Ginsenoside Rh1

20(S)-Ginsenoside Rh1

【化合物】Ginsenoside Rh1

【測定機器】ultra performance liquid chromatography/mass spectrometric method

【対象】動物 (mouse)

【代謝実験】The absolute bioavailability of 20(S)-ginsenoside Rh1 in rats was only 1.01%. After intragastrical administration of ginsenoside Rh1, two mono-oxygenated metabolites were detected from the urine, bile, liver tissue, and intestinal tract content, while the de-glucosylated product, 20(S)-protopanaxatriol, was only found in the contents of the intestinal tract. An in vitro incubation study confirmed that the CYP450-catalyzed mono-oxygenation, the intestinal bacteria mediated de-glucosylation, and the gastric acid mediated hydration reaction were the main metabolic pathways of 20(S)-ginsenoside Rh1. [Lai et al., *Planta Med.*, 75: 797–802 (2009)]

【代謝パラメータ】

The main pharmacokinetic parameters of Rh1 after i. v. (5 mg/kg) and i. g. (50mg/kg) administrations (mean \pm S.D., n = 4).

i. v.		i. g.	
Parameters	Rh1	Parameters	Rh1
$T_{1/2_{\alpha}}$ (h)	0.07 ± 0.04	$T_{1/2_{eta}}(\mathrm{h})$	0.43 ± 0.08
$T_{1/2\beta}$ (h)	0.41 ± 0.05	CL/F(L/h)	67.68 ± 9.63

Vd(L)	0.13 ± 0.04	AUC_{0-t} (mg/L · h)	0.13 ± 0.01
<i>CL</i> (L/h)	0.67 ± 0.11	$AUC_{0-\infty}$ (mg/L ·h)	0.14 ± 0.01
AUC_{0-t} (mg/L · h)	1.50 ± 0.23	T_{max} (h)	1.00 ± 0.00
$AUC_{0-\infty}$ (mg/L · h)	1.52 ± 0.24	$C_{ m max}$ (mg/L)	0.05 + 0.01
MRT_{0-t} (h)	0.50 ± 0.05	F(%)	1.01 ± 0.03
$MRT_{0-\infty}$ (h)	0.58 ± 0.06		

CL/F: apparent oral total body clearance (where F represents bioavailability); CL: total clearance; C_{\max} : the maximum plasma concentration; F: bioavailability; MRT: mean retention time; $T_{1/2}$: the absorption half-life time; $T_{1/2}$: the apparent elimination half-life time; T_{\max} : the time to reach C_{\max} ; Vd: the apparent volume of distribution. [Lai et al., $Planta\ Med.$, 75: 797–802 (2009)]

【参考文献】

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Li Lai, Haiping Hao, Yitong Liu, Chaonan Zheng, QiongWang, Guangji Wang, Xijin Chen, Characterization of pharmacokinetic profiles and metabolic pathways of 20(*S*)-ginsenoside Rh1 in vivo and in vitro. *Planta Med.*, **75**: 797–802 (2009).