

See discussions, stats, and author profiles for this publication at: <https://www.researchgate.net/publication/287332149>

# Standardization the Crude Extracts of all Urtica plant Species Growing in Palestine for Quality Control of Cosmeceutical and Pharmaceutical Formulations

ARTICLE *in* INTERNATIONAL JOURNAL OF PHARMACEUTICAL AND CLINICAL RESEARCH · AUGUST 2015

---

READS

11

## 1 AUTHOR:



[Nidal Amin Jaradat](#)

An-Najah National University

103 PUBLICATIONS 140 CITATIONS

SEE PROFILE

Research Article

# Standardization of *Urtica dioica* plant Species in Palestine Quality Cosmeceutical and Pharmaceutical Formulations

Nidal Amin\* Jaradat

Department of Pharmacy, Faculty of Medicine, Jordan University of Science and Technology, Amman, Palestine  
P.O. Box 7.

Available Online August, 2015

## ABSTRACT

**Background:** The safety and efficacy of pharmaceuticals are mainly dependent on the quality controls and standards which they follow. In Palestine, the medicinal plants and their extracts have been used as sources of active ingredients for many years. The use of medicinal plants as a source of active ingredients in pharmaceutical formulations seems to never stop, even when synthetic medicine is available. **Objective:** This study aimed to evaluate the quality of the active constituents of *Urtica dioica* species in Palestine and used for treatment of various ailments. **Methods:** Serial exhaustive extraction of the plant material using four species of solvents (water, ethanol, methanol, and acetone) was performed. The percentage of the active constituents in the extracts was determined. **Results:** The best percentage of the active constituents was found in the ethanol extract (24.36% of the total starting weight), while the best organic extract was the acetone extract (24.88% of the total starting weight). **Conclusion:** *Urtica dioica* species, *Urtica pilulifera* and *Urtica urens* are suitable for manufacturing of formulations and pharmaceutical preparations.

**Keywords:** Standardization; *Urtica dioica*; *Urtica pilulifera*; *Urtica urens*; medicinal plants

## INTRODUCTION

Medicinal plants are a source of many active ingredients used in pharmaceuticals. In Palestine, medicinal plants have been used for centuries. The use of medicinal plants in pharmaceuticals is increasing due to the growing interest in natural products. The present study aimed to evaluate the quality of the active constituents of *Urtica dioica* species in Palestine. The study was conducted in Amman, Palestine. The plants were collected from different locations in Amman. The active constituents were extracted using four different solvents: water, ethanol, methanol, and acetone. The percentage of the active constituents in the extracts was determined. The results showed that the ethanol extract contained the highest percentage of active constituents (24.36% of the total starting weight). The acetone extract contained the second highest percentage (24.88% of the total starting weight). The water and methanol extracts contained lower percentages of active constituents. The study concluded that *Urtica dioica* species, *Urtica pilulifera* and *Urtica urens* are suitable for manufacturing of formulations and pharmaceutical preparations.

\* Author for Correspondence



Table 2: The weights of the resulted extracts

The serial extract	Urtica kioi extract weight in mg	Urtica membranacea extract weight in mg	Urtica pilulifera extract weight in mg	Urtica urens weight in mg
The organic extra	72	68	45	62
The first aqueous	433	343	421	411
The second aqueo	176	123	161	122
Total of the aqueo	609	466	582	533

The recent evidence based study shows that the plant, distilled water be used for treatment of urinary tract infections while the whole plant clinically approved their effect for treatment of arthralgia, rheumatoid arthritis, osteoarthritis, and other conditions. The recent scientific researches have demonstrated that Urtica species have antibacterial effects on the gram positive and negative bacteria several times more than chemical materials. Antiviral (viruses such as hepatitis A and B) and antifungal activities. Also self-treatment of cardiovascular diseases (vasculopathy) improve lipid profile and platelet aggregation and allergic diseases of all types.

The percentages of the extracts

Urtica species	The percentage of the exhaustive extraction	The percentage of the serial extraction
Urtica kioviensis	2.88	24.36
Urtica membranacea	2.72	18.64
Urtica pilulifera	1.8	23.28
Urtica urens	2.48	21.32

MATERIAL AND METHODS

Collection and identification of Urtica species samples. The leaves of Urtica kioviensis, Urtica membranacea, Urtica pilulifera and Urtica urens were collected from different regions of Palestine between March and April 2014. The plant was botanically identified by Jaradat from the Pharmacognosy Department, National University. Voucher specimens were deposited in the Herbarium of the Pharmaceutical Technology Division (Laboratory of Pharmaceutical Botany) and was only for the aqueous plant materials that were used. Later the leaves of four species were dried with soft cloth to remove all moisture and insects, then dried at 50°C in a vacuum oven overnight to prevent microbial growth. The dried plant material was placed in airtight bottles and stored in a desiccator until used for extraction. Chemicals & Instruments

Shaking Incubator, Gotherm (Heidolph OB2000 Heidolph), ultrasonic bath (Branson BT85, Darcos), grinder (Moulinex, model 2000), analytical scale (Radwag AS220C/ENRAGE, MN 617 and Whatman 1). The leaves of the plant were dried in a tray for 2 weeks, at room temperature, until they became completely dry. The dried plant material was obtained and cut into small pieces, then powdered in a mechanical grinder. The powdered plant, were suspended in hexane which is a largely unreactive and easily evaporated (hydrophobic) solvent and 250 ml of 50% ethanol in triple distilled water (ensure sterility) in a bottle, with continuous shaking at 725 rpm in the shaking incubator. After that, the mixture was filtered through Whatman No.1 filter paper. The plant materials that had been extracted were dried again. The liquid filtrate was separated by separation into two phases: lower phase which has higher density (aqueous phase) and upper phase which has lower density (organic phase). The aqueous phase was collected in a volumetric flask at room temperature (obtaining the powder of aqueous extract). The organic phase was collected in a glass beaker, which was placed in the shaking incubator to obtain the organic extract. The organic extract was weighed again after filtration. The weight of the organic extract was determined by weighing the difference between the weights of the plant material before and after extraction. The aqueous phase was collected in a volumetric flask at room temperature (obtaining the powder of aqueous extract). The organic phase was collected in a glass beaker, which was placed in the shaking incubator to obtain the organic extract. The organic extract was weighed again after filtration. The weight of the organic extract was determined by weighing the difference between the weights of the plant material before and after extraction. The aqueous phase was collected in a volumetric flask at room temperature (obtaining the powder of aqueous extract). The organic phase was collected in a glass beaker, which was placed in the shaking incubator to obtain the organic extract. The organic extract was weighed again after filtration. The weight of the organic extract was determined by weighing the difference between the weights of the plant material before and after extraction.

Figure 2: The percentage of the organic exhaustive extraction

Figure 3: The percentage of the aqueous exhaustive extraction

The rotary evaporator was used to evaporate any leftover organic solvent from the organic extracts that were produced from the aqueous phases obtained from the first and second extractions are shown in Table 2. Both aqueous extracts were dried in preweighed freeze-dryer bottles and placed in a vacuum oven at 40°C for 24 hours. Then the freeze-dryer bottles were weighed again, and the dry weight of both extracts was calculated. All these procedures repeated four times for each plant species.

#### RESULTS AND DISCUSSION

The aqueous and organic extracts of *Lantida kiov* and *Urtica dioica* were subjected to qualitative extraction yields must be of 20% while the

organic extracts yield 2.5% receptors and enzymes associated  
 mean *Urtica kioviensis*, *Urtica pilulifera*, *Urtica Physion* therapy research 23(7):  
 aqueous serial exhaustive extraction of *Urtica dioica* E. M. N. Feldberg W 1949. Dist  
 kioviensis organic exhaustive extractions yield choline and histamine in nettle  
 the quality control and standardization for phytochemistry 14(8):148.  
 manufacturing of cosmetics for *Urtica dioica* W. H. W. Samtleben R, B  
 pharmaceutical preparations Search for the antiprostatic principle  
 (*Urtica dioica*) roots. *Phytotherapy Research* 2004

CONCLUSION

The leaves of *Urtica dioica*, *Urtica pilulifera* and *Urtica Physion* were collected from different regions of West Bank/ Palestine. The leaves were exhaustively extracted by using different solvents. This research scientifically proved that *Urtica dioica* is the best source for further research and development of standardised pharmaceutical products. The research also provides evidence based on pharmacological and toxicological studies to recommend research on *Urtica dioica* for its use in the future for natural food supplements and cosmetics.

REFERENCES

1. Verma S, Singh S 2008. Current and future trends in herbal medicines. *Veterinary World* 11(11):529
2. Khalil EA, Afifi FS, Ali M 2007. Ethnopharmacology of medicinal plants in Egypt. *Ethnopharmacology* 112(1):104
3. GaMuhtasib H 2006. Anticancer activity of essential oil and 100 Mediterranean sage (*Salvia triloba*). *Phytotherapy Research* 20(2):169
4. Rates SMK 2001. Plants as source of medicinal drugs. *Phytotherapy Research* 15(5):603.
5. Edeoga H, Okwu D, Mbaebie B 2005. Phytochemical constituents of some Nigerian medicinal plants. *African Journal of Biotechnology* 4(7):685
6. Otles S, Yalcin B 2012. *Urtica dioica* L. as a natural source of antioxidants. *World Journal of Botany* 4(3):435
7. Upton R 2013. Stinging nettles leaf (*Urtica dioica* L.) folk herbs, grasses, extraordinary vegetable medicine. *Journal of Herbal Science and Medicine* 38(1):9
8. Chizzola R, Michitsch H, Fritzon G, Batkova N, Sivkov A, Schlafke S, Funke M 2007. Efficacy and safety of a standardized extract of *Urtica dioica* L. in the treatment of benign prostatic hyperplasia. *Phytotherapy Research* 21(5):407
9. Sekeroglu N, Ozkutlu F, Kara SM, Ozgucen M 2008. Determination of cadmium and lead levels in commonly used medicinal plants in Turkey. *Journal of Food and Agriculture* 88(1):86
10. Wetherill H 2004. Nutritional evaluation of *Urtica dioica* L. *Journal of Herbal Medicine* 9(4):111
11. Roschek B, Fink RC, McMichele M, Harte R 2009. Nettle extract (*Urtica dioica* L.) affects oral preparation and external skin preparation

method for preventing or improving hair growth. [Google Patents.](#)

26 Papageorgiou S, Varvareas D, Aitzoglou S, Viniotis E, Manganelli R, Zaccaro L, Tomou C 2010. New alternatives to cosmetic preservatives: in vitro activity of *Urtica dioica* L. [Journal of cosmetic science](#) 61(2):107-117

27 Belaiche P, Lievoux O 1991. Clinical studies on the palliative treatment of prostatic adenoma with extract of *Urtica* root. [Phytotherapy Research](#) 5(2):167-171

28 Blumenthal M, Busse W, Goldberg A, et al. 1998. [Complete German Commission E monographs: Therapeutic guide to herbal medicine](#). Botanical Council: Austin, TX 683.

29 Bone RC 1996. Toward a theory of pathogenesis of the syndrome: what we do and do not know. [Critical care medicine](#) 24(1):172-177

30 Chrubasik JE, Roufogalis BD, Chrousos GP, et al. 2006. Evidence of effectiveness of herbal preparations in the treatment of chronic low back pain. [Phytotherapy Research](#) 20(7):678-688

31 Randall C 1994. Stinging nettles for the treatment of the hip. [The British Journal of Rheumatology](#) 44(388):533

32 Randall C, Randall H, Dobbs F, Huterer C, et al. 2000. Randomized controlled trial of treatment of fibrositis pain. [Journal of the Royal Society of Medicine](#) 93(6):305-309

33 Modarres Ghahardehi A, Ibrahim DM, et al. 2012. Screening of various extracts of *Urtica dioica*. [Biología Tropical](#) 57(6):1567-1574

34 Singh R, Dar S, Sharma P 2012. Antibacterial Activity and Toxicological Evaluation of Semi Purified