

Fig. S1. Induction curves for YFP-CheR and CheB-CFP for single gene mutants in the capillary assay. (A) Fluorescence intensity of the YFP-CheR proteins as a function of the concentration of the inducer IPTG used in capillary experiments. The grey line is the best fitted Hill function to the data points: $A \cdot [\text{IPTG}]^N / ([\text{IPTG}]^N + B^N) + C$, $A=1239.0$, $B=41.1 \mu\text{M}$, $C=84.4$ and $N=1.5$ with $R^2=0.99$. (B) Fluorescence intensity of the CheB-CFP proteins as a function of the concentration of the inducer aTc used in capillary experiments. The grey line is the best fitted Hill function to the data points: $A \cdot [\text{aTc}]^N / ([\text{aTc}]^N + B^N) + C$, $A=6299.3$, $B=12.3$, $C=24.4$ and $N=5.3$ with $R^2=0.92$.

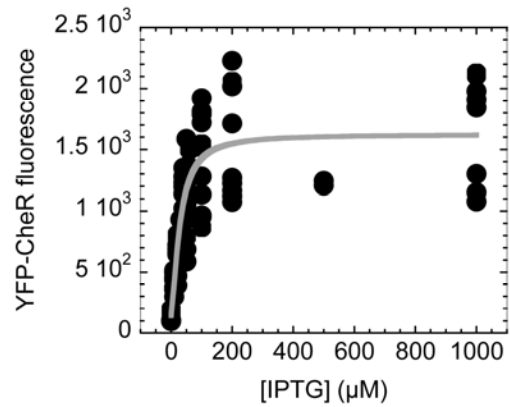


Fig. S2. The fluorescent intensity of YFP-CheR as a function of [IPTG] in capillary experiments with $\Delta cheR$ - $\Delta cheB$ cells.

We measured the fluorescence intensity of YFP-CheR from $\Delta cheR$ - $\Delta cheB$ cells in the capillary experiments. The grey line is the best fitted Hill function to the data points: $A \cdot [IPTG]^N / ([IPTG]^N + B^N) + C$, $A=1480.4$, $B=28.8 \mu M$, $C=143.3$ and $N=1.5$ with $R^2=0.76$.

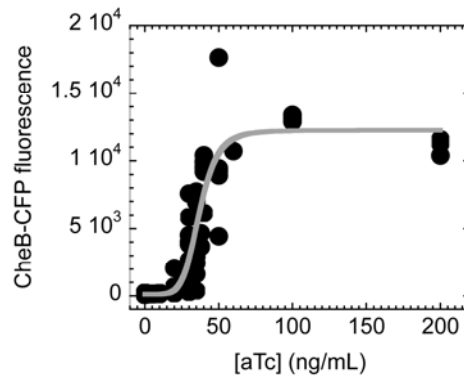


Fig. S3. The fluorescent intensity of CheB-CFP as a function of [aTc] in capillary experiments with Δ CheR- Δ CheB cells.

We measured the fluorescence intensity of CheB-CFP from Δ *cheR*- Δ *cheB* cells in the capillary experiments. The grey line is the best fitted Hill function to the data points: $A \cdot [aTc]^N / ([aTc]^N + B^N) + C$, $A=12129$, $B=37.0$, $C=123.7$ and, $N=6.0$ with $R^2=0.83$.