Binding of the Escherichia coli response regulator CheY to its target measured in vivo by fluorescence resonance energy transfer

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In Escherichia coli chemotaxis, signaling depends on modulation of the level of phosphorylation of CheY, a small protein that couples receptors and flagellar motors. Working in vivo, we used fluorescence resonance energy transfer (FRET) to measure the interaction of CheY–P with its target, FliM. Binding of CheY–P to FliM was found to be much less cooperative than motor switching; however, under the conditions of our experiment, most of the FliM appeared to be in the cytoplasm. We studied signal processing times in the chemotaxis pathway by measuring the changes in CheY–P binding to FliM on flash release of caged chemoeffectors. Following sudden addition of attractant, the amount of CheY–P bound to FliM decayed exponentially with a rate constant of about 2 s⁻¹. Following sudden addition of repellent, FliM occupancy increased with a rate constant of about 20 s⁻¹. Using these data, we were able to construct a simple model for the chemotactic pathway and to estimate values of rate constants for several key reactions.

Materials and Methods

Plasmids and Strains. Fusions to CheY and FliM of yellow and cyan fluorescent protein (YFP and CFP, CLONTECH) were made as described (2, 9). The chey-cyfp fusion was cloned into pTrc99A (Amp⁺, Pharmacia) under an isopropyl β-D-thiogalactoside (IPTG)-inducible promoter, yielding pVS18. The ecfp-fliM fusion was cloned into pBAD33 (Cm⁺) under an arabinose-inducible promoter (12), yielding pVS31. The strains used for the FRET experiments were derived from strain DFB228 (ΔfliM), the gift of David Blair (University of Utah, Salt Lake City). VS120 (ΔcheY ΔfliM) and VS121 (Δ[cheY-cheZ] ΔfliM) were made by in-frame deletions of the corresponding chemotaxis genes (2). Strain VS125 [Δ2206(tap-cheZ) ΔfliM] was made by P1-transduction of the ΔfliM strain from RP2893 [Δ2206(tap-cheZ)], the gift of Sandy Parkinson (University of Utah, Salt Lake City). Growth with 34 μg/ml chloramphenicol and 100 μg/ml ampicillin was used for selection. Because the CheY/FliM interaction always was tested in a ΔcheY ΔfliM background, strain VS120 is referred to as wild type and only additional background mutations are indicated in the text. Strain VS125 is referred to as cheR cheB cheZ. YFP and CFP proteins were expressed as C-terminal fusions to 6×His tag from pVS47 and pVS48 plasmids, respectively, and purified using Ni-NTA spin kit (Qiagen, Valencia, CA).

Data Acquisition and Analysis. Cells were grown and prepared for FRET as described (9). The levels of arabinose and IPTG induction were 100 μg·ml⁻¹ and 0.05 mM, respectively, unless specified otherwise. We monitored fluorescence from a field of 300–500 cells in each experiment, and repeated each experiment at least three times by using different cell cultures. Fluorescence
signals in cyan and yellow channels were detected using two photon-counting photomultipliers (H7421-40, Hamamatsu, Bridgewater, NJ) whose outputs were converted to analog signals by ratemeters (RIS-375, Rowland Institute). For the flow experiments the signals were filtered by 8-pole low-pass Bessel filters (3384, Krohn-Hite, Avon, MA) with 2 Hz cutoff frequency and sampled at 5 Hz by a computer data-acquisition system [National Instruments (Austin, TX) LABVIEW 5.1 and Macintosh G3]. For the flash experiments using wild-type cells the signals were filtered with 200 Hz cutoff frequency and sampled at 500 Hz, whereas for the flash experiments using cheZ cells the signals were filtered with 10 Hz cutoff frequency and sampled at 20 Hz.

FRET was calculated as fractional change in CFP fluorescence, as described in the text. Alternatively, changes in yellow emission could be computed from changes in the ratios of yellow and cyan emission, as described (9). The two methods gave essentially the same results (data not shown).

**Curve Fitting.** Fits were made to a multisite Hill model of the form $F_{\text{max}}Y^H/(Y^H + K_{H}^H)$, where $F_{\text{max}}$ is the value of FRET at saturating CheY-YFP concentrations, $Y$ is the concentration of CheY-YFP, $H$ is the Hill coefficient, and $K_{H}^H$ is the apparent dissociation constant (Fig. 2A), or of the form $M_0 + (\Delta M_{\text{max}} A^H/(A^H + K_{H}^H))$, where $M_0$ is the initial FliM occupancy, $\Delta M_{\text{max}}$ is the change in FliM occupancy observed at saturating stimulation, $A$ is the step in attractant concentration, $H$ is the Hill coefficient, and $K_{H}^H$ is the value for the half-maximal response (Fig. 3). Decays following release of caged aspartate were fit to exponentials, $\left[ M_0 - M_1 \right] \exp\left[ -k(t - \tau) \right] + M_1$, where $M_0$ is the initial FliM occupancy or initial CheY–P level, $M_1$ is the final FliM occupancy or final CheY–P level, $k$ is the decay rate, $t$ is the time after the flash, and $\tau$ is a delay before the onset of the response.

**Concentration Measurements.** The cellular concentrations of CheY-YFP and CFP-FliM were determined by sonication of suspensions of nonfluorescent cells of similar density. Cell number per milliliter of culture was determined by plating on nonselective LB plates, incubating overnight, and counting colonies. The average volume of the cell cytoplasm was determined as described (13), and was found to be $1.4 \times 10^{-15}$ liters. The average intracellular protein concentration was then calculated by dividing the amount of CheY-YFP or CFP-FliM fusion protein per milliliter cell culture by the number of cells per milliliter of cell culture and the cytoplasmic volume. In a standard experiment, with cells grown in the presence of 100 μg/ml−1 arabinose and 0.05 mM IPTG, the concentration of CFP-FliM was 5.8 ± 0.6 μM, and the concentration of CheY-YFP was 17.9 ± 1.5 μM. The concentration of CheY-YFP at other induction levels was determined by comparing the levels of YFP fluorescence. The level of expression of CFP-FliM in our standard experiment, necessary for optimum complementation of motility, was about three times higher than the level of FliM in the wild type (1.7–2.0 μM; refs. 14 and 15). Applying fractionation by ultracentrifugation, Zhao et al. (15) found that in wild-type cells, most of FliM (~75%) is present in either functional or incomplete flagellar motors. Using a similar approach, we estimated that at the expression level used in our experiments about 20–30% of CFP-FliM is present in such motors. The balance appears to be in the cytoplasm (data not shown).

**Flash Release of Caged Chemoeffectors.** NPE-caged l-aspartate (N-[1-(2-nitrophenyl)-ethyl]oxycarbonyl]aspartic acid, Calbiochem), nitroindolyl-caged l-aspartate (the gift of John Corrie, National Institute for Medical Research, Mill Hill, London), and NPE-caged proton (2-hydroxyphenyl-1-(2-nitrophenyl)ethyl phosphate, Molecular Probes) were photolyzed by use of a xenon flash lamp [Chadwick–Helmuth (El Monte, CA) 35S driven by model 238 power supply, 200 Ws, 50 μs]. The flash lamp was mounted in an Oriel (Stamford, CT) housing (66055) equipped with a fused silica condensing lens (60076) and a Schott UG-5 filter (330 nm peak). The light was fed to the microscope with a liquid light guide (Oriel 77554) and focused onto the flow cell with a 25-nm fused silica lens (Oriel 41220). The light was reflected by a UV-enhanced aluminum mirror (Edmund Scientific, Barrington, NJ) and entered the flow cell from below, through a fused quartz bottom window. The focused spot was about 4 mm wide, much larger than the field of view. A UV-blocking filter (Chroma Technology, Brattleboro, VT) was inserted into the light path above the lower dichroic mirror to prevent UV light from reaching the photomultipliers. The UV flash generated an artifact present in the absence of cells or caged chemoeffectors (a pulse sensed by the photomultipliers that decayed in about 10 ms). This artifact was measured separately before each experiment and subtracted from the data. The efficiency of release of caged l-aspartate was calibrated using the amplitude of the FRET response in cheR cheB cells (9).

In a typical experiment, 200 μM caged l-aspartate in tethering buffer supplemented with 5 mM DTT was drawn into the flow cell, and the cells were allowed to equilibrate for 5 min under slow flow (50 μl·min−1). The flow was stopped, and ~10 μM of aspartate was released by a single flash. After data collection, the solution was exchanged by flow, and the procedure was repeated 10–20 times with the same cell culture. The experiment was repeated at least three times with different cell cultures. Release of NPE-caged l-aspartate produced a FRET response identical to that produced by nitroindolyl-caged l-aspartate, which is released much more rapidly; therefore, the rate of release is not limiting at these concentrations. Stimulation with 2.5-mM caged proton was performed similarly, except that the concentration of phosphate in the tethering buffer was reduced to 0.1 mM. The resulting pH change was calibrated by using the pH-sensitivity of the fluorescence of 0.05 μM fluorescein.

**Modeling Response Kinetics.** A system of linear differential equations was used describing CheA autophosphorylation, phospho-transfer from phosphorylated CheA to CheY, binding of CheY–P to FliM, binding of CheY–P to CheZ, dephosphorylation of CheY–P by CheZ, and autodephosphorylation of CheY–P, but excluding adaptation reactions. The activity of chemotaxis pathway was expressed as the fraction of active CheA (16). The system of equations was solved using the NDSolve function of MATHEMATICA 3.0 (Wolfram Research, Champaign, IL). The complete description of the model including rate constants and protein concentrations is published as supporting information on the PNAS web site, www.pnas.org.

**Results**

CheY–P Interaction with FliM Can Be Studied in Vivo by Using FRET. The experimental scenario is shown in Fig. 1. Here, cells are attached to the coverslip of a flow cell and attractant is added or removed. The addition of attractant lowers the level of CheY phosphorylation and leads to a decrease in energy transfer, and the removal of attractant has the opposite effect (Fig. 1A). Addition of attractant to wild-type cells (Fig. 1B) resulted in a rapid decrease in FRET followed by gradual recovery due to adaptation (receptor demeth-
Bleaching YFP with a 532-nm laser (9) also can reveal any non-phosphorylation-dependent interactions. The two methods gave the same result as addition of a saturating amount of attractant, except at high concentrations of CheY-YFP, where a fraction of the CFP-FliM fluorescence was quenched in absence of CheY-YFP phosphorylation. At the highest CheY-YFP concentration we have used, 45 μM, about 3% of CFP-FliM fluorescence was quenched. This quenching probably was due to the formation of heterodimers of CFP and YFP, observed recently at high protein concentrations (20). The FRET efficiency predicted for the CFP/YFP heterodimer is \( \approx 90\% \) (19). Thus, in our experiment, \(<4\%\) of CFP-FliM is expected to be bound by CheY-YFP in a phosphorylation-independent manner.

**CheZ Does Not Play a Role in Signal Amplification.** Fig. 3 shows initial changes in FliM occupancy deduced from initial changes in FRET following addition and removal of α-methylaspartate (MeAsp) by flow, as defined in ref. 9, in a sequence of steps of increasing size, for wild-type cells (Fig. 3A) and cheZ cells (Fig. 3B). The values for \( K_{1/2} \) and the Hill coefficient obtained for the
wild type (see Fig. 3 legend) indicate that the interaction of CheY–P with FliM does not show a significantly higher cooperativity than the phosphorylation-dependent interaction of CheY with CheZ, observed earlier (9), which gave \( K_{1/2} = 2.6 \pm 0.5 \text{ M} \) for attractant addition and \( 7.4 \pm 0.4 \text{ M} \) for attractant removal, with \( H = 1.3 \pm 0.1 \). Because the \( K_{1/2} \) values for the interaction of CheY with FliM are only 2-fold higher in the absence of CheZ, our results confirm the previous observations (9, 23) that CheZ does not play an important role in signal amplification in chemotaxis.

Fig. 3 also illustrates the advantage of using FRET to study chemotactic response. As in Fig. 2B, the unshaded areas indicate the range of FliM occupancies over which one would expect to see a motor response. Whereas the FRET assay is able to distinguish responses to the stimuli of up to 100 \( \mu \text{M} \) MeAsp, the motor would saturate at concentrations above 2 \( \mu \text{M} \) for wild-type cells and above 10 \( \mu \text{M} \) for cheZ cells. Moreover, with the motor assay, the response of cheZ cells to addition of attractant would appear less sensitive because of the high initial FliM occupancy, and the response to removal of attractant would not be detected at all.

**Time-Resolved Study of Kinetics of the Chemotactic Response.** By using flash-stimulated release of caged attractants or repellents, we were able to make measurements of signal processing times in *E. coli* chemotaxis, as shown in Fig. 4. In wild-type cells stimulated with a saturating amount of the attractant, aspartate (Fig. 4A), there was an initial delay (65 ± 9 ms), followed by a rapid decay (with a first order rate constant, \( k = 2.0 \pm 0.1 \text{ s}^{-1} \)).

The delay was determined by the intersection of the exponential fit to the decay with the prestimulus baseline (Fig. 4A Inset). The rate constant is smaller than that measured in similar experiments following changes in behavior of free-swimming cells (7). As argued above, this might be due to the nonlinear nature of the motor response, because a saturating counterclockwise motor bias (shaded area) is reached sooner than zero FliM occupancy, which gives a lower value of the half-time of the response, and thus a higher value of apparent rate constant. The decrease of FliM occupancy in cheZ cells after stimulation was much slower (Fig. 4B), with \( k = 0.059 \pm 0.001 \text{ s}^{-1} \), as expected, given the absence of the phosphatase.

The delay was not detected, because the filter used to smooth the data were set to a longer time span.

Given the relationship between FliM occupancy and the total concentration of CheY–P (Fig. 2A) we also could compute the corresponding CheY–P decay curves (not shown). This gave us slightly larger rate constants, \( k = 2.2 \pm 0.1 \text{ s}^{-1} \) and \( 0.085 \pm 0.001 \text{ s}^{-1} \), corresponding to decay half times of \( 0.52 \pm 0.32 \text{ s} \) and 8 s, respectively. Assuming that the off rate of CheY–P from the FliM complex is fast relative to rate of dephosphorylation, these
would be the half times for CheZ-catalyzed and spontaneous dephosphorylation, respectively.

Stimulation of wild-type cells by flash photolysis of a caged proton, a repellent, Fig. 4C, revealed no measurable delay (<20 ms). The maximum FliM occupancy was reached within 70–100 ms. This is a good agreement with the value of ~50 ms obtained for free-swimming cells (6). From the time required to reach ~65% \( (1 - (1/2)) \) of the maximal repellent response (~45 ms), the first-order rate constant of the repellent response was estimated as ~22 s\(^{-1}\).

**Discussion**

Physiological responses of chemotactic *E. coli* cells have been studied extensively before, and many biochemical assays have been carried out on the reactions involved in the pathway. The use of FRET has allowed us to take these studies one step further to investigate responses at the level of the response regulator, CheY-P. In our previous study (9), we used FRET between CheZ-CFP and CheY-YFP to infer the activity of the receptor kinase complex. Here, we used FRET between CFP-FliM and CheY-YFP to study their binding and its changes on chemotactic stimulation. We found that the binding of phosphorylated CheY-YFP to CFP-FliM in the cell is much less cooperative than the motor response. This finding agrees with allosteric transition models, where the binding function is generally less cooperative than the state function (24, 25). However, given that only a fraction of CFP-FliM in our experiment is incorporated into functional flagellar motors, the agreement with the allosteric models could be fortuitous. To make a precise comparison between binding of CheY\(^-\)P to the motor and motor switching, a better test would be to measure the binding affinities of phosphorylated CheY-YFP to CFP-FliM localized on a single motor. We hope that such a measurement will be possible using the FRET technique.

We combined FRET with flash release of caged chemoeffectors to determine signal processing times in the pathway. In wild-type cells, response times for changes in FliM occupancy are determined by several sets of rates: (i) rates of receptor-linked CheA autophosphorylation, CheY phosphorylation by CheA\(^-\)P, and CheY\(^-\)P dephosphorylation by CheZ; (ii) on- and off-rates of CheY\(^-\)P binding to FliM; and (iii) CheY\(^-\)P diffusion from receptor-kinase clusters on the cell pole across the cytoplasm to the flagellar motors or to cytoplasmic FliM. The phosphotransfer from CheA\(^-\)P to CheY is very fast (26), so the response to repellent (CheA-activating) stimuli is presum-ably limited by the rate of CheA autophosphorylation. From the kinetics of the repellent response, the rate constant for autophosphorylation of activated CheA should be >22 s\(^{-1}\), in good agreement with *in vitro* measurements (27) of 20–30 s\(^{-1}\). The response time to the sudden addition of attractant is determined in wild-type cells by CheZ-dependent dephosphorylation of CheY, giving the rate constant of ~2 s\(^{-1}\). The 65-ms delay between stimulation and the onset of the response might be due, in part, to the ~50-ms delay between CheY\(^-\)P binding to CheZ and CheZ activation observed *in vitro* (28). Our data also allowed us to estimate the on- and off-rates of CheY\(^-\)P binding to FliM. The time course of the repellent response implied an on-rate constant of >4 \( \times 10^6 \) M\(^{-1}\)s\(^{-1}\) and, given the average \( K_d \) of CheY\(^-\)P binding to FliM of 3.7 \( \mu \)M, an off-rate of >15 s\(^{-1}\). This is close to the values for the on-rate constant, 3 \( \times 10^4 \) M\(^{-1}\)s\(^{-1}\) (diffusion-limited), and the off-rate constant, 10 s\(^{-1}\), assumed in a recent model for motor switching (25).

More detailed modeling was performed using a set of linear differential equations describing phosphotransfer and binding reactions in the chemotaxis pathway. Using the rate estimates

\[ \Delta \text{pH} = -0.45 \]

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made above and values from the literature (see Materials and Methods and supporting information), we were able to reproduce the responses observed to addition of attractant or repellent shown in Figs. 4A and C. The corresponding model values appear in Fig. 5A and B. Note that adaptation was not included in our model; hence, the difference between Figs. 4C and 5B.

The model allowed us to further refine the estimates of reaction rates required to match the observed response kinetics. To match the rate of increase in FliM occupancy observed on addition of repellent (Fig. 5B), the rate constant for autophosphorylation of activated CheA had to be set to 50 s\(^{-1}\), higher than the value estimated above. The delay predicted for this response was ≈5 ms, below the time resolution of our experiment. To match the rate constant of decay in FliM occupancy observed on addition of attractant (≈2 s\(^{-1}\); Fig. 5B), we had to set the concentration of CheZ in the cytoplasm to 1.1 μM. The total concentration of CheZ in the cell is 4–12 μM, depending on strain and growth conditions (13), but most of the CheZ is localized to receptor clusters (2), which might justify the lower value for the cytoplasmic concentration. Our simulation predicted a short (≈20 ms) delay in response to attractant because of a lag in dissociation of CheY-P from FliM. Given the additional 50-ms delay in CheZ activation mentioned above, this might explain the observed delay of ≈65 ms.

The reactions in the chemotaxis pathway appear to be as fast as needed to work within the diffusion limit. The diffusion coefficient for CheY-GFP has been measured (8) (4.6 μm\(^2\)s\(^{-1}\)), and the diffusion coefficient for CheY was estimated to have a similar value (ref. 29; of order 10 μm\(^2\)s\(^{-1}\)). Thus, the time required for CheY-P to diffuse ≈1 μm from a receptor cluster near one pole to the flagellar motors, distributed at random along the sides of the cell, is of order \(x^2/2D\) ≈ 50 ms. This is the same order of magnitude as the time required for the repellent response. If CheY is phosphorylated at one pole and CheY-P decays with the rate constant measured for the attractant response, \(k \approx 2 s^{-1}\), then its concentration will not be uniform over the length of the cell. At steady state, the distance from the pole over which the concentration of CheY-P falls by \(1/e \approx 1/3\), is \((D/k)^{1/2} \approx 2 \mu m\), as observed (29). Thus, given the ultrasensitivity of the motor response shown in Fig. 2B, the rotational bias might vary significantly with the distance from the pole. This probably sets an upper limit on the allowable rate for CheY-P hydrolysis in the cytoplasm. We know that the disparity in rotational bias is not very great, because fully adapted wild-type cells spend most of their time swimming at top speed (30), which occurs only when all of the filaments spin counterclockwise in one coherent bundle, while some tumble involve clockwise rotation of all of the flagella (31).

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