CHEMISTRY AND BIOLOGICAL ACTIVITIES OF VIBSANE-TYPE DITERPENOIDS

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Abstract – This review focuses on the structural diversity, biological activities and synthesis of vibsane-type diterpenoids. Vibsane-type diterpenoids are considered to be rarely occurring natural products because they have been found exclusively in a few Viburnum species such as V. awabuki, V. odoratissimum, and V. suspensum. These diterpenoids are further classified into 11-membered ring, 7-membered ring, and rearranged (neovibsanin) types, and therefore, their chemical diversity forms a unique chemical library. We describe the absolute stereochemistry of the typical 11-membered ring vibsanins B and F, a variety of vibsane-type diterpenoids, the chemical correlations between the three subtypes, their biological activities, and the synthesis of vibsanin F and neovibsanin B.

INTRODUCTION

The genus Viburnum consists of about 150 species of shrubs or small trees that were previously included in the family Caprifoliaceae. However, recent classifications based on molecular phylogeny have put them in the family Adoxaceae. They are distributed in the temperate Northern Hemisphere, with a few species extending into tropical regions in South America and Southeast Asia, and about 15 species are distributed in Japan. There is a long history of the folk medicinal use of Viburnum species. For example, the dried bark of V. opulus, which is known as "Cramp Bark," is used to alleviate painful menstrual cramps as well as a sedative. Native American women took black haw (V. prunifolium) to treat the menopause and menstrual cramps. The genus Viburnum has been documented to contain a variety of compounds such as iridoids, terpenoids, and aromatic compounds. Among the chemical contents of Viburnum species, vibsane-type diterpenoids are considered to be characteristic of the Viburnum species because they have not been found in other higher plants. In this review, we focus on the structural diversity, biological activities and synthesis of vibsane-type diterpenoids.
1. Vibsane-type diterpenoids

In 1980, Kawazu reported the isolation of vibsane-type diterpenoids from the leaves of *Viburnum awabuki*, and they were shown to consist of a unique fumulane skeleton with an additional C5 unit (Figure 1).

![Figure 1. Carbon skeletons of fumulane and vibsane](image)

However, there has been little interest in their chemical structure and biological activity since their discovery. Since 1996, we have continued to investigate the vibsane-type diterpenoids that are specific to *Viburnum* species and their biological activities, resulting in the discovery of about 60 new diterpenoids. They possess unique structures, some of which have unexpected chemical reactivity and interesting biological activity. The first vibsane-type diterpenoids that were reported by Kawazu were vibsanins A

![Figure 2. Vibsanins A–F, which were isolated from Viburnum awabuki by Prof. Kawazu](image)
17), B (1), and F (3), which possess an 11-membered ring, and vibsanins C (2) – E (16), which have a 7-membered ring (Figure 2). Their stereochemistry has remained unexplored except for that of vibsanin E (16). Since these diterpenoids consist of a new carbon skeleton, we have proposed the new term of “Vibsane” for diterepenoids possessing a fumulane carbon framework with an additional isoprene unit.

2.1. The stereochemistry of vibsanins B (1) and C (2)

It was necessary to determine the absolute stereochemistry of vibsanins B (1) and C (2) before discussing the structures of the newly isolated vibsane-type diterpenoids. The $^1$H NMR of 1 showed two kinds of broad signals at room temperature, but a pair of the sharp signals was observed at 0°C. This phenomenon indicated that vibsanin B (1) is present in solution as two conformational isomers.

![Figure 3](image)

**Figure 3.** The two conformers, 1a and 1b, elucidated by NOESY, and the lowest energy conformers CT and BC obtained by MM2 calculations. The arrows show the NOE

Two conformers, 1a and 1b, were elucidated on the basis of NOESY data and $J$ values (Figure 3). The main conformer, 1a, demonstrated the NOE correlations shown in Figure 3 and had a high $J_{8,9}$ (9.3 Hz) value. These NMR data suggested that 1a adopts a chair-like conformation for the sequential bonds from C-5 to C-10 with a dihedral angle of 180° between C8-H and C9-H and takes a *transoid* geometry for the $\alpha,\beta$-unsaturated ketone at C4–C6, whereas 1b consists of a boat conformation and has a *cisoid* form according to NOE analysis and its small $J_{8,9}$ (2.2 Hz). Additionally, 1a and 1b were consistent with the two most stable conformers, CT and BC, found by the MM2 calculations. In the course of the VT (variable temperature) experiments for 1 in DMSO-$d_6$, we found that 1 induced an irreversible change at
110°C. To elucidate this thermal transformation in detail, a solution of \textbf{1} in toluene was refluxed for 1 h to give rise to four products, which eventually were found to correspond to 7-membered ring vibsanin C (\textbf{2}) (85.9%) and its stereoisomers \textbf{2a} (11.2%), \textbf{2b} (1.6%), and \textbf{2c} (0.2%) (Scheme 1). Vibsanin C (\textbf{2}) and 5-\textit{epi}-vibsanin C (\textbf{2b}) are natural products, but \textbf{2a} and \textbf{2c} have been not found in nature. The formation of the four 7-membered ring vibsanins can be ascribed to the \textit{oxy}-Cope rearrangement of \textbf{1}. The major product, vibsanin C (\textbf{2}), and its 5-epimer, \textbf{2b}, which contains a $\Delta^{8,9} E$ olefin, are considered to rearrange through the \textbf{CT} and \textbf{BC} conformers, respectively; whereas, \textbf{2a} and \textbf{2c}, which contain $\Delta^{8,9} Z$ olefin, are presumably transformed through the \textbf{CC} and \textbf{BT} conformers, which were found to be within 6 kcal/mol of the global minimum energy by MM2 calculations (Figure 4).

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{schematic.png}
\caption{Scheme 1. \textit{Oxy}-Cope rearrangement of vibsanin B (\textbf{1})}
\end{figure}

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{conformers.png}
\caption{\textbf{CC} and \textbf{BT} conformers leading to \textbf{2a} and \textbf{2c}, respectively}
\end{figure}
Thus, the absolute configuration of the 11-membered ring vibsanin B (1) was completely correlated with that of the 7-membered ring vibsanin C (2) via an oxy-Cope rearrangement. Next, the enol ester of 2 was saponified under basic conditions, which was followed by an intramolecular aldol condensation reaction to give rise to the aldehyde 2d, which was converted to the bromophenyl carbamate 2e. X-ray crystallographic analysis of 2e unambiguously established the absolute 5S, 10R, and 11S configurations of vibsanin C (2) (Scheme 2). This result means that vibsanin B (1) has chiral centers of 7R, 8R, and 11S. Thus, the absolute structures of 1 and 2, which were previously unsolved, have been established (Figure 5).
2.2. Determination of the absolute configuration of vibsanin F (3) by synthesis

Although vibsanin F (3), which was isolated from the leaves of *V. awabuki* in 1980, belongs to the simplest 11-membered ring form of the vibsane-type diterpenoids, its stereochemistry has never been solved. We have decided to unambiguously determine the absolute stereochemistry of 3 via its asymmetric synthesis. Vibsanin F (3) has three chiral centers, among which C-7 and C-11 are anticipated to be 7S and 11S, respectively, based on those of vibsanin B (1) as shown in Figure 6. However, the chirality of C-6 may be 3a (6S) or 3b (6R).

![Possible structures of 3a and 3b for vibsanin F (3)](image)

Figure 6. Possible structures of 3a and 3b for vibsanin F (3)

First, we selected 3a as the first synthetic target. The synthetic procedures used are outlined in Schemes 3 and 4. Asymmetric epoxidation of the allyl alcohol 4 by the Sharpless protocol provided 5, which had 6S and 7S chiral centers that corresponded to those of 3a. Regioselective epoxidation of 5 with m-chloroperbenzoic acid exclusively gave the diepoxyd 6 as a diastereomeric mixture. The primary hydroxyl group of 6 was converted to its triflate, with which a large excess of the dianion of methyl acetoacetate was reacted at 0°C to give rise to 7 in high yield. The introduction of a 4-methyl-3-pentenyl unit using sodium hydride and 15-crown-5 in DMSO gave rise to the precursor 8 in good yield, which was required for the subsequent palladium-catalyzed macrocyclization. Subjecting 8 to the Tsuji-Trost reaction using 10 mol% Pd(PPh₃)₄ in DMSO afforded the sole product 9 in a moderate yield. The stereoselective formation of 9 can be explained by assuming that transition states A and B, as shown in Figure 7, are involved in this cyclization. If the nucleophilic displacement of the π-allylpalladium intermediate with the anion of the β-ketoester moiety proceeds through a product-like transition state, transition state A, which should lead to the more stable product 9 with a pseudoequatorial C6 unit and a pseudoaxial methyl group, probably favors over transition state B, producing a less stable product with the opposite stereochemistry at the quaternary center.
Scheme 3. Reagents and conditions: a) Ti(OiPr)$_4$, L-(-)-DET, TBHP, MS4A, CH$_2$Cl$_2$, -35°C, 99% (92% ee); b) MCPBA, CH$_2$Cl$_2$, 0°C, 91%; c) Tf$_2$O, Et$_3$N, THF, -78°C, then 5 eq. the dianion of methyl acetoacetate generated with NaH and nBuLi, THF, 0°C, 94%; d) NaH, 15-crown-5, DMSO, then 5-iodo-2-methylpent-2-ene, 77%; e) 10 mol% Pd(PPh$_3$)$_4$, DMSO, 90°C, 60%.

Figure 7. Possible transition states, A and B, for palladium-catalyzed cyclization of 8

Although the 11-membered ring was diastereoselectively constructed, the Z-geometry of the trisubstituted olefin in 9 has to be converted to E-geometry. First, the alcohol 9 was protected with TBDMSCl, before the reduction of both carbonyl groups with LiAlH$_4$ to give rise to the diol 10, which was then mesylated (Scheme 4). The resultant dimesylate was subjected to elimination of the secondary mesylate under basic conditions, giving rise to the monomesylate 11 in a moderate yield over three steps. Subsequent reductive demesylation of 11 was successfully achieved using a NaBH$_4$-DMPU system, resulting in the formation of 12 in good yield. The aldehyde 13, which was derived from 12, was treated with AIBN and thiophenol
to produce the desired E-olefin 14. Finally, the conjugate aldehyde in 14 was reduced by the Luche protocol to afford 3a. The $^1$H NMR of 3a, however, was not identical to that of natural vibsanin F (3). Thus, each epoxide ring of the synthetic product 3a and vibsanin F was reduced with LiAlH$_4$, resulting in the preparation of the same diol 15. All the spectroscopic data for both diols were identical to each other, and therefore, the absolute configuration of vibsanin F has been established to be 3b with 6R, 7S, and 11S forms.

Scheme 4.  Reagents and conditions: a) TBDMSCl, Et$_3$N, 4-DMAP, CH$_2$Cl$_2$, 74%; b) LiAlH$_4$, THF, 0°C; c) MsCl, Et$_3$N, 4-DMAP, CH$_2$Cl$_2$, 0°C; d) DBU, toluene, 120°C, 60% over three steps; e) NaBH$_4$, DMPU, 55°C; f) TBAF, THF, 100%; g) Dess–Martin periodinane, Et$_3$N, CH$_2$Cl$_2$, 100%; h) PhSH, AIBN, benzene, 90°C, 48%; i) NaBH$_4$, CeCl$_3$, MeOH, 0°C, 56%.

2.3. 11-Membered ring vibsane-type diterpenoids

Vibsane-type diterpenoids consist of three sub-types, 11-membered ring, 7-membered ring, and the rearranged types. These diterpenoids occur exclusively in V. awabuki, V. odratissimum, V. suspensum, and V. sieboldi. The 11-membered ring and 7-membered ring vibsane-type diterpenoids are common to these plants. Most 11-membered ring vibsanins consist of a fumulane-like skeleton containing a β, β-dimethylacrylate group at the C-8 position except for vibsanin F (3). Vibsanin A (17) and vibsanins P (18) – T (22), which contain oxidatively modified C6 units at the C-11 position, were isolated from both
Japanese and Taiwanese *V. odoratissimum*.

Since the conformations of the 11-membered ring vibsanins A and P – T, which bear a 6,7-epoxide ring, are fixed, analysis of their NMR can be performed normally. On the other hand, vibsanin B (1) and vibsanols A (23) and B (24), which contain a cross-conjugated diene, show complex NMR signals due to the presence of several kinds of conformational isomers. Therefore, careful structure elucidation should be performed by a combination of VT NMR experiments and MM2 calculations.\(^{11}\)

![Figure 8. 11-Membered ring vibsanins](https://example.com/figure8.png)

**Figure 8.** 11-Membered ring vibsanins

### 2.4. 7-Membered ring vibsane-type diterpenoids

The 7-membered ring vibsanins are made up of a diverse range of compounds, which possess two ketones and an isoprene unit attached to the 7-membered ring. There are two basic types, the (5S,10R) and (5R,10R) stereoisomers. Vibsanins C (2), G (25), H (27), and 18-O-methylvibsanin G (26), which belong to the (5S,10R) type, occur commonly in *Viburnum* species.\(^{17,18}\) Vibsanins I (30), J (31), K (32), and 18-O-methylvibsanin K (33)\(^{19}\) as well as 14,15-epoxyvibsanin C (34) were isolated from *V. awabuki* in Tokushima\(^{17,19}\) and *V. awabuki* in Taiwan, respectively. On the other hand, 5-epi-vibsanins C (35), H (36), I (38), K (39) and their 18-O-methyl and/or 15-O-methyl congeners 37 and 40\(^{20}\) have been found in all
**Viburnum** species except for *V. suspensum*. Vibsanin M (41), which possesses a $\Delta^{4,5}$ double bond\(^{18}\) and the bicyclic vibsanin N (46)\(^{21}\) were isolated from *V. odoratissimum* and *V. awabuki* in Taiwan. Furthermore, aldovibsans A (42) – C (45) were isolated from *V. odoratissimum*.\(^{22, 18}\)

![Chemical structures](image.png)

**Figure 9.** 7-Membered ring vibsanins

Additionally, interesting tricyclic 7-membered vibsanins are known (Figure 10). Vibsanin E (16) and 16-hydroxyvibsanin E (47),\(^{21}\) which were isolated from *V. awabuki*, have an ether bond between C-15 and C-18; whereas, cyclovibsans A (48) and 15-O-methylcyclovibsans A (49), 15-O-methylcyclovibsans B (50), and 3-hydroxy-15-O-methylcyclovibsans A (51), which are composed of a tricyclo[6.3.2.0\(^{3,6}\)]tridecane skeleton, contain C-C bonds between C-18 and C-16 or C-17. Cyclovibsans have no oxygen atoms between C-18 and C-15.\(^{24}\)
Figure 10. 7-Membered ring tricyclic vibsanins

Scheme 5. Conversion of vibsanin C (2) to vibsanin E (16) and a plausible biosynthetic pathway for cyclovibsanins

Vibsanin E (16) can be readily converted from vibsanin C (2) via a cationic process (a) by treating it with BF₃OEt₂ as shown in Scheme 5.³³ On the other hand, another plausible biosynthetic pathway for cyclovibsanins is (b) shown in Scheme 5, in which a proton is eliminated from one of two methyl groups in 52 to give rise to 52a. Dehydration produces the exomethylene ketone 52b, the C-4 carbonyl group of which is then protonated to trigger cyclization through the cationic intermediate 52b, resulting in the formation of a tricyclic framework such as that seen in the cyclovibsanins.

Furanovibsanins, a diverse range of 7-membered ring vibsanins, which are presumed to be produced by two ketones at the C-4 and C-7 positions, have also been found (Figure 11). The examples shown are furanovibsanin A (53) and its 3-O-methyl congener 54, which is derived from 3-hydroxyvibsanin E (47). The additional examples are furanovibsanin B (55) and its 7-epimer 56, and furanovibsanins C (57) – G.
(61). These diterpenoids were isolated from *V. awabuki* collected in Tokushima.$^{25}$

![Diagram of diterpenoids](image)

**Figure 11.** 7-Membered ring furanovibsanins

### 2.5. Rearranged vibsane-type diterpenoids (neovibsanins)

The rearranged vibsane-type diterpenoids (neovibsanins), which contain a β, β-dimethylacrylate ester and an isoprene unit-substituted cyclohexene ring core fused to a tetrahydrofuran ring, are rarely occurring natural products (Figure 12). In 1996, the first rearranged vibsane-type diterpenoids, neovibsanins A (62) and B (63), were isolated from *V. awabuki*. Recently, we have ascertained the presence of neovibsanin (86) in the fresh leaves of *V. awabuki* and also found that neovibsanin is changed to neovibsanins A (62) and B (63) when kept in methanol at room temperature, indicating that neovibsanins A and B are artifacts derived from neovibsanin. Since neovibsanins A (62) and B (63) were reported, a number of rearranged vibsane-type diterpenoids have been found in *V. awabuki*, *V. suspensum*, and *V. sieboldi*, and have become known as a characteristic compound group of the *Viburnum* species as summarized in Figure 12.
Figure 12. Rearranged vibsane-type diterpenoids (neovibsanins)
Among them, neovibsanin C (64) is the first example of a natural product with a macrocyclic structure formed through an endo-peroxide group. The unusual structure of 64 was established by converting it from neovibsanin B (63) as outlined in Scheme 6. The reduction of 64 with Zn in EtOH-AcOH resulted in the formation of the diol 83, which was treated with methanol under acidic conditions to give rise to the methyl acetal 84. The acetal 84 was also derived from neovibsanin B (63) by photosensitized oxidation, followed by reduction of the formed peroxy group [7-epi-neovibsanin D (66)]. In addition, when 66 was treated with pTsOH in anhydrous benzene, 64 was generated in good yield, presumably by an acetal exchange reaction. Thus, the structure of neovibsanin C (64) including its absolute configuration was established.

Scheme 6. Chemical correlation of neovibsanin B to neovibsanin C
Neovibsanins H (67) and I (69), and their 2-O-methyl congeners 68 and 70 are neovibsanins that do not possess an acetal group. Similar compounds containing an additional 6-membered ring [neovibsanins F (71) and its 14-epimer 73, and 14-epi-18-oxoneovibsanin F (74)] were isolated from *V. suspensum*.

Spirovibsanin A (82) is the first example of a nor-vibsane-type diterpenoid. Recently, neovibsanins J (79), K (80), and P (81), in which two hydroxy groups at the C-4 and C-18 positions are involved in acetal formation on the C-7 carbonyl, were found.

Scheme 7. Photochemical reaction of vibsanin B (1)

It has been shown that a 7-membered ring vibsanin C (2) can be derived from vibsanin B (1), an 11-membered ring vibsane-type diterpenoid, by thermal oxy-Cope rearrangement. However, no formation of neovibsanins was observed even upon heating 1 under acidic or basic conditions. It should be noted
that neovibsanins A (62) and B (63) can be produced from vibsanin B (1) by photochemical reaction. Surprisingly, irradiation of vibsanin B (1) in benzene with a high pressure Hg lamp afforded 87, which possesses a neovibsanin-framework, in 4% yield, along with 85 (18%) and 2c (27%) (Scheme 7). These compounds have been neither found in natural sources. When the photochemical reaction of 1 was carried out in MeOH, neovibsanins A (62) and B (63) were produced in 12% and 20% yields, respectively, in addition to 88 (9%) and 89 (8%) as over-reacted products. The generation of 88 and 89 was presumably due to a series of fragmentation and cationic cyclizations triggered by the methanolysis of a β,β-dimethylacryl ester group. Furthermore, irradiation of 1 in 50% aqueous MeOH for 1 h directly yielded neovibsanin (86). This photochemical reaction of 1 forces the E/Z isomerization of the double bond at C-5 to generate (5Z)-vibsanin B (85). The MM2 calculations for 85 and 1, in which the C12–C17 side chain was replaced with a t-butyl group, were performed using MacroModel® and provided the most stable conformers, 1-(5Z) and 1-(5E), for each molecule as shown in Figure 13. In the case of 1-(5Z), the distance between the C-4 carbonyl and the C-7 OH is 1.65 Å, whereas 1-(5E) has a distance of 4.98 Å as depicted in Figure 13. This means that a 1,7-hydrogen shift from the OH group at C-7 to the carbonyl at C-4 occurs in 1-(5Z), but not in 1-(5E). This hydrogen shift not only causes a breakage of the C-7/C-8 bond as well as cyclization between C-10 and C-4, leading to the neovibsanin-framework, but also undergoes another oxy-Cope rearrangement via 1a to give rise to a 7-membered ring (8Z)-10-epi-vibsanin C (86), which is not formed by the thermal oxy-Cope rearrangement of vibsanin B (1).

![Figure 13](image-url)

**Figure 13.** The most stable conformers, 1-(5Z) and 1-(5E), for (5Z)-vibsanin B (85) and vibsanin B (1), respectively, obtained by MM2 calculation.
Taking the aforementioned results into consideration, we wish to propose plausible biosynthetic mechanisms for neovibsanins, as outlined in Scheme 8. The biosynthesis of all neovibsanins starts from the key intermediate A, which is derived from vibsanin B (I). The C-18 hydroxy group undergoes 1,4-addition to yield B. In the case of route a, a hemiacetal E is generated, leading to neovibsanins A–D. Likewise, route b gives the intermediate allyl cation C prior to dehydration, which is trapped by some nucleophiles such as water to afford neovibsanins H–I (route c) and causes cationic cyclization to give D, resulting in the formation of neovibsanins F–G (route d). Additionally, neovibsanins J–K are presumably converted from the key intermediate A via the production of a bicyclic acetal of the C-7 carbonyl containing C-18 and C-4 hydroxy groups. However, a question still remains about how the key intermediate A is generated from vibsanin B (I) by enzyme-catalyzed reaction.

Scheme 8. Plausible biosynthetic pathways for the formation of various neovibsanins from vibsanin B (I)

3. Biological activities of vibsane-type diterpenoids
The leaves of V. awabuki have been used as a fish poison for catching fishes in Okinawa and Southeast Asia for a long time. In 1980, Kawazu reported the first isolation of a piscicidal compound, vibsanin A, and a plant growth inhibitor, vibsanin B (I). Later, many vibsane-type diterpenoids were reported,
among which some diterpenoids such as vibsain C (2), 5-epi-vibsain C (35), vibsanol A (23), and vibsains K (32) and P (18) exhibited significant and/or moderate cytotoxicity against tumor cells. Additionally, vibsain B (1) and neovibsain F (71) showed moderate toxic activity in a brine shrimp lethality assay.

3.1 Neurotrophic activities of neovibsains

A remarkable pathological symptom of Alzheimer’s disease (AD) is the loss of neuronal cells in the brain. Correspondingly, the overall strategy for treatment of AD is to prevent neuronal death or to produce new neuronal cells in the degenerative regions. Neurotrophins, such as nerve growth factor (NGF), brain-derived neurotrophic factor (BDNF), neurotrophin 3 (NT-3) and glial cell line-derived neurotrophic factor (GDNF), are recognized as important regulatory substances in the nervous system. Thus, neurotrophins are expected to have therapeutic efficacy for the treatment of AD. However, they can not cross brain-blood barrier because of the properties of its high molecular polypeptide and are easily metabolized by peptidases under physiological conditions. To address this issue, considerable efforts have been made to find small molecules that have neurotrophic properties or are capable of enhancing the action of NGF in appropriate cell populations. Rat pheochromocytoma (PC12) cells have been used as a good in vitro model of neuronal differentiation. After stimulation with NGF, PC12 cells differentiate to extend neurites and develop the characteristics of sympathetic neurons. For example, two iridoids, picrosides I and II and some clerodane-type diterpenoids, 6α,7α-dihydroxyannonene, and 7α,20-dihydroxyannonene have been demonstrated to show neurite outgrowth-promoting activity in NGF-mediated PC12 cells. Also some synthetic compounds, N-benzyloxycarbonyl-Leu-Leu-leucinal (ZLLLal), AIT-082, SR57746, and Aroclor 1254 have been reported to accelerate the action of NGF in PC12 cells. Recently, we have found that neovibsain (86), and neovibsains A (62) and B (63) have neurotrophic properties. Namely, they promote neurite outgrowth of NGF-mediated PC12 cells at concentrations ranging from 10 μM to 40 μM.

Evaluation was carried out for PC12 cells neurite outgrowth according to a previously reported experiment procedure. As shown in Figure 14, neovibsain (B), neovibsain A (C) and neovibsain B (D) significantly promoted neurite outgrowth from NGF (10 ng/mL)-treated PC12 cells at 40 μM. Among three compounds, neovibsain A seems less potent than neovibsain and neovibsain B. However, all of them had no effect on morphology of PC12 cells in the absence of NGF. Additionally, other vibsane-type diterpenoids such as 7-membered and 11-membered ring subtypes have not been so far found to show neurotrophic properties in PC12 cells.
PC12 cells were cultured in DMEM / 2% HS + 1% FBS and treated by A: NGF 10 ng/mL, B: Neovibsain 40 μM + NGF 10 ng/mL, C: Neovibsain A 40 μM + NGF 10 ng/mL, D: Neovibsain B 40 μM + NGF 10 ng/mL.

Quantitative analysis of the percentage of cells with neurites and the neurite length extending from the cell bodies (Figure 15) showed that neovibsain (86) and neovibsain A (62) significantly increased the percentage of PC12 cells bearing neurites and the neurite length compared with those of NGF-mediated PC12 cells at concentrations ranging from 5 to 40 μM, and, in the degree of activity, 62 (Figure 15, C and D) was likely to be less that 86 (Figure 15, A and B). On the other hand neovibsain B (63) promoted efficiently the neurite outgrowth from of NGF-mediated PC12 cells in a dose-dependent manner at concentrations from 10 to 40 μM (Figure 15, E and F). In comparison of the percentage of cells with neurites and the average neurite lengths (Figure 16), neovibsain B (63) seemed to be a more potent NGF-potentiatior among three compounds. This result consists with morphological evaluation. It is assumed from these results that a stereochemistry on the acetal carbon may be related with affecting neurite outgrowth activity.
Figure 15. Neurite outgrowth-promoting activities of neovibsanin, neovibsanin A, and neovibsanin B in PC12 cells

PC12 cells were cultured in DMEM / 2% HS + 1% FBS with or without 10 ng/mL NGF and different concentrations of neovibsanin, neovibsanin A, and neovibsanin B for 48 h. PC12 cells were fixed and quantified for the percentage of cells bearing neurites and the primary neurite length. Over 40 fields were randomly selected under microscope for analysis of the percent of cells with neurites. At least 200 cells were selected for calculating the neurite length. Data were expressed as means ± SE. ***$P < 0.001$ compared with NGF only by one-way ANOVA followed by Bonferroni post hoc means comparison. ###$P < 0.001$ vs. control by Student's t-test. A, B: neovibsanin, C, D: neovibsanin A, E, F: neovibsanin B.
Figure 16. Comparison of neurite outgrowth-promoting activities of neovibsanin, neovibsanin A, and neovibsanin B in PC12 cells

The method was the same as that in Figure 15. Data were expressed as means ± SE. Difference between groups was tested with Student's t-test. ***P < 0.001 compared with NGF only.

4. Synthesis of neovibsanin B

These unique molecular architectures and significant biological activities have strongly motivated organic chemists to devote their efforts to the syntheses of the vibsane-type diterpenoids. So far, the vibsane-type diterpenoids whose total syntheses have been achieved are only three molecules, i.e. (±)-2-O-methylneovibsanin H (68), (±)-neovibsanin B (63), and (±)-vibsanin E (16). From a synthetic point of view, the stereocontrol in the successive stereogenic centers involved in these diterpenoids has been challenging. Williams’s pioneering synthetic studies have addressed the issue of diastereoselective creation of the stereocenters involved in vibsane natural products. In fact, through many approaches toward the total syntheses of these molecules, the synthetic efforts have resulted in formation of diastereomers of the natural products (Figure 17), i.e. (6S)-vibsanin F (3a), (±)-5,10-bis-epi-vibsanin E (16a), (±)-5,14-bis-epi-spirovibsanin A (82a), and
(±)-4,5-bis-epi-neovibsanins A (62a) and B (63a).  

This review is focused on the synthesis of neovibsanin B (63), which is not only a significant neurotrophic mimic but also the most challenging molecule among vibsane natural products. In 2009, Imagawa and Nishizawa reported the first synthesis of (±)-neovibsanin B (63). They employed an intramolecular Diels-Alder reaction of 91 at 200 °C in dimethyl imidazolidinone (DMI) as a solvent for constructing the core cyclohexenone ring of neovibsanin, leading to a mixture of 92a and 92b (9:1) in 58% yield. DMI plays a crucial role in accelerating this reaction rate and thus makes it suitable for scale-up. Further manipulation of Diels-Alder adducts 92a and 92b involves introduction of a hydroxymethyl group by Baylis-Hillman reaction with formaldehyde to provide the key intermediate 95 (Scheme 9). It should be noted that 1,2-addition of nucleophiles to the C-4 ketone was dominated by addition of the less hindered undesired face to afford exclusively products having the undesired stereochemical disposition at C-4. Imagawa and Nishizawa overcame this steric drawback by devising tactic of using the oxygen of 2,4-dimethoxybenzyl (2,4-DMMP) group at C-10 to coordinate with the organolithium reagent and deliver the propargyl group from the same face. Reaction of excess lithio ethylpropiolate 96 with a toluene solution of 95 at -78°C successfully proceeded to give rise to the

**Figure 17.** Synthesized diastereomers of vibsane family and neovibsanin B (63)
Scheme 9. Reagents and conditions: a) (COCl)$_2$, benzene, reflux; b) TBSOCH=CHCH=CH$_2$, MeLi, CH$_2$Cl$_2$, DME, -20°C, 79%; c) DMI, 200°C, 58%; d) DIBAL, THF, -70°C to rt, 89%; e) Bu$_3$SnO, toluene, reflux, and then 2,4-DMPMCl, TBAI, toluene, reflux; f) Dess-Martin reagent, CH$_2$Cl$_2$, 0°C, 70%; g) Bu$_3$P, HCHO, aq. MeOH/CHCl$_3$, rt, 84%; h) TBSCl, imidazole, CH$_2$Cl$_2$, 0°C, 99%.

Adduct 97 with the correct stereochemistry in 87% yield as a single diastereomer. 1,2-Addition of 96 to the C-4 ketone in 95 was completely controlled by the coordination of the two oxygen atoms of 2,4-DMPM group with 96, which is shown in Figure 18, to give 97 having the desired β-configuration at C-4. The triple bond of 97 was reduced with Red-Al to the α,β-unsaturated ester 98, which was in turn treated with TBAF for the deprotection of the TBS group, thereby triggering the subsequent Michael addition and lactonization to give rise to a tricyclic lactone 99 having the desired stereochemistry in good yield. The Deprotection of 2,4-DMPM in 99 by DDQ oxidation was troublesome due to the acidity of in situ formed DDQH. To avoid acidic conditions, this oxidation was carried out in a two phase system of CH$_2$Cl$_2$ and NaCl saturated-phosphate buffer to give a desired alcohol, which was protected again as the TBS group, giving rise to 100 in good yield. The resulting 100 was reacted with Tebbe reagent, and then treated with PPTS in methanol afforded a mixture of 102a and 102b (1:4.5), which were readily separated by HPLC. The major 102b was converted to the aldehyde 103, which was treated with KHMDS to generate a potassium enolate. This was in situ trapped with 3,3-dimethylacryloyl chloride completing the first total synthesis of (+)-neovibsanin B (63).

Recently, we reported efficient construction of the chiral all-carbon quaternary center with a vinyl moiety that would permit post-functional group manipulation by the conjugate addition of lithium divinyl cuprate to (4S,2′E)-3-(6′-TBDPS-3′-methylhex-2′-enoyl)-4-phenyloxazolidin-2-one (115), and demonstrate that this method provides a versatile chiral quaternary carbon source 117 for the synthesis of natural products by its use to the synthesis of (+)-bakuchiol (Scheme 11).
Scheme 10. Reagents and conditions: a) Red-Al, THF, -78°C, 87%; b) TBAF, THF, rt, 87%; c) DDQ, CH\textsubscript{2}Cl\textsubscript{2}/phosphate buffer, NaCl, 0°C, 83%; d) TBSCl, imidazole, 0°C to rt, 99%; e) Tebbe reagent, pyridine, THF-toluene; f) PPTS, MeOH, 0°C, 91%; g) TBAF, THF, rt, 99%; h) SO\textsubscript{3}pyridine, Et\textsubscript{3}N, DMSO, rt, 89%; i) KHMDS, THF, then 3,3-dimethylacryloyl chloride, -78°C, 60%.

Figure 18. Chelated control of the 2,4-DMPM group with 96
Scheme 11. Enantioselective construction of the chiral all carbon quaternary center at C-11

Scheme 12. Reagents and conditions: a) 30% \( \text{H}_2\text{O}_2 \), LiOH, THF-H\( \text{H}_2\text{O} \), 0°C; b) EtOH, EDC, DMAP, CH\( _2\text{Cl}_2 \), rt; c) LiAlH\( _4 \), THF, 0°C; d) PCC, celite, CH\( _2\text{Cl}_2 \), 0°C; e) CBr\( _4 \), PPh\( _3 \), CH\( _2\text{Cl}_2 \), 0°C; f) \( \text{n-ButLi} \), THF, -78°C, then (CH\( _2\text{O})\_n \), rt; g) Bu\(_3\)SnH, AIBN, THF, reflux; h) I\(_2\), CH\( _2\text{Cl}_2 \), 0°C; i) TBSCl, DMAP, Et\(_3\)N, CH\( _2\text{Cl}_2 \), rt.

We have decided to apply this methodology to create the C-11 chiral quaternary carbon involved in vibsane natural products. As the first target, we selected the Imagawa-Nishizawa intermediate 95 that is the core framework of neovibsanin natural products. The asymmetric 1,4-addition reaction of
(CH₂=CH)₂Cu(CN)Li₂ to 118 bearing (R)-4-phenyl-2-oxazolidinone was employed to give 119 as a diastereomeric mixture of 95 (11S):5 (11R) in good yield. Each diastereomer was readily separated by silica-gel chromatography. The optically pure (11S)-119 was converted to (2Z,11S)-122 according to the procedures outlined in Scheme 12.

Previously, we reported that the modified Negishi palladium(0)-catalyzed carbonylative cyclization of (±)-122 provided the cyclohexene-1-one derivative (±)-125, which corresponds to the cyclohexene ring of neovibsanin. We examined this reaction in detail (Table 1). First, the reaction was performed using 5 mol % PdCl₂(PPh₃)₂ and Et₃N (1.5 equiv) in MeCN/PhH (1:1) containing 4 equiv of MeOH at 100°C in an autoclave, which gave rise to the desired diastereomeric mixture of 123 in 11% yield along with ca. 50% of 125 containing a small amount of 124 (6%) (Table 1, entry 1). On the other hand, the addition of an excess amount (48 equiv) of MeOH to this reaction system dramatically increased the yield of 123 to 54%, contaminated with 2% of the noncyclic ester 124 (entry 2). The use of high pressure (8 MPa) was found to be ineffective at suppressing the generation of 124 (entry 3), but low temperature (60°C) was able to decrease the generation of 124 (entry 4). After several trials, we found that the following reaction

<table>
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<th>entry</th>
<th>base (1.5 equiv)</th>
<th>solvent</th>
<th>MeOH (equiv)</th>
<th>CO (MPa)</th>
<th>temp. (°C)</th>
<th>123 (10R:10S)⁺</th>
<th>122</th>
<th>124</th>
<th>125 (%)</th>
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<tr>
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<td>100</td>
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<td>60</td>
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<td>6 (1.4:1)</td>
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<td>24 (1.6:1)</td>
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<td>9</td>
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⁺ Ratio was determined by ¹H NMR spectroscopy in CDCl₃ (300 MHz).
conditions; 24 equiv of MeOH, 4 MPa CO, temperature of 60°C, led to the formation of 123 alone in ca. 70% yield as a diastereomeric mixture (10R:10S = 2.6:1) (entry 5). Each diastereomer of 123 was readily separated by silica gel column chromatography. It should be noted that 125 was exclusively generated in high yield when 1,2-dioxane was used as solvent (entry 6).

With (10R,11S)-123 in hand, we focused on the last few steps for the synthesis of Imagawa-Nishizawa’s intermediate 95 (Scheme 13). Treatment of (10R,11S)-123 with n-Bu₄NF containing acetic acid gave 126, and the resultant hydroxy group was oxidized by Swern oxidation to its aldehyde, which was subjected to Wittig olefination to give the dimethyl olefin 127 in 69% yield over two steps. Reduction of 127 with DIBAL-H provided the cyclic hemiacetal 128 and diol 129 in 29 and 26% yields, respectively. The cyclic hemiacetal 128 was reduced with NaBH₄ to give 129.

Scheme 13. Reagents and conditions: a) n-Bu₄NF, AcOH, THF, rt; b) (COCl)₂, DMSO, -78°C to 10°C, then Et₃N; c) Me₂CHP⁺Ph₃I⁻, n-BuLi, THF, 0°C; d) DIBAL-H, CH₂Cl₂, -78°C; e) NaBH₄, EtOH, rt; f) 2,4-DMPM-trichloroacetoimidate, 10 mol% CSA, CH₂Cl₂, -20°C; g) Dess-Martin periodinane, CH₂Cl₂, rt.
The selective protection of the primary alcohol in 129 using the reaction conditions, 2,4-DMPM-trichloroacetimidate in the presence of 10 mol % CSA gave rise to the desired 2,4-DMPM-ether 130 in 45% yield. Finally Dess-Martin oxidation of 130 afforded the Imagawa-Nishizawa’s intermediate (+)-95 as an optically active form ([α]D +20.1 (c 1.05, MeOH)) in 95% yield. Thus, the first enantiocontrolled formal synthesis of (+)-neovibsanin B was accomplished.53

CONCLUSION
Since Prof. Kawazu’s first report in 1980, over 80 vibsane-type diterpenoids have been found, and the new term of “Vibsane” has now become accepted. These diterpenoids not only show a rich structural diversity, but also occur specifically in a few Viburnum species. Thus, they are very interesting natural products from chemical and taxonomic points of view, and moreover, their chemical diversity makes them a valuable chemical library. As neovibsanins A and B have been found to exhibit interesting neurotrophic activity, it is expected that new biological activities will be discovered among the vibsane-type diterpenoids. Recently, vibsane-type diterpenoids have attracted much attention from organic chemists due to their unique structures and important biological activities.54-60 We hope that this review has stimulated much attention in synthetic studies of these diterpenoids and that comprehensive reviews of synthetic studies of the vibsane-type diterpenoids will appear in the near future.

ACKNOWLEDGEMENTS
We thank our colleagues for their dedication toward these vibsane projects, whose names are listed in the literature cited in references. These works were supported by Grant-in-Aids for Scientific Research from the Ministry of Education, Culture, Sports, and Technology of Japan and the Open Research Fund from the Promotion and Mutual Aid Corporation for Private School of Japan.

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