

# Response surface modeling and optimization of ultrasound-assisted extraction of three flavonoids from tartary buckwheat (*Fagopyrum tataricum*)

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## ABSTRACT

**Background:** Buckwheat (*Fagopyrum* spp., Polygonaceae) is a widely planted food crop. Flavonoids, including quercetin, rutin, and kaempferol, are the main bioactive components in tartary buckwheat (*Fagopyrum tataricum* (L.) Gaertn). From the nutritional and pharmacological perspectives, flavonoids have great value in controlling blood glucose and blood pressure levels, and they also have antioxidant properties. **Objective:** To optimize the conditions for extraction of quercetin, rutin, and kaempferol from *F. tataricum*. **Materials and Methods:** A combination of ultrasound-assisted extraction (UAE) and response surface methodology (RSM) was used for flavonoid extraction and yield assessment. The RSM was based on a three-level, three-variable Box-Behnken design. **Results:** Flavonoids were optimally extracted from *F. tataricum* by using 72% methanol, at 60°C, for 21 minutes. Under these conditions, the obtained extraction yield of the total flavonoids was 3.94%. **Conclusion:** The results indicated that the UAE method was effective for extraction of flavonoids from tartary buckwheat.

**Key words:** Buckwheat, flavonoids, *Fagopyrum tataricum*, response surface methodology, ultrasound-assisted extraction

## INTRODUCTION

Buckwheat (*Fagopyrum* spp., Polygonaceae) is widely planted as a food crop. Tartary buckwheat (*Fagopyrum tataricum* (L.) Gaertn), an edible and medicinal crop, is becoming increasingly popular because of its benefits to the human body.<sup>[1-3]</sup> Tartary buckwheat is receiving widespread attention as a functional food,<sup>[4]</sup> and a number of commercial buckwheat products are now being produced and distributed. Buckwheat contains many beneficial components, such as, flavonoids, fagopyrins, and D-*chiro*-inositol.<sup>[5,6]</sup> Such components have been reported to help control blood glucose<sup>[7]</sup> and blood pressure levels.<sup>[8]</sup> Moreover, buckwheat (*Fagopyrum esculentum* Moench) may be useful in the treatment of cancer.<sup>[9]</sup> Considerable research done currently on buckwheat is focused on its health efficacy and on component extraction processing.<sup>[10-13]</sup> The amount of total flavonoids (quercetin, rutin, and

kaempferol) contained in *F. tataricum* is reported to be far higher than that in common buckwheat (*F. esculentum* Moench).<sup>[14]</sup> Rutin and quercetin are the most intensely studied flavonoids in tartary buckwheat, due to their functions and high concentrations.<sup>[15-18]</sup>

At present, various extraction techniques have been developed for the extraction of flavonoids from tartary buckwheat, including, ultrasound-assisted extraction (UAE), microwave-assisted extraction, and oscillation extraction.<sup>[19-21]</sup> The UAE method is a simple, rapid extraction technique, with high extraction efficiency, which is attributed to the effect of acoustic cavitation produced in the solvent by the passage of an ultrasound wave.

Response surface methodology (RSM) is a powerful statistical technique that is useful when optimizing processes in the fields of medicine and nutrition. It has been reported that RSM can be used to optimize complex processes used to extract compounds from plants.<sup>[22-27]</sup> To our knowledge, there are no reports on the use of RSM to optimize extraction conditions for quercetin, rutin, and kaempferol in buckwheat. In this study, we have focused on establishing a rapid and convenient method for extracting

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and quantifying three of the flavonoids (quercetin, rutin, and kaempferol) present in *F. tataricum*. The method uses UAE and RSM, which is based on a three-level, three-variable (extraction time, extraction temperature, and methanol concentration) Box-Behnken design (BBD). The results should be helpful in the further utilization of flavonoids from tartary buckwheat.

## MATERIALS AND METHODS

### Chemicals and reagents

Quercetin, rutin, and kaempferol used as reference standards were purchased from the National Institute for the Control of Pharmaceutical and Biological Products (Beijing, China). Acetonitrile and methanol (high performance liquid chromatography (HPLC) grade) were purchased from Fisher Scientific Co. (USA). All others chemicals and solvents used in the study were of analytical grade.

### Plant material

The samples of *F. tataricum* were harvested from the experimental farm of Chengdu University, Chengdu, Sichuan, China, in November 2011. The species identification was authenticated by Professor Zhao Gang (Chengdu University). The obtained tartary buckwheat seeds were dried, ground, and then passed through the sieve screen. The powder obtained from the 20 and 40 mesh sieve screens was subjected to UAE extraction.

### Ultrasound-assisted extraction

Ultrasound-Assisted Extraction was performed by mixing 0.1 g of the sieved, dried powder with 25 ml of a predetermined concentration of ethanol in a single conical flask, followed by weighing and ultrasonic extraction for a predetermined time, at a predetermined temperature. The ultrasonic device parameters were 200 W of power at a frequency of 50 kHz. Following extraction, the mixture was cooled to room temperature. Subsequently, the extract was filtered and the filtrate collected for HPLC assessment.

### Experimental design

The preliminary ranges of the extraction variables, namely, methanol concentration ( $X_1$ ), extraction time ( $X_2$ ), and extraction temperature ( $X_3$ ), were established by using a single-factor test. Subsequently, a Box-Behnken factorial design (BBD; Design-Expert software, version 7.1.6, Stat-Ease, Minneapolis, MN, USA) with three levels and three variables was applied, to determine the best UAE conditions for optimizing flavonoid yield. The adjusted R-squared ( $R^2$ ) values along with the  $F$ -test results and probability ( $p$ ) values were used to evaluate the results of the model equations.

### High performance liquid chromatography analysis

The UAE-obtained *F. tataricum* extracts were passed through 0.45  $\mu\text{m}$  filters and then placed in an HPLC autosampler vial for immediate HPLC analysis. The rutin, quercetin, and kaempferol reference standard solution was prepared by dissolving rutin, quercetin, and kaempferol in 70% methanol. The HPLC system was comprised of two Shimadzu LC-20A pumps and a Shimadzu LC-20A autosampler (Kyoto, Japan). A Diamonsil-ODS  $C_{18}$  (250 mm  $\times$  4.6 mm  $\times$  5  $\mu\text{m}$ ) column was used. The temperature of the column was 30°C. Separation was performed by using a mixture of acetonitrile and distilled water containing 0.3%  $\text{H}_3\text{PO}_4$ , with a gradient elution: 0–8 minutes (20% acetonitrile), 8–13 minutes (20–40% acetonitrile), 13–29 minutes (40% acetonitrile), 29–29.1 minutes (40–20% acetonitrile), and 29.1–30 minutes (20% acetonitrile). The flow rate was set at 1 ml/minute. The eluent was obtained after the column was sent to a UV/VIS detector (Shimadzu, Kyoto, Japan). The detector wavelength was set at 365 nm.

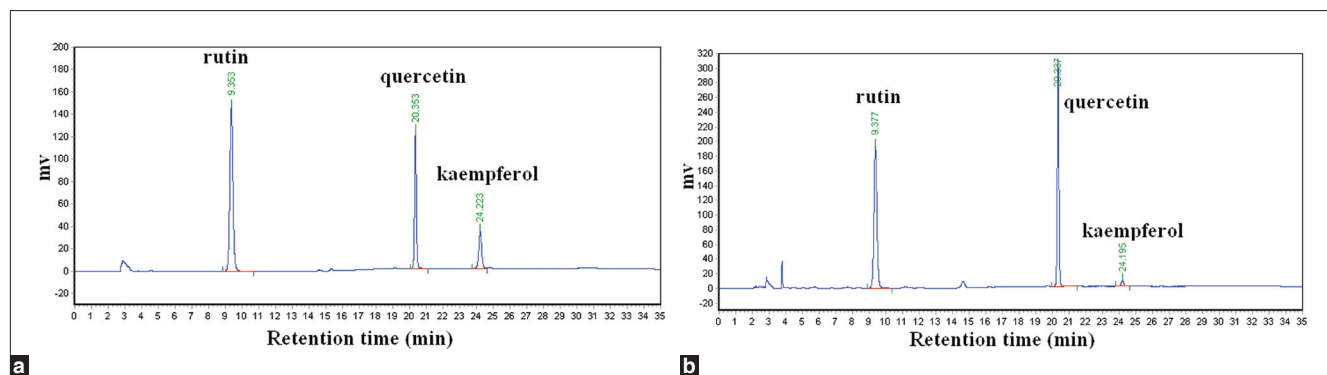
## RESULTS AND DISCUSSION

### Chromatographic results

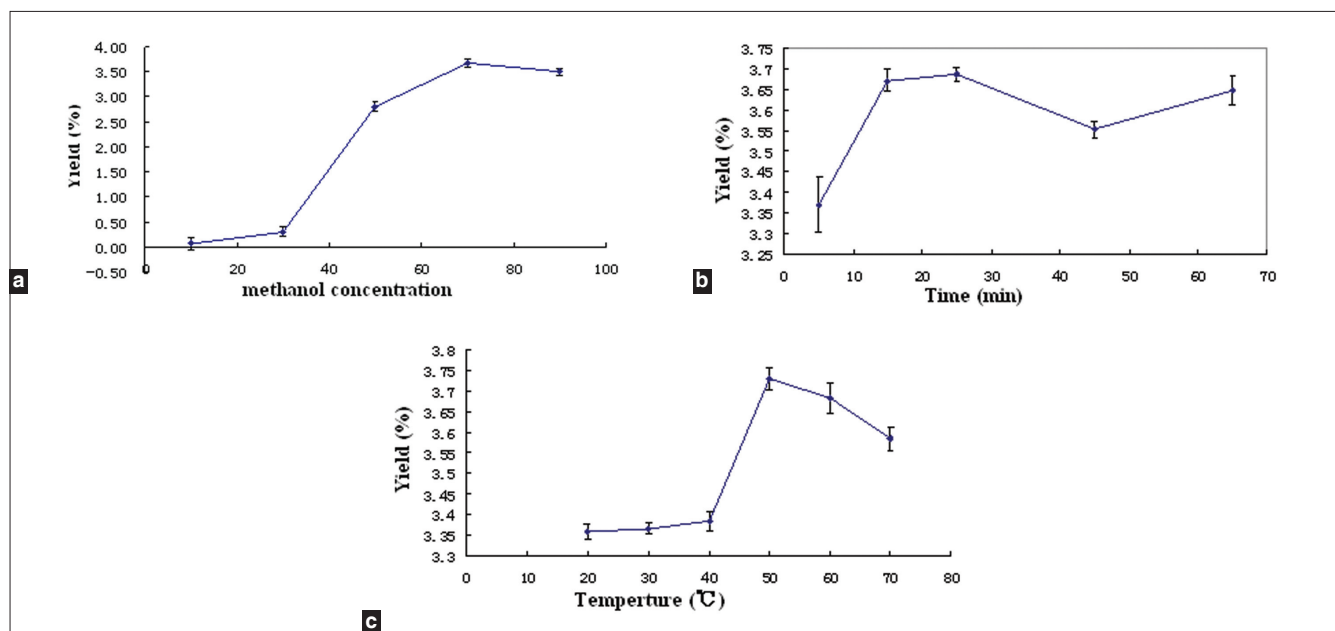
Chromatography images from the reference standard flavonoids and from the UAE-extracted tartary buckwheat sample are shown in Figure 1. Chromatographic results from the *F. tataricum* sample [Figure 1b] show that quercetin, rutin, and kaempferol were separated well, with a retention time of 9.337 minutes, 20.337 minutes, and 24.195 minutes, respectively. The total flavonoid yield was the total of the individual yields of the three assessed flavonoids: quercetin, rutin, and kaempferol.

### Selection of solvent

Flavonoids are normally extracted with methanol and ethanol. In this study, we attempted to determine the extraction solvent that produced the highest flavonoid yield. Our results indicated that the extraction yield of total flavonoids was higher when methanol was used as the extraction solvent (3.15%) than when ethanol (3.04%) was used. Moreover, methanol concentration was important to investigate the effect of methanol concentration on flavonoid yield; five different methanol concentrations were compared [Figure 2a]. When the methanol concentration was increased from 10 to 50%, the yields of quercetin and kaempferol reached their peak, but the yield of rutin was low. This could be because rutin could degrade quickly in low methanol concentrations due to the rutin-degrading enzyme contained in tartary buckwheat.<sup>[28]</sup> When the methanol concentration increased from 50 to 70%, the rutin yield reached a maximum. The yield slightly decreased at methanol concentrations over 70%. Hence,



**Figure 1:** The HPLC profiles of quercetin, rutin, kaempferol standard substance (a) and tartary buckwheat extraction (b)



**Figure 2:** The effects of the extraction parameters on flavonoids yield: (a) Effect of methanol concentration on flavonoids yield. Other conditions were 50 °C extraction temperature and 25 minutes extraction time. (b) Effect of extraction time on flavonoids yield. Other conditions were 70% methanol and 50 °C extraction temperature. (c) Effect of extraction temperature on flavonoid yield. Other conditions were 70% methanol and 25 minute extraction time

70% methanol was chosen for testing in the subsequent optimization experiments.

### Extraction time

Extraction time is another important factor that influences extraction yield, as usually, extraction yield increases with extraction time. In this study, UAE extraction was carried out for different durations (5 – 65 minutes). The extraction yield increased markedly when the duration increased from five minutes to 15 minutes [Figure 2b]. However, the yield remained approximately constant at durations from 15 minutes to 65 minutes. On the basis of these results, the extraction time was set at 15 minutes in the optimization experiments.

### Extraction temperature

Temperature could also influence the extraction yield,

as an increase in temperature could accelerate the extraction speed.<sup>[24]</sup> In this study, five different extraction temperatures were tested [Figure 2c]. The results indicated that the extraction yield increased markedly at extraction temperatures of 40°C and 50°C, but slightly decreased at temperatures above 50°C and higher. We speculated that, due to the actions of the rutin-degrading enzyme, the speed of rutin degradation may be accelerated when the temperature was lower than 50°C. Hence, 50°C was chosen for testing in the subsequent optimization experiments Figure 2.

### Optimization of flavonoid yield

In order to determine the best extraction conditions, we performed parameter optimization through RSM. The RSM approach was based on the three levels and three variables described in Table 1. During our assessment, 12

factorial experiments were performed, along with three zero-point tests, to allow error estimation. Table 2 shows the results of those 15 experiments with the total flavonoid yield ranging from 1.88 to 3.92%. The results indicated that the best yield was obtained by performing UAE for 21 minutes at 60°C with a methanol concentration of 72%. The regression equation between the flavonoid yield (Y) and variables  $X_1$ ,  $X_2$ , and  $X_3$  was:

$$Y = 3.73 + 0.37 X_1 + 0.30 X_2 + 0.050 X_3 + 0.43 X_1 X_3 - 0.17 X_2 X_3 - 0.32 X_1^2 - 0.077 X_2^2 - 0.71 X_3^2$$

The analysis of variance (ANOVA) results for the regression model are shown in Table 3. The *Model F*-value of 70.30 implied that the model's relationships were significantly relative to the noise in the data, as there was only a 0.01% chance that a *Model F*-value this large could occur due to noise. The model's *Lack of Fit F*-value of 1.98 implied the model's lack of fit was not significant relative to pure error, as there was a 35.35% chance that a *Lack of Fit F*-value this large could occur due to noise. The non-significance of the *Lack of Fit F*-value indicated the validity of the regression model. The adjusted R-square for the equation was close to unity ( $R^2 = 0.9922$ ), indicating a high correlation between the observed and predicted values. The Design-Expert statistic 'Adeq Precision' was a measure of the model's signal-to-noise ratio, and a ratio greater than 4 indicated adequate model discrimination. The ratio obtained from our model was 28.781, which indicated model adequacy. Moreover, a low coefficient of variance (2.78) indicated a high degree of precision in the experimental values. In conclusion, the model (equation 1) was suitable for extracting flavonoid from tartary buckwheat.

Three-dimensional response surface plots are presented in Figure 3. These results differed from those of the single-factor-test. The RSM results indicated that an increase in extraction temperature improved the extraction yield, which was not shown by the single-factor-test results. A possible explanation for the difference was that at a high methanol concentration, the extracted rutin was relatively stable, and as a result, with an increase in temperature, the extraction speed increased. On the basis of the RSM results, an increase in methanol concentration from 50 to 72% improved the extraction yield. However, when the methanol concentration was more than 72%, a slight decline in the response was observed. In the RSM plots in Figure 3, extraction times of over 21.4 minutes did not have an obvious effect on the extraction yield. A possible explanation for the result was that an increase in extraction time could accelerate rutin degradation during extraction, resulting in a lower yield.

The maximum extraction yield of the three flavonoids

**Table 1: The three levels of the three variables in the RSM assessment**

Independent variables	Levels		
Time ( $X_1$ )	5	15	25
Temperature ( $X_2$ )	40	50	60
Methanol concentration ( $X_3$ )	50	70	90

**Table 2: Response surface design and experimentally obtained data**

Test order	Variable levels			Flavonoid yield (%)
	$X_1$	$X_2$	$X_3$	
1	5	40	70	2.74
2	25	40	70	3.38
3	5	60	70	3.28
4	25	60	70	3.92
5	5	50	50	2.69
6	25	50	50	2.65
7	5	50	90	1.88
8	25	50	90	3.58
9	15	40	50	2.37
10	15	60	50	3.38
11	15	40	90	2.86
12	15	60	90	3.17
13	15	50	70	3.69
14	15	50	70	3.69
15	15	50	70	3.81

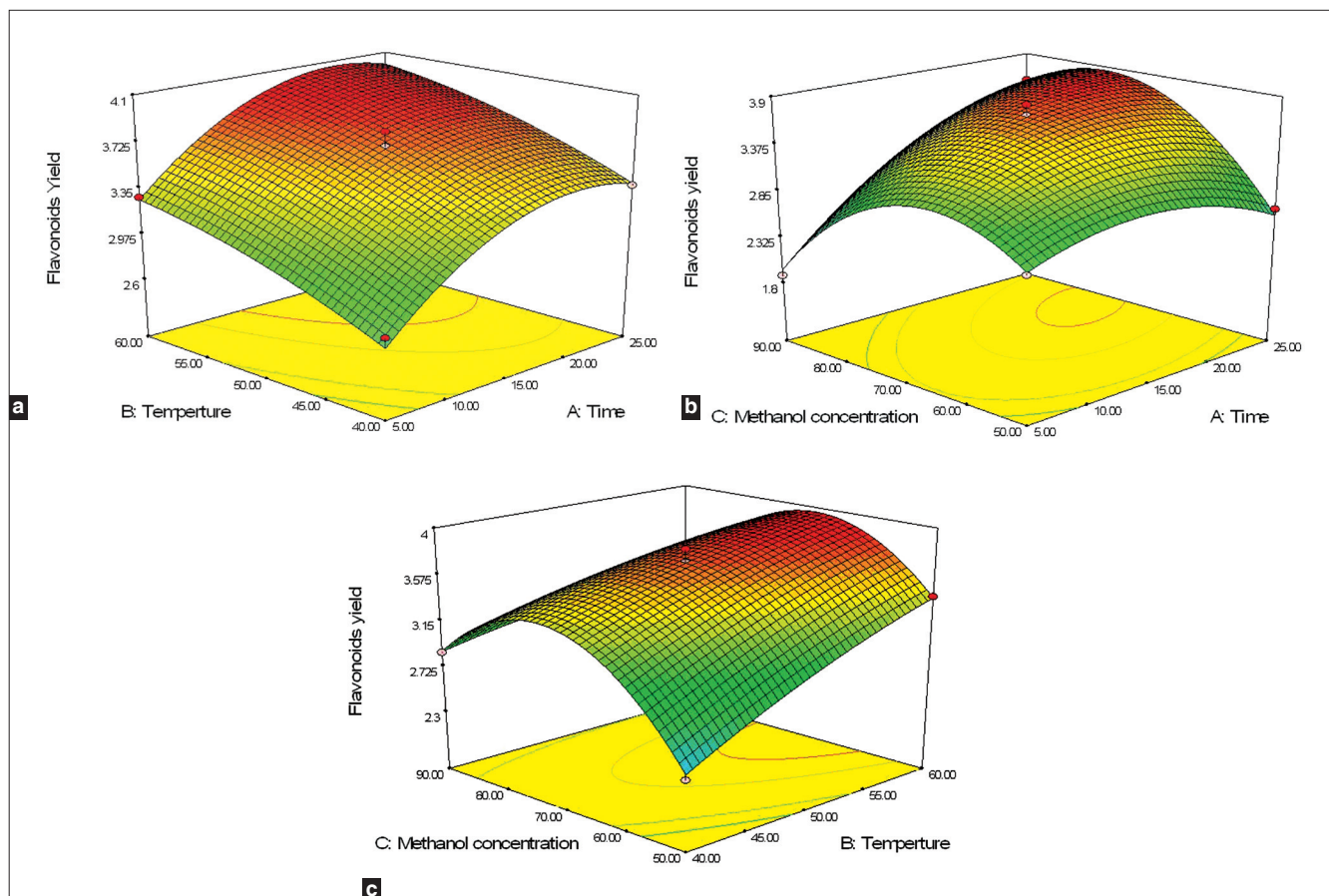
**Table 3: Analysis of variance (ANOVA) results for regression equation 1**

SD	SS	DF	MS	F-value	p value
<b>Model</b>	<b>4.82</b>	<b>9</b>	<b>0.54</b>	<b>70.30</b>	<b>&lt; 0.0001</b>
$X_1$	1.08	1	1.08	141.98	< 0.0001
$X_2$	0.72	1	0.72	94.61	0.0002
$X_3$	0.020	1	0.020	2.63	0.1659
$X_1 X_2$	0.000	1	0.000	0.000	1.0000
$X_1 X_3$	0.76	1	0.76	99.46	0.0002
$X_2 X_3$	0.12	1	0.12	16.10	0.0102
$X_1^2$	0.38	1	0.38	50.46	0.0009
$X_2^2$	0.022	1	0.022	2.91	0.1485
$X_3^2$	1.85	1	1.85	242.87	< 0.0001
<b>Lack of Fit</b>	<b>0.028</b>	<b>3</b>	<b>0.0095</b>	<b>1.98</b>	<b>0.3535</b>

SD: sources of deviation; SS: sum of squares; DF: degree of freedom; MS: mean square

combined was calculated by the Design-Expert software. The conditions that provided the highest percentage of extraction of total flavonoids were, a methanol concentration of 72%, extraction time of 21 minutes, and a temperature of 60°C. For these conditions, the corresponding theoretical maximum yield was 4.06%. To confirm the theoretical results, three parallel experiments were carried out under those optimized conditions. The average actual extraction yield obtained from the experiments was 3.94%, very close to the predicted results [Table 4]. By using UAE with these optimized conditions (extraction time, 21 minutes; temperature, 60°C; and





**Figure 3:** Response surface graphs for the effects of extraction time, extraction temperature, and methanol concentration on the flavonoid yield of tartary buckwheat: (a) Temperature and Time. (b) Time and methanol concentration. (c) Temperature and methanol concentration

**Table 4: Theoretical yield of total flavonoids based on RSM-derived optimum extraction conditions and actual yield from modified experimental conditions**

	Extraction time (min)	Temperature °C	Methanol (%)	Yield (%)
Optimum conditions	21.44	60	72.19	4.06
Modified conditions	21	60	72	3.94 ± 0.062

solvent, 72% methanol), the yield of total flavonoids (%) in three varieties of tartary buckwheat were as follows:  $3.98 \pm 0.057$  in Chuanqiao 1,  $3.87 \pm 0.065$  in Xiqiao 1, and  $4.04 \pm 0.063$  in Miqiao 1.

In conclusion, a new optimization method based on a combination of UAE and RSM was investigated for the extraction of total flavonoids from tartary buckwheat. The RSM method was based on a three-level, three-variable (extraction time, extraction temperature, and methanol concentration) BBD. The maximum extraction yield of total flavonoids was obtained by performing UAE with

72% methanol at 60°C, for 21 minutes. Under these conditions, the experimental yield of total flavonoids was 3.94%, close to the theoretical yield of 4.06%. The results indicated that the UAE-RSM approach was effective for maximizing the extraction of total flavonoids from tartary buckwheat.

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**Conflict of Interest:** None declared.

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