DnaK and DnaJ heat shock proteins participate in protein export in *Escherichia coli*

Jadwiga Wild,1 Elliot Altman,2 Takashi Yura,3 and Carol A. Gross4

1Department of Bacteriology, University of Wisconsin–Madison, Madison, Wisconsin 53706 USA; 2Elliot Altman, Department of Biology, University of Utah, Salt Lake City, Utah 84112 USA; 3Institute for Virus Research, Kyoto University, Kyoto, Japan

In *Escherichia coli* secreted proteins must be maintained in an export-competent state before translocation across the cytoplasmic membrane. This function is carried out by a group of proteins called chaperones. SecB is the major chaperone that interacts with precursor proteins before their secretion. We report results indicating that the DnaK and DnaJ heat shock proteins are also involved in the export of several proteins, most likely by acting as their chaperones. Translocation of alkaline phosphatase, a SecB-independent protein, was inhibited in *dnaK*− and *dnaJ*− mutant strains, suggesting that export of this protein probably involves DnaK and DnaJ. In addition, DnaK and DnaJ play a critical role in strains lacking SecB. They are required both for viability and for the residual processing of the SecB-dependent proteins LamB and maltose-binding protein (MBP) seen in *secB* null strains. Furthermore, overproduction of DnaK and DnaJ permits strains lacking SecB to grow in rich medium and accelerates the processing of LamB and MBP. These results suggest that under conditions where SecB becomes limiting, DnaK and DnaJ probably substitute for SecB and facilitate protein export. This provides the cell with a mechanism to overcome a temporary imbalance in the secretion process caused by an abrupt expansion in the pool of precursor proteins.

[Key Words: DnaK, Hsp70, protein export, chaperones, alkaline phosphatase, SecB]

Received March 23, 1992; revised version accepted April 22, 1992.
with GroES [Fayet et al. 1989], and DnaK which usually acts with DnaJ and GrpE (Alfano and McMaken 1989; Dodson et al. 1989; Straus et al. 1990; Liberek et al. 1991]. Two pieces of evidence suggest that HSPs also perform a chaperone function in protein export. First, the GroEL/GroES HSP complex appears to be the primary chaperone for the export of β-lactamase. groEL and groES mutants are defective in processing β-lactamase in vivo [Kusukawa et al. 1989] and the GroEL and GroES proteins are required for in vitro transport of this protein [Bochkareva et al. 1988; Laminet et al. 1990]. Second, HSPs can substitute for the SecB chaperone and are required for cellular viability in the absence of a functional SecB protein [Altman et al. 1991].

Interestingly, there has been no evidence that DnaK plays a role in the export of normal proteins, although increased levels of DnaK (or GroEL and GroES) facilitate export of a LamB–LacZ hybrid protein [Phillips and Silhavy 1990]. The lack of involvement of DnaK in protein export in prokaryotes was surprising because Hsp70s, the DnaK homologs in eukaryotic cells, participate in several aspects of protein transport. Mitochondrial Hsp70, together with cytoplasmic Hsp70, is involved in the initial step in mitochondrial import [Kang et al. 1990]. In addition, a cytoplasmic Hsp70 [Deshaiies et al. 1988] and dnaJ homologs [Sadler et al. 1989; Atencio and Yaffe 1992] are required for protein transport through the endoplasmic reticulum.

In this report we have further investigated the role of DnaK in protein export. We find that DnaK and DnaJ are involved in export of AP, a SecB-independent protein. Furthermore, we show that secB null mutant strains require DnaK and DnaJ for viability and for export of MBP and LamB, two SecB-dependent proteins.

Results

DnaK and DnaJ participate in the export of AP

Although AP was the first periplasmic enzyme shown to be synthesized in a precursor form [Inouye and Beckwith 1977], its translocation pathway has not yet been fully established. AP is transported via the SecA export pathway [Oliver and Beckwith 1981], but secB mutations have little effect on the rate of processing of pre-AP to the mature protein [Kumamoto and Beckwith 1985; Gannon et al. 1989] except at 30°C [Kusukawa et al. 1989], and no other chaperone has been implicated in its export.

Recently, we described dominant dnaK mutations that result in partially functional DnaK proteins [Wild et al. 1992]. We used the three mutations with the strongest phenotypes [EK171, GD229, and GD341] to determine whether the export of AP was dependent on DnaK. Whereas processing of AP in wild-type cells was almost complete after a 10-sec chase, two of the dnaK mutant strains (EK171 and GD229) processed AP very slowly [Fig. 1]. The third mutant (GD341) exhibited a wild-type processing rate [data not shown]. The greatest defect in processing was exhibited by EK171, which required 5 min to process 50% of the pre-AP to the mature form. In GD229, 30% of AP remained unprocessed even after a 2.5-min chase [Fig. 1]. These results suggest a role for DnaK in the export of AP.

DnaK often works in concert with two other HSPs, DnaJ and GrpE. DnaJ and GrpE not only modulate the activity of DnaK [Liberek et al. 1991] but may also interact directly with the target proteins [Alfano and McMaken 1989; Dodson et al. 1989; Zylicz et al. 1989; S. Wickner et al. 1991]. We therefore determined whether dnaJ and grpE mutations affected AP processing. We found that the dnaJ mutant HQ33 exhibited a defect in AP processing [Fig. 1]. Although the effect of the dnaJ mutation was not as severe as those exhibited by dnaK mutations, these results suggest that DnaJ is involved in the processing of AP. In contrast, grpE280 [Saito and Uchida 1977; Ang et al. 1986] did not exhibit a defect in AP processing [Fig. 1].

We have also examined the kinetics of processing of pre-RBP, a periplasmic protein that is believed, by a variety of criteria, to be exported independently of SecB [Collier et al. 1990]. Neither the dnaK, dnaJ, and grpE mutations, described above, nor the groEL30 or groEL140 mutations affect processing of this protein [data not shown]. Either export of pre-RBP is chaperone independent, or a different chaperone is involved in maintaining the translocation-competent form of this precursor.

DnaK and DnaJ are required in strains lacking SecB

The growth and protein export properties of secB::Tn5, a strain that lacks SecB, have been characterized extensively. Two of its properties, which may be related, are

Figure 1. Processing of alkaline phosphatase in mutant strains. Cells growing exponentially in glycerol medium at 30°C were pulse-labeled for 15 sec with Tran35S-label [ICN], chased with nonradioactive amino acids for indicated times, immunoprecipitated with anti-AP antibody, and subjected to electrophoresis on 10% SDS–polyacrylamide gels to display the unprocessed [pre-AP] and mature [AP] forms of alkaline phosphatase.

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surprising. First, SecB-dependent proteins are still exported, albeit at reduced rates. Second, the growth defect of secB strains is manifested only in rich medium (Kumamoto and Beckwith 1985). This growth defect is suppressed when HSPs are overproduced by increasing expression of σ22. This observation suggests that some HSPs can substitute for the SecB function by acting as “backup chaperones” (Altman et al. 1991). The level of HSPs in the secB null mutant may be sufficient to permit growth and protein export in minimal but not rich medium.

We considered the possibility that DnaK and DnaJ might be the HSPs substituting for the SecB function. If so, severe dnaK and dnaJ mutations should be lethal when combined with a secB null mutation. This proved to be the case. It was impossible to construct a secB::Tn5 strain that contained a ΔdnaJ allele, although the isogenic secB+ strain was perfectly viable when deleted for dnaJ [data not shown]. Likewise, when the secB::Tn5 insertion was transduced into the GD229 and EK171 dnaK mutants, 10-fold fewer transductants were recovered. All of the transductants, unlike the recipient strain, grew in rich medium. The simplest explanation of these results is that the transductants had acquired a suppressor mutation that simultaneously permitted viability of the secB::Tn5 dnaK strain in minimal and growth in rich medium. Consistent with this interpretation, when these double mutant strains were retransduced to dnaK++, the resulting secB::Tn5 transductants grew in rich medium. In contrast, a secB::Tn5 grpE280 double mutant strain was viable and retained typical SecB phenotypes. These results suggest that DnaK and DnaJ are required for the viability of a secB::Tn5 strain.

To further investigate the role of the DnaK and DnaJ proteins in cells lacking SecB, we used a secB::Tn5 strain in which all synthesis of DnaK and most synthesis of DnaJ was under control of the IPTG-inducible PlacUV5 operon and, in addition, carried the pAKL1 plasmid allowing IPTG-inducible expression of the dnaKJ operon controlled by PlacUV5 carried on pAKL1. At A450 = 0.2, cells were filtered to remove IPTG and resuspended in medium lacking IPTG for continued growth. A450 of the secB::Tn5 strain CAG13522 (∗) and the secB+ strain CAG13553 (○) is shown as a function of time after removal of IPTG.

Overproduction of DnaK and DnaJ partially substitutes for lack of SecB

Since we showed that DnaK and DnaJ were required for the viability of a secB null strain on minimal medium, we tested whether overproduction of DnaK and DnaJ would suppress the lethality of secB::Tn5 on rich medium. To achieve overproduction, we used a strain that retained the chromosomal copy of the dnaKJ operon and, in addition, carried the pAKL1 plasmid allowing IPTG-inducible expression of the dnaKJ genes. Upon induction of DnaK and DnaJ synthesis, the secB::Tn5 strain was able to grow in rich medium with 100% plating ef-
Figure 3. Processing of LamB and MBP in strains depleted for DnaK and DnaJ. The secB::Tn5 strain CAG13522 (A) and the isogenic secB+ strain CAG13553 (B), grown and depleted for DnaK and DnaJ as described in Fig. 2, were labeled with Tran35S-label for 30 sec and chased for 5 min with nonradioactive amino acids immediately before removal of IPTG and at 1, 2, 3, and 4 doublings (>6 doublings for secB+ strain) after removal of IPTG. Samples were processed to detect and quantify the precursor and mature forms of LamB, MBP, and OmpA.

ciency [Fig. 4], but with a growth rate slower than that of secB+ (data not shown). These results suggest that increased amounts of DnaK and DnaJ partially substitute for the function of SecB. In agreement with previous results, the rich medium growth defect of the secB::Tn5 strain was not relieved by overproduction of DnaK alone [Altman et al. 1991]. Suppression of rich medium growth defect of secb::Tn5 by the overproduction of DnaK and DnaJ was strain dependent and occurred in MC1061 (used throughout this study) and MG1655 [Bachmann 1987] but not in C600 [Bachmann 1987] or MC4100 [Casadaban 1976]. We have no clear explanation for the strain dependence we observe. However, because the secB null mutation is deleterious for growth, different strain backgrounds may accumulate different low-level suppressors. These suppressors may affect the ability of DnaK and DnaJ to compensate for the secB::Tn5 secretion defect.

Increased amounts of DnaK and DnaJ presumably permitted growth of the secB::Tn5 strain in rich medium because their overproduction compensated more effectively for the lack of SecB. In this case, overproduction of DnaK and DnaJ should result in an increase in the rate of export of at least some SecB-dependent proteins. The processing of two SecB-dependent proteins, LamB [Fig. 5A,B] and MBP [Fig. 6A–C] was facilitated by overproduction of DnaK and DnaJ, whereas that of OmpA was not [Fig. 7]. The time required to process half of the precursor to the mature form \( t_{1/2} \) was 3.6 min in secB::Tn5 cells; upon overproduction of DnaK and DnaJ, the \( t_{1/2} \) decreased to 2.4 min [Fig. 5A,B]. For pre-MBP, the processing kinetics were biphasic [Fig. 6A,B]. The second phase of the reaction \( t_{1/2} = 50 \text{ min} \) corresponded to the degradation of pre-MBP, as there was not a corresponding increase in the absolute amount of the mature form during this time period [Fig. 6A]. By subtracting the degradation component of the reaction, the processing component was observed directly [Fig. 6C]. In the case of MBP, overproduction of DnaK and DnaJ not only accelerated the initial rate of processing but also permitted it to proceed for a longer time. The net result is that almost twice as much MBP was exported when DnaK and DnaJ were overproduced [Fig. 6C]. Such a result is not surprising. If cells with higher levels of DnaK and DnaJ maintain pre-MBP in an export-competent state for a longer time than those with lower levels, effective processing can occur over a longer time frame. Thus, overproducing DnaK and DnaJ not only will affect the rate of export but also, and possibly more significantly, the amount of the precursor protein that is transported to its proper location.

Figure 4. Growth of a secB::Tn5 strain in rich medium after overproduction of DnaK and DnaJ. CAG13471 [secB::Tn5 pAKL1] grown overnight at 30°C in glycerol/maltose/Amp/IPTG medium was diluted 10-fold and plated on LB–Amp plates with [right] or without [left] 0.2 mM IPTG to induce the dnaKJ operon carried on pAKL1. Control platings indicate that the induced culture exhibited 100% plating efficiency, whereas the uninduced culture had a plating efficiency of \( <10^{-6} \).
Discussion

Our results establish that DnaK and DnaJ participate in the process of protein export in E. coli. Both proteins play a primary role in the export of AP and a secondary role in the export of some SecB-dependent proteins. These results complement previous studies that have established a role for Hsp70 proteins and homologs of DnaJ in several aspects of the secretion process in eukaryotic cells [De-Shaies et al. 1988; Sadler et al. 1989; Kang et al. 1990; Atencio et al. 1992]. Our results to date using the grpE280 mutant do not indicate a role for GrpE either in export of AP or in processing of SecB-dependent proteins. However, this strain still retains some GrpE function, as judged by the fact that deletion mutants of grpE are inviable [Fayet et al. 1989]. Therefore, additional approaches will be required to determine whether GrpE is also involved in protein export.

DnaK and DnaJ participate in the export of AP

The delay in processing pre-AP to the mature form in dnaK and dnaJ mutants implicates DnaK and DnaJ in AP export. DnaK and DnaJ could participate directly in AP secretion by being the chaperones that maintain the precursor protein in a transport-competent state. Although direct evidence will be required to establish whether DnaK and DnaJ interact with pre-AP, our results are consistent with this possibility. First, not all dnaK mutations affect AP export. Thus, impaired processing is unlikely to be a consequence of global changes in gene expression, such as increased expression of the HSPs, characteristic of all dnaK mutants examined to date. Second, the extent of the processing defect is allele specific, as would be expected if the mutant proteins affect the interaction of DnaK with AP in different ways. Finally, the processing defect is specific to AP and does not result from “jamming” the export machinery, as several other proteins [RBP, MBP, LamB, and OmpA] exhibited normal kinetics of export in the mutant strains [data not shown].

Figure 5. Processing of LamB upon overproduction of DnaK and DnaJ in a secB::Tn5 strain. secB::Tn5 and secB+ strains growing exponentially in glycerol/maltose/Amp/IPTG medium were pulse-labeled for 30 sec with Tran3S-label, chased with nonradioactive amino acids for the time periods indicated, and processed to detect and quantify precursor and mature LamB. [A] Conversion of pre-LamB to LamB. [Top] Disappearance of pre-LamB upon overproduction of DnaK and DnaJ [CAG13471 (secB::Tn5 pAKL1)]; [bottom] disappearance of pre-LamB without overproduction of DnaK and DnaJ [CAG13469 (secB::Tn5 pB10a)]. [B] The kinetics of pre-LamB processing. The fraction of pre-LamB protein present in CAG13471 [●] and CAG13469 [○] at each time point was calculated from the data in Fig. 5A and plotted as a function of the time of chase.

Figure 6. Processing of MBP upon overproduction of DnaK and DnaJ in a secB::Tn5 strain. Strains CAG13471 and CAG13469, grown and labeled as in Fig. 5, were processed to detect and quantify precursor (pre-MBP) and mature MBP. [A] Conversion of pre-MBP to MBP. [Top] Disappearance of pre-MBP upon overproduction of DnaK and DnaJ [CAG13471 (secB::Tn5 pAKL1)]; [bottom] disappearance of pre-MBP without overproduction of DnaK and DnaJ [CAG13469 (secB::Tn5 pB10a)]. [B] Kinetics of disappearance of pre-MBP. The fraction of pre-MBP present in CAG13471 [●] and CAG13469 [○] at each time point was calculated from the data in [A] and plotted as a function of the time of chase. The slow phase of the curves [t1/2 ~ 50 min] represents degradation of pre-MBP as there is no increase in the absolute amount of MBP. [C] Kinetics of processing of pre-MBP. To determine the processing component of the disappearance of pre-MBP at early times, the degradation component [determined by back-extrapolating the late time points in Fig. 6B] was subtracted. Disappearance of pre-MBP protein resulting only from processing in CAG13471 [●] and CAG13469 [○] is plotted as a function of the time of chase.
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Figure 7. Processing of OmpA in secB::Tn5 strain overproducing DnaK and DnaJ. Disappearance of pre-OmpA in strain CAG13471 [lanes 1–4] and CAG13469 [lanes 5–8] grown and labeled as in Fig. 5, and processed to detect and quantify pre-precursor [pre-OmpA] and mature OmpA.

The AP precursor in the cytoplasm may be a favored substrate for binding DnaK because it is more unfolded than other precursor proteins such as pre-MBP [Derman and Beckwith 1991]. The dominant-negative DnaK variants could be defective in releasing the bound AP precursors. Kusukawa et al. [1989] noted that export of AP is retarded in the secB::Tn5 mutant at 30°C but not at 42°C. Thus, at low temperature, when only basal levels of DnaK and DnaJ are present, SecB may participate in AP export as well. If DnaK does play a chaperone role, it will be interesting to determine whether DnaK, like SecB, helps target precursor proteins to the translocon or functions solely to maintain them in an export-competent conformation.

The role of DnaK and DnaJ in a strain lacking SecB

We have established a role for the DnaK and DnaJ proteins in the viability and protein export process of cells lacking SecB. In reciprocal experiments, we showed that overproduction of DnaK and DnaJ allowed a secB null strain to grow in rich medium and accelerated the export rates of MBP and LamB. Using a secB::Tn5 strain in which all expression of dnaK and most expression of dnaJ was driven by the IPTG-inducible lacUV5 promoter, we were able to show that depleting DnaK and DnaJ reduced export of MBP and LamB, impaired growth rate, and reduced the viability of these cells dramatically. No such defects in export or growth were observed in secB+ cells. Among other possibilities, the decrease in growth rate could be a consequence of “jamming” the protein export machinery several doublings after specific export defects first become evident. In this case, accumulation of pre-OmpA, observed only when the growth rate of the culture declined, could result from jamming the protein export machinery. Taken together, these data indicate that DnaK and DnaJ can partially substitute for several of the functions of SecB.

Our depletion experiments indicate that strains lacking SecB require elevated levels of DnaK and DnaJ to be able to grow in minimal medium. In these experiments, DnaK and DnaJ were initially supplied at a level sixfold higher than that in wild-type cells [data not shown]. Secretion defects were already observed two doublings after shut-off of DnaK and DnaJ synthesis when the amount of DnaK is still greater than that in wild-type strains. These findings are in accord with our observation that the amount of DnaK and DnaJ is about five- to sixfold higher in the secB::Tn5 mutant than in the isogenic wild-type strain [J. Wild, E. Altman, W. Walter, and C. Gross, in prep.].

Our experiments suggest an important role for DnaK and DnaJ in secB+ cells even though depleting DnaK and DnaJ had little effect on the export of SecB-dependent proteins in secB+ cells during steady-state growth. The high rate of HSP synthesis in a secB null strain indicates that the rate of synthesis of DnaK and DnaJ is responsive to the accumulation of precursor proteins. Most likely, the accumulated precursor proteins deplete DnaK and DnaJ from the cell, preventing them from carrying out their normal function of negatively regulating the synthesis of σ52. The consequent accumulation of σ52 results in increased expression of the HSPs. This points to a very interesting scenario. There is currently no evidence that the rate of synthesis of secB is sensitive to accumulation of precursor proteins. Thus, if SecB became limiting, as it would be, for example, immediately after massive induction of some transport systems, precursor proteins would accumulate and the cell could respond by overproducing DnaK and DnaJ to facilitate the transport of these proteins.

The role of DnaK and Hsp70 as regulators of the heat shock response has been widely studied [Lindquist 1986; Stone and Craig 1990, Straus et al. 1990]. There is increasing evidence that DnaK [together with DnaJ and GrpE] acts as a cellular thermometer by regulating the stability, synthesis, and activity of σ52 in response to the availability of its substrates (for review, see Craig and Gross 1991). Our studies on the involvement of DnaK in protein secretion suggests that it may actually have a much wider sensory role in the cell. The ability of the pool of DnaK and DnaJ to change in response to varying levels of its substrates may not only maintain a reasonable rate of protein secretion under diverse sets of conditions but may also allow a number of other cellular processes involving DnaK to respond to environmental change [e.g., flagella formation, proteolysis, and replication of the E. coli, λ, P1, and F plasmid DNAs [for review, see Georgopoulos et al. 1990; Gross et al. 1990]]. From this point of view, DnaK may be a master regulator that functions to coordinate the activity of a wide variety of proteins to maintain balanced growth in a fluctuating environment.

Materials and methods

Bacterial strains, plasmids, and genetic methods

All strains employed were derivatives of MC1061 [araD139 Δ[ara-lac]7679 ΔlacX74 galK galU mcrB hsdR rpsL thi] [Casadaban and Cohen 1980]. CAG13469 is secB::Tn5 pB10a, CAG13471 is secB::Tn5 pAKL1, CAG13553 is secB+ ΔdnaK1 thrTn10 pAKL1, CAG13522 is secB::Tn5 ΔdnaK1 thrTn10 pAKL1. The original secB::Tn5 insertion is MM152 [Kumamoto and Beckwith 1985]. Strains used in experiments on AP processing contained the phoS21 mutation [obtained as BW3890 from B. Wanner], which synthesizes AP constitutively.

The pB10a vector is a pBR322 derivative carrying the lacIq gene and the IPTG-inducible lacUV5 promoter, followed by the
XhoI and HindIII restriction endonuclease cloning sites. pNRK411 (kindly provided by N. Kusukawa) was derived by fusing 900 bp of the 5' end of dnaK to placUV5 on pBlOa. pAKL1 is a pBl0a derivative carrying the promoterless dnaKJ operon fused to the IPTG-inducible lacUV5 promoter. It was constructed by subcloning an EcoRI fragment of pW4 containing the 5' end of dnaK and the entire dnaJ gene [Wild et al. 1992] into pNRK411. In pAKL1, all synthesis of dnaK is under control of PlacUV5; however, a low level of the dnaJ transcription originates from a weak, internal promoter (Ohki et al. 1986).

Cells were grown in LB or in M9 minimal medium supplemented with 0.5% glycerol and all amino acids except L-methionine and L-cysteine. When indicated, glycerol medium was supplemented with 0.2% maltose, 0.2 mM IPTG, and 50 μg/ml of ampicillin [Amp]. Transformations and phase P1 transductions were as described (Silhavy et al. 1984). The dnaJ and dnaK mutations were isolated by selecting for mutants exhibiting elevated HSP synthesis [Wild et al. 1992] and are designated with a one-letter code. The first letter denotes the amino acid in the wild-type strain, the second the mutational change, and the number indicates the location of the change within the protein.

Depletion of DnaK and DnaJ

CAG13553 [secB+ ΔdnaKJ thrTn10 pAKL1] and CAG13522 [secB−:Tn5 ΔdnaKJ thrTn10 pAKL1] were grown at 30°C to A600 = 0.2 in glycerol medium supplemented with maltose, IPTG, and Amp. To shut off the induced synthesis of DnaK and DnaJ from the lacUV5 promoter that drives expression of the dnaKdnaJ operon on pAKL1, IPTG was removed by filtering the cultures onto sterile nitrocellulose filters and washing the cells with 15 volumes of glycerol medium. Washed cells were resuspended in 10 ml of prewarmed glycerol medium supplemented with maltose and Amp and grown at 30°C with aeration. Samples were withdrawn periodically for growth measurements and labeling. Immunoprecipitations with DnaK antisera indicated that no further DnaK synthesis was detectable after removal of IPTG. Preexisting DnaK and DnaJ were diluted by growth in the absence of IPTG.

Radiolabeling, immunoprecipitation, and electrophoresis

One-milliliter samples of cells growing exponentially at 30°C in glycerol medium were pulse-labeled with 60 μCi/ml of Tran35S-label (ICN; 1000 Ci/m mole), chased with a nonradioactive mixture of l-methionine and l-cysteine (200 μg/ml), and precipitated with TCA as described (Grossman et al. 1987). The precursor and mature forms of LamB, MBP, OmpA, AP, and RBP were immunoprecipitated [Ito et al. 1981], resolved by electrophoresis on 10% SDS-polyacrylamide gels [Laemmli 1970], detected by autoradiography, and quantified with an Ambis Scanner interfaced with an IBM computer. Percent exported protein was calculated as mature protein/(mature + precursor protein). Antisera against AP were obtained from 5 Prime–3 Prime, Inc. (West Chester, PA).

Acknowledgments

We thank B. Wanner for supplying strain BW3890, N. Kusukawa for pNRK411, and E. Craig, H. Echols, A. Kamath, and K. Ito for comments on the manuscript. Antisera against OmpA, RBP, and MBP were kindly provided by J. Beckwith. This work was supported by U.S. Public Health Research grant GM36278 from the National Institutes of Health to C.A.G.

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Genes Dev. 1992, 6:
Access the most recent version at doi:10.1101/gad.6.7.1165

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