

Biotechnological potential of marine natural products*

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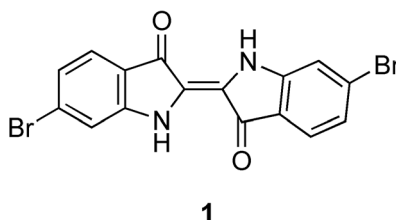
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Abstract: The number of marine natural products (MNPs) that have been applied to biotechnological industry is very limited, although nearly 20 000 new compounds were discovered from marine organisms since the birth of MNPs in the early 1970s. However, it is apparent that they have a significant potential as pharmaceuticals, cosmetics, nutraceuticals, research tools, and others. This article focuses on selective antitumor metabolites isolated from marine sponges and tunicates and their modes of action, as well as promising candidates for nontoxic antifoulants discovered from marine organisms.

Keywords: antifoulants; antitumor compounds; biotechnology; marine natural products; pharmaceuticals; research tools.

INTRODUCTION

Tyrian purple (**1**), a brilliant purple dye documented in texts dating from about the 16th century B.C., was produced from the hypobranchial glands of marine gastropod mollusks of the superfamily Muricacea [1]. This was man's first large-scale marine biotechnological industry originating in Tyre, a Phoenician city. However, the application of marine natural products (MNPs) to industry started quite recently, because of man's limited access to marine resources during virtually all of recorded history. MNPs are potential sources of pharmaceuticals, cosmetics, nutraceuticals, agrochemicals, and others. Their application to drugs has been explored since the early 1970s, but only a small number of MNPs have been applied to pharmaceuticals [2]. Recently, the increasing numbers of MNPs have been developed for nutraceuticals [3] and cosmetics [4]. It should be noted that many MNPs, especially toxins (e.g., tetrodotoxin) have been used as research tools (reagents) and contributed enormously to the progress of life science. Surprisingly, only nereistoxin isolated from a polychaete was applied to an agrochemical [5]. This article focuses on selective antitumor compounds that we isolated from Japanese marine invertebrates and their modes of action, as well as MNPs applicable to nontoxic antifoulants.

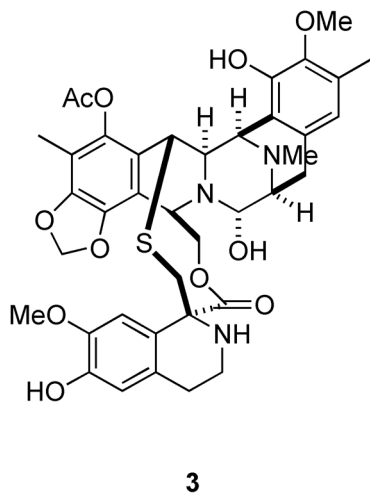
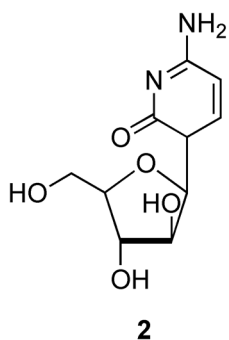


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ANTITUMOR COMPOUNDS

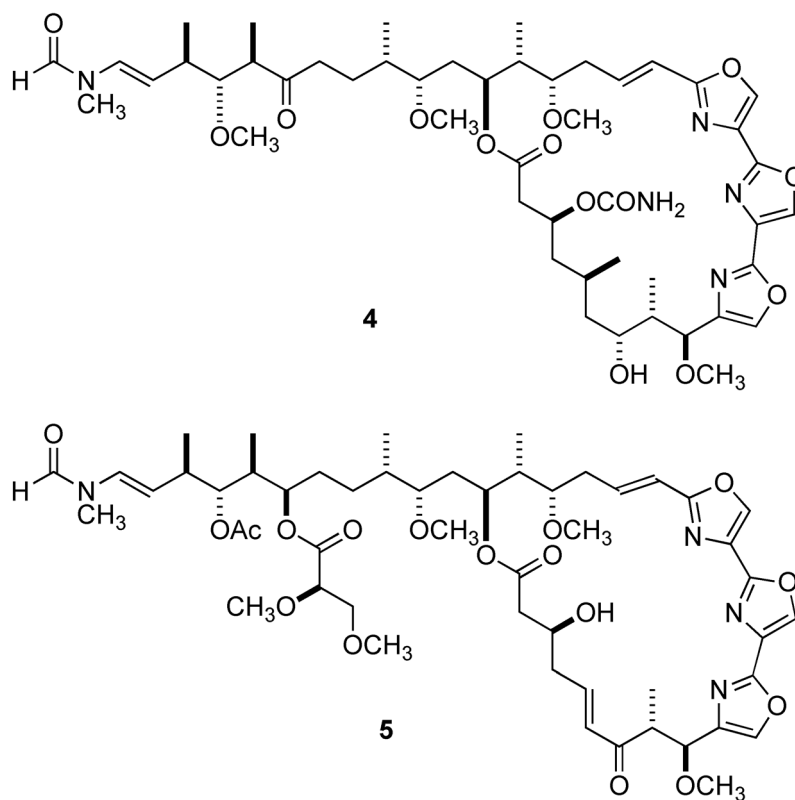
Bergmann's discovery of novel nucleosides, namely, spongouridine and spongothymidine, from a Caribbean sponge in the early 1950s led to the development of an anticancer drug, Ara-C (cytarabine) (**2**) [2]. Being stimulated by this event, significant efforts toward the development of anticancer drugs from marine organisms have been made since the early 1970s, which led to the isolation of a large array of compounds having potent cytotoxicity as well as novel structures [6]. More than 30 MNPs and their derivatives have entered clinical trials as anticancer drugs. Finally, yondelis/ectenascidin 734 (**3**), discovered from a Caribbean tunicate, has been approved for treatment of soft tissue sarcoma in the European Union in 2007; however, clinical trials of several MNPs have been discontinued [7].

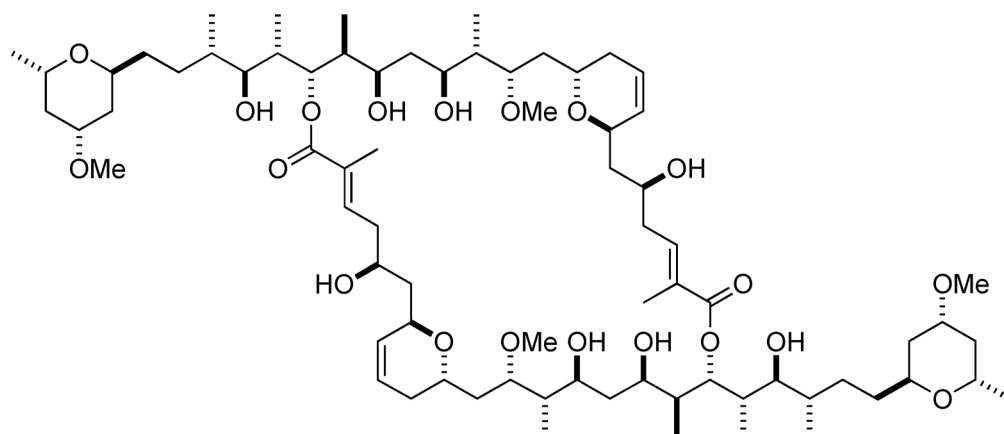


For nearly 30 years, we have been involved in the discovery of antitumor leads from Japanese marine invertebrates, especially sponges and tunicates, by employing cell-based and enzyme inhibition assays, which resulted in the isolation of a number of structurally novel and highly potent cytotoxic metabolites [8]. Although none of them have entered clinical trials, a handful of compounds whose modes of action have been elucidated are used as research tools (reagents).

Kabiramide C (**4**), isolated from egg masses of a nudibranch of the genus *Hexabranhus* (Mollusca; Opisthobranchia), is an unprecedented trisoxazole-containing macrolide with polyketide

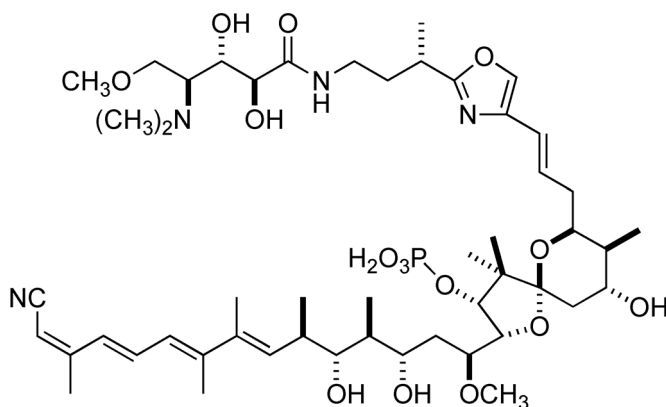
side chain [9]. Later, mycalolide B (**5**), a macrolide with essentially the same framework, was isolated from sponges of the genus *Mycale* [10]. These compounds showed strong cytotoxicity along with potent antifungal activity, but their *in vivo* antitumor activity was disappointing due to high toxicity. The mode of action study revealed that these macrolides bind to G-actin in a molar ratio of 1:1, thereby inhibiting actin polymerization as well as sequestering liberated actin monomers, resulting in depolymerization of actin filaments [11]. Recently, the crystal structure of the complex between kabiramide C and G-actin was solved [12]; the polyketide side chain, particularly the *N*-methyl vinylformamide group, plays a crucial role in binding to the actin. Interestingly, kabiramide C mimics the action of gelsolin, an actin-binding protein. An unusual macrodiolide, bistheonellide A (**6**) derived from a marine sponge *Theonella* sp., showed biological activities similar to those of the trisoxazole-containing macrolides. In fact, it binds to G-actin in a molar ratio of 1:2 [13]. A similar mode of action was observed for several MNPs [12].





6

Calyculin A (**7**) is an extraordinary metabolite of polyketide/non-ribosomal peptide synthase isolated from the Japanese marine sponge *Discodermia calyx*. It is not only highly cytotoxic (IC_{50} 0.74 ng/ml against L1210 leukemia cells) but also strongly antifungal. Calyculin A showed potent *in vivo* antitumor activity (T/C 246 % at 15 μ g/kg against Ehrlich ascites tumor cells; mice); however, unfortunately, it is highly toxic to mice [14]. Later, calyculin A was demonstrated to be a potent cancer promoter which was then found to be due to selective inhibition of protein phosphatases (PPs) 1 and 2A [15]. Reversible serine/threonine phosphorylation regulates numerous biological functions, including cellular signal transduction. Therefore, selective inhibitors of protein kinases or PPs are important tools for life science research. In fact, natural protein PP inhibitors have contributed greatly to our understanding of biological processes [16]. Structure–activity relationship study using natural and chemically transformed calyculins disclosed that 17-phosphate, 11- and 13-hydroxyl, and hydrophobic tetraene moieties are all necessary for binding to PP1 [17]. This finding was also supported by the crystal structure of PP1/calyculin A complex where calyculin A binds to the catalytic region of the enzyme [18].



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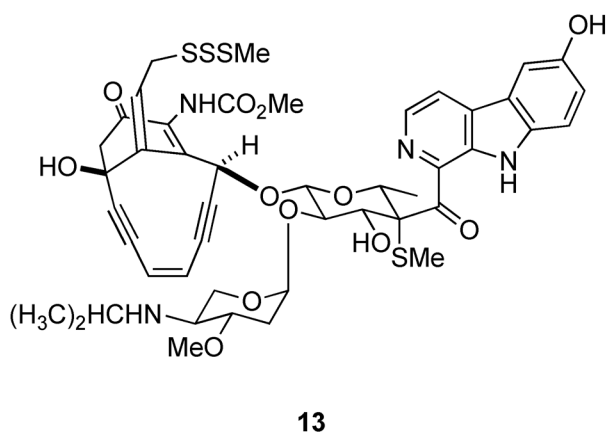
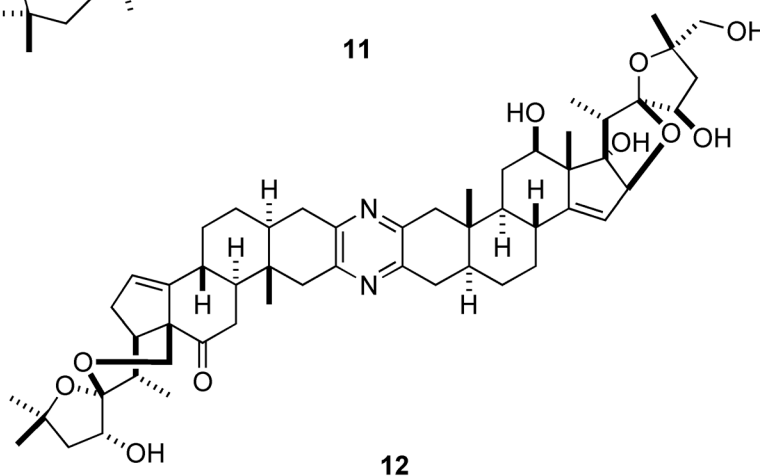
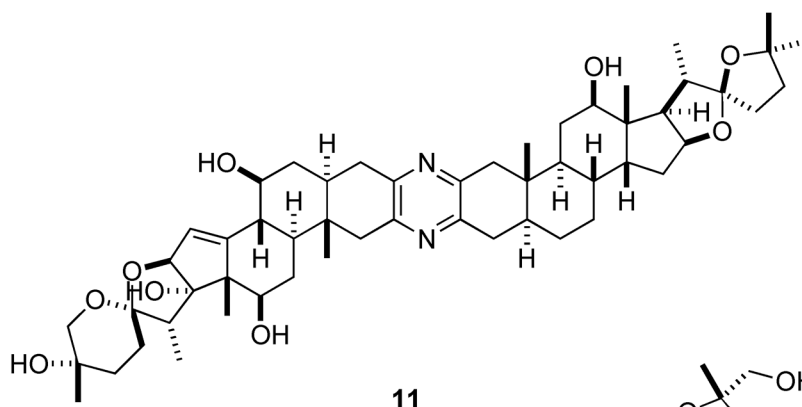
13-Deoxytedanolide (13-DT) (**8**), an 18-membered macrolide isolated from marine sponges of the genus *Mycale*, shows not only potent cytotoxicity (IC_{50} < 0.14 ng/ml against P388 leukemia cells)

Chemical structure of compound **8**, a complex macrocyclic ketone. The structure features a large ring with multiple stereocenters indicated by wedges and dashes. Key functional groups include a terminal alkene, a methyl ketone, a hydroxyl group, a methoxy group (OMe), and several other carbonyl groups. The molecule is labeled **8** at the bottom.

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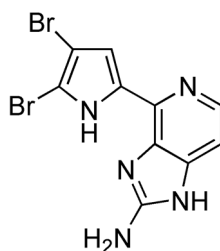
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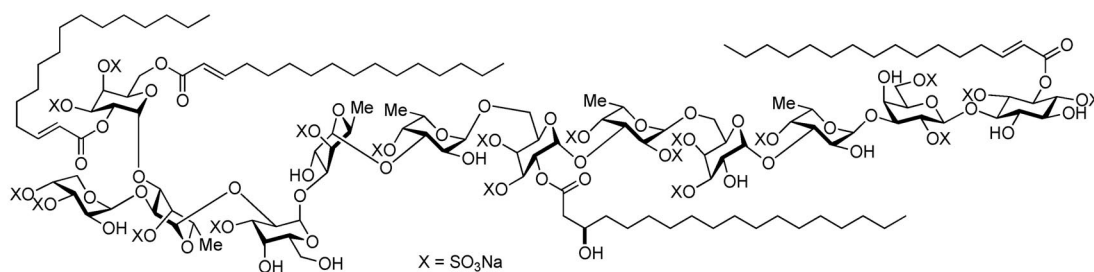


ENZYME INHIBITORS

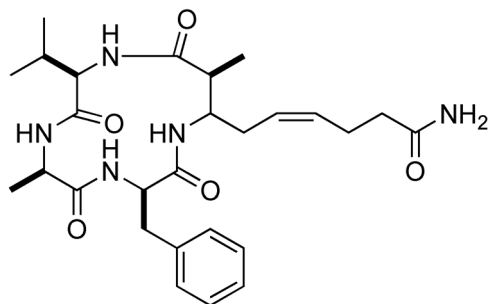
Matrix metalloproteinases (MMPs), a family of Zn-containing endopeptidases, have been a target for cancer chemotherapy [28]. Among a number of inhibitors we isolated from marine sponges, the most interesting inhibitor is ageladine A (**14**), a fluorescent oroidin derivative, discovered from the marine sponge *Agelas nakamurai*; it not only inhibits various MMPs at low concentration but also angiogenesis [29].

**14**

Telomerase is a reverse transcriptase that adds telomere sequences (TTAGGG) onto the 3'-end of chromosomes and expressed in 90 % human tumor cells, but not in normal cells [30]. Therefore, its inhibitors are potential anticancer drugs; indeed, some synthetic inhibitors have been successful in cancer chemotherapy. Axinelloside A (**15**), isolated from the marine sponge *Axinella imfundebula*, is an unprecedented highly sulfated lipopolysaccharide, comprised of 12 sugars, 19 sulfates, and 4 fatty acids [31]. It inhibits telomerase at IC_{50} 2 μ g/ml; perhaps sulfate groups are important for the activity.

**15**

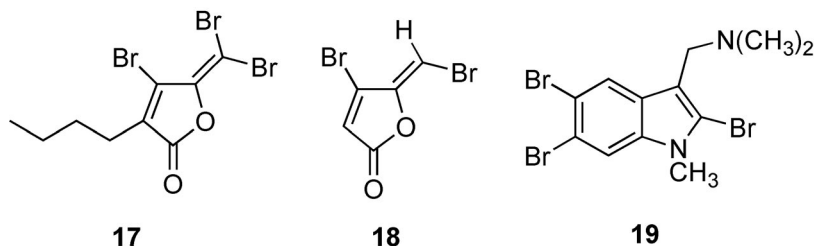
Histone deacetylases (HDACs) remove acetyl groups from acetylated lysine residues of histone, and its inhibitors show antitumor activity and some inhibitors have entered clinical trials [32]. Azumamide A–E [azumamide A (**16**)] are cyclic tetrapeptides containing a new amino acid isolated from the marine sponge *Mycale izuensis* collected in southern Japan [33]. They inhibit a human leukemia HDAC at low concentrations. Importantly, they inhibit vascular formation in ES cells.

**16**

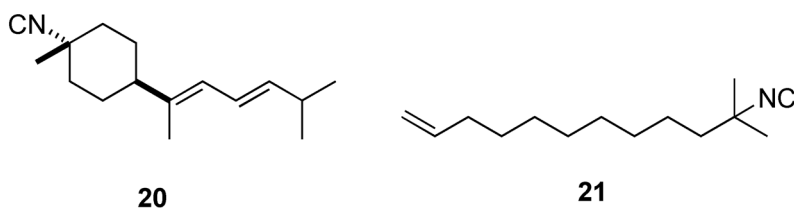
NONTOXIC ANTIFOULANTS

Macrofouling on man-made submersible structures, namely, ships' hulls, cooling systems for power plants, and aquaculture facilities, cause enormous economic losses. Combats against fouling organisms started when humans built boats. However, protection of ships' hulls from fouling goes back more than 2000 years. Various toxic materials have been used for control of fouling organisms, among which organotin compounds, especially tributyltin (TBT) and tributyltin oxide (TBTO), are remarkably effective. However, the adverse effects of organotin compounds on marine organisms became evident in Europe and other regions by the mid-1980s, and the International Maritime Organization (IMO) decided to ban TBT/TBTO coating on ships completely from 2008. Cupuric oxide-based paints and bio-icide-based "booster" paints have been introduced as alternative paints; however, these paints also showed environmental problems in recent years [34]. Therefore, development of nontoxic or environmentally friendly antifoulants has been urgently required.

An approach to nontoxic antifouling strategy is to apply chemical defense of sessile marine organisms; they maintain their body surfaces clean. In fact, such important antifouling compounds as furanones, pukalide, and renillafulin were isolated from sessile organisms [35]. Furanones (e.g., **17**), isolated from the Australian red alga *Delisea pulchra*, are potent bacterial quorum sensing inhibitors, but the mechanism of their antifouling activity against macrofouling is unknown. Furanone (**18**), a synthetic analog, is under development for antifoulants and other products [36]. 2,5,6-Tribromo-1-methylgramine (**19**) found from the Japanese bryozoan *Zoobotryon pellucidum* showed remarkable antifouling activity, and more than 150 analogs and derivatives were evaluated for antifouling activity; 5,6-dichloro-1-methylgramine was selected as a candidate for nontoxic antifoulant. Since the panel tests showed promising results, a large-scale, 5-year field test has been under way in Japan [37].



The Fusetani Biofouling Project explored nontoxic antifouling compounds from sessile marine invertebrates using "cyprid assay", which resulted in isolation of more than 65 antifouling compounds, including 35 new compounds. The most promising antifouling compounds we discovered are sesqui- and diterpenoid isocyanides isolated from the marine sponge *Acanthella cavernosa* and nudibranchs of the family Phyllidiidae [38]. 4-Isocyanotheonellin (**20**) was selected as a model compound for development of nontoxic antifoulants, and more than 70 isocyanotheonellin analogs and much simpler isocyanides were synthesized and evaluated by the cyprid assay. Finally, a simple alkyl isocyanide (**21**) was selected and evaluated its antifouling activity in the field. A preliminary field test showed promising results [39].



CONCLUSION

Marine invertebrates, especially sponges and tunicates, are rich sources of bioactive small molecules applicable to biotechnological industry. However, the most serious obstacle for their industrial applications is the supply problem (the problem of supplying sufficient material for drug development and others); most of the promising MNPs are contained at very small concentrations, usually 10^{-4} – 10^{-6} % of wet tissue, and have complex structures [40]. Thus, detailed studies on structure–activity relationships, modes of action, and others required for industrial applications cannot be carried out. How do we solve this problem? Chemical synthesis is not mature yet to synthesize kilogram-scale of complex molecules; perhaps one of the solutions is “diverted total synthesis” [41]. Aquaculture of sponges, bryozoans, and tunicates has been attempted to produce anticancer drugs, which is not realistic for large-scale production of drugs [42]. Perhaps cloning and heterologous expression of biosynthetic enzymes of drug candidates is most promising [43], since many drug candidates seem to be produced by symbiotic micro-organisms that are uncultured [44].

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