

Investigation the distributions and pharmacokinetics of five *Rhubarb* anthraquinones in rabbits and rats

Abstract

The main effective components of *Rhubarb* are anthraquinones (AQs), most of which are glycoside and others are free. The concentrations of AQs derivatives (rhein, aloe-emodin, emodin and chrysophanol) in plasma and homogenate were assayed with a high performance liquid chromatography (HPLC) method. The pharmacokinetic parameter and distributions of *rhubarb* AQs in rabbits or rats were obtained after given different formulas respectively. The elimination of AQs was fit to a two-compartment model in rats and rabbits. There were no significant differences in the main pharmacokinetic parameters between with rhein and AQs in rats. AQs were distributed in kidney, liver, blood, heart progressively. The AQs were mainly existed as rhein *in vivo* and excreted by kidney. To the formulas containing *rhubarb*, rhein could be used as a probe for pharmacokinetic study of *Rhubarb* AQs *in vivo*.

Keywords: *rhubarb*, anthraquinones, pharmacokinetics, distribution, chrysophanol, aloe-emodin, *in vivo*

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Introduction

Rhubarb (Dahuang in Chinese), a well-known Chinese herbal medicine, has long been used in the world and is officially listed in Chinese Pharmacopoeia which containing three species, *Rheum palmatum* L., *Rheum officinale* Bail and *Rheum tanguticum* Maxim. ex Balf.¹ It is also formally listed in European and Japanese Pharmacopoeia.^{2,3} This important herbal medicine has been discerned for centuries in traditional medicine for its multiple pharmacological actions include antibacterial, laxative, antispasmodic and hemostatic.^{4,5} The main effective components of *Rhubarb* are five anthraquinones, namely rhein, emodin, aloe-emodin, chrysophanol and physcion. Most of which are glycoside. More than 16 pure compounds which about 3-5% constituents of *Rhubarb* have been separated. These are the basis for the quality control of *rhubarb* and other plant drugs from *Rheum*, *Cassia* and *Polygonum* genera. The chemical structures of these AQs and internal standard (IS) are shown in Figure 1. We designed the experiment for tentative discussing the pharmacokinetic parameters and the excretion of *Rhubarb* AQs *in vivo*.

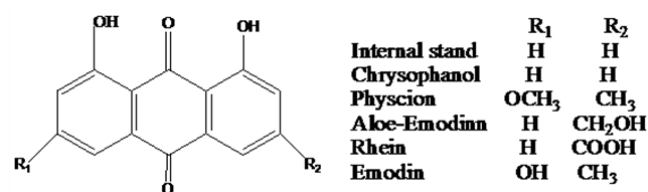


Figure 1 The chemical structures of rhein, aloe-emodin, emodin, internal standard, chrysophanol and physcion.

The methods commonly used for the separation and determination of all or some of the five major AQs in *rhubarb* are micellar electrokinetic chromatography (MEC),^{6,7} capillary zone electrophoresis⁸ and high performance liquid chromatography (HPLC).⁹⁻¹⁸

In our studies, we established a reliable and rapid method for simultaneous assay of five AQs in *rhubarb* on plasma and tissue homogenate and its related preparations using HPLC. Because of its sensitivity and speediness, HPLC is obtaining considerable attention in recent years for pharmaceutical and biomedical analysis. The results of our studies provided a reference for investigation of pharmacokinetics

and distributions and of relations between pharmacokinetics, distributions and clinical efficacy of *Rhubarb* anthraquinones in human body.

Materials and methods

Materials

Rhein, Aloe-emodin, Emodin, Chrysophanol and physcion were obtained from the national institutes for food and drug control, 1,8-dihydroxyanthraquinone (internal standard, IS), acetonitrile, methanol, phosphoric acid, sulfuric acid, and ethyl ester were of analytical grade.

Subjects

6 healthy rabbits, males, weight 2.0kg - 3.0kg, and 30 male and 30 female SD rats, weight 150g-200g were purchased from School of Pharmacy, Chongqing Medical University, and R.P China. They were individually housed in stainless steel cages and acclimatized for 3 days prior to experiments which were kept on a 12 h light /dark cycle, allowed free access to water prior to oral gavage and without drinking during the experiment.

Method of oral gavage administration

WPY granule formula composed of *Rhubarb*, Ginseng, Betel nuts and Pharbitic was a novel drug which was used to treat acute pancreatitis. The particles contain Aloe-emodin 7.9824mg/g, rhein 12.2679mg/g, Emodin 7.3886mg/g, Chrysophanol 9.4768mg/g and Physcion 6.3278mg/g. The WPY dried powder was adjusted with water to equivalent as 0.763g original *rhubarb* herbs including rhein 4.50mg/mL. Preserved in refrigerator for rabbit and rats oral gavage.

The extract solution of *Rhubarb* was prepared as follow: *Rhubarb* was extracted twice with 70% ethanol, filtrated; the filtrate was concentrated to the relative density of 1.05 to 1.10. Which contained Aloe-emodin 4.0097mg/g, rhein 11.8765mg/g, Emodin 3.7676mg/g, Chrysophanol 3.3462mg/g and Physcion 2.3001mg/g? The liquid extract solution was adjusted to equivalent as 0.833g of native herbs including rhein 4.51mg/mL. Preserved in refrigerator for rabbits and rats oral gavage.

6 Healthy rabbits and 60 healthy SD rats were divided randomly into two groups, respectively. Blood was gotten and preserved in a tube which had added sodium heparin after rabbits had oral gavage of the extract solution of *Rhubarb* and the compounding formula solution of WPY respectively. Plasma and heart, liver and kidney tissues samples were preserved in refrigeration at -20°C till analysis within two weeks.

Preparation for plasma and tissues samples

Each biological sample was added 100mL internal standard solution and 1.0mL of 15% sulfuric acid and mixed. First, the samples were placed in $70\pm 2^{\circ}\text{C}$ water bath for 30min to hydrolysis conjunctive anthraquinones. Then, the samples were added 5.0mL ethyl ether to extract *Rhubarb* anthraquinones, after vigorous vortexing 3min, the samples were centrifuged at $10,000\times g$ for 10 min at -4°C , the supernatant was drawn and dried with nitrogen at 40°C . Finally, the clear supernatant was reconstituted with 100mL mobile phase and 50mL supernatant was injected into the HPLC instruments.

Analytical method

The analytical method was established as follow: HPLC instrument with a UV detector (Shimadzu LC-2010) and Hypersil C18 column (250mm \times 4.6mm i.d. Dalian Elite Analytical Instruments Co., Ltd. China) were used for analysis. The mobile phase components were methanol- acetonitrile-water (10:50:40, v: v: v, adjusted to pH 2.8 with phosphoric acid), the detective wavelength was 225nm. The flow-rate was 1.0mL/min. Plasma and homogenate concentrations of anthraquinones were measured by using the established method. The ratios of peak areas of aloe-emodin, rhein, emodin and chrysophanol to I.S. were plotted versus the concentrations of aloe-emodin, rhein, emodin and chrysophanol.

Statistical and Pharmacokinetic analysis

The pharmacokinetic parameters in rabbits and rats, intergroup comparison were calculated by DAS version 2.0 which bought from the Center for Drug Clinical Evaluation of Anhui Medical University, China. Statistical analysis was performed with the two-sided t-test and paired samples test. As the plasma pharmacokinetic parameter of rhein after oral gavage displayed a typical bi-exponential decline, the plasma concentration-time data was fitted into the classical two-compartment first-order open model ($C=Ae^{-\alpha t}+Be^{-\beta t}$). The plasma exposure (area under the plasma tetrandrine concentration-time curve (AUC_{0-24h}), clearance (Cl_{0-24h}), mean transit time (MTT_{0-24h}) and terminal elimination half-life ($t_{1/2}$) were calculated.

Results and discussion

Anthraquinones existed form of before and after oral gavage administration

Most of the five anthraquinones are glycoside in *Rhubarb*, In order to determination the total anthraquinones we added 1.0mL of 15% sulfuric acid in each samples and placed it in $70\pm 2^{\circ}\text{C}$ water bath for 30min to hydrolysis conjunctive anthraquinones. The exits forms of anthraquinones in the extract solution of *Rhubarb* and WPY compounding formula solution before rabbits and rats oral gavage were shown in Figure 2 and that after oral were shown in Figure 2A-2F. From the chromatogram we can see that anthraquinones can be determined under the separation conditions and there were no interference.

Pharmacokinetics after oral gavage administration

Plasma was gotten from ear vein after rabbits had oral gavage at 0,

0.25, 0.5, 1, 2, 3, 4, 6, 9, 12, 24 hours. The concentration of rhein in rabbits plasma vs time curves were shown in Figure 3.

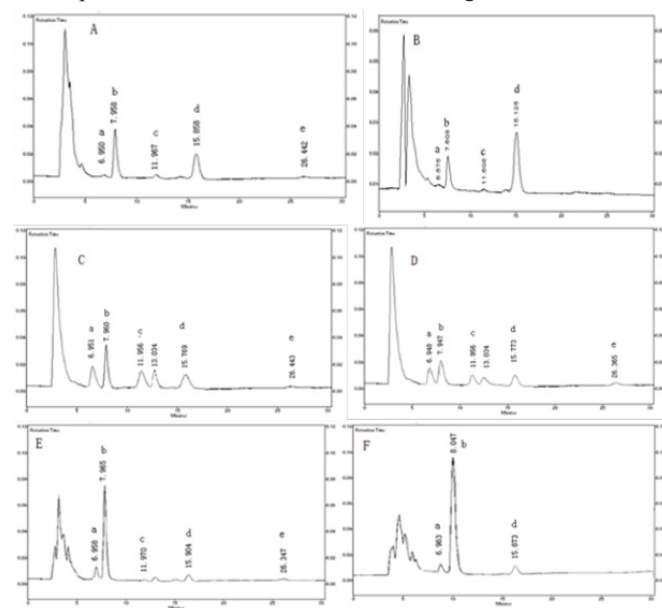


Figure 2A: A plasma sample taken from a rabbit after receiving an oral gavage of the extract solution of *Rhubarb*;

Figure 2B: A plasma sample taken from a rabbit after oral gavage of WPY,

Figure 2C: The extract solution of *Rhubarb* before oral administration;

Figure 2D: WPY before oral administration.

Figure 2E: A plasma sample taken from a rat after receiving an oral gavage of the extract solution of *Rhubarb*;

Figure 2F: A plasma sample taken from a rat after oral gavage of WPY.

Figure 2 The chromatography of *Rhubarb* anthraquinones in rabbits and rats plasma were achieved on a Hypersil C18 column (10mm, 4.6mm \times 250mm i.d.).

The mobile phase consisted of methanol, acetonitrile and water (10:50:40, v/v, pH2.8) at a flow rate of 1.0mL/min, and the detective wavelength was 225nm.

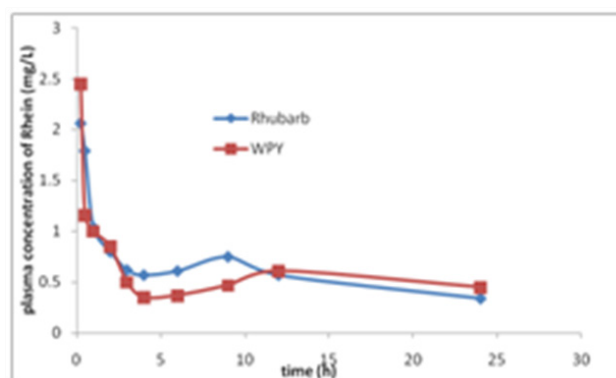


Figure 3 The curves of mean rhein concentration-time in rabbits plasma after oral gavage the extract solution of *Rhubarb* or WPY.

The pharmacokinetic parameters in rabbits after oral gavage of the extract solution of *Rhubarb* or compounding formula (WPY) were shown in Table 1. From Table 1 we can see that there were no significant differences ($p<0.05$) in the metabolism procedures of *Rhubarb* anthraquinones after rabbits had oral gavage different formulas. Plasma was obtained after rats had oral gavage of different formula at 5min, 0.083, 0.25, 0.5, 1, 2, 3, 4, 6, 12, 24hours. The concentration-time curves of rhein and total AQS in rat's plasma were shown in Figure 4. We can know that rhein increased and others decreased in rabbits and rats after oral gavage of different formulas

containing *Rhubarb*. The elimination of total AQs was fit to a two-compartment model in rats. The Pharmacokinetic parameters of rhein and its comparison result were listed in Table 2&3 and Figure 5. The comparison results of pharmacokinetic parameters of rhein in rats and in rabbits plasma were shown in Figure 5. From Figure 5 and Table 4 we can see that there were no significant differences in the main pharmacokinetic parameters in rats after oral gavage of different formulas containing *Rhubarb*. That is, the metabolism procedures of *Rhubarb* anthraquinones in different formulas can be described with rhein. From Figure 5 we can see that there were no significant differences between rats and rabbits after oral gavage of different formulas which containing *Rhubarb* anthraquinones from the metabolism procedures. Pharmacokinetic study on single-*rhubarb* and compound formula of anthraquinone derivatives showed that the metabolism in rabbits and rats are consistent with two-compartment model; the pharmacokinetic parameters calculated with rhein can represent the behalf of *Rhubarb* Anthraquinones metabolism in the biological process. The results of biological samples study can extrapolated that *Rhubarb* AQs in the metabolic process *in vivo* may be carried out as Figures 6&7 shown. Combination of oxidation and metabolism. Oxidation is the 3-methyl (Chrysophanol) have been oxidized to hydroxymethyl (aloe-emodin) and carboxyl (rhein).

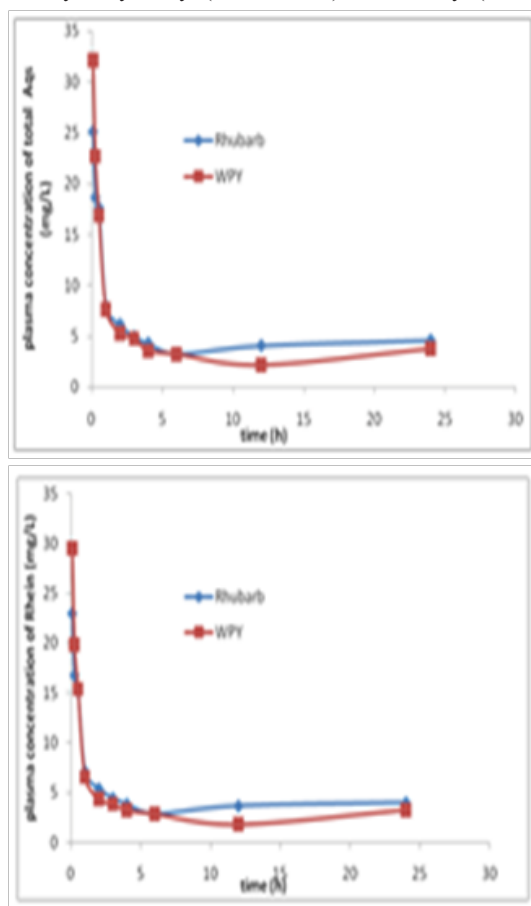


Figure 4 The curves of rhein or total AQs concentration-time in rats plasma after oral gavage of the extract solution of *Rhubarb* or WPY.

Tissue distribution study on rats after oral gavage

Tissue homogenates were obtained after rats had oral gavage different formula at 1, 2, 3, 4, 6, 12 hours. The concentrations of rhein in different tissues were obtained by HPLC and the data were

shown in Table 5. From Table 5, we can see that total AQs and rhein were distributed in kidney, liver, blood and heart progressively. The anthraquinones were mainly existed as rhein *in vivo* and were excreted by the kidneys. The studies of distributions of *Rhubarb* anthraquinones were performed on 30 healthy rats after oral gavage of two formulas respectively. The results showed that *Rhubarb* anthraquinones in rats were distributed in kidney, liver, blood, heart progressively. It confirmed that the anthraquinones were mainly existed in rhein *in vivo* and excreted by kidney. The metabolic pathway of emodin may be described as Figure 7 shown. The metabolic processes can explain why the biological sample with a high concentration of rhein, and that of aloe-emodin and chrysophanol with low even do not detected.

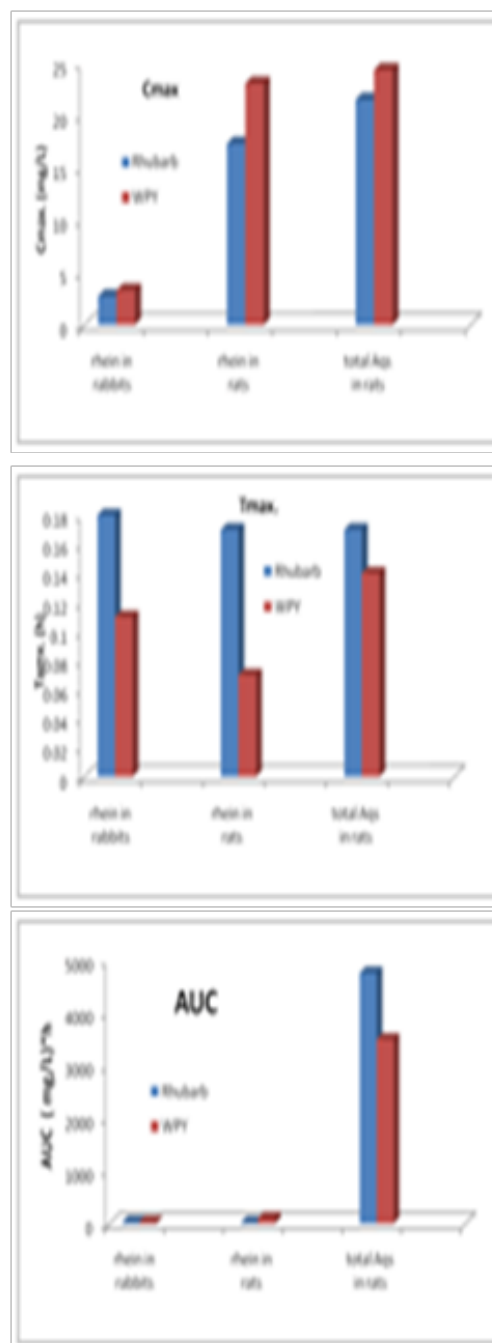


Figure 5 The comparison of the main pharmacokinetic parameters in rats and rabbits.

Table 1 The pharmacokinetic parameters in rabbits after oral gavage of the extract solution of *Rhubarb* and WPY respectively

Pharmacokinetic parameters compartment model	Extract solution of rhubarb two compartment	WPY two compartment
Weighted Coefficient	1	1
A (mg/L)	4.1728±3.3881	6.4958±4.0837
α (1/np)	2.4617±1.0525	4.6488±4.2997
B (mg/L)	0.7331±2.2870	0.6419±0.1647
β (1/np)	0.0247±0.0782	0.0250±0.0272
K _a (1/hr)	6.5799±7.2936	13.4752±8.6081
V/F _(g) (10ml/kg)/(mg/L)	2.9922±4.0194	2.0427±2.4699
t _{1/2(a)} (hr)	0.2816±1.3778	0.1491±0.4757
t _{1/2(b)} (hr)	28.1055±16.2102	27.7543±22.8011
t _{1/2(k_a)} (hr)	0.1053±0.2860	0.0514±0.0210
K ₂₁ (1/hr)	0.5573±0.2609	0.6301±0.2036
K ₁₀ (1/hr)	0.1089±0.0520	0.1843±0.1670
K ₁₂ (1/hr)	1.8202±0.8233	3.8594±4.3104
AUC (mg/ml)*hr	30.6755±4.5517	26.5689±8.6120
CL _(s) (1ml/100g)/hr/(mg/L)	0.3260±0.0437	0.3764±0.5739
T _(peak) (hr)	0.2800±3.1190	0.1275±0.0232
C _{max} (mg/L)	2.0453±1.2900	2.9512±1.7254

Table 2 Pharmacokinetic parameters with rhein or total AQs in rats after oral gavage of the extract solution of *Rhubarb* and WPY respectively

Pharmacokinetic parameters	Extract solution of rhubarb rhein	Total of AQs	WPY rhein	Total of AQs
Compartment Model	Two compartment			
Weighted Coefficient	1/C2			
A (mg/L)	22.5419±20.8715	22.1741±10.3235	28.8972±13.7067	39.5559±6.6164
α (1/np)	1.4575±0.9118	1.3214±0.7375	1.9421±0.3740	2.3534±0.5111
B (mg/L)	3.1455±0.4490	3.7652±1.5683	3.2091±1.2954	4.1697±0.8693
β (1/np)	-0.0119±0.0142	-0.0082±0.0407	0.0192±0.0336	0.0281±0.0071
K _a (1/hr)	18.6264±39.2112	43.8505±1.9578	44.0966±89.1147	19.3712±38.2309
V/F _(g) (1ml/100g)/ (mg/L)	0.0418±0.0134	0.0396±0.0222	0.0324±0.0377	0.0257±0.0062
t _{1/2(a)} (hr)	0.4756±0.3074	0.5246±1.1632	0.3569±0.1153	0.2945±0.1140
t _{1/2(b)} (hr)	-58.1627±76.5560	-84.771±10.081	36.0917±29.4595	24.631±122.1592
t _{1/2(k_a)} (hr)	0.0372±0.0635	0.0158±0.0182	0.0157±0.0131	0.0358±0.0079
K ₂₁ (1/hr)	0.1814±0.0671	0.1899±0.1395	0.2193±0.0586	0.2769±0.0170
K ₁₀ (1/hr)	-0.0958±0.1280	-0.0569±0.4931	0.1701±0.3426	0.2391±0.1146
K ₁₂ (1/hr)	1.3599±0.9407	1.1801±0.9179	1.5720±0.0371	1.8654±0.5882
AUC (mg/L)*hr	26.0291±310.2810	4753.93±5752.36	181.248±115.889	3494.16±6416.72
CL _(s) (1ml/100g)/hr/(mg/L)	-0.0040±0.0053	-0.0023±0.0303	0.055±0.0256	0.0061±0.0039
T _(peak) (hr)	0.1583±0.1170	0.1698±0.1083	0.0746±0.0511	0.1364±0.0300
C _{max} (mg/L)	19.7035±1.2258	21.5728±2.1430	27.0116±13.6160	24.3986±5.3003

Table 3 The results of paired samples test of pharmacokinetic parameters in rabbits (rhein)

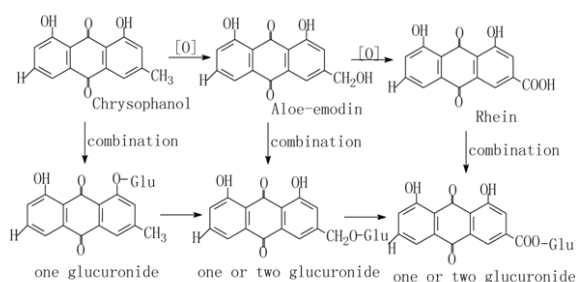
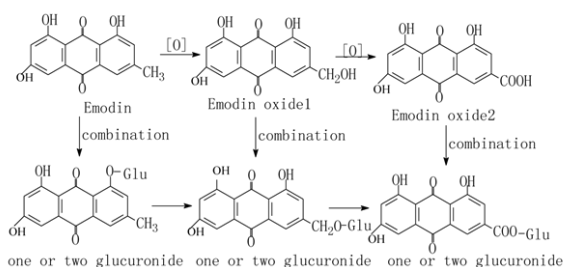
No.	AUC	T _{max}	C _{max}
	Extract Solution of Rhubarb WPY	Extract Solution of Rhubarb WPY	Extract Solution of Rhubarb WPY
\bar{x}	33.16	34.4 0.18	0.11 2.78
SD	4.55	25.29 0.15	0.02 1.29
2-tailed t-test	t=17.86	t=3.33 t=3.07	t=3.61 t=5.29
t-test point value	t _{0.05} =4.303 t _{0.001} =31.599		
Paired t-test	0.144	0.149	1.525
Paired t-test P value	0.895	0.294	0.405

Table 4 The results of paired samples test of pharmacokinetic parameters in rats (total AQs)

No.	AUC	T _{max}	C _{max}
	Extract Solution of Rhubarb WPY	Extract Solution of Rhubarb WPY	Extract Solution of Rhubarb WPY
\bar{x}	4753.93	3494.16 0.17	0.14 21.57
SD	5752.36	26.07 0.11	0.03 2.14
2-tailed t-test	t=2.02	t=-1.33 t=3.85	t=11.00 t=24.68
t-test point value	t _{0.05} =4.303 t _{0.001} =31.599		
Paired t-test	1.784	1.025	3.085
Paired t-test P value	0.172	0.381	0.054

Table 5 The data of rhein concentration in 30 rats after oral gavage of the extract solution of *Rhubarb* and WPY respectively

	Samples	Time (h)/Concentration (mg/kg)				
		1	2	4	6	12
Extract Solution of <i>Rhubarb</i>	Homogenates of heart	1.42	5.59	4.25	1.36	4.03
		4.79	6.38	1.81	1.55	0.28
		20.64	9.76	0.49	0.56	0.35
	Homogenates of liver	2.49	2.67	1.96	1.83	2.78
		4.56	5.56	3.86	0.9	2.95
		0	10.1	0.88	1.09	1.64
	Homogenates of kidney	22.1	21.12	16.51	20.8	15.58
		28.81	12.77	7.4	1.07	1.79
		11.73	15.33	2.13	1.14	0.14
WPY	Homogenates of heart	3.46	2.55	2.68	4.38	0.62
		6.36	2.53	3.88	0.83	0.39
		3.47	8.02	0.69	0.17	0.93
	Homogenates of liver	7.07	6.38	2.42	5.99	3.45
		2.58	0.93	4.04	2.63	4.18
		1.07	8.02	1.19	0.86	1.16
	Homogenates of kidney	35.71	15.06	13.23	14.77	11.84
		7.36	3.91	14.41	2.6	2.71
		14.67	11.38	2.24	0.83	1.42

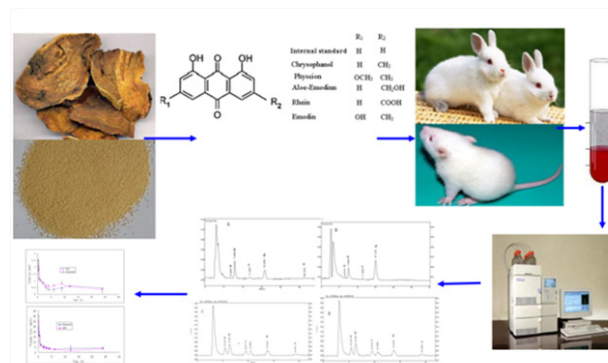
**Figure 6** The metabolic pathway of chrysophanol in vivo after rats oral gavage of different preparation containing *Rhubarb*.**Figure 7** The metabolic pathway of emodin in vivo after rats oral gavage of different preparation containing *Rhubarb*.

In view of the complexity of Chinese herbal compound, a comprehensive picture of what all of its chemical constituents of the absorption, distribution, metabolism and excretion of the law is very difficult. To clarify the effectiveness of components can be detected and their dynamic changes *in vivo*, the efficacy of Chinese medicine composed of clear material basis, Compatibility of the law reform and clinical forms are useful medication.

Conclusion

The results of our studies could provide references for investigation of pharmacokinetics and distributions and of relations between pharmacokinetics, distributions and clinical efficacy

of *Rhubarb* anthraquinones in human body, and provide a new thinking method for exploit and application of the traditional Chinese medicine preparations. And to the preparations containing *Rhubarb*, rhein could be used as a probe for pharmacokinetic study of anthraquinones *in vivo* (Figure 8).

**Figure 8** Graphical Abstract.

Acknowledgments

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Conflicts of interest

The authors declare no conflict of interest.

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None.

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