

The apolipoprotein N-acyl transferase Lnt is dispensable for growth in *Acinetobacter* species

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Abstract

Directing the flow of protein traffic is a critical task faced by all cellular organisms. In Gram-negative bacteria, this traffic includes lipoproteins. Lipoproteins are synthesized as precursors in the cytoplasm and receive their acyl modifications upon export across the inner membrane. The third and final acyl chain is added by Lnt, which until recently was thought to be essential in all Gram-negatives. In this report, we show that *Acinetobacter* species can also tolerate a complete loss-of-function mutation in *lnt*. Absence of a fully functional Lnt impairs modification of lipoproteins, increases outer membrane permeability and susceptibility to antibiotics, and alters normal cellular morphology. In addition, we show that loss of *lnt* triggers a global transcriptional response to this added cellular stress. Taken together, our findings provide new insights on and support the growing revisions to the Gram-negative lipoprotein biogenesis paradigm.

INTRODUCTION

The Gram-negative cell envelope is composed of an inner membrane (IM) and an outer membrane (OM), with the aqueous periplasm in between [1]. The peptidoglycan cell wall is contained within the periplasm [2]. Biogenesis of the cell envelope relies on the activity of multiple proteins, many of which function as part of multi-subunit molecular machines [3–5]. For example, delivery of lipopolysaccharide (LPS) to the outer leaflet of the OM requires the Lpt machine [3]. Outer membrane protein (OMP) assembly requires the Bam complex [6], while lipoproteins are sorted between the IM and OM by the Lol transporter [7]. In model Gram-negative organisms like *Escherichia coli*, the functions of these assembly factors are essential under standard laboratory conditions [8]. However, there is increasing evidence that the requirement for these factors varies among different Gram-negative species and under particular conditions [9–12].

Lipoproteins are synthesized as precursors in the cytoplasm and then targeted for export across the IM [7]. These precursors then undergo an ordered series of modifications to install the lipid moieties. First, the diacylglycerol transferase Lgt adds two acyl chains to a conserved cysteine residue

adjacent to the amino-terminal signal sequence [13]. Next, the signal sequence is cleaved by the signal peptidase LspA [14]. Finally, the third acyl chain is added to the amino terminus of the cysteine residue by the apolipoprotein N-acyl transferase Lnt [15]. In *E. coli*, lipoproteins with an aspartate residue at position +2 will be retained at the IM [16, 17]; in other cases, the +3 or +4 residue guides IM retention [18]. Lipoproteins destined for the OM are routed there through the Lol pathway [19].

The Lol pathway consists of an IM platform composed of the LolCDE proteins, which functions as an ABC transporter [20]. Fully acylated lipoproteins are routed to LolCDE. If destined for the OM, lipoproteins are then delivered to the periplasmic carrier LolA which carries them to the OM receptor LolB for assembly into the OM [21, 22]. Lipoproteins can remain anchored to the inner leaflet of the OM, or in some cases may flip to the outer leaflet of the OM [23]. This flipping mechanism is an area of active research interest [4, 19]. Recently, the classical model of how Lol functions has been challenged [12]. In some mutant strains of *E. coli*, the need for both LolA and LolB can be bypassed upon activation of an envelope stress response [12]. Grabowicz and Silhavy also propose the existence of a redundant

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Abbreviations: ABC, ATP-binding cassette; IM, Inner membrane; LPS, lipopolysaccharide; OM, outer membrane; PCR, polymerase chain reaction.

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Three supplementary tables are available with the online version of this article.

machinery for lipoprotein trafficking, and that the true role of Lol is to ensure that the OM does not become overloaded with lipoproteins [12]. However, the identity of this novel machinery has not yet been revealed.

In *E. coli*, all three lipoprotein modification factors – Lgt, Lsp, and Lnt – as well as the Lol machinery, are essential for growth under standard laboratory conditions [8]. Until recently, this was thought to be the case for all Gram-negative bacteria. However, this no longer appears to be the case. The first demonstration that *lnt* is dispensable for growth came from work in *Francisella* and *Neisseria* species [10]. Here, we show that Lnt is not essential but important for growth in *Acinetobacter baylyi* or *Acinetobacter baumannii*. These two organisms have recently emerged as models for studying complex genetic interactions and pathogenesis of drug-resistant microbial infections, respectively [24, 25]. We also found that the impact of crippling lipoprotein biogenesis by mutating *lnt* is broad. Without Lnt, the bacterial cell envelope becomes more permeable and modification of OM lipoproteins is impaired. Given the central importance of lipoproteins during cell envelope biogenesis, we reasoned that the absence of Lnt would impair the flow of lipoprotein traffic and thus trigger a cellular stress response. By using RNA-seq and quantitative PCR, we identified nearly 80 genes whose expression is altered in the absence of *lnt*. Taken together, our findings support a key role for Lnt in lipoprotein maturation, as well as providing a foundation for future studies of stress responses in *Acinetobacter* species. Defining the lipoprotein biogenesis pathways in *A. baylyi* and *A. baumannii* is of fundamental importance to elucidating the biology of these important Gram-negative microbes.

METHODS

Strains and growth conditions

All strains of *Acinetobacter* were grown in LB (Lennox, Fisher Scientific); *A. baylyi* strains were grown at 30°C, while *A. baumannii* strains were grown at 37°C. *A. baumannii* AB5075-UW wild-type and *lnt* mutant (*lnt*153::T26) strains were obtained from the University of

Washington [26]. To determine colony morphology, overnight cultures of each strain were serially diluted and then aliquots were spread on the surface of LB agar or MacConkey agar plates. Plates were incubated overnight and photographed the next morning. For liquid growth curves, overnight cultures of each strain were diluted in fresh LB to an optical density (OD) at 600 nm of 0.05. Cultures were incubated with intermittent shaking, and OD measurements were taken every 45 min. All strains were grown in triplicate.

Construction of an *A. baylyi lnt::kan* insertion-deletion mutant and complementation experiments

To construct the *A. baylyi lnt* mutant, we used an overlap-extension PCR strategy using the primers listed in Table S1 (available in the online version of this article) [27]. Briefly, approximately 1 kb of upstream and downstream sequences flanking *lnt* (ACIAD0415) were amplified by colony PCR from the *A. baylyi* wild-type reference strain ADP1 (obtained from ATCC). A 795 bp kanamycin-resistance cassette was amplified from plasmid pIM1445 (gift from Ichiro Matsumura) [28]. These three products were joined together in a final PCR using the outermost upstream and downstream flanking primers, resulting in the *lnt::kan* insertion-deletion allele. This PCR fragment was purified using a QIAGEN PCR purification kit and then used to transform wild-type *A. baylyi*, which is naturally competent [25]. Correct incorporation of the insertion-deletion allele was confirmed by PCR analysis (Fig. 1b, c) and DNA sequencing. To construct an *lnt* complementation strain, the full-length sequence of *lnt* (Table S2) was cloned into the BamHI site of pWH1266 (ATCC) by custom gene synthesis (GenScript, Piscataway, NJ). The resulting plasmid pLnt was transformed into *A. baylyi* by electroporation using a Bio-Rad MicroPulser.

Immunoblot analysis

Bacterial cultures were grown until mid-exponential phase when 1 ml samples were harvested by centrifugation. Cell pellets were then re-suspended in SDS-PAGE sample buffer in a volume (in ml) equal to the OD_{600 nm} divided by seven. After boiling for 10 min, 10 µl of each sample was resolved

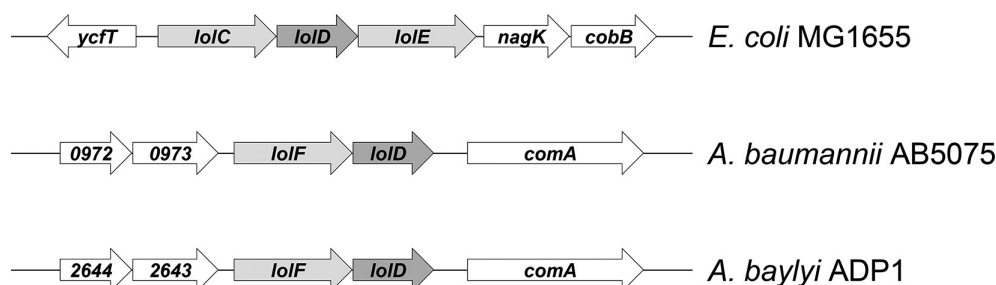


Fig. 1. Organization of Lol ABC transporter genes in Gram-negative bacteria. In *E. coli*, the inner membrane (IM) Lol ABC transporter is organized in the standard *lolCDE* format. In *Acinetobacter* species, the ABC transporter is organized in a *lolFD* module. Genes that flank *lolFD* in *A. baumannii* AB5075-UW and *A. baylyi* ADP1 are listed by number.

by SDS-PAGE. Proteins were transferred to a nitrocellulose membrane and probed with polyclonal rabbit antibodies against BamD, RpoD and SecY. Antibodies were generated against His-tagged, purified full-length proteins (BamD and RpoD) or peptides (DNVALARFFKANEGC, SecY) by GenScript (Piscataway, NJ). Blots were probed with antibodies that were diluted 1:1000 in Blocking Buffer (LI-COR Biosciences) overnight at 4°C. Primary antibodies were detected using IRDye 800CW donkey anti-rabbit antibodies (LI-COR Biosciences) prior to imaging with a LI-COR Odyssey Fc imager according to the manufacturer's protocols.

Scanning electron microscopy

Wild-type and *lnt* mutant *A. baylyi* strains were grown overnight on LB plates. Plugs of individual colonies were harvested and then fixed with 3% glutaraldehyde (in 0.1 M phosphate buffer, pH 7.2) for one hour at room temperature. The glutaraldehyde solution was removed and then samples were washed three times with phosphate buffer. Samples were then dehydrated in an ascending ethanol series (30, 50, 70, 80, 85, 90 and 95% ethanol for 10 min each and three 100% ethanol washes for 15 min each). After dehydration, the samples were dried using a Samdri-795 critical-point drier. Samples were secured onto aluminum stubs with double-sided adhesive tabs and were gold-coated using an EMS-550 sputter coater. All samples were visualized using an FEI Quanta 250 scanning electron microscope at 30 kV.

Disc diffusion antibiotic sensitivity assays

Aliquots of 100 µl from overnight cultures of each strain were spread onto the surface of LB agar plates and allowed to dry. Antibiotic-containing discs (BD BBL Sensi-Discs) were placed on the agar surface, and zones of growth inhibition around the discs were measured the next day. Discs contained the following amounts of antibiotics: bacitracin (10 units), novobiocin (30 µg), vancomycin (30 µg) and polymyxin B (300 units).

RNA-seq analysis

All RNA-seq experiments were conducted with Cofactor Genomics (St. Louis, MO), using the following workflow.

Ribosome-depletion and library preparation

Triplicate cultures of both wild-type and *lnt* mutant *A. baylyi* were grown in LB at 30°C and harvested during mid-exponential phase. Total RNA was extracted using the RNeasy kit (Qiagen) following the manufacturer's instructions. Total RNA was processed for library construction by Cofactor Genomics (<http://cofactorgenomics.com>, St. Louis, MO) as follows. Species-specific rRNA probes were hybridized to total RNA to remove any ribosomal RNA, and the resulting ribosome-depleted RNA was then fragmented. First-strand cDNA synthesis was performed using reverse transcriptase and random primers in the presence of Actinomycin D, followed by second-strand cDNA synthesis with DNA polymerase I and RNase H. Double-stranded cDNA was end-repaired and A-tailed for subsequent

adaptor ligation. Indexed adaptors were ligated to the A-tailed cDNA. Enrichment by PCR was performed to generate the final cDNA sequencing library. The library was sequenced as single-end 75 base pair reads on an Illumina NextSeq 500 following the manufacturer's protocols.

Quality control and data analysis

Initial quality control was performed by Cofactor Genomics (<http://cofactorgenomics.com>, St. Louis, MO). Raw sequence data in FASTQ format were assessed for quality using FastQC (<http://www.bioinformatics.babraham.ac.uk/projects/fastqc/>) and ribosomal RNA content using sortmeRNA (<http://bioinfo.lifl.fr/RNA/sortmerna/>). NovoAlign (Novocraft) was used to align reads to a set of transcript sequences. Alignments to the genome were performed using STAR (<https://github.com/alexdobin/STAR>). Only unique alignments to the genome were allowed. The genome alignment loci from all samples were combined and clustered to generate contiguous read coverage. The reads per kilobase of transcript per million mapped reads (RPKM) expression value was calculated for each sample and used as the basis for expression comparison and statistical analysis. The resulting comparative expression data were visualized in ActiveSite (Cofactor Genomics). All raw data were deposited via SRA (SRP133498).

Quantitative RT-PCR (qPCR)

RNA was extracted as described above. cDNA was synthesized using the iScript reverse transcriptase kit (Bio-Rad). The primers used for expression analysis are listed in Table S2. qPCR was carried out using POWER SYBR Green Supermix on a StepOnePlus thermocycler (Applied Biosystems), with the 16S rRNA gene as the reference. Reactions were set up according to the manufacturer's protocols using 500 nM primers and 2 µl of the cDNA template (diluted 1:10). Relative expression was determined using the comparative cycle threshold ($\Delta\Delta C_T$) method [29]. The cycling conditions used were as follows: amplification stage – 95°C for 10 min and then 40 amplification cycles of 95°C for 30 s, 60°C for 1 min and 72°C for 30 s; melting curve stage – 95°C for 15 s, 60°C for 1 min and 95°C for 15 s. No-template and no-reverse transcriptase reactions served as the negative controls. All reactions were carried out in triplicate, using cDNA derived from triplicate cultures.

RESULTS AND DISCUSSION

Acinetobacter lnt mutants are viable, but exhibit growth defects

Based on the work of LoVullo *et al.*, in organisms where the LolCDE ABC transporter is instead organized as LolFD, *lnt* is dispensable for growth [10]. Examination of the *Acinetobacter baylyi* and *Acinetobacter baumannii* genomes revealed the existence of this LolFD format (Fig. 1 and [10]). Therefore, we hypothesized that *lnt* mutants would be viable in *Acinetobacter* spp. To test this hypothesis, we created an insertion-deletion mutant in the *A. baylyi lnt* gene ACIAD0415 [27]. We obtained kanamycin-resistant

colonies and confirmed integration of the insertion-deletion allele into the chromosome by PCR and DNA sequence analysis (Fig. 2). Compared to wild-type *A. baylyi*, colonies of the *lnt* mutant were much smaller when grown on LB agar plates (Fig. 3a, upper panels). The *lnt* mutant also displayed a growth defect when cultured in broth (Fig. 3c). Notably, a previous study reported that *lnt* is essential in *A. baylyi* [30]. The mutants constructed in this study were cultured using a defined minimal medium. In the experiments we report here, the *A. baylyi lnt* mutant was generated on a rich medium and displays a profound growth defect. We have found that growth of the *lnt* mutant on minimal medium is severely impaired; normal-sized colonies appear only after 3 days of incubation, so the *lnt* mutant could have been overlooked and incorrectly labelled as essential for growth.

To determine whether *lnt* was dispensable in other species of *Acinetobacter* that possess a LolFD module, we searched the *A. baumannii* AB5075-UW transposon mutant library collection browser and found three confirmed insertions within the *lnt* gene (<http://www.gs.washington.edu/labs/manoil/baumannii.htm>) [26]. Similar to our observations with *A. baylyi*, the *lnt* mutant of *A. baumannii* AB5075-UW showed a growth defect on LB agar and a diminished generation time in liquid culture (Fig. 3b, d). We also observed that the *lnt* mutants have a profound growth defect when plated on MacConkey agar (Fig. 3a, b, lower panels). MacConkey agar contains bile salts, a detergent which impairs the growth of Gram-negative bacteria with defective outer membranes [31]. Though wild-type *A. baylyi* displayed

reduced growth on MacConkey, colonies of the *lnt* mutant failed to appear even after extended incubation. Similarly, the *A. baumannii lnt* mutant was also unable to grow on MacConkey. These findings suggest that the function of *lnt*, while important, is not required to sustain growth of *Acinetobacter* species under normal laboratory conditions. However, when bacteria are subjected to membrane-disrupting detergents like bile salts, the function of *lnt* is vital.

The OM permeability barrier is compromised in the absence of *lnt*

The growth defects associated with the loss of *lnt* under standard laboratory conditions are consistent with disruption of an important cellular process – namely, the final acylation step of lipoproteins prior to their sorting between the IM and OM. It has already been shown in *Francisella* that without *lnt*, trafficking of diacylated lipoproteins to the OM is diminished [10]. Many OM lipoproteins are critical for establishing the barrier function of the OM. Indeed, this barrier function is compromised in both *Francisella* and *Neisseria lnt* mutants as evidenced by slightly increased sensitivity to different antibiotics [10]. Given these findings, we expected the OM to be severely compromised in the *Acinetobacter lnt* mutants. Indeed, we found that both *Acinetobacter lnt* mutants were more sensitive to antibiotics than their wild-type strains (Fig. 4). The disparity was particularly notable for *A. baylyi*. The *lnt* mutant was extremely sensitive to both bacitracin and vancomycin, which disrupt cell wall synthesis, and novobiocin, which

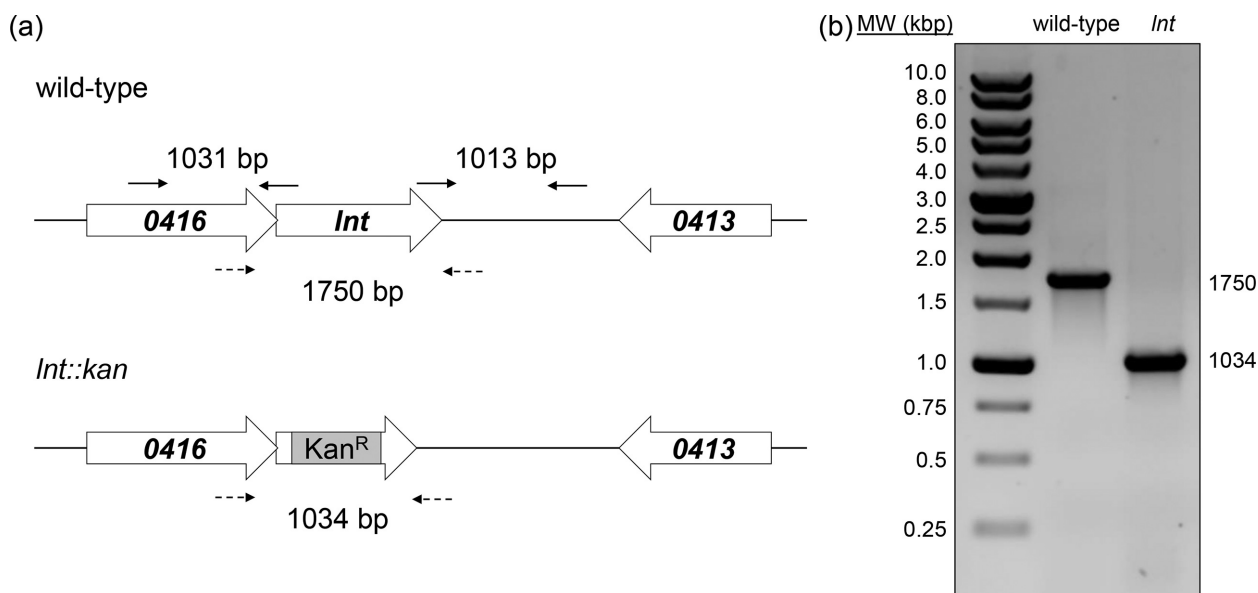


Fig. 2. Construction of an *A. baylyi lnt* mutant. (a) Genomic region of *lnt* (top panel) and location of primers (solid arrows) used to generate an *lnt::kan* insertion-deletion allele (bottom panel) as described in the Methods. Mutant construction was confirmed by PCR analysis of products amplified using primers (dashed lines) flanking *lnt* or *lnt::kan*. (b) Products were amplified from chromosomal DNA of wild-type or *lnt* mutant cells by colony PCR and resolved by agarose gel electrophoresis.

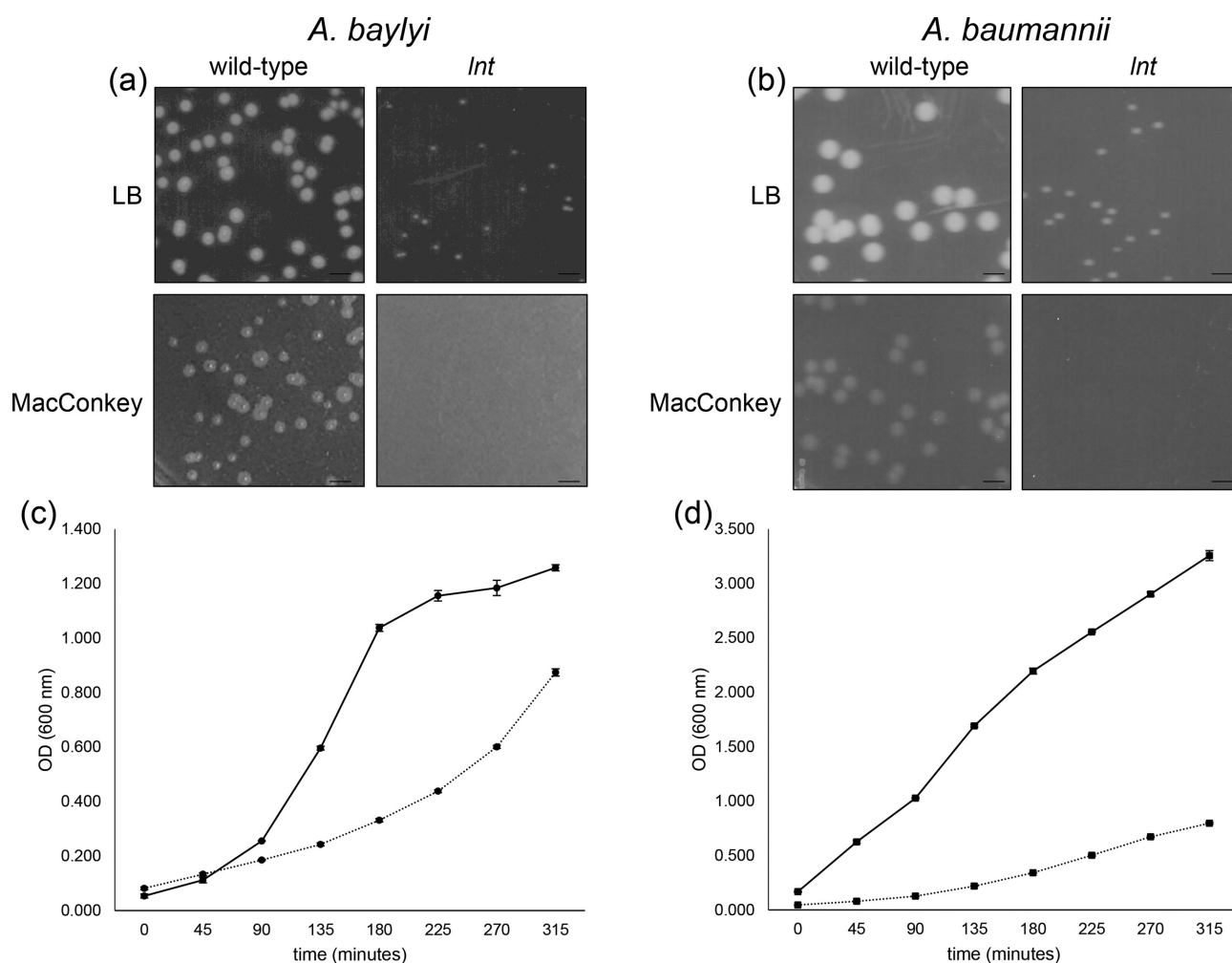


Fig. 3. Growth of *A. baylyi* and *A. baumannii* *lnt* mutants is impaired relative to wild-type. Overnight cultures of the indicated *A. baylyi* (a) or *A. baumannii* (b) strains were serially diluted and plated on either LB (upper panels) or MacConkey (lower panels) agar plates. Scale bar=3.5 mm. Growth of *A. baylyi* (c) and *A. baumannii* (d) wild-type (solid line) and *lnt* mutant (dashed line) strains was monitored by optical density (OD) at 600 nm. Shown is a representative experiment of strains grown in triplicate, where error bars indicate standard deviation from the mean.

targets DNA gyrase (Fig. 4a $P \leq 0.05$) [32–34]. In the case of *A. baumannii*, the difference in antibiotic sensitivity between wild-type and *lnt* mutant strains was considerably less pronounced; however, the *lnt* mutant was significantly more sensitive to bacitracin and novobiocin ($P \leq 0.05$) and vancomycin ($P \leq 0.067$) (Fig. 4b). Interestingly, only the *A. baumannii* *lnt* mutant displayed significantly increased sensitivity to polymyxin B ($P \leq 0.05$), which is a membrane-disrupting agent [35]. We do not fully understand the differential in sensitivities between the *A. baylyi* and *A. baumannii* strains. The *A. baumannii* *lnt* mutant contains a transposon insertion at nucleotide position 929 (of 1560). This mutation should disrupt the active site of *lnt* based on previous studies [36, 37]. The *A. baylyi* *lnt* mutant is disrupted by an antibiotic resistance cassette, replacing nucleotides 17–1554 (of 1560). However, *A. baumannii* AB5075-UW is a multidrug-resistant clinical that

produces a capsule which likely contributes to the increased antibiotic resistance profile shown in Fig. 4 [38]. A similarly diverse spectrum of antibiotic sensitivities was also noted among *lnt* mutants in *Francisella* and *Neisseria* species [10]. One of the most important functions of the Gram-negative OM is to serve as a barrier against large antibiotics. The fact that the *lnt* mutants displayed increased sensitivity to various antibiotics indicates a defective OM, particularly in *A. baylyi*. Given the stronger sensitivity phenotypes associated with the *A. baylyi* *lnt* mutant, we conducted our additional analyses with this mutant.

Proper acylation of OM lipoprotein BamD requires *lnt* in *A. baylyi*

To confirm that the *A. baylyi* *lnt* mutant cannot properly acylate precursor lipoproteins, we investigated the OM

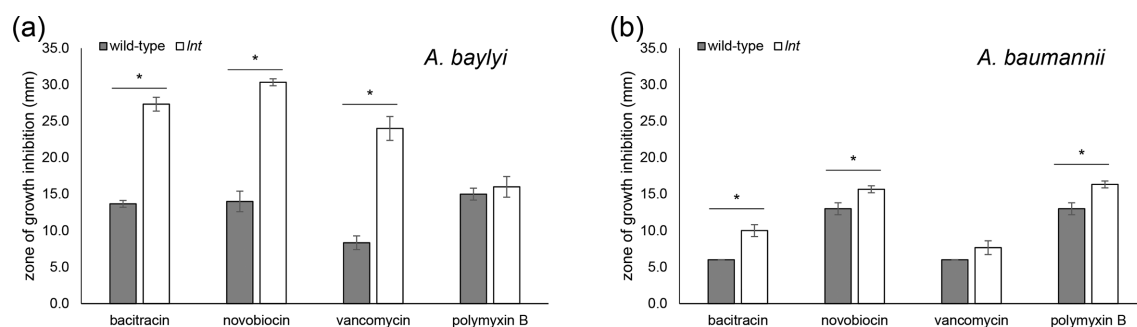


Fig. 4. In the absence of *lnt*, the integrity of the outer membrane (OM) is compromised. Overnight cultures of wild-type and *lnt* mutant for *A. baylyi* (a) and *A. baumannii* (b) strains were spread onto the surface of an LB agar plate. After allowing the cultures to dry, filter discs impregnated with the indicated antibiotics [bacitracin (10 units), novobiocin (30 µg), vancomycin (30 µg) and polymyxin B (300 units)] were placed on the agar surface. Zones of growth inhibition (in mm) were measured after overnight growth at 30 °C. Bar charts represent the mean of triplicate measurements, with error bars indicating standard deviation from the mean. *, $P \leq 0.05$ by Student's *t*-test.

lipoprotein BamD (Fig. 5). BamD is the only essential lipoprotein component of the Bam complex, which is required for assembly of OMPs in the Gram-negative OM [39]. Without functional BamD, OMP assembly is impaired which destabilizes the OM [40]. We probed whole-cell extracts of wild-type and *lnt* mutant *A. baylyi* with antibodies raised against BamD. In the absence of *lnt*, BamD migrated at a lower molecular weight than in the wild-type parent (Fig. 5a). This is consistent with failure to add the third and final acyl chain to the diacylated BamD precursor. To complement the *lnt* mutant, we used the *Acinetobacter* shuttle vector pWH1266 to express full-length *lnt*. We found that migration of BamD to a higher molecular weight was restored (Fig. 5a). The larger BamD species in the wild-type and complemented mutant runs at about 16 kDa, while the smaller BamD species in the *lnt* mutant runs at about 15 kDa. We also noticed that the overall level of BamD was slightly decreased in the *lnt* mutant. Given the central importance of BamD, this finding could partially explain the growth and permeability defects exhibited by the *lnt*

mutant. For comparison, we blotted for the cytoplasmic protein RpoD (Fig. 5b) and the IM protein SecY, demonstrating no migration or expression differences (Fig. 5c).

Cells of the *lnt* mutant have structural defects

To gain a better picture of the cell envelope defects observed in the *lnt* mutants, we examined wild-type and mutant cells using scanning electron microscopy (Fig. 6). Wild-type cells appeared as short rods with a smooth surface, typical of *A. baylyi* (Fig. 6a). We found that the overall shape and size of the *lnt* mutant cells was similar to wild-type (Fig. 6b). However, the surface of the *lnt* mutant cells evinced a rougher surface and the presence of outer membrane vesicles (OMVs). These OMVs varied in size, ranging from 70 to 200 nm in diameter, and were consistently observed across multiple fields of view. Notably, increased production of OMVs has been associated with a general response to cell envelope stress [41]. For SEM analysis, we examined microbial cells that were grown under optimal conditions, devoid of major stress stimuli. However, given the important role

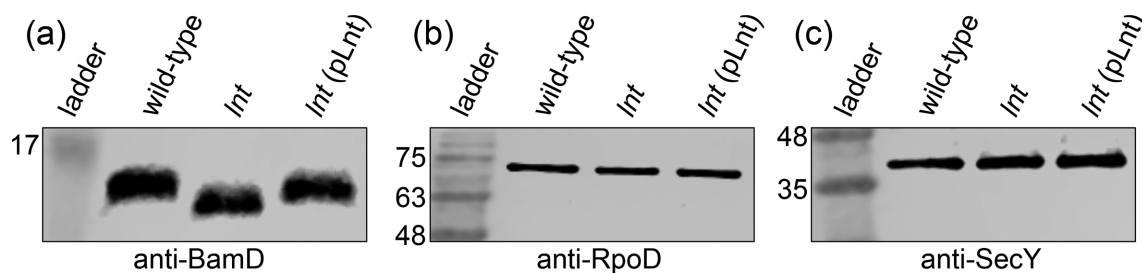


Fig. 5. *lnt* is required for full maturation of OM lipoprotein BamD. Whole-cell protein extracts derived from an equal amount of wild-type, *lnt* mutant or complemented mutant (*lnt* (pLnt)) *A. baylyi* cells were resolved by SDS-PAGE on a 12 % resolving gel and then probed using the indicated antibodies during an immunoblot. (a) BamD migrates at a lower molecular weight in the *lnt* mutant as compared to wild-type. (b) Cytoplasmic protein RpoD and (c) IM protein SecY displayed similar gel migration in wild-type and *lnt* mutant strains. Numbers indicate molecular weight in kDa.

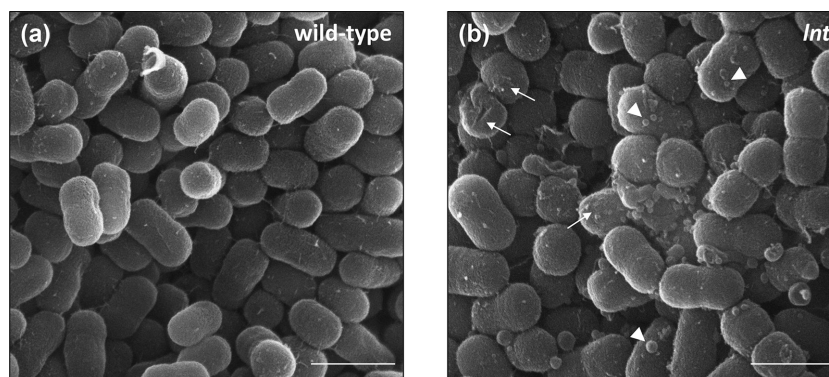


Fig. 6. Surface roughness and OMV production are increased in *lnt* mutant cells. Scanning electron microscopy images of wild-type and *lnt* mutant *A. baylyi* cells grown on LB agar plates. Arrows indicate rough surface, and arrowheads denote OMVs. Scale bar=1 μ m. Shown are representative fields from at least 6 different sample preparations.

played by lipoproteins in cell envelope biogenesis, it stands to reason that the *lnt* mutant cells would exhibit signs of stress even under the most favourable conditions. Accordingly, we postulated that the *lnt* mutant could be used as a tool to investigate cellular stress responses. In this case, the stress response in question would be triggered by a specific defect in lipoprotein biogenesis. We therefore assessed whether or not the loss of Lnt is sufficient to alter normal patterns of gene expression.

The gene expression profile of *A. baylyi* changes in response to *lnt* mutation

We examined global changes in gene expression using RNA-seq (Fig. 7). Total RNA was collected from cultures of wild-type and *lnt* mutant strains of *A. baylyi* grown under stress-free conditions (30 °C in LB broth). We considered genes with a greater than five-fold increase in expression as being induced, while genes with expression levels reduced by five-fold as being repressed ($P \leq 0.01$ as determined by Welch's *t*-test [42]). By this metric, 57 genes were upregulated in the *lnt* mutant relative to the wild-type, while 21 were repressed (Table S3). Many of these genes encode proteins with predicted roles in central metabolism, and thus their connection to envelope stress is not immediately obvious (Fig. 7a). Eleven genes have no predicted function. However, there are 17 genes associated with cell envelope biosynthesis (2), protein secretion (5), membrane proteins (6) or stress responses (4).

We selected a subset of these differentially regulated genes for confirmation by qPCR, some of which have previously been implicated in transcriptional responses to interference with lipoprotein transport [43]. This includes *hslJ*, *lola* and *rcnB*. We confirmed that expression of *lola* was upregulated in the *lnt* mutant (Fig. 7b). Given that LolA is thought to shuttle lipoproteins between the IM and OM, it is not surprising to see its expression increased. If tri-acylation is required for efficient delivery of lipoproteins to the OM, perhaps an overproduction of LolA increases delivery of di-

acylated lipoproteins. Indeed, it has already been shown that *lola* expression increases in response to defective lipoprotein sorting via the Rcs stress response and upon depletion of *lolCDE* [44, 45]. Additionally we observed that expression of *hslJ*, which encodes a putative periplasmic heat shock chaperone, is increased in the *lnt* mutant. Interestingly, a previous study also found that in *E. coli* expression of *hslJ* was upregulated upon treatment with the LolCDE inhibitor known as compound 2 [43]. This study also discovered that treatment with compound 2 induced expression of *lola* and *lolB*, as well as stress response factors *degP*, *rcsA*, and *cpxP* [43]. Thus, interference with the machinery of lipoprotein biogenesis, by either chemical or genetic means, can induce transcription of stress response genes. The response to defects in the early stages of lipoprotein maturation is an important component of how Gram-negatives counter disruptions in OM biogenesis by adjusting expression of a similar suite of genes [46–48].

We also identified differentially regulated genes with no obvious connection to lipoprotein biogenesis. For example, both *yjiM* and *rcnB* were strongly induced in the *lnt* mutant, while *cynT*, a predicted carbonic anhydrase, was repressed. Additional studies are required to fully understand the contribution of such a diverse subset of genes toward mitigating defects in lipoprotein biogenesis, particularly so that directly related factors can be separated from indirect effects. Given this wide range of targets, it seems that bacteria have a suite of genes that are used to mitigate stress caused by alteration to the typical lipoprotein composition. It is possible that this suite of genes is also activated during other conditions that trigger cell envelope stress, such as the use of membrane and cell wall-targeting antibiotics. Toward that end, it will be important to monitor gene expression when *Acinetobacter* is challenged with a different stress to the cell envelope, such as interference with OMP biogenesis or LPS assembly. Further studies will be needed to characterize the specifics of such transcriptional responses; in the meantime, *lnt* disruption may serve as an important tool to fully classify stress

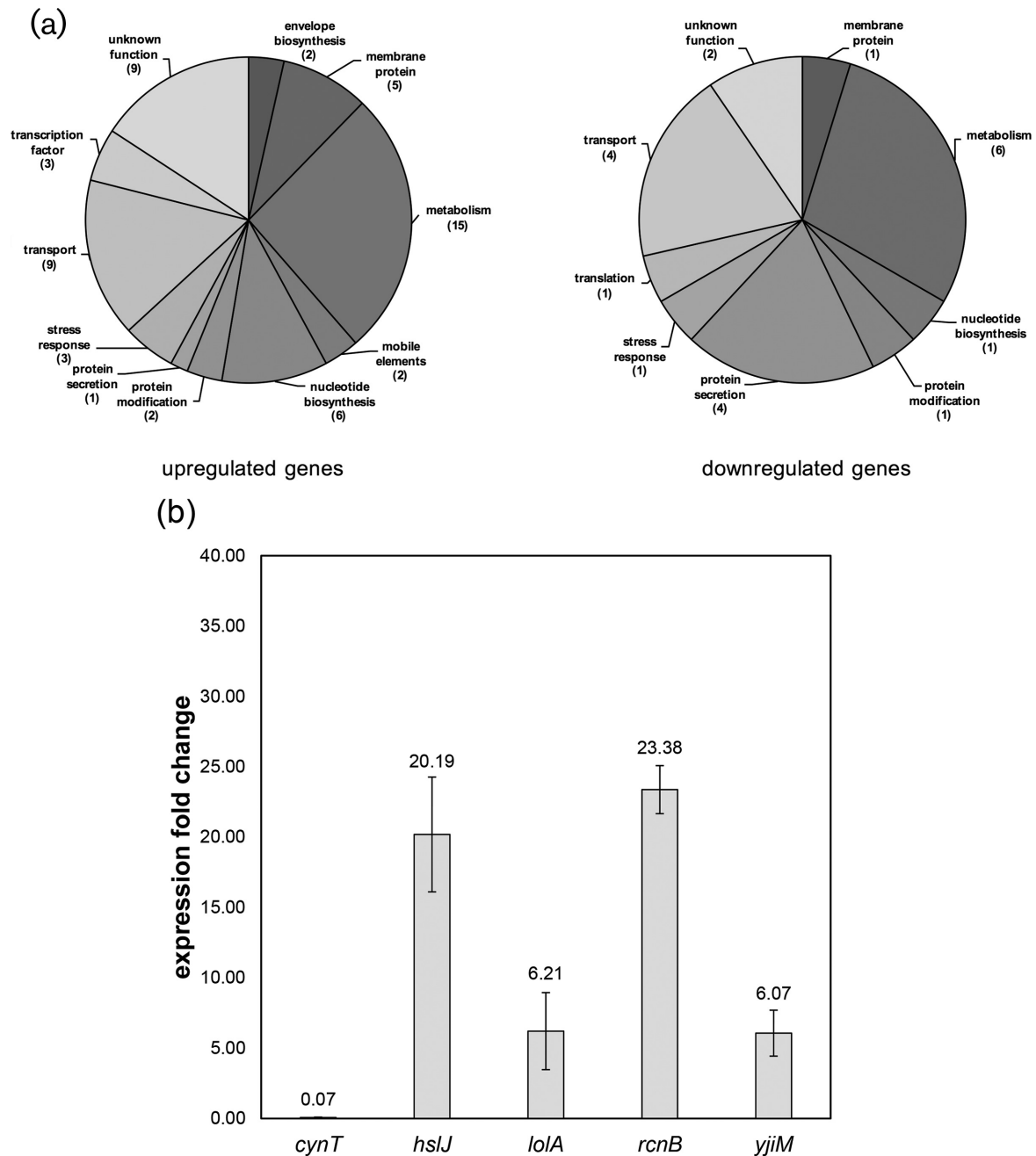


Fig. 7. Differential gene expression is induced in the *lnt* mutant. (a) Transcriptional profile of *lnt* mutants shows gene upregulation (57 genes) and downregulation (21 genes). Pie charts summarize the data found in Table S3, using a differential expression threshold of five-fold, $P \leq 0.01$. (b) qPCR confirmation of a subset of genes with altered expression when *lnt* function is disrupted. Changes in expression of the indicated genes in *A. baylyi* are represented as fold change in the *lnt* mutant as compared to wild-type, using the 16S rRNA gene as the housekeeping control gene. Numbers above the bars are the average values of biological triplicates, with error bars indicating standard deviation from the mean.

responses that are activated to compensate for defects in lipoprotein maturation.

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Conflicts of interest

The authors declare that there are no conflicts of interest.

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