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ORIGINAL ARTICLES

Synthesis and antitumor activities of naturally occurring oleanolic acid triterpenoid saponins and their derivatives

pp. 1-15

Qingchao Liu, Hongchun Liu, Lei Zhang, Tiantian Guo, Peng Wang, Meiyu Geng and Yingxia Li*

Cytotoxic activities IC_{50} values between 3.2-12.9 μM

Development of peptidomimetic boronates as proteasome inhibitors

Nicola Micale*, Roberta Ettari, Antonio Lavecchia, Carmen Di Giovanni, Kety Scarbaci, Valeria Troiano, Silvana Grasso, Ettore Novellino, Tanja Schirmeister and Maria Zappalà

pp. 23-34

Pyrazole derivatives as inhibitors of arachidonic acid-induced platelet aggregation

 $Serkan\ Levent,\ Burcu\ \zetaalışkan,\ Murat\ \zetaiftçi,\ Yeşim\ \"{O}zkan,\ \dot{I}dil\ Yenicesu,\ H\"{u}seyin\ \ddot{U}nver\ and\ Erden\ Banoglu^*$

pp. 42-53

A structure–activity relationship study based on the 1,5-diarylpyrazole-3-carboxamide scaffold for inhibition of arachidonic acid-induced platelet aggregation were performed.

SAR of starting compound (I) Inhibition of AA-induced platelet aggre 61 X = N, $\text{IC}_{50} = 15 \text{ nM}$ 62 X = C, $\text{IC}_{50} = 5.7 \text{ nM}$

Synthesis and antimicrobical evaluation of a novel class of 1,3,4-thiadiazole: Derivatives bearing 1,2,4-triazolo[1,5-a] pyrimidine moiety

pp. 54-61

Yin Luo, Shuai Zhang, Zhi-Jun Liu, Wu Chen, Jie Fu*, Qing-Fu Zeng and Hai-Liang Zhu*

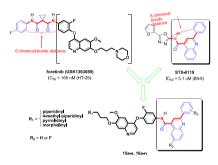
New 1,2,4-triazolo[1,5-a]pyrimidine derivatives have been synthesized and evaluated for their antimicrobial activity. Most of them exhibited good activities and some could be comparable to positive controls.

Design, synthesis and antitumour activity of bisquinoline derivatives connected by 4-oxy-3-fluoroaniline moiety

pp. 62-73

Sai Li, Qiang Huang, Yajing Liu, Xiaolong Zhang, Shuang Liu, Chao He and Ping Gong*

A series of bisquinoline derivatives connected by 4-oxy-3-fluoroaniline moiety were synthesized and evaluated for their cytotoxic activities. Six potent compounds were further examined for their c-Met kinase activity.



Synthesis, antifungal and cytotoxic activities of 2-aryl-3-((piperidin-1-yl)ethyl)thiazolidinones

pp. 74–80

Alice Kunzler, Patrícia D. Neuenfeldt, Adriana M. das Neves, Claudio M.P. Pereira, Gabriela H. Marques, Patrícia S. Nascente, Maureen H.V. Fernandes, Silvia O. Hübner and Wilson Cunico*

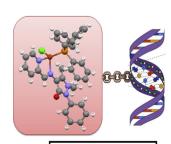
The studies of synthesis, antifungal and cytotoxic activities of 2-aryl-3-((piperidin-1-yl)ethyl)thiazolidinones are reported.

Designing, structural elucidation, comparison of DNA binding, cleavage, radical scavenging activity and anticancer activity of copper(I) complex with 5-dimethyl-2-phenyl-4-[(pyridin-2-ylmethylene)-amino]-1,2-dihydro-pyrazol-3-one Schiff base ligand

pp. 81-89

Subbaiyan Sathiyaraj, Krishnan Sampath, Ray J. Butcher, Raghavaiah Pallepogu and Chinnasamy Jayabalakrishnan*

The Schiff base ligand and copper(I) complex were synthesized and characterized crystallographically. Further, the compounds were evaluated for their pharmacological properties.



 $IC_{50} = 3.04 - 19.25 \,\mu\text{M}$

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