RESEARCH ARTICLE

Vinyl halogenated fatty acids display antibacterial activity against clinical isolates of methicillin-resistant Staphylococcus aureus

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ABSTRACT

Methicillin-resistant Staphylococcus aureus is a pathogen responsible for skin and wound infections, pneumonia, and bloodstream infections. Serious attention is needed because Methicillin-resistant S. aureus is also resistant to many other commonly used antibiotics. This study explores the potential of vinyl halogenated fatty acids as antibacterial agents. Specifically, the total synthesis of vinyl halogenated fatty acids was performed to investigate their antibacterial activity against clinical isolates of methicillin-resistant S. aureus. The novel synthesis of the vinyl halogenated fatty acids was carried out by treating either 2-dodecynoic acid or 2-hexadecynoic acid with an allyl halide and 5 mol\% of bis(benzonitrile)palladium (II) chloride as catalyst. Our results demonstrate that vinyl halogenated fatty acids displayed significant antibacterial activity against clinical isolates of methicillin-resistant S. aureus and low cytotoxicity against eukaryotic Vero Cells. Moreover, it was demonstrated that vinyl brominated fatty acids could disrupt the S. aureus plasma membrane and inhibit the expression of the \textit{norB} gene.
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1. Introduction

Methicillin-resistant Staphylococcus aureus (MRSA) is a pathogen responsible for skin and wound infections, pneumonia, and bloodstream infections. The Centers for Disease Control and Prevention (CDC) describes MRSA as a serious concern that requires prompt attention and rapid action. MRSA is responsible for approximately 323,700 infections in hospitals resulting in 10,600 deaths and an estimated $1.7B in total healthcare costs in the United States. Serious attention is needed because MRSA is resistant to many other commonly used antibiotics. This scenario becomes complicated by the lack of a big pharma pipeline to develop new compounds that target superbugs.

Synthetic unsaturated fatty acids (uFA) are attractive candidates to become next-generation antibacterial agents for treating MRSA infections. Some benefits of synthetic uFA include their high antibacterial efficacy and selectivity against MRSA and their straightforward synthesis that can be performed in just a few synthetic steps. In addition, synthetic uFAs are potential antibacterial agents because they appear to have multiple mechanisms of action, making it more difficult for bacteria to develop resistance to these drugs. Synthetic uFAs can directly kill multi-drug resistant bacteria at very low concentrations (i.e., at micromolar and/or nanomolar levels).

Many reports address the antibacterial properties of synthetic uFA against Gram-positive bacteria. For example, Konthikamee et al. reported the antibacterial activity of 2-alkynoic FA against Gram-positive bacteria, including Staphylococcus aureus. In 2014, Sanabria et al. performed the first study to determine the structural characteristics needed to prepare antibacterial 2-alkynoic FA. The compound that displayed the best overall antibacterial activity towards Gram-positive S. aureus (MIC = 15.5 μg/mL) was the 2-hexadecynoic acid (2-HDA). Moreover, it was significantly effective against methicillin-resistant S. aureus (MRSA) (MIC = 15.6 μg/mL) and clinical isolates of MRSA (MIC = 3.9 μg/mL). In another study, Sanabria et al. reported that 2-HDA displayed bactericidal activity against five CIMRSA strains (MIC = 0.24-0.49 μg/mL), which were resistant to Ciprofloxacin (Cipro), indicating that 2-HDA was the most effective treatment against CIMRSA.

The potential of both naturally occurring and synthetic halogenated FA against Gram-positive bacteria have been reviewed. Specifically, it was reported that FA containing triple bonds, double bonds, and either bromine or chlorine atoms in their chemical structures are pivotal in their biological activity against Gram-positive bacteria, including S. aureus. In 2021, Latifah et al. reported that brominated C18 and C20 FA isolated from Xestospongia testudinaria displayed antibacterial activity against Vibrio harveyi, Vibrio parahaemolyticus, and the Vibrio alginolyticus. The above studies emphasize the halogen moiety in the FA’s antibacterial activity. Furthermore, in another study performed by Metrangolo and Resanti, it was reported that incorporating halogen atoms into drug candidates improves the drug’s intracellular delivery because of an increment in its lipophilicity, which improves its internalization through lipid membranes and tissues. Several studies in medicinal chemistry utilize halogen substitution to improve drug pharmacokinetics. For this reason, halogenated FA holds promise as antibacterial agents.

A recent study reveals that the antimicrobial activity of peptoids, specifically the brominated analogs, displayed up to a 32-fold increase in the activity against S. aureus than its non-halogenated counterparts. Another study demonstrated that the bromination of Marula seed oil improved the inhibition activity against S. aureus. We recently reported the synthesis of novel vinyl halogenated FA (VHFA) and demonstrated that these FA displayed antileishmanial activity against Leishmania infantum promastigotes.

In this study, we explored the potential of VHFA as antibacterial agents. Herein, we also report the total synthesis of VHFA and the antibacterial activity against clinical isolates of MRSA. The antibacterial 2-allyl-3-bromo-2E-dodecenoic acid (1a), 2-allyl-3-chloro-2E-dodecenoic acid (2a), 2-allyl-3-bromo-2E-hexadecenoic acid (1b), and 2-allyl-3-chloro-2E-hexadecenoic acid (2b) were synthesized as displayed in Scheme 1.
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![Diagram](https://example.com/diagram.png)

**Scheme 1.** Total synthesis of the antibacterial VHFA 1a-1b and 2a-2b.

2. Materials and Methods

2.1 Instrumentation

The glassware was flame dried under Ar or N₂ gas and assembled with a magnetic bar, stirrer, and septum. THF was distilled over CaH₂ and dried over Na and benzophenone before use. Hexane, ether, and ethyl acetate were purified using simple distillation. DMI and other reagents were purchased from Aldrich and Acros Organics. All reactions were analyzed by ¹H NMR (300, 400, or 700 MHz) and ¹³C NMR (75, 100, or 176 MHz) using a Bruker DPX-300, Bruker Advance DRX-400, or Bruker Ascend 700 spectrometer. The samples were dissolved in 99.8% chloroform-d (CDCl₃), and the solvent signals at 7.26 (¹H) and 77.0 (¹³C) ppm were used as internal standards for hydrogen and carbon, respectively. The IR spectra were measured either on a Bruker Tensor 27 FT-IR spectrometer or a Thermo Nicolet 1S5.

2.2 Synthesis of vinyl halogenated fatty acids

2.2.1. 2-Allyl-3-bromo-2E-dodecenoic acid (1a)

To 0.75 g (3.82 mmol) of 2-dodecenoic acid and bis(benzonitrile) palladium (II) chloride (0.073 g, 0.19 mmol, 5 mol%), was added dropwise, under a N₂ atmosphere, 33.0 mL (380 mmol) of allyl bromide at 0 °C. The reaction was kept for 24 h. The crude product was purified with silica gel column chromatography eluting with hexane and ethyl acetate (9:1) to afford 1a (0.76 g, 0.78 mmol) as a yellow/orange oil in a 73% yield. IR (neat) λmax: 3700-2700 (O-H), 3082, 2954, 2923, 2853, 1683 (C=O), 1640 (C=C), 1602 (C=C), 1456, 1437, 1405, 1281, 1225, 1123, 1039, 990, 915, 722, 669 (C-Cl) cm⁻¹. ¹H-NMR (CDCl₃ 300 MHz) δ (ppm) 5.85 (1H, ddt, Jtrans = 17 Hz, Jcis = 10 Hz, Jcis = 6.2 Hz, H-2), 5.11 (2H, brd, H-3), 3.32 (2H, brd, H-1'), 2.89 (2H, brt, H-4), 1.67 (2H, m, H-5), 1.28 (12H, s, H-6 to H-12), 0.91 (3H, t, J = 6.7 Hz, -CH₃); ¹³C-NMR (CDCl₃, 75 MHz) δ (ppm) 171.59 (s, C = O), 147.06 (d, C-3), 135.58 (t, C-3'), 129.35 (s, C-2), 116.37 (d, C-2'), 40.40 (t), 38.84 (t), 31.86 (t), 29.43 (t), 29.31 (t), 29.27 (t), 28.92 (t), 28.74 (t), 28.20 (t), 14.07 (q).

2.2.2. 2-Allyl-3-chloro-2E-dodecenoic acid (2a)

To 0.75 g (3.82 mmol) of 2-dodecenoic acid and bis(benzonitrile)palladium(II) chloride (0.073 g, 0.5 mol%) was added dropwise 33.0 mL (380 mmol) of allyl chloride under N₂ at 0 °C. The reaction mixture was stirred for an additional 12 h. The crude product was purified with silica gel column chromatography eluting with hexane and ethyl acetate (9:1) to afford 2a (0.76 g, 0.78 mmol) as a yellow/orange oil in a 73% yield. IR (neat) λmax: 3700-2700 (O-H), 3082, 2954, 2923, 2853, 1683 (C=O), 1640 (C=C), 1602 (C=C), 1456, 1437, 1405, 1281, 1225, 1123, 1039, 990, 915, 722, 669 (C-Cl) cm⁻¹. ¹H-NMR (CDCl₃ 300 MHz) δ (ppm) 5.85 (1H, ddt, Jtrans = 17 Hz, Jcis = 10 Hz, Jcis = 6.1 Hz, H-2'), 5.10 (2H, m, H-3'), 3.32 (2H, brd, H-1'), 2.90 (2H, brt, H-4), 1.67 (2H, m, H-5'), 1.30 (12H, s, H-6 to H-12), 0.91 (3H, t, J = 6.6 Hz, H-12), ¹³C-NMR (CDCl₃, 75 MHz) δ (ppm) 171.59 (s, C-1'), 153.15 (d, C-3), 135.59 (t, C-3'), 126.65 (s, C-2'), 116.10 (d, C-2'), 38.02 (t), 35.53 (t), 31.86 (t), 29.43 (t), 29.31 (t), 29.26 (t), 28.88 (t), 28.20 (t), 25.28 (t), 14.06 (q).

2.2.3. 2-Allyl-3-bromo-2E-hexadecenoic acid (1b)

To 1.00 g (3.96 mmol) of 2-hexadecenoic acid and bis(benzonitrile)palladium (II) chloride (0.0759 g, 0.2 mmol, 5 mol%) were added 32.2 mL (396 mmol) of allyl bromide under argon at 0 °C. The solution was stirred for 24 h. The crude...
reaction mixture was filtered through celite and rotoevaporated. The crude product was purified using silica gel column chromatography eluting with hexane/ether (B:2) affording 1b (0.494 g, 1.51 mmol) as a light-yellow solid (mp 34-36 °C) in 38% yield. IR (neat) \( \nu_{\text{max}} = 3500-2500 \) (O-H), 2957, 2916, 2847, 1684 (C=O), 1638 (C=C), 1609, 1458, 1433, 1402, 1281, 1227, 918, 725, 640, 609 (C-Br) cm\(^{-1}\). \(^1\)H-NMR (CDCl\(_3\)) 700 MHz) \( \delta \) (ppm) 5.85 (1H, m, H-2'), 5.15 (1H, ddt, \( J_{\text{trans}} = 17.6 \) Hz, \( J_{\text{gem}} = 1.5 \) Hz, \( J_{\text{allyl}} = 1.3 \) Hz, H-3'\( \text{trans} \)), 5.10 (1H, ddt, \( J_{\text{cis}} = 10.1 \) Hz, \( J_{\text{gem}} = 1.3 \) Hz, \( J_{\text{allyl}} = 1.3 \) Hz, H-3'\( \text{cis} \)) 3.34 (2H, br d, H-1'), 2.99 (2H, br t, H-4), 1.67 (2H, quint, H-5), 1.30 (18H, s, H-6-15), 0.90 (3H, m, H-16); \(^{13}\)C-NMR (CDCl\(_3\)) 176 MHz) \( \delta \) (ppm) 170.73 (s, C-1), 146.94 (d, C-2), 133.31 (t, C-3'), 129.33 (s, C-3), 116.38 (d, C-2'), 40.40 (t), 38.87 (t), 36.64 (t), 31.94 (t), 29.70 (t), 29.68 (t), 29.65 (t), 29.52 (t), 29.38 (t), 29.35 (t), 28.93 (t), 28.77 (t), 22.71 (t), 14.13 (q).

2.2.4. 2-Allyl-3-chloro-2E-hexadecenoic acid (2b)

To 1.00 g (3.96 mmol) of 2-nonadecenoic acid and bis(benzonitrile)palladium(II) chloride (0.0759 g, 0.2 mmol, 5% mol), were added 32.2 mL (396 mmol) of allyl chloride under a nitrogen atmosphere at 0 °C. The solution was further stirred at this temperature for 24 h. The crude mixture was also filtered through celite and the solvent rotoevaporated. The crude product was purified using silica gel column chromatography and eluting with hexane/ethyl acetate (9:1) affording 2b (0.992 g, 2.41 mmol) as a light-yellow solid (mp 32-34 °C) in 67% yield. IR (neat) \( \nu_{\text{max}} = 3500-2500 \) (O-H), 2594, 2594, 2516, 2848, 1680 (C=O), 1639 (C=C), 1607, 1462, 1445, 1281, 1231, 910, 723, 669 (C=Cl) cm\(^{-1}\). \(^1\)H NMR (CDCl\(_3\)) 400 MHz) \( \delta \) (ppm) 5.88 (1H, m, H-2'), 5.15 (1H, ddt, \( J_{\text{trans}} = 17.2 \) Hz, \( J_{\text{gem}} = 1.7 \) Hz, \( J_{\text{allyl}} = 1.6 \) Hz, H-3'\( \text{trans} \)), 5.10 (1H, ddt, \( J_{\text{cis}} = 10.1 \) Hz, \( J_{\text{gem}} = 1.8 \) Hz, \( J_{\text{allyl}} = 1.4 \) Hz, H-3'\( \text{cis} \)) 3.34 (2H, br d, H-1'), 2.92 (2H, br t, H-4), 1.69 (2H, quint, H-5), 1.31 (18H, m, H-6-15), 0.93 (3H, m, H-16); \(^{13}\)C NMR (CDCl\(_3\)), 100 MHz) \( \delta \) (ppm) 172.04 (s, C-1), 153.26 (d, C-2), 133.56 (t, C-3'), 126.72 (s, C-3), 116.07 (d, C-2'), 38.82 (t), 38.03 (t), 35.50 (t), 31.92 (t), 31.58 (t), 29.68 (t), 29.65 (t), 29.63 (t), 29.49 (t), 29.36 (t), 28.89 (t), 28.21 (t), 22.62 (t), 14.05 (q).

2.3. Bacterial strains and growth conditions

The Gram-positive Staphylococcus aureus (ATCC 29213) and the Gram-negative Escherichia coli (ATCC 25922) were obtained from the American Type Culture Collection (Manassas, VA, USA). In addition, CIMRSA strains were generously given by a community hospital in San Juan, Puerto Rico (USA). Stock cultures were preserved on blood agar (TSA with 5% sheep blood, Remel, and Oxoid Microbiology Products, Lenexa, KS). Inoculation of a single colony in 5 mL of Trypticase Soy Broth (TSB, BD Diagnostic Systems, Franklin Lakes, NJ) was carried out to prepare suspension cultures that were incubated for 16-18 h at 37 °C. Before preparing susceptibility assays, bacteria cells were resuspended in TSB and visually standardized using 0.5 McFarland standard, providing an equivalent concentration of 1.0 x 10^8 colony-forming units (CFU)/mL.

2.4. Susceptibility testing

Susceptibility tests were conducted as described by Sanabria-Rios et al. Briefly, VHFA stock solutions were prepared using 100 % DMSO. Stock solutions were serially diluted with sterile TSB, and 100 µL of each dilution were transferred to a flat-bottomed microplate well preincubated with 10 µL of TSB containing 4-5 x 10^3 CFU. Each well was inspected using a spectrophotometer at 620 nm using a positive control well (containing the bacterial inoculated TSB but not the VHFA treatment) and a negative control well (containing sterile TSB without VHFA solution) for further comparisons. The minimum inhibitory concentration (MIC) was the concentration at which the test compound did not show turbidity in the well after incubation for 16-18 h at 37 °C.

2.5 Cytotoxicity assays

Vero Cells ATCC CCL-81.5 (Kidney African Green Monkey, Manassas, VA, USA) cells were cultured in DMEM (ATCC, Manassas, VA, USA) supplemented with 10% FBS (Hyclone, Waltham, MA, USA). Vero cells were seeded at a density of 6.4 x 10^4 cells/mL (25,000 cells/well) of medium in a 96-well plate (0.2mL/well) in triplicate (N = 3). After 24 h, cells were treated at various concentrations. Controls containing 1 % DMSO were included in the experimental design for all assays. After 48 h of exposition, the medium containing the FA understudy was removed, washed with PBS 1X (Hyclone, GE Healthcare Life Sciences, UT, USA) and MTS [3-(4,5-dimethylthiazol-2-yl)-5-(3-carboxy-methoxyphenyl)-2-(4-sulfophenyl)-2H-tetrazolium] solution (Promega, Wisconsin, USA) was added to each well (0.2mL/well) and incubated for 2h. Absorbance was read at 490 nm
with a spectrophotometer (Bio-Rad x Mark Microplate Spectrophotometer). Cell viability was quantitatively assessed by the color change associated with the bio-reduction of the MTS compound. The % of viable cells was determined based on the control.

2.6 Methicillin-resistant S. aureus kinetic growth assays

We followed a modified version of a previously reported procedure.® CIMRSA XIII strain was grown to an optical density of 1.0 (at 600 nm) and diluted 30,000-fold in TSB. A 10 μL aliquot of diluted bacteria was added to each well of a flat-bottom 96-well plate containing 100 μL of TSB with the appropriate concentration (4 x MIC) of either palmitic acid, 1a, or Cipro. All FA dilutions contained 1% DMSO. TSB medium containing 1% DMSO was used as a control with no treatment. The plate was incubated at 37 °C for 20 h using a Thermo Scientific Varioskan Lux plate reader and then read at 600 nm.

2.7. Permeability studies of the HVFA on the methicillin-resistant S. aureus cell

CIMRSA XIII was grown in 5 mL of TSB to an optical density of 1.0 at 600 nm. A 45 mL sterile TSB was inoculated with 5 mL of CIMRSA XIII and treated with a VHFA stock solution to reach a final concentration of 4 x MIC. All VHFA solutions contained 1% DMSO. TSB medium containing 1% DMSO was used as a control with no treatment. Palmitic acid was used as the negative control. CIMRSA XIII cultures containing 1% DMSO and FA were centrifuged at 5,000 rpm at 4°C for 20 minutes. The resulting pellets were washed three times with 1 x PBS and resuspended with 1,000 μL of the same aqueous medium. A 10 μL of bacterial suspension was diluted with 990 μL of 1 x PBS to obtain a 1:100 bacterial suspension. To the 1:100 bacterial suspension, 1,000 μL of 1:500 of 1% fluorescein sodium salt (Sigma Aldrich, St. Louis, MO, USA) in 1 x PBS was added and incubated at 4°C for 8-10h. After incubation, a 60 μL of the 1:100 bacterial suspension was fixed in a microscope slide by adding 40 μL of a solution containing 3:1 ethanol-glacial acetic acid and 4% glycerol. The fixed bacteria were washed three times with 1 x PBS, and the excess of the aqueous solution was removed with a wipe. The fixed bacteria were visualized by fluorescence microscopy (Motic, AE31E Trinocular Inverted Microscope, Richmond, Canada) using a FITC filter.

2.8 q RT-PCR

A 100 mL of TSB was inoculated with 20 mL of TSB containing CIMRSA XIII and subsequently treated with the corresponding treatments (Cipro, 1b, or 2b) at their GI50s. A 25 mL of each cultured TSB was pelleted, and RNA was extracted using the AllPrep® Bacterial DNA/RNA/Protein Kit (Qiagen, California, USA) following the manufacturer's instructions. cDNA synthesis was prepared using the Power SYBR™ Green RNA-to C® 1-Step Kit (Applied Biosystems, CA, USA) with the following primers: 5’-ATGGAAAAAAGCCGTCAGAGAG-3’ (forward primer) and 5’-AAACATGATTGTCGAAATAGC-3’ (reverse primer) according to the manufacturer’s instructions. The resulting cDNA templates were quantitatively amplified in a Step One Real-Time PCR System (Applied Biosystems, California, USA). The cycling parameters included: initial denaturation at 95°C for 4 min, followed by 40 cycles of denaturation (95°C, 45s), annealing (57°C, 45s), and final extension (72°C, 55s), and final extension at 72°C for 2 min. Efflux pump genes' relative expression was calculated using the ΔΔCT method, in which the amount of the desired cDNA was normalized to an endogenous reference (housekeeping gene 16S rRNA) and relative to an in vitro calibrator, is given by the variable 2^-ΔΔCT, where Ct is the cycle number of the detection threshold.

3. Results

The synthesis of the VHFA was performed as described in the literature.6,20,21 Once the synthesis of VHFA 1 and 2 was carried out, we determined the antibacterial activity against CIMRSA strains, as described by Sanabria-Rios et al.7 Results from antibacterial activity tests are displayed in Table 1. In this study, CIMRSA XIII was selected as a bacterial model for further mechanistic studies because this strain was very resistant to Ciprofloxacin, a broad-spectrum antibiotic used to treat bacterial infections.22
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### Table 1. Antibacterial activity of 1a-1b and 2a-2b against several S. aureus strains and E. coli.

| Compounds | MIC/GI<sub>50</sub> ± SEM (µg/mL) | Bacterial strains |
|-----------|-----------------------------------|-------------------|
|           | E. coli (ATCC 25922) | S. aureus (ATCC 29213) | CIMRSA XI | CIMRSA XII | CIMRSA XIII | CIMRSA XIV | CIMRSA XV | CIMRSA XVI |
| 1a        | >1000 | >128 | 62.5/27.5 ± 1.2 | 250/95.1 ± 6.0 | 62.5/46.8 ± 1.2 | 62.5/30.4 ± 1.2 | 62.5/24.4 ± 1.1 | 62.5/42.5 ± 1.1 |
| 1b        | >1000 | 62.5/45.4 ± 3.3 | 7.81/5.4 ± 1.1 | 15.6/10.0 ± 1.3 | 7.81/4.7 ± 1.9 | 15.6/9.3 ± 1.5 | 15.6/9.1 ± 1.1 | 7.81/4.1 ± 1.1 |
| 2a        | >1000 | 15.6/6.6 ± 1.2 | 31.3/13.3 ± 12 | 62.5/47.7 ± 1.3 | 31.3/17.8 ± 1.2 | 31.3/6.7 ± 1.2 | 125/64.4 ± 1.1 | 125/12.1 ± 1.2 |
| 2b        | >1000 | 125/86.5 ± 1.1 | 31.3/16.4 ± 1 | 62.5/54.0 ± 1.4 | 62.5/21.8 ± 1.4 | 125/72.4 ± 1.1 | 62.5/40.1 ± 1.1 | 62.5/22.6 ± 1.1 |
| 2-HDA<sup>b</sup> | >1000 | 15.6/6.2 ± 0.9 | 0.49/0.30 ± 0.03 | 0.49/0.25 ± 0.03 | 0.24/0.16 ± 0.02 | 0.49/0.34 ± 0.03 | 0.49/0.27 ± 0.01 | 0.49/0.30 ± 0.03 |
| Cipro<sup>b</sup> | <0.008 | 0.25/0.14 ± 0.01 | 7.8/5.7 ± 1.1 | 31.3/17.2 ± 1.0 | 62.5/30.4 ± 1.4 | 0.49/0.35 ± 0.04 | 7.8/4.7 ± 1.1 | 7.8/2.3 ± 1.1 |

<sup>a</sup>Experiments were performed in triplicate (N = 3). GI<sub>50</sub> values were obtained from dose-response curves.

<sup>b</sup>These results were obtained from Sanabria et al. 2020<sup>7</sup>

Table 1 shows that HVFA were particularly active against Gram-positive bacteria, and no cytotoxic effect was observed in the Gram-negative E. coli. Also, it can be appreciated that 1b is a 4- to 8-fold better antibacterial agent than 1a, 2a, and 2b in CIMRSA XIII. It is imperative to point out that CIMRSA XIII is the most resistant MRSA strain to Cipro than the other S. aureus strains displayed in Table 1<sup>7</sup>.

To assess the cytotoxicity of the VHFA that displayed the highest inhibitory effect towards the CIMRSA strains, 1b, and its chlorinated counterpart, 2b, their cytotoxic effect on the growth of Vero Cells ATCC CCL-81.5 (Kidney African Green Monkey, Manassas, VA, USA) was tested using the MTS approach. The experiments were performed in triplicate (N = 3), as described below.

**A)**

**B)**

Figure 1. Graph bars showing the cytotoxicity of the VHFA 1b (Figure 1A, red box) and 2b (Figure 1B, blue box) against Vero Cells ATCC CCL-81.5. Vero cells were treated with either 1b or 2b at concentrations ranging from 0.0001 to 100 µg/mL. MTS reagent was added to cells after 48 h incubation. Experiments were performed in triplicates (N=3). Graphs analyses were performed with GraphPad Prism v. 6.01 (GraphPad Software).
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**Figure 1** shows that 1b was not cytotoxic against eukaryotic Vero Cells at a concentration ranging from 0.0001 to 10 μg/mL. Additionally, only 19% of cytotoxicity was observed when Vero Cells were treated with 100 μg/mL. Similar behavior was observed when Vero Cells were treated with 100 μg/mL of 2b, where only 20% of cytotoxicity was obtained.

Kinetic growth assays were performed to determine whether CIMRSA XIII is adapted to metabolize 1b (**Figure 2**). These assays were performed as described above.

**Figure 2**. CIMRSA XIII growth curve showing its log-phase in TSB when treated with 1b (31.2 μg/mL), Cipro (250 μg/mL), and palmitic acid (4 mg/mL). CIMRSA XIII treated with 1% DMSO was used as control. Experiments were performed in three biological replicates (N =3).

**Figure 2** reveals that 1b displayed similar cytotoxic activity to Cipro against CIMRSA XIII. Also, it was observed that palmitic acid was less effective in inhibiting the growth of CIMRSA XIII than 1b. The difference in O.D.600 values between the palmitic acid and control curves can be explained by the slight turbidity when preparing TSB solutions containing palmitic acid. Both 1b and Cipro arrested the growth of CIMRSA XIII within 600 minutes of its addition. The effectiveness of 1b as an antibacterial agent is comparable with the Cipro curve at 31.2 μg/mL. In these assays, Cipro required a concentration of 250 μg/mL to inhibit the growth of CIMRSA XIII. Altogether, these results agree with those in Table 1 and demonstrate the potency of 1b as an antibacterial agent in ciprofloxacin-resistant *S. aureus*.

Permeability studies on CIMRSA XIII were performed to acquire further knowledge about the mechanism of action of 1b. The CIMRSA XIII cells used in these studies were visualized by fluorescence microscopy using a FITC filter. The results are shown in **Figure 3**. It can be observed that the CIMRSA XIII treated with 1% DMSO and palmitic acid did not permeabilize fluorescein inside the bacterial cell (see yellow arrows). On the other hand, the CIMRSA XIII treated with 1b showed no well-defined membranes, the bacterium’s size was reduced, and fluorescein was observed inside the bacterium. These results suggest that 1b kills CIMRSA XIII, possibly disrupting the cell membrane.
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Continuing the search to find insight into the mechanism of action of 1b in terms of its inhibitory effect on MRSA strains, we studied its effect in expressing the norB gene in <i>S. aureus</i>. The norB gene regulates the NorB protein's expression, an efflux pump that creates resistance to quinolones and other agents in <i>S. aureus</i>. In addition, NorB is a surface protein that provides integrity to the <i>S. aureus</i> plasma membrane. Therefore, q RT PCR was used to determine the inhibitory effect of 1b and its chlorinated analog 2b in the expression of the norB gene in <i>S. aureus</i>. Results from this study are displayed in Figure 4.

Results in Figure 4 demonstrate that both 1b and 2b significantly decreased the expression of norB when CIMRSA XIII was treated with these compounds. Another aspect observed in Figure 4 is
that Cipro significantly overexpresses the norB gene, implying that CIMRSA is producing sufficient NorB to regulate the internalization of Cipro into the bacterial cell. Overexpression of norB in CIMRSA XIII is consistent with findings previously reported in the literature.24

4. Discussion

In 2020, it was determined that 2-HDA was effective in inhibiting the growth of CIMRSA strains compared with other structural isomers of this uFA.27 In this study, we explored the antibacterial activity of VHFA in six MRSA strains that are also resistant to Ciprofloxacin. Our results demonstrate that 1b displayed significant antibacterial activity against the six CIMRSA strains in Table 1. Also, our results show that a cis double bond, a vinyl group, and a bromine atom are important for the antibacterial activity against CIMRSA. Another relevant structure-activity relationship (SAR) information that can be detached from Table 1 is that the presence of chlorine atoms decreases the antibacterial activity of VHFA against CIMRSA. Also, it can be observed that compounds 1a, 1b, 2a, and 2b were not active against the Gram-negative E. coli. This phenomenon can be due to the difference in bacterial cell wall composition. Gram-positive bacteria are absent of an outer membrane but are surrounded by thicker layers of peptidoglycan than Gram-negative bacteria.25 The lack of bioactivity of VHFA 1 and 2 in E. coli agrees with other studies reporting the selectivity of FA towards Gram-positive bacteria.6,7

The cytotoxic effect of HVFA 1b and 2b on Vero Cells’ growth was also assessed using the MTS approach to determine the selectivity of these compounds in prokaryotic cells. Results in both Table 1 and Figure 1 demonstrate that 1b and 2b were highly selective in S. aureus than in Vero Cells. Furthermore, our results demonstrated that HVFA are less cytotoxic in eukaryotic cells than 2-HDA which highlights their potential as antibacterial agents.

Results in Figure 2 demonstrate that 1b is a potent growth inhibitor since this compound displayed similar growth behavior to 2-HDA.7 In addition, 1b displayed similar kinetic growth to palmitoleic acid (16:1 Δ9), a FA that has shown a cytotoxic effect on the growth of S. aureus by rapid membrane depolarization, disruption of the bacterial plasma membrane, and the release of critical proteins in the medium.26 Thus, Figure 2 supports the idea that 1b can disrupt the plasma membrane in S. aureus.

In order to better understand the mechanism of action of 1b, we performed permeability studies involving CIMRSA XIII (Figure 3). Our results demonstrate that 1b promotes fluorescein internalization inside the bacterial cell. This finding supports our hypothesis that membrane disruption occurs when MRSA is treated with 1b.

When results in Figures 3 and 4 are analyzed, these suggest that VHFA provoke disruption of the S. aureus plasma membrane by inhibiting the expression of norB, which has repercussions on the biosynthesis of the NorB efflux pump. NorB is a surface protein that confers resistance to some antibiotics, such as hydrophilic fluoroquinolones, biocides, hydrophobic fluoroquinolones, and tetracycline.27 To some extent, 1b and 2b are internalizing the S. aureus plasma membrane, thus affecting the norB gene expression, as observed in Figure 4.

5. Conclusions

We further investigated how bromine and chlorine atoms affect the antibacterial activity of VHFA in CIMRSA. This study showed that VHFA containing a bromine atom is pivotal for their antibacterial activity against CIMRSA strains resistant to Cipro. Moreover, it was outlined that vinyl brominated FA can disrupt the S. aureus plasma membrane, likely by inhibiting the expression of the norB gene. This study will undoubtedly impact the understanding of the antibacterial properties of halogenated FA and highlight 1b as a novel agent against nosocomial bacterial infections.

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References

1. Centers for Disease Control and Prevention. *Antibiotic resistance threats in the United States, 2019*. CS298822-A; Centers for Disease Control and Prevention: Atlanta, GA, 2019, 95-96.

2. Gade N.D., M.S. Q. Fluoroquinolone therapy in *Staphylococcus aureus* infections: Where do we stand? *J Lab Physicians*. 2013;5(2):109-112. DOI: https://dx.doi.org/10.1177/1574891112801619728

3. Walker DD, David MZ, Catalano D, Daum R, Gluth MB. In vitro susceptibility of ciprofloxacin-resistant methicillin-resistant *Staphylococcus aureus* to ototopical therapy. *Otolaryngology*. 2018;158(5):923-939. DOI: https://doi.org/10.1177/1574891112801619728

4. Spetalnick BM, Powers JS. Eradication of methicillin-resistant *Staphylococcus aureus* colonization with ciprofloxacin. *J Am Geriatr Soc*. 1990;38(3):389-390. DOI: https://doi.org/10.1111/j.1532-5415.1990.tb03532.x

5. Desbois AP. Potential applications of antimicrobial fatty acids in medicine, agriculture and other industries. *Recent Pat Antiinfect Drug Discov*. 2012;7(2):111-122. DOI: https://doi.org/10.2174/157489112801619728

6. Sanabria-Rios DJ, Rivera-Torres Y, Maldonado-Domínguez G, et al. Antibacterial activity of 2-alkynoic fatty acids against multidrug-resistant bacteria. *Chemistry and physics of lipids*. Feb 2014;178:84-91. DOI: https://doi.org/10.1016/j.chemphyslip.2013.12.006

7. Sanabria-Rios DJ, Morales-Guzman C, Mooney J, et al. Antibacterial activity of hexadecenoic acid isomers toward clinical isolates of multidrug-resistant *Staphylococcus aureus*. *Lipids*. 2020;55(2):101-116. DOI: https://doi.org/10.1002/lipd.12213

8. Konthikamee W, Gilbertson JR, Langkamp H, Gershon H. Effect of 2-alkynoic acids on in vitro growth of bacterial and mammalian cells. *Antimicrob Agents Chemother*. November 1, 1982 1982;22(5):805-809. DOI: https://doi.org/10.1128/AAC.22.5.805

9. Sanabria-Rios DJ, Rivera-Torres Y, Rosario J, et al. Chemical conjugation of 2-hexadecenoic acid to C5-curcumin enhances its antibacterial activity against multi-drug resistant bacteria. *Bioorg Med Chem Lett*. 2015;25(22):5067-5071. DOI: https://doi.org/10.1016/j.bmcl.2015.10.022

10. Dembtsky VM, Srebnik M. Natural halogenated fatty acids: Their analogues and derivatives. *Prog Lipid Res.* 2002;41(4):315-367. DOI: https://doi.org/10.1016/S0163-7827(02)00003-6

11. Hirsh S, Carmely S, Kashman Y. Brominated unsaturated acids from the marine sponge *Xestospongia* sp. *Tetrahedron*. 1987;43(14):3257-3261. DOI: https://doi.org/10.1016/S0040-4020(01)90293-5

12. Bourguet-Kondracki ML, Rakotoarisoa MT, Martin MT, Guyot M. Bioactive bromopolyacetylenes from the marine sponge *Xestospongia testudinaria*. *Tetrahedron Lett.* 1992;33(2):225-226. DOI: https://doi.org/10.1016/0040-4020(92)88056-8

13. Latifah LA, Seokamto NH, Tahir A. New antibacterial activities of brominated C18 and C20 fatty acids isolated from marine sponge *Xestospongia testudinaria* against shrimp pathogenic bacteria. *RASÂYAN J Chem*. 2021;14(1):460-465. DOI: https://doi.org/10.31788/RJC.2021.141599

14. Priimagi A, Cavallo G, Metrangolo P, Resnati G. The halogen bond in the design of functional supramolecular materials: Recent advances. *Acc Chem Res*. 2013;46(11):2686-2695. DOI: https://doi.org/10.1021/ar400103r

15. Wilcken R, Zimmermann MO, Lange A, Joerger AC, Boeckler FM. Principles and applications of halogen bonding in medicinal chemistry and chemical biology. *J Med Chem*. 2013;56(4):1363-1388. DOI: https://doi.org/10.1021/jm3012068

16. Xu Z, Yang Z, Liu Y, et al. Halogen bond: Its role beyond drug–target binding affinity for drug discovery and development. *J Chem Inf Model*. 2014;54(1):69-78. DOI: https://doi.org/10.1021/ci400539q

17. Hernandes ZM, Cavalcanti TSM, Moreira MDR, de Azevedo Junior FW, Leite LAC. Halogen Atoms in the Modern Medicinal Chemistry: Hints for the Drug Design. *Curr Drug Targets*. 2010;11(3):303-314. DOI: https://doi.org/10.2174/138945010790711996

18. Molchanova N, Nielsen J, Sørensen K, et al. Halogenation as a tool to tune antimicrobial activity of peptoids. *Sci Rep*. 2020;10:14805. DOI: https://doi.org/10.1038/s41598-020-71771-8

19. Ibrahim JS, Adamu HM, Shakeeda OI. Antibacterial activity of Marula [Sclerocarya Birrea]
and brominated Marula seed oil. IJISRT.

20. Carballeira NM, Alequín D, Lotti Diaz LM, et al. synthesis of a novel brominated vinylic fatty acid with antileishmanial activity that effectively inhibits the Leishmania topoisomerase IB enzyme mediated by halogen bond formation. Pure Appl Chem 2019;91(8):1405-1416.
DOI: https://doi.org/10.1515/pac-2018-1113

21. Carballeira NM, Sanabria D, Cruz C, et al. 2,6-Hexadecadiynoic acid and 2,6-nonadecadiynoic acid: Novel synthesized acetylenic fatty acids as potent antifungal agents. Lipids. 2006;41(5):507-511.

22. Fass RJ. Ciprofloxacin. Best use of this new broad-spectrum antibiotic. Postgrad Med. Jun 1990;87(8):117-131.
DOI: https://doi.org/10.1080/00325481.1990.11704677

23. Steinbuch KB, Fridman M. Mechanisms of resistance to membrane-disrupting antibiotics in Gram-positive and Gram-negative bacteria. 10.1039/C5MD00389J. Med Chem Comm. 2016;7(1):86-102.
DOI: https://doi.org/10.1039/C5MD00389J

24. Kwak YG, Truong-Bolduc QC, Bin KH, et al. Association of norB overexpression and fluoroquinolone resistance in clinical isolates of Staphylococcus aureus from Korea. J Antimicrob Chemother. 2013;68(12):2766-2772.
DOI: https://doi.org/10.1093/jac/dkt286

25. Silhavy TJ, Kahne D, Walker S. The bacterial cell envelope. Cold Spring Harb Perspect Biol. 2010;2(5):a000414-a000414.
DOI:10.1101/cshperspect.a000414

26. Parsons JB, Yao J, Frank MW, Jackson P, Rock CO. Membrane disruption by antimicrobial fatty acids releases low-molecular-weight proteins from Staphylococcus aureus. J Bacteriol 2012;194(19):5294-5304.
DOI:10.1128/jb.00743-12

27. Truong-Bolduc QC, Dunman PM, Strahilevitz J, Projan SJ, Hooper DC. MgrA is a multiple regulator of two new efflux pumps in Staphylococcus aureus. J Bacteriol. 2005;187(7):2395-2405.
https://doi.org/10.1128/JB.187.7.2395-2405.2005