Novel 4-substituted phenyl-2,2′-bichalcophenes and aza-analogs as antibacterial agents: a structural activity relationship

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Abstract: Antibiotic resistance is a major health problem; therefore, new antibacterial agents will need to be continuously developed. A series of novel bichalcophenes has been tested and found to have antimicrobial activity against selected bacteria. Due to the promising antimicrobial effects of these 4-substituted phenyl bichalcophene derivatives, the study reported here was launched to examine the interaction between novel bichalcophenes and tetracycline. The minimum inhibitory concentration values for all bichalcophenes were between 8 and 64 µM. Many of the bichalcophenes had synergistic activity that increased the inhibitory effect of tetracycline against bacterial growth, as indicated by the fractional inhibitory concentration index. The post-antibiotic effects of the novel bichalcophenes were determined. Many bichalcophenes were able to elongate the period required for bacteria to recover and grow after a brief exposure to tetracycline. Escherichia coli did not develop resistance to many bichalcophenes over a period of 7 days. A structural activity relationship could be characterized, as monocationic derivatives were more active than the corresponding mononitriles. The presence of a pyridyl group and/or furan ring reduced the activity, while the presence of a phenyl or thiophene ring enhanced the antibacterial activity. Our results suggest that bichalcophenes could be useful to elevate the shelf life of many antibiotics.

Keywords: synergic interaction, fractional inhibitory concentration, post-antibiotic effect, resistant variants

Introduction

Since the discovery of penicillin, many antibiotics have been developed to control bacterial infections.1 The use of antibacterial agents along with improved sanitary conditions suggested that industrialized nations had won the war against pathogenic microbes. However, over the past few years, bacterial resistance to antibiotics has developed and resistance to multiple antimicrobial agents has become a major health problem.2 Many factors affect the rate of development of resistance, including excessive use and misuse of antibiotics.3 Due to the development of antibiotic resistance in virtually all clinically important pathogens, new antibacterial agents will need to be continuously developed.

Aromatic dicaticonic compounds, such as pentamidine, furamidine, and their analogs, have been reported to have broad-spectrum antimicrobial effects against fungi and bacteria.4,5 The antimicrobial activity of pentamidine has been recognized for over 50 years and pentamidine remains the only aromatic diamidine that has been significantly used in the drug industry.6 However, due to the poor oral bioavailability and unfavorable side-effects of many diamidines, numerous new compounds have
been synthesized aimed at improving these deficiencies.\textsuperscript{7} Thiophene-, furan-, and nitrile-containing compounds have been synthesized and shown to exhibit a broad-spectrum antimicrobial activity.\textsuperscript{8–12} Recently, a series of type I dicatonic bichalcophenes (Figure 1) and aza-analogs of furamidine have been reported to be antiprotozoal agents.\textsuperscript{11,13}

A series of type II 4-substituted phenyl bichalcophenes (Figure 1) have been synthesized and tested for their antimicrobial activity against the growth of several bacteria.\textsuperscript{14} Due to the promising antimicrobial effects of these 4-substituted phenyl bichalcophene derivatives, the study reported here was launched to examine the interaction between type II novel bichalcophenes and tetracycline by calculating the fractional inhibitory concentration (FIC) of each combination against selected bacteria. Moreover, the effect of the bichalcophenes on the suppression of bacterial growth that persists after a brief exposure of \textit{Escherichia coli} to tetracycline (known as “post-antibiotic effect” [PAE]) was determined. The increased PAE in the presence of bichalcophenes could offer an alternative way for extension of the useful lifespan.

![Figure 1](image-url)
of antibiotics.\textsuperscript{15} The rationale behind the selection of tetracycline was to test the potency of novel bichalcophenes to enhance the efficacy of an antibiotic that has lost efficiency and against which many bacteria have acquired resistance. In addition, to investigate the stability of the tested compounds as antimicrobial agents, the development of spontaneous resistance to bichalcophenes was investigated.

**Materials and methods**

**Chemicals**

All reagents were purchased from Sigma-Aldrich (St Louis, MO, USA) except where indicated in the specified methods. Ten bichalcophene derivatives were used throughout the study (Figure 1). Bichalcophenes were sterilized by filtration through a 0.2 µm filter (Nalgene, Rochester, NY, USA).

**Bacterial strains**

The tested bacterial strains were purchased from the American Type Culture Collection (ATCC; Manassas, VA, USA). The antibacterial assays were performed against standard strains of Gram-positive (Staphylococcus aureus ATCC 25923 and Bacillus megaterium ATCC 14591) and Gram-negative (E. coli ATCC 25922, Pseudomonas aeruginosa ATCC 27853, and Klebsiella pneumoniae ATCC 700603) bacteria. All procedures were approved by the Committee of Scientific Research Ethics of King Faisal University.

**Minimum inhibitory concentration (MIC) of the novel bichalcophene derivatives**

The MIC of the novel bichalcophenes was determined according to the Clinical and Laboratory Standards Institute guidelines.\textsuperscript{16} An inoculum density of $5 \times 10^5$ colony forming units (CFU)/mL of each organism was dispensed into the wells of a 96-well microtiter plate. The bichalcophenes were serially (1–128 µM) diluted in the wells followed by the addition of bacteria. The plates were incubated at 37°C for 18 hours and the absorbance was recorded at 600 nm. The MIC value was recorded as the lowest concentration at which no growth was observed. All MIC values were determined on three independent experiments.

**Synergic interaction between the tested bichalcophenes and tetracycline**

A combination of each bichalcophene at 0.25 or 0.5 × MIC and tetracycline (0.25–128 µg/mL) was prepared and inoculated with $5 \times 10^5$ CFU/mL of each strain. The plates were incubated at 37°C for 18 hours, and the lowest antibiotic concentration in combination with bichalcophene that prevented the development of growth was regarded as the MIC*. The FIC was calculated as follows:\textsuperscript{17} FIC of tetracycline = MIC* (combination of tetracycline + bichalcophene)/MIC of tetracycline alone. An FIC ≤ 0.5 indicated synergy, an FIC > 0.5–4.0 indicated indifference, and an FIC > 4 indicated antagonism.

**PAE**

The *E. coli* inocula equivalent to $10^6$ CFU/mL were incubated at 37°C for 30 minutes with tetracycline (1 × MIC), each bichalcophene (0.25 × MIC), or a combination of each bichalcophene and tetracycline (0.25 × MIC + 1 × MIC, respectively). After 30 minutes, bacteria were diluted 1000-fold with Mueller-Hinton broth (MHB) to eradicate the antibiotic and/or bichalcophene. A 100 µL sample of each treatment was obtained at time zero and every hour thereafter until turbidity developed, and spread onto nutrient agar for total viable cell count after incubation at 37°C for 24 hours. The PAE was calculated as described by Craig and Gudmundsson:\textsuperscript{18} PAE = T − C, in which T is the time required for the viable cell count of the exposed bacteria to increase by a factor of $10_\text{m}$, from the initial time immediately after washing, and C is the corresponding time for cells unexposed to either antibiotic or bichalcophenes. Each assay was carried out in triplicate. To avoid overgrowth of the control culture, a 100-fold dilution was prepared using MHB at every time interval.

**Detection of bichalcophenes-resistant variants**

To evaluate the development of spontaneous resistance in *E. coli* against bichalcophenes or tetracycline, growth assays were performed using *E. coli* in the presence or absence of each bichalcophene or tetracycline. Twenty milliliters of cells in the logarithmic growth phase were centrifuged, and the cells were resuspended in 2 mL of MHB. The inoculum density was adjusted to $10^6$ CFU/mL. Different concentrations of each bichalcophene (3, 2, 1 × MIC) were added. All plates were incubated at 37°C and the absorbance was recorded at 600 nm daily for 7 days.\textsuperscript{19}

**Statistical analysis**

Statistical analyses were performed using analysis of variance followed by Fisher’s protected least significant difference multiple-range test. Differences were considered significant at $P$ values < 0.05.

**Results**

The bichalcophenes had broad-spectrum activities against the tested bacteria. Compound 5B was the most effective
against all bacteria, with an MIC value at 8 µM (2.7 µg/mL). Compounds 1A and 2A were alike in inhibiting the growth of bacteria, with an MIC value at 32 µM. Gram-positive bacteria were more sensitive to Compounds 3A, 4A, and 5A (MIC 32 µM) than Gram-negative strains (MIC 64 µM). Similarly, Compounds 3B and 4B were most effective in inhibiting the growth of Gram-positive bacteria (MIC 16 µM), while the MIC value against Gram-negative bacteria was 32 µM. Compounds 1B and 2B showed the same pattern, with MIC values at 16 µM against *S. aureus*, *B. megaterium*, and *E. coli* and 32 µM against *P. aeruginosa* and *K. pneumoniae* (Table 1).

The interaction of each bichalcophene at a concentration of 0.25 or 0.5 × MIC with various concentrations of tetracycline (0.25–128 µg/mL) was examined (Table 2). The reported MIC values for tetracycline were 4 µg/mL against *B. megaterium*, 8 µg/mL against *S. aureus*, *E. coli*, and *P. aeruginosa*, and 16 µg/mL against *K. pneumoniae* (Table 1). With the exception of Compound 2A, all bichalcophenes at 0.25 × MIC had a synergic effect on the activity of tetracycline against *K. pneumoniae* and antagonized the activity of tetracycline against the growth of all other tested bacteria (FIC > 4). With the exception of Compounds 2A and 3A against *S. aureus* and *P. aeruginosa* and 4A and 4B against *P. aeruginosa*, all bichalcophenes at 0.5 × MIC acted synergistically with tetracycline against the growth of all bacterial strains under investigation (Table 2). Compounds 1A, 1B, and 3B at 0.25 × MIC were similar in potentiating the activity of tetracycline against all tested bacteria except *S. aureus* (FIC = 2). Indifference (FIC > 0.5–4.0) was shown with Compound 5A at the lowest concentration against *S. aureus* and *B. megaterium*, while it increased the activity of tetracycline against Gram-negative bacteria. Compounds 2B and 5B at both concentrations investigated were the most effective in inhibiting the growth of all tested bacteria when used in combination with tetracycline, with both achieving additive effects, as indicated by the FIC values (0.063–0.500). The effect of Compound 3A on the activity of tetracycline varied with the tested bacteria, with indifferent effects reported with *S. aureus* (at both concentrations) and *E. coli* (only at low concentration), and an antagonistic activity against the growth of *P. aeruginosa* at both concentrations and a synergic effect was shown against *B. megaterium* and *K. pneumoniae*. Synergic effect was reported with Compound 4B against the growth of *B. megaterium*, *E. coli*, and *K. pneumoniae*, and an antagonistic effect against *P. aeruginosa*. Compound 4A did not affect the tetracycline activity against *B. megaterium*, *E. coli*, or *S. aureus* only at 0.25 × MIC and antagonized the tetracycline activity when used against *P. aeruginosa* at both 0.25 and 0.5 × MIC (Table 2).

The estimated PAE for tetracycline was 2 hours. Using bichalcophenes 1A–4A and 4B resulted in PAEs of 2 hours. The recorded PAE for 5A, 1B, and 2B was 3 hours. Compounds 3B and 5B were more effective in suppressing *E. coli* growth for 5 and 6 hours, respectively. The combination of Compound 1A or 5A with tetracycline resulted in a PAE of 3 hours. The combination of 3A, 2B, or 4B with tetracycline had similar effects, elevating the PAE value to 4 hours. Compounds 1B, 3B, and 5B seemed the most effective at extending the suppression time of bacterial growth, with PAEs of 6, 7, and 7 hours, respectively. However, Compounds 2A and 4A had no effect on PAE value as compared to unexposed control cells (Figure 2).

On investigation of the development of resistant variants, a high frequency of resistance development was detected with tetracycline after 2 days of incubation at 1 × MIC (8 µg/mL).

### Table 1 Minimum inhibitory concentration (MIC) of the bichalcophene derivatives against different bacterial strains

<table>
<thead>
<tr>
<th>Tested compound µM (µg/mL)</th>
<th>MIC* Staphylococcus aureus</th>
<th>Bacillus megaterium</th>
<th>Escherichia coli</th>
<th>Pseudomonas aeruginosa</th>
<th>Klebsiella pneumoniae</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tetracycline (mg/mL)</td>
<td>8</td>
<td>4</td>
<td>8</td>
<td>8</td>
<td>16</td>
</tr>
<tr>
<td>1A</td>
<td>32 (7.5)</td>
<td>32 (7.5)</td>
<td>32 (7.5)</td>
<td>32 (7.5)</td>
<td>32 (7.5)</td>
</tr>
<tr>
<td>1B</td>
<td>16 (4.6)</td>
<td>16 (4.6)</td>
<td>16 (4.6)</td>
<td>32 (9.2)</td>
<td>32 (9.2)</td>
</tr>
<tr>
<td>2A</td>
<td>32 (7.6)</td>
<td>32 (7.6)</td>
<td>32 (7.6)</td>
<td>32 (7.6)</td>
<td>32 (7.6)</td>
</tr>
<tr>
<td>2B</td>
<td>16 (5.4)</td>
<td>16 (5.4)</td>
<td>16 (5.4)</td>
<td>32 (10.7)</td>
<td>32 (10.7)</td>
</tr>
<tr>
<td>3A</td>
<td>32 (8.0)</td>
<td>32 (8.0)</td>
<td>64 (16.0)</td>
<td>64 (16.0)</td>
<td>64 (16.0)</td>
</tr>
<tr>
<td>3B</td>
<td>16 (4.9)</td>
<td>16 (4.9)</td>
<td>32 (9.9)</td>
<td>32 (9.9)</td>
<td>32 (9.9)</td>
</tr>
<tr>
<td>4A</td>
<td>32 (8.1)</td>
<td>32 (8.1)</td>
<td>64 (16.1)</td>
<td>64 (16.1)</td>
<td>64 (16.1)</td>
</tr>
<tr>
<td>4B</td>
<td>16 (5.5)</td>
<td>16 (5.5)</td>
<td>32 (11.0)</td>
<td>32 (11.0)</td>
<td>32 (11.0)</td>
</tr>
<tr>
<td>5A</td>
<td>32 (8.6)</td>
<td>32 (8.6)</td>
<td>64 (17.1)</td>
<td>64 (17.1)</td>
<td>64 (17.1)</td>
</tr>
<tr>
<td>5B</td>
<td>8 (2.7)</td>
<td>8 (2.7)</td>
<td>8 (2.7)</td>
<td>8 (2.7)</td>
<td>8 (2.7)</td>
</tr>
</tbody>
</table>

**Note:** Data are expressed as the means of three independent experiments.
Table 2 Synergic interaction between the bichalcophene derivatives and tetracycline against the tested bacterial strains

<table>
<thead>
<tr>
<th>Compound</th>
<th>Tested bacterial strains</th>
<th>Staphylococcus aureus</th>
<th>Bacillus megaterium</th>
<th>Escherichia coli</th>
<th>Pseudomonas aeruginosa</th>
<th>Klebsiella pneumoniae</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>0.25 × MIC</td>
<td>0.5 × MIC</td>
<td>0.25 × MIC</td>
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<td>0.5 × MIC</td>
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<tr>
<td></td>
<td>MIC*</td>
<td>FIC</td>
<td>MIC*</td>
<td>FIC</td>
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<td>FIC</td>
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<tr>
<td>1A</td>
<td>16</td>
<td>2</td>
<td>4</td>
<td>0.5</td>
<td>4</td>
<td>0.5</td>
</tr>
<tr>
<td>1B</td>
<td>16</td>
<td>2</td>
<td>4</td>
<td>0.5</td>
<td>4</td>
<td>0.5</td>
</tr>
<tr>
<td>2A</td>
<td>64</td>
<td>8</td>
<td>8</td>
<td>1</td>
<td>128</td>
<td>32</td>
</tr>
<tr>
<td>2B</td>
<td>2</td>
<td>0.25</td>
<td>2</td>
<td>0.25</td>
<td>4</td>
<td>0.5</td>
</tr>
<tr>
<td>3A</td>
<td>32</td>
<td>4</td>
<td>8</td>
<td>1</td>
<td>1</td>
<td>0.25</td>
</tr>
<tr>
<td>3B</td>
<td>16</td>
<td>2</td>
<td>1</td>
<td>0.125</td>
<td>1</td>
<td>0.25</td>
</tr>
<tr>
<td>4A</td>
<td>16</td>
<td>2</td>
<td>2</td>
<td>0.25</td>
<td>16</td>
<td>4</td>
</tr>
<tr>
<td>4B</td>
<td>16</td>
<td>2</td>
<td>2</td>
<td>0.25</td>
<td>16</td>
<td>4</td>
</tr>
<tr>
<td>5A</td>
<td>16</td>
<td>2</td>
<td>1</td>
<td>0.125</td>
<td>16</td>
<td>4</td>
</tr>
<tr>
<td>5B</td>
<td>1</td>
<td>0.125</td>
<td>1</td>
<td>0.125</td>
<td>1</td>
<td>0.25</td>
</tr>
</tbody>
</table>

Notes: MIC* = MIC of tetracycline in combination with each bichalcophene (0.25 or 0.5 × MIC). FIC = MIC* of tetracycline + tested bichalcophene/MIC of tetracycline alone. FIC = 0.5 indicates synergy, FIC > 0.5–4.0 means indifference, and FIC > 4 indicates antagonism. Data are expressed as the means of two independent experiments.

Abbreviations: FIC, fractional inhibitory concentration; MIC, minimum inhibitory concentration.

Discussion

The emergence of antibiotic resistance in bacteria is a serious problem facing humanity. The rate at which this resistance develops exceeds the rate of antibiotic development and production. Many antibiotics are now ineffective and the efficiency of many others is declining. Therefore, there must be a continuous and extensive effort to develop novel antibiotics that have lower toxicity and are more effective than their corresponding monocationic derivatives. The MIC values of monocationic derivatives used in this study were between 8 and 32 µM, while those of mononitriles were between 32 and 64 µM. The reported antimicrobial activity of the bichalcophenes can be attributed to the introduction of cyanogroups in the monocationic derivatives. The MIC values of monocationic derivatives used in this study were between 8 and 32 µM, while those of mononitriles were between 32 and 64 µM. The reported antimicrobial activity of the bichalcophenes can be attributed to the introduction of cyanogroups in the monocationic derivatives.
Figure 2. The post-antibiotic effect of bichalcophenes (0.25 × MIC), tetracycline (1 × MIC), or a combination of bichalcophenes and tetracycline on *Escherichia coli* American Type Culture Collection 25922. The control curve represents the growth of bacteria unexposed to either bichalcophenes or tetracycline. (A) Compounds 1A and 1B; (B) Compounds 2A and 2B; (C) Compounds 3A and 3B; (D) Compounds 4A and 4B; (E) Compounds 5A and 5B.

Notes: Data are expressed as the mean of three independent assays. Standard error of the mean has been omitted for clarity.

Abbreviations: CFU, colony forming unit; cpd, compound; MIC, minimum inhibitory concentration.

In a dose-dependent manner, all bichalcophenes under investigation, except Compounds 2A, 3A, and 4A, had synergic interactions with tetracycline against almost all strains 1B and 2B; 3B and 4B; and 3A, 4A, and 5A. In killing the bacteria, the bichalcophenes behaved like many broad-spectrum antibiotics and were equally effective against both Gram-negative and Gram-positive bacteria; their mechanism may be through inhibition of the synthesis of nucleic acids and/or protein. In a previous study, bichalcophenes were found to cause the degradation of bacterial DNA, but further studies are required to determine the mechanism(s) of action through which these antibacterial agents may act.
tested except *S. aureus* (Table 2). Compounds 2B and 5B were the most effective and significantly elevated the tetracycline activity against all organisms, while 2A, 3A, and 4A were the least effective. These compounds are all mononitriles and have at least one furan ring and two of them have pyridyl rings. Compound 2A (mononitrile bifuran with pyridyl ring) antagonized the actions of tetracycline against almost all bacterial strains investigated. Similar to our MIC findings, the monocationic bichalcophenes (2B, 3B, 4B, and 5B) were more effective than the corresponding mononitrile derivatives (2A, 3A, 4A, and 5A). This differentiation in action was evident in *P. aeruginosa*, where 2A, 3A, and 4A antagonized the inhibitory actions of tetracycline (Table 2). Taken together, the bichalcophenes used in this study can be arranged in descending order according to their ability to potentiate tetracycline activity against bacteria: 2B and 5B > 3B, 4B, and 1A > 4B and 5A > 3A > 4A > 2A. From this, it is clear that the monocationic derivatives were the most effective.

The in vitro pharmacodynamic properties of bichalcophenes were investigated by studying the PAE against *E. coli.*

**Figure 3** Development of spontaneous resistant variants of *Escherichia coli* toward bichalcophenes or tetracycline (3, 2, and 1 × MIC) over 7 days. (A) Compound 1A; (B) Compound 1B; (C) Compound 2A; (D) Compound 2B; (E) Compound 3A; (F) Compound 3B; (G) Compound 4A; (H) Compound 4B; (I) Compound 5A; (J) Compound 5B; (K) tetracycline.

**Notes:** Data are expressed as the mean of three independent assays. Standard error of the mean has been omitted for clarity.

**Abbreviation:** MIC, minimum inhibitory concentration.
Any agent that induces a long antibiotic PAE may secure longer dosing intervals without losing the antibiotic’s efficacy.26 E. coli exposed to either tetracycline or most bichalcophenes was able to recover and grow 2–3 hours post-exposure. The low value obtained in the present study after exposure to tetracycline has been previously reported.27 Two compounds (3B and 5B) resulted in a remarkable delay in bacterial growth (PAE) of up to 5–6 hours after exposure; not surprisingly, these compounds were monocationic phenyl bichalcophenes. With the exception of 3A, all mononitrile bichalcophene (phenyl or pyridyl) derivatives when combined with tetracycline had no significant effect on the PAE. Combining monocationic phenyl bichalcophenes (1B, 3B, and 5B) with tetracycline resulted in a significant 6–7-hour delay in bacterial growth. The presence of a pyridyl ring in Compounds 2B and 4B reduced the ability of these monocationic derivatives, resulting in a 4-hour delay in PAE (Figure 2).

In the search for new antibacterial agents, it is not enough to prove their efficacy; it is also necessary to investigate the rate of resistance development toward these new agents. In our study, after only 1 day of incubation, E. coli developed resistance to all mononitrile derivatives (1A–5A), monocationic pyridyl bichalcophenes (2B and 4B), and tetracycline at the low concentration of 1 × MIC. Similar findings have previously been reported for tetracycline.28 At the highest concentration of 3 × MIC, the bacteria were not able to develop resistance against almost all bichalcophenes after 7 days of incubation, but overcame tetracycline and started to grow after 5 days. In contrast, the bacteria failed to develop resistance toward the monocationic phenyl bichalcophenes (1B, 3B, and 5B), even after 7 days, at all concentrations investigated (Figure 3).

In conclusion, in our study, a structural activity relationship was evident, as the presence of mononitrile or pyridyl groups and, to a lesser extent, the furan ring, reduced antibacterial activity, while the presence of a phenyl, monocationic, or thiophene ring clearly enhanced antibacterial activity. Recently, the National Cancer Institute Developmental Therapeutics Program examined these bichalcophenes for anticancer activity in 58 cell lines and found that monocationic bichalcophenes were more active than the corresponding aza-analogs and that 1B, 4B, and 5B were the most active compounds (unpublished data). These bichalcophenes not only have antibacterial properties but can also potentiate the activity of antibiotics currently in use. As it is unlikely that bacteria could develop resistance to these novel compounds, their possible clinical effectiveness is suggested. Two of these bichalcophenes (1A and 1B) have been tested in mice and found to be not toxic.29 Treatment of methicillin-resistant Staphylococcus aureus-infected mice with these bichalcophenes resulted in significant reductions in the viable bacterial count in blood, liver, kidney, and spleen. The structural activity relationship in the present study was also confirmed in vivo, as Compound 1B was superior to 1A and sometimes both provided better antistaphylococcal agents than vancomycin against methicillin-resistant S. aureus pathogenesis.29


