

Single Intramuscular-dose Toxicity of Water soluble Carthmi-Flos herbal acupuncture (WCF) in Sprague-Dawley Rats

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Key Words

Carthamus tinctorious L., *Cathami Semen*, Carthmi-Flos herbal acupuncture, Water soluble Carthmi-Flos herbal acupuncture, pharmacopuncture, intramuscular toxicity test

Abstract

Objectives: This experiment was conducted to examine the toxicity of Water soluble Carthmi-Flos herbal acupuncture (WCF) by administering a single intramuscular dose of WCF in 6-week-old, male and female Sprague-Dawley rats and to find the lethality dose for WCF.

Methods: The experiment was conducted at Biototech according to Good Laboratory Practices under a request by the Korean Pharmacopuncture Institute. This experiment was performed based on the testing standards of "Toxicity Test Standards for Drugs" by the Ministry of Food and Drug Safety. Subjects were divided into 4 groups: 1 control group in which normal saline was administered and 3 test groups in which 0.1, 0.5 or 1.0 mL of WCF was administered; a single intramuscular dose

was injected into 5 males and 5 females in each group. General symptoms and body weights were observed/measured for 14 days after injection. At the end of the observation period, hematological and clinical chemistry tests were performed, followed by necropsy and histopathological examinations of the injected sections.

Results: No mortalities were observed in any group. Also, symptoms, body weight, hematology, clinical chemistry and necropsy were not affected. However, histopathological examination of the injected part in one female in the 1.0-mL group showed infiltration of mononuclear cells and a multi-nucleated giant cell around eosinophilic material.

Conclusion: Administration of single intramuscular doses of WCF in 3 groups of rats showed that the approximate lethal dose of WCF for all rats was in excess of 1.0 mL, as no mortalities were observed for injections up to and including 1.0 mL.

1. Introduction

Pharmacopuncture refers to be in the limelight acu-

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puncture therapy that treat diseases by using techniques such as warming (Chunghwa), alcohol extraction or pressing to extract drugs from one or more oriental medicines and then injecting them into specific acupoints related to the disease and adjusting meridian functions [1]. *Carthamus tinctorious L.* is an annual safflower belonging to the family *Compositae*. *Cathami Semen* is the seed of the safflower and is composed of 20%-30% fatty oil (acids), whose main constituents are glycerides, linoleic acid, and oleic acid, as well as serotonin, serotonin conjugate, and serotobenin. *Carthamus tinctorious L.* contains Safflower-yellow cathamin, which reportedly extends blood vessels, excites the uterus, and reduces blood cholesterol. In oriental medicine, *Carthamus tinctorious L.* has warmness, which can affect detoxification, pain relief and blood circulation [2]. Carthmi-Flos herbal acupuncture (CF), a representative meridian pharmacopuncture medicine, is an oriental medicine made by extracting and refining oil from *Cathami Semen*.

CF was prepared in a sterile laboratory at the Korean Pharmacopuncture Institute (KPI). After the seeds had been cleaned, ground and peeled, they were screw pressed on a prepared mat. Rough unwanted residue was removed, and natural precipitation was induced. The extract from a 3-step filter was packaged with nitrogen [3]. Water soluble Carthmi-Flos herbal acupuncture (WCF) was prepared by homogenizing CF, water for injection (WFI) and an emulsifier.

Pharmacopuncture with *Carthamus tinctorious L.* is a representative pharmacopuncture method whose efficacy was demonstrated by many studies. Various studies on the effects of pharmacopuncture with *Carthamus tinctorious L.* in treating painful diseases, such as anti-pain effects in an arthritis model on experimental animals [4], pain-relief effects in degenerative knee arthritis patients [5], and effects in reducing backache according to the Oswestry Disability Index [6], were conducted, as were studies on immune-function and blood-vessel effects, including its anti-carcinogenic activity [7] and its effect on suppressing blood clots induced by endotoxin [8]. Other studies addressed hematological diseases (the treatment of alopecia areata by using pharmacopuncture with *Carthamus tinctorious L.*) [9], and gynecological diseases (the treatment of oligomenorrhea with pharmacopuncture and *Carthamus tinctorious L.*) [10]. Thus, the effects of pharmacopuncture combined with *Carthamus tinctorious L.* have been demonstrated in a variety of different fields.

Lim *et al.* studied the safety of pharmacopuncture with *Carthamus tinctorious L.* against skin irritation and eye irritation [11], and An *et al.* verified its safety through acute and subacute toxicity experiments [12]. Because pharmacopuncture with *Carthamus tinctorious L.* is expected to

show gradual diversification of clinical use as an Oriental medicine, studies on toxicity, dose, and side effects must be pursued because pharmacopuncture inserts foreign substances into the human body. In this study, single intramuscular-dose toxicity tests were performed on rats to verify the safety and to identify any side effects of WCF prepared using *Carthamus tinctorious L.* Preliminary study was performed at Biototech by non-Good Laboratory Practices (GLP). The 6 rats applied WCF 0.1 mL dose. No rats died or showed abnormal finding. From this pilot study, main study's WCF dose determined to 1.0 mL. These tests were performed by Biototech according to GLP and the Standards of the Ministry of Food and Drug Safety as requested by the KPI. There was no conflict of interest.

2. Materials and Methods

The WCF was provided by the KPI. WCF was developed to increase the treatment effects by increasing absorptivity and to reduce the side effects by reducing residue of existing CF. WCF was prepared in three steps: In step 1, CF, WFI and primary emulsifier were mixed together and homogenized for 30 minutes by using a high-speed homogenizer. Emulsifier was added in step 2 and homogenized at 30-80°C. In step 3, the temperature was reduced, and the mixture was homogenized to yield WCF (Fig. 1)

Sprague-Dawley rats, purchased from Orientbio Inc., Korea, were used as experimental animals. The rats, 24 males and 24 females, were 5-weeks old with body weights of 111.8-135.8 g and 107.4-127.9 g, respectively. The animals were given a 7-day grace period to acclimatize before any

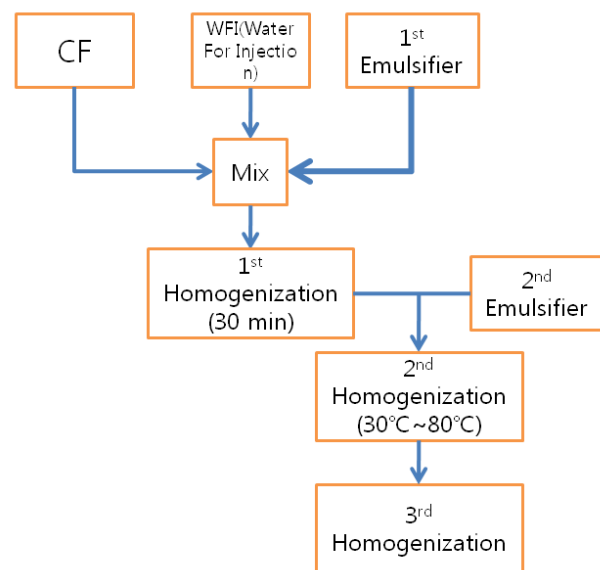


Figure 1 Flow diagram showing the manufacturing process of WCF pharmacopuncture.

measurements. General symptoms and body weights were observed to verify that no problems existed with the animals. During the grace period, individual subjects were marked, and at the end of the grace period, 20 males and 20 females closest to the standard body weight were selected and randomly divided into 4 groups of 5 male rats and 5 female rats so that the mean weights of the four groups were uniform.

Solid feed (Teklad Certified Irradiated Global 18% Protein Rodent Diet 2918C) and filtered tap water (Cheongju tap water filtered using a filter water sterilizer and irradiated with ultraviolet rays) were freely available. The breeding environment was as follows: a temperature of 20.0°C-22.8°C, relative humidity of 48.5%-65.8%, 10- to 15-hour ventilation, 12-hour/day lighting cycle, and illuminance of 150-300 Lux.

Intramuscular dose was selected because WCF is expected to be clinically applied to muscles. The clinical dose of WCF was 1.0 mL per application, as no fatalities had been observed at doses up to this value in preliminary tests (Biototech Study No.: B12873P) on male and female rats. Thus, under consultation with the test's requester, 1.0-mL/animal was configured as the high dose. Medium and low doses were 0.5- and 0.1-mL/animal; the same amount of saline solution, 1.0-mL/animal, was injected into the control group. (Table 1)

Injection was performed using a disposable syringe (1 mL, 26 G). Single doses were injected into the left femoral muscle for the low-dose and the medium-dose groups. For the control and the high-dose groups, single injections, 0.5 mL/site, were done into the left and the right femoral muscles. On the day of injection (day 0), clinical symptoms and fatalities were observed at 30 minutes and at 1, 2, 4 and 6 hours after injection. From day 1 to day 14, general symptoms were observed once a day. Body weights were measured (CP3202S scale) on days of injection (before injection) and on days 3, 7 and 14. For the hematology test, blood was collected from the abdominal aorta on day 15

after applying anesthesia by isoflurane. Hematological analyses were carried out by placing 1 mL of collected blood into a tube containing ethylenediaminetetraacetic acid, which was then analyzed using a blood corpuscle analyzer (ADVIA 120, SIEMENS, Germany). For the coagulation test, about 2 mL of collected blood was placed in a tube containing 3.2% sodium citrate and centrifuged for 10 minutes at 3,000 rpm, after which the blood plasma was collected. Measurement was done using a coagulation time analyzer (Coapresta 2000, SEKISUI, Japan). The prothrombin time and the activated partial thromboplastin time (APTT) were measured. After completing the hematology tests, for the clinical chemistry test, we centrifuged the remaining blood for 10 minutes at 3,000 rpm and collected the blood serum. A clinical chemistry analyzer (7180, Hitachi, Japan) and an electrolyte analyzer (AVL9181, Roche, Germany) were used.

Visual inspection of all body organs and tissues was performed on all animals after necropsy. Body organs and tissues were extracted and fixed in 10% neutral buffered formalin, and histopathological observations were carried out.

Body-weight, hematology and clinical chemistry results were tested using SAS (version 9.3, SAS Institute Inc., USA). Bartlett tests were performed for homo-scedasticity (significance level: 0.05). For homo-scedasticity, one-way analysis of variance tests were performed to yield the significance (significance level: 0.05), and multiple Dunnett's *t*-tests were carried out (significance level: 0.05 on both sides and 0.01). For hetero-scedasticity, significance (significance level: 0.05) was checked using the Kruskal-Wallis test.

3. Results

In this study, no mortalities were caused by injecting intramuscular doses of WCF. Accordingly, computation of

Table 1 Groupings of the rats

Group	WCF injection [mL]	Saline-solution injection [mL]	Number of animals [serial number]	
			Male	Female
G1: control group	0	1.0	5 [1101-1105]	5 [2101-2105]
G2: low-dose group	0.1	0	5 [1201-1205]	5 [2201-2205]
G3: mid-dose group	0.5	0	5 [1301-1305]	5 [2301-2305]
G4: high-dose group	1.0	0	5 [1401-1405]	5 [2401-2405]

the LD₅₀ was impossible; however, a single intramuscular dose of WCF at 1.0 mL/animal caused no mortalities. (Table 2)

No abnormal clinical symptoms (type of toxicity symptom, time of expression, time of recovery, etc.) were observed on the day of injection (day 0) at 30 minutes and at 1, 2, 4 and 6 hours after injection, as was the case for days 1 to 14 after injection. All groups showed a continued increase in body weight, but the differences among the groups were not significant. (Fig. 2. Table 3, 4, 5)

Hematological analyses on the rats' blood showed WCF to have no effect. However, APTT (sec) mean of the G3 female group showed a statistically significant change ($P < 0.01$),

but it was a miscellaneous change with no dose dependence, clinical meaning or toxicological meaning. Clinical chemistry analyses showed no significant differences for any of the measured items between the normal saline and the WCF groups. In no male or female experimental group were visual abnormalities observed. (Table 6, 7, 8)

On the histopathological tests, one female in the high-dose group showed infiltration of mononuclear cells and a multi-nucleated giant cell around eosinophilic materials caused by WCF in an area similar to a cross section of muscular fiber, and 8 eosinophilic materials were clustered at a local part. No abnormalities were observed in any other subjects. (Table 9)

Table 2 Mortality

Group	WCF injection [mL]	Saline-solution injection [mL]	Mortality [%] [% = dead/tested]	
			Male	Female
G1	0	1.0	0% [0/5]	0% [0/5]
G2	0.1	0	0% [0/5]	0% [0/5]
G3	0.5	0	0% [0/5]	0% [0/5]
G4	1.0	0	0% [0/5]	0% [0/5]

Table 3 Clinical signs on day 0

Group	WCF injection [mL]	Saline-solution injection [mL]	Sex	Hours after dosing					Clinical signs
				0.5	1	2	4	6	
G1	0	1.0	Male	5	5	5	5	5	NOA*
			Female	5	5	5	5	5	NOA
G2	0.1	0	Male	5	5	5	5	5	NOA
			Female	5	5	5	5	5	NOA
G3	0.5	0	Male	5	5	5	5	5	NOA
			Female	5	5	5	5	5	NOA
G4	1.0	0	Male	5	5	5	5	5	NOA
			Female	5	5	5	5	5	NOA

*NOA: No observable abnormality

Table 5 Body weights

Group	WCF injection [mL]	Saline-solution injection [mL]	Sex	Weight in grams at 0, 3, 7, 14 days				Gain 0-14	
				Mean, S.D, N(number).	0 (before dose)	3	7		14
G1	0	1.0	Male	Mean	195.4	221.7	260.6	316.5	121.1
				S.D.	5.0	8.5	10.5	16.4	14.1
				N	5	5	5	5	5
			Female	Mean	163.6	176.6	193.9	218.8	55.2
				S.D.	6.9	10.0	8.8	14.3	8.9
				N	5	5	5	5	5
G2	0.1	0	Male	Mean	192.4	219.6	257.1	306.1	113.7
				S.D.	5.1	6.1	9.8	13.5	9.8
				N	5	5	5	5	5
			Female	Mean	163.6	175.2	189.9	214.8	51.1
				S.D.	6.5	5.6	8.6	9.5	6.8
				N	5	5	5	5	5
G3	0.5	0	Male	Mean	194.1	220.8	256.8	308.5	114.4
				S.D.	8.4	8.8	11.3	21.0	14.7
				N	5	5	5	5	5
			Female	Mean	163.5	176.7	190.3	217.8	54.4
				S.D.	8.8	9.9	12.3	13.6	7.2
				N	5	5	5	5	5
G4	1.0	0	Male	Mean	192.4	216.3	249.7	295.7	103.3
				S.D.	7.6	8.2	10.7	18.0	15.9
				N	5	5	5	5	5
			Female	Mean	162.5	178.3	191.6	217.9	55.3
				S.D.	8.4	7.4	7.4	8.8	3.9
				N	5	5	5	5	5

Table 6 Hematology findings

Group	WCF injection WCF [mL]	Saline-solution injection [mL]		Male	Female	Abnormality value
G1	0	1.0	No abnormalities found	✓	✓	No abnormalities
G2	0.1	0	No abnormalities found	✓	✓	No abnormalities
G3	0.5	0	No abnormalities found	✓		Female - APTT: Mean 14.4 ($P < 0.01$: Dunnett's <i>t</i> -test)
G4	1.0	0	No abnormalities found	✓	✓	No abnormalities

Table 7 Clinical chemistry findings

Group	WCF injection WCF [mL]	Saline-solution injection [mL]		Male	Female
G1	0	1.0	No abnormalities found	√	√
G2	0.1	0	No abnormalities found	√	√
G3	0.5	0	No abnormalities found	√	√
G4	1.0	0	No abnormalities found	√	√

Table 8 Necropsy findings

Sex	Male				Female			
Group	G1	G2	G3	G4	G1	G2	G3	G4
Dose (mL/animal)	0	0.1	0.5	1.0	0	0.1	0.5	1.0
Number of animals	5	5	5	5	5	5	5	5
No abnormalities found	5	5	5	5	5	5	5	5

Table 9 Histopathological change findings

Organ	Sex	Male				Female			
Injection site	Group	G1	G2	G3	G4	G1	G2	G3	G4
Injection site	Dose (mL/animal)	0	0.1	0.5	1.0	0	0.1	0.5	1.0
Injection site	Number of animals	5	5	5	5	5	5	5	5
Injection site	No abnormalities found	5	5	5	5	5	5	5	4
Injection site	Remarkable findings	0	0	0	0	0	0	0	1*

* Infiltration, mononuclear cells and multinucleated giant cells around eosinophilic materials. Grade - minimal

accumulation of CF in the human body from long-term therapy [16]. The safety of CF was reported in [12] based on a toxicity test of CF, but this was only a short-term study on acute and subacute toxicity. Chronic toxicity tests on CF must be carried out in the future.

Although histological abnormalities caused by WCF were found in a subject of the high-dose group in this experiment, according to [17], no histological abnormalities were observed in Sprague-Dawley rats treated with CF 3 times a week for 1-2 weeks. However, the group with 4 weeks of therapy showed histological ambiguity regarding the boundary of muscular tissues, observation of flares, and induction of inflammations [17]. A histological study with a greater number of subjects and a longer period is deemed necessary to examine long-term or high-dose effects of CF or WCF.

The safety of CF was demonstrated, but its side effects were presented in several studies. However, studies on the safety and the side effects of WCF are lacking. Thus, the toxicity and the safety of WCF must be investigated, and

long-term histological studies based on large numbers of subjects are deemed necessary.

5. Conclusion

The results of single intramuscular-dose toxicity tests with WCF on 4 groups of Sprague-Dawley rats, 3 experimental groups (0.1 mL/animal, 0.5 mL/animal and 1.0 mL/animal) and 1 control group (1.0 mL/animal of saline solution), are as follows:

1. In all groups no mortalities occurred.
2. There was no abnormalities in clinical symptoms, body-weight, hematology tests, clinical chemistry tests, necropsy, histopathological tests.
3. In 1.0 mL/animal dose of WCF, we could not found toxicity of WCF for both males and females Rats.

This study do not show WCF's safety of human body. To examine the safety of WCF, we need to carry out testing to check whether or not the toxicity would have negative ef-

fects on human body.

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References

1. Korean Pharmacopuncture Institute. [Pharmacopunctureology]. Seoul: Elsevier Korea LLC; 2008. Chapter 3, Chapter Types of Pharmacopuncture; p. 21-3. Korean.
2. Yook TH, Song BY, Sin MS, Kim BH, Park JJ, Yun JH. [Effects on the local thermal change following herbal acupuncture on D.I.T.I.]. *The Acupuncture*. 2000;17(3):57-68. Korean.
3. Kang KS, Lee JS, Kwon GR. [Clinical studies on the cases of side effect of CF herbal acupuncture treatment]. *Pharmacopuncture*. 2001;4(2):65-71. Korean.
4. Lee HT, Lee HI. [Effect of Cathami Flos aqua-acupuncture on arthritis induced by injecting Freund's complete adjuvant]. *The Journal of East-West Medicine*. 1998;23(3):6-20. Korean.
5. Yoon MY, Cho NG, Hur TY, Cho EH, Lee OJ, Kim KS. [Clinical study on effect of Carthmi - Flos herbal acupuncture therapy on osteoarthritis in knee joint]. *The Acupuncture*. 2002;19(2):189-200. Korean.
6. Cho TS, Son IS, Kwon HY, Youn HM, Jang KJ, Song CH, et al. [The effect of CF herbal acupuncture by Oswestry disability index]. *Pharmacopuncture*. 2002;4(3):15-20. Korean.
7. Oh CS, Lee H, Yim YK, Seong NK. [Influence on the anti-cancer and immune response improvement of herbal-acupuncture with Carthami Flos infusion solution into Chung-wan (CV12)]. *The Acupuncture*. 2004;21(5):205-18. Korean.
8. Kim YT, Han SW. [The effects of aqua-acupuncture with *Hirudo nipponica* Whitman and *Carthamus tinctorius* L. on the intravascular coagulation induced by endotoxin in rats]. *The Acupuncture*. 1997;14(1):464-77. Korean.
9. Kim KW, Lee MK, Lim SC, Jung TY, Han SW, Seo JC. [A case of alopecia areata treated with bee venom and Carthami Flos herbal acupuncture]. *Pharmacopuncture*. 2004;7(2):91-6. Korean.
10. Kim KS, Yang SI. [A clinical study of two patient with oligomenorrhea treated Carthami Flos of aqua-acupuncture]. *Pharmacopuncture*. 2005;8(3):107-13. Korean.
11. Lim SBN, Kang DC. [The study of irritation and toxicity of Carthami oil aquapuncture solution applied topically to the skin and the eye mucous membrane]. *Pharmacopuncture*. 2000;3(1):53-63. Korean
12. An CS, Kwon KR, Lee SG. [The study on acute and sub-acute toxicity and sarcoma-180 anti-cancer effects of carthami tinctor-fructus herbal-acupuncture (CF)]. *Pharmacopuncture*. 2002;5(1):7-26. Korean
13. Choi SW, Park PB, Oh SJ. [A case report of carpal tunnel syndrome with raynaud's phenomenon treated by bee venom and Carthami Flos pharmacopuncture]. *Pharmacopuncture*. 2009;12(1):103-8. Korean
14. Kim SW, Jeong HW, Wei TS, Cho MR, Yun YC. [Experimental effects of aqua-acupuncture of Carthami Flos on the regional cerebral blood flow and mean arterial blood pressure in rats]. *The Acupuncture*. 2004;21(2):167-82. Korean
15. Woo DS, Lee KM, Lee BH, Lim SC, Jung TY, Seo JC. [The scavenging effect on nitric oxide in Carthami Flos herbal-acupuncture solution]. *Pharmacopuncture*. 2007;10(2):25-30. Korean
16. Lee JY, Kang HM, Lim CS, Kwon KR. [Report on the side-effects of pharmacopuncture lubricants (CF, JsD)]. *Pharmacopuncture*. 2007;10(3):137-142. Korean
17. Kim BW, Kwon KR. [The side effect of lubricants pharmacopuncture in rat tissues]. 2010;13(1):87-92. Korean.