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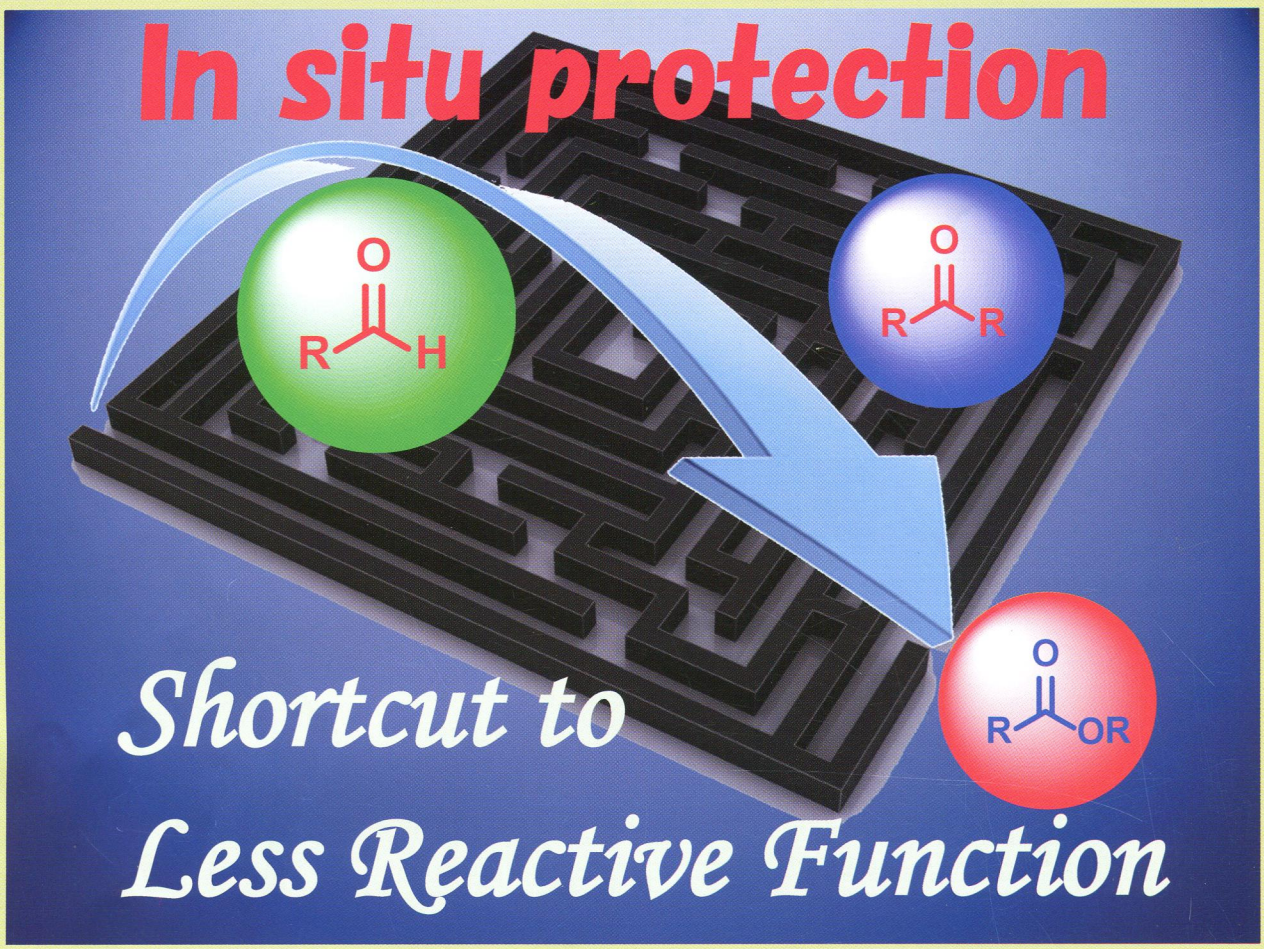

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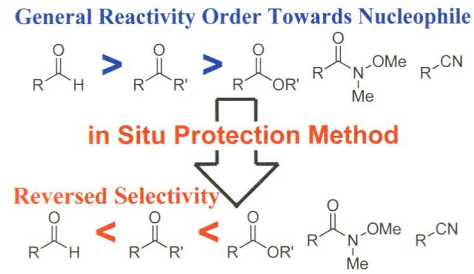

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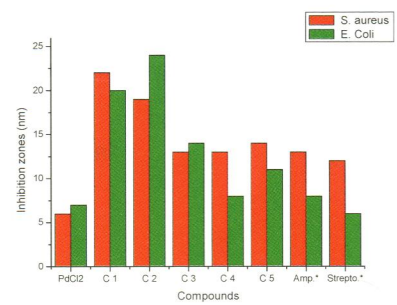

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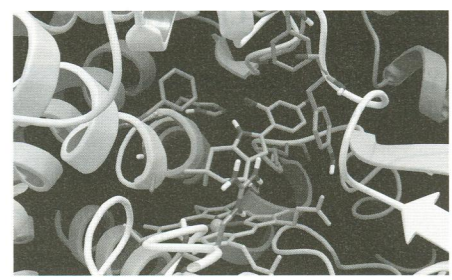

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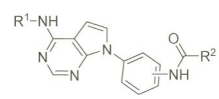

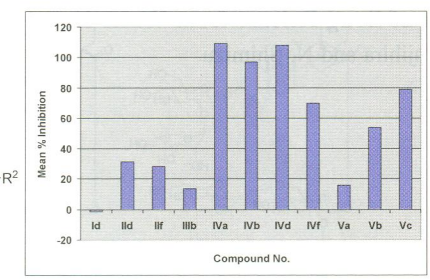

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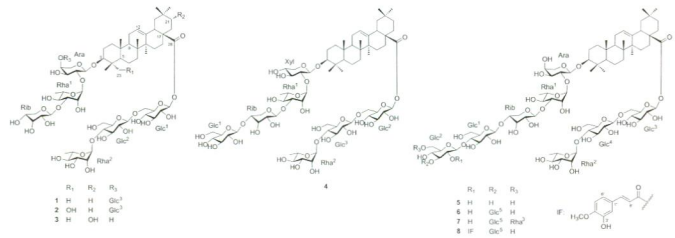

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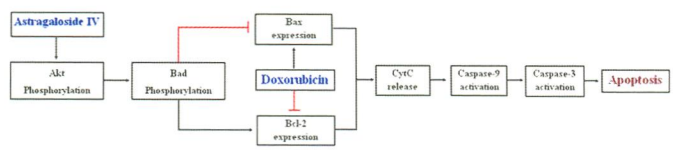

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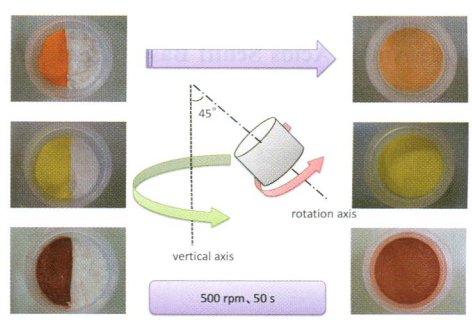

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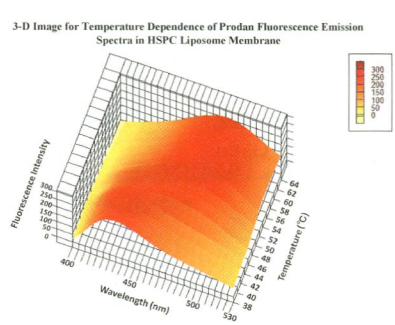

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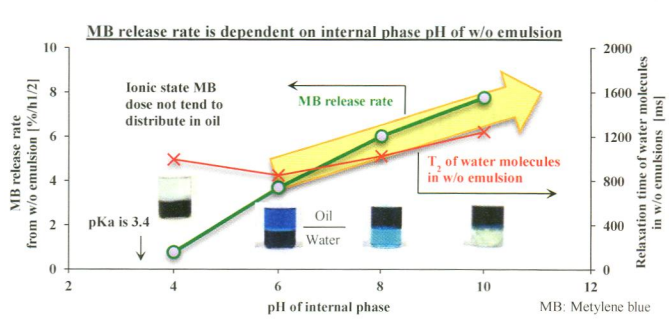

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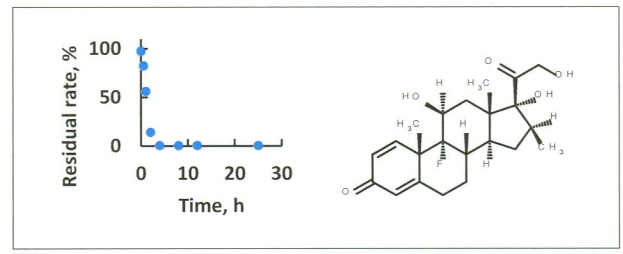

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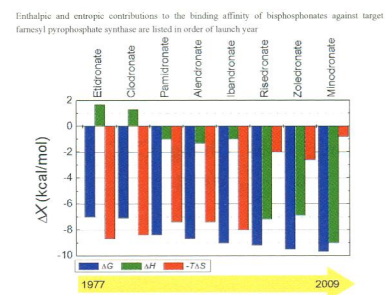

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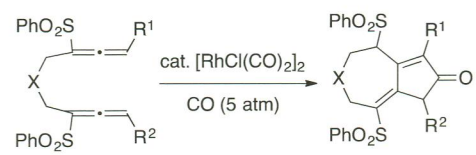

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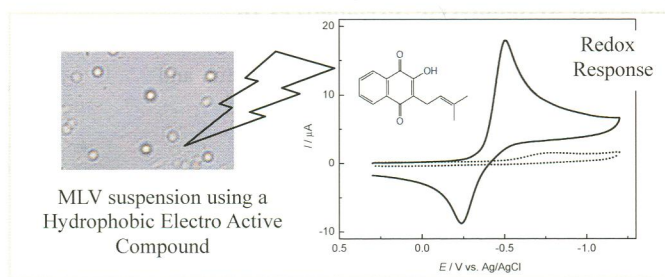

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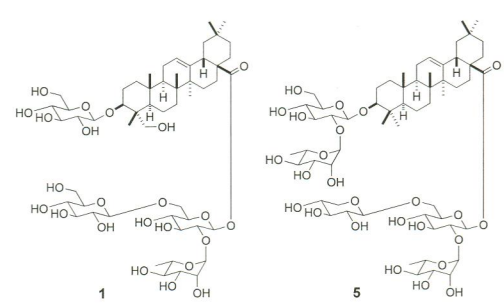

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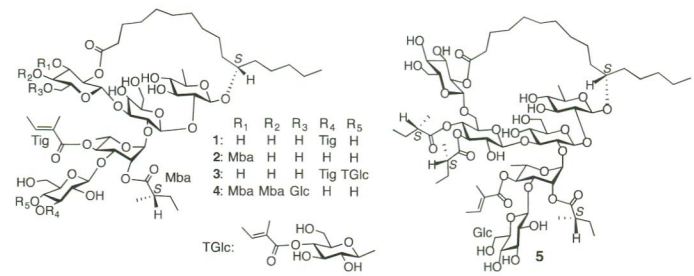

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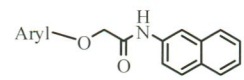

Aryl = Anthraquinone, Xanthone, Carbazole

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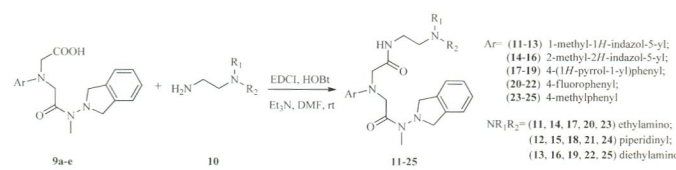
Design and Synthesis of Amide Derivatives as *S*-Adenosyl-L-Homocysteine Hydrolase Inhibitors

X. Tan, P. Wang, S. Nian, and G. Wang



Compound **14** displayed significant SAHase inhibitory activity.

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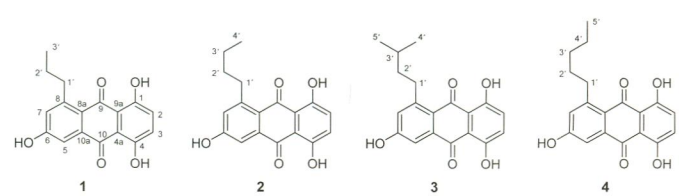
Z.-Y. Wu, W. Fang, L.-Q. Shi, Z.-Y. Wan, Y. Min, and K.-M. Wang



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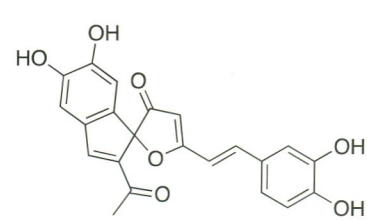

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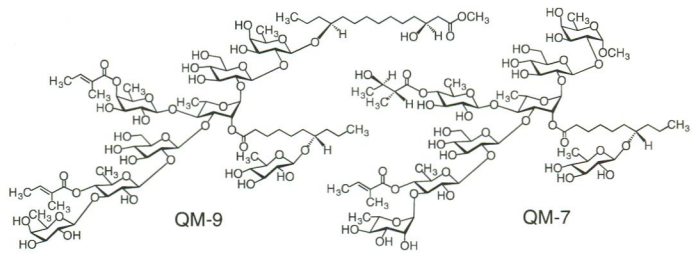
K. Akiyama, K. Yamamoto, T. Mineno, M. Okawa, J. Kinjo, H. Yoshimitsu, T. Nohara, and M. Ono



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About the cover: This review discusses *in situ* protection techniques on carbonyl chemistry. In general, intrusive multistep protection-deprotection detour is required to transform less reactive functions selectively in the presence of more reactive ones. *In situ* protection methodologies are useful and powerful one-pot techniques which enable to shortcut to the transformation of less reactive functions. In the present paper, various methodologies for selective transformation of carbonyl groups accomplishing reversed selectivity, such as a selective alkylation of ketones and esters in the presence of aldehydes, are reviewed. See the review by Yahata and Fujioka on page 1 of this issue.