

International Journal of Life Sciences and Review (IJLSR)

Received on 28 June, 2015; received in revised form, 28 July, 2015; accepted, 29 July, 2015; published 31 July, 2015

Document heading

doi: 10.13040/IJPSR.0975-8232.IJLSR.1 (7).238-42

MARINE DRUGS FROM SPONGES AND THEIR USES – A REVIEW

Meenakshi Mohan*

BDS, Saveetha Dental College, Chennai, Taminadu, India

ABSTRACT: The marine world is largely unexplored that harbors most of the biodiversity. In recent years, marine natural products have yielded a considerable number of drug candidates. Marine microorganisms, whose genetic and biochemical diversity became a rich source of novel chemical entities for the discovery of more effective drugs. Marine microbes specially marine sponges are playing a unique contribution for human health and well-being. In addition to the primary metabolites (amino acids, nucleotides and vitamins), they also contribute many secondary metabolites which constitute 50% of the pharmaceuticals. Drugs derived from the marine natural products are being developed for treating cancers, immune suppressive disorders and resistant microbial species. The need to augment production of these marine compounds to prepare various drugs through tissue culture and mariculture and their uses in human population has been stressed on this article.

Keywords: Marine Sponges, Pharmaceuticals, Microbes, Metabolites.

Correspondence to Author:

Meenakshi Mohan

BDS, Saveetha Dental College, Chennai, Taminadu, India

E-mail: drmeena.mohan23@gmail.com

INTRODUCTION: Man has taken advantage of nature's ability to produce remedies to treat infection, inflammation, pain, and many other diseases. The oceans are a rich source of both biological and chemical diversity. They cover more than 70% of the earth's surface and contain more than 200,000 described species¹. The first living organisms appeared in the sea more than 3500 million years ago and evolutionary development has equipped many marine organisms with the appropriate mechanisms to survive in hostile and extreme conditions in terms of temperatures, salinity and pressure, as well as overcoming the effects of mutation, and pathogens^{2,3}. A relatively small number of marine organisms have already yielded thousands of novel chemical compounds⁴.

It is estimated that several species of marine microorganisms are yet to be discovered and described⁵. The history of using marine products for therapeutics began from the Chinese seaweed-based recipes for a number of disorders such as pain, abscesses, menstrual difficulties and cancer⁶. The marine sponges and other microorganisms has been a vast source of natural compounds covering a wide range of bioactivity such as photoprotective, antihelmintic, antibacterial, anticoagulant, antifungal, anti-inflammatory, antimalarial, antiprotozoal, antituberculosis, antiviral and other miscellaneous mechanisms of action^{7,8}. Sponges, belongs to the phylum Porifera and are among the oldest multicellular organisms and show relatively little differentiation and tissue coordination^{9,10}.

Marine sponges are sessile invertebrates with a wide variety of colours, shapes, and consistencies. Sponges have strategies to defend themselves against foreign prokaryotic and eukaryotic organisms, by production of secondary metabolites that repel them^{11,12}. In fact, marine sponges are

	<p>QUICK RESPONSE CODE</p>
	<p>DOI: 10.13040/IJPSR.0975-8232.IJLSR.1(7).238-42</p>
<p>Article can be accessed online on: www.ijlsr.com</p>	
<p>DOI link: http://dx.doi.org/10.13040/IJPSR.0975-8232.IJLSR.1(7).238-42</p>	

among the richest sources of interesting chemicals produced by marine organisms.

Antibacterial Activity:

This resistance has rapidly spread, and the infections caused by *Staphylococcus aureus* and other resistant strains of pathogenic bacteria are currently a considerable problem. Even vancomycin, which was the last resource for the

treatment of infections by methicillin-resistant *S. aureus* (MRSA), recently has been rendered ineffective¹³. Thus, the discovery and development of new antibiotics has become a high priority in biomedical research. Marine sponge crude extracts present a high incidence of antibacterial activity against terrestrial pathogenic bacteria^{14, 15}.

TABLE 1: ANTIBACTERIAL PROPERTY OF FEW MARINE SPONGES

Compound	Compound class	Species/order	Reference
Discodermins B, C, and D	Cyclic peptide	Discodermia kiiensis/ Lithistida	(16)
Topsentasterol sulfates A–E	Sulfated sterol	Topsentia sp./Halichondrida	(17)
Arenosclerins A, B, and C	Alkylpiperidine Alkaloid	Arenosclera brasiliensis/ Haplosclerida	(18)
Axinellamines B–D	Imidazo-azoloimidazole Alkaloid	Axinella sp./Halichondrida	(19)

Antiinflammatory Compounds:

Acute inflammations in the human body can result from microbial infection, physical damage, or chemical agents²⁰. The antiinflammatory sponge products are selective inhibitors of specific enzymes of a range of diseases, like psoriasis or

rheumatic arthritis. The currently used nonsteroidal antiinflammatory drugs often fail to control the disease and present important side effects such as risk of gastrointestinal bleeding and renal complications²¹.

TABLE 2: ANTI INFLAMMATORY PROPERTY OF FEW MARINE SPONGES

Compound	Compound class	Species/order	Reference
Manoalide	Cyclohexane sesterterpenoid	Luffariella variabilis/ Dictyoceratida	(22)
Dysidotronic acid	Drimane sesquiterpenoid	Dysidea sp./ Dendroceratida	(23)
Ircinin-1 and -2	Acyclic sesterterpenoid	Ircinia oros/ Dictyoceratida	(24)
Petrosaspongiolides M-R	Cheilantane sesterterpenoid	Petrosaspongia nigra/ Dictyoceratida	(25)
Spongidines A-D	Pyridinium alkaloid	Spongia sp./ Dictyoceratida	(26)
Topsentin	Bis-indole alkaloid	Topsentia genitrix/ Halichondrida	(27)

Antimalarial Compounds: Several sponge-derived antimalarial compounds have been discovered during the last decade. New antimalarial drugs are needed to cope with the increasing

number of multidrug-resistant Plasmodium strains that cause malaria. Plasmodium falciparum has become resistant against chloroquinone, pyrimethamine, and sulfadoxine²⁸.

TABLE 3: ANTI MALARIAL PROPERTY OF FEW MARINE SPONGES

Compound	Compound class	Species/order	Reference
Axisonitrile-3	Sesquiterpenoid isocyanide	Acanthella klethra/ Halichondrida	(28)
Manzamine A	Manzamine alkaloid, diterpene isocyanates	Haplosclerida Cymbastela hooperi/ Halichondrida	(29)
Kalihinol A	Isonitril-containing kalihinane diterpenoid	Acanthella sp./ Halichondrida	(30)

Antiviral Compounds:

Sponges are also a rich source of compounds with antiviral properties. The high number of HIV-

inhibiting compounds discovered does not reflect greater potential of sponges to fight AIDS

compared with other viral diseases, but rather the interest of many researchers. The strong focus on screening for anti-HIV activity has led to discovery

of numerous compounds, but the mechanism of inhibition is still poorly characterized.

TABLE 4: ANTI VIRAL PROPERTY OF FEW MARINE SPONGES

Compound	Compound Class	Species/Order	Reference
Dragmacidin F	Indole alkaloid	Halicortex	(31)
Papuamides C and D	Cyclic peptide	Theonella mirabilis, T. swinhoei/Lithistida	(32)
Mololipids	Tyramine lipid	Verongida	(33)
Haplosamates A and B	Sulfamated steroid	Xestospongia	(34)
Hamigeran B	Phenolic macrolide	Hamigera tarangaensis/Poecilosclerida	(35)

Immunosuppressive Compounds: In addition to their potential for treatment of cancer, nitric oxide synthetase inhibitors down regulate T-cells are, suppressing the immune system, and they diminish

the fierceness of migraine attacks. Immune system suppression is desired in cases of hypersensitivity to certain antigens (e.g., allergies) or organ transplantations.

TABLE 5: IMMUNOSUPPRESSIVE PROPERTY OF FEW MARINE SPONGES

Compound	Compound Class	Species/Order	Reference
Simplexides	Glycolipid	Plakortis simplex/Homosclerophorida	(36)
Polyoxygenated Sterols	Sterol	Dysidea sp./Dendroceratida	(37)
Contignasterol	Oxygenated Sterol	Petrosia contignata/Haplosclerida	(38)

Muscle Relaxants: Disturbances in neuromuscular communication resulting from stress cause permanent muscle activation.

TABLE 6: MUSCLE RELAXANT PROPERTY OF FEW MARINE SPONGES

Compound	Compound Class	Species/Order	Reference
1-Methylisoguanosine	Nucleoside analogue	Tedania digitata / Poecilosclerida	(39)
Xestospongin C	Macrocyclic bis-oxaquinolizidine	Haplosclerida	(40)

CONCLUSION: Sponge-microbial associations are found to be very specific in the production of particular bioactive compounds. However, the mutual mechanism between host and the microbial associate, in compound production is not well understood. The easiest and best way for commercial production of these compounds are either by culturing the host and/or the associated microbe under controlled conditions. But, the ability of the symbiont to produce the compound consistently for several generations in culture media has to be tested and standardized. Understanding the optimum ecological conditions which drives the sustainable production of bioactive compounds from sponges and their microbial associates would help in formulating

various production strategies. Adopting different cultivation strategies and metagenomic approaches would be the need of the hour in discovering new genes, enzymes and natural products and in enhancing the commercial production of marine drugs.

REFERENCES:

1. Winston, J.E.,: The systematist' perspective. In: Biomedical Importance of Marine Organisms. D.G. Fautin, ed., California Academy of Sciences, San Francisco. 1988; pp. 1-6.
2. Macdougall, J.D. A short history of Planet Earth; John Wiley Eds.: New York, 1996; p. 5.

3. Argulis, L., Schwartz, K. Five Kingdoms, an illustrated guide to the phyla of life on Earth; W.H. Freeman & Company: New York, 1982; pp. 16-17.
4. Ireland, C.M., B.R. Copp, M.P. Foster, L.A. McDonald, D.C. Radisky and J.C. Swersey: Biomedical potential of marine natural products. In: Marine Biotechnology, Vol. 1: Pharmaceutical and Bioactive Natural Products. Plenum Press, New York. 1993; pp. 1-43.
5. Malakoff, D.: Extinction on the high seas. *Scienc*, 1997; 277, 486-488.
6. Ruggieri, G.D. Drugs from the sea. *Science* 1976, 194, 491-496.
7. Pallela, R.; Yoon, N.-Y.; Kim, S.K. Anti-photoaging and photoprotective compounds derived from marine organisms. *Mar. Drugs* 2010, 8, 1189–1202.
8. Mayer, A.M.; Rodríguez, A.D.; Berlinck, R.G.; Hamann, M.T. Marine pharmacology in 2005–6: Marine compounds with anthelmintic, antibacterial, anticoagulant, antifungal, anti-inflammatory, antimalarial, antiprotozoal, antituberculosis, and antiviral activities; affecting the cardiovascular, immune and nervous systems, and other miscellaneous mechanisms of action. *Biochim. Biophys. Acta* 2009, 1790, 283–308.
9. Bergquist, P.R. In *Sponges*. Hutchinson and Co. Ltd., London, UK, 1978; p. 268.
10. Simpson, T.L. In *The cell biology of sponges*. Springer-Verlag, New York, NY. 1972; p. 662.
11. Sarma, A.S.; Daum, T. and Müller, W.E.G. in *Secondary Metabolites from Marine Sponges—Akademie gemeinnütziger Wissenschaften zu Erfurt*. Ullstein-Mosby Verlag, Berlin. 1993; p. 168.
12. Proksch, P. *Toxicon*. 1994; 32, 639-655.
13. Rice, L.B. *Am. J. Infect. Control*. 2006; 34(5), S11-19.
14. Amade, P.H.; Pesando, D. and Chevolut, L.. *Oceanography And Marine Biology, An annual review*. *Mar. Biol.*, 1986; 70, 223-228.
15. Amade, P.G.; Chariou, G.; Baby, C. and Vacelet, J. Antibiotic-resistant bacteria inhibited by extracts and fractions from Brazilian marine sponges. *Mar. Biol.*, 1987; 94, 271-275.
16. Matsunaga S, Fusetani N, Konosu S. Bioactive marine metabolites, VII: structures of discodermins B, C, and D, antimicrobial peptides from the marine sponge *Discodermia kiiensis*. *Tetrahedron Lett* 1985; 26, 855–856.
17. Fusetani N, Matsunaga S, Matsumoto H, Takebayashi Y. Cyclotheonamides, potent thrombin inhibitors, from a marine sponge *Theonella* sp. *J Am Chem Soc*. 1990 112, 7053–7054.
18. Torres YR, Berlinck RGS, Nascimento GGF, Fortier SC, Pessoa C, De Moraes MO. Antibacterial activity against resistant bacteria and cytotoxicity of four alkaloid toxins isolated from the marine sponge *Arenosciera brasiliensis*. *Toxicon*. 2002; 40, 885–891.
19. Urban S, De Almeida Leone P, Carroll AR, Fechner GA, Smith J, Hooper JNA, Quinn RJ. Axinellamines A–D, novel imidazo-azolo-imidazole alkaloids from the Australian marine sponge *Axinella* sp. *J Org Chem* 1999; 64, 731–735/190.
20. Tan P, Luscinskas FW, Homer-Vanniasinkam S. Cellular and molecular mechanisms of inflammation and thrombosis. *Eur J Endovasc Surg* 1997; 17, 373–389.
21. De Rosa S. “Mediterranean marine organisms as source of new potential drugs”. In: *Natural Products in the New Millennium: Prospects and Industrial Applications*, Rauter A, Palma FB, Justino J, Araujo ME, Santos SP, eds. (The Netherlands: Kluwer Academic Publishers) 2002; pp 441461.
22. Bennet CF, Mong S, Clark MA, Kruse LJ, Crooke ST. Differential effects of manoilide on secreted intracellular phospholipases. *Biochem Pharmacol*. 1987; 36, 2079–2086.
23. Giannini C, Debitus C, Posadas I, Paya M, DAuria MV. Dysidotronic acid, a new and selective human phospholipase A2 inhibitor from the sponge *Dysidea* sp. *Tetrahedron Lett*. 2000; 41, 3257–3260.
24. Cimino G, De Stefano S, Minale L, Fattorusso E. Ircinin 1 and 2, linear sesterterpenes from the marine sponge *Ircinia oros*. *Tetrahedron*. 1972; 28, 333–341.
25. Randazzo A, Debitus C, Minale L, Pastor PG, Alcaraz MJ, Paya M, Gomez-Paloma L. Petrosaspongiolides M-R: new potent and selective phospholipase A2 inhibitors from the New Caledonian marine sponge *Petrosaspongia nigra*. *J Nat Prod*. 1988a; 61, 571–575.
26. Jacobs RS, Koehn FE, Gunasekera SP. Topsentin, a unique phospholipase A2 inhibitor [abstract]. Presented at the Japan–US Seminar on Bioorganic Marine Chemistry 1994.
27. De Carvalho MS, Jacobs RS. Two-step inactivation of bee venom phospholipase A2 by scalaradial. *Biochem Pharmacol*. 1991; 42, 1621–1626.
28. Angerhofer CK, Pezzuto JM, König GM, Wright AD, Stichter O. Antimalarial activity of sesquiterpenes from the marine sponge *Acanthella klethra*. *J Nat Prod*. 1992; 55, 1787–1789.
29. König GM, Wright AD, Angerhofer CK. Novel potent antimalarial diterpene isocyanates, isothiocyanates, and isonitriles from the tropical marine sponge *Cymbastela hooperi*. *J Org Chem*. 1996; 61, 3259–3267.
30. Miyaoka H, Shimomura M, Kimura H, Yamada Y, Kim H-S, Wataya Y. Antimalarial activity of kalahinol A and new relative diterpenoids from the Okinawan sponge, *Acanthella* sp. *Tetrahedron*. 1998; 54, 13467–13474.
31. Cutignano A, Bifulco G, Bruno I, Casapullo A, Gomez-Paloma L, Riccio R. Dragmacidin F: a new antiviral bromoindole alkaloid from the Mediterranean sponge *Halicortex* sp. *Tetrahedron* 2000; 56, 3743–3748.
32. Ford PW, Gustafson KR, McKee TC, Shigematsu N, Maurizi LK, Pannell LK, Williams DE, De Silva ED, Lassota P, Alien TM, Van Soest R, Andersen RJ, Boyd MR. Papuamides A–D, HIV-inhibitory and cytotoxic depsipeptides from the sponges *Theonella mirabilis* and *Theonella swinhoei* collected in Papua

- New Guinea. *J Am Chem Soc.* 1999; 121, 5899–5909.
33. Ross SA, Weete JD, Schinazi RF, Wirtz SS, Tharnish P, Scheuer PJ, Hamann MT. Mololipids, a new series of anti-HIV bromotyramine-derived compounds from a sponge of the order Verongida. *J Nat Prod.* 2000; 63, 501–503.
 36. Costantino V, Fattorusso E, Mangoni A, Di Rosa M, Ianaro A. Glycolipids from sponges, VII: simplexides, novel immunosuppressive glycolipids from the Caribbean sponge *Plakortis simplex*. *Bioorg Med Chem.* 1999; Lett 9, 271–276.
 37. de Leone PA, Redburn J, Hooper JNA, Quinn RJ. Polyoxygenated Dysidea sterols that inhibit the binding of [I125] IL-8 to the human recombinant IL-8 receptor type A. *J Nat Prod.* 2000; 63, 694–697.
 38. Takei M, Burgoyne DL, Andersen RJ. Effect of contignasterol on histamine release induced by antiimmunoglobulin E from rat peritoneal mast cells. *J Pharm Sci.* 1994; 83, 1234–1235
 34. Qureshi A, Faulkner DJ. Haplosamates A and B: new steroidal sulfamate esters from two haplosclerid sponges. *Tetrahedron.* 1999; 55, 8323–8330.
 35. Wellington KD, Cambie RC, Rutledge PS, Bergquist PR. Chemistry of sponges, 19: Novel bioactive metabolites from *Hamigera tarangaensis*. *J Nat Prod.* 2000; 63, 79–85.
 39. Quinn RJ, Gregson RP, Cook AF, Bartlett AF. Isolation and synthesis of 1-methylisoguanisine, a potent pharmacologically active constituent from the marine sponge *Tedania digitata*. *Tetrahedron Lett.* 1980; 21, 567–568.
 40. De Smet P, Parys JB, Callewaert G, Weidema AF, Hill E, De Smedt H, Erneux C, Sorrentino V, Missiaen L. Xestospongins C is an equally potent inhibitor of the inositol 1,4,5-triphosphate receptor and the endoplasmic-reticulum Ca²⁺ pumps. *Cell Calcium.* 1999 26, 9–13

How to cite this article:

Mohan M: Marine Drugs from Sponges and Their Uses – A Review. *Int J Life Sci Rev.* 2015; 1(7) 238-42: .doi:10. 13040/ IJPSR.0975-8232.IJLSR.1(7).238-42.

All © 2015 are reserved by International Journal of Life Sciences and Review. This Journal licensed under a Creative Commons Attribution-NonCommercial-ShareAlike 3.0 Unported License.

This article can be downloaded to **ANDROID OS** based mobile. Scan QR Code using Code/Bar Scanner from your mobile. (Scanners are available on Google Playstore)