Molecular mechanisms governing different pharmacokinetics of ginsenosides and potential for ginsenoside-perpetrated herb-drug interactions on OATP1B3

Rongrong Jiang, Jiajia Dong, Xiuxue Li, Feifei Du, Weiwei Jia, Fang Xu, Fengqing Wang, Junling Yang, Wei Niu, Chuan Li

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- Supporting Information Appendix S3 -

Time courses for accumulation of the ppt-type ginsenosides in HEK293 cells expressing human OATP1B3 or rat Oatp1b2

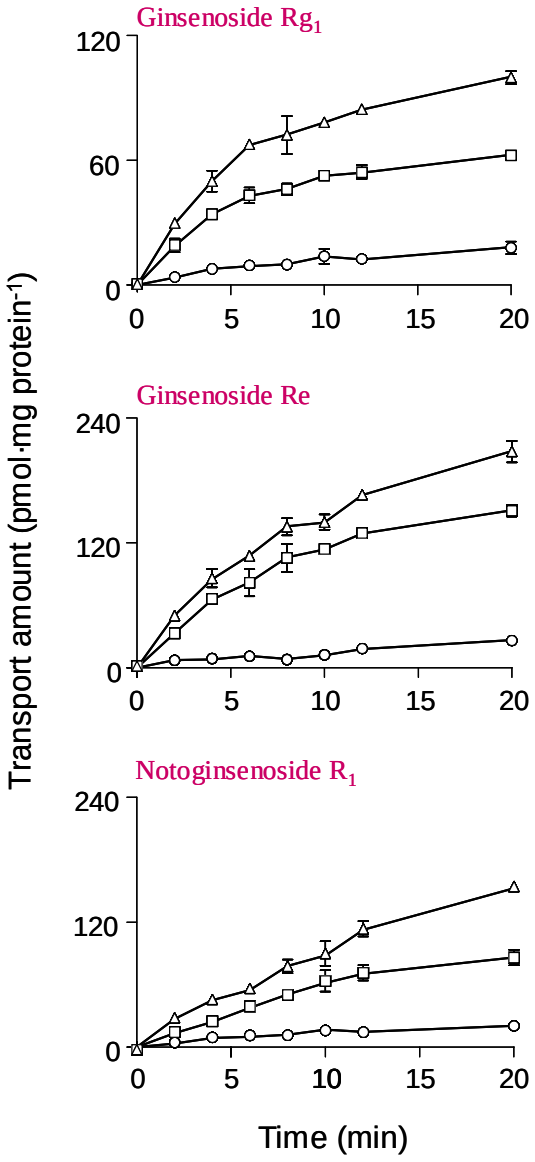

Figure 1

Time courses for accumulation of ginsenoside Rg_1, ginsenoside Re and notoginsenoside R_1 in HEK293 cells expressing pcDNA vector (circle), human OATP1B3 (square) or rat Oatp1b2 (triangle). The initial concentrations of ginsenosides used for incubation were 50 μM. Data represent the means and standard errors and each time point was performed in triplicate.