

عنوان مقاله:

Identification of novel antimicrobial peptide from Asian sea bass (*Lates calcarifer*) by in silico and activity characterization

محل انتشار:

بیستمین کنگره بین المللی میکروب شناسی ایران (سال: 1398)

تعداد صفحات اصل مقاله: 1

نویسندگان:

Behrouz Taheri - *Department of Medical Laboratory Sciences, School of Paramedicine, Ahvaz Jundishapur University, of Medical Sciences, Ahvaz, Iran*

Mohsen Mohammadi - *The Persian Gulf Marine Biotechnology Research Center, the Persian Gulf Biomedical Sciences Research Institute, Bushehr University of Medical Sciences, Bushehr, Iran*

Mona Roozbehani - *PhD Students of Medical Parasitology School of Medicine, Iran University of Medical Sciences, Tehran, Iran*

Iraj Nabipour - *The Persian Gulf Marine Biotechnology Research Center, the Persian Gulf Biomedical Sciences Research Institute, Bushehr University of Medical Sciences, Bushehr, Iran*

Nilloofar Momenzadeh - *The Persian Gulf Marine Biotechnology Research Center, the Persian Gulf Biomedical Sciences Research Institute, Bushehr University of Medical Sciences, Bushehr, Iran*

خلاصه مقاله:

Introduction and Objectives: The global crisis of antibiotic resistance increases the demand for the new promising alternative drugs such as antimicrobial peptides (AMPs). Accordingly, we have described a new, previously unrecognized effective AMP, named dicentracin-like, from Asian sea bass and characterized its antimicrobial activity by comparison with moronecidin. **Materials and Methods:** Gene expression analysis demonstrated the expression of dicentracin-like peptide in tissues of the immune system such as the skin and the head kidney, which is an important endocrine and lymphoid organ. Moronecidin and dicentracin-like exhibited a higher antibacterial activity against gram-positive bacteria relative to gram-negative ones, while both peptides showed a greater binding ability to gram-negative bacteria compared to gram-positive ones. This contradiction between antibacterial activity and binding affinity may be related to the outer membrane from gram-negative bacteria. Compared with moronecidin, dicentracin-like peptide showed more potent binding ability to all gram-positive and gram-negative bacteria. In addition, dicentracin-like peptide exhibited a high antibacterial activity against the investigated microorganisms, except against *Staphylococcus aureus*. A direct relationship was found between the binding affinity/cationicity and the antibiofilm activity of the peptides wherein, an elevation in pH corresponded to a decrease in their antibiofilm property. Time kill kinetics analysis against clinical *Acinetobacter baumannii* isolate indicated that bactericidal effect of dicentracin-like and moronecidin at inhibitory concentration (1XMIC) was observed after 4 and 6 hours, respectively, while bactericidal effect of both AMPs at concentration of 2XMIC was observed after 2 hours. Dicentracin-like peptide showed higher inhibitory activity at subinhibitory concentration (1/2XMIC), relative to moronecidin. Compared with moronecidin, dicentracin-like peptide possessed greater binding affinity to bacteria at high salt concentration, as well as at alkaline

pH; In addition, dicentracin-like exhibited a higher antibiofilm activity in comparison to moronecidin even at alkaline pH. Hemolytic analysis against human RBC revealed that hemolytic activity of moronecidin was more potent than that of dicentracin-like, which is consistent with its greater non-polar face hydrophobicity. Conclusions: In the present study, In Silico comparative sequence analysis and antimicrobial characterization led to identify a new, previously ... unrecognized antimicrobial function for named dicentr

کلمات کلیدی:

Antimicrobial peptide, *Acinetobacter baumannii*, *Staphylococcus aureus*, in silico analysis

لینک ثابت مقاله در پایگاه سیویلیکا:

<https://civilica.com/doc/987254>

