

RESEARCH COMMUNICATION

Latent ClpX-recognition signals ensure LexA destruction after DNA damage

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The DNA-damage response genes in bacteria are up-regulated when LexA repressor undergoes autocatalytic cleavage stimulated by activated RecA protein. Intact LexA is stable to intracellular degradation but its auto-cleavage fragments are degraded rapidly. Here, both fragments of LexA are shown to be substrates for the ClpXP protease. ClpXP recognizes these fragments using sequence motifs that flank the auto-cleavage site but are dormant in intact LexA. Furthermore, ClpXP degradation of the LexA-DNA-binding fragment is important to cell survival after DNA damage. These results demonstrate how one protein-processing event can activate latent protease recognition signals, triggering a cascade of protein turnover in response to environmental stress.

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Protein degradation plays a critical role in allowing cells to adjust to changing conditions. Intracellular proteolysis is an essential component of many response pathways that permit bacteria to survive or recover from DNA damage, heat shock, or attack by reactive oxygen species (Mizusawa and Gottesman 1983; Gerth et al. 1998; Roberston et al. 2002). In addition to stress-related regulatory functions, proteases serve to degrade damaged proteins as well as undamaged proteins that the cell no longer needs under a new set of environmental conditions. In each of these examples, one can ask the same question: what mechanisms allow intracellular proteases to degrade the appropriate substrates at the right time?

In bacteria, many intracellular proteases (e.g., ClpXP, ClpAP, and HslUV) divide the tasks of substrate recognition and proteolysis between a hexameric AAA⁺ ATPase and a multisubunit peptidase with an internal degradation chamber (for review, see Gottesman 1996). The ATPase (ClpX, ClpA, or HslU) recognizes specific substrates, unfolds these proteins, and translocates the denatured polypeptide into the peptidase (ClpP or HslV) chamber for degradation (Kim et al. 2000). In many instances, the ATPase components recognize substrates via specific peptide motifs, often at their N or C termini

(Levchenko et al. 1995; Gonciarz-Swiatek et al. 1999; Flynn et al. 2003). For example, the *ssrA* degradation tag, an 11-residue C-terminal peptide sequence, targets proteins for degradation by ClpXP and ClpAP (Gottesman et al. 1998). For some substrates, such as the stationary-phase sigma factor (σ^S), accessory proteins are needed for efficient degradation, allowing regulation of proteolysis through the synthesis or modification of these factors (Becker et al. 1999; Zhou et al. 2001). Here we provide evidence for an alternative mechanism of regulated degradation: We find that complete degradation of the LexA repressor requires an initiating cleavage event that is regulated in response to DNA damage.

The SOS regulatory system in *Escherichia coli* controls the cellular response to DNA damage. Under normal conditions, LexA repressor dimers negatively regulate the expression of genes involved in DNA repair, replication, and cell division (for review, see Freidberg et al. 1995). After DNA damage, single-stranded DNA is exposed and is bound by the RecA protein. This event activates RecA, which then stimulates the auto-cleavage of LexA at a site between its N-terminal DNA-binding domain and its C-terminal dimerization domain (Philzicky and Roberts 1981; Little 1984). The resulting N- and C-terminal fragments are then degraded rapidly (Little 1983a). The C-terminal fragment is stabilized ~10-fold in Lon-defective cells (Little 1983b), but the protease that degrades the N-terminal fragment has not previously been identified.

In recent proteomic experiments, we identified tryptic fragments from *E. coli* proteins that were trapped in vivo within a proteolytically inactive ClpP variant (ClpP^{trap}) and found that some peptides originated from LexA (Flynn et al. 2003). Western blots of the undigested trapped proteins indicated that the two auto-cleavage fragments of LexA represented the dominant captured forms. This result suggested that these auto-cleavage fragments might be ClpXP substrates. Here, we show that ClpXP degrades the auto-cleavage fragments of LexA, but not full-length LexA, both in vivo and in vitro. Recognition of these fragments for destruction occurs via peptide signals created or exposed by the initial auto-cleavage event. The use of these latent recognition signals allows specific recognition and degradation of the LexA fragments by ClpXP at the biologically appropriate time—after LexA has undergone RecA-stimulated self-cleavage in response to DNA damage.

Results and Discussion

The LexA auto-cleavage fragments are ClpXP substrates

Capture experiments using ClpP^{trap} were performed in *E. coli* exposed to the DNA-damaging agent nalidixic acid. The ClpP^{trap} is an inactive (S97A) affinity-tagged variant of ClpP that, when expressed in cells, allows the capture and purification of physiological ClpXP substrates (Flynn et al. 2003). A Western blot of the trapped material revealed two bands that cross-reacted with anti-LexA antibodies and had the same electrophoretic mobilities as the LexA^{1–84} and LexA^{85–202} fragments gener-

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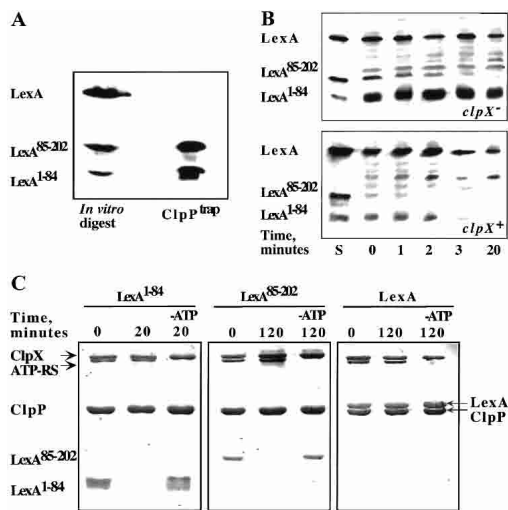


Figure 1. LexA auto-cleavage fragments are captured and degraded by ClpXP. (A) Western blot of the LexA species captured by the ClpP^{trap} and generated by auto-cleavage in vitro. (B) The proteolytic stability of LexA, LexA¹⁻⁸⁴, and LexA⁸⁵⁻²⁰² in vivo was determined in *clpX*⁻ (MC4100) and *clpX*⁺ (SG22101) strains bearing plasmid pJL42. LexA, LexA¹⁻⁸⁴, and LexA⁸⁵⁻²⁰² were identified by comparison with auto-cleaved LexA (lane S). It is difficult to see the continued disappearance of LexA, as expected from auto-processing, because the Western blots are mildly overexposed to allow observation of the low levels of LexA⁸⁵⁻²⁰². (C) ClpXP degradation of purified LexA, LexA¹⁻⁸⁴, and LexA⁸⁵⁻²⁰² in vitro. The band marked ATP-RS is creatine kinase.

ated by auto-cleavage of purified LexA (Fig. 1A). No full-length LexA was detected in the trapped material, although we observed ~20 ng of each cleavage fragment and could detect as little as 1 ng of LexA protein. Furthermore, when purified LexA was added to a sample of the trapped material and prepared for Western blotting, this protein was efficiently detected, indicating that full-length LexA could have survived this procedure. Trapping required ClpX, as the LexA fragments were not detected in material isolated from *clpX*⁻ cells (Flynn et al. 2003). We conclude that ClpX recognizes the two auto-cleavage fragments but not full-length LexA.

To determine the stability of the various forms of LexA in vivo, we induced the DNA-damage response with nalidixic acid, blocked protein synthesis, and measured the half-lives of intact LexA and the two cleavage fragments. The N-terminal LexA¹⁻⁸⁴ fragment was rapidly degraded in a wild-type strain (half-life 2–5 min) but was not degraded to any detectable extent in an otherwise isogenic *clpX*⁻ strain (Fig. 1B). The C-terminal LexA⁸⁵⁻²⁰² fragment was also unstable in wild-type cells (half-life ~1 min) but was only stabilized modestly in the *clpX*⁻ cells (half-life ~2 min), presumably because Lon protease also contributes to its degradation (Little 1983b). Full-length LexA appeared to be reasonably stable to degradation in the wild-type and *clpX*⁻ strains, although it was susceptible to continued auto-cleavage (Fig. 1B).

To determine if the LexA fragments were indeed

ClpXP substrates, we assayed for degradation in vitro. Purified LexA, LexA¹⁻⁸⁴, and LexA⁸⁵⁻²⁰² were incubated with ClpXP and an ATP-regenerating system, and degradation was assayed by SDS-PAGE (Fig. 1C). ClpXP degraded both fragments but not LexA in ATP-dependent reactions. Hence, both auto-cleavage fragments of LexA are ClpXP substrates, whereas the full-length protein is not.

The new C-terminal sequence of LexA¹⁻⁸⁴ targets it to ClpXP

RecA-stimulated auto-cleavage of LexA creates a Val 82–Ala 83–Ala 84–COOH sequence at the end of the N-terminal fragment, which is very similar to the Leu–Ala–Ala–COOH sequence at the end of the *ssrA* degradation tag (Tu et al. 1995; Keiler et al. 1996). These residues of the *ssrA* tag are the principal determinants of recognition by ClpX (Flynn et al. 2001). To test the importance of the C-terminal residues of LexA¹⁻⁸⁴ for ClpXP degradation, we constructed a variant (LexA^{1-84DD}) with aspartic acids replacing both alanines. In ClpXP degradation assays in vitro (Fig. 2A), LexA^{1-84DD} was not degraded to any detectable extent, whereas LexA¹⁻⁸⁴ was degraded with a half-life of ~10 min. Thus, these alanines play a major role in targeting the N-terminal LexA fragment for degradation by ClpXP.

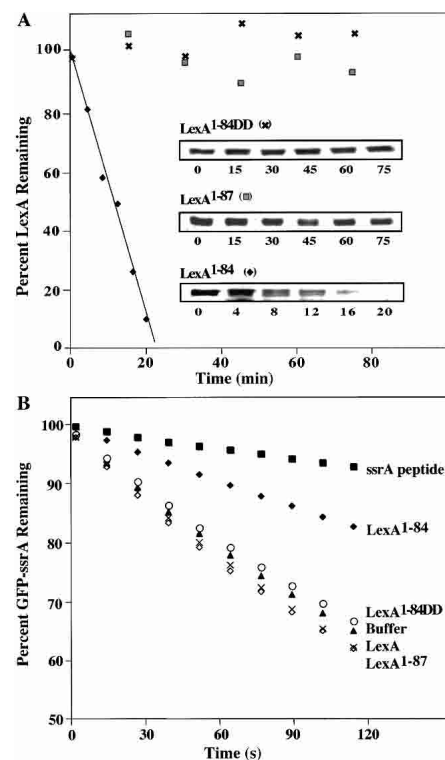


Figure 2. The new C terminus of LexA¹⁻⁸⁴ directs its degradation. (A) A time course of ClpXP degradation of LexA¹⁻⁸⁴, LexA¹⁻⁸⁷, and LexA^{1-84DD} was quantified after SDS-PAGE. (B) ClpXP degradation of GFP-*ssrA* was measured by loss of fluorescence at 511 nm in the presence of various forms of LexA at 20 μM. An effective inhibitor of GFP-*ssrA* degradation, the *ssrA* peptide (20 μM), is included for comparison.

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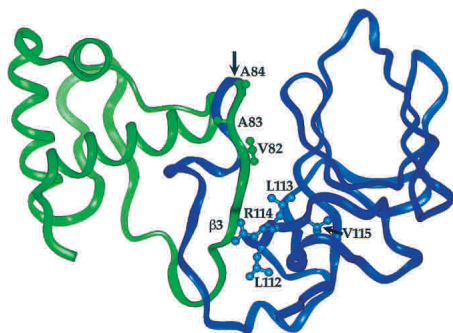


Figure 3. Location of latent ClpX-recognition signals in the structure of full-length LexA. The N-terminal domain is green and the C-terminal domain is blue. The proposed ClpX-recognition site in the C-terminal domain is labeled. The cleavage site is indicated by an arrow. (Structure from Luo et al. 2001.)

To test whether these alanines need to be at the C terminus for recognition, we constructed and purified a variant, LexA¹⁻⁸⁷, extended by the next three residues (Gly 85–Glu 86–Pro 87) of intact LexA. ClpXP did not degrade this variant (Fig. 2A), revealing that the Val 82–Ala 83–Ala 84 sequence must be at the C-terminal end for efficient degradation. LexA¹⁻⁸⁴ also inhibited degradation of another ClpXP substrate, GFP bearing an *ssrA* degradation tag, suggesting that LexA¹⁻⁸⁴ and *ssrA*-tagged proteins compete for ClpX recognition or other processes involved in degradation (Fig. 2B). In contrast, LexA^{1-84DD}, LexA¹⁻⁸⁷, and full-length LexA failed to inhibit GFP-*ssrA* degradation. Together, these results demonstrate that LexA¹⁻⁸⁴ carries a functional ClpX-recognition signal, whereas LexA^{1-84DD}, LexA¹⁻⁸⁷, and the full-length protein lack an accessible or functional signal.

The Val 82–Ala 83–Ala 84 sequence is within a folded region in the LexA crystal structure, and the side chain of Val 82 is buried and packs against residues in the C-terminal domain (Luo et al. 2001; Fig. 3). As a consequence, it seems likely that auto-cleavage results in exposure and increased flexibility of the Val 82–Ala 83–Ala 84 sequence, allowing unimpeded interactions with ClpX. However, as the LexA¹⁻⁸⁷ fragment is not a ClpXP substrate, improved accessibility of ClpX to the Val 82–Ala 83–Ala 84 sequence is not sufficient to explain how auto-cleavage activates proteolysis of this domain. Based on peptide studies, the α -COOH group of the *ssrA* degradation tag has been shown to be an important determinant for ClpX recognition (Kim et al. 2000). The resistance of LexA¹⁻⁸⁷ to degradation strongly suggests that the α -COOH group of Ala 84 is similarly important for recognition of LexA¹⁻⁸⁴. Thus, our results indicate that auto-cleavage of the LexA Ala 84–Gly 85 peptide bond directly creates an essential portion of the degradation signal for the resulting N-terminal fragment.

Sequences that target the C-terminal LexA fragment to ClpXP

Although most LexA⁸⁵⁻²⁰² molecules seem to be degraded by Lon protease in vivo (Little 1983b), our experi-

ments show that LexA⁸⁵⁻²⁰² is also degraded by ClpXP (Fig. 1B,C). To search for peptide sequences within LexA⁸⁵⁻²⁰² that might mediate ClpX recognition, we tested which regions of this fragment bound to ClpX (Fig. 4A). We prepared a covalent array of synthetic 12-residue peptides of the LexA⁸⁵⁻²⁰² fragment, each sharing a 10-residue overlap with its sequence neighbors. The peptide filter was incubated with ClpX and washed, and bound ClpX was detected using an anti-ClpX antibody. Peptides corresponding to the first 26 amino acids of LexA⁸⁵⁻²⁰² bound ClpX poorly. In contrast, a cluster of adjacent peptides containing sequences from 103 to 126 bound ClpX reasonably well. Hence, this region of the LexA sequence may contain a ClpX-recognition signal.

To investigate the function of this sequence, we constructed two fusion proteins consisting of LexA residues 85–103 or 85–126 attached to the Arc-st11 protein, a variant of the Arc repressor with a stabilizing C-terminal sequence (Milla et al. 1993). ClpXP degraded the LexA⁸⁵⁻¹²⁶ fusion protein with a half-life (50 ± 9 min) similar to the half-life for LexA⁸⁵⁻²⁰² (60 ± 3 min; Fig. 4B). This result shows that sequence information between residues 85 and 126 of LexA is sufficient to target a protein to ClpXP for degradation. In contrast, the LexA⁸⁵⁻¹⁰³ fusion protein was degraded substantially

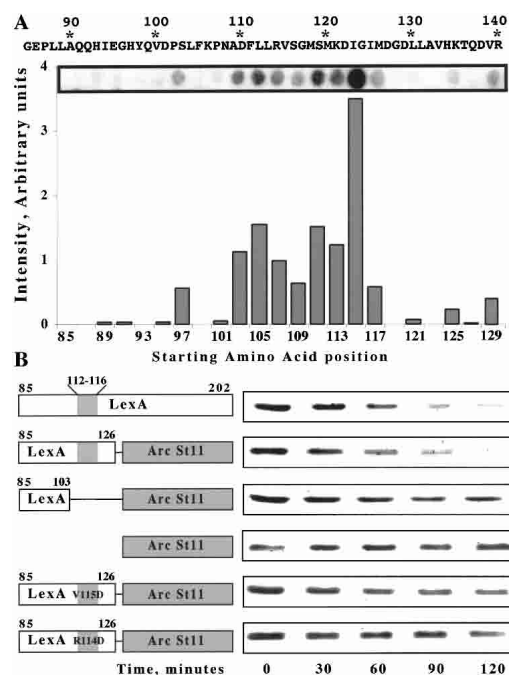


Figure 4. Identification of residues in LexA⁸⁵⁻²⁰² important for ClpXP recognition. (A) An array of LexA peptides was incubated with purified ClpX protein, and bound ClpX was detected using anti-ClpX antibody. The chart shows the relative intensity of ClpX binding to individual sequences. The starting amino acid positions of the 12-residue peptides are listed below the chart. The signal intensity of the peptide beginning at residue 115 may be artificially high. In other experiments, we have observed high signals from different peptides with the same three initial residues (VSG). (B) Degradation of LexA⁸⁵⁻²⁰² and LexA-Arc-st11 fusion proteins by ClpXP in vitro.

more slowly ($t_{1/2} = 184 \pm 24$ min), indicating that the most important determinants for ClpXP degradation are located between residues 104 and 126, and not between residues 85 and 103.

The 104–126 region contains a sequence (Leu 112–Leu 113–Arg 114–Val 115–Ser 116) with some similarity to a peptide motif proposed to function as a ClpX-recognition sequence (Flynn et al. 2003). To test the importance of the LexA sequence motif, we constructed LexA^{85–126} fusion proteins containing the Arg 114 → Asp or Val 115 → Asp mutations and found that both slowed ClpXP degradation ($t_{1/2} = 118 \pm 22$ and 122 ± 30 min, respectively; Fig. 4B). Taken together, these data support the conclusion that ClpX recognizes a region of the polypeptide chain about 30 residues distal from the new N terminus of LexA^{85–202}. Other sequences, however, may also contribute to ClpX recognition of LexA^{85–202}.

Unlike LexA^{1–84}, the ClpX-targeting signal on the C-terminal domain is not directly adjacent to the site of auto-cleavage. To account for the fact that intact LexA, which contains the same peptide sequences as LexA^{85–202}, does not interact with ClpX, we propose that auto-cleavage disrupts the structure of the C-terminal domain in some fashion, helping to expose the 112–116 peptide signal. Although we do not know the structure of the isolated C-terminal domain, it is reasonable to propose that the 112–116 region will be more flexible and exposed after cleavage. In the structure of the full-length protein, a β -strand (β 3) from the N-terminal domain forms an integral part of the C-terminal domain and contacts the proposed recognition signal. The loss of these contacts following cleavage may result in exposure of this signal to ClpXP (Fig. 3).

ClpX helps cells survive DNA damage

To test the importance of ClpX in the overall cellular response to DNA damage, we assayed cell survival after exposure to increasing doses of UV irradiation (Fig. 5A). These experiments showed that *clpX*⁻ cells were more sensitive to UV irradiation than their wild-type counterparts, with the effect being most pronounced at the highest UV doses. When exposed to a UV dose of 200 J/m², *clpX*⁻ cells had a roughly 10-fold lower survival frequency than wild-type cells. Because the N-terminal DNA-binding domain of LexA can function as a repressor on its own (Schmidt-Dorr et al. 1991), one interpretation of this result is that ClpXP degradation of this fragment may be required to allow maximal expression of one or more DNA-damage inducible gene products, thereby improving survival following near-lethal UV doses. Alternatively, the loss of ClpX could perturb the levels of proteins other than the LexA^{1–84}, thus affecting survival.

To determine if excess levels of LexA^{1–84} causes increased UV sensitivity, we tested the effect of expressing LexA^{1–84} and LexA^{1–84DD} on the percentage of cells surviving a fixed UV dose. We assayed survival of wild-type cells containing plasmid-borne genes for either LexA^{1–84} or LexA^{1–84DD} under the control of the native LexA pro-

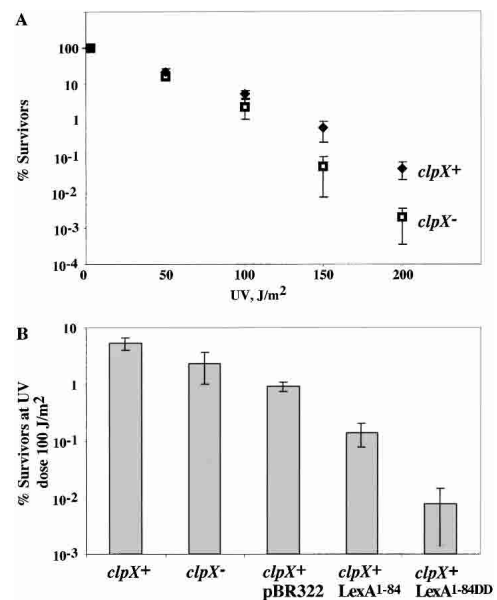


Figure 5. Cells accumulating LexA^{1–84} are UV sensitive. (A) Survival after UV irradiation of *clpX*⁺ (MC4100) and *clpX*⁻ (SG22101) cells. The percentage of surviving cells is plotted against UV dose. Each point is the average (\pm S.D.) of three trials. (B) Survival of cells containing the empty parent vector or plasmids directing expression of LexA^{1–84} or LexA^{1–84DD} was compared at a UV dose of 100 J/m².

moter (Fig. 5B); Western analysis revealed that these fragments were modestly overexpressed (two- to fourfold, with LexA^{1–84DD} accumulating to higher levels than LexA^{1–84}) compared with the chromosomal-encoded LexA (data not shown). Cells expressing LexA^{1–84DD} had ~10-fold lower survival than those with the LexA^{1–84}-producing plasmid. Furthermore, both strains were more sensitive to UV irradiation than cells bearing the empty plasmid. Thus, accumulation of the LexA DNA-binding domain is deleterious to cell survival after DNA damage.

These data reveal that timely destruction of LexA^{1–84} following damage-induced auto-cleavage is likely to play an important biological role. We suggest that failure to degrade the LexA^{1–84} fragment following DNA damage results in the retention of some repressor activity and a consequent failure to fully induce one or more DNA-damage genes. This residual repressor activity, in turn, may account for our finding that ClpX improves bacterial survival following near-lethal UV doses. Our observation that cells expressing even modest levels of the nondegradable LexA^{1–84DD} are more sensitive to UV irradiation than those expressing LexA^{1–84} supports this theory, although we cannot currently rule out more indirect explanations. In preliminary work, we did not detect reproducible differences in expression of the LexA-controlled *sulA*, *recA*, and *uvrA* genes following DNA damage in wild-type versus ClpX-defective cells (data not shown), but there are more than 20 genes under LexA control and some may be more sensitive than others to repression mediated by the LexA^{1–84} fragment.

Changes in environmental conditions often result in

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the modification of regulatory proteins by phosphorylation or proteolytic cleavage. Similarly, protein-binding partners often change during the progression of a biochemical pathway. The results presented here show that ClpXP degrades the two LexA cleavage fragments but not the intact protein. By mapping recognition signals in LexA¹⁻⁸⁴ and LexA⁸⁵⁻²⁰², we show that auto-cleavage activates otherwise dormant protease recognition signals in both fragments. Our analysis of the mechanisms used to target LexA to ClpXP thus highlights how changes in protein primary or tertiary structure can be coupled to the exposure of recognition signals for a destructive protease, thereby triggering a cascade of protein turnover in response to environmental change.

LexA is a member of a protein family that includes several phage repressors that also undergo RecA-mediated auto-cleavage (Sauer et al. 1982; Little 1984; Eguchi et al. 1988). It will be interesting to see if ClpXP plays a role in degrading the auto-cleavage fragments of these proteins. The fragments of unrelated proteins that undergo proteolytic processing events may also be targeted for degradation by a similar mechanism. For example, RseA is an antisigma factor that is sequentially cleaved by two membrane proteases in response to periplasmic stress (Alba et al. 2002). The N-terminal cytoplasmic domain of RseA released by cleavage is captured by ClpXP^{trap} (Flynn et al. 2003) and has been proposed to contain a Val-Ala-Ala-COOH sequence identical to the signal that targets LexA¹⁻⁸⁴ for ClpXP degradation (Alba et al. 2002). Thus, we suspect that dormant degradation signals like those we have characterized in LexA will be used to couple the destruction of other proteins to changes that occur in the intracellular or extracellular environment.

Materials and methods

Strains and plasmids

Strains MC4100 and SG22101 (MC4100 *clpX::kan*) were used in UV survival assays. The pBR322-based plasmid pJWL42 contains wild-type *lexA* under the control of its native promoter (Markham et al. 1981). Plasmid pJWL228 was a construct for overexpression of LexA and consisted of wild-type *lexA* under the control of the T7 promoter (Shepley and Little 1996). Plasmids pSBN15 and pSBN24 were used for overexpression of LexA¹⁻⁸⁴ and LexA¹⁻⁸⁷, respectively and were derivatives of pJWL228. Stop codons were inserted in the coding sequence after LexA residue 84 or 87, respectively, using a Quickchange kit (Stratagene). Plasmid pSBN16 was derived from pJWL228 for overexpression of the N-terminal fragment of LexA with aspartates replacing the two terminal alanines (LexA^{1-84DD}) using Quickchange. The plasmid for the expression of LexA⁸⁵⁻²⁰² (pSBN17) was constructed by PCR amplification of the sequence coding for amino acids 85–202 from pJWL228. The resulting PCR product was digested with *Bam*HI and *Nde*I and ligated into *Bam*HI/*Nde*I-digested pET11a. Plasmid pSBN19 was created for the expression of a fusion of LexA amino acids 85–126 to Arc-st11 (LexA⁸⁵⁻¹²⁶-Arc). It was made by amplifying the region of LexA coding for amino acids 85–126, digesting with *Nde*I and *Nhe*I, and ligating into *Nde*I/*Nhe*I-digested pET11a-arc-st11 (Flynn et al. 2003). The resulting N-terminal

sequence was *MGEPLLAQQHIEGHYQVDP**SLFKPNADFLLRVSGMSMKDIGIMASMGK* (LexA in italics, Arc in bold). Plasmid pSBN20 consisted of LexA residues 85–103 fused to Arc-st11 using an oligonucleotide cassette. The N-terminal sequence of this fusion was *MGEPLLAQQHIEGHYQVDP**SMGK*. Plasmids pSBN21 and pSBN22 were R114D and V115D variants, respectively, of the LexA portion of pSBN19 and were constructed using the Quickchange kit. Plasmids pSBN2 and pSBN23 are variants of pJL42 for the expression of LexA¹⁻⁸⁴ and LexA^{1-84DD}, respectively, under the control of the native promoter and were constructed as per pSBN15 and pSBN16. All constructs were verified by DNA sequencing.

Proteins/peptide array

Purification of GFP-ssrA, Arc-st11, ClpX (Flynn et al. 2003 and references therein), and ClpP (Kim et al. 2000) have been described. LexA⁸⁵⁻²⁰², LexA, LexA¹⁻⁸⁴, and LexA^{1-84DD} were purified from overproducing strains essentially as described for LexA (Little et al. 1994). A LexA peptide array was prepared by the MIT biopolymers facility using an Abimed instrument. Peptides interacting with ClpX were detected as described (Flynn et al. 2003).

Trapping/detection of LexA

Trapping was carried out essentially as described using strain JF162 (Flynn et al. 2003) except that cultures were treated with 50 µg/mL of nalidixic acid for 2 h before expression of the ClpP^{trap} was induced. ClpP cross-reacts with the LexA antibody and therefore was removed before Western blotting. Samples were dialyzed against 8 M urea for 6 h, then incubated with Ni-NTA beads (Quiagen) with rocking for 1 h. Supernatant was removed, dialyzed against LexA storage buffer, concentrated using a centricon YM-3 concentrator (Microcon), and used for Western blotting (see following). In a control experiment, this treatment did not affect our ability to detect full-length LexA, indicating that the protein is stable under these conditions.

Degradation in vitro and in vivo

For in vitro degradation, ClpX₆ (0.3 µM), ClpP₁₄ (0.8 µM), and an ATP-regenerating system (4 mM ATP, 50 µg/mL creatine kinase, 2.5 mM creatine phosphate) were incubated in buffer NB (50 mM Tris-HCl, 100 mM KCl, 10 mM MgCl₂, 1 mM DTT) at 30°C for 2 min. Substrate (10 µM) was added, and samples were removed at specific times, added to SDS loading buffer, and frozen on liquid nitrogen. After heating at 100°C for 5 min, samples were analyzed by 15% (N-LexA, C-LexA, Arc fusions) or 12.5% (LexA) SDS-PAGE. Gels were visualized by staining with Sypro orange (Molecular Probes) and scanning on a Molecular Dynamics Fluorimager 595. Imagequant (Molecular Dynamics) was used to quantify degradation. Inhibition of GFP-ssrA degradation was measured as described using 0.5 µM GFP-ssrA and 20 µM inhibitor (Flynn et al. 2001).

For in vivo degradation, samples were prepared from MC4100 and SG22101 cells containing plasmid pJL42. Cultures at an OD₆₀₀ of 0.2 were treated with 50 µg/mL nalidixic acid (Sigma) and incubated with shaking at 37°C for 30 min, then chloramphenicol (Sigma) was added to 100 µg/mL to stop protein synthesis. For each time point, cells were harvested by centrifugation, resuspended in SDS loading buffer, and rapidly frozen on liquid nitrogen. Cells were lysed by boiling and centrifuged in a microfuge at top speed for 5 min, and the resulting extract was separated by 15% SDS-PAGE. Western blots were performed with an ECF Western blotting kit in accordance with the manu-

facturer's guidelines (Amersham) using rabbit polyclonal LexA antibody at a 1:5000 dilution (John Little). Blots were imaged using a Molecular Dynamics Fluorimager 595. LexA fragments were prepared from purified LexA as described (Little 1984).

UV survival assays

Cells were grown to an OD₆₀₀ of 0.5 in Luria broth (LB), gently pelleted, washed once, and then resuspended in an equal volume of 0.85% saline. Suspensions were irradiated at a UV dose of 1.5 J/m²/sec for set times using a 15W G15T8 germicidal lamp (GE). The UV intensity was measured using a UVX radiometer (UVP). Appropriate dilutions were plated on LB agar plates (or LB agar plus 100 µg/mL ampicillin, as appropriate) and colonies were counted after 24 h. The decreased plating efficiency of cells carrying the pBR322 plasmid, compared with the nonplasmid-containing cells, was attributed to the need to maintain selection for the plasmid, based on parallel platings on media lacking antibiotic. Each assay was done at least in triplicate.

Acknowledgments

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Latent ClpX-recognition signals ensure LexA destruction after DNA damage

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