

# Novel Chimerical Endolysins with Broad Antimicrobial Activity Against Methicillin-Resistant *Staphylococcus aureus*

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Due to their bacterial lytic action, bacteriophage endolysins have recently gained great attention as a potential alternative to antibiotics in the combat of Gram-positive pathogenic bacteria, particularly those displaying multidrug resistance. However, large-scale production and purification of endolysins is frequently impaired due to their low solubility. In addition, a large number of endolysins appear to exhibit reduced lytic efficacy when compared with their action during phage infection. Here, we took advantage of the high solubility of two recently characterized enterococcal endolysins to construct chimeras targeting *Staphylococcus aureus*. The putative cell wall binding domain of these endolysins was substituted by that of a staphylococcal endolysin that showed poor solubility. Under appropriate conditions the resulting chimeras presented the high solubility of the parental enterococcal endolysins. In addition, they proved to be broadly active against a collection of the most relevant methicillin-resistant *S. aureus* epidemic clones and against other Gram-positive pathogens. Thus, fusion of endolysin domains of heterologous origin seems to be a suitable approach to design new potent endolysins with changed and/or extended lytic spectrum that are amenable to large-scale production.

## Introduction

AFTER DECADES OF ALMOST exclusive use of antibiotics in the treatment of infectious diseases caused by pathogenic bacteria, the emergence of multiresistant bacterial strains combined with a slowdown in the discovery of new classes of antibiotics is currently viewed as a major public health problem.<sup>23,78,79</sup> Among the different bacterial pathogens that have acquired antibiotic resistance *Staphylococcus aureus* is still one of the most serious threats worldwide, being able to develop resistance to virtually all classes of antibiotics.<sup>19</sup> Methicillin-resistant *S. aureus* (MRSA) has been a leading cause of nosocomial infections for some decades and more recently has also emerged as a community-associated pathogen.<sup>5,12,30,40</sup> *S. aureus* is responsible for a wide number of infectious diseases ranging from skin and soft tissues to fatal bloodstream infections,

endocarditis, meningitis, and bovine mastitis in dairy herds.<sup>34,45</sup> For all these reasons, there is an urgent need to develop new antibacterials that ensure elimination of multidrug-resistant pathogenic bacteria, particularly MRSA strains.

Among the different approaches to obtain new antibacterials, one that has recently caught great attention is the exploitation of bacteriophage endolysins.<sup>27,29,48,59</sup> Endolysins are bacterial cell wall-hydrolyzing enzymes that promote host cell lysis in the end of the lytic cycle of double-stranded DNA phages, thus allowing efficient release of the viral progeny to the extracellular medium.<sup>71</sup> Despite the fact that endolysins have been evolutionarily designed to operate from the inside of infected cells, when exogenously applied as purified recombinant proteins they have been shown to promote lysis of different Gram-positive pathogenic bacteria, including *S. aureus*.<sup>47,55,58,80</sup>

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Most endolysins have a modular organization with a conserved N-terminal catalytic domain (CD) and a more diverse C-terminal cell wall binding domain (CWBD).<sup>28,32</sup> However, several *S. aureus* phages produce endolysins with two CDs in their N-terminus, such as those from phages K,  $\phi$ 11 and  $\phi$ MR11. These lytic enzymes present a CHAP (cysteine, histidine-dependent amidohydrolases/peptidases) domain followed by an Amidase-2 domain (*N*-acetylmuramoyl-L-alanine amidase), where the CHAP seems to be the most effective in inducing lysis.<sup>54,58,63</sup> Another feature commonly reported for *S. aureus* phage endolysins is their poor solubility when overexpressed in *Escherichia coli*, frequently leading to low yields of soluble protein.<sup>9,31,51,58</sup>

We have experienced this insolubility problem when attempting to purify a broad-host-range anti-staphylococcal endolysin available in our lab, Lys87.<sup>11</sup> Here, we describe a new strategy to overcome this problem that consisted of the construction of new chimerical endolysins composed of the Lys87 CWBD fused to the highly soluble CDs of two *Enterococcus faecalis* phage endolysins, Lys168 and Lys170.<sup>62</sup> By employing this strategy, complemented with optimized expression conditions, we obtained two highly soluble lytic enzymes that are broadly active against a large cohort of *S. aureus* strains, including MRSA, and against other staphylococcal, streptococcal, and enterococcal species. To our knowledge, this corresponds to the first report of chimeras comprising domains of *S. aureus* and *E. faecalis* phage endolysins.

## Materials and Methods

### Bacteria, culture media, and growth conditions

The *E. coli* cloning strain XL1-Blue MRF' and its derivatives were grown at 37°C with aeration in Luria-Bertani (LB) medium.<sup>68</sup> The *E. coli* expression strain CG61<sup>72</sup> and its derivatives were grown in LB in the same conditions, except that incubation temperature was 28°C before induction of protein production and 16°C afterward. When appropriate, LB medium was supplemented with kanamycin (40 µg/ml) and/or ampicillin (100 µg/ml) for plasmid selection. Lytic action of chimerical endolysins was assayed in 200 bacterial clinical isolates (Table 1, Supplementary Tables S1 and S4; Supplementary Data are available online at [www.liebertonline.com/mdr](http://www.liebertonline.com/mdr)). Table 1 corresponds to a panel of 30 MRSA and 13 Methicillin-sensitive *S. aureus* (MSSA) typed strains. Table S1 lists 100 *S. aureus* isolates from Technophage's collection, 42 MRSA, 29 MSSA, and 29 with unknown methicillin susceptibility, which were obtained from different Portuguese community and hospital settings between 2005 and 2008. Recombinant lytic enzymes were also tested in clinical isolates of other species, namely against *Staphylococcus epidermidis* (n=10), *Staphylococcus haemolyticus* (n=9), *Staphylococcus saprophyticus* (n=7), *Enterococcus faecalis* (n=12), *Enterococcus faecium* (n=10), and *Streptococcus pyogenes* (n=9) (Technophage collection, Table S4). The growth media for these bacteria were purchased from Biokar Diagnostics, Beauvais, France. *Staphylococcus* and *Enterococcus* species were cultured in Brain-Heart Infusion and *Streptococcus pyogenes* in Todd Hewitt broth supplemented with 2% of yeast extract (THY). Bacteria were grown at 37°C, with aeration, except for *Enterococcus* that was grown without agitation. When necessary, 1.5% or 0.7% agar was added

to these culture media to obtain solid or soft-agar plates, respectively. *E. faecalis* and *S. aureus* phages were propagated and purified by standard methods,<sup>15,42</sup> either in soft-agar media or liquid broth supplemented with CaCl<sub>2</sub> and MgCl<sub>2</sub> (5 mM each).

### Identification and bioinformatics analysis of phage endolysins

Genomic DNA from *E. faecalis* and *S. aureus* phages was extracted from CsCl-purified lysates<sup>77</sup> and their complete nucleotide sequence determined (service purchased to Macrogen). DNA homology searches were carried out with BLASTN,<sup>81</sup> using the NCBI non-redundant nucleotide sequences database. Recognition of phage genome putative genes was performed by integrating the results obtained with GeneMark.hmm and MetaGeneAnnotator web software.<sup>10,56</sup> Endolysins Lys168 and Lys170 from *E. faecalis* phages F168/08 and F170/08, respectively, were characterized previously.<sup>62</sup> Identification of *S. aureus* phage F87s/06 endolysin gene was based on BLASTP homology searches<sup>4</sup> against the NCBI non-redundant protein sequences database, using the deduced gene product, and on prediction of protein functional domains using NCBI's CDD<sup>52</sup> and Pfam (<http://pfam.janelia.org/search>). Assignment of putative linkers connecting protein functional domains was performed with SVM<sup>25</sup> and the SVM-joint output. Multiple protein sequence alignments were performed with ClustalW2.<sup>45</sup>

### Construction and cloning of lys168-87 and lys170-87 chimerical genes

The coding sequences of the N-terminal regions of the enterococcal phage endolysins Lys168 and Lys170, which included CHAP and Amidase-2 CDs, respectively<sup>62</sup> (Fig. 1A and Supplementary Fig. S1A), were polymerase chain reaction (PCR) amplified from the corresponding phage DNA using a high-fidelity *Pfu* DNA Polymerase (Fermentas Molecular Biology Tools, Thermo Scientific). The sequence encoding the C-terminal region of the staphylococcal phage endolysin Lys87 (Fig. 1A and Supplementary Fig. S1A), and that harbored the putative CWBD, was similarly amplified by PCR in a separate reaction. The forward primers used for CD amplification carried an *Nco*I restriction site in their 5' end, whereas the reverse primer employed in CWBD amplification contained an *Xma*I site. The reverse and forward primers used in CD and CWBD amplification, respectively, carried a 26-bp complementary segment in their 5' end. This allowed fusing the coding sequence of each enterococcal endolysin CD to that of the CWBD of Lys87 by the technique of overlap-extension by polymerase chain reaction (OE-PCR),<sup>37</sup> using the CD forward and CWBD reverse primers and a mixture of the initial PCR products as template. The resulting PCR products, corresponding to the chimerical genes *lys168-87* and *lys170-87* (Fig. 1B and Supplementary Fig. S1B), were purified using the High Pure PCR Product Purification Kit (Roche Applied Science), double-digested with *Nco*I and *Xma*I, and ligated to the equally digested pI-VEX2.3d expression vector (Roche Applied Science). This vector is designed to drive the expression of cloned genes under the control of the phage T7  $\phi$ 10 promoter and to allow the production of the corresponding proteins C-terminally fused to a hexahistidine tag. The *E. coli* strain XL1-Blue MRF'

was transformed with these ligations as described previously.<sup>14</sup> Transformants were selected in the presence of 100 µg/ml ampicillin and screened for the presence of the desired recombinant plasmids by PCR, using insert and vector complementary primers. Plasmid DNA from positive clones was extracted (Pure Link Quick Plasmid Miniprep Kit; Invitrogen) and the correct DNA structure confirmed by endonuclease restriction and DNA sequencing (Macrogen). The constructs pSF168-87 and pSF170-87 are pIVEX2.3d derivatives carrying *lys168-87* and *lys170-87*, respectively.

#### *Production and purification of the chimerical endolysins Lys168-87 and Lys170-87*

*E. coli* strain CG61, which overproduces phage T7 RNA polymerase upon temperate upshift<sup>72</sup> was transformed with plasmids pSF168-87 and pSF170-87 and transformants selected at 28°C in presence of 100 µg/ml ampicillin and 40 µg/ml kanamycin. Production of soluble and active Lys168-87 and Lys170-87 by CG61 transformants was first confirmed by growing them over a dense lawn of autoclavated staphylococcal cells, incorporated in soft-agar LB medium, and checked for the presence of lysis halos around *E. coli* colonies (Supplementary Fig. S2).

Selected clones of each chimera were grown at 28°C until an optical density at 600 nm (OD<sub>600</sub>) of 0.8–1.0, after which protein production was induced by changing cultures to a shaking water bath set to 42°C. After 30 min, cultures were transferred to an incubator at 16°C and agitated for an additional period of 14 hr. Cells from induced cultures were pelleted by centrifugation (8,000 g, 30 min, 4°C) and resuspended in 1/50 volume of lysis buffer (20 mM Hepes, 500 mM NaCl, 20 mM imidazole, 1% glycerol, and 1 mM DTT [pH 6.5]) supplemented with 1× Complete Mini EDTA-free Protease Inhibitor Cocktail (Roche Applied Science). Cells were kept on ice and disrupted by sonication (Vibra Cell MS2T, Sonic Materials) by performing about 10 bursts of 1 min (amplitude 50, pulse 3, 30–40 W) intercalated with pauses of 1 min. Insoluble material was sedimented by centrifugation (20,000 g, 30 min, 4°C). The supernatant corresponding to the total soluble extract was filtered through 0.22 µm and the chimerical enzymes purified by affinity chromatography using HisTrap™ HP columns (GE Healthcare) coupled to an AKTA-Prime system (GE Healthcare). The column and elution buffers had the same composition of the lysis buffer, except that the imidazole concentration in the elution buffer was 500 mM. Eluted fractions were analyzed by sodium dodecyl sulfate polyacrylamide gel electrophoresis and Coomassie blue staining.<sup>44</sup> Chimeras pure fractions were pooled, concentrated, and changed to an imidazole-free, phosphate-based endolysin buffer (50 mM phosphate-Na, 500 mM NaCl, 25% glycerol, and 1 mM DTT [pH 6.5]) using HiTrap™ Desalting columns (GE Healthcare). Protein concentrations were determined by the Bradford method (Bio-Rad Laboratories) using bovine serum albumin as standard. The enzymes were divided into small aliquots and kept at –20°C.

#### *Evaluation of the lytic action of chimerical endolysins against bacterial pathogens*

The ability of chimeras Lys168-87 and Lys170-87 to induce lysis of clinical strains from different bacterial species was evaluated by two different assays. When tested against a

large number of bacterial isolates, these were individually cultured as indicated above until an OD<sub>600</sub> of 0.8–1.0. Cells were recovered by centrifugation and resuspended in 1/100 volume of fresh medium. A 100 µl sample of these cellular suspensions was diluted in 10 ml of incorporation buffer (25 mM phosphate-Na and 250 mM NaCl [pH 6.5]) supplemented with 0.7% agar and poured in a Petri dish. Four quantities of the purified chimeras (10, 5, 1, and 0.2 µg, in 10 µl final volume) were spotted on each bacterial lawn and after overnight incubation at 37°C checked for the presence of lysis halos. These were evaluated and scored (– to + + +) according to their relative diameter and transparency.

Bacterial cell lysis was also studied in liquid medium. Selected strains were grown until an OD<sub>600</sub> of 0.3–0.4, centrifuged, and cells recovered in 1/2 volumes of a phosphate buffer (25 mM phosphate-Na and 250 mM NaCl [pH 6.5]). One-milliliter cell suspensions were challenged with 10 µg/ml of each chimera separately or with a mixture of both enzymes, each at 5 µg/ml. The OD<sub>600</sub> variation was followed over time. At the end of each assay the surviving colony forming units (CFU)/ml were determined. Negative controls were equally prepared except that endolysin buffer was added instead of endolysin.

## Results

### *Production of chimerical endolysins targeting S. aureus*

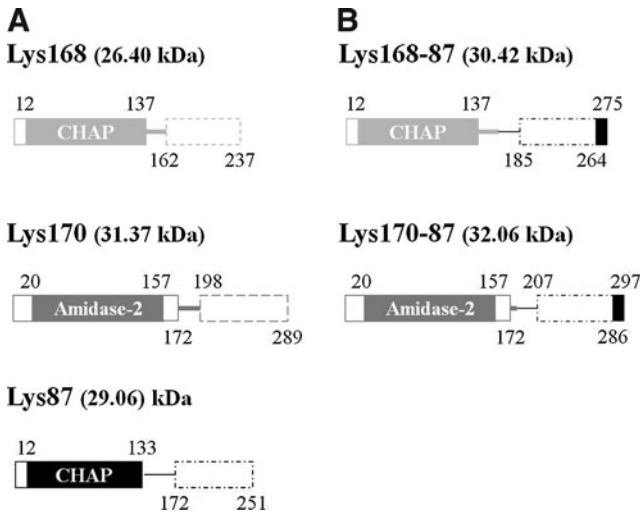
We have recently characterized three phage endolysins, two from *E. faecalis* phages F170/08 and F168/08 and another from *S. aureus* phage F87s/06.<sup>62,11</sup> *In silico* analysis of the endolysin from phage F170/08, that is, Lys170 identified a CD of the Amidase-2 family in its amino-terminal region (Fig. 1A and Supplementary Fig. S1A). This family includes zinc amidases that have N-acetylmuramoyl-L-alanine amidase activity.<sup>13</sup> The same analysis performed with endolysins Lys168 and Lys87 from phages F168/08 and F87s/06, respectively, revealed a CD of the CHAP family,<sup>7,46,64</sup> also located in the enzymes' N-terminal region (Fig. 1A and Supplementary Fig. S1A). Enzymes from this family have been shown to cleave different amide bonds in the peptidoglycan mesh, frequently displaying amidase or endopeptidase activity.

Lys170 and Lys168 could be easily produced in *E. coli* in its soluble form, whereas Lys87 showed a high propensity to precipitate as inclusion bodies in all tested conditions.<sup>11,62</sup> The enterococcal endolysins exhibited a narrow lytic spectrum with their activity being basically restricted to *E. faecalis* cells.<sup>62</sup> This feature might result from CWBD binding to a specific epitope present on the cell wall of *E. faecalis* cells and/or dependence of the CD on CWBD binding for activity.<sup>49</sup> Lys170 showed better lytic performance when both enterococcal endolysins were tested against a large number of *E. faecalis* clinical strains.<sup>62</sup>

Having this in mind we have envisaged that replacing the CWBD of Lys170 and Lys168 by that of Lys87 would result in chimerical endolysins (Fig 1B and Supplementary Fig. S1B) maintaining the high solubility properties of the parental enterococcal enzymes and the ability to recognize and act on *S. aureus* cells conferred by the parental Lys87 CWBD.

The coding sequences of the functional domains composing each chimerical endolysin were PCR-amplified, fused by





**FIG. 1.** Schematic representation of parental (**A**) and resulting chimerical endolysins (**B**). N-terminal catalytic domains (CD) are depicted as filled rectangles with indication of functional families (CHAP and Amidase-2). Dashed rectangles delimit the C-terminal region that must contain the putative CWBD. Linker regions connecting CD and CWBD are indicated by a line with the filling code and thickness of parental endolysins. The black rectangle at the end of each chimera represents the C-terminal hexahistidine tag used for affinity chromatography purification. The amino acidic coordinates of functional and linker domains and the total length of each endolysin are indicated above and below the schemes. CWBD, cell wall binding domain; CHAP, cysteine, histidine-dependent amidohydrolases/peptidases.

OE-PCR, and cloned in the *E. coli* expression vector pIVEX2.3d, which allowed production of the chimeras C-terminally fused to a hexahistidine tag (Fig. 1B, Supplementary Fig. S1B, and Materials and Methods). The ability of the chimerical enzymes Lys168-87 and Lys170-87 to lyse *S. aureus* cells was initially confirmed by growing the resulting protein-expressing *E. coli* clones on a dense lawn of autoclaved staphylococcal cells and checking for the presence

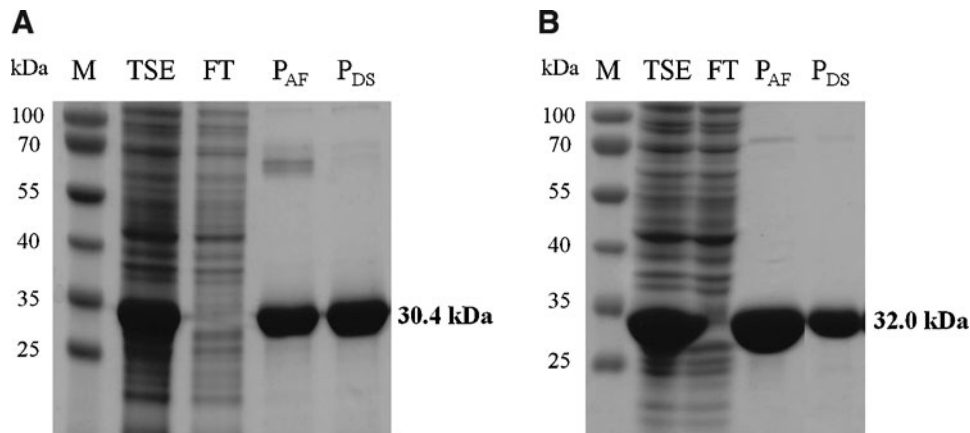
of lysis halos around the *E. coli* colonies (Supplementary Fig. S2).

By employing standard protein expression conditions (37°C, 3–4 hr production after culture induction) with selected clones we could obtain much higher soluble amounts of the chimerical endolysins than those we had previously obtained with Lys87. However, the quantity of soluble Lys168-87 and Lys170-87 did not match the levels we achieved with the parental enterococcal endolysins, as we anticipated (not shown). This could be overcome though by simply incubating the induced cultures at 16°C during 14 hr. In these conditions we achieved similar or higher soluble quantities of the chimeras when compared with the native enterococcal enzymes. The chimeras were subsequently purified from total soluble extracts by affinity chromatography using nickel columns. Pure fractions were pooled and subjected to a desalting step for protein exchange to an imidazole-free, sodium phosphate-based buffer (Fig. 2). Average yields were 20 µg and 30 µg of pure Lys168-87 and Lys170-87, respectively, per ml of induced culture.

#### *Lytic action of the chimerical endolysins against clinical S. aureus*

The lytic action of Lys168-87 and Lys170-87 against *S. aureus* was first assessed using a panel of 100 clinical isolates from Technophage's collection (Supplementary Table S1), which were isolated from different infection contexts and obtained both from Portuguese community and hospital settings. This panel included 42 isolates identified as MRSA. The lytic capacity of each chimera was evaluated by spotting four different amounts of pure enzyme (10, 5, 1, and 0.2 µg) on a dense lawn of viable cells from each tested isolate, which was produced by incorporating cells from exponentially growing cultures in a soft-agar, phosphate-buffered medium (see Materials and Methods). The chimera-induced lysis was qualitatively evaluated by scoring the relative diameter and turbidity of the lysis halos produced after overnight incubation at 37°C (Supplementary Table S2).

For the highest tested protein amount (10 µg) we observed that both chimeras were able to produce lysis halos in more than 90% of the isolates (Supplementary Fig. S3). For each



**FIG. 2.** Sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE) analysis of Lys168-87 (**A**) and Lys170-87 (**B**) purification. M, molecular weight marker; TSE, total soluble extract loaded in the affinity column; FT, flowthrough of the affinity column; P<sub>AF</sub>, eluted fraction of the affinity peak; P<sub>DS</sub>, eluted fraction of the desalting peak.

TABLE 1. DETAILED DESCRIPTION OF TYPED *STAPHYLOCOCCUS AUREUS* CLINICAL STRAINS

Strain ID	Origin	Isolation date	spa typing Ridom	MLST (ST)	CC*	SCCmec type	Reference
MRSA, <i>n</i> = 30							
N315	Japan	1982	t002	5	5	II	43,60
USA100	United States	1995–2003	t002	5	5	II	53
JP1	Japan	1987	t002	5	5	II	2,60
BM18	United States	1989	t002	5	5	IVa	18,60
HDE288	Portugal	1996	t311	5	5	VI	60,67
COB3	Colombia	1996	t1107	5	5	IV <sub>NT</sub>	33,60
USA300	United States	1995–2003	t008	8	5	IVa	53
GRE120	Greece	1993	t036	8	5	IVh	1
JH1	United States	2000	t002	105	5	II	75
JH9	United States	2000	t002	105	5	II	75
ANS46	Australia	1982	t037	239	5	III	24,36,60
DEN907	Denmark	2001	t037	239	5	NT	26,36
HDC2	Portugal	1992	t421	239	5	IIIB	36,69
HPV107	Portugal	1992	t051	247	5	IA	60,70
BK1953	United States	1995	t051	247	5	IA	60,65
PER184	Spain	1991	t121	247	5	IA	21,60
BK793	Egypt	1961	t008	250	5	I	41,60
COL	United Kingdom	1965	t051	250	5	I	60
MW2	United States	1998	t128	1	15	IVa	6
USA400	United States	1995–2003	t558	1	15	IVa	53
HAR22	Finland	2002	t032	22	22	IVh	35,66
HGSA146	Portugal	2003	t032	22	22	IVh	1
DEN2294	Denmark	2001	t019	30	30	IVc	26
DEN2946	Denmark	2001	t975	30	30	IVc	26
USA200	United States	1995–2003	t018	36	30	II	53
HAR24	Finland	2002	t018	36	30	II	35,66
HAR38	Belgium	1995	t004	45	45	IVa	35,66
WIS	Australia	1995	t123	45	45	V	39
GRE14	Greece	1998	t044	80	80	IV	1
DEN114	Denmark	2001	t044	80	80	IVc	26
MSSA, <i>n</i> = 13							
HSA29	Portugal	1992–1993	t002	5	5	NA	3
NCTC8325	United Kingdom	1943	t211	8	5	NA	16,57
DCC403	Portugal	1996–1997	t148	615	5	NA	3
Draftees728	Portugal	1996–1997	t213	12	12	NA	3
IPOP33	Portugal	2001	t1071	9	15	NA	3
DCC192	Portugal	1996–1997	t084	15	15	NA	3
DCC750	Portugal	1996–1997	t078	25	15	NA	3
IPOP58	Portugal	2001	t189	188	15	NA	3
Draftees721	Portugal	1996–1997	t790	22	22	NA	3
DCC1060	Portugal	1996–1997	t012	30	30	NA	3
IPOP34	Portugal	2001	t884	34	30	NA	3
DCC457	Portugal	1996–1997	t330	45	45	NA	3
IPOP74	Portugal	2001	t645	121	121	NA	3

\*Clonal complexes were determined using the E-burst software <http://saureus.mist.net/>, last accessed on January 25, 2012.

NA, not applied; NT, non-typeable; MLST (ST), multilocus sequence typing (sequence type); SCCmec, Staphylococcal Chromosomal Cassette *mec*; MRSA, methicillin-resistant *S. aureus*; MSSA, methicillin-sensitive *S. aureus*; CC, clonal complex.

protein quantity though, Lys168-87 generally produced larger and more transparent lysis halos than Lys170-87 (Supplementary Table S2). The narrowing of the lytic spectrum of this chimera as the spotted protein amount decreased was more pronounced than that observed with Lys168-87 (Supplementary Fig. S3). Interestingly, these results were basically opposite to those we obtained with the parental Lys168 and Lys170, where the latter exhibited better lytic performance against *E. faecalis*.<sup>62</sup>

Although both chimerical endolysins revealed to be very effective in lysing this panel of *S. aureus* clinical isolates (*n*=100), these were not typed and therefore, the diversity present in this group was unknown. To have a more precise

view on the lytic potential of the chimeras against *S. aureus*, the enzymes were similarly tested in a panel composed of diverse and typed MRSA and MSSA strains (Table 1). This strain collection was previously characterized in terms of their genetic background and included representatives of the most relevant MRSA pandemic clones from different parts of the world and representatives of the dominant of MSSA clones.

Remarkably, the chimerical endolysins lysed about 70% to 100% of this panel of typed *S. aureus* clinical strains, depending on the amount of tested enzyme (Fig. 3). Lys168-87 again showed a better lytic performance when compared with Lys170-87 since 0.2 µg of the former were

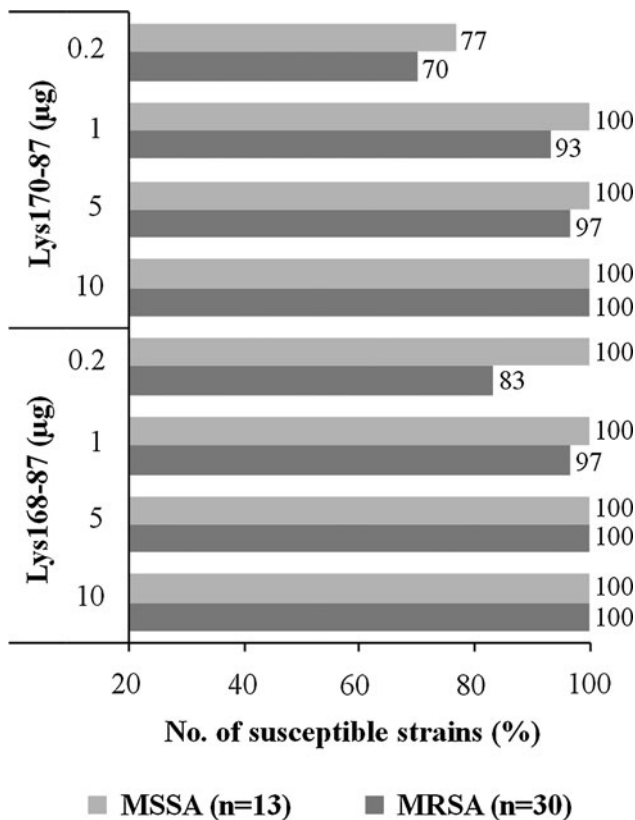


FIG. 3. Susceptibility of a panel of diverse and typed *Staphylococcus aureus* clinical strains to the lytic action of Lys168-87 and Lys170-87. Typed MRSA and MSSA strains (Table 1) were challenged by the spot assay (see text) with the indicated amounts of each chimera. The percentage of strains that developed a lysis halo after overnight incubation at 37°C is plotted as a function of each enzyme amount. A detailed description of the susceptibility of each strain to both chimeras is presented in Supplementary Table S3. MRSA, Methicillin-resistant *S. aureus*; MSSA, Methicillin-sensitive *S. aureus*.

sufficient to produce a lysis halo in 83% of the tested MRSA strains, whereas Lys170-87 lysed only 70%. A detailed description of the lytic activity of the endolysins in each particular *S. aureus* strain is shown in Supplementary Table S3.

#### *Lys168-87 and Lys170-87 have a synergistic lytic effect in liquid media*

The chimerical endolysins were also able to induce lysis of viable *S. aureus* cells in dense liquid suspensions, as shown in the example of Fig. 4, for the MRSA epidemic clone USA200. When applied separately, each endolysin caused a drop of cell suspension turbidity to about 40% of the initial value, after 1 hr incubation, which translated into a corresponding decrease of the initial CFU/ml. However, if simultaneously added to cultures, each at half of the concentration when used separately, the chimerical enzymes decreased the cell suspension turbidity to almost 5% of the initial value and eliminated ~99% of the initial CFU/ml (Fig. 4). These results indicate a synergistic effect of both endolysins in the effective lysis and killing of *S. aureus*.

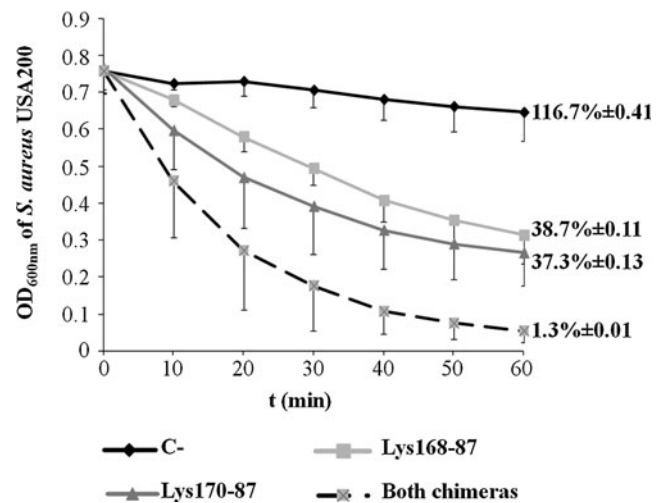


FIG. 4. Lytic action of Lys168-87 and Lys170-87 against *S. aureus* strain USA200 in liquid media. The graphic shows the change of cell suspensions OD<sub>600</sub> over time, after adding each chimera separately (10 µg/ml) or in combination (5 µg/ml each). Endolysin buffer was added to the negative control (C-) instead of the enzymes. Values are the means of three independent experiments with indication of standard deviation. The values on the right side indicate the percentage of the initial CFU/ml after 60 min of enzymes' action and the corresponding standard deviation. OD, optical density.

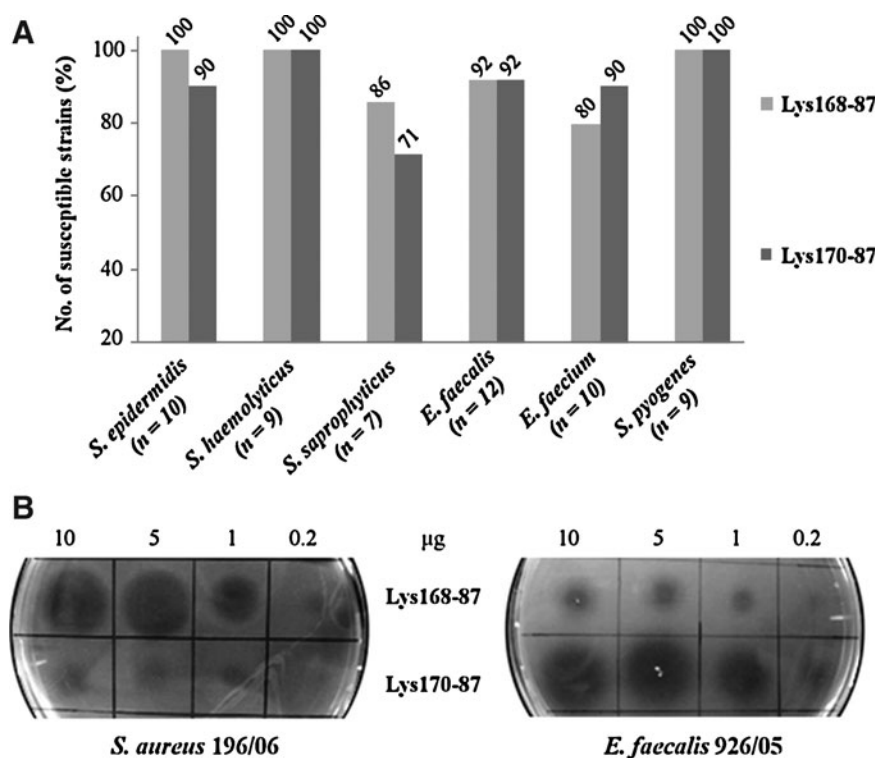
#### *Activity of the chimerical endolysins against other Gram-positive pathogenic bacteria*

As mentioned before, the parental enterococcal endolysins Lys168 and Lys170 showed a clear preference for *E. faecalis* when tested against a representative group of typed strains of this species and of *E. faecium*. When comparing their lytic action, Lys170 showed to be more effective than Lys168.<sup>62</sup> By replacing their putative CWBD by that of the staphylococcal endolysin Lys87 we could efficiently retarget their activity toward *S. aureus*, as initially envisaged (see above). Next, we studied how the chimeras acted on Gram-positive pathogenic bacteria other than *S. aureus* (Supplementary Table S4). When tested against the indicated *E. faecalis* and *E. faecium* clinical isolates, the chimerical enzymes essentially reproduced the lytic profile displayed by the parental enterococcal endolysins,<sup>62</sup> with Lys170-87 producing larger and clearer lysis halos compared with Lys168-87 (Fig. 5 and Supplementary Table S5). Note that the *E. faecalis* and *E. faecium* isolates tested here were not genetically characterized and the vast majority of them have been previously shown to be susceptible to both Lys168 and Lys170.<sup>62</sup>

Interestingly, and in contrast to the parental enterococcal endolysins, the chimeras were also able to cause lysis of clinical isolates of other staphylococcal pathogens and Group A streptococci (Fig. 5A and Supplementary Table S5). Qualitatively, Lys168-87 demonstrated equal or superior lytic action than Lys170-87 against all tested bacteria, except for the enterococcal isolates, where Lys170-87 showed to be more powerful (Fig. 5B and Supplementary Table S5).

## Discussion

A survey of the literature indicates that researchers frequently have to deal with the poor solubility of different



**FIG. 5.** Lytic action of Lys168-87 and Lys170-87 in other Gram-positive bacterial pathogens. **(A)** Bar plot showing the percentage of isolates from each species (Supplementary Table S4) that developed a lysis halo after spotting 10 µg of each chimera. A detailed description of the susceptibility of each isolate to this and lower quantities of each chimera is presented in Supplementary Table S5. **(B)** Representative lysis halos for the indicated amounts of each chimera in a dense lawn of viable cells from *S. aureus* and *E. faecalis* isolates 196/06 and 926/05, respectively.

endolysins, including those of *S. aureus* phages, when aiming their production and purification.<sup>9,31,51,58,74,80</sup> This is also a recurrent problem in our lab and generally the available technical approaches to increase solubility, or to refold proteins recovered from inclusions bodies, are time consuming and poorly effective.

We propose here that construction of chimeras, where we take advantage of the heterologous, highly soluble endolysin functional domains, may represent a solution to obtain large amounts of active lytic enzymes with retargeted and/or extended lytic spectrum relative to parental endolysins. As far as we know, this is the first time that such approach is employed with the specific objective of obtaining highly soluble lytic enzymes targeting *S. aureus*. There is though, at least one report where a similar strategy was attempted to solve the insolubility problem of an *S. aureus* phage endolysin. In this case the putative CD of the endolysin P16 from phage P68 was fused to the inferred CWBD of the virion-associated lysin P17, also from the same phage. However, the resulting chimera remained insoluble and accumulated as inclusion bodies.<sup>51</sup>

The construction of chimerical enzymes as a mean to improve the lytic efficacy or change the lytic spectrum, both *in vitro* and *in vivo*, toward relevant pathogens has been previously described. Recently, a chimerical endolysin (ClyS) assembling a CD and a CWBD from different *S. aureus* phages, and in which the latter displayed *Staphylococcus*-specific binding, was shown to lyse different *S. aureus* MRSA and MSSA strains and several coagulase-negative *Staphylo-*

*coccus* species.<sup>17</sup> This chimera revealed to be more effective than mupirocin for skin decolonization of MRSA and MSSA.<sup>61</sup> In another example of homologous fusion, the CHAP domain of LysK, the endolysin from *S. aureus* phage K, was fused to the CWBD of the bacteriocin lysostaphin from *Staphylococcus simulans*.<sup>38</sup> The recombinant endolysin PRF-119 exhibited antimicrobial activity against a great number of MSSA and MRSA clinical isolates, with an average MIC<sub>90</sub> of 0.391 µg/ml.

Heterologous fusion constructs where the native CWBD of the streptococcal LambdaSa2 endolysin Cpl-7 was replaced by that of either lysostaphin or LysK resulted in a approximately fivefold increase in staphylolytic activity mediated by the Cpl-7 CD, while maintaining significant streptolytic activity.<sup>8</sup> The results suggested that the CWBD of *S. aureus* lytic enzymes may not always be staphylococcal-specific. Yet in another example, chimeras composed of the streptococcal phage endolysin B30, or of its CHAP CD only, fused to the mature form of lysostaphin were shown to display lytic specificity for streptococcal pathogens and *S. aureus*.<sup>22</sup>

In the work reported here, we have fused the CD of two highly soluble *E. faecalis* phage endolysins, Lys168 and Lys170, to the CWBD of the staphylococcal phage endolysin Lys87, which showed high propensity to precipitate in the form of inclusion bodies. Our rationale was that the CD of the enterococcal enzymes would confer high solubility to the resulting chimerical endolysins. In addition, the fact that Lys168 and Lys170 CDs were of the CHAP and Amidase-2



families, respectively, increased the chances of these domains to act on the *S. aureus* cell wall, if targeted by the Lys87 CWBD. Actually, the Lys168 CHAP domain, as deduced by Pfam analysis, shares 98% identity with that of the endolysin of *S. aureus* phage SAP6.<sup>62</sup> Lys170 Amidase-2 CD should confer N-acetylmuramoyl-L-alanine amidase activity as this was the activity experimentally determined for ORF9, an endolysin that is virtually identical to Lys170.<sup>62,76</sup> Amidases cleave the amide bond that links the N-acetyl muramic acid of glycan strands to the L-alanine residue of peptide stems. This bond and linked residues are common to the vast majority of bacterial cell wall peptidoglycans, including that of *S. aureus*.<sup>73</sup>

The chimeras thus constructed proved to efficiently lyse *S. aureus* clinical strains. When tested at their highest concentration each recombinant endolysin was able to induce lysis in more than 96% of 143 *S. aureus* clinical isolates. The lytic efficiency of each chimera varied among the different isolates, as judged by the relative size and transparency of the lysis halos (Supplementary Tables S2 and S3). Remarkably, all the 30 typed strains composing a panel representative of the most relevant MRSA epidemic clones (Table 1) showed to be susceptible to both chimerical enzymes. Very few anti-staphylococcal endolysins have been tested in such a representative group of MRSA typed strains.

Also interesting was the fact that these chimerical endolysins not only essentially maintained the lytic performance of the parental enterococcal endolysins when tested against *Enterococcus*,<sup>62</sup> but were also extended in their lytic spectrum being active against other *Staphylococcus* species and *Streptococcus pyogenes*. This suggests that the Lys87 CWBD may target a bacterial cell wall epitope that is common to several Gram-positive bacterial species, while that of the parental enterococcal endolysins binds to a ligand predominantly found on enterococcal cell wall. This apparent wider target range of the CWBD of anti-staphylococcal lytic enzymes was also observed for lysostaphin and LysK, as mentioned above. Thus, Lys168-87 and Lys170-87 can be added to the small list of broadly active, phage-derived lytic enzymes.<sup>8,22,80</sup>

A curious difference was observed when we compared the lytic efficiency of Lys168 and Lys170 with that of the derived chimeras Lys168-87 and Lys170-87. When tested against their natural target, *E. faecalis*, Lys170 generally produced larger and more transparent lysis halos than Lys168. However, this scenario was inverted when Lys168-87 and Lys170-87 were assayed against *S. aureus* (see illustrative example in Fig. 5B). This appears to indicate that the lytic performance of a given endolysin is not a mere function of the affinity and binding of the CWBD to the cell wall. In fact, other parameters such as the affinity of the CD to its substrate may influence lytic activity.<sup>49</sup>

Lys168-87 and Lys170-87 could also lyse liquid cell suspensions of viable *S. aureus*. In addition, they showed a clear synergistic effect when simultaneously added to cells (Fig. 4). To the best of our knowledge, such synergy between phage-derived hydrolytic enzymes in the effective lysis and killing of *S. aureus* has never been reported. Several studies though have demonstrated the occurrence of synergy between *S. aureus* phage endolysins and bacteriocins such as nisin<sup>31</sup> or lysostaphin<sup>9</sup> or between endolysins and conventional antibiotics.<sup>17,20,51,63</sup>

In conclusion, our results indicate that the engineering of chimerical endolysins from heterologous functional domains can be a good strategy to obtain large quantities of soluble and highly effective peptidoglycan hydrolases, either with narrow or broad lytic spectrum. The efficacy of the chimeras Lys168-87 and Lys170-87 is currently under study in animal models of *S. aureus* infection.

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### Author Disclosure Statement

Technophage has proprietary rights over *E. faecalis* phages F168/08, F170/08, and *S. aureus* phage F87s/06, their encoded endolysins Lys168, Lys170, and Lys87, and the derived chimerical enzymes Lys168-87 and Lys170-87. All authors reviewed and approved the present article before submission. All authors declare that there are no conflicts of interest. All authors contributed to the elaboration of the article. S.F., D.P. and C.C. were responsible for endolysin cloning, production, and purification; SF executed all experiments evaluating lytic action of chimerical endolysins; F.A.S., C.L., C.C., and S.L. were responsible for phage isolation, purification, and extraction of phage DNA; C.L. and S.L. performed phage genomic analysis; C.M. was responsible for handling and molecular typing of *S. aureus* typed clinical strains. H.deL., P.C.-S., M.G., M.P., and C.S.-J. were responsible for the experimental design and work supervision; C.S.-J. coordinated the collaborative work of the involved laboratories.

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