초음파 공정을 이용한 Prunus persica 추출물의 면역효능 증진

Enhancement of Immune Activities of *Prunus persica* Extracts by Ultrasonification Extraction

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Objectives

This study, we were to enhance whitening effects of seeds of *Prunus persica* extracts by ultra high pressure extraction process

Materials and Methods

Materials

Prunus persica was extracted by water extraction at $100\,^{\circ}$ C and $60\,^{\circ}$ C, 70% ethyl alcohol extraction at $60\,^{\circ}$ C and ultrasonification extraction at 130 KHz for 30 minutes at $60\,^{\circ}$ C.

o Methods

In order to measured immune activities, we performed MTT assay, SRB assay and T and B cell growth ratio.

Results

Prunus persica was extracted at 60°C and 130 KHz for 30 min. The extraction yield was 26.1% (v/v) which was higher than that from convenational extraction using water at 100°C for 12 hours. All of extracts at a concentration of 1.0 mg/ml showed relatively low cytotoxicity on human normal kidney cell (HEK293) in range of 15 ~ 25%. The inhibition ratios of several cancer cell lines such as human lung cancer cell A549, human gastric cancer cell AGS and human hepatocarcinoma cell Hep3B were measured using the sulforhodamine–B assay. The ultrasonification extracts of *Prunus persica* showed the highest cancer cell (A549,AGS,Hep3B) growth inhibition ratio as 77.3%, 79,1% and 75.9%, respectively. The immune B and T cell growth was improved by the ultrasonification extracts of *Prunus persica* up to $1.2 \times 10^4 \text{cells/mL}$ and $1.0 \times 10^4 \text{cells/mL}$, respectively. The extract prepared also greatly increased the secretion of both IL–6 and TNF–α from ultrasonification process. This results can conclude that ultrasonification process effectively released active biomaterials which could important role in enhancing immune activity in the body.

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Table 1. The extraction yields of *Lithospermum erythrorhizon* according to different extraction processes.

Prunus persica						
Extraction condition [‡]	WE100	WE60	EE	UE		
Yields(%, w/w) [†]	20.2	17.3	17.5	26.1		

Mean values±SD from triplicate separated experiments are shown. Mean with difference letter (A–C) within extraction yields are significantly different at p < 0.05.

^{*}WE100: water extraction at 100℃, WE60: water extraction at 60℃ EE: 70% ethyl alcohol extraction at 60℃ UE: ultrasonification extraction at 130 KHz.

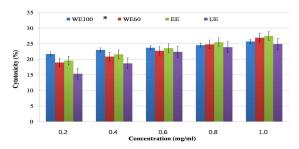


Fig. 1. The cytotoxicity of *Prunus persica* according to different extraction processes.

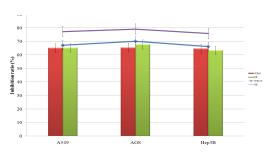


Fig. 2. The cancer cell growth ingibition ratio of *Prunus persica* according to different extraction processes.

Mean values \pm SD from triplicate separated experiments are shown. Mean with difference letter (A–D) within same concentration are significantly different at p < 0.05 and mean with difference letter (a–c) within same sample are significantly different at p < 0.05.

Table 1. The immune cell growth of *Prunus persica* according to different extration processes.

Prunus persica						
	WE 100	WE 60	EE	UE		
B cell	1.1×10 ⁴	1.0x10 ⁴	1.1×10 ⁴	1.2×10 ⁴		
T cell	0.9×10 ⁴	0.9×10 ⁴	0.8×10 ⁴	1.0×10 ⁴		

^{*}WE100: water extraction at 100℃, WE60: water extraction at 60℃ EE: 70% ethyl alcohol extraction at 60℃ UE: ultrasonification extraction at 130 KHz.

Reference

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Mean values \pm SD from triplicate separated experiments are shown. Mean with difference letter (A–D) within same concentration are significantly different at p < 0.05 and mean with difference letter (a–e) within same sample are significantly different at p < 0.05.

^{*}WE100: water extraction at 100°C; WE60: water extraction at 60°C; EE: 70% ethyl alcohol extraction at 60°C; UE: ultrasonification extraction at 130 KHz.

^{*} See the Fig. 1. for abbreviation.