

CheX Is a Phosphorylated CheY Phosphatase Essential for *Borrelia burgdorferi* Chemotaxis

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Motility and chemotaxis are believed to be important in the pathogenesis of Lyme disease caused by the spirochete *Borrelia burgdorferi*. Controlling the phosphorylation state of CheY, a response regulator protein, is essential for regulating bacterial chemotaxis and motility. Rapid dephosphorylation of phosphorylated CheY (CheY-P) is crucial for cells to respond to environmental changes. CheY-P dephosphorylation is accomplished by one or more phosphatases in different species, including CheZ, CheC, CheX, FliY, and/or FliY/N. Only a *cheX* phosphatase homolog has been identified in the *B. burgdorferi* genome. However, a role for *cheX* in chemotaxis has not been established in any bacterial species. Inactivating *B. burgdorferi cheX* by inserting a *flgB-kan* cassette resulted in cells (*cheX* mutant cells) with a distinct motility phenotype. While wild-type cells ran, paused (stopped or flexed), and reversed, the *cheX* mutant cells continuously flexed and were not able to run or reverse. Furthermore, swarm plate and capillary tube chemotaxis assays demonstrated that *cheX* mutant cells were deficient in chemotaxis. Wild-type chemotaxis and motility were restored when *cheX* mutant cells were complemented with a shuttle vector expressing CheX. Furthermore, CheX dephosphorylated CheY3-P *in vitro* and eluted as a homodimer in gel filtration chromatography. These findings demonstrated that *B. burgdorferi* CheX is a CheY-P phosphatase that is essential for chemotaxis and motility, which is consistent with CheX being the only CheY-P phosphatase in the *B. burgdorferi* chemotaxis signal transduction pathway.

Bacteria move toward or away from environments that are favorable or unfavorable, respectively, to enhance their survival (reviewed in references 5, 63, and 65). When this movement is in response to chemicals, the process is termed chemotaxis. Flagella or periplasmic flagella, depending upon their location in a cell, are responsible for locomotion in many species of bacteria. Regulation of flagellar rotation and chemotaxis has been studied most extensively in *Escherichia coli* and *Salmonella enterica* serovar Typhimurium, and phosphorylation of the response regulator CheY plays an important role in regulating the swimming pattern of cells (reviewed in references 5, 8, 12, 60, 63, and 65). The concentration of phosphorylated CheY (CheY-P) determines whether a cell runs or tumbles. In the absence of attractants, the concentration of CheY-P is relatively high, and CheY-P diffuses to and binds the flagellar switch protein FliM, switching flagellar rotation from a default counterclockwise (CCW) state to a clockwise (CW) rotation. CW rotation of one or more flagella disrupts flagellar bundles, causing cells to tumble and reorient direction during the next run (37, 64). Although CheY-P autodephosphorylates, *E. coli* CheZ is required for efficient CheY-P dephosphorylation, allowing rapid responses to the environment (53). Thus, functionally reducing CheY-P in null mutants of *cheA* (encoding the protein that transfers phosphate to CheY) or *cheY*

results in cells with constant running phenotypes. In contrast, functionally elevating CheY-P in *cheZ* mutants results in cells that constantly tumble; all of these mutants are also nonchemotactic (1, 6, 43).

Borrelia burgdorferi is the flat-wave, motile spirochete that causes Lyme disease. Chemotaxis and motility are believed to be important in the pathogenesis induced by *B. burgdorferi* (10, 13, 36, 45). Motility of this bacterium is attributed to 7 to 11 periplasmic flagella that are attached near each end of the cell cylinder and overlap in the center of the cell (3, 21, 27, 39, 51). These flagella reside in the periplasmic space between the outer membrane sheath and cell cylinder and are involved in both cell morphology and motility. Inactivating the gene encoding the major periplasmic flagellar protein, FlaB, results in cells that are rod-shaped and nonmotile (39, 51).

Relatively little is known about spirochete chemotaxis. Several defined compounds, as well as undefined substances such as serum, are chemoattractants for *Spirochaeta aurantia*, *B. burgdorferi*, and *Treponema denticola* (2, 26, 29, 34, 57). A membrane potential appears to play a role in the chemotaxis signal transduction pathway of *S. aurantia* (24). Transcriptional and genomic analysis of several spirochete species reveals motility and chemotaxis gene clusters/operons. Many of these genes show sequence similarity to *E. coli* and *S. enterica* serovar Typhimurium motility and chemotaxis genes (15, 16, 44, 54). The *B. burgdorferi* genome contains multiple homologs of several motility and chemotaxis genes, e.g., there are two *cheA*, three *cheY*, three *cheW*, and two *cheB* genes. *B. burgdorferi* lacks a *cheZ* homolog (15), which appears to be restricted to β -

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and γ -proteobacteria (63). Some bacterial species that lack *cheZ* contain multiple copies of CheY, and one or more CheY protein(s) may function as a phosphate sink to sequester phosphate from a CheY that binds flagellar motors (25, 55, 56, 59).

In bacteria lacking *cheZ*, other chemotaxis genes, such as *cheC*, *cheX*, and *fliY* or *fliY/N*, have recently been shown to encode proteins which catalyze dephosphorylation of CheY-P and therefore may have a functional role analogous to that of *cheZ* in *E. coli* (31, 42, 62, 63). Except for *fliY*, which is exclusively found in gram-positive bacteria, these genes are present in *Thermotoga maritima* as well as the vast majority of chemotactic bacteria whose genomes have been sequenced to date. Many of these species are human pathogens (42, 61, 62). *B. burgdorferi* CheX shares approximately 25% amino acid sequence identity with *Bacillus subtilis* CheC, as well as with *T. maritima* CheC and CheX (31). In *B. subtilis*, a *cheC* mutant has decreased flagellar switching frequency (i.e., longer CCW and CW rotations), but the flagellar rotational bias (CCW versus CW) is unaltered (52). In addition, CheC possesses an enzymatic activity that weakly dephosphorylates CheY-P and increases significantly in the presence of CheD (62). Recently, Park et al. (42) determined the structure and biochemical activity of CheC and CheX of *T. maritima*. Both of these proteins possess phosphatase activity towards CheY-P, with CheX having a higher specific activity. The homodimeric state of CheX was postulated to increase its phosphatase activity (42).

Although *T. maritima* CheX has been shown to dephosphorylate CheY-P, the importance of this protein in chemotaxis or motility has not yet been determined in any bacterial species. To investigate the function of CheX in *B. burgdorferi*, we inactivated *cheX* by targeted mutagenesis, and the phenotypes of the mutant, as well as the complemented strains (*cheX*⁺), were analyzed. *cheX* mutant cells exhibited a unique motility phenotype, i.e., they constantly flexed and were unable to translate. In addition, *cheX* mutant cells were nonchemotactic. Biochemical analyses confirmed that *B. burgdorferi* CheX is indeed a phosphatase that efficiently dephosphorylates CheY-P in vitro. Taken together, these results indicate that *cheX* is an essential CheY-P phosphatase in the *B. burgdorferi* chemosensory system.

MATERIALS AND METHODS

Bacterial strains and growth conditions and *cheX* inactivation and complementation. High-passage, avirulent *B. burgdorferi* strain B31A, the nonmotile *flaB* mutant, and their growth conditions have been described previously (39). Targeted inactivation of *cheX* (486 base pairs; Bb0671) was achieved using an *flgB-kan* cassette (7, 39). PCR amplification of *cheX*, construction of the inactivation plasmids, and electroporation of linear *cheX-kan* DNA into competent cells were carried out as described previously (39, 49). Briefly, *cheX* DNA was PCR amplified using primers (5'-3') CheX-F (GGGGAGCTGATTGTTTGG AAG) and CheX-R (CCTTTGCCCTATCTAATGGT) and then ligated into the pGEM-T Easy vector (Promega), yielding pCheX-Easy. *flgB-kan* was similarly PCR amplified as reported previously (39), except the AgeI restriction sequences were replaced with XmaI. The XmaI-restricted *flgB-kan* DNA was then inserted into a unique XmaI site within *cheX* (located 92 base pairs from the *cheX* ATG start codon) of the pCheX-Easy vector, yielding pCheX-kan. Restriction mapping indicated that the direction of transcription of *kan* was the same as that of *cheX*. Approximately 2.8 kb of *cheX-kan* DNA was PCR amplified for electroporation. To complement *cheX:kan*, the coding sequence of *cheX* was PCR amplified with primers (5'-3') CheX/com/F (CATATGAGAATAGATTATAT AGAG) and CheX/com/R (AAGCTTCAAACCCTCTCTTATTG) and ligated into a plasmid (pGEM-T Easy) containing the *B. burgdorferi* *flgB* promoter,

using NdeI and HindIII restriction sites. The *flgB-cheX* DNA was then inserted into shuttle vector pKFSS1 (14) using HindIII restriction digestion, yielding pKFcheX. This shuttle vector and the backbone of pKFSS1 vector, pBSV2, have been successfully used to complement targeted genes in this and other laboratories (14, 67; M. Motaleb, C. Li, and N. Charon, unpublished). Twenty micrograms of the resultant plasmid, pKFcheX, was electroporated into the *cheX:kan* mutant cells. Growth medium contained 350 μ g/ml kanamycin and 80 μ g/ml streptomycin.

Dark-field microscopy. Video sequences were taken with a Basler A600fm digital camera at a resolution of 640 by 480 pixels and a speed of 50 frames per second. Dark-field images at a magnification of 100 \times on the camera sensor were produced with Zeiss optics and a modified Chadwick-Helmuth stroboscope, as previously described (22). Individual frames were selected with MoviePlayer and edited in Photoshop.

Protein preparation and antibody production. To express CheX, the gene encoding the protein sequence was PCR amplified (without the ATG codon) using primers (5'-3') F RcheX (GGATCCAGAATAGATTATATAGAGC) and CheX/com/R (AAGCTTCAAACCCTCTCTTATTG); the amplified DNA was then cloned into the BamHI-HindIII-restricted pQE30 vector (QIAGEN Inc.) and overexpressed with a six-His tag in *E. coli* M15(pREP4) by using 0.1 M isopropyl- β -D-thiogalactoside. The His-tagged protein was purified as recommended by the manufacturer (QIAGEN Inc.) and dialyzed against 25 mM NaCl, 50 mM Tris-HCl, pH 8.5 (enzyme studies) or pH 7.0 (antibody production). Approximately 400 μ g of purified His-CheX was used to immunize rats to produce CheX-specific antiserum (Strategic Biosolutions, Newark, DE). Amino-terminal His-tagged CheY3 and CheA2 were prepared using similar procedures. Rat or rabbit polyclonal antisera against recombinant CheY3 or CheA2, respectively, were produced in a similar manner and are described elsewhere (M. Motaleb, R. Bakker, C. Li, and N. Charon, unpublished data). Antibody specificities were confirmed by Western blotting using their respective targeted mutants constructed in our laboratory (32, 40).

Gel electrophoresis and Western blot analysis. Sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) and Western blotting with an enhanced chemiluminescent detection method (Amersham Pharmacia) were carried out as reported previously (19). The concentration of proteins in cell lysates was determined by a Bio-Rad protein assay kit. Unless otherwise noted, 10 μ g of lysate protein was subjected to SDS-PAGE and Western blotting using specific antibodies. Monoclonal antibodies kindly provided by other investigators included the following: anti-FlaB (H9724) by A. Barbour (University of California, Irvine, CA), anti-FlaA by B. Johnson (Centers for Disease Control and Prevention, Atlanta, GA), and anti-DnaK by J. Benach (SUNY, Stony Brook, NY). Specific reactivity to *B. burgdorferi* FlaA, FlaB, and DnaK has been reported previously (4, 11, 20, 39, 40).

Swarm plate and capillary tube chemotaxis assays. Swarm plate chemotaxis assays were performed as described previously (32, 39). Approximately 10⁶ washed cells in a 5- μ l volume were spotted onto 0.35% (wt/vol) agarose plates containing Barbour-Stoener-Kelly II (BSK-II) medium diluted 1:10 in Dulbecco's phosphate-buffered saline. Capillary tube assays were carried out as reported previously (32), with some modifications (2). Briefly, cells were centrifuged and resuspended in a motility buffer (57) containing 1% (wt/vol) bovine serum albumin and 1% (wt/vol) methylcellulose (400 mesh). Capillary tubes filled with attractant (0.1 M glucosamine) or motility buffer controls were sealed and inserted into microcentrifuge tubes containing resuspended cells. Tubes were incubated for 2 h at 33°C in a humidified chamber, after which solutions were expelled from the capillary tubes. To determine the concentration of *B. burgdorferi* cells in capillary tubes, aliquots of expelled cells were mixed with 10 mM HEPES, pH 7.4, containing cell-permeant nucleic acid fluorescent dye Syto61 (Molecular Probes) and analyzed by flow cytometry. Data were acquired and analyzed with a FACSCalibur using CellQuest Pro software (Becton Dickinson).

CheY-P dephosphorylation assays. Phosphotransfer and dephosphorylation experiments were performed as reported previously (6). Briefly, CheA2 (200 pmol) was autophosphorylated by incubation with TKM buffer (50 mM Tris-HCl, pH 8.5, 50 mM KCl, 5 mM MgCl₂), and 0.3 mM [γ -³²P]ATP (5,000 Ci/mmol, 10 mCi/ml; from MP Biomedicals) in a total volume of 0.1 ml at room temperature for 30 min. Unincorporated [γ -³²P]ATP was removed using Micro Bio-spin 6 chromatography columns (Bio-Rad). Twenty picomoles of autophosphorylated [³²P]CheA2-P was then added to premixed TKM buffer containing 150 pmol of CheY3, with or without CheX (0 to 20 pmol), for 1 min. The reactions were stopped with 4 \times SDS-PAGE sample buffer (50 mM Tris-HCl, pH 6.8, 100 mM dithiothreitol, 2% [wt/vol] SDS, 0.1% [wt/vol] bromophenol blue, 10% [vol/vol] glycerol, and 5% [vol/vol] 2-mercaptoethanol), and proteins were separated by SDS-PAGE (15% acrylamide). The gels were dried and subjected to analysis on a PhosphorImager system (Storm 860; Molecular Dynamics).

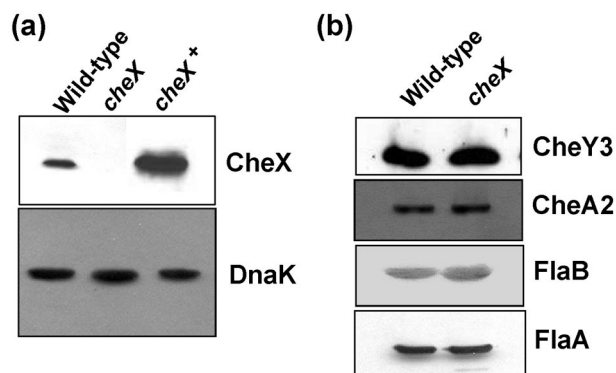


FIG. 1. (a) Western blot analysis of CheX expression in wild-type, *cheX* mutant, and *cheX*⁺ cells. Cell lysates from wild-type, *cheX* mutant, and *cheX*⁺ cells were analyzed by SDS-PAGE and Western blotting with anti-CheX antiserum (top panel). The overexpression of CheX in the *cheX*⁺ strain was likely due to the strong *flgB* promoter and multicopy nature of the shuttle vector used. As an internal control, 2 μ g of each cell lysate was probed with anti-DnaK antibodies (bottom panel). (b) Western blot analysis of CheY3, CheA2, FlaB, and FlaA expression in wild-type and *cheX* mutant cells. Cell lysates (2 μ g for FlaB) from wild-type and *cheX* mutant cells were probed with antibodies specific for CheY3, CheA2, FlaB, and FlaA.

Gel filtration chromatography. Purified His-CheX (225 μ g in 200 μ l 150 mM NaCl, 50 mM Tris, pH 7.4) or protein standards (bovine serum albumin [66 kDa], carbonic anhydrase [29 kDa], and cytochrome *c* [12.4 kDa]; Sigma molecular weight marker kit for gel chromatography) were applied to a Superdex 75 HR 10/30 gel filtration column and eluted at 0.3 ml/min with 150 mM NaCl, 50 mM Tris, pH 7.4, using Pharmacia LCC 501 Plus Controller and P500 pump. Eluted proteins were detected by monitoring absorbance at 280 nm.

RESULTS

Construction and complementation of *cheX* strains. *B. burgdorferi cheX* is located in a motility/chemotaxis operon consisting of *flaA*, *cheA2*, *cheW3*, *cheX*, and *cheY3* genes that is driven by a σ^{70} promoter (17–19, 32, 33). Western blot analysis (see below) indicates that all of the genes in this operon are expressed in growing cells (32, 40). To investigate the function of *cheX*, *cheX* was inactivated by targeted mutagenesis using a *kan* cassette as described in Materials and Methods and reference 39. Western blot analysis confirmed *cheX* inactivation, using antiserum from rats immunized with purified CheX. Figure 1a shows that *B. burgdorferi* CheX antiserum reacted with an 18-kDa protein in wild-type cells, as expected, and that this 18-kDa protein is absent in the *cheX* mutant. PCR analysis confirmed the insertion of *kan* into the targeted gene (data not shown). Together, these results indicate that CheX synthesis is inhibited by inserting *kan* into *cheX*. Since *cheX* is located within an operon, *kan* insertion within *cheX* might have caused an alteration of downstream gene expression, thus complicating interpretation of any phenotype associated with *cheX* mutant cells. However, several lines of evidence indicate that a polar effect due to the inactivation of *cheX* is unlikely. First, Western blot analysis demonstrated that the expression of CheY3, which is immediately downstream of *cheX*, was essentially identical in *cheX* mutant and wild-type cells (Fig. 1b). Second, targeted mutation of *cheY3* results in cells that constantly run (M. Motaleb, R. Bakker, C. Li, and N. Charon, unpublished data), which is markedly different from the swim-

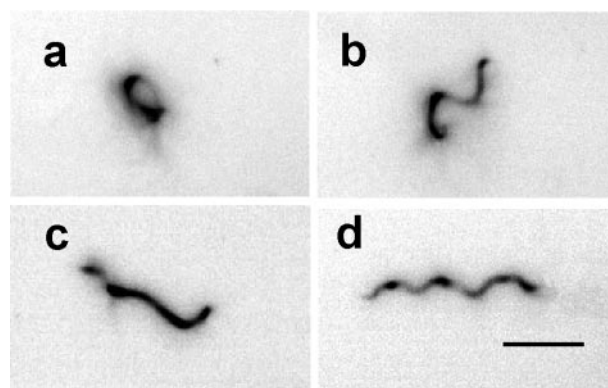


FIG. 2. Dark-field images with contrast reversed of *cheX* mutant and wild-type *B. burgdorferi* cells. (a to c) *cheX* mutant cells illustrating their distorted morphological forms. Cells change their shape from one form to another. (d) Morphological form of wild-type cell during translational motility interval. This form is rarely seen in *cheX* mutant cells. The bar represents 5 μ m.

ming behavior of *cheX* mutant cells (see below). Finally, complementing *cheX* mutant cells (*cheX*⁺) with a shuttle vector expressing CheX restored wild-type swimming (see below), indicating that the phenotype of *cheX* mutant cells was not due to an unexpected secondary alteration. Western blot analysis demonstrated that CheX is expressed in the *cheX*⁺ strain (Fig. 1a).

Altered swimming in *cheX* mutant cells. Because CheX shares amino acid sequence similarity with the recently characterized CheY-P phosphatases CheC of *B. subtilis* and CheX of *T. maritima* (31, 42, 62), it was important to characterize the phenotypes associated with *B. burgdorferi cheX* mutant cells. Video analysis of swimming cells demonstrated that wild-type *B. burgdorferi* ran, paused (stopped or flexed), and reversed; they had a regular wave-like morphology during the interval when they translated in either BSK-II or BSK-II supplemented with 1% methylcellulose (Fig. 2d). The latter compound, which causes the medium to be gel-like, has been shown to promote enhanced translation of *B. burgdorferi* and other spirochetes (10, 23, 30). For *B. burgdorferi*, a “flex” is considered to be when cells bend in the middle or appear distorted in BSK-II medium (23). In contrast to wild-type cells, *cheX* mutant cells had a distorted morphology (Fig. 2a to c). Such cells appeared to be locked into the flexing mode. Although *cheX* mutant cells were motile, as the ends of the cells gyrate, they did not translate in BSK-II with or without methylcellulose. As expected, the wild-type swimming behavior and morphology were restored in *cheX*⁺ cells (data not shown). The possibility that the *cheX* mutation caused altered expression of motility genes, resulting in a constant flexing phenotype, was investigated. Previous studies in this laboratory demonstrated that periplasmic flagella are crucial for *B. burgdorferi* morphology and motility (39). The levels of expression of major and minor flagellins, FlaB and FlaA, were similar in wild-type and *cheX* mutant cells, as shown by Western blotting (Fig. 1b). Furthermore, microscopic examination of *cheX* mutant cells demonstrated that both ends of the cell gyrate at a given time. This observation indicates that both bundles of the periplasmic flagella are rotating (data not shown). Therefore, the constant

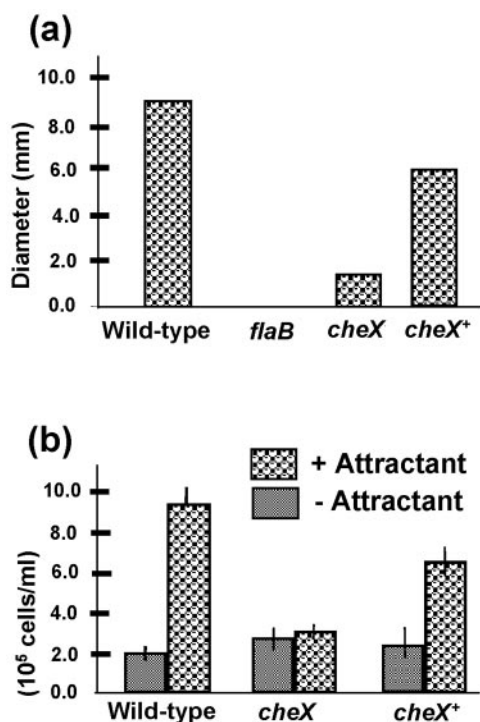


FIG. 3. Chemotaxis assays. (a) Swarm plate assays were performed using BSK-II medium diluted 1:10 with phosphate-buffered saline. Wild-type, *cheX* mutant, and *cheX*⁺ cells were spotted onto agarose plates and incubated for 3 days. The results are expressed as swarm diameter after subtracting the diameter of a nonmotile *flaB* mutant. (b) Capillary tube chemotaxis assay coupled with flow cytometry. Cells from each strain shown were incubated with capillary tubes filled with attractant or no attractant as described in Materials and Methods. After 2 h of incubation, the concentration of cells in each tube was determined using flow cytometry. At least three independent experiments were performed with each strain. Results are expressed as the average number of cells/ml \pm standard deviation.

flexing phenotype in *cheX* mutant cells can be attributed solely to inactivation of *cheX*.

Altered chemotaxis in *cheX* mutant cells. CheC and especially CheX are considered to be functional analogues of *E. coli* CheZ, based on their biochemical properties as CheY-P phosphatases in *B. subtilis* and *T. maritima* (42, 62, 63). If *cheX* is an essential *B. burgdorferi* CheY-P phosphatase, a null mutant in *cheX* should result in cells with a nonchemotactic behavior, such as *E. coli cheZ* mutants (6, 43). To test this hypothesis, swarm plate and capillary tube chemotaxis assays were performed with the wild-type, *cheX* mutant, and *cheX*⁺ cells. The swarm diameter of *cheX* mutant cells was significantly reduced compared to wild-type cells (Fig. 3a). Furthermore, the swarm diameter of *cheX*⁺ cells was similar to that of the wild type, indicating that deficient chemotaxis of *cheX* mutant cells was due to *cheX* inactivation and not an unknown mutation elsewhere. Noticeable growth defects were not observed in *cheX* mutant cells, indicating the decreased swarming phenotype was not due to a growth defect (data not shown). The swarm diameter likely depends on the precise cell run/pause/reverse frequency (50, 66). Therefore, excess CheX synthesized in the *cheX*⁺ cells (Fig. 1a) would likely alter the run/pause/reverse frequency and modestly reduce swarm size,

as seen in Fig. 3a. In addition, capillary tube chemotaxis assays were performed using glucosamine as an attractant. Glucosamine is a chemoattractant for *S. aurantia* (26) and *B. burgdorferi* (2). Whereas the wild-type and *cheX*⁺ strains were chemotactic toward glucosamine, the *cheX* mutant cells failed to respond to this compound (Fig. 3b). These capillary tube results support those from the swarm plate assays and further indicate that *cheX* mutant cells are nonchemotactic.

CheX is a CheY-P phosphatase. Recently, CheC and CheX of *B. subtilis* and *T. maritima* have been shown to dephosphorylate CheY-P in vitro (42, 62, 63). Amino acid sequence alignment indicates that the functional residues of CheC and CheX of *T. maritima* are conserved in *B. burgdorferi* CheX (references 31 and 42 and data not shown). Therefore, the effect of *B. burgdorferi* CheX on CheY-P was determined in vitro. Although the three CheYs of *B. burgdorferi* each possess all of the conserved functional residues of a response regulator (data not shown), initial studies focused on the CheA2-CheY3 pathway for the following reasons: *cheA2* and *cheY3* are located in the same operon as *cheX*, and *cheA2* and *cheY3* are the only sensor kinase and response regulator genes, respectively, that exhibit detectable nonchemotactic phenotypes when inactivated (32; M. Motaleb, R. Bakker, C. Li, and N. Charon, unpublished data). After overexpression and purification of *B. burgdorferi* CheA2 and CheY3, optimal conditions (pH, time course, and KCl concentration) for CheA2 autophosphorylation with [γ -³²P]ATP were determined. As shown in Fig. 4a, CheA2 autophosphorylated (lane 1) and effectively transferred its phosphoryl group to CheY3 (lane 3), confirming that CheA2 is a phosphodonor for CheY3. CheY3 was not phosphorylated by ATP itself (not shown). CheX dephosphorylated [³²P]CheY3-P, in a concentration-dependent manner, when added to a reaction mixture of [³²P]CheA2-P and CheY3 (Fig. 4a, lanes 4 to 6, and b). However, the stability of [³²P]CheA2-P was unaffected by even a sevenfold molar excess of CheX (Fig. 4a, lane 2, and data not shown). These results indicate that CheX of *B. burgdorferi* is a functional CheY-P phosphatase in the *B. burgdorferi* chemotaxis signal transduction pathway.

CheX is a dimer. CheX and CheC share some sequence similarity; however, a major difference between these two proteins is that *T. maritima* CheC is a monomer whereas CheX is a dimer (42). Furthermore, the dimeric state of *T. maritima* CheX was proposed to be important for the stronger phosphatase activity exhibited by CheX (42). The monomer/dimer status of *B. burgdorferi* CheX was investigated by gel filtration chromatography to further validate that this protein is properly classified as a CheX rather than a CheC. The calculated mass of purified CheX is ~19 kDa, including the six-His tag; however, all of the CheX applied to a Superdex column eluted as an ~40-kDa protein (Fig. 5), indicating that *B. burgdorferi* CheX is a homodimer, consistent with classification as a CheX protein.

DISCUSSION

The motility and chemotaxis system of the Lyme disease spirochete *B. burgdorferi* is unique and complex. Motility and chemotaxis are important for many pathogenic organisms to colonize and/or cause disease, including *Vibrio cholerae*, *S. enterica* serovar Typhimurium, *Helicobacter pylori*, *Brachyspira*

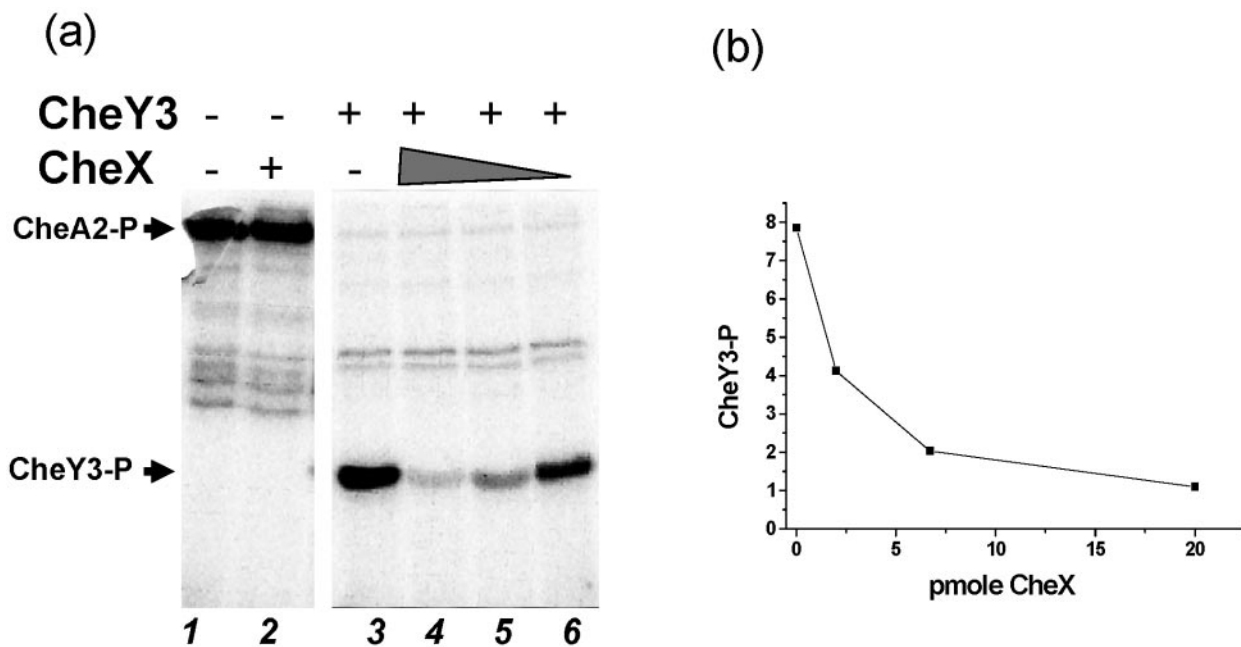


FIG. 4. CheY3-P dephosphorylation assay. (a) Twenty picomoles of autophosphorylated $[^{32}\text{P}]\text{CheA2-P}$ was incubated alone (lane 1), with 20 pmol CheX for 15 min (lane 2), or with 150 pmol CheY3 for 1 min with no CheX (lane 3), 20 pmol CheX (lane 4), 6.3 pmol CheX (lane 5), or 2.1 pmol CheX (lane 6). $[^{32}\text{P}]\text{CheY3-P}$ did not significantly autodephosphorylate during this time period. Arrows indicate the positions of $[^{32}\text{P}]\text{CheA2-P}$ and $[^{32}\text{P}]\text{CheY3-P}$. (b) The intensity of $[^{32}\text{P}]\text{CheY3-P}$ bands as a function of CheX concentration is depicted as relative phosphor-image units.

hyodysenteriae, and possibly *T. denticola* (9, 28, 38, 41, 46). In *T. denticola*, mutation of chemotaxis genes *cheA*, *dmcA*, or *dmcB* produced cells that were unable to penetrate eukaryotic cell monolayers (35, 36). Several studies suggest that motility and chemotaxis are also important for *B. burgdorferi* to penetrate mammalian hosts (36, 47, 48). In addition, during *B. burgdorferi*'s life cycle, it shuttles between mammalian and tick hosts, and it swims in viscous gel-like media where most other bac-

teria fail to swim (10, 30). Multiple variants of chemotaxis and motility proteins may be necessary for these spirochetes to transit to the different hosts and adapt to these very different environments (10, 15, 32). The goal of the present investigation was to determine the extent to which *cheX* is important for *B. burgdorferi* chemotaxis and motility and to begin to characterize the properties of this protein.

CheX represents a recently identified family of protein phosphatases involved in catalyzing the dephosphorylation of CheY-P (42, 62, 63), a reaction essential for rapid chemotactic responses (53). While CheZ is a CheY-P phosphatase in *E. coli* and other β - and γ -proteobacteria (63), in other species CheY-P dephosphorylation is mediated by CheC, CheD, FliY, and/or CheX (42, 62, 63). *B. burgdorferi* CheX shares amino acid sequence similarity to CheX, CheC, and FliY, which have recently been reported to be CheY-P phosphatases of *B. subtilis* and *T. maritima* (42, 62). However, *fliY* and *cheC* were not identified in the *B. burgdorferi* genome (see below). While other spirochetes, e.g., *T. denticola*, *Treponema pallidum*, and *Leptospira interrogans*, contain homologs of *fliY* in addition to *cheX* or *cheC* (16, 44, 54), *B. burgdorferi* is the only spirochete whose genome sequence has been published that does not contain a *fliY* homolog (15). In fact, the only known potential CheY-P phosphatase in the *B. burgdorferi* genome is CheX, suggesting that this protein plays a critical role in *B. burgdorferi* chemotaxis.

Inactivating *cheX* (Fig. 1a) produced *B. burgdorferi* cells that constantly flexed (Fig. 2a to c) and were nonchemotactic (Fig. 3), demonstrating an essential role of CheX in *B. burgdorferi*'s chemosensory pathway. Neither the constantly flexing phenotype nor the loss of chemotaxis in *cheX* mutant cells appeared

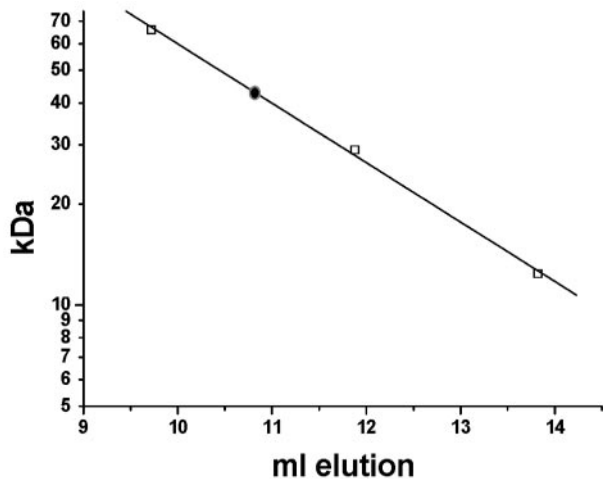


FIG. 5. CheX chromatography on Superdex 75. The elution positions of protein standards bovine serum albumin (66 kDa), carbonic anhydrase (29 kDa), and cytochrome *c* (12.4 kDa) from a Superdex 75 column are depicted as open squares; the position of CheX elution is indicated by a filled circle.

to be attributed to polar effects or secondary mutations: first, no appreciable alteration in levels of expression of the motility proteins FlaB or FlaA or the chemotaxis proteins CheY3 or CheA2 was observed (Fig. 1) (*cheY3* is immediately downstream of *cheX*, and *flaA* and *cheA2* are upstream of *cheX*); second, both flagellar bundles appear to be rotating; and finally, the wild-type swimming behavior and chemotactic ability were restored in *cheX*⁺ complemented cells. In *B. burgdorferi*, *cheX* inactivation likely removes the only CheY-P phosphatase, resulting in highly elevated CheY-P and disruption of the chemosensory signal transduction pathway. Most bacteria that do not contain a *cheZ* gene have multiple presumptive CheY-P phosphatases, and inactivating one of those phosphatases may have less dramatic effects than those observed in *B. burgdorferi* *cheX* mutant cells. For example, in *B. subtilis*, inactivating either of the two CheY-P phosphatase genes, *fliY* or *cheC*, altered flagellar rotation, indicating increased levels of CheY-P. The *fliY* mutant exhibited a stronger phenotype than the *cheC* mutant, and a *cheC fliY* double mutant had the strongest phenotype, consistent with the highest levels of CheY-P (62). Recently, Sim et al. (58) reported that *T. denticola* CheX interacted with itself and CheA in a yeast two-hybrid system. They also postulated that *cheX* is likely to be a phosphatase in *T. denticola* (58); however, the effects of inactivating *cheX* in *T. denticola* have not yet been reported. All these observations are consistent with the hypothesis that *B. burgdorferi* relies solely on CheX for CheY-P dephosphorylation. Furthermore, a double mutant of *cheY3* and *cheX* would be expected to have a constantly running phenotype (as a single *cheY3* mutant) instead of a constantly flexing phenotype (such as the *cheX* mutant cells). Consistent with this prediction, a double mutation that inactivated both *cheX* and *cheY3* did exhibit a constantly running phenotype (M. Motaleb and N. Charon, unpublished).

In vitro phosphorylation studies confirmed that CheX stimulates dephosphorylation of *B. burgdorferi* CheY3-P (Fig. 4), and gel filtration studies indicate *B. burgdorferi* CheX is a homodimer (Fig. 5). Park et al. (42) proposed to differentiate CheX and CheC, based in part on CheX being smaller than CheC and on the presence of a conserved Gly residue in CheX that is involved in forming a β -sheet important for CheX dimerization. Based on these criteria, *B. burgdorferi* CheX is correctly classified as a CheX rather than a CheC. While results of the CheA2-CheY3-CheX pathway have been presented herein (Fig. 4), additional studies suggest that CheX also dephosphorylates CheY1-P in the CheA1-CheY1-CheX pathway (M. Motaleb, C. Li, N. Charon, and M. Miller, unpublished data). Recent reports indicate that the mechanism of *T. maritima* CheX-mediated dephosphorylation of CheY-P is different from that of *E. coli* CheZ (42, 68). *B. burgdorferi* and *T. maritima* CheX share ~25% amino acid sequence identity but no sequence similarity to CheZ. The ability of *T. maritima* CheX to form homodimers was proposed to be one factor in the higher phosphatase activity of CheX relative to CheC (42). *B. burgdorferi* CheX is also a homodimer. Taken together, this information suggests *B. burgdorferi* and *T. maritima* CheX may dephosphorylate CheY-P by similar mechanisms. Determining the structure of a *B. burgdorferi* CheX/CheY3-P complex should provide more definitive information about the mecha-

nism of *B. burgdorferi* CheX-mediated CheY-P dephosphorylation.

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