutants were constructed by transformation of the WT strain derivatives as previously described (Martin et al., 2000) using precompetent cells treated at 37°C during 30 min with the synthetic competence stimulating peptide 1 (CSP1) at the concentration of 100 ng/10^8 cells to induce competence. Cells were then plated on THY-agar supplemented with 3% (vol/vol) defibrinated horse blood and incubated for 120 min at 37°C. Selection of transformants was then performed by adding a THY-agar overlay containing the appropriate antibiotic (streptomycin 200 µg.mL^{-1}, kanamycin 250 µg.mL^{-1}, spectinomycin 100 µg.mL^{-1}) and further incubated for either 16 or 30 h at 37°C. The strains expressing an ectopic copy of *ubk*, *gfp-ubk*, *gfp-ubkK36R* or *gfp-ubkY58F* at the amiF treR locus under the control of the maltose inducible promoter P_M (Guiral et al., 2006) were grown in THY or in C + Y medium containing either glucose or 1% maltose.

When transformation assays were conducted to compare the ability of different strains to transform different DNAs carrying *ubk* deletion, CSP was used to induce competence at a saturating concentration of 100 ng/10^8 cells and a genomic control DNA (either DNA from the ΔphpP-stkP-spcrepsL, or the Δspr1424-kan-rpsL mutant) was used as a reference in order to standardize the transforming ability of the recipient strains.

The *E. coli* XL1-Blue strain was used as a host for cloning. *E. coli* BL21 (DE3) strain was used as a host for overexpression. Luria-Bertani (LB) broth and agar supplemented with appropriate antibiotic (tetracycline 15 µg.mL^{-1}, ampicillin 100 µg.mL^{-1}) were used for routine growth at 37°C. Strains used in this study are listed in Supplementary Table 1.

**Allelic Replacement Mutagenesis**

To construct pneumococcus mutants (gene deletions, *gfp-ubk* fusions or site-directed mutagenesis), we used a two-step procedure based on a bicistronic *kan-rpsL* cassette called Janus (Sung et al., 2001) or a *spc-rpsL* cassette that we constructed on purpose called Janus2. The *spc-rpsL* Janus2 cassette was constructed by amplification of the *aad9* gene from the pR412 plasmid (a gift from J.P. Claverys’ laboratory) and fused to the *rpsL* gene. A strain containing the *kan-rpsL* cassette in its genome was then transformed with the PCR fusion fragment that allowed the substitution of the *kan-rpsL* cassette with the *spc-rpsL* cassette. The *spc-rpsL* cassette was then PCR amplified from the genomic DNA of this strain with the same primers used for the *aad9* gene and cloned into a third-plasmid cloning vector that was inserted into the genome of the pneumococcal strain to be mutated.
primers as for the kan-rpsL cassette and used in the two-step Janus technique. Throughout this study, gene mutagenesis (ubk or spr1397) or fusions with gfp (ubk) were constructed at each native chromosomal locus. The mutated or fusion genes were thus expressed under the control of the native promoter and represented the only source of protein. Description of primers used for the construction of strains is provided in Supplementary Table 2. The full procedure is described in Fleurie et al. (2012). Briefly, the Janus or Janus2 cassette is used to replace the gene of interest. After transformation and selection, this confers resistance to kanamycin or spectinomycin and dominant streptomycin sensitivity (KanR–StrS or SpcR–StrS) in an initial WT rpsL1 genetic background. Then, a DNA fragment flanked on each end by sequences homologous to the upstream and downstream regions of the gene of interest is used to transform and substitute the KanR–StrS or SpcR–StrS in the KanR or SpcR strains. Integration of the fragment is selected in final non-polar markerless mutant strains by streptomycin resistance.

The gene encoding the monomeric GFP was from Martin et al. (2010). For allelic replacement of spr1397, which is an essential gene, by spr1397 D121A, we used a one-step procedure without any selection and direct enzymatic digestion of a PCR fragment amplified from genomic DNA of each selected clone after transformation. Briefly, the introduction of the D121A mutation was concomitant with the introduction of an AluI restriction site in the recombinant PCR fragment covering upstream to downstream regions of spr1397. Characterization of the AluI restriction profile of the PCR fragments covering the spr1397 region of transformants led to identify the AluI restriction site. The presence of the suppressive D121A mutation in these transformants was further confirmed by direct DNA sequencing of the spr1397 locus.

Complementation and/or Introduction of an Ectopic Copy of ubk

To obtain ectopic copies of ubk, gfp-ubk, gfp-ubk K36R or gfp-ubk Y58F at the amiI/treR locus under the control of the maltose inducible promoter P_M (Guiral et al., 2006), we cloned an amplified copy of ubk in the pCEP plasmid that was transformed directly and integrated in the WT strain to generate the ubk+/-gfpCEP strain (Guiral et al., 2006). We did the same with the gfp gene that was integrated into the WT reference strain to generate the gfpCEP strain. From these two strains PCR fusion fragments using primers designed to introduce single mutations were generated. These PCR fragments (gfp-ubk, gfp-ubkK36R or gfp-ubkY58F) contained the upstream and downstream amiI and kan-treR regions of the site of integration of the CEP platform respectively. Transformation of the Δubk mutant with these PCR fragments allowed obtaining the Δubk− gfp-ubkCEP, Δubk− gfp-ubkK36RCEP and Δubk− gfp-ubkY58FCEP strains from which the GFP-UbK fusion proteins were purified. When necessary, the absence of any rearranged copy of ubk in the genome of the transformants was controlled by PCR with the 13/14 primer pair internal to ubk (Supplementary Table 2).

Construction of Plasmids

DNA fragments coding for UbK were obtained by PCR using the chromosomal DNA of our reference WT strain as template. To introduce the different mutations corresponding to UbK K36R, UbK Y58F and UbK Y58E, we used each time a specific primer overlapping the mutated region. When introducing a single mutation, chimera DNA fragments using primers introducing the mutation were amplified. Then, these chimera DNA fragments were used as a primer for a second round of PCR to amplify the whole sequence. The fragments were then cloned between the NdeI and PstI, BamH1 and HindIII, and NdeI and BamH1 restriction sites of the pT7.7 (Cortay et al., 1994), pQE30 and pETphos plasmids respectively. The nucleotide sequences of all DNA fragments were sequenced to ensure error-free amplification. Plasmids and primers used in this study are listed in Supplementary Tables 1, 2 respectively.

Whole Genome Sequencing

For each bacterial strain (WT and two independent ubk mutants), extracted genomic DNA was sheared (1 µg) following the manufacturer instructions using the S220 focused ultrasonicator (Covaris) to obtain a fragment distribution length from 100 bp to 1 kb, with an average peak around 400 bp. Each fragmented DNA material was then used to build a barcoded library using the Ion Xpress Plus gDNA (genomic DNA) Fragment Library Preparation kit (Thermofisher) following the protocol of the kit. Libraries were then size-selected using the E-gel Electrophoresis system (Invitrogen) in order to select fragments from 350 to 450 bp in length. Libraries were qualified according to the concentration and distribution profile using the TapeStation 2200 (Agilent). Diluted libraries (26 pm) were mixed at an equimolar range and were amplified through emulsion PCR using the Ion PGM Template OT2 400 kit (Thermofisher). Finally, the enriched libraries were loaded into a 316v2 chip and sequenced on the Ion PGM sequencer with the Ion PGM HiQ chemistry.

Protein Purification

Recombinant plasmids overproducing the UbK, UbK-K36R, UbK-Y58F, and UbK-Y58E proteins were transformed into BL21 (DE3) E. coli strain. The transformants were grown at 37°C until the culture reached an OD600 = 0.5. Expression was induced with IPTG to a final concentration of 0.5 mM and incubation was continued for 3 h. After 3 h culture at 37°C, cells were harvested and resuspended in buffer A (Tris-HCl 50 mM, pH 7.5; NaCl 300 mM; DTT 1 mM; imidazole 10 mM; glycerol 10%) containing 10 mg.L−1 of lysozyme and 6 mg.L−1 of DNase I and RNase A, and sonicated. After centrifugation at 15000 g for 30 min, the supernatant was loaded onto a Ni-NTA agarose column (Qiagen) and extensively washed with buffer A supplemented with 30 mM imidazole. Samples were eluted with buffer A supplemented with 300 mM imidazole. The fractions corresponding to the pure protein were pooled and dialyzed against the following buffer: HEPES 50 mM, pH 7.5; NaCl 100 mM; DTT 1 mM; MgCl2 1 mM; glycerol 10%. The protein concentrations were determined using a Coomassie Assay Protein Dosage Reagent (Uptima) and aliquots were stored at −80°C.
Cells were incubated in a hypotonic buffer (0.1 M Tris-HCl, 1 mM S. pneumoniae 150 mM NaCl, 0.5 mM EDTA 0.1% Triton X100 buffer and eluted activities are calculated from the Beer-Lambert law using the disappearance of NADH, followed at 340 nm. Specific oxidized to NAD+. One mole of ATP hydrolyzed is directly converted to one mole of NADH by lactate dehydrogenase (LDH) (Jault et al., 1991). One mole of ATP hydrolysis couples the regeneration of ATP from the ADP produced to the conversion of phosphoenolpyruvate (PEP) to pyruvate by pyruvate kinase (PK) and the conversion of pyruvate to lactate by lactate dehydrogenase (LDH) (fault et al., 1991). One mole of ATP hydrolyzed is directly converted to one mole of NADH oxidized to NAD+. The ATP hydrolysis activity is monitored by the disappearance of NADH, followed at 340 nm. Specific activities are calculated from the Beer-Lambert law using the molar extinction coefficient of NADH (6200 mol.L⁻¹.cm⁻¹) and expressed in nmol of ATP.mg⁻¹ of UbK.min⁻¹. Mixtures were prepared as described by Karst et al. (2009).

**Nano LC-MS/MS Analysis of Purified UbK**

Purified UbK was in-gel digested using trypsin as described in Borchert et al. (2010) and peptide mixture was desalted using C18 StageTips (Rappsilber et al., 2007). Analysis of peptides was done on a Proxeon Easy-LC system (Proxeon Biosystems) coupled to an LTQ-Orbitrap-Elite mass spectrometer (Thermo Fisher Scientific) equipped with a nanoelectrospray ion source (Proxeon Biosystems) as described previously (Conzelmann et al., 2013).

Mass spectra were analyzed using the software suite MaxQuant, version 1.3.0.5 (Cox and Mann, 2008). Extracted peak lists were submitted to database search using the Andromeda search engine (Cox et al., 2011) to query target-decoy databases consisting of 35,214 protein entries from *S. pneumoniae* serotype 2 strain D39/NCTC 7466, and 2,029 protein entries from *S. pneumoniae* strain ATCC BAA-255, both obtained from Uniprot and against the sequence of the His-tagged protein UbK and 248 commonly observed contaminants. Trypsin was defined as protease and two missed cleavage sites were allowed. Acetylation at the N-terminus, oxidation of methionine and phosphorylation on serine, threonine, and tyrosine were set as variable modifications. Carbamidomethylation of cysteine was set as fixed modification. Initial precursor mass tolerance was set to six parts per million (ppm) at the precursor ion level and 20 ppm at the fragment ion level. Spectra of modified peptides were manually validated.

**Microscopy Techniques**

GFP fusion of wild-type UbK or mutants were visualized by fluorescence microscopy using *S. pneumoniae* cell culture at OD₅₅₀ nm = 0.1. Slides were visualized with a Nikon TiE microscope fitted with an Orca-CMOS Flash4 V2 camera with a 100 × 1.45 objective. Images were collected using NIS-Elements (Nikon). Images were analyzed using the software ImageJ and the plugin MicrobeJ (Ducret et al., 2016).

For TEM, *S. pneumoniae* cells were grown at 37°C to an OD of 0.1 in THY medium. Samples were then collected, centrifuged and fixed overnight with 5% glutaraldehyde in 0.1 M cacodylate buffer (pH 7.5) at 4°C. Postfixation with 1% osmium tetroxide in cacodylate buffer was carried out for 1 h at room temperature. These fixed cells were dehydrated using a graded series of ethanol and embedded in LR-white at 60°C for 48 h. Ultrathin sections (60 nm) were obtained using a Leica UC7 microtome and were counterstained with uranyl acetate and lead citrate (Reichert Ultrastainer, Leica, Germany). Samples were examined with a Philips CM120 transmission electron microscope equipped with a Gatan Orius SC200 CCD camera.

For SEM, cells were grown at 37°C to an OD of 0.1 in THY medium, spotted onto poly-lysine coverslips, washed with phosphate-buffered saline, and fixed in 0.18 M cacodylate buffer (pH 7.6) containing 2% glutaraldehyde. All samples were then prepared as described by Karst et al. (2009).

1http://rsb.info.nih.gov/ij/
dehydrated with a graded series of ethanol, passage in HMDS (1,1,1,3,3,3-hexamethyldisilazane) and finally air dried. The dried samples were covered with a 10 nm-thick gold/platinum layer (MED20, Baltec). Samples were then observed with a Quanta 250 FEG (FEI) scanning electron microscope.

DATA AVAILABILITY
The genomic data generated can be found in the NCBI Sequence Read Archive (PRJNA562556).

AUTHOR CONTRIBUTIONS
AP, SG, CF, CGa, and JT conducted all the experiments of cell biology, genetics, purified proteins, and performed the ATP hydrolysis assays and western blot analysis. CC performed the transmission electron microscopy images. MF-W and BM analyzed the data together with SG, J-MJ, and CGr. SG and CGr wrote the manuscript. All authors edited the manuscript.

REFERENCES
Akochy, P. M., Bernard, D., Roy, P. H., and Lapointe, J. (2004). Direct glutaminyl-tRNA biosynthesis and indirect asparaginyl-tRNA biosynthesis in Pseudomonas aeruginosa PAO1. J. Bacteriol. 186, 767–776. doi: 10.1128/jb.186.3.767-776.2004
Beilharz, K., Novakova, L., Fadda, D., Branny, P., Massidda, O., and Veening, J. W. (2012). Control of cell division in Streptococcus pneumoniae by the conserved Ser/Thr protein kinase StkP. Proc. Natl. Acad. Sci. U.S.A. 109, E905–E913. doi: 10.1073/pnas.1119172109
Berge, M. J., Mercy, C., Mortier-Barriere, L, Vannieuwenhze, M. S., Brun, Y. V., Grangeasse, C., et al. (2017). A programmed cell division delay preserves genome integrity during natural genetic transformation in Streptococcus pneumoniae. Nat. Commun. 8:1621. doi: 10.1038/s41467-017-01716-9
Bidnenko, V., Shi, L., Kobir, A., Ventroux, M., Pigeonneau, N., Henry, C., et al. (2013). Bacillus subtilis serine/threonine protein kinase YabT is involved in spore development via phosphorylation of a bacterial recombina. Mol. Microbiol. 88, 921–935. doi: 10.1111/mmi.12233
Borchert, N., Dieterich, C., Krug, K., Schutz, W., Jung, S., Nordheim, A., et al. (2013). Interplay of the serine/threonine-kinase StkP and the paralogs DivIV A and GpsB in pneumococcal cell elongation and division. Mol. Microbiol. 89, 1794–1805. doi: 10.1021/pr101068r
Campbell, T. L., Ederer, C. S., Allali-Hassani, A., and Brown, E. D. (2007). Isolation of a novel essential ATPase of E. coli. J. Bacteriol. 189, 3318–3321. doi: 10.1128/jb.01130-07
Canova, M. J., and Molle, V. (2014). Bacterial serine/threonine protein kinases in host-pathogen interactions. J. Biol. Chem. 289, 9473–9479. doi: 10.1074/jbc.R113.529917
Conzelmann, M., Williams, E. A., Krug, K., Franz-Wachtel, M., Macke, B., and Jekely, G. (2013). The neuropeptide complement of the marine annelid Platynereis dumerilii. BMC Genomics 14:906. doi: 10.1186/1471-2164-14-906
Cortay, J. C., Negre, D., Scabrel, M., Rasseiere, T. M., Vartak, N. B., Reizer, J., et al. (1994). In vitro asymmetric binding of the pleiotropic regulatory protein, FruR, to the ace operator controlling glyoxylate shunt enzyme synthesis. J. Biol. Chem. 269, 14885–14891.

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SUPPLEMENTARY MATERIAL
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Cox, J., and Mann, M. (2008). MaxQuant enables high peptide identification rates, individualized p.p.b.-range mass accuracies and proteome-wide protein quantification. Nat. Biotechnol. 26, 1367–1372. doi: 10.1038/nbt.1511
Cox, J., Neuhauer, N., Michalski, A., Scheltema, R. A., Olsen, J. V., and Mann, M. (2011). Andromeda: a peptide search engine integrated into the MaxQuant environment. J. Proteome Res. 10, 1794–1805. doi: 10.1021/pr101065x
Dworkin, J. (2015). Ser/Thr phosphorylation as a regulatory mechanism in bacteria. Curr. Opin. Microbiol. 24, 47–52. doi: 10.1016/j.mib.2015.01.005
Elas, J. E., and Gygi, S. P. (2010). Target-decoy search strategy for mass spectrometry-based proteomics. Methods Mol. Biol. 604, 55–71. doi: 10.1007/978-1-60761-444-9_5
Fleurie, A., Cluzel, C., Guiral, S., Freton, C., Galisson, F., Zanella-Cleon, I., et al. (2012). Mutational dissection of the S/T-kinase StkP reveals crucial roles in cell division of Streptococcus pneumoniae. Mol. Microbiol. 83, 746–758. doi: 10.1111/j.1365-2958.2011.07962.x
Fleurie, A., Lesterlin, C., Manuse, S., Zhao, C., Cluzel, C., Lavergne, J. P., et al. (2014a). MapZ marks the division sites and positions FtsZ rings in Streptococcus pneumoniae. Nature 516, 239–262. doi: 10.1038/nature13966
Fleuvie, A., Manuse, S., Zhao, C., Campo, N., Cluzel, C., Lavergne, J. P., et al. (2014b). Interplay of the serine/threonine-kinase StkP and the paralogs DivIV A and GpsB in pneumococcal cell elongation and division. PLoS Genet. 10:e1004275. doi: 10.1371/journal.pgen.1004275
Freiberg, C., Wieland, B., Spaltmann, F., Ehrt, K., Brotz, H., and Labischinski, H. (2001). Identification of novel essential Escherichia coli genes conserved among pathogenic bacteria. J. Mol. Microbiol. Biotechnol. 3, 483–489.
French, C. T., Lao, P., Loraire, A. E., Matthews, B. T., Yu, H., and Dybvig, K. (2008). Large-scale transposon mutagenesis of Mycoplasma pneumoniae. Mol. Microbiol. 69, 67–76. doi: 10.1111/j.1365-2983.2008.06262.x
Grangeasse, C. (2016). Rewiring the pneumococcal cell cycle with serine/threonine- and tyrosine-kinases. Trends Microbiol. 24, 713–724. doi: 10.1016/j.tim.2016.04.004
Grangeasse, C., Cozzone, A. J., Deutscher, J., and Mijakovic, I. (2007). Tyrosine phosphorylation: an emerging regulatory device of bacterial physiology. Trends Biochem. Sci. 32, 86–94. doi: 10.1016/j.tibs.2006.12.004
Guiral, S., Henard, V., Laaberki, M. H., Granadel, C., Prudhomme, M., Martin, B., et al. (2006). Construction and evaluation of a chromosomal expression
Johnsbo, O., and Havarstein, L. S. (2009). Pneumococcal LytR, a protein

Jault, J. M., Di Pietro, A., Falson, P., and Gautheron, D. C. (1991). Alteration of

Havarstein, L. S., Martin, B., Johnsborg, O., Granadel, C., and Claverys, J. P. (2006).

Karst, J. C., Foucher, A. E., Campbell, T. L., Di Guilmi, A. M., Stroebel, D., Mangat,

Mangat, C. S., and Brown, E. D. (2008). Known bioactive small molecules probe

Lee, J. J., Kan, C. M., Lee, J. H., Park, K. S., Jeon, J. H., and Lee, S. H.

Martin, B., Granadel, C., Campo, N., Henard, V., Prudhomme, M., and Claverys, J. P. (2010). Expression and maintenance of ComD-ComE, the two-component signal-transduction system that controls competence of

Streptococcus pneumoniae. Mol. Microbiol. 75, 1513–1528. doi: 10.1111/j.1365-

Havarstein, L. S., Havarstein, L. S., Martin, B., Johnsbo, O., Granadel, C., and Claverys, J. P. (2006). New insights into the pneumococcal fratricide: relationship to clumping and identification of a novel immunity factor. Mol. Microbiol. 59, 1297–1307.

Henriques, M. X., Rodrigues, T., Carido, M., Ferreira, L., and Filipe, S. R. (2011). Synthesis of capsular polysaccharide at the division septum of Streptococcus pneumoniae is dependent on a bacterial tyrosine kinase. Mol. Microbiol. 82, 515–534. doi: 10.1111/j.1365-2958.2011.07828.x

Henriques-Normark, B., and Normark, S. (2010). Commensal pathogens, with a focus on Streptococcus pneumoniae, and interactions with the human host. Exp. Cell Res. 316, 1408–1414. doi: 10.1016/j.yexcr.2010.03.003

Hunt, A., Rawlins, J. P., Thomaides, H. B., andErrington, J. (2006). Functional analysis of 11 putative essential genes in Bacillus subtilis. Microbiology 152, 2895–2907. doi: 10.1099/mic.0.29152-0

Jadeau, F., Bechet, E., Cozzone, A. J., Deleage, G., Grangeasse, C., and Combet, C. (2008). Identification of the idsynergistic bacterial protein-tyrosine kinase (BY-kinase) family signature. Bioinformatics 21, 2427–2430. doi: 10.1093/bioinformatics/btn662

Jadeau, F., Grangeasse, C., Shi, L., Mijakovic, I., Deleage, G., and Combet, C. (2012). BYKDb: the bacterial protein Tyrosine kinase database. Nucleic Acids Res. 40, D231–D234. doi: 10.1093/nar/gkr915

Jault, J. M., Di Pietro, A., Falson, P., and Gautheron, D. C. (1991). Alteration of apparent negative cooperativity of ATPase activity by alpha-subunit glutamine 173 mutation in yeast mitochondrial F1. Correlation with impaired nucleotide interaction at a regulatory site. J. Biol. Chem. 266, 8073–8078.

Johnsbo, O., and Havarstein, L. S. (2009). Pneumococcal LytR, a protein from the LytR-CpsA-Psr family, is essential for normal septum formation in Streptococcus pneumoniae. J. Bacteriol. 191, 5859–5864. doi: 10.1128/JB.00631-10

Nguyen, H. A., El Khoury, T., Guiral, S., Laaberki, M. H., Candusso, M. P., Galisson, F., et al. (2017). Expanding the kinome world: a new protein kinase family widely conserved in bacteria. J. Mol. Biol. 429, 3056–3074. doi: 10.1016/j.jmb.2017.08.016

Nouriikjan, Y., Kjos, M., Mercy, C., Cluzei, C., Morlot, C., Niorot-Gros, M. F., et al. (2015). Autophosphorylation of the bacterial tyrosine-kinase CpsD connects capsule synthesis with the cell cycle in Streptococcus pneumoniae. PLoS Genet. 11:e1005518. doi: 10.1371/journal.pgen.1005518

Pearlman, S. M., Serber, Z., and Ferrell, J. E. Jr. (2011). A mechanism for the evolution of phosphorylation sites. Cell 147, 934–946. doi: 10.1016/j.cell.2011.08.052

Rappuolff, J., Mann, M., and Ishihama, Y. (2007). Protocol for micro-purification, enrichment, pre-fractionation and storage of peptides for proteomics using StageTips. Nat. Protoc. 2, 1896–1906. doi: 10.1038/nprot.2007.261

Reinstein, J., Schlitching, L., and Wittinghofer, A. (1990). Structurally and catalytically important residues in the phosphate binding loop of adenylate kinase of Escherichia coli. Biochemistry 29, 7451–7459. doi: 10.1021/bi00484a004

Soulat, D., Jault, J.-M., Geourjon, C., Gouet, P., Cozzone, A. J., Grangeasse, C. (2007). Tyrosine-kinase Yac from Escherichia coli possesses an ATPase activity regulated by autophosphorylation. FEMS Microbiol. Lett. 274, 252–259. doi: 10.1111/j.1574-6968.2007.00841.x

Standish, A. J., Salim, A. A., Capon, R. J., and Morona, R. (2013). Dual inhibition of DNA polymerase PolC and protein tyrosine phosphatase CpsB uncovers a novel antibiotic target. Biochem. Biophys. Res. Commun. 430, 167–172. doi: 10.1016/j.bbrc.2012.11.049

Standish, A. J., Whitfield, J. J., and Morona, R. (2014). Tyrosine phosphorylation enhances activity of pneumococcal autolysin LytA. Microbiology 160, 2745–2754. doi: 10.1099/mic.0.08074-0

Sun, X., Gu, F., Xiao, C. L., Yin, X. F., Ge, R., Zhang, L. H., et al. (2010). Phosphoproteomic analysis reveals the multiple roles of phosphorylation in pathogenic bacterium Streptococcus pneumoniae. J. Proteome Res. 9, 275–282. doi: 10.1021/pr900612v

Sung, C. K., Li, H., Claverys, J. P., and Morrison, D. A. (2001). An rpsl cassette, janus, for gene replacement through negative selection in Streptococcus pneumoniae. Appl. Environ. Microbiol. 67, 5190–5196. doi: 10.1128/aem.67.11.5190-5196.2001

Tacke, K. (2017). Global Priority List of Antibiotic-Resistant Bacteria to Guide Research, Discovery and Development of New Antibiotics. WHO Report, 1–7. Geneva: WHO.

Tedpylakov, A., Obmolovga, G., Tordova, M., Thanki, N., Bonander, N., Eisenstein, E., et al. (2002). Crystal structure of the YeiE protein from Haemophilus
Pelletier et al. UbK, a Tyrosine-Autokinase of Streptococcus pneumoniae

**Proteins** 48, 220–226. doi: 10.1002/prot.10114

Thanassi, J. A., Hartman-Neumann, S. L., Dougherty, T. J., Dougherty, B. A., and Pucci, M. J. (2002). Identification of 113 conserved essential genes using a high-throughput gene disruption system in *Streptococcus pneumoniae*. *Nucleic Acids Res.* 30, 3152–3162. doi: 10.1093/nar/gkf418

Ty pas, A., Banzh af, M., Gross, C. A., and Vollmer, W. (2011). From the regulation of peptidoglycan synthesis to bacterial growth and morphology. *Nat. Rev. Microbiol.* 10, 123–136. doi: 10.1038/nrmicro2677

Thanassi, J. A., Hartman-Neumann, S. L., Dougherty, T. J., Dougherty, B. A., and Pucci, M. J. (2002). Identification of 113 conserved essential genes using a high-throughput gene disruption system in *Streptococcus pneumoniae*. *Nucleic Acids Res.* 30, 3152–3162. doi: 10.1093/nar/gkf418

Typas, A., Banzhaf, M., Gross, C. A., and Vollmer, W. (2011). From the regulation of peptidoglycan synthesis to bacterial growth and morphology. *Nat. Rev. Microbiol.* 10, 123–136. doi: 10.1038/nrmicro2677

Vilcheze, C., Molle, V., Carrere-Kremer, S., Leiba, J., Mourey, L., Shenai, S., et al. (2014). Phosphorylation of KasB regulates virulence and acid-fastness in *Mycobacterium tuberculosis*. *PLoS Pathog.* 10:e1004115. doi: 10.1371/journal.ppat.1004115

Vincent, C., Doublet, P., Grangeasse, C., Vaganay, E., Cozzone, A. J., and Duclos, B. (1999). Cells of *Escherichia coli* contain a protein-tyrosine kinase, Wzc, and a phosphotyrosine-protein phosphatase, Wzb. *J. Bacteriol.* 181, 3472–3477.

Wright, D. P., and Ulijasz, A. T. (2014). Regulation of transcription by eukaryotic-like serine-threonine kinases and phosphatases in Gram-positive bacterial pathogens. *Virulence* 5, 863–885. doi: 10.4161/21505594.2014.983404

Yoon, H. S., Lee, M. H., Xiong, J., and Golden, J. W. (2003). Anabaena sp. strain PCC 7120 hetY gene influences heterocyst development. *J. Bacteriol.* 185, 6995–7000. doi: 10.1128/jb.185.23.6995-7000.2003

Zalacain, M., Biswas, S., Ingraham, K. A., Ambrad, J., Bryant, A., Chalker, A. F., et al. (2003). A global approach to identify novel broad-spectrum antibacterial targets among proteins of unknown function. *J. Mol. Microbiol. Biotechnol.* 6, 109–126. doi: 10.1159/000076741

Zucchini, L., Mercy, C., Garcia, P. S., Cluzel, C., Gueguen-Chaignon, V., Galisson, F., et al. (2018). PASTA repeats of the protein kinase StkP interconnect cell constriction and separation of *Streptococcus pneumoniae*. *Nat. Microbiol.* 3, 197–209. doi: 10.1038/s41564-017-0069-3

**Conflict of Interest Statement:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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