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Honokiol-Inspired Analogs as Inhibitors of Oral Bacteria

Amy E. Solinski[†], Cristian Ochoa[‡], Young Eun Lee[‡], Thomas Paniak[‡], Marisa C. Kozlowski^{*‡,iD}, and William M. Wuest^{*‡,iD}

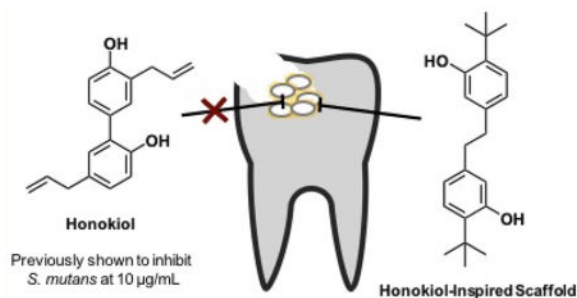
[†]Department of Chemistry, 1515 Dickey Drive, Emory University, Atlanta, Georgia 30322, United States

[‡]Department of Chemistry, Roy and Diana Vagelos Laboratories, University of Pennsylvania, Philadelphia, Pennsylvania 19104, United States

Abstract

The oral microbiome is a complex ecological niche where both commensal and pathogenic bacteria coexist. Previous reports have cited that the plant isolate honokiol is a potent inhibitor of *S. mutans* biofilms. Herein we report a cross-coupling method that provides access to a concise library of honokiol-inspired analogs. Through this work we determined that the inhibitory activity of honokiol is highly dependent on the growth conditions. Further, we identify a series of analogs that display significant potency against oral bacteria leading to the discovery of a potent antimicrobial.

Graphical abstract



*Corresponding Authors. marisa@sas.upenn.edu. wwuest@emory.edu.

ORCID

Marisa C. Kozlowski: 0000-0002-4225-7125

William M. Wuest: 0000-0002-5198-7744

A.E.S., C.O., and Y.E.L. contributed equally.

ASSOCIATED CONTENT

Supporting Information

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Synthetic characterization and biological assay procedures (PDF)

The manuscript was written through contributions of all authors. All authors have given approval to the final version of the manuscript. The authors declare the following competing financial interest(s): W.M.W. and M.C.K. have filed a patent on the technology disclosed.

Keywords

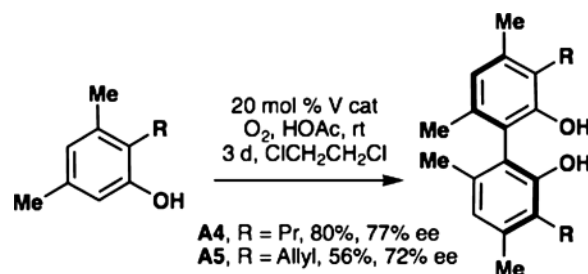
oxidative coupling; antibiotic; oral microbiome; natural product

In the past decade studies have correlated that healthy human microbiomes play a role in combating “modern plagues” such as diabetes, obesity, Crohn’s, and celiac disease.¹ The oral microbiome, a specific microbial niche residing in the oral cavity of humans, has gained attention due to the complexity in which the microorganisms interact both in commensal and pathogenic manners. This ecosystem is home to various species of bacteria that have adapted to live in the microaerophilic and/or anaerobic environments that exist surrounding the teeth, gingiva, and tongue.^{2,3} Scientists have turned to chemical biology to develop new tools to study these systems and understand them at the molecular level. Interest in *S. mutans* has grown since it has been highly associated with the formation of carries via environmental acidification which leads to the corrosion of tooth enamel (Figure 1A).^{4,5} Early colonizers such as commensal *Streptococcus sanguinis* and *Streptococcus gordonii* are responsible for early plaque formation by anchoring adhesion proteins to the pellicle of the tooth and producing glucan polymers that constitute the matrix of dental plaque. *S. mutans* is able to invade this matrix, form microcolonies, and eventually develop into a mature biofilm that is responsible for tooth decay via acidification.^{6,7} Another lesser known and more harrowing disease that has been associated with *S. mutans* biofilm growth is infective endocarditis, or inflammation of the inner tissues of the heart.⁸ *S. mutans* has the capability to nest itself in the heart as a mature biofilm and block the blood supply to the inner heart tissues causing inflammation. To date, few natural products have been reported to be effective inhibitors of the oral pathogen *S. mutans*. One such example is the natural product carolacton which has attracted the attention of our group as well as the Kirshning and Wagner-Döbler laboratories.^{9–11} Carolacton specifically targets *S. mutans* cells as they transition into a biofilm. In contrast, the phenolic natural product honokiol has received attention due to the reportedly potent inhibitory activity against *S. mutans* (Figure 1B).^{12,13} Although isolated and first reported in 1982 from the bark or seeds of a magnolia tree, honokiol has been used as a therapeutic in Chinese, Japanese, and Korean traditional herbal remedies for centuries.^{14,15} Previously, our group developed a concise synthesis of honokiol via oxidative phenolic coupling.¹⁶ In this report we leverage this method to develop a focused library of honokiol-inspired analogs to better understand the structure–activity relationship against oral bacteria.

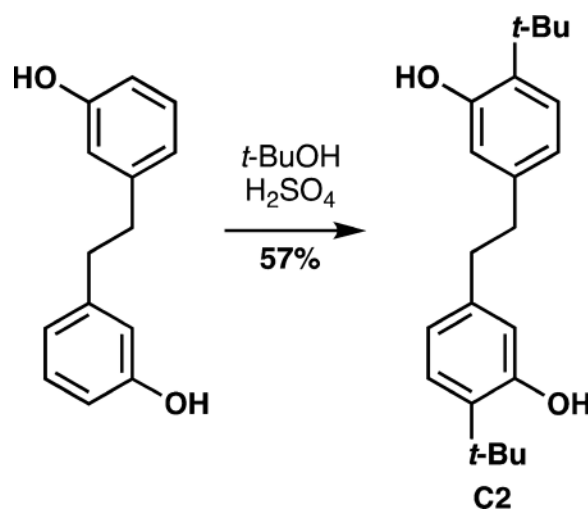
Our group has developed an expedited method to access this natural product scaffold.¹⁷ Accordingly, we sought to apply this method in a general sense for two reasons: 1) to demonstrate the scope of this method for uniting aryl moieties and 2) to provide a library of analogs to answer specific structure–activity relationship questions. The analog design was organized into three groups based on the scaffold (Figure 2). Group A mimics the biaryl architecture of the natural product honokiol, Group B focuses on the naphthalene scaffold, and Group C examines the necessity of the biaryl linkage.

As mentioned previously, the oxidative coupling reactions developed in our lab were used to synthesize the specific congeners of the general subclasses outlined in Figure 2. A vanadium-catalyzed phenol homocoupling was used to assemble **A4** and **A5** (eq 1).¹⁸

Selective cross-coupling of two different phenols was accomplished with a chromium catalyst developed previously.¹⁶ Table 1 illustrates how the technique was used to rapidly assemble an array to investigate structure–activity relationships; in these cases no optimization of the yields was performed as the bioactivity was the focus. To investigate an alternate biaryl union, **C2** was prepared by Friedel–Crafts alkylation of the parent bisphenol (eq 2). The analogs described in Table 1 are all congeners of the parent structures in Figure 2.



(1)



(2)

At the beginning of our investigation we were interested in comparing the inhibitory activity of honokiol (**1A**) to that of our newly synthesized analogs. Minimum inhibitory concentration (MIC) assays, minimum biofilm inhibitory concentration (MBIC) assays, and minimum bactericidal concentration (MBC) assays were undertaken. We initially performed the MIC assays in a 5% CO₂-supplemented environment to promote growth of *S. mutans* in an environment that most closely mimics a healthy oral cavity. The MIC of honokiol was determined to be 250 μM (66.6 μg/mL), which was in stark contrast to the literature value of 10 μg/mL (Table 2). After revisiting the original procedures, we recognized that the original assays were completed in an aerobic environment, which precludes the growth of *S. mutans*.¹² As expected, when aerobic conditions were employed in the assay, the potency of

honokiol increased to 125 μM (33.3 $\mu\text{g/mL}$). These results demonstrate that although *S. mutans* growth is inhibited by honokiol, the overall efficacy of the compound will be less under physiological conditions. Furthermore, our studies show that honokiol is unable to inhibit biofilm growth (Figure S3) and was not bactericidal at concentrations of 250 μM or lower.

Undeterred by these findings, we sought to evaluate the bioactivity of our honokiol-inspired analogs against a panel of representative oral bacteria via MIC, MBIC, and MBC assays (Table 2 and Figure S3). Out of the 26 honokiol analogs (see Figures S1 and S2 for all structures) four compounds showed significant inhibition at low concentrations ($16 \mu\text{M}$). Analogs C2, B5, B8, and B11 were the most impressive with MIC values of 2 μM , 8 μM , 16 μM , and 16 μM , respectively, against planktonic *S. mutans*. The analogs were also tested against two commensal strains that are early colonizers: *S. gordonii* and *S. sanguinis*. Generally, the MICs for these commensal strains mirrored the values for the pathogenic *S. mutans* hinting at a broad-spectrum inhibitory mechanism.

Biofilms are of particular importance to the oral cavity as they protect the bacteria and allow them to outcompete other colonizers by decreasing the local pH, thereby causing enamel erosion and other pathologies resulting from the acidified environment.¹⁹ For these reasons, the analogs were tested for their ability to inhibit the formation of *S. mutans* biofilm (Figure S3). Initial reports identified honokiol as a biofilm inhibitor; however in our hands, the MBIC values were within one dilution of the MIC. Based on these results, it is likely that biofilm inhibition is an effect of the inherent toxicity of the compounds to the planktonic bacteria and not by a biofilmspecific mechanism. Honokiol and the analogs tested herein all potently deterred the formation of biofilms when the cells were grown in the presence of sucrose, albeit at the previously determined MIC values. This likely indicates that the compounds are targeting the bacteria in a general fashion and do not show any preferential killing to biofilms.

We next sought to determine if the compounds were working in a bacteriostatic or bactericidal manner. Toward this end, a regrow analysis was completed to determine the MBC values of the active analogs against *S. mutans* (Figure S4). The MBC values reported refer to the concentration at which there is a 3-Log reduction in CFU count corresponding to 99.9% bacterial cell death (Table 2). Analog **C2** had the lowest MBC value at 4 μM confirming that the molecule is bactericidal.²⁰ Analogs **B8** and **B11** were also shown to be bactericidal; however, the MIC and MBC of analog **B5** differs by four dilutions hinting at a bacteriostatic mechanism. These findings suggest that compounds **C2** and **B5** may be inhibiting the growth of *S. mutans* by different mechanisms.

Research has focused mainly on the antitumor, antifungal, and anti-inflammatory activities of honokiol resulting in the identification of various cellular targets.^{21–24} In contrast, the mechanism of action of honokiol, or the derivatives reported, in these oral pathogens has been elusive. Therefore, future work will focus on determining the mechanism by which these molecules elicit their response.

Our initial interest in the natural product honokiol was 2-fold: 1) to probe the bioactivity profile of the compound and 2) as a means to showcase our newly developed synthetic method to access biaryl scaffolds. By extending the aryl connectivity with a two-carbon linker, the potency of the lead compound increased from 250 μM to 2 μM . Intriguingly, the phytochemical dihydrostillbene Batatasin-III, which possesses an analogous scaffold has been studied for its antifouling abilities and has ties to ancient Chinese herbal remedies along with honokiol (Figure 3).²⁵ Based on these chemical similarities, it is possible that the extended scaffold has unrealized biological activity against cariogenic agents and warrants further study. Future work will include a more expansive analog library that will address these hypotheses and will be reported in due course.

In conclusion, the work presented herein highlights the importance of performing biological testing at physiologically relevant conditions. Our results have demonstrated that the bioactivity of honokiol may be overstated. However, our curiosity surrounding the natural product led to the serendipitous discovery of a highly potent, bactericidal analog, **C2**, with an MIC value of 2 μM (66 ng/mL). This compound serves as an exciting starting point for future translational studies which may be of particular interest to the oral care industry based on its simple structural architecture and potent bioactivity.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

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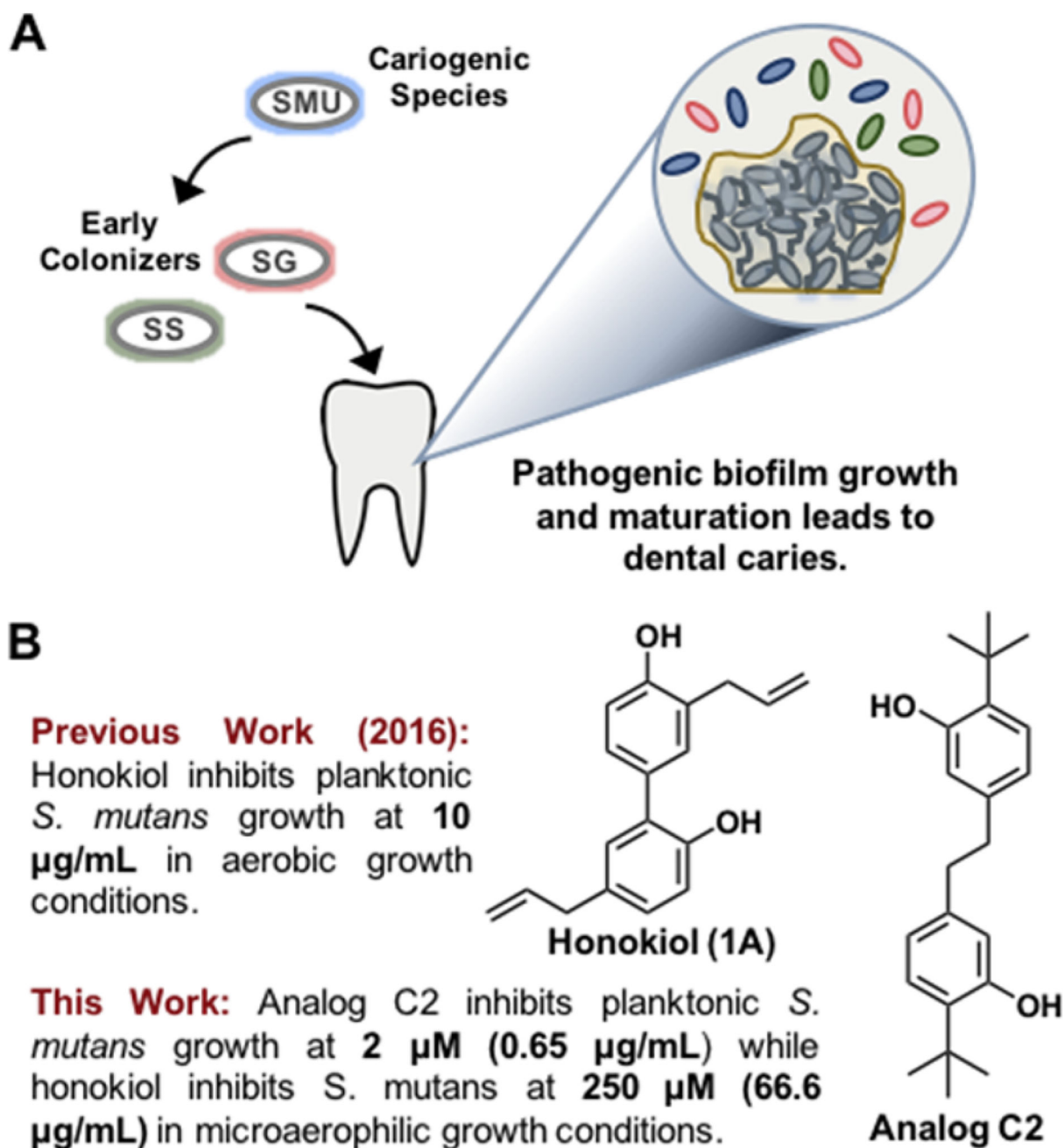


Figure 1.

A) Early colonizers *S. sanguinis* and *S. gordonii* allow cariogenic *S. mutans* to form biofilms on the surface of the tooth by adhering to the pellicle. **B)** The natural product honokiol has been previously reported to inhibit *S. mutans* growth. Here we demonstrate that analog C2 is a more potent bactericidal agent against oral microbiome bacteria.

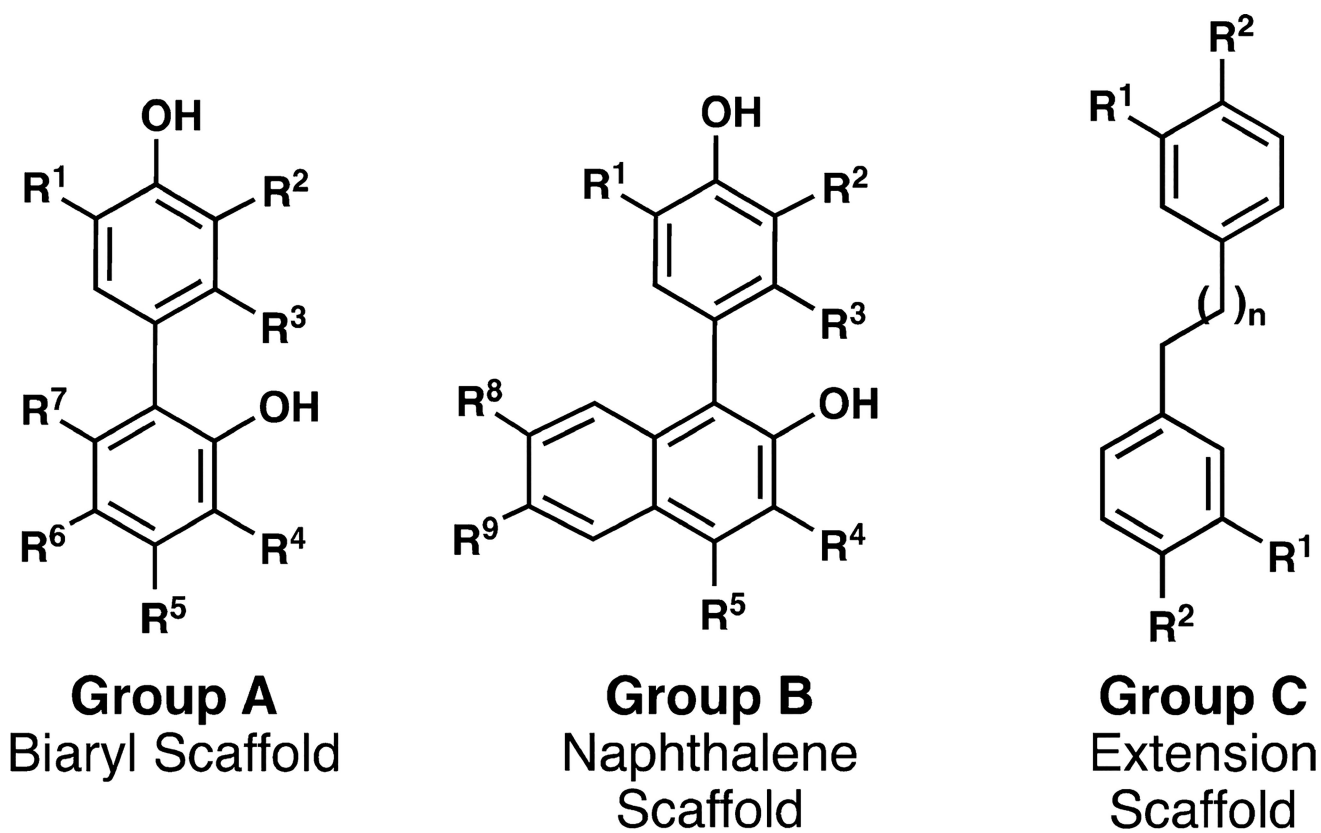


Figure 2.

Analogs are classified in three groups. Group A = biaryl scaffold; Group B = naphthalene scaffold; Group C = extension scaffold.

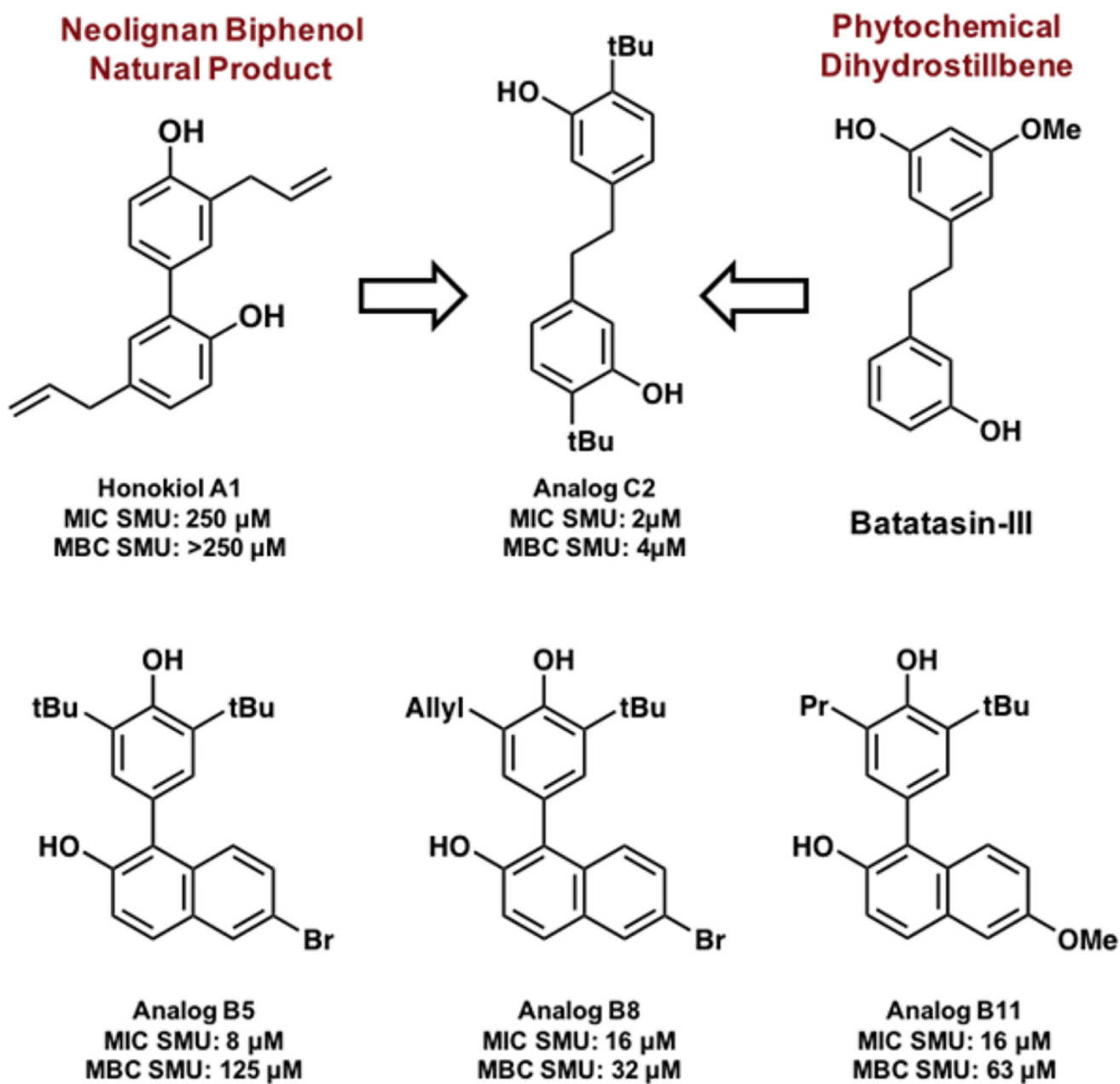
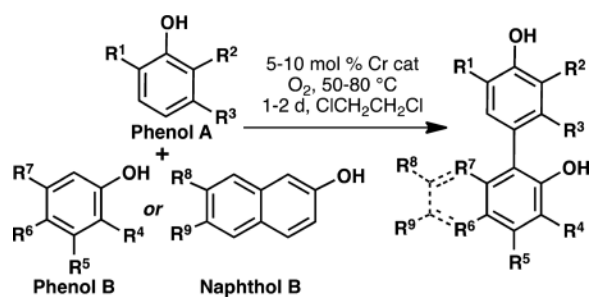


Figure 3.
Summary of biologically active honokiol-inspired analogs. Comparison of the structure of C2 and the natural product batatasin- III.

Table 1

Cr-Salen Catalyzed Cross-Couplings



product	phenol A R ¹ , R ² , R ³	phenol B/naphthol B R ⁴ , R ⁵ , R ⁶ , R ⁷ / R ⁸ , R ⁹	yield (%)
A2	<i>t</i> -Bu, <i>t</i> -Bu, H	Me, HO, H, Me	38
A3	Me, Me, H		63 ^{a,b,c}
A6	allyl, <i>t</i> -Bu, OH	H, Me, Me, Me	33
A7	<i>t</i> -Bu, Me, H	Me, H, H, <i>i</i> -Pr	74 ^{a,b}
A8	allyl, <i>t</i> -Bu, OH	H, H, Me, H	28
A9	<i>i</i> -Pr, <i>i</i> -Pr, H	H, Me, Me, Me	18
A10	allyl, allyl, H	Me, H, H, <i>i</i> -Pr	4 ^b
A11	<i>t</i> -Bu, <i>t</i> -Bu, H	Me, OH, H, Me	29 ^d
B1	Me, Me, H	H, H	65
B2	MeO, MeO, H	H, H	72
B3	<i>t</i> -Bu, <i>t</i> -Bu, H	H, H	83 ^a
B4	<i>t</i> -Bu, <i>t</i> -Bu, H	HO, H	88
B5	<i>t</i> -Bu, <i>t</i> -Bu, H	H, Br	24
B6	<i>i</i> -Pr, <i>i</i> -Pr, H	H, Br	72
B7	allyl, <i>t</i> -Bu, H	H, MeO	49
B8	allyl, <i>t</i> -Bu, H	H, Br	85
B9	Me, <i>t</i> -Bu, H	H, H	13
B10	<i>i</i> -Pr, <i>i</i> -Pr, H	MeO ₂ C, HO	56
B11	<i>n</i> -Pr, <i>t</i> -Bu, H	H, MeO	18 ^a ,
B12	<i>n</i> -Pr, <i>t</i> -Bu, H	H, Br	93 ^e
B13	<i>i</i> -Pr, <i>i</i> -Pr, H	H, pyrazole	28
B14	allyl, <i>t</i> -Bu, H	H, OH	21 ^d

^aSee ref 16.

^b*para-para* Coupling.

^cHomo coupling.

^dTrimer from two molecules of phenol A and one of phenol/naphthol B.

^eObtained by hydrogenation of **B7** or **B8**.

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Table 2 Summary of MIC (*S. mutans*, *S. sanguinis*, and *S. gordonii*) and MBC (*S. mutans*) Values for Analogs^a

Analogs	MIC <i>S. mutans</i>	MIC <i>S. sanguinis</i>	MIC <i>S. gordonii</i>	MBC <i>S. mutans</i>
1	250	125	125	-
2	32	32	32	63
3	>250	>250	>250	-
4	>250	>250	>250	-
5	>250	>250	>250	-
6	>250	>250	>250	-
7	32	32	16	32
8	>250	125	125	-
9	>250	>250	>250	-
10	>250	250	250	-
11	>250	>250	>250	-
1	>250	>250	>250	-
2	>250	>250	>250	-
3	63	63	63	-
4	125	125	125	-
5	8	4	16	125
6	>250	>250	>250	-
7	125	63	125	-
8	16	8	16	32
9	32	32	32	32
10	32	8	8	32
11	16	8	16	63
12	>250	250	250	-
13	>250	>250	>250	-
14	63	16	32	-
1	>250	125	125	-

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Analog	MIC <i>S. mutans</i>	MIC <i>S. sanguinis</i>	MIC <i>S. gordonii</i>	MBC <i>S. mutans</i>
2	2	2.5	1.25	4

^aMIC and MBC values were completed in biological triplicate. MBC assays were completed for analogs with MIC values <32 µM.