TITLE: BIOPROSPECTION FOR ANTIBIOTIC RESISTANCE IN STRAINS OF *Vibrio* AND *Shewanella* ISOLATED FROM MARINE SPONGES

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ABSTRACT:

Antibiotic resistance is a global public health problem. The qnr and bla_{OXA-48} genes encode resistance mediators to quinolones and carbapenems, respectively, and were been detected in the chromosome of Shewanella strains. Genes encoding similar Onr proteins were detected in strains of Vibrio. Both genera are Gram-negative bacilli, found in marine environments and are often isolated from marine sponges. Because they are filter feeders, marine sponges can accumulate environmental contaminants such as antibiotics, which might induce the selection of associated bacteria expressing determinants of resistance to these substances. Thereby, strains carrying genes conferring resistance to antibiotics specially in mobile genetic elements, might transfer these genes to pathogenic bacteria, thus underlining the importance of resistance encountered in bacteria associated with marine sponges. The aims of the present study were to determine the susceptibility profile to antibiotics (SPA) and to verify the presence of antibiotic resistance genes in strains of Vibrio and Shewanella isolated from marine sponges collected in Cabo Frio, Brazil and Marseille, France. Previously, a total of 108 strains, 69 Vibrio sp. and 39 Shewanella sp., were isolated and identified by rrs gene sequencing. SPA determination was done using the Kirby-Bauer antimicrobial disk diffusion procedure on Mueller Hinton Agar plates. Preliminary results were obtained for 41 strains (23 Vibrio sp. and 18 Shewanella sp.). These strains were tested for 21 different antibiotics. Twenty-one strains (51.2%, 11 Vibrio sp. and 10 Shewanella sp.), showed concomitant resistance to cefepime, aztreonam and ceftazidime. Interestingly, one Shewanella sp. was multiresistant (to β -lactams, aminoglycosides and quinolones). Overall, 25 (60.9%) strains were resistant to aztreonam; 24 (58.5%) to ceftazidime; 21 (51.2%) to cefepime; 9 (21.9%) to nalidixic acid; 7 (17.0%) to erythromycin; 7 (17.0%) to cefoxitin; one (2.4%) to cephalexin; one (2.4%) to tobramycin, amikacin and gentamycin; and one (2.4%) to ciprofloxacin and ofloxacin. Only 5 (12.2%) strains were sensitive to all antibiotics tested. In the future, we will use PCR to detect antibiotic resistance genes and perform plasmid profile analysis of Vibrio and Shewanella strains. We expect to contribute to the understanding of the roles of so-called native resistance genes in natural microbial communities.

Keywords: antibiotic resistance, Vibrio, Shewanella, marine sponges

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