

Review Article	2455	Immunocyte membrane-derived biomimetic nano-drug delivery system: a pioneering platform for tumour immunotherapy Yuan-ye Zeng, Qing Gu, Dan Li, Ai-xue Li, Rong-mei Liu, Jian-ying Liang and Ji-yong Liu
Articles		
Neuropharmacology	2474	Ginsenoside Rg1 mitigates cerebral ischaemia/reperfusion injury in mice by inhibiting autophagy through activation of mTOR signalling Zhi-chao Xi, Han-gui Ren, Lin Ai, Yuan Wang, Meng-fan Liu, Yu-fei Qiu, Ji-ling Feng, Wang Fu, Qian-qian Bi, Feng Wang and Hong-xi Xu
	2487	Carotid artery transplantation of brain endothelial cells enhances neuroprotection and neurorepair in ischaemic stroke rats Yi-ting Du, Zhi-guang Pan, Bin-chi Chen and Feng-yan Sun
	2497	ErbB4 deficiency exacerbates olfactory dysfunction in an early-stage Alzheimer's disease mouse model Xian-hua Deng, Xing-yang Liu, Yi-hua Wei, Ke Wang, Jun-rong Zhu, Jia-jun Zhong, Jing-yuan Zheng, Rui Guo, Yi-fan Zhu, Qiu-hong Ye, Meng-dan Wang, Ying-jie Chen, Jian-quan He, Ze-xu Chen, Shu-qiong Huang, Chong-shan Lv, Guo-qing Zheng, Sui-feng Liu and Lei Wen
	2513	Transplant of fecal microbiota from healthy young mice relieves cognitive defects in late-stage diabetic mice by reducing metabolic disorders and neuroinflammation Xian-xi Ye, Qiao-ying Jiang, Meng-jun Wu, Qing-huai Ye and Hong Zheng
	2527	Trametinib, an anti-tumor drug, promotes oligodendrocytes generation and myelin formation Ying Yang, Na Suo, Shi-hao Cui, Xuan Wu, Xin-yue Ren, Yin Liu, Ren Guo and Xin Xie
Cardiovascular Pharmacology	2540	In vivo proximity proteomics uncovers palmdelphin (PALMD) as a Z-disc-associated mitigator of isoproterenol-induced cardiac injury Cong-ting Guo, Blake D. Jardin, Jun-sen Lin, Rachele L. Ambrose, Ze Wang, Lu-zi Yang, Neil Mazumdar, Fu-jian Lu, Qing Ma, Yang-po Cao, Can-zhao Liu, Kai-long Li, Xu-jie Liu, Feng Lan, Ming-ming Zhao, Han Xiao, Er-dan Dong, William T. Pu and Yu-xuan Guo
	2553	DLK1 promoted ischemic angiogenesis through notch1 signaling in endothelial progenitor cells Ya-yu You, Ning Zhang, Zhuo Wang, Zhe-hui Yin, Qin-yi Bao, Shu-xin Lei and Xiao-jie Xie
	2567	Geniposide alleviates heart failure with preserved ejection fraction in mice by regulating cardiac oxidative stress via MMP2/SIRT1/GSK3 β pathway Yan-lu Han, Teng-teng Yan, Hua-xin Li, Sha-sha Chen, Zhen-zhen Zhang, Meng-yao Wang, Mei-jie Chen, Yuan-li Chen, Xiao-xiao Yang, Ling-ling Wei, Ya-jun Duan and Shuang Zhang
Hepatic and Renal Pharmacology	2579	SGLT2 inhibitors ameliorate NAFLD in mice via downregulating PFKFB3, suppressing glycolysis and modulating macrophage polarization <i>Open</i> Xia-fang Lin, Xiao-na Cui, Jin Yang, Ya-fei Jiang, Tian-jiao Wei, Li Xia, Xin-yue Liao, Fei Li, Dan-dan Wang, Jian Li, Qi Wu, De-shan Yin, Yun-yi Le, Kun Yang, Rui Wei and Tian-pei Hong
	2598	Chromodomain Y-like (CDYL) inhibition ameliorates acute kidney injury in mice by regulating tubular pyroptosis Ting Xiang, Ling-zhi Li, Jin-xi Li, Xin-yun Chen, Fan Guo, Jing Liu, Yi-ting Wu, Lin Lin, Rui-han Xu, Hui-ping Wang, Liang Ma and Ping Fu
Inflammation and Immunopharmacology	2611	G protein-coupled receptor kinase 2 as a novel therapeutic target for gland fibrosis of Sjögren's syndrome Ru-hong Fang, Zheng-wei Zhou, Rui Chu, Qiu-yun Guan, Feng He, Ming-li Ge, Pai-pai Guo, Hua-xun Wu, Ling-li Yao, Wei Wei, Yang Ma and Qing-tong Wang
Chemotherapy	2625	Oncoprotein LAMTOR5-mediated CHOP silence via DNA hypermethylation and miR-182/miR-769 in promotion of liver cancer growth Xue Wang, Qian-qian Li, Yan-xin Tang, Ye Li, Lu Zhang, Fei-fei Xu, Xue-li Fu, Kai Ye, Jia-qi Ma, Shi-man Guo, Fang-yuan Ma, Zhi-yu Liu, Xu-he Shi, Xian-meng Li, Hui-min Sun, Yue Wu, Wei-ying Zhang and Li-hong Ye

- 2646 Recombinant human adenovirus type 5 promotes anti-tumor immunity *via* inducing pyroptosis in tumor endothelial cells
Zhi-ming Wang, Meng-kai Li, Qing-ling Yang, Shi-xin Duan, Xin-yi Lou, Xin-yi Yang, Ying Liu, Yu-wen Zhong, Yu Qiao, Zi-shu Wang, Lei Sun and Feng Qian
- 2657 Adaptor protein CEMIP reduces the chemosensitivity of small cell lung cancer *via* activation of an SRC-YAP oncogenic module
Xiao-ju Shen, Hui-lan Wei, Xiao-cheng Mo, Xiao-xiang Mo, Li Li, Jing-chuan He, Xin-yu Wei, Xiao-jun Qin, Shang-ping Xing, Zhuo Luo, Zhi-quan Chen and Jie Yang
- 2672 Isotoosendanin inhibits triple-negative breast cancer metastasis by reducing mitochondrial fission and lamellipodia formation regulated by the Smad2/3-GOT2-MYH9 signaling axis
Jing-nan Zhang, Ze Zhang, Zhen-lin Huang, Qian Guo, Ze-qi Wu, Chuang Ke, Bin Lu, Zheng-tao Wang and Li-li Ji

Correction

- 2684 Publisher Correction: E3 ubiquitin ligase UBR5 modulates circadian rhythm by facilitating the ubiquitination and degradation of the key clock transcription factor BMAL1
Chun-yan Duan, Yue Li, Hao-yu Zhi, Yao Tian, Zheng-yun Huang, Su-ping Chen, Yang Zhang, Qing Liu, Liang Zhou, Xiao-gang Jiang, Kifayat Ullah, Qing Guo, Zhao-hui Liu, Ying Xu, Jun-hai Han, Jiajie Hou, Darran P O'Connor and Guoqiang Xu

Cover

Diagram of the proposed mechanisms by which CEMIP reduces the chemotherapy sensitivity of SCLC cells. CEMIP, an adaptor protein, interacts with SRC and YAP. CEMIP binds to SRC, allowing autophosphorylation of Y416 and activation of SRC, which facilitates the interaction between YAP and activated SRC, resulting in increased phosphorylation of Y357, protein stability, nuclear accumulation and transcriptional activation of YAP, thus reducing the chemotherapeutic sensitivity of SCLC cells. The SRC inhibitor dasatinib and the YAP inhibitor verteporfin display synergistic antitumor effects on SCLC both in vitro and in vivo. See the article in pages 2657–2671.

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