Expression and structure of the Chlamydia trachomatis DksA ortholog

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One sentence summary: Expression of the protein DksA in Chlamydia trachomatis is regulated with the bacterium’s developmental cycle. Ectopic expression of DksA affects generation of infectious bacterial progeny.

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Abstract

Chlamydia trachomatis is a bacterial obligate intracellular parasite and a significant cause of human disease, including sexually transmitted infections and trachoma. The bacterial RNA polymerase–binding protein DksA is a transcription factor integral to the multicomponent bacterial stress response pathway known as the stringent response. The genome of C. trachomatis encodes a DksA ortholog (DksA_Ct) that is maximally expressed at 15–20 h post infection, a time frame correlating with the onset of transition between the replicative reticulate body (RB) and infectious elementary body (EB) forms of the pathogen. Ectopic overexpression of DksA_Ct in C. trachomatis prior to RB→EB transitions during infection of HeLa cells resulted in a 39.3% reduction in overall replication (yield) and a 49.6% reduction in recovered EBs. While the overall domain organization of DksA_Ct is similar to the DksA ortholog of Escherichia coli (DksA_Ec), DksA_Ct did not functionally complement DksA_Ec. Transcription of dksA_Ct is regulated by tandem promoters, one of which also controls expression of nrdR, encoding a negative regulator of deoxyribonucleotide biosynthesis. The phenotype resulting from ectopic expression of DksA_Ct and the correlation between dksA_Ct and nrdR expression is consistent with a role for DksA_Ct in the C. trachomatis developmental cycle.

Keywords: Chlamydia, transcription, riboswitch, morphological transition

Introduction

Chlamydia trachomatis is a human-adapted bacterial obligate intracellular parasite and different pathotypes (aka serovars) of the pathogen cause distinct diseases. Urogenital serovars of C. trachomatis are a major cause of sexually transmitted infections (Stamm 1999, Rowley et al. 2012), while ocular serovars can cause trachoma (Burton and Mabey 2009), a leading cause of preventable blindness most prevalent in developing areas with poor sanitation. During infection of cultured host cells, C. trachomatis transitions between two physiologically distinct cell forms: the infectious but non-replicative elementary body (EB) and the replicative but noninfectious reticulate body (RB) (Shaw et al. 2000). While transition between cell forms is critical for replication and generation of infectious progeny, little is known about the trigger(s) or regulatory mechanism(s) that influence and control chlamydial morphological differentiation. Temporally restricted expression of different classes of genes during the chlamydial developmental cycle (Shaw et al. 2000, Belland et al. 2003) and recent genetic analysis identifying several genes with apparent functions in chlamydial development (Brothwell et al. 2016) suggest that EB→RB→EB transitions can occur in response to various stimuli and that response mechanisms controlling chlamydial development are redundant.

Chlamydia trachomatis has undergone genome reduction at the expense of reduced metabolic capacity (Stephens 1998). As a result, numerous essential nutrients are scavenged from the host cell during infection to allow for pathogen replication. In addition to biosynthetic deficiencies, certain regulatory pathways appear reduced to a single component of the canonical machinery. Consequently, C. trachomatis may have evolved to use these single components or ‘orphan regulators’ in place of certain canonical regulatory cascades.

In several bacteria, including Escherichia coli (Magnusson et al. 2007, Vinella et al. 2012, Kessler et al. 2017), Pseudomonas aeruginosa (Perron et al. 2005) and Legionella pneumophila (Dalebroux et al. 2010), DksA is a relatively small (<160 residue) RNA polymerase (RNAS)-binding protein and global transcriptional regulator that integrates responses to diverse stimuli, including nutrient availability (Paul et al. 2005), via a multicomponent response mechanism referred to as the stringent response (Cashel and Gallant 1969, Cashel et al. 1996, Potrykus and Cashel 2008, Traxler et al. 2008). The C. trachomatis genome encodes a DksA ortholog,
Results and discussion

Native expression of DksA<sub>Ct</sub> is maximal at the intermediate-late stages of the chlamydial developmental cycle

During infection of HeLa cells, C. trachomatis L2 undergoes a predictable developmental cycle in which log phase RBs initiate transition to the EB form approximately 18 h post infection (hpi) (Grieshaber et al. 2018). To determine whether DksA<sub>Ct</sub> expression is temporally regulated, expression of the native protein was determined with RB–EB transitions in the chlamydial developmental cycle. A similar expression pattern was previously reported for DksA in C. pneumoniae (Mukhopadhyay et al. 2006), indicating that DksA expression profiles are conserved between pathogenic Chlamydia species. DksA<sub>Ct</sub> was detected at moderately reduced levels at 30 and 45 hpi. Western blot analysis of EBs purified by density gradient centrifugation after sonication confirmed expression of DksA<sub>Ct</sub> in this cell form (data not shown). The dynamic expression pattern of DksA<sub>Ct</sub> is unique compared with the intermediate-late stages of the chlamydial developmental cycle (Grieshaber et al. 2018). To determine whether the RB–EB transition reduces both replication and production of infectious progeny

Overexpression of DksA<sub>Ct</sub> to RB–EB transition reduces both replication and production of infectious progeny

To test whether early expression of dksA<sub>Ct</sub> affects replication and/or production of infectious EBs, C. trachomatis was transformed with the inducible riboswitch construct pBOMB4-E-Riboswitch-dksA<sub>Ct</sub> (R-dksA<sub>Ct</sub>). Additionally, to ensure any phenotype observed upon induction of the R-dksA<sub>Ct</sub> construct is not a nonspecific effect of protein expression, HeLa cultures were infected with C. trachomatis transformed with p2TK2-SW2: E-Riboswitch-Clover (R-Clover) (Grieshaber et al. 2022) and expression of Clover, a variant for green fluorescent protein, used as a control. Replication was measured via analysis of GE in samples harvested at 30 hpi in the absence or presence of 0.5 mM theophylline, added at the time of infection. Induced expression of DksA<sub>Ct</sub> resulted in a 39.3% reduction in GE compared with non-induced control cultures (Fig. 2A). No significant effect on replication was observed with ectopic expression of Clover. These data suggest that the negative impact of early expression of DksA<sub>Ct</sub> on
replication is specific to DksA activity and is not due to nonspecific effects of induced protein expression.

To assess whether early expression of dksA results in a statistically significant reduction in C. trachomatis intracellular replication and recovered IFUs. The effect of theophylline was confirmed via western blot or fluorescence microscopy, respectively (Fig. S2, Supporting Information).

**Figure 2.** Ectopic expression of DksA results in reduction in C. trachomatis intracellular replication and recovered IFUs. The effect of DksA expression on C. trachomatis growth and EB generation was tested by theophylline-induced ectopic expression during infection of HeLa cells. (A) Infection of HeLa cells with R-dksA in the presence of theophylline resulted in a 39.3% (P = 0.0115, n = 3) reduction in GE when compared with the non-induced R-dksA control, and 48.2% (P = 0.0012, n = 3) and 44.1% (P = 0.0036, n = 3) reduction in GE when compared with non-induced or induced R-Clover, respectively. Induction of R-Clover expression did not significantly affect measured GE (P = 0.7931, n = 3) when compared with non-induced controls. (B) IFU assays conducted with samples normalized to GE showed a 49.6% (P < 0.0001, n = 3) and 13.6% (P = 0.0428, n = 3) reduction in recovered IFUs for induced R-dksA and R-Clover, respectively, when compared with non-induced control (set to 100%). The number of recovered IFUs following induction of R-dksA was 36% less than that observed with induced R-Clover (P = 0.004, n = 3). P-values were calculated using one-way analysis of variance (ANOVA) with Tukey's post hoc test.

**DksA retains major structural characteristics of DksA proteins**

Amino acid sequence identity is commonly used as an indicator for predicting conservation of protein function (Tian and Skolnick 2003, Capra and Singh 2007, Krissinel 2007). Amino acid sequence alignment of DksA from various chlamydial species and multiple orthologs for which the crystal structures have been determined (DksA, DksA(N) and DksA(L)) indicated high sequence conservation within the Chlamydia genus, and low sequence identity between the Chlamydia proteins and those encoded by genes in other bacteria (Fig. 3). Among human-adapted Chlamydia species, dksA showed the highest identity with C. pneumoniae (90% identity at the amino acid level). In addition to apparent high conservation across the Chlamydia genus, homologs of dksA can also be found in nonpathogenic Proteobacteria amoebophila UWE25 and Proteochlamydia naegleriophila KNiC (Clark et al. 2016). Despite low (~10%) primary sequence identity (Perederina et al. 2004, Rutherford et al. 2007), the transcription elongation factor GreA (GreA) has been shown to assume the physiological function of DksA in E. coli ΔdksA ΔppGpp mutants (Vinella et al. 2012), indicating that sequence identity does not consistently correlate with conserved function in this family of transcription factors.

Sequence analyses of DksA proteins from several bacteria has shown that while extensive diversity exists in the primary amino acid sequences, all proteins contain a conserved four-cysteine (C4) zinc-finger motif, CxxC-(x17)-CxxC, or a two-cysteine (C2) or one-cysteine (C1) zinc-independent motif, CxxS/T-(x17)-C/S/TxxA, with the first cysteine always conserved (Perederina et al. 2004, Henard et al. 2014). While the C4 proteins are sometimes referred to as DksA and the C2 and C1 proteins as DksA2 (Furman et al. 2013), here we will refer to all three groups as DksA proteins. The genomes of Chlamydia species all contain a single less frequently observed C1-DksA ortholog. Structures have been determined for two C4-DksA proteins, Escherichia coli (DksA) (Perederina et al. 2004) and Agrobacterium fabrum (DksA), and one C2-DksA protein, P. aeruginosa (DksA) (Furman et al. 2013). Prototypically, DksA proteins contain two domains that interact with RNAP: an N-terminal, two helix, coiled-coil domain that has been shown to interact with the α′ rim helices outside the RNA secondary channel (Perederina et al. 2004, Mandel et al. 2013).
Table 1

| Organism | Identity |
|----------|----------|
| Ct       | 100%     |
| Cp       | 90.3%    |
| Cs       | 96.0%    |
| Cm       | 91.0%    |
| Ec       | 16.2%    |
| Pa       | 25.7%    |
| Af       | 19.5%    |

Figure 3. Primary sequence alignment of DksA proteins. Amino acid sequences of DksA proteins were assessed by comparing proteins with solved crystal structures and those of pathogenic chlamydiae. High sequence identity was observed for DksA between chlamydial species. Ct = C. trachomatis; Cp = C. pneumoniae; Cs = C. suis; Cm = C. muridarum; Ec = E. coli; Pa = P. aeruginosa; and Af = A. fabrum. The four-cysteine residues of the zinc-finger domain in the E. coli and A. fabrum proteins are indicated by arrowheads.

Lennon et al. (2012, Parshin et al. (2015)). This interaction has high specificity, and it has been demonstrated in E. coli that a single point mutation in the C-terminal α-helix can result in loss of both binding and activity in vitro (Parshin et al. 2015). To facilitate a better general understanding of C1-DksA proteins and further elucidate the significance of widespread DksA conservation in Chlamydia (Domman and Horn 2015), we solved the 3D crystal structure of DksA_Ct. Despite the relatively low resolution (2.95 Å) of the X-ray diffraction data, the final model had excellent refinement statistics (Table 1). Two protein molecules are observed in the asymmetric unit with the structure of each protomer having a backbone atom (N-Cα-C = O) root-mean-standard deviation of 0.33 Å between both chains (Fig. 4A). Superposition of the protein structures show that the topology of all four DksA protein structures are similar with four α-helices organized into two domains. The relative orientation of the two domains overlap in all four structures despite DksA_Ct and DksA_Ec containing two or one cysteine, respectively, and no zinc ion, further establishing that zinc is not required for function of DksA proteins (Furman et al. 2013). Instead, the cysteine and zinc content of DksA may impact its capacity to respond to reactive oxygen and nitrogen species (Henard et al. 2014, Crawford et al. 2016). Because all pathogenic chlamydiae appear to express a highly homologous C1-form of DksA, the protein may have evolved in these pathogens to maintain a functional structure in (intracellular) environments where zinc is not abundant. What is different between the four structures is the region towards the turn between α1 and α2 in the coiled coil domain. In the crystal structure of DksA_Ct (C1-DksA), electron density is missing (E27-V62) and in the Nuclear Magnetic Resonance (NMR) structures for DksA_Af (C4-DksA) there is no convergence of the calculated structures (K29-L58). These observations in crystal (Oldfield et al. 2013) and NMR (Konrat 2014) structures may be correlated to intrinsically disordered regions. On the other hand, the loop between α1 and α2 is significantly smaller for DksA_Pa (chain C, R50-A58) and DksA_Ec (F69-P70), averaging eight and two residues, respectively. To explore the possibility that this difference could be attributed to varying degrees of intrinsic disorder we analyzed all four primary amino acid sequences of the proteins depicted in Fig. 4A using IUPred2A (Mészáros et al. 2018, Erdős and Dosztányi 2020). Alignment of these sequences starting with the first residue of the first N-terminal α-helix identifies a region of ~20 residues between the first two N-terminal α-helices with IUPred scores above 0.5, the threshold for disorder, for three of the four proteins (~10 residues for DksA_Pa showed scores below the cutoff) (Fig. S3, Supporting Information). We speculate that in monomeric DksA proteins the tip of the coiled coil domain is disordered with the helical regions at the C-terminal and N-terminal ends of α1 and α2, respectively, being transient in nature. This structural fluidity may be necessary for this domain to dock with the secondary channel of RNAP (Garner et al. 1999). These regions are more helical in the crystal structures of DksA_Ec and DksA_Pa, perhaps only due to crystal packing contacts that restrict motion and stabilize the helical state (Oldfield et al. 2013). Figure 4B is a cartoon representation of one of the two structures in the asymmetric unit showing that the protein contains...
Figure 4. Structure of DksA<sub>Ct</sub>. The 3D structure of DksA<sub>Ct</sub> was derived by X-ray diffraction. (A) Superposition of cartoon representations of both molecules in the asymmetric unit of DksA<sub>Ct</sub> (6PTG-B, marine and cyan, C1-DksA) with the crystal structure of DksA<sub>Ec</sub> (4IJJ-C, red, C2-DksA), DksA<sub>Ec</sub> (1TJL-F, orange, C4-DksA) and the NMR solution structure of DksA<sub>Af</sub> (2KQ9-Model_1, gray, C4-DksA) using the PDBeFold server (https://www.ebi.ac.uk/msd-srv/ssm/). The zinc ion in DksA<sub>Af</sub> is shown (magenta). (B) The overall domain structure of DksA<sub>Ct</sub> is consistent with that of other proteins known to bind the minor groove of RNA polymerase.

four α-helices with the N-terminal two helices forming a coiled coil domain and the two C-terminal helices part of a C-terminal globular domain containing a C1-motif. In both chains, electron density was insufficient to model the eight residues of the N-terminal tag, one or two residues at the C-terminus, and the region between L26(A)/E27(B) and V62, residues between the two helices of the coiled coil domain. No electron density corresponding to zinc or any other metal was observed.

**DksA<sub>Ct</sub> does not functionally complement DksA<sub>Ec</sub>**

Complementation of dksA-deficient E. coli has been conducted successfully with genes encoding C4-, C2- and C1-DksA proteins (Blaby-Haas et al. 2011, Pal et al. 2012, Lennon et al. 2014). For example, the C1-DksA ortholog of the α-proteobacterium R. sphaeroides shares 42% identity with DksA<sub>Ec</sub>, and has been shown to function alone and synergistically with ppGpp (Lennon et al. 2014). Divergent primary sequences, unresolved structural domains and the unique biology of C. trachomatis prompted analysis to determine whether dksA<sub>Ct</sub> can functionally complement the prototypical E. coli ortholog dksA<sub>Ec</sub>. DksA functionality can be tested based on the role of DksA in regulating amino acid synthesis in E. coli during growth on minimal M9 medium (Jude et al. 2003). dksA complementation in E. coli has shown that the dksA genes of Vibrio cholerae (Pal et al. 2012), R. sphaeroides (Lennon et al. 2014) and P. aeruginosa (Blaby-Haas et al. 2011) can functionally complement dksA<sub>Ec</sub>. In our hands, P. aeruginosa dksA (dksA<sub>1Pa</sub>) resulted in partial rescue of an E. coli mutant unable to express dksA<sub>Ec</sub>, while complementation with dksA<sub>Ct</sub> did not restore the growth defect (Fig. 5). Moreover, complementation with P. aeruginosa dksA2 (dksA<sub>2Pa</sub>) did not result in recovery of the growth defect exhibited by E. coli lacking
Figure 5. dksACt cannot functionally complement dksAEc. To test whether dksACt could functionally complement the dksA gene of the model organism E. coli, dksACt was used to complement a dksAEc deletion mutant. (A) Complementation of E. coli ΔdksA with dksACt, dksA1Pa or dksA2Pa plated on minimal medium. Complementation with dksAEc or an empty vector were used as controls. (B) Arabinose induction of pBAD18::dksACt was confirmed via western blotting of strains following growth in Luria-Bertani (LB) broth.

Table 1. X-ray diffraction data for DksA.Ct.

| Parameter                  | Value |
|----------------------------|-------|
| PDB ID                    | 6PTG  |
| Space group               | P3_21 |
| **Diffraction data**      |       |
| a (Å)                     | 83.42 |
| b (Å)                     | 83.42 |
| c (Å)                     | 99.03 |
| α, β, γ (°)               | 90, 120 |
| Matthews coefficient (Å^3Da⁻¹) | 3.32 |
| Solvent content (%)       | 63%   |
| Resolution range (Å)      | 50–2.95 (3.03–2.95) |
| Mean I/σ (I)              | 20.54 (3.12) |
| No. of observed unique reflections | 8668 (628) |
| Completeness (%)          | 98.8 (98.9) |
| Multiplicity              | 6.1 (6.2) |
| Rmerge^2                  | 0.068 (0.625) |
| CC 1/2                    | 0.999 (0.890) |
| **Refinement**            |       |
| Resolution                | 50–2.95 (3.13–2.95) |
| No. of used reflections   | 8651 (1271) |
| Rwork                     | 0.1823 (0.2636) |
| Rfree                     | 0.2264 (0.3279) |
| RMSD bonds (Å)            | 0.004 |
| RMSD angles (°)           | 0.54 |
| Mean β factor overall (Å²) | 73.8 |
| Mean β factor protein (Å²) | 74.1 |

**MolProbity model analysis**

| Clash score, all atoms  | 2.77  |
| Ramachandran favored (%) | 100   |
| Ramachandran outliers (%) | 0     |
| Rotamer favored (%)     | 100   |
| Rotamer outliers (%)    | 0     |
| MolProbity score        | 1.07  |

The dksACt expression is regulated from promoter sites both upstream of and within ndrR

Gene linkage and co-expression can be related to the function of two or more genes in a coordinated biological process. In all chlamydial genomes analyzed, dksA is located immediately downstream of ndrR, encoding the NrdA/B repressor NrdR, a negative regulator of deoxyribonucleotide synthesis. lspA, encoding a lipoprotein signal peptidase, is located immediately downstream of dksACt. The genomic localization and organization of dksA with ndrR and lspA in chlamydial genomes, and intergenic regions of only 5–24 bp, are consistent with their control by a single promoter upstream of ndrR. This conserved orientation provides a potential link between the role of dksA and the metabolic function of ndrR in nucleic acid metabolism. Previously published deep sequencing data indicate that ndrR, dksA and lspA in C. pneumoniae are transcribed as a polycistronic mRNA (Albrecht et al. 2011) and suggest that the same may be true in C. trachomatis (Albrecht et al. 2010).

To determine whether dksACt could be regulated from an independent promoter downstream of the promoter controlling expression of the ndrR-dksACt-lspA operon, C. trachomatis was transformed with transcriptional reporter constructs based on the pBOMB vector (Bauler and Hackstadt 2014) containing sequences 830 or 469 bp upstream of dksACt linked to GFP-LVA (Fig. 6A). Constitutive expression of mCherry was used as a positive control for transformation. GFP was detected regardless of construct configuration suggesting that dksACt can be transcribed from promoters both upstream of and within ndrR (Fig. 6B). No signal was detected from the empty vector control.
Expression of \textit{dksACt} is controlled by two promoters. The possibility that \textit{dksACt} expression can be controlled independently of \textit{nrdR} was tested using a transcriptional reporter construct. (A) Schematic representation of the \textit{C. trachomatis} genome with sequences upstream of \textit{dksACt} containing potential promoters, and the reporter construct used to test expression. (B) Fluorescence micrographs (×200 magnification, 45 hpi) reveal expression of GFP-LVA regardless of promoter region used. Constitutive mCherry expression was used to confirm transformation. No signal from GFP-LVA was observed in the empty vector control. Equivalent reporter expression was observed in both mixed populations of transformants under antibiotic selection (\(n = 3\)) and in clonally isolated transformants (\(n = 1\)).

To independently verify the existence of two unique transcripts containing \textit{dksACt}, total RNA was extracted from \textit{C. trachomatis}-infected HeLa cells at 20 and 30 hpi and northern blotting performed with oligonucleotide probes complementary to \textit{dksACt}. Blots indicate that \textit{dksACt} is expressed as two transcripts with a prominent band at \(\sim 1.0\) kb and a less prominent band at 1.5 kb (Fig. 7A). This pattern is consistent with polycistronic regulation via the 1.5 kb fragment, as the distance from the published TSS of \textit{nrdR} to the \textit{lspA} stop codon is 1477 bp (Albrecht et al. 2010).

Characterization of the 1.0 kb fragment required additional verification due to the similar sizes of \textit{nrdR} (465 bp) and \textit{lspA} (504 bp). Therefore, we utilized neural network-based prediction software (Reese 2001) to search for sites within the identified operon region that may constitute an alternative promoter. Four sites were identified, one of which correlated directly with the TSS previously determined for \textit{nrdR}. Of the other three predicted TSSs, two were located within \textit{nrdR} and one within \textit{dksACt}. Using the predictions as a guide, 5’RACE was conducted using gene specific primers (GSPs) designed to bind within \textit{dksACt} (Fig. 7B; Fig. S4, Supporting Information). Sequencing of 5’RACE fragments indicate a promoter site \(\sim 80–150\) bp upstream of the \textit{dksACt} ORF, which directly correlates with the 783880–783925 region predicted to contain the alternative promoter.

Transcript levels for \textit{dksACt} and the adjacent genes \textit{nrdR} and \textit{lspA} were determined by RNA-Seq at time points correlating with early, mid and late stages of the chlamydial developmental cycle (Fig. 7C). Data obtained by RNA-Seq support independent regulation of \textit{dksACt} through a 5–6-fold increase in \textit{dksACt} reads when compared with \textit{nrdR}. Additionally, the expression pattern of \textit{DksA_Ct} (Fig. 1) correlates with the temporal pattern of transcript abundance observed in the RNA-Seq data set. As a final confirmation, quantitative reverse transcription polymerase chain reaction (RT-qPCR)-based transcriptional analysis conducted using cDNA extracted at 30 hpi indicated a 3–4-fold increase in \textit{dksACt} transcript when compared with a region of \textit{nrdR} upstream from the 5’RACE site (Fig. S5, Supporting Information). This increase
in transcript abundance corroborates data obtained by RNA-Seq, further supporting transcriptional regulation for dksA_C from two independent promoters.

**Conclusions**

DksA has been described as one of eight transcription factors conserved across the phylum Chlamydiaceae (Domman and Horn 2015). Based upon data from previously studied C4-, C2- and C1-DksA orthologs, we predicted that DksA_C is maintained as a protein associated with RB–EB transitions in the chlamydial developmental cycle. Herein, we showed that DksA_C is expressed maximally at the RB–EB transition point (i.e. 15–20 hpi), a unique expression pattern compared with the stable expression observed for DksA_L (Paul et al. 2004, Brown et al. 2002). Ectopic overexpression of DksA_C prior to RB–EB transition produced a defect in both replication and EB generation. We found that dksA_C does not functionally complement dksA_L with respect to recovery of replication during growth on minimal medium. DksA_C has an overall domain structure similar to other C2- and C4-type DksA proteins despite containing only one cysteine residue. Comparing our C1-type DksA structure to the structure of other C2- and C4-type DksA proteins we speculate that the ‘tip’ of the coiled coil domain may be transiently structured/intrinsically disordered to assist binding to the secondary channel of RNAP. Expression of DksA_C is likely regulated by tandem promoter regions, a regulatory scheme also described for other genes in this pathogen (Lambden et al. 1990, Rosario and Tan 2012). The phenotype resulting from induced DksA_C expression prior to RB–EB transition and the correlation between dksA_C and nrdR expression is consistent with a role for DksA_C in the C. trachomatis developmental cycle. It should be mentioned that DksA_C can function in ways that could be beneficial to C. trachomatis even in the absence of a stringent response, including resolution of transcription conflicts (Tehranchi et al. 2010) and protection of phleomycin-induced double-stranded breaks in DNA (Sivaramakrishnan et al. 2017). Therefore, selective pressure to maintain dksA in chlamydiae could be related to such alternative functions.

It will be valuable to pursue loss of function phenotypes in future studies via deletion of dksA_C (Mueller et al. 2017, Keb et al. 2018, Keb and Fields 2020) or CRISPR interference (Ouellette et al. 2021) in case dksA_C is essential to C. trachomatis intracellular replication and/or developmental transitions. Mechanistic understanding of how DksA_C functions in C. trachomatis biology will require analysis of whether the protein directly influences pathogen gene expression by binding to RNAP akin to prototypical DksA proteins, or whether DksA_C is associated with a different mechanism of action, such as interfering with the binding of another protein to RNAP, thus indirectly affecting transcriptional regulation.

**Experimental procedures**

**Organisms and cell culture**

*Chlamydia trachomatis* serovar L2 (LGV 434/Bu) was grown in the *Homo sapiens* cervix adenocarcinoma cell line HeLa or the murine fibroblast cell line McCoy (ATCC; Manassas, VA) cultured at 37°C with 5% CO₂ in RPMI 1640 (HyClone, Logan, UT) supplemented with 10% FetalPlex serum complex (Gemini; Sacramento, CA) and 10 μg/mL Gentamicin (MilliporeSigma; St Louis, MI).

**Maintenance, purification and quantification of *Chlamydia***

*Chlamydia trachomatis* RB and EB cell forms were purified by density gradient centrifugation as previously described (Griehaber et al. 2018). Chlamydial EBs were stored at −80°C in sucrose-phosphate-glutamate (Bovarnick et al. 1950) or K36 (Weiss 1965) buffer until use. Crude preparations of EBs for use in transformations were generated as previously described (Mueller et al. 2017). *Chlamydia trachomatis* was quantified by measurement of GE by SYBR green-based qPCR using a CFX96 or CFX384 real time PCR Detection System (BioRad, Hercules, CA) and primers targeting the hctA gene of *C. trachomatis*.

**Plasmid construction and transformation of *Chlamydia***

PCR primers used in this study are listed in Table S1 (Supporting Information). The vector pBOMB3cdMCI::incD-GFPLVA (Bauler and Hackstadt 2014) was linearized by PCR to remove the incD cassette using Q5 High-Fidelity DNA polymerase (New England Biolabs; Ipswich, MA). For dksA transcriptional fusion constructs, two inserts were generated spanning regions of 469 and 830 base pairs upstream from the annotated start site of dksA. For the E-Riboswitch dksA construct, dksA was inserted into pBOMB4 with an E-Riboswitch (Topp et al. 2010) driven by a T5 promoter to generate pBOMB4-E-Riboswitch-dksA_C. All DNA inserts were PCR-amplified using Phusion DNA polymerase (New England Biolabs; Ipswich, MA) and primer sets were generated according to the In-Fusion HD Cloning (Takara Bio, USA) guidelines. Linearized pBOMB4 and pBOMB4::GFPLVA were DpnI treated, then ligated to each insert and transformed into Stellar competent cells using the In-Fusion HD Cloning Kit (Takara Bio, USA). After incubation overnight at 37°C on LB agar plates containing 100 μg/mL carbenicillin (LBCarb) (Chem-Impex International; Wood Dale, IL), colonies were selected for replating and screening via PCR. Plasmids were isolated using the Purelink Quick Plasmid Miniprep Kit (Invitrogen; Carlsbad, CA) and sequenced for insert confirmation. The confirmed plasmids were transformed into methyltransferase-deficient *E. coli* K-12 ER2925 competent cells and incubated overnight. Positive colonies were subcultured in 50 mL LBCarb broth at 37°C, and unmethylated plasmids were isolated using the GenElute HP Endotoxin Free Plasmid Maxiprep Kit (MilliporeSigma; St Louis, MO). Eluted plasmids were either concentrated to ~500 ng/μL using the DNA Clean and Concentrator-100 kit (Zymo Research; Irvine, CA) or via sodium acetate and isopropanol-mediated DNA precipitation. Unmethylated plasmids purified from *E. coli* were transformed into *C. trachomatis* essentially as described (Mueller et al. 2017). For some transformations of *C. trachomatis*, density gradient purified EBs (1 × 10⁸ GE/well) were mixed with 2 μg of plasmid DNA. Cultures identified to have positive transformants as determined by fluorescence microscopy, were passaged two additional times to remove residual penicillin G susceptible bacteria and increase the titer of transformed cells. Clonal populations were generated via extraction of *C. trachomatis* from inclusions by micromanipulation repeated four times under limiting dilution. Plasmid DNA was obtained from isolated clones and transformed into *E. coli* where five colonies expressing mCherry were sequenced for confirmation of clonality.

**Sequence analysis**

Amino acid sequence alignments were done using the ClustalW (Thompson et al. 1994) alignment tool using MEGA software
**Escherichia coli complementation assay**

Solid M9 medium was generated using Difco 5× M9 Minimal Salts, (Becton, Dickinson and Company, Franklin Lakes, NJ) supplemented with glycerol (0.4% v/v), MgSO₄ (2 mM), CaCl₂ (0.1 mM) and agar (1% w/v). Sterile filtered l-arabinose inducer was added at a final concentration of 0.1% (w/v) as indicated. *Escherichia coli* ΔdksA (strain: JW0141-1) was obtained from the Keio Knockout Collection, Yale University [Baba et al. 2006]. dksAEc, dksACt, dksAIth or dksA2Pa was ligated into the pBAD18 vector and transformed into E. coli ΔdksA. *Escherichia coli* K-12 was used as a positive control. Bacteria were plated and incubated for 24 h at 37°C, then imaged using a ChemiDoc Imaging System (Bio-Rad, Hercules, CA).

**Analysis of DksAΔ expression**

Protein lysates for analysis of DksAΔ expression by western blotting were obtained from 10–15 HeLa cell cultures (T-175) infected with *C. trachomatis* serovar L2 genome. 

RNA extraction and northern blotting

HeLa cells were established in T-175 cell culture flasks and infected at an MOI of 300; bacteria were enriched by density gradient centrifugation using a 30% diatrizoate meglumine (MD-76R) pad essentially as described [Howard et al. 1974, Grieshaber et al. 2018]. Bacteria were enumerated by area of *C. trachomatis* heat denatured at 100°C for 10 min. Twelve % sodium dodecyl-sulfate polyacrylamide gel electrophoresis (SDS–PAGE) gels were loaded with the equivalent of 2 × 10⁶ GE of each sample per well and electrophoresed at 25 mA for 40–60 min, then transferred to a 0.45 um PVDF membrane using a Trans-Blot Turbo transfer apparatus (Bio-Rad, Hercules, CA). The membrane was blocked in PBS + 0.1% Tween-20 (PBS-T) and 3% nonfat dry milk overnight at 4°C. The membrane was probed with a custom protein-A purified anti-DksAΔ antibody raised in rabbits against a truncated recombinantly expressed DksAΔ antigen in combination with a goat anti-rabbit poly-HRP secondary antibody (Invitrogen; Carlsbad, CA). The membrane was treated with the SuperSignal West Femto luminol and peroxide solution (ThermoFisher; Waltham, MA) and imaged using a ChemiDoc Imaging System (Bio-Rad; Hercules, CA). Animal use was approved by the Institutional Animal Care and Use Committee at Washington State University.

**5′ RACE**

DNA was isolated at 24 hpi and converted into cDNA tagged with the SMARTer II A Oligonucleotide using random hexamers as described in the SMARTer RACE 5′/3′ Kit (Takara Bio USA, Mountain View, CA). cDNA libraries were diluted as needed, and 5′RACE PCR reactions were run using three separate GSPs designed to target the intragenic dksAΔ sequence (Table S1, Supporting Information). PCR products were electrophoresed on a 1% TAE gel stained with SYBR Safe DNA Gel Stain (Invitrogen; Carlsbad, CA), and bands were excised for DNA isolation using the Nucleospin Gel and PCR Cleanup Kit (Macherey-Nagel; Düren, Germany). Fragments were then cloned using the CloneJET PCR cloning kit (ThermoFisher; Waltham, MA) and plasmids were isolated from positive transformants using the Nucleospin Plasmid Kit (Macherey-Nagel; Düren, Germany). Plasmids were sequencing using pJet1.2 vector specific primers (Eurolabs Genomics). Clones were analyzed using NCBI BLAST against the *C. trachomatis* serovar L2 genome. Sequences not containing both the GSP and Universal Primer Mix regions, indicative of incomplete sequences, were excluded from the analysis.

**PCR and RNA-Seq**

cDNA for both conventional PCR analysis of polycistronic RNA and RT-qPCR applications was generated using the SuperScript III First-Strand Synthesis kit (Invitrogen; Carlsbad, CA) with RNA isolated at 20 and 30 hpi. Negative controls not containing reverse transcriptase were generated for each time point. Primers specific to mdr and IspA (Table S1, Supporting Information) were utilized in conventional PCR to amplify cDNA for operon determination. For transcriptional analysis, *C. trachomatis* serovar L2 genomic DNA was extracted from density gradient purified EBs using phenol/chloroform precipitation. A dilution series of genomic DNA was generated and used for calculation of primer efficiencies and for calculation of relative copy number. Primer efficiencies between 85% and 110% were deemed acceptable for direct comparison of transcript levels. RNA-Seq data was recovered from a previously published data set [Grieshaber et al. 2018].

**Ectopic expression of DksA in *C. trachomatis***

HeLa cells were grown to confluency in T-25 cell culture flasks. The culture medium was then replaced with RPMI-1640 supplemented with 10% (v/v) FetalPlex and 10 µg/mL Gentamicin. Each condition was tested in duplicate with two cultures supplemented with tagged ssDNA oligonucleotide probe (Table S1, Supporting Information) was designed to bind dksA mRNA according to the ULTRAhyb Oligo Buffer (Invitrogen; Carlsbad, CA) recommendations. The RiboRuler high range RNA ladder and 2x loading dye (ThermoFisher; Waltham, MA) were used to assess mRNA size. The membrane was cut to match the exact size of the denaturing gel, and RNA transferred via capillary action for 2 h. The membrane was incubated in a UV crosslinker ( Hoefer; Holliston, MA) for 1 min, repeated 3 times. The membrane was pre-hybridized with rotation in a 50 mL conical tube for 1 h in ULTRAhyb Oligo Buffer at 40°C. The probe was added to 3.5 mL buffer at a concentration of ~600 pM and hybridized at 40°C for 17–20 h. The membrane was washed in the NorthernMax low stringency solution twice for 5 min at RT, and once at hybridization temperature for 2 min. The washed membrane was treated with the Chemiluminescence Nucleic Acid Detection Module Kit (ThermoFisher; Waltham, MA) according to the manufacturer’s instructions, and imaged using a ChemiDoc Imaging System (Bio-Rad; Hercules, CA).
0.5 mM theophylline and the other two serving as controls. Theophylline was added to the culture medium prior to inoculation. To facilitate comparative analysis, each flask was inoculated to reach equivalent levels (50–75%) of infection using bacteria carrying the pBOMB4::E-Riboswitch-dksA<sub>Ct</sub> or pZTK2-SW2::E-Riboswitch-Clover (Grieshaber et al., 2022) vectors and incubated at 37°C in a 5% CO<sub>2</sub> incubator for 30 h. Induction of pBOMB4::E-Riboswitch-dksA<sub>Ct</sub> results in expression of an untagged DksA<sub>Ct</sub> protein. The medium was then removed, and HeLa cell monolayers were washed with K-36 buffer. Infected host cells were disrupted by vigorous scraping in K-36 buffer and then host cell debris was removed from the mixture via centrifugation at 100 × g for 3 min at 4°C. The supernatant was removed, and Chlamydia was pelleted at 20000 × g for 15 min at 4°C. Pelleted bacteria were resuspended in 30 μL SPG and frozen at −80°C. GEs were enumerated using qPCR and IFU assays were conducted using HeLa cells grown to confluency in 12-well plates with inocula normalized to GE.

**Microscopy**

Micrographs were acquired using a Leica DMi8 inverted microscope (Leica Microsystems; Buffalo Grove, IL) equipped with an X-Cite LED light source and DMC2900 camera. *Chlamydia trachomatis* IFU assays were done using methanol fixed samples stained with a fluorescein isothiocyanate-tagged anti-MOMP antibody (PA1-73073 Thermo Fisher; Waltham, MA) at 1/250 dilution in PBS buffer. Images were analyzed and processed using ImageJ (The National Institute of Health; Bethesda, MD). Adjustments of contrast or signal intensity were applied to the entire image.

**Cloning of dksA<sub>Ct</sub>**

The dksA<sub>Ct</sub> gene (Gene ID: 5858340) was amplified from the genomic DNA of *C. trachomatis* serovar L2 (LGV 434/ Bu) (NCBI: 471472) and inserted into the expression vector BG1861 (a derivative of pET14b) (Myler et al. 2009) at a site containing an uncleavable, 8-residue N-terminal tag (MAHHHHHHH). Using a heat shock method, the recombinant plasmid was used to transform *E. coli* BL21(DE3)-R3-pRARE2 cells (gift from SGC Toronto). From these transformed cells, stocks were prepared (~1 mL LB media, OD<sub>600</sub> ~0.8) from a single colony and stored at −80°C in glycerol (~15%) solution until preparation of samples for structural analysis.

**Crystallization, X-ray data collection and structure refinement of DksA<sub>Ct</sub>**

For X-ray crystallographic analysis, DksA<sub>Ct</sub> was expressed and purified following standard SSGCID protocols (Bryan et al. 2011). The protein was concentrated to 24.6 mg/mL in 25 mM HEPES, 500 mM NaCl, 5% (v/v) glycerol, 2 mM DTT, and 0.025% sodium azide (v/v), pH 7.5. Initial crystallization conditions were searched using 0.4 μL drops in XJR crystallization trays (Rigaku Reagents; Bainbridge Island, WA) and several commercial screens: JCSG+, JCSG-Top96, Wizard 1/2, Wizard 3/4 (Rigaku Reagents; Bainbridge Island, WA), Crystal Screen, IndexHR (Hampton Research; Aliso Viego, CA), MCSG-1, MCSG-2 (Microlytic/Anatrace; Maumee, OH), Morpheus 1 and 2, PACT (Molecular Dimensions; Holland, OH). Initial crystal hits were optimized starting from CrystalScreen, condition B5: 200 mM lithium sulfate, 25% (v/v) PEG 4000, 100 mM Tris, pH 8.5. The crystals were cryo-protected with 15% ethylene glycol and vitrified in nylon loops by plunging into liquid nitrogen. Diffraction data up to 2.95 Å resolution were collected at the Advanced Photon Source (APS), with a Rigaku R-AXIS RAPID detector. Diffraction data statistics are summarized in Table 1. The structure could not be solved using the standard SSGCID molecular replacement strategies. However, with MR-Rosetta (Terwilliger et al. 2012) as implemented in Phenix (Adams et al. 2011), a dimer solution could be found that was based on PDB entries 4UIJ_A, 1TJL_H, 2KQ9_A, 2KGO_A and 5W1S_N. The structure was then completed with iterative refinement cycles in phenix refine (Zwart et al. 2008), and real space modeling in Coot (Emsley et al. 2010). The quality of the model was assessed with tools built into Coot and phenix refine, such as MolProbity (Williams et al. 2018). The coordinates and structure factors were deposited in the PDB with code 6PTG. Diffraction images are available at proteindiffraction.org.

**Statistical analysis**

Statistical analyses were conducted using Prism (GraphPad Inc.; San Diego, CA) software. Statistical analysis and normalization of read counts for RNA-Seq data (Fig. 7) was done using DESeq2 in R (Love et al. 2014). Normalization from Dseq2 was used for transcript visualization in Integrated Genomics Viewer (Robinson et al. 2011).

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**Supplementary data**

Supplementary data are available at FEMSPD online.

**Contributions**

CM, HY, GWB, JA, TC, NG, SG and AO designed and performed experiments and analyzed data. CM and AO wrote the manuscript.
CM, HY, GWB, JA, TC, NG, SG and AO edited the manuscript. AO designed the study.

**Data availability**

The data underlying this article are available via Protein Data Bank and proteinidiffract.org and can be accessed with PDB identifier 6PTG.

**Conflict of interest statement.** The authors have no conflicts of interest to declare.

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