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- 740 A structure-based virtual screening identifies a novel MDM2 antagonist in the activation of the p53 signaling and inhibition of tumor growth
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- 751 Longitudinal and time-to-event modeling for the survival of advanced pancreatic ductal adenocarcinoma patients
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Absorption, Distribution, Metabolism, and Excretion

- 759 Circulating metabolites of *Borneolum syntheticum* (Bingpian) ameliorate atherosclerosis in ApoE^{-/-} mice via inhibiting macrophage foam-cell formation
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- 777 Intestinal human carboxylesterase 2 (CES2) expression rescues drug metabolism and most metabolic syndrome phenotypes in global Ces2 cluster knockout mice *Open*
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Correction

- 794 Author Correction: CaIPF14030 negatively modulates intracellular ATP levels during the development of azole resistance in *Candida albicans*
Xin-ming Jia, Ying Wang, Jun-dong Zhang, Hong-yue Tan, Yuan-ying Jiang and Jun Gu

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The constituents of *Borneolum syntheticum* (Bingpian), borneol and isoborneol, are efficiently absorbed but undergo extensive first-pass hepatic metabolism, producing borneol-2-O-glucuronide, isoborneol-2-O-glucuronide, and camphor. These metabolites work synergistically to inhibit macrophage foam-cell formation, reduce atherosclerotic lesions, and enhance plaque stability, suggesting oral Bingpian's potential as a therapeutic agent for atherosclerosis. See the article in pages 759–776.

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