







Biological activities of marine invertebrates extracts from the northeast brazilian coast

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Abstract

This paper reports the *in vitro* antiproliferative effects, antiprotozoal, anti-herpes and antimicrobial activities of 32 organic extracts of 14 marine sponges and 14 corals collected in northeast Brazilian coast. The ethanolic extracts of the sponges *Amphimedon compressa* and *Tedania ignis*, and the acetone extract of *Dysidea* sp. showed relevant results concerning the antiproliferative effects against A549, HCT-8, and PC-3 cell lines by sulforhodamine B assay, but also low specificity. Concerning the antiprotozoal screening, the ethanolic extract of *Amphimedon compressa* and the acetone and ethanolic extracts of *Dysidea* sp. were the most active against *Leishmania amazonensis* and *Trypanosoma cruzi* expressing β -galactosidase in THP-1 cells. In the preliminary anti-HSV-1 (KOS strain) screening, the ethanolic extracts of the sponges *Amphimedon compressa*, *Haliclona* sp. and *Chondrosia collectrix* inhibited viral replication by more than 50%. The most promising anti-herpes results were observed for the ethanolic extract of *Haliclona* sp. showing high selective indices against HSV-1, KOS and 29R strains (SI > 50 and > 79, respectively), and HSV-2, 333 strain (IS > 108). The results of the antibacterial screening indicated that only the ethanolic extract of *Amphimedon compressa* exhibited a weak activity against *Enterococcus faecalis*, *Pseudomonas aeruginosa* and *Escherichia coli* by the disk diffusion method. In view of these results, the extracts of *Amphimedon compressa*, *Tedania ignis* and *Dysidea* sp. were selected for further studies aiming the isolation and identification of the bioactive compounds with antiproliferative and/or antiprotozoal activities. The relevant anti-herpes activity of the ethanolic extract of *Haliclona* sp. also deserves special attention, and will be further investigated.

Keywords: marine invertebrates, antiproliferative effects, anti-herpes activity, antiprotozoal activity, antimicrobial activity.

Atividades biológicas de extratos de invertebrados marinhos da costa nordeste do Brasil

Resumo

Este artigo reporta as atividades *in vitro* antiproliferativa, antiprotozoária, anti-herpética e antimicrobiana de 32 extratos orgânicos provenientes de 14 esponjas marinhas e 14 corais coletados no litoral nordestino brasileiro. Os extratos etanólicos das esponjas *Amphimedon compressa* e *Tedania ignis*, e o extrato acetônico de *Dysidea* sp. demonstraram resultados promissores em relação aos efeitos antiproliferativos frente as linhagens celulares A549, HCT-8, PC-3 pelo método da sulforrodamina B, mas sem especificidade. Em relação à atividade antiprotozoária, os extratos etanólico de *Amphimedon compressa* e acetônico e etanólico de *Dysidea* sp. apresentaram atividade contra *Leishmania amazonensis* e *Trypanosoma cruzi* através do método de expressão de β -galactosidase em células THP-1. Na investigação preliminar de atividade antiviral frente ao vírus *Herpes simplex* tipo 1 (cepa KOS), os extratos etanólicos das esponjas *Amphimedon compressa*, *Haliclona* sp. e *Chondrosia collectrix* inibiram mais de 50% da replicação viral. O extrato etanólico da esponja

Haliclona sp. demonstrou resultados promissores para atividade anti-herpética com altos índices de seletividade para as cepas KOS (IS >50) e 29R (IS >79) frente ao VHS-1 e cepa 333 (IS >108) frente ao VHS-2. O extrato etanólico da esponja *Amphimedon compressa* exibiu uma pequena atividade contra *Enterococcus faecalis*, *Pseudomonas aeruginosa* and *Escherichia coli* pelo método de difusão em disco. De acordo com os resultados apresentados, os extratos das esponjas *Amphimedon compressa*, *Tedania ignis* e *Dysidea* sp. serão selecionados para futuros estudos de isolamento e identificação dos compostos bioativos para as atividades antiproliferativa e antiprozoária. O extrato etanólico de *Haliclona* sp. será investigado por possuir atividade relevante anti-herpética.

Palavras-chave: invertebrados marinhos, atividade antiproliferativa, atividade anti-herpética, atividade antiprozoária, atividade antimicrobiana.

1. Introduction

Most of the natural products used in therapeutics come from terrestrial organisms. However the marine environment allowed a new approach for the discovery of chemical molecules and biological activities. The marine habitat is a prolific source of bioactive secondary metabolites with potential to treat several diseases (Montaser and Luesch, 2011). A comparative analysis performed by Kong and coworkers showed that marine natural products are superior to terrestrial natural products in terms of chemical novelty and significant bioactivity. The analysis, which compared molecular scaffolds reported in the Dictionary of Natural Products (DNM) to those in the Dictionary of Marine Natural Products (DMNP), showed that approximately 71% of the molecular scaffolds in the DMNP were exclusively found in marine organisms (Kong et al., 2010).

Despite the large number of natural products isolated from marine sources in the last decade that show high bioactivity for a variety of diseases, only few have the chance to get to the end of drug discovery pipeline (Blunt et al., 2010, 2017). Nevertheless, worldwide bioprospecting efforts have not ceased, and marine invertebrates are amongst the top group of organisms that have contributed for this fact with a larger number of new molecules entries in Natural Products libraries (Jha and Xu, 2004; Blunt et al., 2008). Leal and coworkers showed that between 2000 and 2009, a great number of natural products from marine invertebrates was isolated from the phyla Porifera (47%) and Cnidaria (33%) (Leal et al., 2012).

Sponges (phylum Porifera) are primitive filter-feeders, about 9,000 species already described, but it is thought that there are more than 20,000 marine sponges. They can produce high levels of cytotoxic compounds, the majority coming from the secondary metabolism protecting them against predation, overgrowth by fouling organisms, and/or competition for space (Perdicaris et al., 2013). It was proved that marine sponges produce an enormous array of antitumor, antiviral, anti-inflammatory, immunosuppressive, antibiotic, and other bioactive molecules that have potential for therapeutic use. Many bioactive metabolites from sponges proved to be inhibitors of certain enzymes, which often mediate or produce mediators of intracellular or intercellular messengers involved in diseases pathogenesis (Sagar et al., 2010; Villa and Gerwick, 2010; Frota et al., 2012).

The phylum Cnidaria is a large, diverse and ecologically important group of marine invertebrates that includes over

11,000 existing species (Daly et al., 2007). Over 3,000 marine natural products have been described and the researches have been demonstrated that compounds produced by cnidarians are more than just toxins and venoms (Rocha et al., 2011). The class Anthozoa currently includes over 7,500 valid species (about 2/3 of all known cnidarian species) (Daly et al., 2007). Within the Anthozoa, the soft corals or octocorals are the ones, which have contributed with the highest number of promising bioactive marine compounds, although other groups, such as Actiniaria (sea anemones) and Scleractinia (hard corals), have also yielded relevant compounds (Štrukelj et al., 2000; Fontana et al., 1998; Miyaoka et al., 2006; Meyer et al., 2009).

Despite the great biodiversity present along the Brazilian coastline, an important source of potential biologically active compounds, chemical analysis and bioprospection screening of Brazilian marine organisms is still incipient (Almeida et al., 2014). To this point, the chemistry of marine natural products in Brazil has been focused on sponges, tunicates and brown algae (Berlinck et al., 2004). Concerning the chemical studies on octocorals, from the approximately 107 species reported in Brazil to the present date only twenty have been studied, thirteen of them by Brazilian research groups (Almeida et al., 2014).

So far, most of the bioprospection studies on marine organisms extracts conducted in Brazil were focused on evaluating their cytotoxic and antimicrobial activities (Seleguim et al., 2007; Monks et al., 2002; Rangel et al., 2001). In addition, the collection range of these organisms on the Brazilian coast is concentrated in the south and southeast regions of the country (Seleguim et al., 2007; Monks et al., 2002; Rangel et al., 2001). Based on these data and the urgent requirement to discover new molecules for the treatment of neglected tropical diseases, such as Chagas disease and leishmaniasis, we evaluated the *in vitro* antiproliferative, antiprozoal, anti-herpes, and antimicrobial activities of 32 organic extracts obtained from 14 marine sponges and 14 coral species from the northeastern Brazilian coast.

2. Materials and Methods

2.1. Collection of the marine organisms

The marine sponges and corals were collected in June 2016 at Ponta de Pedra beach, Goiana, Pernambuco State (07°33'38" S; 35°00'09" W) and Ponta de Mangue beach, Maragogi, Alagoas State (09°02'46" S; 35°12'12" W),

Brazil (Table 1). They were frozen and taken to the Laboratory of Natural Products in Santa Catarina State. The sponges and corals were identified by two of the authors (U.S.P. and C.D.P, respectively), and are kept in the Porifera Collection of the *Universidade Federal de Pernambuco* (UFPEPOR) and in the Cnidaria Collection of the *Centro Acadêmico de Vitória* of the *Universidade Federal de Pernambuco* (GPA). The organisms used for the analyzes are all colonial and we collected a portion of them. They have a high vegetative growth and they can regenerate the lost portion, for this reason the populations of these organisms are not affected by the sampling.

2.2. Preparation of the extracts

The extracts of marine sponges were obtained using different organic solvents. Initially, an acetone (A) extraction was performed for seven days. After, the same samples were extracted with ethanol (E) followed by dichloromethane (D). All the extracts were firstly evaporated to dryness in a rotary evaporator, kept in the freezer and submitted to an evaluation by ¹H NMR to look for chemical similarities. Pharmacological screenings were performed with all ethanolic

extracts, whereas only some acetone and dichloromethane extracts, which showed different ¹H NMR profiles to those of ethanolic extracts, were pharmacologically evaluated. The corals analyzed in this work were only extracted with ethanol for seven days.

2.3. Antibacterial assays

The microorganism strains tested came from the American Type Culture Collection (ATCC): *Enterococcus faecalis* (ATCC 29212), *Pseudomonas aeruginosa* (ATCC 27853) and *Escherichia coli* (ATCC 25922). The antibacterial activity was evaluated by the disk diffusion method as previously described by de Oliveira et al. (2005) with minor modifications. Briefly, filter paper disks (6 mm) were impregnated with 20 µl of the extracts (100 mg/mL) and then placed on Mueller-Hinton agar plates (HIMEDIA®), which were inoculated with these microorganisms according to the standard protocol described by the Clinical Laboratory Standard Institute (CLSI, 2012). The plates were incubated at 35 °C (± 1 °C) and after 18 h the diameters of the inhibition zones were measured. Filter-paper disks containing DMSO were used as negative control and no

Table 1. Marine organisms collected from Northeastern Brazilian coast.

Species	Family	Collection site
Sponges		
<i>Amphimedon compressa</i>	Niphatidae	Ponta de Pedra beach (07°33'38" S; 35°00'09" W)
<i>Chondrilla caribensis</i>	Chondrillidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Chondrosia collectrix</i>	Chondrosiidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Cinachyrella</i> sp.	Tetillidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Cladocroce caelum</i>	Chalinidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Desmaysamma anchorata</i>	Desmaciidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Dysidea etheria</i>	Dysideidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Dysidea</i> sp.	Dysideidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Echinodictyum dendroides</i>	Raspailiidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Haliclona implexiformis</i>	Chalinidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Haliclona melana</i>	Chalinidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Haliclona</i> sp.	Chalinidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Scopalina ruetzleri</i>	Scopalinidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Tedania ignis</i>	Tedaniidae	Ponta de Pedra beach (07°33'38" S; 35°00'09" W)
Corals		
<i>Agaricia humilis</i>	Agariciidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Carijoa riisei</i>	Clavulariidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Favia grävada</i>	Faviidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Millepora alcicornis</i>	Milleporidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Millepora brasiliensis</i>	Milleporidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Montastrea cavernosa</i>	Faviidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Muriceopsis sulphurea</i>	Plexauridae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Mussismilia harttii</i>	Mussidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Mussismilia hispida</i>	Mussidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Palythoa caribaeorum</i>	Zoanthidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Plexaurella grandiflora</i>	Plexauridae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Porites asteroides</i>	Poritidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Scolymia wellsi</i>	Mussidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)
<i>Siderastrea stellata</i>	Siderastreidae	Ponta de Mangue beach (09°02'46" S; 35°12'12" W)

inhibition was observed. Standard antibiotic disks were selected according to the sensitivity of the microorganism tested: Ampicillin (10 µg), Oxacillin (1 µg), Ceftazidime (30 µg) and Fluconazole (25 µg).

2.4. Antiprotozoal assays

The trypanocidal and leishmanicidal assays were performed as previously described by Lima et al. (2016). Human monocytic leukemia cell line THP-1 (ATCC TIB202) was grown in RPMI-1640 medium without phenol red (Sigma-Aldrich, St. Louis, MO, USA) supplemented with 10% fetal bovine serum (FBS, Life Technologies, USA), 12.5 mM HEPES, Glutamax (2 mM), and the antibiotics penicillin (100 U/ml) streptomycin (100 µg/ml), at 37 °C in a humidified 5% CO₂ atmosphere. *L. amazonensis* MHOM/BR/77/LTB0016 promastigotes expressing β-galactosidase were grown at 26 °C in Schneider's insect medium (Sigma Chemical, St. Louis, MO, USA) supplemented with 5% heat inactivated FBS and 2% human urine. For the antiprotozoal screening against intracellular *L. amazonensis* amastigotes, THP-1 cells (4.0x10⁴ per well) were cultivated in 96 well plates in RPMI-1640 medium supplemented as described previously, and treated with 100 ng/ml of phorbol 12-myristate 13-acetate (PMA, Sigma-Aldrich) for 72h at 37 °C in a 5% CO₂ to allow THP-1 cells differentiation into non-dividing macrophages (Schwende et al., 1996). Four days old culture promastigotes (4.0x10⁶ parasites/ml) were washed with phosphate buffered saline (PBS), pH 7.4, and incubated in RPMI-1640 medium supplemented with 10% heat-inactivated human B+ serum for 1h at 34°C to parasite opsonization. THP-1 cells were incubated with a parasite/cell ratio of 10:1 for 3h at 34 °C and 5% CO₂. Thereafter, non-internalized parasites were removed by one wash with PBS, and infected cells were incubated at 34 °C and 5% CO₂ with 180µl of complete supplemented RPMI-1640 medium for another 24h to allow the transformation of promastigotes into intracellular amastigotes. The β-galactosidase *T. cruzi* Tulahuen strain (Buckner et al., 1996) was kindly provided by the Laboratory of Cellular and Molecular Parasitology (Centro de Pesquisas René Rachou, FIOCRUZ, Belo Horizonte, MG, Brazil). Culture-derived trypomastigotes raised from infected L929 cell line were used to infect differentiated THP-1 cells (4.0x10⁴ cells/well) in 96-well microplates in a parasite:cell ratio of 3:1 and incubated overnight at 37 °C and 5% CO₂. The medium containing non-internalized parasites was removed by PBS washing and replaced with 180 µl of fresh complete medium.

Extracts were solubilized in 1% dimethyl sulfoxide (DMSO) and serially diluted (50 µg/ml to 1.56 µg/ml). Infected cell layer was treated by addition of 20 µl of each sample in triplicate followed by incubation for 48h at 34 °C and 5% CO₂. After treatment, cells were carefully washed with PBS and incubated for 16h at 37 °C with 250 µl of chlorophenol red-β-D-galactopyranoside (Sigma-Aldrich) at 100 µM and 0.1% Nonidet P-40 (Amresco, Solon, Ohio, USA). Optical densities were read at 570/630 nm in an Infinite M200 TECAN, Austria. The concentration of each

sample that reduced parasite viability by 50% (IC₅₀), when compared to untreated controls, was estimated by non-linear regression of concentration-response curves. Amphotericin B (Bristol-Myers. Squibb, Princeton, NJ, USA) and benzimidazole (Sigma-Aldrich) were used as positive controls for leishmanicidal and trypanocidal assays, respectively, and 1% DMSO as negative control. To evaluate cell toxicity, THP-1 cells were seeded (4.0 x10⁴ per well) in 96-well microplates and samples at concentration ranging from 500 µg/ml to 15.6 µg/ml or 1% DMSO (negative control) were added, and cells were incubated at 37°C and 5% CO₂ for 72h. Cell viability was evaluated by the colorimetric 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) method and the optical densities were read at 540 nm in a TECAN Infinite M200 microplate reader immediately after the dissolution of formazan crystals with DMSO. The CC₅₀ was defined as the concentration of each sample that reduced the absorbance of treated cells by 50% when compared with cell control.

2.5. Anti-herpes assays

VERO cells (from African Green monkey kidneys) were obtained from the American Type Culture Collection (ATCC: CCL81), and were grown in Eagle's minimum essential medium (MEM; Cultilab, Campinas, Brazil), supplemented with 10% FBS (Gibco, Carlsbad, CA, USA), and maintained at 37 °C and 5% CO₂ in a humidified atmosphere. HSV-1 (KOS and 29R strains, which are sensitive- and acyclovir-resistant strains, respectively, Faculty of Pharmacy, University of Rennes I, Rennes, France) and HSV-2 (333 strain, Department of Clinical Virology, Göteborg University, Göteborg, Sweden) were propagated in the permissive VERO cells. Viral stocks were titrated as based on plaque forming units (PFU), counted by plaque assay as previously described (Burlinson et al., 1992) and stored at -80 °C. To perform the anti-herpes screening, the viral plaque number reduction assay was used. Firstly, the cytotoxicity of the extracts was determined by sulforhodamine B (SRB) assay (Vichai and Kirtikara, 2006). In brief, VERO cells (2.5x10⁴ cells per well) were exposed to different concentrations of the samples for 48 h. After incubation, 10% trichloroacetic acid (TCA) was added to each well to fix the cells and then washed and stained with SRB. The protein-bound dye was dissolved in 10 mM Tris-Base [(tris(hydroxymethyl) aminomethane] solution and optical densities (OD) were read at 510 nm on a spectrophotometer Spectra Max M2 (Molecular Devices, USA). The 50% cytotoxic concentration (CC₅₀) was defined as the concentration that reduced cell viability by 50% when compared to untreated controls. Thereafter, the potential antiherpes activity was screened by plaque number reduction assay as described previously by Boff et al. (2016). Briefly, confluent cell monolayers (2.5x10⁵ cells per well) were infected with approximately 100 PFU of each virus strain for 1 h at 37°C. Treatments were performed by adding non-cytotoxic concentrations of the extracts for VERO cells after viral infection (post infection treatment). Cells were then washed with

phosphate-buffered saline (PBS) and overlaid with MEM containing 1.5% carboxymethylcellulose (CMC; Sigma-Aldrich) in the presence or absence of different concentrations of the compounds, and incubated for 48 h. Cells were fixed and stained with naphthol blue-black (Sigma-Aldrich) and viral plaques were counted by using a stereomicroscope. The results are presented as percentages of viral inhibition or by IC_{50} values, which were defined as the concentration of each sample that inhibited viral plaque formation by 50% (IC_{50}), when compared to untreated controls. The ratio between CC_{50} and IC_{50} values was calculated to obtain the selectivity index (SI) of each sample. Acyclovir (ACV, Sigma-Aldrich) was used as positive control to HSV-1 (KOS strain) and HSV-2 (333 strain) and as negative control to HSV-1 (29-R strain) replication.

2.6. Antiproliferative assays

Non-small cells of human lung cancer cells (A549, ATCC: CCL 195) were grown in Eagle's minimum essential medium (MEM; Cultilab, Campinas, Brazil); human ileocecal adenocarcinoma cells (HCT-8, Adolfo Lutz Institute, São Paulo, SP, Brazil) were grown in RPMI-1640 medium, Cultilab); and no hormone-sensitive human prostate adenocarcinoma cells (PC-3, German Collection of Microorganisms and Cell Cultures, DSMZ, Braunschweig, Germany) were cultivated in Dulbecco's Modified Eagle's Medium (DMEM, Cultilab) supplemented with 1% glutamine. All cells were supplemented with 10% FBS, and maintained at 37°C and 5% CO_2 . The antiproliferative screening was performed by sulforhodamine B assay (SRB) (Vichai and Kirtikara, 2006) as described above (item 2.5). A549, HCT-8 and PC3 cells were seeded in 96-well plates (2.5×10^4 cells per well) and exposed to

different concentrations of the extracts for 48 h. Paclitaxel was used as positive control. After the incubation period, the 50% inhibitory concentration (IC_{50}) of each compound was defined as the concentration that inhibited cell viability by 50% when compared to untreated controls.

2.7 Statistical analysis

The mean values \pm standard deviations (SD) are representative of two or three independent experiments. For the determination of IC_{50} values, nonlinear regressions of concentration-response curves were used.

3. Results

The results concerning the antiproliferative, antiprotozoal, anti-herpes and antimicrobial screenings of the organic extracts are described below, and four sponge species demonstrated antiproliferative, antiprotozoal, anti-herpes and/or antibacterial activities.

3.1. Antiproliferative effects

Thirty-two organic extracts of 28 species of marine invertebrates (14 sponges and 14 coral species) were screened for antiproliferative effects of the human cancer A549, HCT-8 and PC-3 cell lines. From them, the sponges *Amphimedon compressa* (ethanolic extract), *Dysidea* sp. (acetone extract), and *Tedania ignis* (ethanolic extract) showed the lowest IC_{50} values against the tested cancer cell lines as well as against VERO cells. Despite of the fact that VERO cells are fibroblasts derived from kidneys of healthy monkeys and, consequently, a comparison with cancer cell lines is not adequate, the obtained results seem to indicate that these sponge extracts are active but present low specificity (Table 2).

Table 2. Antiproliferative effects of marine invertebrate extracts (sponges and corals) on non-small cells of human lung cancer (A549), human colon carcinoma (HCT-8), human prostate cancer (PC-3), and Vero (healthy fibroblasts from kidneys of African Green monkey) cell lines, after 48h of treatment by the sulforhodamine B assay. Data are presented as IC_{50} values ($\mu\text{g/mL}$) and 95% confidence intervals obtained by non-linear regression.

Species	Extracts	Cell lines			
		A549	HCT-8	PC-3	VERO
Sponges		IC_{50} (95% confidence interval)			
<i>Amphimedon compressa</i>	E	8.06 (6.1 - 11.5)	7.12 (5.2 - 13.6)	8.06 (6.1 - 11.5)	9.9 (7.2 - 13.6)
<i>Chondrilla caribensis</i>	A	>500	>500	>500	>500
<i>Chondrilla caribensis</i>	E	>500	>500	>500	>500
<i>Chondrosia collectrix</i>	E	420.4 (305.5 - 578.6)	405.7 (336.4 - 489.1)	420.3 (265.2 - 566.2)	412.6 (352.8 - 482.6)
<i>Chondrosia collectrix</i>	D	476.5 (361.6 - 627.8)	>500	>500	>500
<i>Cinachyrella</i> sp.	E	>500	383.2 (269.2 - 545.4)	394.8 (183.4 - 549.8)	340.6 (291.6 - 397.9)
<i>Cladocroce caelum</i>	E	316.2 (133.99 - 400)	211.8 (184.5 - 243.2)	262.2 (236.9 - 290.3)	391.3 (347.4 - 440.6)
<i>Desmapsamma anchorata</i>	E	>500	>500	>500	>500
<i>Dysidea etheria</i>	E	273 (218.9 - 340.5)	186.3 (164.7 - 210.8)	271.8 (209.0 - 353.5)	221.6 (200.7 - 244.8)

A: acetone extract; E: ethanolic extract; D: dichloromethane extract.

Table 2. Continued...

Species	Extracts	Cell lines			
		A549	HCT-8	PC-3	VERO
Sponges		IC₅₀ (95% confidence interval)			
<i>Dysidea etheria</i>	A	>500	>500	481.9 (400.5 - 579.9)	>500
<i>Dysidea</i> sp.	A	18.70 (10.7 - 31.7)	6.40 (4.9 - 10.5)	10.00 (8.0 - 14.5)	12.20 (8.2 - 18.9)
<i>Dysidea</i> sp.	E	125.5 (91.5 - 172.1)	56.6 (47.2 - 67.96)	105 (76.7 - 154.4)	132.4 (93.4 - 153.6)
<i>Echinodictyum dendroides</i>	E	>500	>500	>500	>500
<i>Haliclona implexiformis</i>	E	434.3 (291.9 - 548.6)	>500	>500	>500
<i>Haliclona melana</i>	E	>500	>500	>500	>500
<i>Haliclona</i> sp.	E	>500	>500	>500	>500
<i>Scopalina ruetzleri</i>	E	369.5 (317.0 - 430.6)	253.1 (215.3 - 297.5)	376.3 (258.8 - 547.2)	381.5 (288.4 - 507.6)
<i>Tedania ignis</i>	E	30.2 (11.26 - 50.85)	16.5 (13.19 - 20.72)	17.8 (14.02 - 22.64)	37.9 (31.77 - 45.42)
Corals		IC₅₀ (95% confidence interval)			
<i>Agaricia humilis</i>	E				
<i>Carijoa riisei</i>	E	181.9 (108.8 - 304.0)	102.6 (89.5 - 117.7)	199.7 (96.32 - 314.1)	172.66 (106.6 - 321.1)
<i>Favia gravida</i>	E	277.9 (196.1 - 393.8)	295.7 (238.0 - 367.2)	454.1 (285.2 - 722.9)	243.1 (180.3 - 312.8)
<i>Millepora alcicornis</i>	E	>500	>500	>500	>500
<i>Millepora brasiliensis</i>	E	>500	>500	>500	>500
<i>Montastrea cavernosa</i>	E	>500	>500	>500	>500
<i>Muriceopsis sulphurea</i>	E	>500	>500	>500	>500
<i>Mussismilia harttii</i>	E	189.7 (126.8 - 283.6)	158.1 (130.0 - 192.1)	254.7 (109.5 - 451.2)	159.6 (132.6 a 192.1)
<i>Mussismilia hispida</i>	E	>500	>500	>500	>500
<i>Palythoa caribaeorum</i>	E	>500	>500	>500	>500
<i>Plexaurella grandiflora</i>	E	129.6 (72.06 - 233.2)	84.45 (74.87 - 95.26)	164.9 (77.92 - 349.1)	107.07 (97.02 - 251.5)
<i>Porites asteroides</i>	E	>500	>500	>500	>500
<i>Scolymia wellsi</i>	E	>500	>500	>500	>500
<i>Siderastrea stellata</i>	E	444.8 (229.4 - 562.5)	382.9 (309.3 - 473.9)	403.7 (301.3 - 556.5)	476.2 (299.4 - 601.6)
Paclitaxel		0.14 (0.06 - 0.18)	0.18 (0.12 - 0.26)	0.57 (0.42 - 0.70)	>500

A: acetone extract; E: ethanolic extract; D: dichloromethane extract.

3.2. Antiprotozoal activity

The antiprotozoal screening against *L. amazonensis* and *T. cruzi* of the same marine invertebrate species tested above showed that the ethanolic extract of *Amphimedon compressa* and the acetone and ethanolic extracts of *Dysidea* sp. presented the most promising results (Table 3).

In view of the promising activity of the ethanolic extract of *Amphimedon compressa* and the acetone and ethanolic extracts of *Dysidea* sp., they were assayed against

L. amazonensis and *T. cruzi* intracellular amastigotes to determine their IC₅₀ values and selective indices (Table 4). Despite of the activity observed and showed in Table 3, the results on the human macrophage cell line THP-1 pointed to a low specificity.

3.3. Anti-herpes activity

The results of the anti-herpes screening indicated that the ethanolic extracts of *Chondrosia collectrix* and *Haliclona* sp. (200 µg/mL) completely inhibited

Table 3. Percentages of growth inhibition of *Leishmania amazonensis* and *Trypanosoma cruzi* caused by marine invertebrate extracts (sponges and corals) at 50 µg/mL.

Species	Extracts	% growth inhibition±SD	
Sponges		<i>Leishmania amazonensis</i>	<i>Trypanosoma cruzi</i>
<i>Amphimedon compressa</i>	E	61.73±3.62	98.20±0.10
<i>Chondrilla caribensis</i>	A	32.68±2.78	2.32±1.25
<i>Chondrilla caribensis</i>	E	17.91±3.40	4.06±0.54
<i>Chondrosia collectrix</i>	E	31.56±0.91	2.70±0.24
<i>Chondrosia collectrix</i>	D	27.26±2.57	0.57±0.05
<i>Cinachyrella</i> sp.	E	ND	1.25±0.47
<i>Cladocroce caelum</i>	E	ND	12.00±0.58
<i>Desmapsamma anchorata</i>	E	35.42±1.27	2.57±0.20
<i>Dysidea etheria</i>	E	ND	14.69 ±1.07
<i>Dysidea etheria</i>	A	3.89±0.52	15.20±1.40
<i>Dysidea</i> sp.	A	80.30±2.15	96.48±0.65
<i>Dysidea</i> sp.	E	64.44±0.74	74.69±2.78
<i>Echinodictyum dendroides</i>	E	29.43±3.80	3.73±0.30
<i>Haliclona implexiformis</i>	E	ND	1.58±0.07
<i>Haliclona melana</i>	E	4.59±0.95	8.81±0.58
<i>Haliclona</i> sp.	E	24.47±6.17	7.47±0.27
<i>Scopalina ruetzleri</i>	E	ND	9.29 ±0.23
<i>Tedania ignis</i>	E	4.17±1.10	16.86±2.04
Species	Extracts	% growth inhibition±SD	
Corals		<i>Leishmania amazonensis</i>	<i>Trypanosoma cruzi</i>
<i>Agaricia humilis</i>	E	6.47±0.24	9.18±1.60
<i>Carijoa riisei</i>	E	10.04±1.82	4.10±0.66
<i>Favia gravida</i>	E	8.75±3.43	ND
<i>Millepora alcornonis</i>	E	16.23±3.43	ND
<i>Millepora brasiliensis</i>	E	23.26±1.82	6.52±2.59
<i>Montastrea cavernosa</i>	E	ND	ND
<i>Muriceopsis sulphurea</i>	E	17.30±0.93	9.66±1.53
<i>Mussismilia harttii</i>	E	3.87±0.82	ND
<i>Mussismilia hispida</i>	E	29.08±3.35	2.00±0.60
<i>Palythoa caribaeorum</i>	E	14.25±3.10	2.93±1.32
<i>Plexaurella grandiflora</i>	E	21.57±2.63	3.93±1.21
<i>Porites asteroides</i>	E	15.15±2.50	4.41±0.65
<i>Scolymia wellsii</i>	E	26.34±3.29	2.51±0.07
<i>Siderastrea stellata</i>	E	21.09±0.81	ND
Amphotericin B (1.84 µg/mL)	-	96.59±1.79	ND
Benznidazole (5.2 µg/mL)	-	ND	90.93±0.57

ND: not detected; A: acetone extract; E: ethanolic extract; D: dichloromethane extract.

Table 4. Leishmanicidal and trypanocidal activities of the extracts of two marine sponges on *Leishmania amazonensis* and *Trypanosoma cruzi* amastigotes in human macrophage THP-1 cell line.

Species	Extracts	THP-1			<i>Trypanosoma cruzi</i>	
		CC₅₀ ± SD	IC₅₀ ± SD	SI	IC₅₀ ± SD	SI
<i>Amphimedon compressa</i>	E	13.96±1.70	18.79±3.52	0.74	26.94±3.87	0.74
<i>Dysidea</i> sp.	A	< 15.60	16.37±2.43	< 0.95	15.00±0.79	< 0.95
<i>Dysidea</i> sp.	E	27.02±1.97	34.80±1.29	0.77	47.48±1.35	0.77
Amphotericin B		24.95± 0.18	0.12±0.07	207		
Benznidazole		130.12±1.34			2.6±0.22	50

A: acetone extract; E: ethanolic extract; CC₅₀ and IC₅₀ (µg/mL); SI: selective index = (CC₅₀/IC₅₀).

HSV-1 (KOS strain) replication. The ethanolic extract of *Amphimedon compressa* and the acetone extract of *Dysidea* sp. were assayed only at very low concentrations (3.12 and 1.56 µg/mL, respectively) due to their high cytotoxicity to VERO cells that are permissive to herpes viruses infection, but even at these low concentrations they showed to be active inhibiting viral replication by almost 50% (Table 5).

In view of these promising results, the ethanolic extracts of the sponges *Amphimedon compressa*, *Haliclona* sp. and *Chondrosia collectrix*, as well as the acetone extract of *Dysidea* sp were further assayed to determine their IC₅₀ and SI values. The results showed that the ethanolic

extract of *Haliclona* sp. demonstrated the highest activity with a SI>50 (Table 6)

Additionally, the ethanolic extract of *Haliclona* sp. was evaluated against other herpesvirus and strains [HSV-1 (29R strain) and HSV-2 (333 strain)]. The results demonstrated that this extract was also able to inhibit HSV-2 replication as well as the other tested strain of HSV-1, 29R, with high selective indices (Table 7). This is an important finding since 29R strain is resistant to acyclovir, the drug of choice to clinically treat herpes infections. In this Table 7, the results obtained with HSV-1 (KOS strain) (originally in Table 6) was inserted to compare the sponge activity against all tested viruses.

Table 5. Percentages of HSV-1 (KOS strain) inhibition after 48h of treatment with marine invertebrate extracts (sponges and corals) by viral plaque number reduction assay.

Species Sponges	Extracts	Tested concentration (µg/mL)	% HSV-1 inhibition±SD
<i>Amphimedon compressa</i>	E	3.12	49.25±5.76
<i>Chondrilla caribensis</i>	A	200	0
<i>Chondrilla caribensis</i>	E	200	0
<i>Chondrosia collectrix</i>	E	100	98.66±0.68
<i>Chondrosia collectrix</i>	D	200	0
<i>Cinachyrella</i> sp.	E	200	0
<i>Cladocroce caelum</i>	E	100	0
<i>Desmapsamma anchorata</i>	E	200	0
<i>Dysidea etheria</i>	E	100	33.97±9.47
<i>Dysidea etheria</i>	A	200	0
<i>Dysidea</i> sp.	A	1.56	42.96±8.38
<i>Dysidea</i> sp.	E	50	13.75±6.26
<i>Echinodictyum dendroides</i>	E	200	3.40±1.04
<i>Haliclona implexiformis</i>	E	200	7.20±2.21
<i>Haliclona melana</i>	E	100	6.47±4.02
<i>Haliclona</i> sp.	E	50	100.00±0.00
<i>Scopalina ruetzleri</i>	E	100	14.19±5.34
<i>Tedania ignis</i>	E	50	40.88±3.88
Species Corals	Extracts	Tested concentration (µg/mL)	% HSV-1 inhibition±SD
<i>Agaricia humilis</i>	E	100	16.95±7.11
<i>Carijoa riisei</i>	E	100	14.45±5.94
<i>Favia gravida</i>	E	100	27.72±8.91
<i>Millepora allicornis</i>	E	100	5.69±3.16
<i>Millepora brasiliensis</i>	E	100	13.13±5.53
<i>Montastrea cavernosa</i>	E	100	6.49±4.62
<i>Muriceopsis sulphurea</i>	E	100	0
<i>Mussismilia harttii</i>	E	50	16.74±9.75
<i>Mussismilia hispida</i>	E	100	0
<i>Palythoa caribaeorum</i>	E	100	0
<i>Plexaurella grandiflora</i>	E	50	39.63±15.64
<i>Porites asteroides</i>	E	100	14.88±4.52
<i>Scolymia wellsi</i>	E	100	0
<i>Siderastrea stellata</i>	E	100	20.52±8.29

A: acetone extract; E: ethanolic extract; D: dichloromethane extract.

Table 6. Anti-herpes (HSV-1, KOS strain) activity after 48 h of treatment with four marine sponge extracts by viral plaque number reduction assay.

Species	Extract	CC ₅₀ IC ₅₀ ±SD		SI
		VERO cells	HSV-1	
<i>Amphimedon compressa</i>	E	9.9 (7.2 a 13.6)	2.40±0.68	4.1
<i>Chondrosia collectrix</i>	E	444.9 (359.3 a 550.8)	60.00±5.35	7.4
<i>Dysidea</i> sp.	A	12.2 (8.2 a 18.9)	2.03±0.71	6.0
<i>Haliclona</i> sp.	E	>500	10.03±2.14	>50

CC₅₀ and IC₅₀ (µg/mL); CC₅₀ (95% confidence interval); SI: selective index=CC₅₀/IC₅₀; A: acetone extract; E: ethanolic extract.

Table 7. Anti-herpes activity [anti-HSV-1 (KOS and 29R strains) and anti-HSV-2 (333 strain)] after 48 h of treatment with the ethanolic extract of *Haliclona* sp. by viral plaque number reduction assay.

Sponge	VERO cells	CC ₅₀ IC ₅₀ ±SD			Selective indices		
		HSV-1 (KOS strain)	HSV-1 (29R strain)	HSV-2 (333 strain)	HSV-1 (KOS strain)	HSV-1 (29R strain)	HSV-2 (333 strain)
<i>Haliclona</i> sp.	>500	10.03±2.14	6.28±2.47	4.59±1.09	>50	>79	>108
Acyclovir	>500	1.23± 0.51	-	3.48±0.76	>406	-	>143

CC₅₀ and IC₅₀ (µg/mL); SI: selective index=CC₅₀/IC₅₀.

3.4. Antibacterial activity

The results of the antibacterial screening against *Enterococcus faecalis*, *Pseudomonas aeruginosa* and *Escherichia coli* indicated that only the ethanolic extract of *Amphimedon compressa* exhibited a weak antibacterial activity with inhibition halos ranging from 7.0 to 8.5 mm.

4. Discussion

The evaluation of antiproliferative effects of 32 marine invertebrate extracts revealed that the ethanolic extracts of *Amphimedon compressa* and *Tedania ignis*, and the acetone extract of *Dysidea* sp showed results that can be considered relevant.

The species *Amphimedon compressa* (formerly *Haliclona rubens*) belongs to the order Haplosclerida. Marine sponges belonging to this order contributed with about 14% of the bioactive compounds isolated from marine sponges (Mehub et al., 2014), and are a rich source of alkaloids with pronounced biological activities (Andersen et al., 1996; Almeida et al., 1997). Considering the previous published results with sponges of this genus, Berlinck and colleagues performed a screening with methanol-soluble crude extracts of sponges collected in southeastern Brazilian coast, and observed that *Amphimedon viridis* (formerly *Haliclona viridis*) displayed potent cytotoxic and hemolytic activities (Berlinck et al., 2004). The same authors reported latter the isolation of a new purine, 1,3-dimethylisoguanine (Chehade et al., 1997) from this species. Another research group reported the isolation of the same compound from the same sponge at the same time. The authors tested it against 26 human cancer cell lines and observed that it displayed the highest cytotoxic activity against an ovarian cancer cell line (Mitchell et al., 1997).

Another paper reported the cytotoxic screening of hydro-methanolic extracts of eight sponges collected in the northwest Brazilian coast (Ceará state), and it was found that the extract of *Amphimedon* sp. showed antiproliferative effects on three tumor cell lines [CEM and HL-60 (human leukemia cells) and HCT-8 (human ileocecal adenocarcinoma cells)], with no effects on the murine melanoma (B-16) tested (Jimenez et al., 2004).

Concerning the antiproliferative effects of *Dysidea* sp. reported here, there is already another work describing the antimicrobial (bacteria and yeasts), antimycobacterial, antileishmania, and cytotoxic screenings of several samples of *Dysidea* sp. collected along the Brazilian southeastern and northeastern coastline, which showed that the crude extracts of four out of the eleven samples tested were strongly active against MCF-7 breast and HCT-8 colon cancer cells (Selegim et al., 2007).

Regarding the leishmanicidal and tripanocidal activities, the extracts of *Amphimedon compressa* and the acetone and ethanolic extracts of *Dysidea* sp. showed promising results. However, both extracts showed very low selectivity indices using THP-1 cell line indicating that they are not specific. There are few reports of antiprotozoal activity of marine sponges and corals in the literature. As example two reports can be cited: firstly, De Paula and co-workers evaluated the *in vitro* activity of five species of Brazilian and Spanish marine sponges against *T. cruzi*, and showed that the acetone extract of *Dysidea avara* collected in Spanish coast presented high activity against trypanomastigote forms (Paula et al., 2015). Secondly, Andrade et al. (2015) investigated the activity of organic and aqueous extracts obtained from the sponge *Amphimedon viridis* collected in southern Brazilian coast against *T. cruzi* and demonstrated

that both extracts displayed concentration-dependent lytic effects on bloodstream trypomastigotes. The aqueous extract was ~19-fold more active than benznidazole used as positive control. Trypomastigotes treated with the aqueous extract and labelled with tetramethylrhodamine led to mitochondrial membrane potential dissipation on almost 80% of the parasites, whereas labelling with 7-aminoactinomycin D induced apoptosis-like death in almost 50% of the parasites. These data suggest a relevant trypanocidal activity of *A. viridis*, which is partially associated with the ability to induce apoptosis-like death (Andrade et al., 2015).

Regarding the anti-herpes activity, the ethanolic extract of *Haliclona* sp. showed the most promising results. When comparing the inhibition of HSV replication, HSV-2 was observed to be more sensitive to *Haliclona* sp. followed by HSV-1-29-R, a resistant ACV strain. Additionally, the SI value showed by *Haliclona* for HSV-2 was > 108, comparable to that found for ACV (>143), used as positive control. Taken together, these results can be considered an advantage among other samples, since the emergence of resistant strains to ACV has been a challenge to treat herpes infections. Most of the antiherpetic drugs available share the same mechanism resulting in cross-resistance and attesting the necessity to find new treatment alternatives (Silva-Mares et al., 2016).

The order Haplosclerida, which includes the sponge genera *Haliclona*, *Xestospongia* and *Amphimedon*, is a rich source of β -carboline alkaloids called manzamines. They were first isolated by Higa and co-workers in 1986 from the Okinawan sponge genus *Haliclona* (Sakai et al., 1986). Until 2016, over 80 manzamine-related alkaloids have been reported from more than 16 marine sponge species belonging to five families (Hu et al., 2003; Peng et al., 2003). Manzamine A, 8-hydroxymanzamine A, and 8-methoxymanzamine A showed moderate activity against HSV-2 replication (Radwan et al., 2012; Palem et al., 2011). Palem et al. (2011) concluded that manzamine A, isolated from the Indo-Pacific sponge *Acanthostrongylophora* sp., displayed potent activity against HSV-1 replication, via targeting the viral gene infectious cellular protein ICP0. Furthermore, aiming to discover new natural product groups that could inhibit virus replication and control HSV-1 infections, they screened ten compounds of different chemical classes isolated from diverse marine sponges. They observed that the anti-HSV-1 activity of 8-hydroxymanzamine A was lower, when compared to that of manzamine A, suggesting that C-8 hydroxy substitution was involved in the anti-herpes activity. Moreover, when manzamine A was chemically transformed to its salt forms, the anti-HSV-1 activity was improved suggesting that augmentation of the overall water solubility can significantly enhance the activity (Jayavardhana et al., 2017). In view of these data and also considering the previous published work of our group (Bianco et al., 2013) that showed the anti-herpes activity of the buthanolic extract of *Haliclona* sp. collected in Aranhas Island, Santa Catarina State, South Brazil, species of this genus should receive special

attention for future works aiming the isolation and structural elucidation of their bioactive metabolites.

The antibacterial screening showed that the only positive result was observed for the ethanolic extract of *Amphimedon compressa*. Interestingly, in the literature there are some previous works reporting, for example, the antibacterial activity of a hydro-methanolic extract of *Amphimedon* sp. against *Staphylococcus aureus* as well as *Pseudomonas aeruginosa* strains (Jimenez et al., 2004). In another study with crude extracts of sponges collected in Rio de Janeiro state, Brazil, the crude extract of *Amphimedon* sp. also showed a weak antibacterial activity against oxacillin-resistant *S. aureus* (Selegim et al., 2007).

5. Conclusion

In this work, 32 different extracts from Brazilian marine invertebrates were screened for their potential antiproliferative, antibacterial, antiprotozoal and anti-herpes activities. The results showed that the sponges *Amphimedon compressa*, *Chondrosia collectrix*, *Tedania ignis*, *Dysidea* sp., and *Haliclona* sp. were the species presenting the most important bioactivities, and deserve further investigations. Special attention should be given to the sponge *Haliclona* sp. regarding its antiherpes activity against all tested strains with emphasis to HSV-1 (29R strain, which is resistant to acyclovir, SI=79) and to HSV-2 (SI=108).

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