Extra-Chromosomal DNA Sequencing Reveals Episomal Prophages Capable of Impacting Virulence Factor Expression in Staphylococcus aureus

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Staphylococcus aureus is a major human pathogen with well-characterized bacteriophage contributions to its virulence potential. Recently, we identified plasmidial and episomal prophages in S. aureus strains using an extra-chromosomal DNA (exDNA) isolation and sequencing approach, uncovering the plasmidial phage φBU01, which was found to encode important virulence determinants. Here, we expanded our extra-chromosomal sequencing of S. aureus, selecting 15 diverse clinical isolates with known chromosomal sequences for exDNA isolation and next-generation sequencing. We uncovered the presence of additional episomal prophages in 5 of 15 samples, but did not identify any plasmidial prophages. exDNA isolation was found to enrich for circular prophage elements, and qPCR characterization of the strains revealed that such prophage enrichment is detectable only in exDNA samples and would likely be missed in whole-genome DNA preparations (e.g., detection of episomal prophages did not correlate with higher prophage excision rates nor higher excised prophage copy numbers in qPCR experiments using whole-genome DNA). In S. aureus MSSA476, we found that enrichment and excision of the prophage φSa4ms into the cytoplasm was temporal and that episomal prophage localization did not appear to be a precursor to lytic cycle replication, suggesting φSa4ms excision into the cytoplasm may be part of a novel lysogenic switch. For example, we show that φSa4ms excision alters the promoter and transcription of htrA2, encoding a stress-response serine protease, and that alternative promotion of htrA2 confers increased heat-stress survival in S. aureus COL. Overall, exDNA isolation and focused sequencing may offer a more complete genomic picture for bacterial pathogens, offering insights into important chromosomal dynamics likely missed with whole-genome DNA-based approaches.

Keywords: bacteriophages, extra-chromosomal DNA, episomal, plasmidial, active lysogeny, virulence factor
INTRODUCTION

Staphylococcus aureus is a Gram-positive human pathogen with an arsenal of virulence factors allowing its successful colonization and infection of hosts (Thammavongsa et al., 2015; Thomer et al., 2016). Prophages and their encoded factors, in particular, play a significant role in the virulence potential and adaptability of the pathogen by means of lysogenic (positive) conversion, or the modulation of bacterial phenotypes by phage-encoded genes. S. aureus Newman, for example, displays severely reduced virulence in a mouse model when cured of its four prophages (Bae et al., 2006). Some prophages in particular have increased roles in the pathogenicity of strains, as they can positively convert cells with additional virulence determinants. The β-hemolysin (hlb)-converting phages of S. aureus carry genes for the well characterized toxins staphylokinase (SAK) and enterotoxin A (SEA) (Coleman et al., 1989), as well as the more recently discovered chemotaxis inhibitory protein of S. aureus (CHIPS) and staphylococcal complement inhibitor (SCIN), which allow bacterial evasion of the innate immune system (van Wamel et al., 2006). Other phages carry toxins such as the pore-forming Panton-Valentine leukotoxin (PVL) or exfoliative toxin A (ETA), the causative agent of staphylococcal scalded skin syndrome (SSSS) (Kaneko et al., 1998; Yamaguchi et al., 2000). Lysogenic conversion can also result from the disruption of host-encoded virulence factors through prophage genome integration (negative conversion). The hlb-converting phages integrate within and disrupt the β-hemolysin encoding gene of S. aureus (negatively converting cells for β-hemolysis), and similarly, lipase (geh)-converting phages, such as φL54a, negatively convert cells for lipase activity by integration within the lipase (geh) gene (Lee and Iandolo, 1986). The disruption of these genes has been shown to affect the colonization and virulence potential of S. aureus (Hu et al., 2012; Katayama et al., 2013), and while hlb-converting phages indeed encode alternative virulence determinants, geh-converting phages do not encode known virulence factors, with the benefits of their carriage unclear.

While positive and negative phage-conversion are well studied events, recent reports have also focused on the excision/integration dynamics and atypical localizations of temperate phages in S. aureus (Goerke et al., 2004, 2006; Utter et al., 2014). In S. aureus isolates from cystic fibrosis and bacteremic patients, genomic alterations driven by temperate phages were found integrated at atypical chromosomal loci, resulting in Hlb+/sak+ phenotypes (cells with an intact hlb gene, but positively converted by prophages encoding sak). Rarely, in some strains, phases appeared to undergo duplication and dual integration, generating populations with a Hlb−/sak2 phenotype (cells with one prophage integrated in hlb and a duplicated prophage integrated in another chromosomal location, with both phases encoding sak) (Goerke et al., 2006). However, such phage mobilization occurred to a significantly lesser degree in nasal isolates from healthy individuals (colonizing but not infectious isolates), indicating selective pressure for Hlb-producing strains in the transition to invasive infection, with atypical prophage integration as a mechanism to allow for dual Hlb and SAK production. Previous work in our laboratory uncovered the plasmidial prophage φBU01, with a DNA sequence containing high homology to known hlb-converting phages. φBU01 did not appear to integrate within the S. aureus chromosome (Utter et al., 2014), suggesting that in addition to atypical chromosomal integration, maintenance of prophages in the extra-chromosomal compartment can result in an Hlb+/sak+ phenotype (Deutsch et al., 2016).

Excision/integration dynamics of temperate phage (or phage-like elements) can also have important roles in other bacterial pathogens. Streptococcus pyogenes strain SF370 contains the episomal phage-like chromosomal island, SpyCIM1 (formerly termed φ370.4; Beres and Musser, 2007), which integrates within the cell’s DNA mismatch repair operon, disrupting transcription of the mutS-mutL genes and consequently increasing the mutation rate of the cell approximately 200-fold (Scott et al., 2008; Hendrickson et al., 2015). The phage-like element was found to be excised at low cellular densities allowing faithful genome replication, but would integrate and increase the cell’s mutation rate at higher cell densities (Scott et al., 2008)—conditions where mutation might be beneficial (i.e., low nutrient availability). SpyCIM1’s temporal dynamics suggest that S. pyogenes is using the phage-like element for its own benefit as a molecular switch at the DNA-level. Other species (Listeria monocytogenes, Bacillus subtilis) have been found to employ similar strategies with their phages, allowing control of activities such as phagosomal escape (Rabinovich et al., 2012) and sporulation (Kunkel et al., 1990; Takemaru et al., 1995), respectively. Feiner et al. (2015) reviewed these and other similar temperate phase-bacteria interactions, where prophage excision/integration generates genomic switches with significant impacts on the host, terming it “active lysogeny.”

Previously, we screened clinical isolates of S. aureus for the presence of rare, cytoplasmically localized prophages using extra-chromosomal DNA (exDNA) enrichment and next-generation sequencing (NGS). In this earlier report (Utter et al., 2014), we identified and sequenced the plasmidial prophage φBU01 from the vancomycin-intermediate S. aureus (VISA) NRS19, and found that the phage encoded multiple virulence determinants; we also uncovered an episomal prophage from VISA NRS26 in the same study. Enrichment and screening of the cytoplasmic compartment revealed that extra-chromosomally localized prophages were fairly widespread in S. aureus, and that such “hidden” elements may alter the virulence potential of a strain. For our current study, we expanded our screen for extra-chromosomal prophages in S. aureus, employing similar NGS methods to screen-by-sequencing an additional 15 diverse, clinical isolates. Unlike our previous work, however, the strains selected for this study had previously sequenced and assembled chromosomes, allowing the classification of prophages uncovered by our method as either episomal (found both integrated and extra-chromosomal in a population) or plasmidial (solely extra-chromosomal). Here, we found extra-chromosomal prophages
present in 5 of 15 strains, but surprisingly, these strains contained only episomal and not plasmidial prophages. Furthermore, we demonstrate that the episomal phages uncovered by our approach are circular DNA elements, and reveal that enrichment and detection of such phages would not occur using conventional whole-genome DNA preparations; PCR-based measurements of excised phage copy numbers and excision rates from whole-genome DNA preparations did not correlate with enrichment in our screening. In addition, we find cytoplasmic localization of φSa4ms, from the S. aureus strain MSSA476, to be growth-phase dependent and that the prophage does not appear to replicate after excision, suggesting it exists in a state of “active lysogeny.” Lastly, we show that excision of φSa4ms alters the promoter sequence and transcription of the stress-response serine protease-encoding htrA2, with promoter alterations affecting heat-stress survival in S. aureus COL. Thus, our extra-chromosomal enrichment and sequencing approach allows the detection of these “active” prophages and other episomal or plasmidial DNA elements, offering greater insights into the virulence potential and genome dynamics of an organism.

MATERIALS AND METHODS

Bacterial Strains and Growth Conditions
Strains used in this study are listed in Supplementary Table S1. S. aureus strains were established from overnight (O/N) cultures grown in Bacto brain heart infusion (BHI), and back-diluted 1:100 into 50 mL BHI without shaking at 37°C unless otherwise noted. For preparation of exDNA, strains were grown as described to an OD600 of 0.6–0.8. Cultures were centrifuged at 4000 rpm for 10 min at 4°C and used immediately or frozen O/N at −20°C. For qPCR studies, strains were back-diluted from O/N cultures and grown to specified optical densities. For PhtrA2-GFP reporter studies, O/N cultures were back-diluted 1:100 and grown at 200 rpm at 37°C to an OD600 = 0.2. Concentrations for antibiotics used are as follows, for Escherichia coli: ampicillin (100 µg/mL); for S. aureus: erythromycin (5 µg/mL), spectINomycin (50 µg/mL). For heat shock studies, plates were incubated at 44°C.

Whole-Genome and exDNA Isolation and Manipulation
Whole-genome DNA isolations (gDNA) were performed using the QIAGEN DNeasy Blood and Tissue Kit, which included an added manufacturer-detailed pretreatment step for Gram-positive bacteria. Enzymatic lysis buffer was composed of lysozyme (100 µg/mL) in 1× phosphate-buffered saline (PBS). exDNA isolation was carried out as previously described (Utter et al., 2014). exDNA samples were visualized on 0.7% agarose 0.5× TAE gels stained with SYBR Safe DNA Gel Stain. Electrophoresis was carried out at 50 V for 1 h in 0.5× TAE, and visualized with UV transillumination. Prior to DNA sequencing, exDNA samples were concentrated as needed using Microcon DNA Fast Flow (EMD Millipore) centrifugal filters, following manufacturer’s directions.

DNA Sequencing of Extra-Chromosomal S. aureus Samples
DNA sample concentrations were measured using the Thermo Fisher Scientific Qubit Fluorometer High Sensitivity DNA kit. Size and quality of DNA was measured using the Agilent Technologies Bioanalyzer High Sensitivity DNA assay. DNA libraries were constructed as per manufacturer’s instructions starting with 1 ng of DNA per sample, using the Illumina Nextera XT DNA Library Preparation Kit and the Nextera XT Index Kit. The completed DNA libraries were quality checked using the Agilent Technologies High Sensitivity DNA assay. Shotgun sequencing of the DNA libraries was performed using the Illumina MiSeq Reagent kit V2 (500 cycle) on the Illumina MiSeq sequencer. Prior to sequencing, libraries were normalized and pooled together to make a multiplexed pooled DNA library at 2 nM concentration. The 2 × 250-paired end sequencing run generated FASTQ files to allow for off-instrument analysis. Raw extra-chromosomal DNA sequencing reads associated with this study are accessible via NCBI BioProject PRJNA475753.

Bioinformatic Sequence Analysis of exDNA Samples
Bioinformatic analysis was performed using CLC Genomics Workbench software unless otherwise described. Extra-chromosomal reads were mapped to respective chromosomal sequences (Supplementary Table S1), and unmapped reads were saved as a separate file, and de novo assembled for identification of possible plasmidial elements. Read mappings were visually examined for regions of increased read depth, corresponding to DNA element enrichment in exDNA samples. Read mappings were then subjected to coverage analysis using CLC Genomics Workbench software to identify regions (minimum length 500 bp) with significantly higher (P < 0.05) coverage distributed throughout integrated prophage genome locations. Prophage and other mobile DNA element regions with high coverage were noted for each sequenced sample.

qPCR Analysis of S. aureus Strains
For qPCR experiments, strains were grown as described for exDNA sequenced samples, or to other desired optical densities. Cultures were divided, with one portion of the culture subjected to the QIAGEN DNeasy Blood and Tissue Kit for gDNA isolation, and the rest undergoing exDNA isolation. Whole-genome DNA samples were used to determine excision rates and phage or plasmid (pSAS1) copy numbers per cell of the bacterial population, while exDNA samples were used only for phage and plasmid copy number measurements. Primer pairs and probe sequences for each target are listed in Supplementary Table S2. All primers and probes were designed and purchased from Integrated DNA Technologies (Coralville, IA, United States). Amplification was carried out using the TaqMan Gene Expression Master Mix (Thermo Fisher Scientific) and the Life Technologies QuantStudio 12K-Flex Instrument following manufacturer's
cycling protocol. A standard curve for each primer-probe set was set up for each experimental run, with amplification efficiencies and linear regression analyzed using QuantStudio Software. All primer-probe sets had efficiencies 90–110% and $R^2 > 0.98$. Excision rates, excised prophage copy numbers, and plasmid copy numbers were calculated by normalizing the targets ($attB$, $attP$, pSAS1, respectively) to $gyrA$. All targets were measured in biological triplicate and technical duplicates. Graphical and statistical analysis was performed using GraphPad Prism, with significance testing done using two-tailed Student's $t$-tests. Bar graphs are presented as mean ± standard error of the mean (SEM).

**Linear DNase and Restriction Endonuclease Treatment of exDNA Samples and End-Point PCR of DNA Targets**

Prior to linear DNase and/or treatment with restriction endonuclease, exDNA samples were treated with PreCR Repair Mix (NEB) to repair nicked DNA. PreCR-treated samples were then either (1) treated with the φS4ms-specific restriction endonucleases PshA1 and PspXI (NEB) following manufacturer’s protocol, (2) treated with Plasmid-Safe-ATP-Dependent DNase (Epicentre), (3) treated with PshA1 and PspXI, then the Plasmid-Safe-ATP-Dependent DNase, or (4) left solely PreCR-treated. Plasmid-Safe-treated samples were treated over the course of 16 h at 37°C, where 2 µL ATP solution and 2 µL DNase were added at 2 and 4 h time points. Samples were then heated to inactivate enzymes prior to PCR. End-point PCR was carried out using target-specific primers (Supplementary Table S2) and the KAPA2G Robust Hotstart Polymerase (KAPA Biosystems) with the following cycling protocol: (1) initial denaturation 95°C, 3 min; (2) denaturation 95°C, 15 s; (3) annealing 60°C, 15 s; (4) extension 72°C, 15s; (5) repeat steps 2–4 thirty three times. Samples were normalized prior to loading individual PCRs to account for dilutions from original sample concentration due to enzyme treatments. PCR products were visualized using 1% Safe-ATP-Dependent DNase, or (4) left solely PreCR-treated.

**Construction and Testing of PhtrA$_2$-GFP Reporter System and htrA$_2$-Complemented Knockouts**

For construction of a PhtrA$_2$-GFP reporter system, 250 bp upstream of the htrA$_2$ gene plus 30 bp of htrA$_2$ were amplified from a MSSA476 gDNA template using Phusion High-Fidelity DNA Polymerase with two primer sets (1) Pint_upstm/P_dwnstms or (2) Pex_upstm/P_dwnstms (Supplementary Table S2) to amplify two versions of the htrA$_2$ promoter and 5' gene region (Pint or Pex). PCR products were gel purified, cut with restriction enzymes KpnI and BamHI (NEB), and further purified. pCN56 vector (Charpentier et al., 2004) was purified using the QIAprep Spin Miniprep Kit (QIAGEN), digested with KpnI and BamHI, dephosphorylated with Antarctic Phosphatase (NEB), and gel purified before ligation using T4 ligase (NEB) with the purified PCR products. Next, 2 µL of the ligation mixture was introduced into One Shot TOP10 Chemically Competent E. coli (Thermo Fisher Scientific) and the resulting colonies screened by PCR and Sanger sequencing (GENEWIZ; South Plainfield, NJ, United States), using primers Pint_upstm_seq/P_dwnstms_seq or Pex_upstm_seq/P_dwnstms_seq for the desired vector insert (Supplementary Table S2). E. coli cultures harboring pCN56 with Pint or Pex inserts were grown and miniprepped, and the purified vectors electroporated into electrocompetent S. aureus RN4220 (Kreiswirth et al., 1983) using a Bio-Rad Gene Pulser with the following settings: 2.5 kV, 25 µF, 100 Ω. Empty pCN56 vector was also introduced into RN4220. Vectors from RN4220 colonies were then transduced, using φNM4y4 (Heler et al., 2015), into S. aureus COL htrA$_3$ (Rigoulay et al., 2005) following an established protocol (Olson, 2016). Colonies were screened via PCR and inserts Sanger sequenced to ensure the correct DNA sequence.

To examine GFP fluorescence in each strain, O/N cultures of constructs were back-diluted 1:100 in 15 mL BHI supplemented with 5 µg/mL erythromycin and 50 µg/mL spectinomycin and grown at 37°C, 200 rpm to an OD$_{600}$ = 0.2. Cultures were then grown for 3 h at 37°C or 44°C with shaking. After 3 h growth, 1 mL of culture was removed, an OD$_{600}$ value measured, and 200 µL of culture was pipetted into a quartz 96-well plate to measure RFUs on a Molecular Devices SpectraMax M5 instrument (485 nm excitation, 515 nm emission). Fluorescence of pCN56 (empty vector) cultures were subtracted as background, and OD$_{600}$ normalized RFUs for each sample were measured. Comparison of construct-GFP fluorescence was calculated as a ratio of Pex/Pint. Graphical and statistical analysis was performed using Prism GraphPad. Bar graphs are presented as mean ± SEM, with significance indicated above each bar. Significance testing was performed using two-tailed ratio paired $t$-tests. The experiment was performed at least in triplicate.

For construction of htrA$_2$-complement vectors, the htrA$_2$ gene and 250 bp upstream of the gene were amplified using the primer set (1) Pint_upstm/full_htrA2_dwnstms or (2) Pex_upstm/full_htrA2_dwnstms (Supplementary Table S2) with Q5 DNA Polymerase (NEB) to generate two DNA fragments containing the full length htrA$_2$ sequence with different 250 bp promoter sequences. PCR products were treated as described above, before being ligated into the pCN35 vector (Charpentier et al., 2004). The ligated vectors and pCN35 empty vector were processed as above, introduced into electrocompetent E. coli DC10B, shuttled into RN4220, transduced into S. aureus COL htrA$_2$, then screened and sequenced as previously described using primers full_htrA2_seq1, full_htrA2_seq2, full_htrA3_seq3, full_htrA2_dwnstms, Pint_upstm_seq, and Pex_upstm_seq (Supplementary Table S2).

To test viability and heat-stress survival of complemented and mutant (empty vector-containing) S. aureus COL htrA$_2$ strains, constructs were spotted in a log-dilution series onto BHI–spectinomycin–erythromycin agar plates and incubated at 37 and 44°C for 24 h following a protocol for determining heat sensitivity from (Rigoulay et al., 2005). Plates were examined...
Deutsch et al. Episomal Prophages in *Staphylococcus aureus*

visually for survival and photographed the following day using a Cell Biosciences Alphalager HP instrument using Alphalager HP software. The experiment was performed in triplicate.

### RESULTS

**Extra-Chromosomal DNA Isolation and Sequencing Reveals the Presence of Episomal Prophage Elements Within *S. aureus* Clinical Isolates**

For this study, we expanded our previously developed exDNA enrichment and sequencing approach to screen *S. aureus* clinical isolates for prophage elements present in the extra-chromosomal compartment of the cell (Utter et al., 2014). We selected 15 clinical isolates sourced from different worldwide geographic regions, containing a diverse array of antibiotic resistances and virulence factors. In addition, these strains had fully sequenced chromosomes, facilitating the classification of any detected prophages as episomal or plasmidial. We enriched and prepared exDNA of these strains as described in Section “Materials and Methods,” and performed paired-end sequencing using the Illumina MiSeq sequencer. exDNA samples were analyzed by first mapping sequencing reads to their corresponding chromosomal sequences, followed by de novo assembly of unmapped reads. Visual examination of chromosomal read mappings revealed areas of increased sequencing coverage, especially over some prophage regions (Figures 1A, B). Coverage analysis transformed visual read-mapping data into histograms and highlighted ≥500 bp regions of the chromosome where read depth was significantly increased (*P* < 0.05). Such regions of higher coverage within integrated prophage genomes indicated the presence of episomal elements (and/or potentially active prophages) enriched in sequencing due to specifically targeting

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**FIGURE 1 |** Read mapping and coverage analysis of MSSA476 extra-chromosomal DNA sequencing. (A, B) Read mappings of sequencing reads for MSSA476 exDNA. φSa4ms (B) has greater read coverage compared to φSa3ms (A) in exDNA sequencing. Selected portions of the read mapping surrounding prophage integration sites for φSa3ms (A) and φSa4ms (B) are shown. Chromosomal positions are labeled at top, and consensus regions are shown below for MSSA476 chromosome sequence and MSSA476 exDNA reads. Locations of φSa3ms (A) and φSa4ms (B) integrated prophage genomes are shown. Extra-chromosomal reads are shown below for each. (C) Coverage analysis histogram from MSSA476 extra-chromosomal DNA sequencing read mapping. Portions of coverage analysis histogram surrounding prophage regions shown for clarity. Chromosomal positions labeled at top of image, and approximate positions of φSa3ms and φSa4ms genomes shown below. Histogram of higher read-coverage regions on the MSSA476 chromosome contains 11 significantly higher coverage regions within the φSa4ms prophage genome location, indicating prophage enrichment in exDNA sequencing. Read-mapping across φSa3ms prophage genome location does not contain any regions of significantly higher coverage, indicating no prophage enrichment by exDNA isolation and sequencing.
and isolating DNA from the extra-chromosomal compartment of the cell (Figure 1C). Read mappings and a coverage analysis histogram are shown for strain MSSA476 as a representative example (Figure 1).

For each strain, we documented whether any prophage elements were enriched in sequencing due to exDNA isolation and classified the strains as: (1) containing enriched prophage elements, (2) not containing enriched prophage elements, or (3) containing unclear prophage element enrichment, when partial but not complete prophage regions had increased coverage detected (Table 1). Surprisingly, de novo assembly of unmapped reads did not reveal any new prophage elements, suggesting no plasmidial prophages were present in the sequenced strains, and that all prophages uncovered in the extra-chromosomal compartment by our approach were episomal elements. Importantly, episomal prophages were identified in one-third (5 of 15) of the staphylococcal strains analyzed. These included: (1) MSSA476, harboring φSa4ms (Sa4-like integrase), (2) NRS143 containing a geh-converting prophage (Sa6-like integrase), (3) BK2529 harboring a prophage with a Sa7-like integrase, as well as two strains each with three episomal prophages. One of these strains, HPV107, contained episomal prophages with Sa2- and Sa3-like integrases, and in addition, one prophage with an unclear integrase type, however, its sequence was homologous to the integrase of a prophage from S. aureus SA268 (Qu et al., 2014). The other strain, NRS22, contained episomal prophages with Sa2-, Sa5-, and Sa7-like integrase sequences. Interestingly, when integrated into the chromosome, the Sa2-like prophage of HPV107 disrupts a 6-phospho-β-galactosidase encoding gene, which to our knowledge is a novel integration site for a S. aureus prophage. The other episomal phages we uncovered had chromosomal integration sites that have been described previously (Bae et al., 2006; Goerke et al., 2006, 2009). In addition to these episomal prophage elements, we also uncovered an enriched integrative and conjugative element (ICE6013-like) typically located within a predicted membrane protein encoding locus in NRS271, however, its existence as an extra-chromosomal element has been previously reported (Table 1) (Smyth and Robinson, 2009; Sansevere et al., 2017). Incidentally, known plasmids, such as pSAS1 in MSSA476, were de novo assembled by using unmapped reads, and while they are not the focus of this current study, they do point to the sensitivity of this exDNA isolation and sequencing method.

### Table 1: Detection of enriched prophages from extra-chromosomal DNA sequencing.

| Strain | Resistances | #Prophage regions | Notes |
|--------|-------------|-------------------|-------|
| **Enriched prophage elements detected** | | | |
| BK2529 | MRSA | 3 | ● One enriched prophage: Sa7-like integrase, intergenic between rpmF and isdB |
| HPV107 | MRSA | 3 | Three enriched prophages: |
| | | | ● Sa2-like integrase, intragenic in RL05_04730 (β-phospho-β-galactosidase) |
| | | | ● Sa3-like integrase, hlb-converting phage (φHPV107.1) |
| | | | ● Unclear integrase type, intergenic between RL05_02285 (tRNA-Ser) and RL05_01940 (enterotoxin) |
| MSSA476 | MSSA | 2 | ● φSa4ms enriched, integrates 30 bp upstream of htrA2 |
| | | | ● φSa3ms not enriched, hlb-converting |
| NRS22 | VISA/MRSA | 4 | Three enriched prophages: |
| | | | ● Sa2-like integrase, intragenic in RK87_02385 (hypothetical protein) |
| | | | ● Sa5-like integrase, intragenic in RK87_04825 (radical SAM) |
| | | | ● Sa7-like integrase, intergenic between rpmF and isdB |
| NRS143 | MSSA | 2 | One enriched prophage: Sa6-like integrase, geh-converting |
| **Unclear enriched prophage elements** | | | |
| NRS2 | VISA/MRSA | 2 | ● Unclear enrichment of Sa7-like integrase prophage, intergenic between rpmF and isdB |
| NRS153 | MSSA | 3 | ● Unclear enrichment of Sa1-like integrase prophage, intergenic between sauB and RK79_06750 (transposon-encoded integrase) |
| NRS387 | MRSA | 1 | ● Unclear enrichment of Sa3-like integrase prophage, hlb-converting |
| **No enriched prophage elements detected** | | | |
| BAA-42 | MRSA | 3 | ● Enriched ICE (ICE6013-like) within RK77_00405 (membrane protein) |
| E2125 | MRSA | 3 |
| HDE288 | MRSA | 3 |
| NRS127 | MRSA | 5 |
| NRS156 | MSSA | 1 |
| NRS158 | MSSA | 2 |
| NRS271 | MRSA | 2 |

Relevant resistances (MSSA, MRSA, VISA, VRSA) and total number of predicted chromosomal prophages for each strain are listed.
qPCR Characterization of MSSA476 Validates Extra-Chromosomal Sequencing Data, and Also Reveals Episomal φSa4ms Is Detectable Only in exDNA Samples

The conditions of our screening-by-sequencing approach did not reveal any non-integrating, plasmidial prophages. However, a number of the clinical strains did contain episomal prophage elements uncovered from the cytoplasmic compartment by specific exDNA isolation (Table 1). Thus, we were curious what insights prophage enrichment, as uncovered by exDNA sequencing, revealed about an individual strain. To accomplish this, we chose to focus on the well-characterized S. aureus strain MSSA476 (Sumby and Waldor, 2003; Holden et al., 2004) and took a qPCR approach to determine the excision rates and copy numbers of its prophages. MSSA476 contains two prophages, φSa3ms and φSa4ms. φSa3ms is an hlb-converting phage, while φSa4ms integrates 30 bp upstream of the serine protease-encoding htrA2. In our sequencing, φSa4ms had significantly increased read depth in read-mappings (i.e., was enriched by exDNA isolation), while φSa3ms did not (Figure 1). Therefore, MSSA476 was ideal to understand the characteristics of both enriched and non-enriched prophages in our screening.

We grew MSSA476 as described in Section “Materials and Methods,” and isolated exDNA and whole-genome DNA (gDNA) from cultures at OD_{600} = 0.35, 0.7, and 1.0, as well as from O/N cultures, and profiled the excision rates and excised copy numbers of the strain’s prophages by qPCR. To determine the excised copy numbers of φSa3ms and φSa4ms, we targeted prophage attachment sites (attP), designing primers to yield a PCR product only when phages were not integrated in the chromosome, and normalized to gene copies of chromosomal DNA gyrase (gyrA) for both exDNA and gDNA samples. Excision rates were determined by targeting phage-free bacterial attachment sites on the chromosome (attB) and normalized to gyrA in gDNA samples. qPCR data for excised copy numbers and excision rates of φSa3ms and φSa4ms at each growth point are listed in Table 2.

We first wanted to validate our extra-chromosomal sequencing results, which suggested φSa4ms may exist as a stable episome in a portion of the MSSA476 population. Sequencing data should have a direct correlation to excised prophage copy numbers, predicting that φSa4ms copy number would be greater than φSa3ms in exDNA samples from OD_{600} = 0.7, a growth point representative of when samples were processed for initial exDNA isolation and sequencing. Indeed, the copy number of excised φSa4ms (copies attP/copy gyrA) was significantly higher than φSa3ms (P = 0.018), supporting our exDNA sequencing data (Table 2 and Figure 2A). Unexpectedly, however, examination of excised prophage copy numbers from gDNA samples revealed that φSa3ms excised copy number was significantly higher than that of φSa4ms (P = 0.0046) (Table 2 and Figure 2B). In addition, excision rate data correlated with the gDNA excised prophage copy numbers; φSa3ms had a significantly higher excision rate (copies attB/copy gyrA) than φSa4ms (P < 0.0001) (Table 2 and Figure 2C). While the gDNA copy number and excision rate data appeared to be in agreement (i.e., higher excision rates correlate with higher excised prophage copy numbers) they were in contrast to copy number results obtained from qPCR of exDNA samples and from the initial sequencing screening. We therefore performed additional experiments to uncover the basis of these apparent conflicting results before analyzing the remaining qPCR data from OD_{600} = 0.35, 1.0, and O/N cultures.

### Extra-Chromosomal DNA Samples Are Enriched for Circular Prophages

To uncover why φSa4ms was enriched by exDNA isolation but not in gDNA preparations, we first compared φSa3ms and φSa4ms prophage genomes for %GC content, which could alter DNA capture efficiency. However, %GC content did not appear to play a role in enrichment differences as φSa3ms and φSa4ms contain almost equivalent %GC content at 33.2 and 33.3%, respectively. Part of the extra-chromosomal enrichment protocol involves alkaline-lysis followed by centrifugation. We hypothesized that this step likely enriches for circular forms of prophage that would remain soluble after alkaline-lysis and not pellet during centrifugation (i.e., would be captured in the exDNA preparation). Linear phage DNA concatemers (contiguous copies of phage genome arising from spontaneously induced lytic cycle phage) would likely pellet with chromosomal DNA and other cellular debris and be removed from exDNA

| Prophage | OD_{600} | Excised prophage copy number (copies attP/copy gyrA) | Excision rate (copies attB/copy gyrA) |
|----------|----------|-----------------------------------------------------|-------------------------------------|
|          |          | exDNA | gDNA |                          |          |
| φSa3ms   | 0.35     | 0.3801 ± 0.050 | 0.109 ± 0.009 | 4.64 × 10^{-4} ± 4.52 × 10^{-5} |          |
|          | 0.7      | 0.3377 ± 0.064 | 0.096 ± 0.005 | 4.80 × 10^{-4} ± 1.10 × 10^{-5} |          |
|          | 1.0      | 0.2653 ± 0.021 | 0.1370 ± 0.013 | 5.05 × 10^{-4} ± 2.73 × 10^{-5} |          |
|          | O/N      | 0.2563 ± 0.014 | 0.1230 ± 0.001 | 9.58 × 10^{-4} ± 2.14 × 10^{-5} |          |
| φSa4ms   | 0.35     | 2.220 ± 0.110 | 0.0840 ± 0.006 | 1.51 × 10^{-4} ± 2.47 × 10^{-5} |          |
|          | 0.7      | 1.257 ± 0.229 | 0.0647 ± 0.001 | 9.91 × 10^{-5} ± 3.13 × 10^{-6} |          |
|          | 1.0      | 0.4270 ± 0.030 | 0.0767 ± 0.008 | 8.44 × 10^{-5} ± 1.14 × 10^{-5} |          |
|          | O/N      | 0.2073 ± 0.019 | 0.0700 ± 0.008 | 8.75 × 10^{-5} ± 8.15 × 10^{-6} |          |

* Asterisk ** indicates culture density representative of samples from extra-chromosomal DNA sequencing. O/N, overnight culture. Values are displayed as mean ± SEM.
Figure 2 | qPCR characterization of MSSA476 excised prophage copy numbers and excision rates at OD600 = 0.7. (A) Prophage copy numbers (copies attP/copy gyrA) from MSSA476 extra-chromosomally enriched DNA (exDNA) samples. Prophage copy number is significantly higher for φSa4ms than φSa3ms, in accordance with sequencing data. (B) Prophage copy numbers from MSSA476 whole-genome DNA (gDNA) samples. Copy number of φSa3ms is significantly higher than φSa4ms. (C) Excision rates of φSa3ms and φSa4ms (copies attB/copy gyrA). Excision rate of φSa3ms is significantly higher than φSa4ms. Bars indicate mean ± SEM. *P < 0.05; **P < 0.01; and ***P < 0.001.

The same linear concatemers, however, would not be excluded in whole-genome preparations, potentially explaining the differences seen in qPCRs of gDNA versus exDNA samples as both single-copy circular and multi-copy concatemer forms of phage DNA contain identical attP sites that can be amplified by PCR.

To test this hypothesis, we designed and performed a selective-depletion experiment of excised φSa4ms in exDNA samples. Specifically, we examined if φSa4ms could be selectively removed from samples only by the combination of φSa4ms-specific restriction endonucleases (linearizing circular prophage forms) followed by linear DNA digestion, and not solely by the exonuclease (linear DNase) treatment. φSa4ms depletion only by sequential endo- and exonuclease treatment would indicate it exists as a closed circular DNA element. Thus, exDNA samples were treated as described in Section “Materials and Methods” and end-point PCR measurements were performed targeting gapdh, the attP sites of φSa3ms and φSa4ms, and the naturally occurring plasmid pSAS1. An agarose gel containing all PCRs is shown in Figure 3. Gapdh (a marker for linear DNA digestion) was depleted after linear DNase treatment, but remained in high abundance after treatment with the φSa4ms-specific restriction endonucleases PshAI and PspXI. φSa3ms attP target was present in each condition tested, indicating that it can also exist in the circular form in extrachromosomally enriched DNA samples, however, in lower abundance. The attP target of φSa4ms is present and in greater abundance than that of φSa3ms in untreated, restriction endonuclease-only and linear DNase-only treated samples. Treatment with φSa4ms-specific endonuclease followed by linear DNase treatment results in the complete loss of φSa4ms target, verifying that φSa4ms is indeed abundant relative to φSa3ms in extra-chromosomally enriched samples, and that this enrichment is most likely due to circular forms of the prophage element (Figure 3). In further support, pSAS1 (a circular plasmid in MSSA476 which does not contain restriction sites for PshAI or PspXI) was not affected by any sample treatment.

**Extra-Chromosomal Localization of φSa4ms Circular Prophage Is a Temporal and Likely Rare Event**

We found episomal, circular copies of φSa4ms in the MSSA476 population. However, this enrichment did not appear to correspond to total excised phage copy numbers in PCRs of gDNA. To better understand the episomal nature of φSa4ms, we next examined our full qPCR data set (Table 2), analyzing changes in φSa3ms and φSa4ms circular copy numbers (via attP in exDNA samples), total excised phage copy numbers (via attP in gDNA samples), and excision rates (via attB in gDNA samples) over the MSSA476 growth cycle. Examination in this manner allowed insights into MSSA476’s prophage dynamics, and in addition, whether φSa4ms may be acting as an active lysogen (i.e., excising into the extra-chromosomal compartment without increases in overall excised copy number). Bar graph visualization of qPCR data from Table 2 is presented in Figure 4. Analysis of excised prophage copy numbers from exDNA samples revealed that for φSa4ms, extra-chromosomal
localization of circular prophage is a temporal event, with the highest prevalence of cytoplasmic prophage carriage at early-logarithmic phase. Excised φSa4ms copy number in exDNA samples decreased significantly from OD$_{600}$ = 0.35 to 1.0 ($P < 0.0001$), while for φSa3ms, there was no significant change in its excised circular copy number over this interval ($P = 0.0982$) (Table 2 and Figure 4A). However, while circular φSa4ms significantly decreased in copy number during logarithmic growth, its total excised phage copy number in the MSSA476 population (as measured in gDNA samples, representing circular and concatemer forms) did not change, nor did that of φSa3ms. φSa3ms and φSa4ms did not have significant differences in total excised phage copy numbers from OD$_{600}$ = 0.35 to 1.0 ($P = 0.118$ and $P = 0.527$, respectively) (Table 2 and Figure 4B), indicating that the excision and presence of circular φSa4ms does not appear to be a precursor to potential phage linear replication. Interestingly, the excision rate of φSa4ms appeared to correlate with its decrease in circular copy number, as it significantly decreased from OD$_{600}$ = 0.35 to 1.0 ($P = 0.036$) (Table 2 and Figure 4C). The φSa3ms excision rate did not significantly change during this interval ($P = 0.2403$), however, it did increase approximately twofold from OD$_{600}$ = 0.35 to the O/N time point (Table 2 and Figure 4C).

While these data indicated that φSa4ms was temporally localized in the cytoplasm as a circular element as well as potentially a lysogen in MSSA476 cells, the prevalence of its stable excision was unclear. We employed qPCR targeting pSAS1 (the naturally occurring circular plasmid in MSSA476) as a reference to understand the level of enrichment imparted by our exDNA preparation procedure. We found that pSAS1 existed on average at one to two copies per cell (copies pSAS1 target/copy gyrA) in gDNA preparations of MSSA476 at OD$_{600}$ = 0.35 and 0.7, but was enriched ~1500-fold to an average of 2322 copies pSAS1/copy gyrA in exDNA samples (data not shown). That φSa4ms copy number in our exDNA qPCR reaches, at a maximum, 2.22 copies attP/copy gyrA (Table 2), suggests that the φSa4ms excision we uncover is a rare event. The percentage of cells with stably excised φSa4ms, as well as its circular copy number per cell, is presently unclear.

### Promoter Alteration by φSa4ms Excision/Integration Affects htrA2 Transcription and Heat-Stress Survival in S. aureus

Since qPCR data indicated φSa4ms could exist as an active prophage, a stable subpopulation of MSSA476 cells may therefore exist with an altered chromosomal sequence, as φSa4ms integrates 30 bp upstream of the stress response serine protease-encoding htrA2. Sumby and Waldor (2003) previously noted the possibility of altered htrA2 transcription by φSa4ms excision/integration, and our data suggested that indeed a subpopulation of MSSA476 cells would harbor htrA2 under an alternative promoter (Figure 5A). We therefore developed a PfhrA2-gfpmut2 reporter system to examine whether GFP fluorescence would be altered when the GFP-encoding gene gfpmut2 was under the control of the φSa4ms-integrated (Pint) or φSa4ms-excised (Pex) promoters. Experiments were performed in a S. aureus COL htrA2 deletion knockout, a well-characterized strain where htrA2 was shown to impact heat-stress survival (Rigoulay et al., 2005). Comparisons of GFP fluorescence at 37 and 44°C showed that in COL htrA2, fluorescence was greater when gfpmut2 was promoted by Pex rather than by Pint, indicating that Pex is a stronger
Deutsch et al. Episomal Prophages in Staphylococcus aureus

**FIGURE 4** | qPCR characterization of MSSA476 prophage excision rates and excised copy numbers from logarithmic and overnight cultures. (A) Excised prophage copy numbers (copies attP/copy gyrA) from MSSA476 extra-chromosomally enriched DNA (exDNA) samples at indicated optical densities. φSa3ms does not show a significant difference in excised copy number from OD$_{600}$ = 0.35 to 1.0, whereas φSa4ms excised copy number significantly decreases during this interval. (B) Prophage copy numbers from MSSA476 whole-genome DNA (gDNA) samples at indicated optical densities. Neither φSa3ms nor φSa4ms have significantly different excised prophage copy numbers comparing OD$_{600}$ = 0.35 and 1.0 samples. (C) Excision rates (copies attB/copy gyrA) of φSa3ms and φSa4ms at indicated optical densities. φSa3ms does not show a significant difference in excision rate comparing OD$_{600}$ = 0.35 and 1.0 samples, whereas a significant decrease exists for φSa4ms. Bars indicate mean ± SEM. *$P < 0.05$ and ***$P < 0.001$. 

**FIGURE 5** | φSa4ms excision/integration alters the promoter and transcription of htrA2. (A) Diagram illustrating φSa4ms integrated and excised states in MSSA476 and the unique DNA regions of the Pint and Pex promoters. Top: When φSa4ms is integrated in the chromosome, htrA2 is promoted by Pint. Bottom: φSa4ms excision joins the Pex region immediately upstream to htrA2, and the gene is promoted by Pex. DNA sequence unique to the Pex region is carried on circular φSa4ms when the phage is excised. (B) Ratio of GFP fluorescence from Pex- versus Pint-promoted gfpmut2 constructs in COL htrA2 deletion strain at 37°C and 44°C. Bars indicate mean ± SEM. Significance as tested by ratio paired t-test indicated above each bar. (C) Dilution series of COL htrA2 deletion strain complemented with htrA2 promoted by Pint (int) or Pex (ex). The symbol “–” indicates COL htrA2 deletion strain containing empty pCN35 plasmid. Constructs are spotted in a log-dilution series and grown at 37°C (left) or 44°C (right) for 24 h. Dilution factor indicated on left of plates; ND, not diluted. COL constructs display equal survival at 37°C, but COL Pex displays 2-log survival above COL Pint at 44°C. Unexpectedly, uncomplemented COL htrA2 shows 1-log survival above COL Pint. COL Pint shows few colonies at 44°C in the undiluted culture spot. *$P < 0.05$ and ***$P < 0.001$.

Due to its reported role in promoting heat-tolerance in COL, we examined if promoter alteration could affect survival of a htrA2-complemented S. aureus COL htrA2 deletion mutant.
Complemented strains, containing the full-length htrA gene under the control of either the Pint or Pex promoter on the pCN35 plasmid or with empty pCN35 (uncomplemented) were grown in a dilution series at 37 or 44°C on agar plates. COL constructs at 37°C did not display any survival differences among the three strains, however, at 44°C, COL Pex displayed approximately 2-log increased survival over COL Pint (Figure 5C, top). Surprisingly, COL Pint survived 1-log worse than uncomplemented COL htrA2, indicating that htrA2-mediated heat-stress survival may depend upon specific promoter sequences and not solely levels of transcription or promoter strength. Environmental conditions may select for cells with one promoter versus another, and φSa4ms's excision/integration dynamics could provide MSSA476 with a potential switching mechanism to create these advantageous subpopulations.

**Episomal Prophages Uncovered in Additional S. aureus Clinical Isolates**

In this study, we uncovered and characterized the episomal dynamics of the φSa4ms prophage in *S. aureus* MSSA476. Our exDNA sequencing screening, however, also uncovered the presence of other likely episomal prophages in additional clinical isolates (Table 1). While we did not characterize the prophage dynamics of these other strains in this study, we were curious if detection of their episomal prophages was also only apparent through exDNA isolation and sequencing, and if their presence would be missed by gDNA-focused approaches. To test this, we performed an inter-strain comparison, examining by qPCR the excision rates and excised prophage copy numbers of hlb-converting phages in BAA-42 (containing φBAA-42.1) and HPV107 (containing φHPV107.1). Sequencing data revealed that only φHPV107.1 and not φBAA-42.1 was significantly enriched by exDNA isolation (Table 1), and qPCR copy number examination of exDNA samples confirmed this result. φHPV107.1 had a higher copy number as compared to φBAA-42.1 (P = 0.0052) in exDNA samples (Table 3 and Figure 6A, left). However, as observed previously with the MSSA476 prophages, copy number data from gDNA samples did not correlate with exDNA sample data. φHPV107.1 and φBAA-42.1 did not have significant copy number differences in gDNA samples (P = 0.3205) (Table 3 and Figure 6A, right), verifying that episomal prophage enrichment is only apparent due to exDNA-focused isolation. Interestingly, when we examined excision rates of the two prophages, we found that in this comparison, φHPV107.1 did indeed have the highest excision rate. The excision rate of φHPV107.1 was significantly higher than φBAA-42.1 (P = 0.0009) (Table 3 and Figure 6B). Overall, this result suggests that like φSa4ms, φHPV107.1 may exist as a circular, episomal DNA element in a higher proportion of the HPV107 population than that of φBAA-42.1 in its respective population. The lytic induction of these prophages in other cells of their populations (and the generation of phage genome concatemers), however, may mask uncovering their rarer episomal natures in qPCRs of gDNA samples.

**DISCUSSION**

This study explored the prevalence of extra-chromosomal prophages in *S. aureus* as an extension of previous work uncovering the plasmidial phage φBU01 in *S. aureus* NRS19 (Utter et al., 2014). We isolated and sequenced the exDNA of
15 clinically relevant *S. aureus* strains with known chromosomal sequences, but unlike our previous study, we did not find any prophages existing as solely plasmidial elements. We did uncover, however, several episomal prophages appearing to exist as circular DNA elements and potential active lysogens that were enriched by our exDNA isolation procedure (Table 1). It therefore seems that the existence of plasmidial prophages in *S. aureus* is uncommon and φBU01 may represent one of these rare prophages. Episomal prophages, on the other hand, appear to be fairly widespread among *S. aureus* isolates and were identified in 33% of strains (5 of 15) examined in this study.

**Strains and Phages Possess Different Mobilization Capacities With Unclear Mechanisms**

We could distinguish strains that carried episomal prophages in the extra-chromosomal compartment from those with no prophages detected, but our study did not reveal why some strains have such a phage-mobilization capacity while others do not. Goerke et al. (2006) demonstrated a similar observation, finding that phages φs80b and φs84b (hlb-converting phages) were integrated within the *hlb* gene in *S. aureus* s64c, but were found to alternate between integrated and extra-chromosomal carriage in strain 8325-4. Both s64c and 8325-4 are phage-cured strains, suggesting that a host-factor (or factors) likely determines phage localization and mobilization capacity within the cell. In a separate manner, phage induction capacity into the lytic cycle has also been observed to depend upon host background, with φSA2mw induced by mitomycin C from *S. aureus* strains MW2 and Newman, but not 8325-4, RN6390, or ISP479c (Wirtz et al., 2009). Our data indicate that the prophages we describe here are localized and harbored within the extra-chromosomal compartment in a manner distinct from lytic excision and replication (Utter et al., 2014; Deutsch et al., 2016). However, it is possible similar host factors may govern both events. The hlb-converting phages we characterized from strains BAA-42 and HPV107 contained high sequence homology over their integration/excision modules; however, they displayed different levels of enrichment in extra-chromosomal sequencing and qPCR characterization, with HPV107 carrying a greater proportion of its hlb-converting phage extra-chromosomally. Host differences similar to those affecting φs80b and φs84b localization may account for the differences in hlb-converting phage localization among our strains, although such factors have yet to be uncovered.

Within individual strains, we also noted differences in phage localization. In MSSA476, φSA4ms—but not φSA3ms—was enriched in sequencing, with the localization of circular, episomal φSA4ms confirmed by qPCR and selective-depletion experiments. φSA3ms and φSA4ms share the same host background, suggesting that a phage factor may be responsible for localization differences, but it is currently unclear which factor(s) could be responsible. Prophage excision requires recombination mediated by integrase; stochastic expression differences of a phage's integrase or excision-related genes may control localization into the extra-chromosomal compartment. If excision occurs without inducing conditions, then phages could be localized in the cytoplasm in a manner consistent with the lysogenic cycle. Feiner et al. (2015) reviewed such phage activity in a range of bacterial species, particularly those whose excision/integration dynamics can act as a molecular switch for the cell, terming the process “active lysogeny.”

**Episomal Phages Are Only Detected in exDNA and Not gDNA Samples**

qPCR characterization of prophage excision rates and excised copy numbers revealed, surprisingly, that episomal prophages are detected only in exDNA samples and that their presence is masked in gDNA preparations. In MSSA476, for example, the overall excised phage copy number as measured from qPCRs of gDNA is relatively low, with φSA3ms and φSA4ms at 0.095 and 0.0647 copies *attP/copy gyrA*, respectively. This would correspond to very low increases in read depth over the φSA3ms and φSA4ms prophage chromosomal locations if only MSSA476 gDNA was sequenced and read-mapped to a MSSA476 reference chromosome (likely 9.5 and 6.5% maximum increases, respectively). Using gDNA, coverage analysis tools would likely not find regions of significantly higher coverage spanning either prophage location, so while φSA3ms copy number was greater than φSA4ms in gDNA samples, neither phage would be found as enriched by this approach, and the episomal nature of φSA4ms would be overlooked. φSA4ms excised phage copy number in exDNA samples, however, is much greater (1.26 copies *attP/copy gyrA* at OD$_{600}$ ~0.7), owing to concurrent enrichment of circular φSA4ms and removal of chromosomal DNA in the exDNA isolation procedure, increasing the *attP/gyrA* ratio. This increase is sufficient for significantly higher read depth across the prophage location when mapping exDNA sequencing reads to the chromosome. The excised copy number of φSA3ms in exDNA samples, while elevated from that seen in gDNA samples (0.34 versus 0.095 copies *attP/copy gyrA*, respectively), was not high enough to have significantly higher read depth or coverage over the φSA3ms genome location in MSSA476 read-mappings. Thus, our exDNA isolation and sequencing approach allows the distinction of episomal elements, even when such elements would be masked in qPCR and sequencing of DNA prepared by other methods.

Additionally, we showed that our exDNA isolation procedure enriches circular DNA elements, allowing the identification of episomal and plasmidial elements, and that whole-genome focused approaches do not impart such selectivity in sample preparations. Higher excision rates and higher total excised phage copy numbers (from gDNA samples) did not correlate with episomal prophage enrichment in our NGS screening. We hypothesize that the reason φSA4ms copy number was lower than that of φSA3ms in gDNA samples, despite the presence of φSA4ms episomes, is likely due to the presence of more linear, lytic-cycle phage genome concatemers of φSA3ms in the MSSA476 population. The greater number of linear phage concatemers in gDNA samples (but not in exDNA) likely masks the low percentage of circular phage elements, preventing their discovery in sequencing or qPCR in the absence...
of exDNA enrichment. This is supported by the fact that the excised prophage copy numbers of φSa3ms and φSa4ms are still very low in exDNA samples despite the potential ∼1500-fold enrichment of circular DNA (i.e., pSAS1 plasmid) by our protocol. Further, this suggests that for MSSA476, both φSa3ms and φSa4ms excised copy numbers are primarily composed of linear elements, which are likely from a small subset of cells whose phages are undergoing spontaneous lytic cycle replication. While both phages in MSSA476 are capable of such lytic events, only φSa4ms showed enrichment for circular prophage elements in its cytoplasm. This result might indicate that for a number of cells in the MSSA476 population, circular φSa4ms prophage is not participating in the lytic cycle, but is perhaps in a state akin to active lysogeny (Feiner et al., 2015). Other individual cells in the MSSA476 population, however, appear to contain φSa3ms or φSa4ms undergoing lytic cycle replication. A model for how these bacteriophages might exist in the MSSA476 population during early-logarithmic phase is presented in Figure 7.

ExDNA Isolation and Sequencing Reveals Potential and Previously Identified Active Prophages

Having better understood the nature of enrichment imparted by exDNA isolation (i.e., enrichment of circular DNA elements), we were able to analyze our full MSSA476 qPCR dataset (Table 2) and uncover the strain’s prophage dynamics. Surprisingly, results revealed that excision of circular φSa4ms into the extra-chromosomal compartment was temporal, with the highest levels of circular prophage recorded in early-logarithmic growth, and that its circular copy number rapidly decreased with increasing cell-density. Furthermore, these decreases occurred without any concurrent increase in overall φSa4ms copy number (as measured by gDNA PCRs), indicating that φSa4ms circular elements were not precursors to production of linear phage genome concatemers and that the prophage was likely not participating in the lytic cycle. In addition, these data show that phage enrichment would likely go undetected in MSSA476 if samples were prepared at OD600 ≥ 1.0. Therefore, it may be beneficial to purify exDNA at multiple time points in future studies, elucidating the entire mobilization dynamics of a bacterial genome over its growth cycle.

The time-course qPCR data also showed that the excision rate and circular copy number of φSa4ms decreased from early to later-log phase, suggesting potential control of this event by the bacterial cell, with some factors either promoting excision at early-logarithmic growth, or perhaps increased re-integration at later growth points. In addition, whether φSa4ms or related circular prophage elements undergo any replication or remain as single copy in this excised state is unclear. Goerke et al. (2006) reported Hlb−/sak2 S. aureus strains with multiple hlb-converting phages integrated in the chromosome. Replication

![FIGURE 7](https://example.com/figure7.png)

**FIGURE 7** | Qualitative model of MSSA476 early-logarithmic culture. At early logarithmic phase, the MSSA476 population contains cells with phages in lytic, lysogenic, and active lysogenic cycles. The vast majority of cells are stably lysogenized (tan) with integrated φSa3ms (blue) and φSa4ms (orange). Subpopulations of cells contain excised prophage elements that appear to be in an active lysogenic cycle (green). Of these cells, those with φSa4ms excised outnumber those with φSa3ms excised. Subpopulations of cells with phage induced into the lytic cycle are pictured in gray. Here, φSa3ms-induced cells outnumber φSa4ms-induced cells. Subpopulations are shown at relative percentages for illustrative purposes, rather than quantitative levels suggested by qPCR and other data.
of excised prophage could be the first step in generating such strains. Our qPCR approach examined bacterial populations as a whole, but cannot address these specific questions; further experiments on the single-cell level are necessary to uncover such exact excision/integration (and potentially replication) dynamics.

The distinction between phages in the lytic versus lysogenic cycle is important when considering the roles of prophages as molecular switches (i.e., active lysogens) or as mechanisms to generate diversity in bacterial populations. Excision by phage induction into the lytic cycle likely leads to cell death, whereas excision in active lysogeny generates stable diversity in the population that can be passed to daughter cells. We found that φSa4ms excision/integration alters the promoter sequence and consequent transcription of htrA2 in a GFP-reporter system and affects survival of S. aureus COL to heat-stress. While qPCR revealed that only a minor proportion of MSSA476 likely carries excised φSa4ms and alternatively promoted htrA2, heat-stress or other environmental factors could select for and expand this subpopulation. Our work suggests the behavior of φSa4ms could generate a small subset of cells in the MSSA476 population better equipped to respond to and survive environmental stress. Studying strains by our extra-chromosomal sequencing approach may be useful for researchers seeking active prophages or similar-acting DNA elements in other species of interest. As further proof of this concept, we performed a “retroactive” exDNA enrichment and sequencing of S. pyogenes SF370, which indeed showed the expected enrichment of sequences for the previously reported episomal phage-like element SpyCIM1, which acts as a switch in the strain’s mismatch repair operon (Scott et al., 2008; Deutsch et al., 2016). Indeed, a similar exDNA enrichment and sequencing approach was used to identify mobile retrotransposons in Arabidopsis and rice (Lanciano et al., 2017).

In addition to φSa4ms, the intergenic prophages we uncovered as enriched in our S. aureus screening included one prophage in HPV107 (unclear integrase type, intergenic between tRNA-Ser and enterotoxin), and two prophages in NRS22 and BK2529 [both with Sa7-like integrases, and intergenic between rpmF (50S ribosomal protein L32) and isdB (iron-regulated surface determinant)]. The potential effects of their extra-chromosomal carriage on hosts are unclear, but phage integration in these loci has been reported previously (Bae et al., 2006; Goerke et al., 2006). The typically intragenic prophages we found enriched in our screening included a geh-converting prophage in S. aureus NRS143, two prophages in HPV107 (an hlb-converting phage, and a 6-phospho-β-galactosidase-integrating phage), and two prophages in NRS22 (one prophage integrated within a hypothetical protein-encoding gene, and one prophage disrupting a radical SAM-encoding gene). In addition, we uncovered an enriched ICE6013-like element, whose localization has been previously reported as both integrated and extra-chromosomal (Smyth and Robinson, 2009; Sansevere et al., 2017). The hlb- and geh-converting phages could potentially act as switches to control the expression of these virulence factors and future work should investigate this possibility. Likewise, the effects of the other intragenic prophages are unclear, however, their enrichment suggests that they could also act as switches to control underlying gene expression. Disruption of genes by the intragenic prophages in NRS22 has been reported previously in other S. aureus strains (Bae et al., 2006). However, to our knowledge, disruption of a 6-phospho-β-galactosidase encoding gene (HPV107) has not been reported previously and represents a novel prophage integration site. The role of the ICE is also unclear, but its potential as a switch is intriguing.

Overall, exDNA isolation and sequencing revealed the epimodal nature of specific typically integrated staphylococcal prophages. Previously, the movement of hlb-converting phages to atypical chromosomal loci was shown to occur in isolates from cystic fibrosis and bacteremic patients with phage mobilization thought to be important for invasive infection. As also noted by (Goerke et al., 2006), the localization of typically integrated prophages into the cytoplasm (without lytic induction) could precede generation of such unique CF and bacteremic S. aureus isolates. Our extra-chromosomal enrichment and sequencing approach may allow characterization of strains for such a mobilization capacity or potential. In addition, uncovering the epimodal nature of other prophages (e.g., geh-converting, 6-phospho-β-galactosidase-disrupting prophages) may help better direct future research efforts and allow discovery of novel processes and events important to S. aureus infection.

CONCLUSION

This study screened the exDNA of 15 clinical S. aureus isolates, uncovering the prevalence of episomal prophages with potential roles in virulence factor expression and regulation. MSSA476 qPCR characterization revealed that prophage φSa4ms can exist as an episomal circular DNA element that does not appear to be a precursor to lytic cycle replication. Importantly, we found that episomal prophages are only detectable in extra-chromosomally enriched DNA samples, and that their presence would have been missed in sequencing or qPCRs of whole-genome DNA samples. Lastly, we showed that φSa4ms may act as a potential phage-molecular switch, as its excision/integration alters the promoter sequence of htrA2, changing its transcription levels and affecting heat-stress survival in S. aureus COL. We believe φSa4ms behavior represents that of an active lysogen, and that the other episomal prophages we identify may exhibit similar activity. Episomal prophages promote the generation of stable, diverse subpopulations with the potential to impact bacterial infection. exDNA enrichment and sequencing should allow the increased discovery of such elements in S. aureus and other bacterial pathogens, and help researchers better assess those genes under control by active lysogeny.

AUTHOR CONTRIBUTIONS

DD, BU, and VF conceived and designed the experiments. DD, BU, KV, HS, and LT performed the experiments. DD, BU, KV, HS, LT, and VF contributed
materials and analyzed the data. DD and VF wrote the paper.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fmicb.2018.01406/full#supplementary-material

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**Conflict of Interest Statement:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest. The opinions, interpretations, conclusions, and recommendations contained herein are those of the authors and are not necessarily endorsed by the FDA.

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