Massively parallel transposon mutagenesis identifies temporally essential genes for biofilm formation in *Escherichia coli*

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Abstract

Biofilms complete a life cycle where cells aggregate, grow and produce a structured community before dispersing to seed biofilms in new environments. Progression through this life cycle requires controlled temporal gene expression to maximise fitness at each stage. Previous studies have focused on the essential genome for the formation of a mature biofilm, but here we present an insight into the genes involved at different stages of biofilm formation. We used TraDIS-Xpress; a massively parallel transposon mutagenesis approach using transposon-located promoters to assay expression of all genes in the genome. We determined how gene essentiality and expression affects the fitness of *E. coli* growing as a biofilm on glass beads after 12, 24 and 48 hours. A selection of genes identified as important were then validated independently by assaying biofilm biomass, aggregation, curli biosynthesis and adhesion ability of defined mutants. We identified 48 genes that affected biofilm fitness including genes with known roles and those not previously implicated in biofilm formation. Regulation of type 1 fimbriae and motility were important at all time points. Adhesion and motility were important for the early biofilm, whereas matrix production and purine biosynthesis were only important as the biofilm matured. We found strong temporal contributions to biofilm fitness for some genes including some which were both beneficial and detrimental depending on the stage at which they are expressed, including *dksA* and *dsbA*. Novel genes implicated in biofilm formation included *ychM* and *truA* involved in cell division, *crfC* and *maoP* in DNA housekeeping and *yigZ* and *ykgJ* of unknown function. This work provides new insights into the requirements for successful biofilm formation through the biofilm life cycle and demonstrates the importance of understanding expression and fitness through time.
Introduction

Bacteria rarely exist planktonically outside of the laboratory and are usually found as part of structured, aggregated communities called biofilms. Clinically, approximately 80% of infections have been suggested to have a biofilm component. Biofilm related infections are complicated by their intrinsic tolerance to antimicrobials, making infections difficult to treat and, often persistent.

Cells within a biofilm grow more slowly than those in planktonic culture and this reduced level of metabolic activity has been associated with tolerance to antimicrobials, allowing biofilms to be typically 10-1000-fold less sensitive to antibiotics than corresponding strains in planktonic conditions. Aside from the clinical setting, there are many useful applications of biofilms, including wastewater treatment and bioprocessing. Biofilms undergo a life cycle that commonly consists of initial attachment to a surface, growth and maturation of the biofilm over time with characteristic production of extracellular matrix components, followed by dispersal of planktonic cells to facilitate colonisation of new surfaces. The switch between planktonic and biofilm lifestyles is driven by environmental stimuli promoting large scale changes in gene expression and regulation that are necessary to support the bacterial community through the life cycle, which is distinct to planktonic growth conditions. Expressing the right genes at the right time and place is critical for efficient production of a biofilm.

The main components of the biofilm extracellular matrix in *E. coli* are the amyloid protein curli, the polysaccharide cellulose and extracellular DNA. Curli is transcribed by the divergent operons csgBAC and csgDEFG, with curli biosynthesis regulated by CsgD. Cellulose biosynthetic machinery is encoded by bcsRQABZC and bcsEFG, and its production is regulated by c-di-GMP. Several genes are known to be involved in the regulation of matrix production, including *ompR*, *cpxR*, *rpoS*, amongst others. Extracellular DNA is also an important component of the biofilm matrix, and the addition of DNase has negatively affected the biomass of biofilms formed by *Pseudomonas aeruginosa*, *Bacillus cereus* and a range of Gram-negative pathogens, including *E. coli*.
Previous studies focusing on identifying the genes and pathways required for biofilm formation in *E. coli* have concentrated on the mature biofilm rather than dissecting events across the life cycle. Various studies have taken a genome wide approach to identifying genes that impact biofilm formation. One assessed biofilm formation of all the mutants in the Keio collection, although analysis is limited to the effect of inactivation. DNA microarrays have also been used to link the presence of different genes with biofilm capacity in panels of isolates. Large scale transposon mutagenesis are high-throughput, sensitive whole genome approaches to link phenotype to genotype methodologies. These methods are limited by an inability to assay essential genes where transposon insertions are not viable. We sought to investigate biofilm formation using TraDIS-Xpress, a method developed recently which allows essential genes to be assayed for their role in survival in test conditions. In this method, gene expression changes caused by an outward-facing inducible promoter incorporated into the transposon are captured as well as traditional essentiality measurements.

This study identified 48 genes that were found to be important at different stages of biofilm formation by *E. coli*. By investigating the genes important across the biofilm life cycle, we were able to get a dynamic view of the main pathways important at different stages of biofilm development. Genes identified by TraDIS-Xpress as important at each stage were then phenotypically validated using defined mutants. Our findings reinforced the importance of adhesion, motility and matrix production in the biofilm, and revealed the importance of genes not previously implicated in biofilm formation. This included genes with roles in cell division (*ychM* and *truA*), DNA housekeeping (*crfC* and *maoP*) and *yigZ* and *ykgJ*, the functions of which have not been elucidated. We identified clear requirements for some pathways at specific points of the biofilm life cycle, furthering our understanding of how biofilms maintain fitness over time.
Results

Analysis of the TraDIS-Xpress data found 48 genes that considerably affected biofilm formation over time in *E. coli*: 42 were identified as being needed to maintain the fitness of a biofilm and expression of 6 genes was predicted to result in reduced biofilm fitness (Figure 1 and Supplementary table 1). The main pathways that were consistently important in the biofilm through all the time points included type 1 fimbriae, curli biosynthesis and regulation of flagella (Figure 1). All other loci identified affected biofilm formation at specific points in the life cycle.

Fimbriae expression and motility are important at all stages of biofilm formation

Only 4 genes were found to be important throughout 12, 24 and 48 hours (figure 2a). These included *fimB* and *fimE*. The recombinase gene *fimB* which helps mediate both ‘ON-to-OFF’ and ‘OFF-to-ON’ switching of fimbriae expression was beneficial for biofilm formation at all time points. Fewer *fimB* mutants were observed in biofilm conditions compared to planktonic, and this number decreased over time. In contrast, inactivation of *fimE*, responsible for only ‘ON-to-OFF’ fimbrial regulation, increased biofilm fitness at all time points. Initially, there were only slightly more *fimE* mutants in biofilm conditions compared to planktonic at 12 hours, but this increased over time with a stark contrast seen between biofilm and planktonic conditions at the 24 and 48 hour time points. The predicted impacts on biofilm formation were phenotypically validated by testing both gene knockout mutants from the Keio collection (which contains two, independent deletion mutants for most genes in *E. coli* BW25113). Analysis of these mutants showed for both that there was no deficit in biofilm biomass (Figure 3a), but both were deficient in cell aggregation (Figure 3b). Together, the TraDIS-Xpress and phenotypic data suggest that the ability to regulate fimbriae expression in a phase-dependent manner is important for fitness of a biofilm, rather than being constrained in a fimbriae ‘ON’ or ‘OFF’ state.

Disruption of *lrhA*, a regulator of motility and chemotaxis, was beneficial for biofilm formation at all time points (Figure 2b). LrhA also has a role in type 1 fimbriae expression through activating expression of *fimE*, but in addition represses flagella-mediated motility. Analysis of the Δ*lrhA*
biofilm showed initial formation of microcolonies occurred faster than the wild-type (Figure 4) but at later time points the biofilms formed by this mutant were less mature than seen with the wild-type. There was no significant change in biomass formed by this mutant (Figure 3a) and mutants appeared less aggregative than the wild-type (Figure 3b). These data suggest that inactivation of \textit{lrhA} impacts both adhesion and aggregation differently at distinct stages of the biofilm life cycle and may result in a benefit to early surface colonisation but with a cost to later maturation.

Expression of the Hha toxin attenuator \textit{tomB} was also found to be consistently important for biofilm formation at 12, 24 and 48 hours (Figure 2b). Consistent with this prediction, the \textit{ΔtomB} mutant biofilm had reduced cell aggregation and curli biosynthesis, and reduced biofilm biomass (Figure 3a,b,c).

\textbf{Regulatory genes are important in the early biofilm}

In the early biofilm, at 12 hours growth, only 13 genes were found to distinguish the planktonic and biofilm conditions. TraDIS-Xpress data indicated that inactivation of transcriptional factor \textit{dksA} promoted biofilm formation at the 12- and 24-hour time points but not in the mature biofilm (Figure 2c). Supporting this, analysis of \textit{ΔdksA} mutant biofilms under flow conditions showed an initial benefit with increased adhesion at both the 12 and 24 hour time points, but reduced microcolony formation at the 48 hour time point, suggesting \textit{dksA} affects biofilm initiation (Figure 4). Inactivation of \textit{ΔdksA} was also seen to reduce cell aggregation, curli biosynthesis and biofilm biomass (Figure 3a,b,c). Expression of \textit{hdfR}, a negative regulator of motility \textsuperscript{37}, was found to be detrimental to biofilm fitness in the early biofilm after 12 and 24 hours growth (Figure 2b), and \textit{ΔhdfR} mutant biofilms had significantly reduced biomass (Figure 3a). In addition, the stress response regulator \textit{marR} \textsuperscript{38} and the 23S rRNA methyltransferase \textit{rlmI} \textsuperscript{39} were both found to be beneficial for biofilm fitness at the 12 hour time point only, and reduced biofilm biomass was found in deletion mutants (figure 3a). These genes have both previously been implicated in biofilm formation \textsuperscript{39-41}, but the effect on early biofilm formation has not been described previously.
Biofilms sampled after 24 hours demonstrate both adhesion and matrix production are important.

More pathways were identified as being important to biofilm formation at 24 hours that at 12 hours. Two genes involved in DNA housekeeping were found to be involved in biofilm formation at this time point. These included *dam*, encoding DNA methyltransferase, insertional activation of which was not tolerated in the 24 hour biofilm, and Δ*dam* mutants were defective in aggregation compared to the wild type (figure 3b). Also, inactivation of *maoP*, involved in Ori macrodomain organisation, was predicted to confer a fitness advantage in the 24 hour biofilm compared to the planktonic condition. TraDIS-Xpress data showed more reads mapped to *maoP* in the biofilm conditions compared to the planktonic at 24 hours suggesting loss of this gene was beneficial. Phenotypic analysis of the Δ*maoP* mutant biofilm did demonstrate a phenotype although in opposition to the prediction, *maoP* mutants were significantly deficient in biofilm biomass production, curli biosynthesis and one mutant displayed reduced aggregation (Figure 3 a,c).

In the 12 and 24 hour biofilms, *dsbA* (encoding disulphide oxidoreductase) was essential with no insertions detected within this gene (Figure 2c). The role of *dsbA* in adhesion to abiotic surfaces and epithelial cells has previously been suggested (Lee et al., 2008, Bringer et al., 2007). Phenotypic validation of the Δ*dsbA* mutant showed a red, dry and rough (*rdar*) phenotype on Congo red plates (Figure 3c), indicative of increased curli biosynthesis. Cell aggregation in the Δ*dsbA* mutant was significantly improved compared to the wild type, implying a role of *dsbA* in inhibiting cell-cell aggregation. Our data showed that *dsbA* is important in the early biofilm, but its deletion appears to be beneficial to the formation of a mature biofilm, according to the Congo red and aggregation data. Supporting this hypothesis, *dsbA* was not essential at the 48 hour time point.

Inactivation of the RNase III modulator *ymdB* was found to reduce fitness in the biofilm, with fewer reads mapping here in biofilm conditions compared to planktonic at both the 24 and 48 hour time points. This follows previous findings that both inactivation and overexpression of *ymdB*
negatively affects biofilm biomass. In concordance with these findings, a ΔymdB mutant had significantly improved cell aggregation and reduced biofilm biomass (figure 3 a,b).

Curli biosynthesis became important by the 24 hour time point as no insertions mapped to csgC, encoding a curli subunit chaperone and more transposon insertions mapped upstream of the curli biosynthesis regulator csgD, likely indicating its increased expression benefitted biofilm formation. At the 48 hour time point, both genes were essential for biofilm formation, which was also the case for the known csgD regulator, ompR, supported by significantly reduced biofilm biomass and reduced aggregation in knockout mutants (figure 3a,b).

The mature biofilm grown for 48 hours requires purine biosynthesis, matrix production, motility and solute transport. There were 38 genes found to be important for fitness of the mature biofilm after 48 hours growth, and 25 of these genes were identified as essential at this time point only. The major pathway implicated in biofilm formation at 48 hours was purine ribonucleotide biosynthesis, with four genes, purD, purH, purL and purE, found to be essential at this time point only. TraDIS-Xpress did not identify mutants in any of these genes in biofilms sampled at 48 hours, whereas several reads mapped to these loci under planktonic conditions, as well as under both biofilm and planktonic conditions earlier at 12 and 24 hours. Visualisation of a ΔpurD mutant biofilm under flow conditions saw poor biofilm formation and no microcolony formation at any time compared to the wild type (Figure 4). Additionally, ΔpurD and ΔpurE mutants were deficient in biofilm biomass production, curli biosynthesis, and ΔpurE also showed reduced cell aggregation (Figure 3 a,b,c), confirming an important role for purine biosynthesis in matrix production and curli biosynthesis in the mature biofilm.

The flagella master regulatory system flhDC was identified as important in the mature biofilm. Biofilms sampled after 48 hours saw fewer flhC mutants, while insertions interpreted as over-expressing flhD increased in numbers both at the 24 and 48 hour time points, compared to
planktonic conditions. No mutants in *flgD* and *fliE*, encoding flagellar filament proteins, were identified at 24 and 48 hours, respectively. It has previously been shown that motility is important for initial biofilm formation, but this may not relate to biomass formation where no differences were seen for ∆*flhD*, ∆*flhC*, ∆*fliE* and ∆*flgD* mutants.

Various pleiotropic transcriptional regulators were also important in the mature biofilm. This included the H-NS antagonist *leuO*. Increased insertions upstream of *leuO* under biofilm conditions after 12 hours growth, as well as no *leuO* mutants in 48 hour biofilms, indicated it was beneficial to biofilm formation. A ∆*leuO* mutant did not aggregate as well as the wild type, and one ∆*leuO* mutant had reduced biofilm biomass (figure 3a,b). The ∆*leuO* mutant biofilm under flow conditions demonstrated an inability to form microcolonies after 48 hours growth (figure 4). The leucine-responsive global regulator *lrp* and a transcriptional regulator responsible for survival under acid stress, *gadW* were also found to have fewer mutants in the 48 hour biofilm compared to the planktonic condition, indicating their importance in the mature biofilm. Reduced biofilm biomass, aggregation and curli biosynthesis were observed for one copy of ∆*lrp*, but no differences in biofilm formation or aggregation were seen for ∆*gadW* mutant biofilms (figure 3a,b,c).

Inactivation of the outer membrane channels *mscL*, *tolC* and *ompF* was not tolerated in the mature biofilm grown for 48 hours. This would indicate the importance of transport in the mature biofilm, however inactivation of *tolC* and *ompF* did not result in a change in biofilm biomass (figure 3a).

Three genes involved in cell division or DNA replication, *crfC*, *ychM* and *truA*, were identified as important in the 48 hour biofilm. Expression of *yhcM*, involved in cell division, was predicted to be essential in the 48 hour biofilm, but this was not reflected in the phenotype of deletion mutants (Figure 3a). The pseudouridine synthase *truA*, also reported to be involved in cell division was found to be essential in the mature biofilm grown for 24 and 48 hours. Although a ∆*truA* mutant showed no change in biofilm biomass or curli biosynthesis, aggregation was significantly improved.
compared to the wild type albeit to a small degree (figure 3b), contrary to the prediction generated by TraDIS-Xpress.
Discussion

We have characterised the essential genome of *E. coli* biofilms across the biofilm lifecycle by using the high throughput transposon mutagenesis screen TraDIS-Xpress. The identification of genes and pathways already described to be involved in biofilm formation validates the efficacy of this experimental model. The early biofilm established 12 hours after inoculation was characterised by genes involved in adhesion. The 24-hour biofilm required both adhesion and matrix production, transitioning into matrix production being of the upmost importance in the mature biofilm after 48 hours. Control of fimbriae expression and motility remained important at each stage of the biofilm life cycle rather than just being involved in initial attachment.

As well as identifying known pathways, TraDIS-Xpress was also able to identify genes not previously known to be involved in biofilm formation, including *yigZ, ykgJ, ychM, maoP, truA, crfC*. We found that expression of *maoP* was detrimental to the fitness of biofilms grown for 24 hours, but a Δ*maoP* mutant biofilm had reduced biofilm biomass and reduced curli biosynthesis compared to the wild type. Chromosomal organisation of the Ori macrodomain requires both *maoP* and *maoS*, however no signal is seen in our data for *maoS*. This warrants further investigation into how chromosomal macrodomain organisation affects biofilm formation. The importance of cell division in the mature biofilm was further supported by our findings of fewer *ychM* and *truA* mutants compared to planktonic conditions, both of which have not before been implicated in biofilm formation. However, deletion of either gene had no effect on biofilm biomass or curli production, and the Δ*truA* deletion mutant had improved aggregation compared to the wild type. Genes involved in cell division are clearly important for the fitness of the mature biofilm, but the essential roles of *ychM* and *truA* in this process are currently unclear but unlikely to relate to crude biomass production.

Expression of *dsbA* and repression of *dksA* was found in this study to benefit early biofilm fitness. Based on previous studies and phenotypic analysis of knockout mutants in this study, we believe the increase in biofilm fitness seen is due to increased adhesion in these mutants. This study
has highlighted the benefit of close temporal gene regulation in the biofilm, where the expression
of certain genes can have a different effect on biofilm fitness at different stages of the biofilm life
cycle. For example, we found that dsbA was important for the early biofilm, but a dsbA mutant
biofilm had increased curli expression and increased aggregation. Expression of dsbA has been
previously found to result in repression of the curli regulator csgD and curli subunit csgA, essential
for optimal fitness of the mature biofilm. Conversely, we found that expression of the
transcription factor dksA was detrimental in the early biofilm, whilst a dksA knockout biofilm had
reduced biofilm biomass, reduced curli biosynthesis and reduced aggregation. The relationship
between dksA expression and curli biosynthesis has been previously characterised in similar
studies. Again, these data show differential expression of important genes at different stages
of the biofilm life cycle is essential for optimising biofilm fitness.

Purine biosynthesis was found to be important in the mature biofilm, through the essentiality of
purD, purE, purL and purH in biofilms grown for 48 hours. Similar findings have previously been
described in another transposon mutagenesis experiment in uropathogenic E. coli. Inactivation
of purine biosynthetic genes was also found to impair biofilm formation in Bacillus cerus, but this
was thought to be due to reduced extracellular DNA in the biofilm matrix. Extracellular DNA is
thought to aid adhesion and has been found to be important in the biofilms of a wide range of
bacterial species. Our data suggest the importance is in the mature biofilm rather than initial
adhesion. A relationship between both purine and pyrimidine biosynthesis and curli production in
the biofilm has been reported. More recently, curli biosynthesis in a purL mutant was
reported to be abrogated through addition of inosine, which is involved in the de novo purine
biosynthetic pathway for production of adenosine monophosphate (AMP) and guanine
monophosphate (GMP). This suggests that nucleotide production itself, rather than the
regulatory effects of the genes involved, affects curli biosynthesis, supporting one hypothesis that
disruption of the purine biosynthetic pathway may directly result in a reduction of cyclic-di-GMP,
known to regulate biofilm formation and affect curli biosynthesis. Quantification of intracellular c-
di-GMP or further investigation of other c-di-GMP-dependent pathways in these mutants would uncover the relationship between these pathways and biofilm formation.

TraDIS-Xpress data suggested that expression of $fimB$ and deletion of $fimE$ was necessary for optimal biofilm fitness at all points in the biofilm, rather than just for initial attachment. Analysis of $\Delta fimB$ and $\Delta fimE$ deletion mutants found no significant change in biofilm biomass and reduced cell-cell aggregation. Previous work has observed a positive correlation between type 1 fimbriae and exopolysaccharide production in the mature biofilm, but the increase in biofilm biomass to support this was not seen in our study. These data suggest that for a population the ability to present cells both with and without fimbriae is beneficial for fitness throughout the life cycle.

Analysis of biofilms under flow conditions found that $\Delta lrhA$ and $\Delta tomB$ mutant biofilms had a similar appearance after 12 hours growth, could potentially indicate a similar role in the biofilm. The role of $lrhA$ in motility regulation has been well documented, and expression of $tomB$ has been seen to reduce motility through repression of $fliA$. We found deletion of either $lrhA$ or $tomB$ resulted in reduced aggregation. Although $\Delta lrhA$ and $\Delta tomB$ deletion mutants shared many similar phenotypes, TraDIS-Xpress data predicted that $tomB$ was beneficial and $lrhA$ was detrimental to biofilm formation at 12, 24 and 48 hours. Previous studies on $\Delta lrhA$ mutant biofilms have reported increased adhesion, aggregation and biomass compared to the wild type. This was not seen in our study, but we did find aggregation of the $\Delta lrhA$ mutant changed over time, with increased aggregates seen in biofilms grown in flow cells after 12 hours (figure 4). However, decreased aggregation was observed in planktonic cultures after 24 hours (figure 3). The initial benefit to biofilm formation resulting from the inactivation of $lrhA$ predicted by the TraDIS-Xpress data agrees with the initial phenotype of the $\Delta lrhA$ biofilm in the flow cells. This may be due to reduced induction of $fimE$ by LrhA, thereby allowing expression of type 1 fimbriae to facilitate adhesion.

We have already described how expression of both $fimB$ and $fimE$ is necessary for optimal fitness of the mature biofilm, and the effect of $lrhA$ on biofilm formation correlates with these findings, with reduced aggregation in $\Delta lrhA$ biofilms after 24 hours (also seen in $fimB$ and $fimE$ mutants) and no
microcolony formation under flow conditions at 24 and 48 hours. Therefore, the importance of *lrhA*

to biofilm formation clearly appears to be time dependent, with the most important role in early

events.

Studies on the effect of *tomB* on biofilm formation have focused on its toxin-antitoxin relationship

with *hha*, which has been found to reduce expression of fimbrial subunit *fimA* and activate

prophage lytic genes causing cell death 67. Deletion of *hha* was found to reduce motility through

*fliDC* and increase curli production through *csgD* 68. We found no obvious benefit to biofilm fitness

with insertional inactivation of *hha*, but this may not be visible in our data due to these mutants

having a functional copy of *tomB*. The role of *tomB* in the biofilm has not previously been

suggested, but we predict *tomB* is involved in regulation of adhesion in the early biofilm and matrix

biosynthesis in the late biofilm.

The relationship between motility and biofilm formation is complex. Although it is widely understood

that motility is crucial for biofilm formation 47,48, it is also true that motility and curli production have

an inverse relationship, where *csgD* directly represses *fliE* 69 and induces c-di-GMP synthesis

through *adrA*, which reduces motility through *ycgR* 70-72. This is often referred to as a lifestyle

‘switch’, where biofilm matrix production represses motility for a motile-to sessile lifestyle transition

73. We found that although insertional inactivation of negative motility regulators *lrhA* and *hdfR*

improved biofilm fitness according to the TraDIS-Xpress data, a Δ*hdfR* deletion mutant actually

had reduced biofilm biomass, and deletion of either *lrhA* or *hdfR* impaired cell aggregation. Our

data found an important role for structural flagella components in the mature biofilm. Previous work

has suggested that flagella filaments are important for initial attachment and adhesion 74, however

we did not find this to be the case, with the expression of flagella filaments only appearing to

increase biofilm fitness in the mature biofilm grown for 48 hours. It appears that regulation of

flagella and motility, rather than their fixed expression or absence, is important for optimal biofilm

fitness.
Previous genome-wide screens on *E. coli* biofilms have identified some of the same genes as this study \(^{26,29,31}\). The TraDIS-Xpress technology used here makes for a more powerful analysis of biofilm formation by predicting roles of changes in gene expression as well as essentiality and by analysing important genes over time. Differences between this work and previous studies may reflect biofilms being grown under different conditions on different surfaces, as these environmental factors greatly affect the pattern of gene expression and gene essentiality in the biofilm \(^{75}\). Analysis of more strains and species, grown on different abiotic and biotic surfaces under a range of environmental conditions may provide a wider list of essential genes for biofilm formation shared amongst a majority of human pathogens, as well as substrate-, condition- and species-specific genes and pathways for specific industrial, clinical and drug-development applications. As well as temporal changes in gene expression, spatial changes have been shown to affect biofilm development \(^{76}\). Integration of the spatial component into this model, to assay how gene expression throughout the biofilm over time affects biofilm fitness, would be the next logical step in furthering our understanding of biofilm development.

This study had revealed important time-specific roles for known and novel genes in biofilm formation. We reveal some pathways have a more important role in the mature biofilm than previously appreciated and identify genes with time dependent conditional essentiality within the biofilm. We also identify potential new candidate genes essential for biofilm formation, which could be targeted for novel anti-biofilm therapies. Further work using high-density transposon mutant libraries across time and in different conditions is likely to further our understanding of biofilm biology.
Methods

Transposon mutant library

The *E. coli* BW25113 transposon mutant library containing over 800,000 distinct mutants that was used in this study has recently described \(^{32}\). The transposon used to construct this library incorporates an outward-transcribing IPTG-inducible promoter.

Biofilm model conditions

The library was used to inoculate parallel cultures of 5 mL LB broth (without salt) with approximately $10^7$ cells. Cultures were grown in 6-well plates containing 40 sterile 5 mm glass beads per well (Sigma, 18406). Two replicates were set up, with or without 1 mM IPTG. Plates were incubated at 30 °C with light shaking for 48 hours. After 12, 24, and 48 hours of incubation, 2 mL of planktonic sample was collected from each culture and 70 beads were taken to constitute the biofilm sample. Beads were washed twice in sterile PBS and vortexed in tubes containing PBS to resuspend cells from the biofilm. Both planktonic and biofilm samples were centrifuged at 2100 x g to form pellets for DNA extraction.

TraDIS-Xpress Sequencing

DNA was extracted from pellets following the protocol described in Trampari, et al. \(^{77}\). A customised sequencing library was prepared to identify transposon insertions. DNA was tagmented using a MuSeek DNA fragment library preparation kit (ThermoFisher) and customised Tn5-i5 and i7 primers were used in PCR for 28 cycles \(^{32}\). DNA fragments of 300-500 bp were size selected using AMPure beads (Beckman Coulter) and nucleotide sequences were generated using a NextSeq 500 and a NextSeq 500/550 High Output v2 kit (75 cycles) (Illumina).

Informatics

Fastq files were aligned to the *E. coli* BW25113 reference genome (CP009273) using the BioTraDIS (version 1.4.3) software suite \(^{78}\) using SMALT (version 0.7.6). Insertion frequencies were determined per gene using tradis_gene_insert_sites within the BioTraDIS toolkit. Insertion
frequencies per gene for each replicate were plotted against each other to determine the experimental error between replicates as well as differences in insertion frequency between control and test conditions (supplementary figure 1). The tradis_comparison.R command (also part of the BioTraDIS toolkit) was also used to determine significant differences ($p < 0.05$, after correction for false discovery) in insertion frequencies per gene between control and test conditions. For all candidate loci, plot files generated by BioTraDIS were also examined manually in Artemis (version 17.0.1) to confirm the results from these two approaches, as well as to identify regions where inserts were under differential selection but did not fall within coding regions of the genome.

Validation experiments

Knockout mutants for genes identified by TraDIS-Xpress data were sourced from the Keio collection and tested for their biofilm-forming abilities. Crystal violet assays, used to assess biofilm biomass production, were undertaken by inoculating $10^4$ of each mutant strain into 200 μL LB broth without salt in a 96-well plate. After 48 hours incubation at 30 °C, the culture was removed, wells were rinsed with water, and the residual biofilms were stained for 10 minutes with 200 μL 0.01% crystal violet. The plate was then rinsed with water to remove the stain and 200 μL 70% ethanol was added to the wells to solubilise the stained biofilm. The optical density (OD) was measured using a FLUOstar Omega plate reader (BMG Labtech) at 590 nM. Cell aggregation was measured by leaving overnight cultures (normalised to an OD$_{600}$ nm of 0.3) on an unagitated surface at room temperature. After 24 hours, the supernatant of each culture was removed by pipetting, diluted in PBS and measured in a plate reader at 600 nm. Biofilm matrix composition was investigated through spotting 10 μL of each mutant (representing $10^5$ CFU) on agar supplemented with 40 μg/mL Congo red (Sigma, C6277) to examine curli production. Plates were incubated at 30 °C for 48 hours and photographed to compare mutant biofilm composition to the wild type.

Adhesion and biofilm architecture were investigated under flow conditions for selected mutants using the Bioflux system. Flow cells were primed with LB broth without salt at 5 dyne/cm$^2$ and seeded with approximately $10^7$ cells. The plate was left at room temperature for 2.5 hours to allow attachment, and subsequently incubated at 30 °C at a flow rate of 0.3 dyne/cm$^2$. After 12, 24 and
419 48 hours, biofilms were visualised with an inverted light microscope and representative images at
420 10x, 20x and 40x magnification were taken at three locations of the flow cell. Experiments were
421 performed in duplicate.
Data availability

Sequence data supporting the analysis in this study has been deposited in ArrayExpress.
Temporary accession number for review E-MTAB-9873.

Code availability statement

All software packages used are described in the methods. No bespoke code was used in this study.

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Author contributions

ERH designed and performed experiments, analysed the data and wrote the paper. AKT and IGC helped design experiments and wrote the paper. MY helped design experiments. JW helped analyse data and wrote the paper. MAW designed the experiments, analysed the data and wrote the paper.

Competing interests

The authors have no competing interests.


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Figures

Figure 1: Genes involved in biofilm formation over time in *E. coli*. Expression of genes in green were found to be beneficial for, and those in red were found to be detrimental to, biofilm formation.

*E. coli*
Figure 2: Mapped reads from TraDIS-Xpress data, plotted with BioTraDIS in Artemis. 

a) Insertion sites in and around *fimB* and *fimE* in planktonic and biofilm conditions after 12 and 48 hours growth. 

b) Insertion sites in and around *hdfR*, *lrhA* and *tomB* in planktonic and biofilm conditions after 24 hours growth. 

c) Insertion sites in and around *dksA* and *dsbA* in planktonic and biofilm conditions after 12 and 48 hours growth. For all plot files, one of two independent replicates is shown and the transposon-located promoter is induced with 1mM IPTG in all conditions shown.
**Figure 3**: Phenotypic validation of selected genes involved in biofilm formation. **a**) Biofilm biomass of single knockout mutants relative to wild type *E. coli* BW25113, measured by crystal violet staining. Two biological and a minimum of two technical replicates were performed for each mutant. **b**) Cell aggregation of single knockout mutants relative to wild type *E. coli* BW25113, measured by OD$_{600\,\text{nm}}$ of the supernatant of unagitated cultures. Points show the ODs of three independent replicates. For both graphs, red points/bars show data from the first KEIO collection mutant of each gene, and blue points/bars show data from the second mutant. Error bars show 95% confidence intervals and the shaded area shows the 95% confidence interval of the wild type. Single asterisks (*) represent a significant difference between one Keio mutant copy and the wild type, and double asterisks (**) denote a significant difference between both Keio mutant copies and the wild type (Welch's *t*-test, *p* < 0.05). **c**) Colonies grown on agar supplemented with Congo red to compare curli biosynthesis between single knockout mutants and the wild type. Images are representative of 2 biological and 2 technical replicates.
a) Biofilm biomass relative to wild type (CV Absorbance 595 nm)

b) Absorbance of supernatant relative to WT (OD600)

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<th>WT</th>
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<th>tomB</th>
<th>maoP</th>
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698
**Figure 4:** Biofilm formation of single knockout mutants on glass analysed under flow conditions after 12 and 48 hours growth. 10x Magnification. Images are representative of two independent replicates. Scale bar indicates 5 µm.

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**Figure 5.** Summary of genes important for biofilm formation by *E. coli* at different stages of development.