

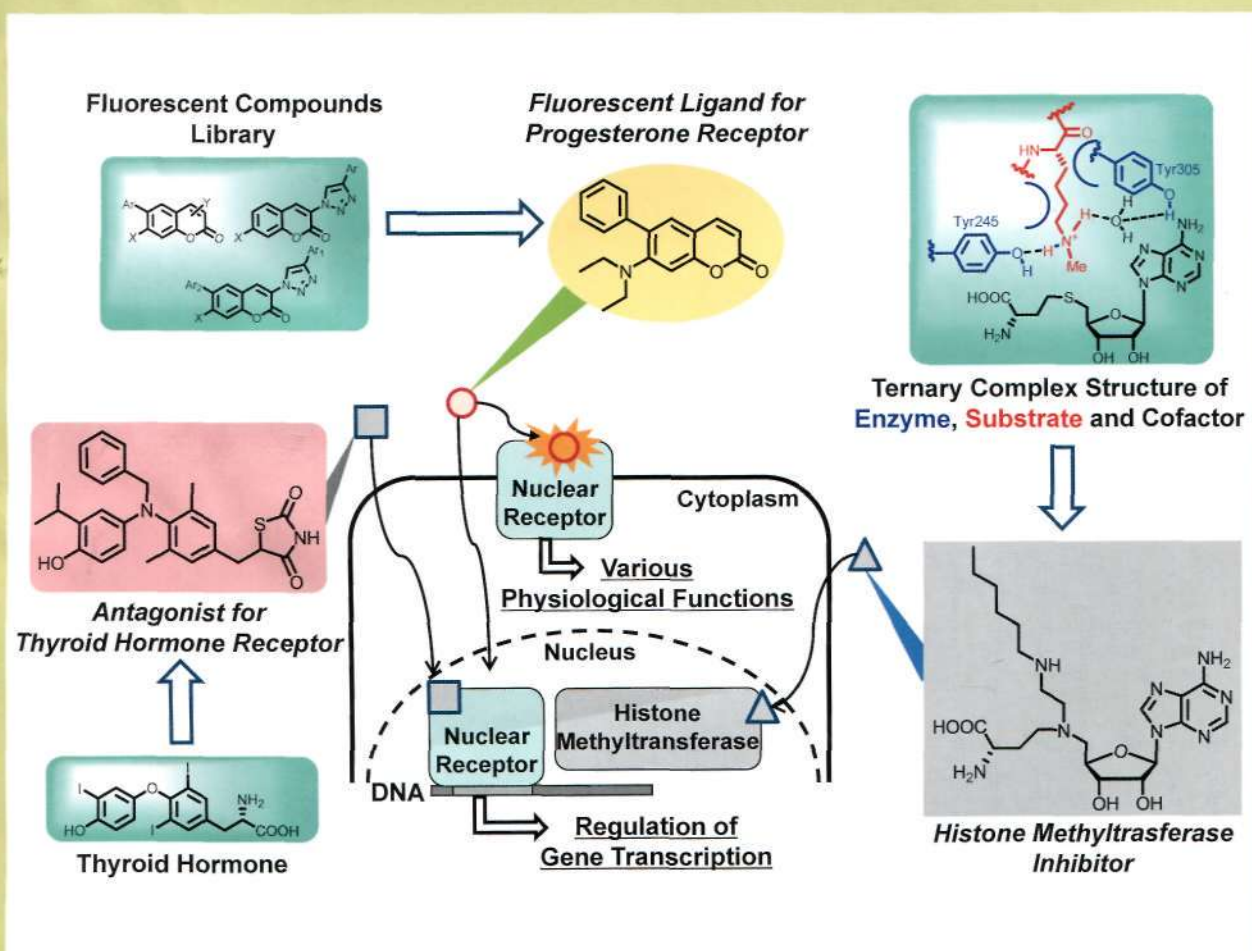
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Vol. 61 No. 2



Development of Functional Molecules for Elucidation of the Physiological Roles of Several Nuclear Receptors and Their Endogenous Ligands

pp. 111-120



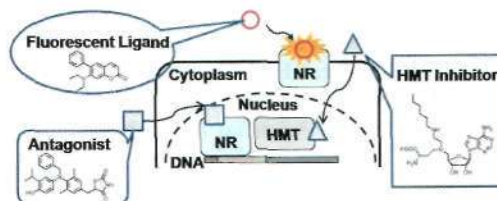
THE PHARMACEUTICAL SOCIETY OF JAPAN

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Review

Development of Functional Molecules for Elucidation of the Physiological Roles of Several Nuclear Receptors and Their Endogenous Ligands

T. Hirano

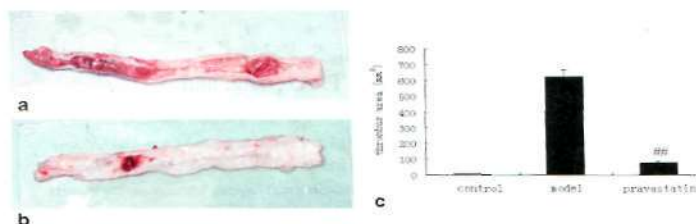


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Regular Articles

Pravastatin Inhibits Plaque Rupture and Subsequent Thrombus Formation in Atherosclerotic Rabbits with Hyperlipidemia

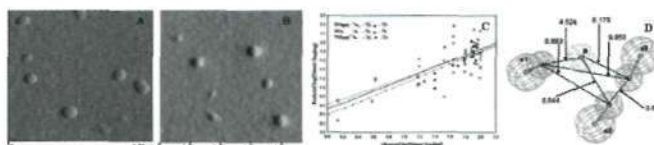
G. Wu, Q. Xie, L. Xu, H. Jiang, Z. Huang, and C. Huang



pp. 121–124

Poly(DL-lactide-co-glycolic acid) Nanoparticle Design and Payload Prediction: A Molecular Descriptor Based Study

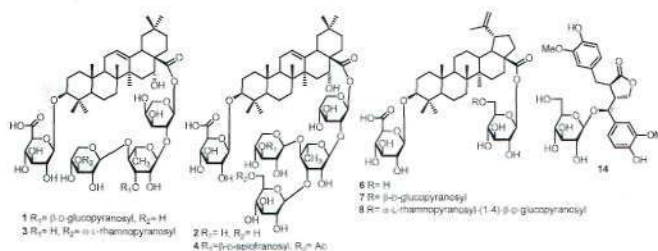
S. Das, P. Roy, M. A. Islam, A. Saha, and A. Mukherjee



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Triterpene Saponins from *Clethra barbinervis* and Their Hyaluronidase Inhibitory Activities

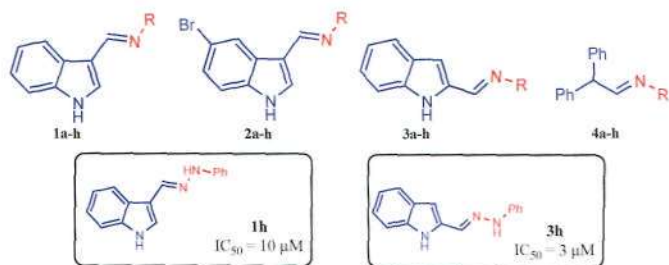
T. Murata, A. Suzuki, N. Mafune, E. Sato, T. Miyase, and F. Yoshizaki



pp. 134–143

Synthesis of Novel Indole Hydrazone Derivatives and Evaluation of Their Antiplatelet Aggregation Activity

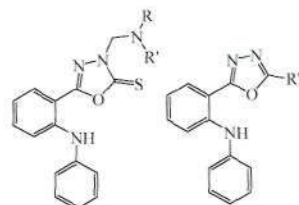
V. Mashayekhi, K. Haj Mohammad Ebrahim Tehrani, S. Amidi, and F. Kobarfard



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Synthesis, Quantitative Structure–Activity Relationship and Biological Evaluation of 1,3,4-Oxadiazole Derivatives Possessing Diphenylamine Moiety as Potential Anticancer Agents

D. E. Abdel Rahman

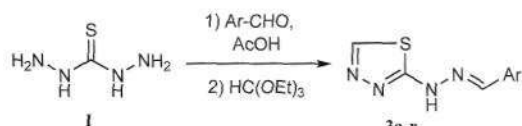


New series of Mannich bases of 5-substituted-1,3,4-oxadiazole-2(3*H*)-thione, 2,5-disubstituted-1,3,4-oxadiazole, 2-substituted thio-5-substituted-1,3,4-oxadiazole were synthesized and evaluated for anti-tumor activity against HT29 and MCF7 cell lines.

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One Pot Synthesis and Biological Activity Evaluation of Novel Schiff Bases Derived from 2-Hydrazinyl-1,3,4-thiadiazole

K. Haj Mohammad Ebrahim Tehrani, S. Sardari, V. Mashayekhi, M. Esfahani Zadeh, P. Azerang, and F. Kobarfard

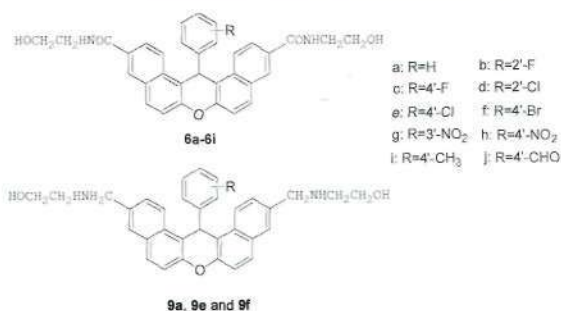


Antiplatelet aggregation activity:		
3l	3-nitrophenyl	IC ₅₀ = 2.6
3o	2-furyl	IC ₅₀ = 2.2
3p	2-thienyl	IC ₅₀ = 3.8
Anti <i>Mycobacterium bovis</i> BCG:		
3g	4-chlorophenyl	MIC = 7.8 μg/ml
3q	3-indolyl	MIC = 7.8 μg/ml

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Design, Synthesis and Anticancer Activity of *N*³,*N*¹¹-Bis(2-hydroxyethyl)-14-aryl-14*H*-dibenzo[*a,j*]xanthenes-3,11-dicarboxamide

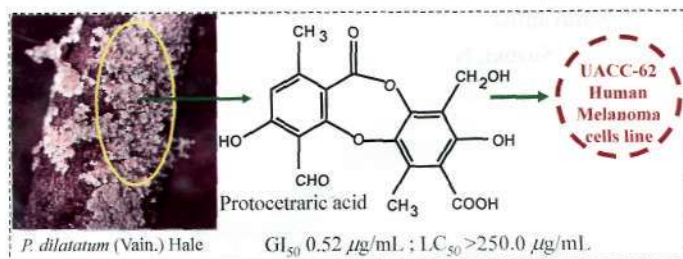
Y. Song, Y. Yang, J. You, B. Liu, L. Wu, Y. Hou, W. Wang, and J. Zhu



pp. 167–175

Cytotoxic Evaluation of Phenolic Compounds from Lichens against Melanoma Cells

L. F. G. Brandão, G. B. Alcantara, M. F. C. Matos, D. Bogo, D. S. Freitas, N. M. Oyama, and N. K. Honda



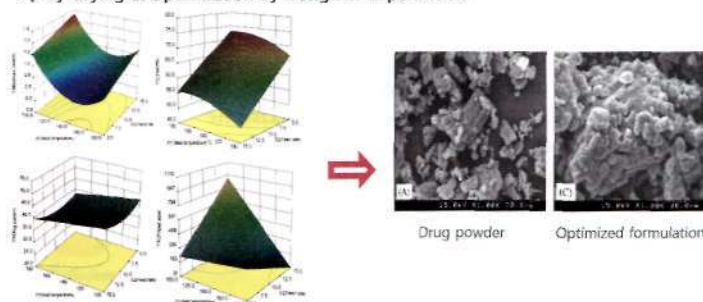
pp. 176–183

Statistical Modeling, Optimization and Characterization of Spray-Dried Solid Self-Microemulsifying Drug Delivery System Using Design of Experiments

N. Marasini, T. H. Tran, B. K. Poudel, H.-G. Choi, C. S. Yong, and J. O. Kim

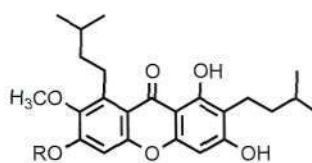
pp. 184–193

Spray-drying & Optimization by Design of Experiments



Potent Activity against Multidrug-Resistant *Mycobacterium tuberculosis* of α -Mangostin Analogs

P. Sudta, P. Jiarawapi, A. Suksamrarn, P. Hongmanee, and S. Suksamrarn

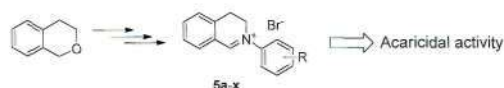


A new series of α -mangostin analogs was prepared and their antimycobacterial activity was evaluated. The 6-*O*-alkyl tetrahydro α -mangostins exhibited highly potent activity against the multidrug-resistant *Mycobacterium tuberculosis* with MICs of 0.78–1.56 $\mu\text{g/mL}$.

pp. 194–203

Synthesis of 2-Aryl-3,4-dihydroisoquinolin-2-ium Bromides and Their *in Vitro* Acaricidal Activity against *Psoroptes cuniculi*

Y.-N. Ma, X.-J. Yang, L. Pan, Z. Hou, H.-L. Geng, X.-P. Song, L. Zhou, and F. Miao



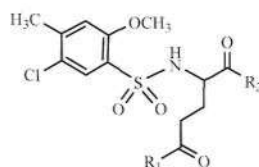
R = H, Me, OH, OMe, NO₂, CF₃, F, Cl, Br, I

Twenty-four 2-aryl-3,4-dihydroisoquinolin-2-ium bromides were synthesized by a new approach and evaluated for *in vitro* acaricidal activity against *Psoroptes cuniculi*. Fourteen compounds showed the stronger activity against *P. cuniculi* than a standard drug ivermectin and their natural model compound.

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Synthesis and Biological Evaluation of Some Substituted-2-*N*-(5-chloro-2-methoxy-4-methylphenylsulphonyl) Glutamic Acid Derivatives against Prostate Cancer Cell Line PC3

G. S. Hassan and D. E. Abdel Rahman

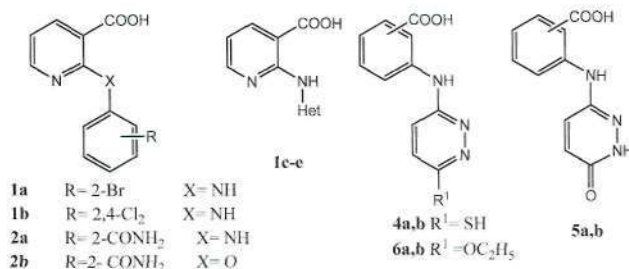


New series of substituted glutamine ($R_1 = \text{NHR}$, $R_2 = \text{OH}$) and glutamic acid diamide, diuride, dihydrazide ($R_1, R_2 = \text{NHR}$) were synthesized from the parent glutamic acid compound ($R_1, R_2 = \text{OH}$) and evaluated for antitumor activity against PC3 cell line.

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Synthesis and Biological Evaluation of New Heteroaryl Carboxylic Acid Derivatives as Anti-inflammatory-Analgesic Agents

K. A. M. Abouzid, N. A. Khalil, E. M. Ahmed, and S. A.-B. Zaitone



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Notes

Effects of Structural and Electronic Characteristics of Chalcones on the Activation of Peroxisome Proliferator-Activated Receptor Gamma

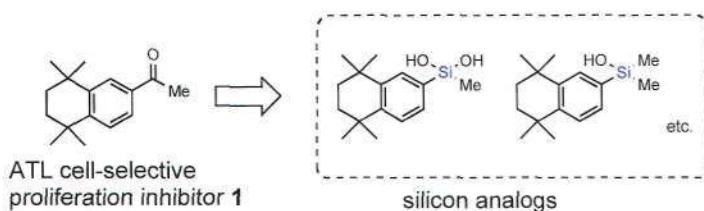
J. T. Schott, C. E. Mordaunt, A. J. Vargas, M. A. Leon, K. H. Chen, M. Singh, M. Satoh, E. L. Cardenas, S. Maitra, N. V. Patel, and H. J. P. de Lijser



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Organosilicon Compounds as Adult T-Cell Leukemia Cell Proliferation Inhibitors

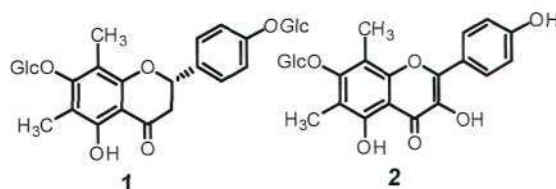
M. Nakamura, Y. Matsumoto, M. Toyama, M. Baba, and Y. Hashimoto



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Diplomorphanins A and B: New C-Methyl Flavonoids from *Diplomorpha canescens*

H. P. Devkota, M. Watanabe, T. Watanabe, and S. Yahara

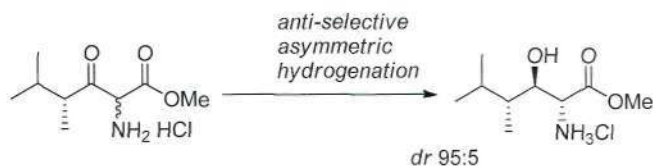


Diplomorphanins A (1) and B (2) were isolated from the aerial parts of *Diplomorpha canescens* along with farrerol 7-*O*- β -D-glucopyranoside (3).

pp. 242–244

Efficient Diastereoselective Synthesis of (2*R*,3*R*,4*R*)-2-Amino-3-hydroxy-4,5-dimethylhexanoic Acid, the Lactone Linkage Unit of Homophymine A

J. Ohtaka, A. Hamajima, T. Nemoto, and Y. Hamada



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