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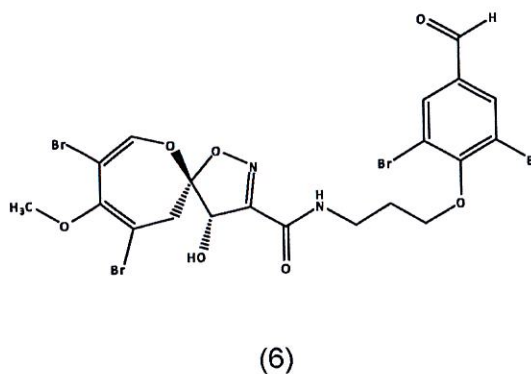
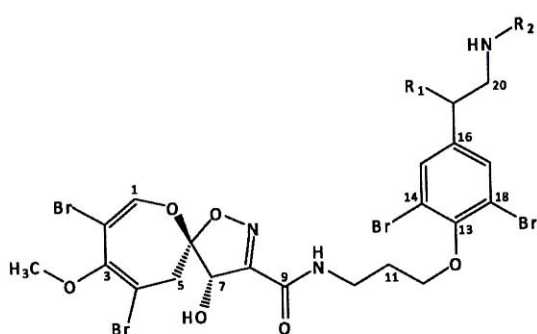
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NEW BROMOTYROSINE DERIVATIVES FROM THE BALINESE MARINE SPONGE *APLYSINELLA STRONGYLATA*

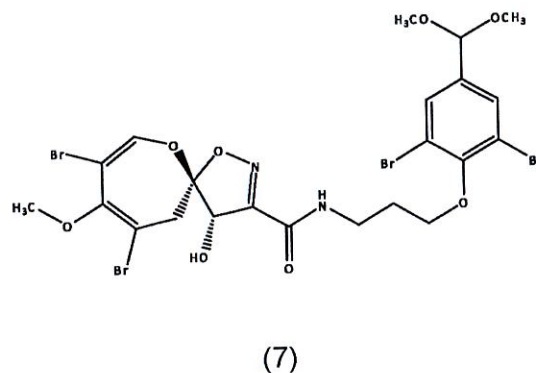
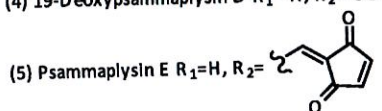
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Sponges of the order Verongida have been a prolific source of bromotyrosine-derived metabolites including those possessing a distinctive spirooxepinisoxazoline (oxepin) moiety.¹ There have been ten oxepin bromotyrosine metabolites identified from marine sponges so far, demonstrating *in vitro* cytotoxicity towards the human colon tumor cell-line, anti-HIV activity against the Haitian RF strain, and activity against *plasmodium falciparum*.¹⁻² A series of known oxepin-containing metabolites named psammaplysin (1, 2-5) together with two new metabolites were isolated from sponge *Aplysinella strongylata*, obtained from Bali.³ An attempt to determine the absolute configuration through a single crystal X-ray study on psammaplysin-A acetamide (2) is underway.



- (1) Psammaplysin A $R_1 = R_2 = H$
 (2) Psammaplysin A acetamide acetate $R_1 = H, R_2 = COCH_3$
 (3) Psammaplysin B $R_1 = -OH, R_2 = H$
 (4) 19-Deoxypsammaplysin D $R_1 = H, R_2 = CO(CH_2)_{11}CH(CH_3)_2$



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- (a) Roll, D. M.; Chang, C. W. J.; Scheuer, P. J.; Gray, G. A.; Shoolery, J. N.; Matsumoto, G. K.; Duyne, G. D. V.; Clardy, J. *J. Am. Chem. Soc.* **1985**, 107,