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Biochemical Pharmacology



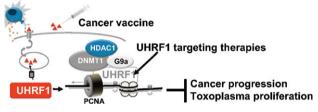


Biochemical Pharmacology, Volume 78, issue 10, 15 November 2009 Contents

COMMENTARY

Drug discovery targeting epigenetic codes: The great potential of UHRF1, which links 1279–1288 DNA methylation and histone modifications, as a drug target in cancers and toxoplasmosis

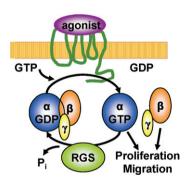
Motoko Unoki, Julie Brunet, Marc Mousli



Regulator of G-protein signaling (RGS) proteins in cancer biology

1289-1297

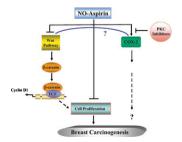
Iillian H. Hurst, Shelley B. Hooks



ANTIBIOTICS AND CHEMOTHERAPEUTICS

Nitro-aspirin inhibits MCF-7 breast cancer cell growth: Effects on COX-2 expression 1298–1304 and Wnt/β-catenin/TCF-4 signaling

Niharika Nath, Rashida Vassell, Mitali Chattopadhyay, Marsel Kogan, Khosrow Kashfi



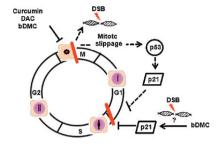
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Curcumin derivatives: Molecular basis of their anti-cancer activity

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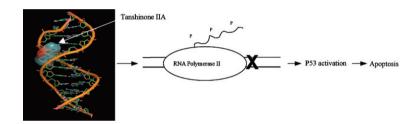
Valentina Basile, Erika Ferrari, Sandra Lazzari, Silvia Belluti, Francesca Pignedoli, Carol Imbriano

Model of the molecular mechanism through which Curcumin, DAC and bDMC induce cell cycle arrest in human colon cancer cells. Bold lines indicate the direct effects of the molecules, dotted lines represent the consequences of the induced mitotic delay.



Tanshinone IIA triggers p53 responses and apoptosis by RNA polymerase II upon DNA 1316–1322 minor groove binding

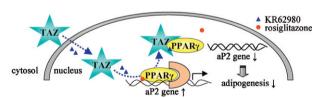
Zhichao Zhang, Jin Gao, Yuanyuan Wang, Ting Song, Jing Zhang, Guiye Wu, Tiantai Zhang, Guanhua Du



METABOLIC DISORDERS AND ENDOCRINOLOGY

Augmentation of PPAR γ -TAZ interaction contributes to the anti-adipogenic activity 1323–1329 of KR62980

Hana Jung, Mi Sook Lee, Eun Jung Jang, Jin Hee Ahn, Nam Sook Kang, Sung-Eun Yoo, Myung Ae Bae, Jeong-Ho Hong, Eun Sook Hwang

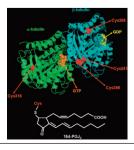


PHARMACOKINETICS AND DRUG METABOLISM

15-Deoxi- $\Delta^{12,14}$ -prostaglandin J_2 is a tubulin-binding agent that destabilizes microtubules and induces mitotic arrest

1330-1339

Claudia Cocca, Jorge Dorado, Enrique Calvo, Juan Antonio López, Angel Santos, Ana Perez-Castillo



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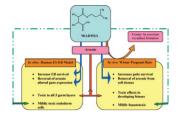
Toxicology

Monoisoamyl dimercaptosuccinic acid abrogates arsenic-induced developmental toxicity in human embryonic stem cell-derived embryoid bodies: Comparison with *in vivo* studies

1340-1349

S.J.S. Flora, Ashish Mehta

The figure summaries the chelation ability of MiADMSA against arsenic when compared in an *in vitro* and *in vivo* model.



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