

## Polar Clustering of the Chemoreceptor Complex in *Escherichia coli* Occurs in the Absence of Complete CheA Function

J. M. SKIDMORE,<sup>1</sup> D. D. ELLEFSON,<sup>2†</sup> B. P. MCNAMARA,<sup>2‡</sup> M. M. P. COUTO,<sup>1</sup>  
A. J. WOLFE,<sup>2</sup> AND J. R. MADDOCK<sup>1\*</sup>

Department of Biology, University of Michigan, Ann Arbor, Michigan 48109-1048,<sup>1</sup> and Department of Microbiology and Immunology, Loyola University Chicago, Maywood, Illinois 60153<sup>2</sup>

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**Bacterial chemotaxis requires a phosphorelay system initiated by the interaction of a ligand with its chemoreceptor and culminating in a change in the directional bias of flagellar rotation. Chemoreceptor-CheA-CheW ternary complexes mediate transduction of the chemotactic signal. In vivo, these complexes cluster predominantly in large groups at the cell poles. The function of chemoreceptor clustering is currently unknown. To gain insight into the relationship between signaling and chemoreceptor clustering, we examined these properties in several *Escherichia coli* mutant strains that produce CheA variants altered in their ability to mediate chemotaxis, autophosphorylate, or bind ATP. We show here that polar clustering of chemoreceptor complexes does not require functional CheA protein, although maximal clustering occurred only in chemotactically competent cells. Surprisingly, in cells containing a minimum of 13 gold particles at the cell pole, a significant level of clustering was observed in the absence of CheA, demonstrating that CheA is not absolutely essential for chemoreceptor clustering. Nonchemotactic cells expressing only CheA<sub>S</sub>, a C-terminal CheA deletion, or CheA bearing a mutation in the ATP-binding site mediated slightly less than maximal chemoreceptor clustering. Cells expressing only full-length CheA (CheA<sub>L</sub>) from either a chromosomal or a plasmid-encoded allele displayed a methyl-accepting chemotaxis protein localization pattern indistinguishable from that of strains carrying both CheA<sub>L</sub> and CheA<sub>S</sub>, demonstrating that CheA<sub>L</sub> alone can mediate polar clustering.**

Bacterial cells sense chemical gradients and modify their swimming behavior accordingly. This behavior, called chemotaxis, depends upon the ability of membrane-bound chemoreceptors (called methyl-accepting chemotaxis proteins [MCPs] or transducers) to communicate with the switch components of flagellar motors to modulate swimming behavior in response to the chemical environment of the cells. In *Escherichia coli*, this communication requires the cooperative effort of the cytoplasmic protein products of six signal transduction genes, *cheA*, *cheW*, *cheR*, *cheB*, *cheY*, and *cheZ* (reviewed in reference 20).

MCPs and the cytoplasmic signaling proteins CheA and CheW interact in a chemosensory ternary complex (5, 6, 10, 25) that, in vitro, forms higher-order structures (17). In vivo, groups of these complexes cluster predominantly at the cell poles (18, 19). Since polar clustering of each protein component requires the presence of the other two (19), this aggregation presumably requires the formation of the ternary complex. Although the methyltransferase (CheR) or methyl-esterase (CheB) interacts with the ternary complex, their activities are not required for clustering (18). Since receptor complexes also form in *Caulobacter crescentus* (1) and *Rhodobacter sphaeroides* (11), clustering of the ternary complexes is thought to play an essential role in chemotaxis signaling, possibly by facilitating signal amplification (8, 19). Although many wild-type cells contain such clusters at only one

of the cell poles, no correlation exists between the location of the cluster and the direction of swimming (3).

The CheA dimer plays a central role in relaying the chemotactic signal from the membrane-bound MCPs to the flagellar switch (reviewed in reference 20). Enteric bacteria synthesize two forms of this histidine kinase, CheA<sub>L</sub> (78 kDa) and CheA<sub>S</sub> (69 kDa) (23), that are translated in frame from two different initiation sites [start(L) and start(S), respectively] (14, 33). Both CheA variants are organized into distinct functional domains (reviewed in reference 29). The N-terminal P1 domain, present in CheA<sub>L</sub> but not in CheA<sub>S</sub>, contains the site of autophosphorylation (His 48) (12). This phosphate is then transferred either to CheY to enhance clockwise signal generation or to CheB to facilitate adaptation (13). The CheA P2 domain assists in the interaction between the phosphodonor site in P1 and CheY (12, 24). The C-terminal domain, MC, appears to play an important role in receiving sensory information from the MCPs (4, 7, 29). Finally, the centrally located transmitter (T) domain contains four highly conserved regions (N, G1, F, and G2) that play a role in the binding and hydrolysis of ATP (4, 29, 34).

Whereas CheA<sub>L</sub> supports chemotaxis in the absence of CheA<sub>S</sub> (30), CheA<sub>S</sub> cannot support chemotaxis on its own. Although CheA<sub>S</sub> can act as a kinase in *trans* (41), it lacks the N-terminal 97 amino acids that include the site of autophosphorylation (12). Despite this, however, most if not all motile enteric bacteria coexpress CheA<sub>L</sub>, CheA<sub>S</sub> and CheZ (23). CheZ interacts directly with CheA<sub>S</sub>, and this interaction enhances the ability of CheZ to aid in dephosphorylating phospho-CheY (21, 22, 37, 38). Thus, it seems likely that CheA<sub>S</sub> plays some important role in chemotaxis distinct from that of CheA<sub>L</sub>.

In this study, we investigated the ability of wild-type and mutant CheA variants to mediate chemoreceptor aggregation in *E. coli*. Here we show that (i) some polar clustering of the

\* Corresponding author. Mailing address: Department of Biology, University of Michigan, 830 North University, Ann Arbor, MI 48109-1048. Phone: (734) 936-8068. Fax: (734) 647-0884. E-mail: maddock@umich.edu.

† Present address: Department of Molecular Microbiology and Immunology, Oregon Health Sciences University, Portland, OR 97201.

‡ Present address: Infectious Diseases, University of Maryland, Baltimore, MD 21201.

TABLE 1. Bacterial strains used in this study

Strain	Relevant genotype	Source or reference
AJW430	<i>cheAK616</i> (Am) <i>recA::cml</i>	41
AJW484	CP366 $\Delta$ <i>cheA::Km</i>	This study
AJW530	AJW430 + pAR1. <i>cheA<sub>S</sub></i>	This study
AJW536	AJW484 <i>cheA<sub>RV</sub>M98L</i> (S <sup>-</sup> ) <i>polA</i> <sup>+</sup> <i>rha</i> <sup>+</sup>	This study
AJW688	AJW1071 + pAR1. <i>cheA<sub>S</sub></i>	This study
AJW689	AJW1071 + pAR1. <i>cheA</i>	This study
AJW768	AJW1071 + pAR1. <i>cheA<sub>RV</sub></i>	This study
AJW774	AJW1071 + pAR1. <i>cheA<sub>RV</sub>98ML</i> (S <sup>+</sup> )	This study
AJW776	AJW1071 + pAR1. <i>cheA<sub>RV</sub>98ML</i> (S <sup>-</sup> )	This study
AJW916	AJW536 + pAR1	This study
AJW917	AJW536 + pAR1. <i>cheA<sub>S</sub></i>	This study
AJW970	AJW1071 + pAR1. <i>cheAG422A</i>	This study
AJW1071	$\Delta$ <i>cheA1643 recA::cml</i>	41
CP366	<i>zig::Tn10 polA</i> (Ts) <i>rha thr</i> (Am)1 <i>leuB6 his-4 metF</i> (Am)159 <i>eda-50 rpsL136</i>	27
KO607	<i>tsr</i> $\Delta$ 7021 ( <i>tar-tap</i> ) $\Delta$ 5201 <i>trg</i> $\Delta$ 100	26
RP437	<i>polA</i> <sup>+</sup> <i>rha</i> <sup>+</sup> <i>thr</i> (Am)1 <i>leuB6 his-4 metF</i> (Am)159 <i>eda-50 rpsL136</i>	28

chemoreceptors occurs in the absence of CheA in cell sections containing sufficient immunogold signal; (ii) CheA<sub>L</sub>, in the absence of CheA<sub>S</sub>, mediates optimal chemoreceptor polarity and clustering; (iii) CheA<sub>S</sub>, in the absence of CheA<sub>L</sub>, supports significant polarity and clustering, although at slightly lower levels than those mediated by CheA<sub>L</sub> and CheA<sub>S</sub> together; and (iv) CheA variants unable to support chemotaxis in vivo or to bind ATP or autophosphorylate in vitro still retain the ability to mediate MCP polarity and clustering. Thus, CheA need not possess all of its domains or all of its functions to promote efficient MCP polarity and clustering.

## MATERIALS AND METHODS

**Bacterial strains, *cheA* alleles, and plasmids.** All strains used in this study are derivatives of *E. coli* K-12 and are listed in Table 1. Strain AJW484 ( $\Delta$ *cheA::Km*), used as a recipient for allele replacements, was constructed as follows. The *cheA* gene on a 2.1-kb *Bam*HI fragment from plasmid pAR1.*cheA* (40) was subcloned into the *Bam*HI site of pUC19. A 1-kb *Eco*RV-*Nru*I fragment was excised from this *cheA* gene and replaced with the *Hinc*II fragment from pUC4K (Pharmacia Biotech, Piscataway, N.J.), which confers kanamycin resistance. The resultant allele, *cheA::Km*, was introduced into the chromosome by homologous recombination in the *polA*(Ts) strain CP366 (27). One recombinant was selected on the basis of its inability to perform chemotaxis in a swarm assay. It was subsequently demonstrated by Southern hybridization to lack the appropriate *cheA* fragment and possess the kanamycin cassette.

Alleles *cheA<sub>RV</sub>*, *cheA<sub>RV</sub>M98L*(S<sup>+</sup>), and *cheA<sub>RV</sub>M98L*(S<sup>-</sup>) express both wild-type CheA<sub>L</sub> and wild-type CheA<sub>S</sub>, both CheA<sub>L</sub>M98L and wild-type CheA<sub>S</sub>, and only CheA<sub>L</sub>M98L, respectively (see Fig. 1). All three alleles carry a translationally silent change in their nucleotide sequence that introduces an *Eco*RV restriction site between the Shine-Dalgarno sequence and the AUG of start(S) that was used to track these alleles during various in vitro and genetic manipulations. Alleles *cheA<sub>RV</sub>M98L*(S<sup>+</sup>) and *cheA<sub>RV</sub>M98L*(S<sup>-</sup>) were constructed by changing the AUG codon of start(S) to UUG and CUC, respectively. All mutations were generated using standard oligonucleotide-directed mutagenesis procedures (15) and were confirmed by dideoxy-chain termination sequencing (31).

Allele *cheA<sub>RV</sub>M98L*(S<sup>-</sup>) was introduced into the chromosome by homologous recombination in the allele replacement strain AJW484 (*cheA*) to produce AJW536. Because *E. coli* cells that express CheA<sub>L</sub> but not CheA<sub>S</sub> perform chemotaxis in motility assays (30), we used this assay to screen for chemotactic recombinants. To avoid phenotypic complications that might arise from the presence of the temperature-sensitive *PolA* protein, we used the generalized transducing phage P1kc (32) to cotransduce the linked *zig::Tn10 polA12*(Ts) *rha* markers to their respective wild-type alleles, using the chemotaxis wild-type strain RP437 (28) as the source of donor DNA. Transductants were selected on the basis of their ability to use rhamnose as a sole carbon source, their sensitivity to tetracycline, and their ability to maintain a ColE1-derived plasmid at 42°C, a phenotype indicative of the wild-type *polA* allele. The chromosome-encoded *cheA* allele was verified by both direct-cycle sequencing (16) across the start(S) region and immunoblot analysis.

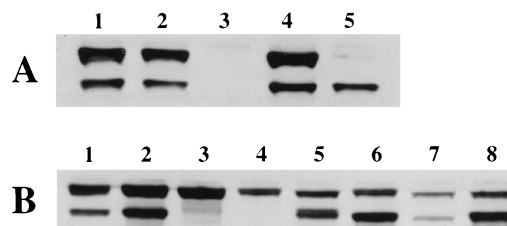


FIG. 1. Immunoblot analysis of strains that carry various *cheA* alleles. Cells were grown in TB at 30°C to an optical density at 600 nm of approximately 0.7. CheA levels were detected using anti-CheA antibodies. (A) Lanes: 1, RP437 (*cheA*<sup>+</sup>); 2, KO607 (*cheA*<sup>+</sup>  $\Delta$ MCP); 3, AJW1071 (*cheA*); 4, AJW689 ( $\Delta$ *cheA* pAR1.*cheA*, uninduced); 5, AJW688 (*cheA* pAR1.*cheA<sub>S</sub>*, 50  $\mu$ M IPTG). (B) Lanes: 1, RP437; 2, AJW768 ( $\Delta$ *cheA* pAR1.*cheA<sub>RV</sub>*, uninduced); 3, AJW776 [*cheA* pAR1.*cheA<sub>RV</sub>M98L*(S<sup>-</sup>), uninduced]; 4, AJW916 [*cheA<sub>RV</sub>M98L*(S<sup>-</sup>), pAR1, 10  $\mu$ M IPTG]; 5, AJW774 [*cheA* pAR1.*cheA<sub>RV</sub>M98L*(S<sup>+</sup>), 10  $\mu$ M IPTG]; 6, AJW917 [*cheA<sub>RV</sub>M98L*(S<sup>-</sup>) pAR1.*cheA<sub>S</sub>*, 10  $\mu$ M IPTG]; 7, AJW430 [*cheAK616*(Am)]; 8, AJW530 [*cheAK616*(Am) pAR1.*cheA<sub>S</sub>*, 10  $\mu$ M IPTG].

Plasmids designed to express various forms of CheA by means of the isopropyl- $\beta$ -D-thiogalactopyranoside (IPTG)-inducible promoter were generated as described previously (41).

**Media and growth conditions.** Cells were grown in tryptone broth (TB) (1% [wt/vol] tryptone, 0.5% [wt/vol] sodium chloride) or Luria broth (LB, 0.5% [wt/vol] yeast extract) at 30°C with aeration. Cell density was monitored spectrophotometrically at 600 nm. Transcription of plasmid-borne *cheA* variants was induced at 10  $\mu$ M IPTG unless otherwise indicated.

**Motility assays.** Aliquots (5  $\mu$ l) of mid-log-phase cells grown in TB supplemented with 50  $\mu$ g of ampicillin per ml and various concentrations of IPTG were spotted onto motility plates (TB, 0.3% agar) in a 30°C humidity chamber, as described previously (39). The diameters of four swarms after 7 h of growth were measured for each strain.

**Immunoelectron microscopy.** A 1/50 dilution of an overnight culture was grown at 30°C for 4 h prior to induction with IPTG. Cells were induced for 1 h at IPTG concentrations that result in about wild-type levels of CheA expression as assayed by immunoblot analysis (11). Cells were fixed and embedded as described previously (11, 18, 19). The antibody was preadsorbed on ice for 15 min with acetone powders prepared from an *E. coli* strain lacking the four major chemoreceptors (KO607) (26). The primary antibody (anti-Tsr) (2) was diluted 1:500 in phosphate-buffered saline-Tween (PBST) plus 2% bovine serum albumin and grids incubated for 1 h in a humidity chamber. The grids were washed three times in PBST and incubated with a 1:30 dilution (in PBST plus bovine serum albumin) of goat anti-rabbit immunoglobulin G coupled with 12-nm-diameter colloidal gold particles (Jackson ImmunoResearch). After being washed in water, the grids were poststained with 1% uranyl acetate.

The positions of colloidal gold particles on longitudinal cell sections were quantified on a Philips CM10 electron microscope at 60kV, as described previously (19). All antibody reactions were performed simultaneously for any given set of data. Cultures were prepared for immunoelectron microscopy, and the localization of MCPs was determined at least twice. Samples were also examined by two different investigators to ensure that the scoring of gold particles was independent of both investigator and sample preparation. Chi-square analysis was performed to analyze differences between data sets. Comparisons reported as either equivalent or different met a probability of  $P \leq 0.05$ .

## RESULTS

The induction of CheA protein from the strains carrying related plasmids used in this study varied significantly. To eliminate variations in CheA abundance that might bias the data interpretation, we first assayed the CheA protein levels produced at different IPTG induction levels by immunoblot analysis. Cells that displayed approximately wild-type levels of CheA protein (Fig. 1) were embedded. The slight variation in the MCP level (as revealed by the relative number of gold particles [see Tables 2 and 3]) also was observed by immunoblot analysis (data not shown).

**MCPs can cluster independently of CheA.** We reported previously that the level of polarity and clustering of MCPs was significantly reduced in the absence of CheA relative to that observed in wild-type cells (from 60 to 80% and from 21 to 81%, respectively) (19). Removal of CheW further diminished

TABLE 2. Spatial distribution of chemoreceptors<sup>a</sup>

Strain	Plasmid-encoded CheA	Total no. of particles	No. of particles in cytoplasm	Membrane particles <sup>b</sup>				No. of membrane particles/section	No. of polar clusters	Size of polar clusters <sup>d</sup>	No. of lateral clusters	Size of lateral clusters <sup>d</sup>
				Polar particles <sup>c</sup>		Lateral particles						
				%	% in clusters	%	% in clusters					
AJW689	CheA <sub>L</sub> and CheA <sub>S</sub>	2,158	104	95	85	5	24	10.3	131	12.7 ± 0.9	5	4.8 ± 0.4
AJW688	CheA <sub>S</sub>	1,483	143	85	68	15	21	6.7	76	10.2 ± 0.6	7	5.7 ± 0.6
AJW1071	None	1,957	137	74	45	26	3	9.1	86	7.0 ± 0.3	3	4.3 ± 0.3

<sup>a</sup> The numbers and positions of gold particles detected by immunomicroscopy from 200 thin sections of cells are shown. The numbers and sizes of gold particle clusters located in either the polar or lateral membrane were determined. The relative size of the clusters ± 1 standard error of the mean is presented. CheA expression was uninduced in strains AJW689 and AJW1071 and induced with 50 μM IPTG in strain AJW688.

<sup>b</sup> The percentages of gold particles at the poles or along the lateral edges and the percentage of polar or lateral gold particles that are clustered are shown.

<sup>c</sup> All pairwise combinations are significantly different ( $P \leq 0.05$ ).

<sup>d</sup> Numbers of particles per cluster.

polar clustering (50% polar gold particles, of which 13% were clustered). To further investigate the requirement for CheA in MCP clustering, we reexamined the localization of the MCPs in AJW1071, a *cheA* deletion strain, and in an AJW1071 transformant that expresses CheA<sub>L</sub> and CheA<sub>S</sub> from a plasmid (strain AJW689). In cells that synthesized both CheA<sub>L</sub> and CheA<sub>S</sub>, the majority of the membrane-associated gold particles clustered at the poles, as observed previously with *E. coli* cells that synthesized CheA<sub>L</sub> and CheA<sub>S</sub> from a chromosomal copy of the wild-type *cheA* allele (strain RP437) (18, 19). The percentage of particles localized to the poles was approximately 95%, and the percentage of those particles in clusters was approximately 85%. Smaller, lateral clusters of gold particles also occurred. In the absence of CheA, the number of polar gold particles decreased to 74% and the clustering percentage reduced to 45% (Table 2). In addition, the average size of the polar clusters decreased from 12.7 gold particles in cells that expressed CheA to 7 gold particles in those that did not (Table 2).

We next examined only cell sections that contained a sufficient number of polar gold particles to generate a potential cluster (by our definition, at least four gold particles) to carefully assess whether the reduction in MCP clustering observed in  $\Delta cheA$  cells was biased by the smaller total number of gold particles observed in these cells. For cells that expressed both CheA<sub>L</sub> and CheA<sub>S</sub> from a plasmid (strain AJW689), 65% of the cell poles that contained four to six gold particles displayed polar clusters and more than 90% of the poles that contained at least seven particles exhibited one or more clusters (Fig. 2). The mean size of the clusters increased as the number of gold particles per pole increased. Cells that expressed both forms of CheA from a chromosomal copy of *cheA* (strain RP437) yielded similar results (data not shown). In contrast, for cells that expressed neither form of CheA (strain AJW1071), the percentage of gold particles in clusters was significantly reduced, even when those poles contained  $\geq 13$  gold particles. Again, the mean cluster size increased as the total number of particles increased; however, many of these poles contained multiple small clusters, resulting in a significantly reduced mean cluster size (8.7 gold particles for strain AJW1071 versus 14.1 for AJW689). These data demonstrate that high levels of clustering occur only in the presence of CheA, even in cells that contain sufficient gold particles at one pole to generate a cluster.

**CheA<sub>S</sub> alone can mediate polar clustering of the MCPs.** To determine whether CheA<sub>S</sub> mediates clustering of chemoreceptor complexes, we examined the immunolocalization patterns of the membrane-bound MCPs in a  $\Delta cheA$  strain that ex-

pressed from a plasmid only CheA<sub>S</sub> (AJW688). These cells mediated a level of chemoreceptor clustering intermediate between those of wild-type and  $\Delta cheA$  cells (Table 2). At wild-type CheA<sub>S</sub> protein levels (induction with 50 μM IPTG [Fig. 1]), the vast majority of the gold particles (85%) localized to the cell poles, although the percentage aggregated into clusters (68%) and the average size of the clusters (10 particles) were somewhat reduced compared to wild-type levels (Table 2). The number of clusters in cell poles containing more than four gold particles was also intermediate between those of cells containing no CheA and those of cells containing both CheA<sub>L</sub> and CheA<sub>S</sub> (Fig. 2). Thus, the nonphosphorylatable CheA<sub>S</sub> protein, which cannot support chemotaxis in vivo, enhances polar localization and polar clustering of MCPs relative to cells without any CheA protein.

**CheA<sub>L</sub> can mediate polar clustering of the MCPs.** Determination of whether CheA<sub>L</sub> alone can mediate polar clustering required a pair of strains that differed only in their ability to synthesize CheA<sub>S</sub>. Because cells translate CheA<sub>L</sub> and CheA<sub>S</sub> in frame, the AUG that encodes start(S) also encodes the amino acid Met 98 within the sequence of CheA<sub>L</sub>. Alleles *cheA<sub>RV</sub>M98L(S+)* (strain AJW776) and *cheA<sub>RV</sub>M98L(S-)* (strain AJW774) were constructed by changing the AUG (Met) codon of start(S) to UUG (Leu) and CUC (Leu), respectively. In *E. coli*, both codons are used with approximately the same frequency; however, the UUG codon can initiate translation of CheA<sub>S</sub> whereas the codon CUC cannot.

When uninduced, cells carrying the plasmid-borne wild-type *cheA<sub>RV</sub>* allele (strain AJW768) synthesized CheA<sub>L</sub> and CheA<sub>S</sub> at levels similar to those produced by isogenic cells carrying the plasmid-borne wild-type *cheA* (strain AJW689) and slightly higher than those produced by cells of the wild-type *E. coli* strain RP437 (Fig. 1). When uninduced, cells carrying the plasmid-borne *cheA<sub>RV</sub>M98L(S-)* synthesized CheA<sub>L</sub>M98L at levels similar to those produced by cells carrying *cheA<sub>RV</sub>*, but they produced no detectable CheA<sub>S</sub>. When exposed to 10 μM IPTG, cells containing *cheA<sub>RV</sub>M98L(S+)* synthesized CheA<sub>S</sub> at levels similar to those produced by cells carrying *cheA<sub>RV</sub>* but somewhat lower levels of CheA<sub>L</sub>M98L (Fig. 1).

Prior to this study, a CheA<sub>L</sub>M98I mutant had been generated by changing the start(S) codon in a manner analogous to our M98L change (30). Because cells bearing the CheA<sub>L</sub>M98I allele were impaired in chemotactic ability, we tested cells carrying the CheA<sub>L</sub>M98L allele for their ability to perform chemotaxis. Cells carrying the wild-type allele *cheA*, the control allele *cheA<sub>RV</sub>*, or the mutant allele *cheA<sub>RV</sub>M98L(S-)* exhibited maximal chemotactic behavior when uninduced; this behavior diminished with increasing IPTG levels (Fig. 3). In

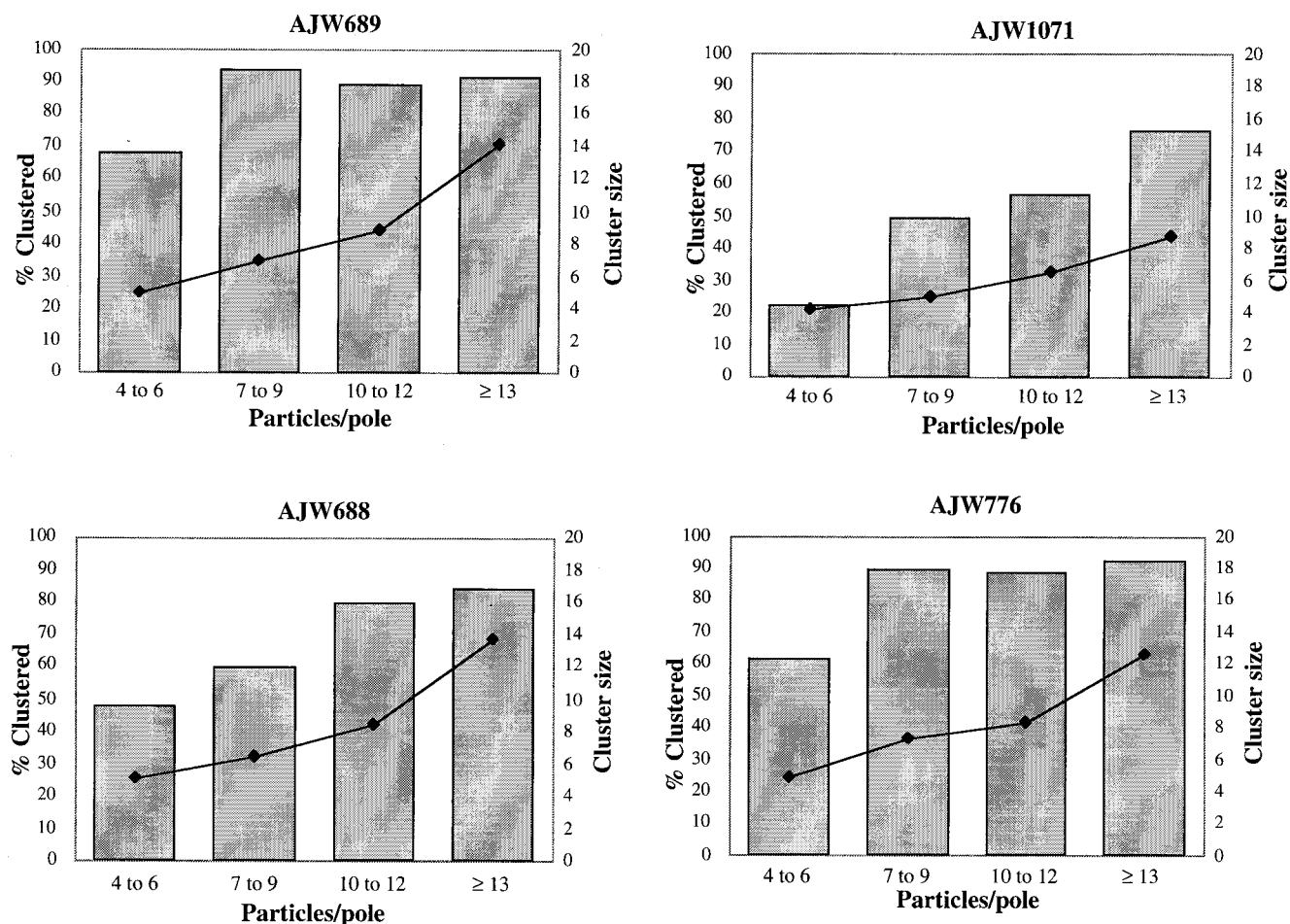


FIG. 2. Clustering potential of CheA variants. The distribution of gold particles in cell sections with at least four gold particles at one cell pole was examined. A minimum of 15 cell poles were included for each category ( $x$  axis). AJW689 ( $\Delta cheA$  pAR1.cheA, uninduced), AJW1071 ( $\Delta cheA$ ), AJW688 ( $\Delta cheA$  pAR1.cheA<sub>S</sub>, 50  $\mu$ M IPTG), and AJW776 [ $\Delta cheA$  pAR1.cheA<sub>RV</sub>M98L(S<sup>-</sup>), uninduced] were used. Grey bars indicate the percentage of gold particles clustered at each cell pole. Black diamonds indicate the mean size of the polar clusters.

contrast, cells carrying the mutant allele *cheA*<sub>RV</sub>M98L(S<sup>+</sup>) exhibited maximum chemotactic behavior when induced by 50  $\mu$ M IPTG.

Cells that synthesized both wild-type CheA<sub>L</sub> and CheA<sub>S</sub> or

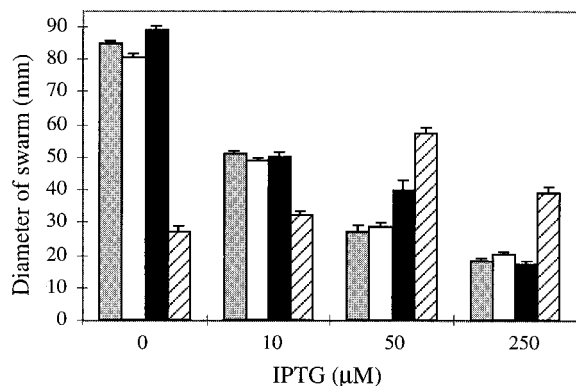


FIG. 3. Chemotactic ability of strains that express various *cheA* alleles. Cells (5  $\mu$ l) were spotted onto semisolid agar, and colony diameters were measured after 7 h of growth at 30°C. Grey bars indicate AJW689 ( $\Delta cheA$  pAR1.cheA), white bars indicate AJW768 ( $\Delta cheA$  pAR1.cheA<sub>RV</sub>), black bars indicate AJW776 [ $\Delta cheA$  pAR1.cheA<sub>RV</sub>M98L(S<sup>-</sup>)], and striped bars indicate AJW774 [ $\Delta cheA$  pAR1.cheA<sub>RV</sub>M98L(S<sup>+</sup>)]. The mean diameter and standard error of the mean for quadruplicate samples are shown.

just CheA<sub>L</sub>M98L alone clustered most of the membrane-associated gold particles at the cell poles (Table 3): 94 and 91% localized to the pole, and 87 and 83% of those polar particles clustered, with mean cluster sizes of 12 and 10 gold particles, respectively. Surprisingly, cells that synthesized CheA<sub>L</sub>M98L and CheA<sub>S</sub> (strain AJW774) yielded lower values (74%, 63%, and 8, respectively) than did cells that synthesized only CheA<sub>L</sub> (strain AJW776 [Table 3]). To explore this observation further, we examined MCP immunolocalization in cells that expressed CheA<sub>L</sub>M98L from a chromosomal location with or without expression of a plasmid-encoded CheA<sub>S</sub> (strains AJW917 and AJW916, respectively). When induced with 10  $\mu$ M IPTG, a concentration that results in CheA levels that approximate those exhibited by strain AJW774 (Fig. 1), the vast majority of the chemoreceptors clustered at the poles in both strains. Thus, expression of CheA<sub>S</sub> in *trans* exerted little or no effect (89 versus 88% of particles localized to the pole; 87 versus 87% of those polar particles clustered, 10 versus 11 gold particles per cluster [Table 3]). Thus, polar clustering of the chemoreceptor complex does not require CheA<sub>S</sub>. Furthermore, under these conditions, moderate levels of CheA<sub>S</sub> apparently neither increase nor decrease the polarity or clustering of the MCPs.

**The carboxyl terminus of CheA is not absolutely required for chemoreceptor clustering.** The C-terminal MC domain of CheA plays a critical role in receiving sensory information

TABLE 3. Spatial distribution of chemoreceptors in cells expressing different CheA variants<sup>a</sup>

Strain	Chromosome-encoded CheA	Plasmid-encoded CheA	Total no. of particles	No. of particles in cytoplasm	Membrane particles <sup>b</sup>				No. of membrane particles/section	No. of polar clusters	Size of polar clusters <sup>d</sup>	No. of lateral clusters	Size of lateral clusters <sup>d</sup>
					Polar particles <sup>c</sup>		Lateral particles						
					%	% in clusters	%	% in clusters					
AJW768	None	CheA <sub>L</sub>	1,901	104	94	87	6	21	9.0	121	12.2 ± 0.8	4	5.5 ± 1.3
		CheA <sub>S</sub>											
AJW776	None	CheA <sub>L</sub> M98L(S-)	2,070	144	91	83	9	23	9.6	143	10.1 ± 0.5	6	6.7 ± 0.9
AJW774	None	CheA <sub>L</sub> M98L(S+)	2,284	286	74*	63*	26	18	10.0	113	8.3 ± 0.4	14	6.8 ± 1.1
AJW916	CheA <sub>L</sub> M98L	None	1,609	121	89	87	11	21	7.4	115	10.0 ± 0.5	5	7.2 ± 1.2
AJW917	CheA <sub>L</sub> M98L	CheA <sub>S</sub>	1,818	118	88	87	12	51	8.5	118	11.0 ± 0.5	15	7.1 ± 0.6
AJW430	CheA <sub>L</sub> K616(Am)	None	1,116	113	81*	77*	19	30	5.0	68	9.2 ± 0.7	10	5.7 ± 0.6
		CheA <sub>S</sub> K616(Am)											
AJW530	CheA <sub>L</sub> K616(Am)	CheA <sub>S</sub>	1,188	119	88	82*	14	9	5.3	73	10.5 ± 0.7	3	4.0
		CheA <sub>S</sub> K616(Am)											
AJW970	None	CheA <sub>L</sub> G422A	2,015	101	87	68*	12	27	9.6	128	8.8 ± 0.5	11	6.2 ± 1.2
		CheA <sub>S</sub> G422A											

<sup>a</sup> The numbers and positions of gold particles detected by immunomicroscopy from 200 thin sections of cells are shown. The numbers and sizes of gold particle clusters located in either the polar or lateral membrane were determined. The relative size of the clusters ± 1 standard error of the mean is presented. CheA expression was uninduced in AJW768, AJW776, and AJW430 and induced at 10 μM IPTG in all other strains.

<sup>b</sup> The percentages of gold particles at the poles or along the lateral edges and the percentage of polar or lateral gold particles that are clustered are shown.

<sup>c</sup> Strains that are significantly different ( $P \leq 0.05$ ) from AJW768 are marked with an asterisk.

<sup>d</sup> Numbers of particles per cluster.

from chemoreceptors (7, 28). To investigate whether clustering of chemoreceptors requires the C-terminal 39 amino acids of CheA, we examined MCP immunolocalization in strain AJW430 [*cheAK616(Am)*] (40). We observed an approximately twofold reduction in the level of chemoreceptor protein in this nonchemotactic strain (Table 3 and data not shown). Despite this reduction in MCP levels, the gold particles clustered moderately at the cell pole (81% polar, 77% of which formed clusters containing a mean of nine gold particles [Table 3]). The addition of wild-type CheA<sub>S</sub> (strain AJW530; 10 μM IPTG) restored chemotactic ability (Fig. 3), presumably due to the formation of functional CheA heterodimers (35, 41), but did not enhance the clustering of the MCPs significantly (Table 3).

**The G1 domain of CheA is not required for chemoreceptor clustering.** The allele *cheAG422A* encodes a single-amino-acid substitution within the highly conserved glycine-rich G1 region of CheA, resulting in a mutant protein that does not support chemotaxis. In vitro, the CheA<sub>L</sub>G422A protein binds ATP poorly and does not autophosphorylate (9, 34). However, cells that synthesized CheA<sub>L</sub>G422A and CheA<sub>S</sub>G422A (strain AJW970) exhibited an enhanced polar clustering of chemoreceptors relative to the *cheA* deletion strain (AJW1071). The majority of the gold particles were polar (87%) and were moderately clustered (68% of the polar particles were clustered, with a mean cluster size of nine particles [Table 3]). Thus, a single-amino-acid change that interferes with nucleotide binding, autophosphorylation, and chemotaxis only slightly reduces the ability of the chemoreceptors to cluster at the poles.

## DISCUSSION

To gain insight into the relationship between chemotactic signaling and clustering of MCP-CheA-CheW ternary complexes, we examined the ability of wild-type and mutant CheA variants to promote chemoreceptor polarity and clustering. We used immunolocalization techniques to determine the cellular location of MCPs in cells that synthesize approximately equal amounts of wild-type or mutant CheA proteins. We found that

the CheA<sub>L</sub> and CheA<sub>S</sub> proteins synthesized from a plasmid-borne wild-type *cheA* allele mediate polar clustering of MCPs approximately as well as reported previously for the same proteins synthesized from the chromosomal locus (19). We also observed that CheA<sub>L</sub>M98L, a functional variant of CheA<sub>L</sub>, suffices to mediate wild-type levels of clustering in the absence of CheA<sub>S</sub> and that CheA<sub>S</sub> alone mediates clustering, albeit at reduced levels.

Cells that synthesized only CheA<sub>L</sub>M98L from the plasmid-borne *cheA<sub>RV</sub>M98L(S-)* allele produced steady-state CheA<sub>L</sub> levels, exhibited chemotactic behavior, and yielded MCP localization patterns indistinguishable from those of cells that synthesized both wild-type CheA<sub>L</sub> and CheA<sub>S</sub>. In contrast, cells that synthesized CheA<sub>L</sub>M98L and CheA<sub>S</sub> from the plasmid-borne *cheA<sub>RV</sub>M98L(S+)* migrated at about one-third the rate of cells that synthesized only CheA<sub>L</sub>M98L. Moreover, they exhibited reduced polar localization and clustering of MCPs. However, the reduction in chemotactic ability and MCP clustering in this strain does not appear to be caused specifically by the presence of CheA<sub>S</sub>. Cells that synthesized CheA<sub>L</sub>M98L from a chromosomal copy of *cheA<sub>RV</sub>M98L(S-)* exhibited chemotactic behavior and yielded MCP localization patterns to identical levels regardless of whether CheA<sub>S</sub> (from a plasmid) was expressed. Thus, although CheA<sub>S</sub> can constitute up to 50% of the total CheA synthesized by wild-type cells (37), it does not seem to be required for either chemotaxis or polar aggregation of MCPs, nor does it seem to interfere with CheA<sub>L</sub>M98L-mediated clustering of these chemoreceptor complexes.

All of the CheA variants examined in this study increased the level of MCP polar clustering over that seen in cells lacking any CheA. However, in all cases, the polar clustering of the MCPs was reduced in comparison to cells that possessed a wild-type CheA<sub>L</sub>. Because CheA<sub>S</sub> lacks most of the P1 domain that contains the site of histidyl phosphorylation, the enhanced aggregation of MCPs cannot require CheA autophosphorylation. In fact, enhanced MCP clustering seems not to require kinase activity at all. A single-amino-acid G422A substitution in CheA<sub>L</sub> and CheA<sub>S</sub> did not eliminate either polar localization or clustering of MCPs, although these mutant proteins exhibit little or no detectable kinase activity (reference 41

and unpublished data). Because these mutant proteins display a considerably reduced capacity to bind ATP and related nucleotides (34), the enhanced MCP aggregation apparently also does not require nucleotide binding by CheA.

The CheA-mediated polar aggregation of the MCPs is also partially independent of the C-terminal 39 amino acids of CheA. The truncated CheA<sub>L</sub>K616(Am) and CheA<sub>S</sub>K616(Am) proteins expressed together also mediated both polar localization and intermediate clustering of MCPs. In vitro, these proteins retain kinase activity, but in vivo, they do not support chemotaxis, presumably because they do not interact properly with MCPs, CheW, or both (7). Thus, it seems likely that the slight reduction in polar aggregation in this mutant results from the diminished capacity of these truncated proteins to form either ternary complexes or a higher-order complex.

At the resolution of immunoelectron microscopy, several CheA variants that are defective in chemotactic signaling clearly support significant levels of chemoreceptor complex clustering. This observation strongly supports the hypothesis that clustering is an integral part of signaling, i.e., that clustering of chemoreceptor complexes occurs prior to rather than in response to signaling. Thus, signaling must occur in the context of these preformed clusters. Although no one has defined the function of MCP clustering, it is reasonable to suppose that aggregation of chemoreceptor complexes contributes to amplification of the chemotactic signal (8, 19). Clearly, additional studies must be performed to clarify the relationship between chemoreceptor complexes and signal amplification.

Finally, it is clear that CheA-independent and therefore ternary-complex-independent aggregation of the MCPs can occur. The lack of maximal clustering in the absence of CheA is not simply due to a reduction in the number of polar gold particles, since  $\Delta cheA$  cells with comparable numbers of polar gold particles were reduced in their cluster number and size compared to wild-type cells. However, the observation that some CheA-independent MCP clustering occurs and that clustering is enhanced by CheA raises the possibility that either (i) the clustering potential of all of the chemoreceptors is greatly enhanced in the presence of CheA or (ii) there are differences in the requirement for CheA in the clustering of the four different chemoreceptors. Methods to detect the individual chemoreceptors must be generated in order to examine these possibilities.

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