Ultrasound-assisted Extraction of Phenolic, Flavonoid and Antioxidant Compounds from *Dodonaea viscose* and Its Green Synthesis of Silver Nanoparticles by Aqueous Extract

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**ABSTRACT**

The *Dodonaea viscose* was commonly known plant in Al-mukhwah area (Al-baha city in Saudi Arabia) by exhibiting various therapeutic pharmacological properties owing to existence of divers polyphenols and flavonoids in its roots and seeds. Ultrasound-assisted extraction was utilized to extract total phenolic and antioxidant compounds from *Dodonaea viscose*. The aim of this study is to inspect the solvent type effects, sonication time and dissipated power on the extraction of total phenolic compounds, total flavonoid compounds (TFC) and on the antioxidant capacity: 2,2′-azinobis-3-ethylbenzothiazoline-6 sulfonic acid (ABTS) and 2,2′-diphenyl-1-picrylhydrazyl (DPPH). The highest TPC was 176.63 mg. g⁻¹ GAE of dry extract), whilst the total flavonoids equal 125.78 mg QE.g⁻¹ of dry extract, the lower value IC₅₀ of DPPH radical scavenging activity was 13.60 µg.g⁻¹ of dry extract and under ABTS radical scavenging assay, IC₅₀ was assessed as 3.58 µg.g⁻¹ of dry extract. These results were observed for the sample extracted with pulsed ultrasound (40%) in Methanol at 15 min which indicated that the extracted compounds contribute to the activity of antioxidant. The natural products which are found in plants extracts, may be considered as a reducing agent as like a stabilizing agent in the formation of metal nanoparticles. Silver nanoparticles (AgNps) were synthesized by aqueous extract through a facile route. Characterizations of nanoparticles were evaluated by the use of UltraViolet-Visible spectroscopy (UV-Vis), Fourier Transform Infra-Red spectroscopy (FT-IR), and Transmission Electron Microscopy. Nanoparticles were found 30.20±19 (nm) by TEM analysis.

**Keywords:** *Dodonaea viscose*, Phenolic and flavonoid content, Antioxidant activity, DPPH and ABTS, Ultrasound mode, Solvent and Sonication time effect, Silver nanoparticles.

**INTRODUCTION**

Sonochemistry may be utilized in different food processing¹⁻³, conservation⁴, removal of pesticides⁵, extraction of different compounds⁶, microencapsulated food⁷, innovative foods⁸, and preparation of ferrite nanoparticles⁹. The superior performance of ultrasound than conventional processes may be associated to the acoustic intensity. Its calculation methods are built on the heat¹¹⁻¹³. The pulse period is the time during which the apparatus is on with a variable off period. A variable on time is offered by some instruments. Common pulse ratio is 1:1.
Pulsed ultrasound is preferred over continuous ultrasound in diverse cases in which heat is contraindicated or in conditions where it is admitted that the beneficial effects are produced by the mechanical activity of ultrasound^{14-16}. 

*Dodonaea viscose* is a plant pertinent to the Sapindaceae family. This family is exploited in conventional remedy for the healing of plural outside fungal infections^{17}. The powder of *D. viscose* leaves utilized to decrease fever characteristics and advantageous for various epidermis disorder.

*Dodonaea viscose* is a frequent medicinal plant utilized in traditional treatment as remedy for fever and hurt in the joints. The plant extract has moderate effects versus most of the microorganism^{17}.

The phenolic compounds can block the oxidation reactions by removed free radical transition states. Cell obtains advanced systems of antioxidants, nevertheless the adequate quantity of phenolic compounds or suppression of the antioxidant enzymes can conduct to oxidative compression^{18}.

Phenolic compounds are the essential antioxidant phytochemicals present in *D. viscose*^{17}. *Dodonaea viscose* phenolics displayed certain biological activities like antioxidant activity^{19,20}, antiallergic and anti-inflammatory activities^{21}, antibacterial activity^{22}, hepatoprotective activity^{23}, nephroprotective activity^{24}, antiproliferative^{25}, chemopreventive and anticancer activities^{26,27}.

The objective of this research was to analyze the effects of solvent type, sonication time and dissipated power on the extraction of TPC, TFC and on the antioxidant capacity: 2,2'-azinobis-3-ethylbenzothiazoline-6 sulfonic acid (ABTS) and 2,2'-diphenyl-1-picrylhydrazyl (DPPH).

Another vital part of this work was the green synthesis of silver nanoparticles. Flavonoids have been reported as the molecules which may contribute in biochemical pathways leading to the biosynthesis of noble metal nanoparticles^{28-30}. Silver has been greatly used for its high antioxidant, antimicrobial, and catalytic properties.

This study investigates a simple and rapid route of Ag NPs preparation by aqueous extract of *Dodonaea viscose*.

### EXPERIMENTAL

#### Plant material

The leaves of *D. viscose* were gathered from the region of Al-mukhwah area in Mai 2018. Dried leaves of this plant sample were ground. In order to extract with a specific solvent (50 mL), 5 g of the powder were placed in a thermostated vessel, which was stirred with a magnetic stir bar (150 rpm). Ultrasound was implemented at the top of the vessel.

Ultrasound power level (three), evaluated in distilled water by the calorimetric method at 25°C, were utilized corresponding to amplitude of 20, 30 and 40%^{10,31} all under pulsed ultrasound (10s ON:10s OFF). After the process of sonication, the suspension gets down and cooled to room temperature and then filtered through Whatman filter paper to reject solid debris. At room temperature, the extract concentrated and dried to get the crude extracts.

The filtered extracts were utilized to determine the TPC, TFC, DPPH and ABTS radical-scavenging activity.

**Fig. 1. Experimental setup: (1) ultrasonic generator, (2) ultrasonic probe, (3) Extraction mixture, (4) magnetic stirrer, (5) probe support**

#### Experimental design

In this current research paper, the objective of singular factor experiments were mainly used to find the optimum conditions for extracting antioxidant phenolic and flavonoids compounds from *Dodonaea viscose*. The four parameters such as extraction time (5,10,15 min.), ultrasound amplitude (20,30 and 40%), ultrasound mode (Continuous and pulsed ultrasound) and extraction solvent (Ethyl acetate, methanol, chloroform) were explored on which
one parameter was varied at a time while the other parameters were fixed.

**Extraction time**

*Dodonaea viscosa* samples were extracted using ethyl acetate and with pulsed ultrasound at amplitude of 20%. The pulse ratio was 1:1 mode with 10s On:10s Off. The time of extraction was varied from 5 to 15 min. (5, 10 and 15 minutes).

**Amplitude of ultrasound**

Using one of three solvents (Ethyl acetate) and for 15 min. of extraction duration with pulsed ultrasound (10:10s) samples were extracted with different amplitudes used in this research (20, 30 and 40%).

**Continuous and pulse ultrasound**

*Dodonaea viscosa* samples were extracted by ethyl acetate with two ultrasound modes: Continuous and pulsed (mode 1:1). The extraction time (15 min.) and the ultrasound power (amplitude of 40%) were used.

**Solvent extraction**

By setting the extraction time at 15 min. with the ultrasound mode (pulsed ultrasound, 10:10 sec.) at an amplitude of 40%, *Dodonaea viscosa* samples were extracted by the use of a variety types of solvents: ethyl acetate, methanol and chloroform.

**Total phenolics**

The TPC in *D. viscose* extracts was determined according to the Folin- Ciocalteu method. Specimen’s solutions of the extracts in the concentration of 1 mg.mL\(^{-1}\) were utilized in the procedure. The reaction blend was obtained by mixing 0.5 mL of specimens solutions of extracts, 2.5 mL of 10% Folin-Ciocalteu’s reagent resolved in water and 2.5 mL 7.5% NaHCO\(_3\). Blank was simultaneously prepared, comprise 0.5 mL methanol, 2.5 mL 10% Folin-Ciocalteu’s reagent resolved in water and 2.5 mL of 7.5% of NaHCO\(_3\). The specimens were eventually brood in a thermostat at 30°C for 90 minutes. Spectrophotometer at \(\lambda_{\text{max}} = 765 \text{ nm}\) was used to measure the absorbance. Specimens were planned three times for each analysis and the average value of absorbance was acquired. The phenol concentrations in the test samples were evaluated from the calibration curve. TPC was referred to as mg Gallic Acid Equivalents. TPC was expressed as mg Gallic acid equivalents per gram sample extract.

**Total flavonoids**

The total Flavonoid content was measured using aluminum chloride. Briefly, 0.5 mL aliquot of sample was blended with 1.5 mL of methanol, 0.1 mL of 1% AlCl\(_3\) and 0.1 mL of CH\(_3\)COOK solution (1 M). Sample was added to 5 mL volumetric flask with distilled water (2.8 mL). The obtained solution was jolted and absorbance at 430 nm was registered after 30 min. of incubation. A standard calibration curve was recorded at 430 nm using different concentrations of C\(_{15}\)H\(_{10}\)O\(_7\) (Quercetin). Total flavonoid content was calculated from the calibration curve as mg QE.g\(^{-1}\) of dry extract.

**DPPH radical scavenging assay**

The DPPH is hydrogen atom transfer processes utilized for antioxidant screening in plant material. The free radical 1,1-diphenyl-2-picyrylhydrazyl (DPPH), which have a dim magenta color, is reduced to the corresponding hydrazine (change of color pale red or yellowish red).

This method was performed according to procedure by (Fenshyur et al., 2005). The solution was designed by resolving 1 mg of the sample in 1 mL of ethanol (98%). Seven solutions with 100, 50, 25, 12.5, 6.25, 3.125, 1.5625 µg/mL concentrations were reduced in ethanol. 0.9 mL Tris-HCl and 1 mL of 0.1 mM DPPH in methanol solution were inserted and incubated at room temperature in the dark (60 minutes). The absorbance was recorded at 517 nm. The antioxidant activity was calculated using the following equation:

\[
\text{Scavenging activity ()}\% = \frac{Abs_{\text{control}} - Abs_{\text{sample}}}{Abs_{\text{control}}} \times 100
\]

The “Inhibitory concentration” or IC\(_{50}\) value (the concentration of antioxidant that makes 50% reduction of the DPPH activity) was also used to assess the antioxidant activity of the plant extract matched to the caliber medicine. The greater the antioxidant activity, the smaller is the amount of IC\(_{50}\).

**ABTS radical scavenging assay**

The antioxidant potential of *D. viscosa* expressed as mg QE.g\(^{-1}\) of dry extract was estimated by the ABTS (2, 2’-azinobis-3-ethylbenzothiozoline-
6-sulphonic acid) method. The solution was prepared by adding equal quantities of 7.4 mM ABTS solution and 2.6 mM K$_2$S$_2$O$_7$ solution. The obtained solution allowed in the dark for 12 h at room temperature and was diluted by blinding 1 mL ABTS solution with 60 mL methanol. The obtained absorbance was 0.8 at 734 nm using the spectrophotometer. Different concentrations of $D$. *viscose* extracts (200, 300 and 400 µg/mL) were permitted to react in a dark condition with 2.94 mL of the ABTS solution for 6 minute. Scavenging ability relative to the reaction was calculated by using the equation Eq.1.

**Green synthesis of Ag NPs**

A 10 g of dried leaves were powdered and soaked in deionized water (100 mL) and boiled for 15 minutes. A 30 mL solution of (0.01 M) AgNO$_3$ was added to 6 mL of the aqueous extract of $D$. *viscose* leaves. Then the mixture was stirred at room temperature. The solution color passed light yellow after 20 min. and gradually a dark brown color appeared which confirmed the Ag NPs formation. The NP deposits were collected after centrifuge the mixture. The de-ionized water was used to wash the AgNps. Lastly, synthesized silver nanoparticles were dried at room temperature (24 h) and kept for further evaluations.

**RESULTS AND DISCUSSION**

**Dissipated power**

The calorimetric method is used to determine the dissipated power. In this research, a temperature probe has been used. The dissipated power, $P_{diss}$, was estimated by using the following equation:

$$P_{diss} = m \times C_p \times \left(\frac{\Delta T}{\Delta t}\right)_{t=0}$$  \hspace{1cm} (2)

Where $m$ the solution mass and $C_p$ the specific heat of water. Measurement of dissipated acoustic power in the reactor was recorded for three amplitudes 20, 30 and 40%$^3\text{a}$. The dissipated power $P_{diss}$ is 25.85 Watt under pulsed ultrasound (The pulse mode is 1:1 with 10/10s). The dissipated power at the same amplitude with continuous mode is 46.27 Watt.

**Extraction time**

The time of extraction is crucial in economizing energy. The extraction efficiency is immensely influenced by the interaction time between solute and solvent. In this research we used 5, 10 and 15 min. as extraction time.

The variation effect of time (5 to 15 min.) on the extraction of TPC, TFC and antioxidant activity of $D$. *viscose* was explored. Findings showed that TPC vary significantly. From 5 to 15 min. the TPC increased to achieve 93.64 mg GAE.g$^{-1}$ Extract (Figure 2).

The obtained results are in accordance with those of Mokrani et al.$^{35}$ and Falleh et al.$^{36}$, which noted that the phenolic content of compounds extracted by Ultrasound increases as a function of time.

As can be noted from Fig. 2, the TFC increased gradually when the extraction time increased from 5 to 15 minutes. This duration allowed extraction of 106.35 mg EQ.g$^{-1}$ Extract. It was observed that the effect of time on TFC is similar to that on TPC.

Figure 3 shows that that the yield of DPPH and ABTS decreased when the time of
The IC_{50} value is the quantity of antioxidant needed to decrease the initial DPPH and ABTS concentration by 50%. The goodness of the antioxidants in the plant material was determined by these values (low values of IC_{50}). Best results of antioxidants were revealed at 15 min. as extraction time; the IC_{50} (DPPH) equal 19.52 µg.g^{-1} extract and IC_{50} (ABTS) equal 5.64 µg.g^{-1} extract.

**Amplitude of ultrasound**

Phenolic and flavonoid compounds were extracted using different ultrasound powers. The results showed that used powers were able to extract phenolics and flavonoids. The TPC was determined as 93.64, 97.07 and 106.34 mg GAE.g^{-1} Extract for 20, 30 and 40% as ultrasound amplitude (Fig. 4). Fig. 4 shows that that the yield of TPC increased gradually when the ultrasound power increase. The TFC was determined as 106.35, 111.21 and 115.01 mg EQ.g^{-1} Extract (Fig. 4). These results confirmed a significant influence of ultrasound amplitude on the content of extracted phenolic and flavonoids compounds. This is consistent with previous findings from Z. Pan et al., (2012).^{37}.

The effect of amplitude (20 to 40%) on the DPPH and ABTS was investigated. Results showed that the extract of D. viscosa at 40% of amplitude exhibited the lowest DPPH and ABTS, as depicted by high IC_{50} values (Fig. 5). The results showed that high intensity level (high amplitude) with pulsed ultrasound significantly benefited the antioxidant.

A positive correlation between the DPPH, ABTS yields and ultrasound Amplitude was observed. This is consistent with previous findings from Helena et al.,^{38}.

**Continuous and pulse ultrasound**

The results reveal that all ultrasound modes were able to extract phenolics and flavonoids. The highest yield of TPC and TFC was registered for pulsed ultrasound mode (106.35 mg GAE.g^{-1} Extract and 115.01 mg EQ.g^{-1} Extract), which represents the important result in this study (using pulsed ultrasound). It was observed that the effect of ultrasound mode on TPC is similar to that on TFC.

Best results of antioxidants were obtained with pulsed mode; the IC_{50} (DPPH) equal 18.41 µg.g^{-1} extract and IC_{50} (ABTS) equal 4.61 µg.g^{-1} extract. These results confirm the significance of using the pulsed ultrasound mode compared to continuous mode. A similar trend was observed in the study of DPPH and ABTS.

The use of pulsed ultrasound to obtain
phenolic and flavonoids may be considered as an applicable technology in sustainable chemistry, which decreases the use of substances, solvents, and makes less the consumption of energy.

Flavonoids are "high level" antioxidants, including flavones, flavanols, isoflavonoles, anthocyanins and condensed tannins. Results of this research are also supporting the presence of flavonoids from the *D. viscose* plant. Fig. 8 revealed that TPC and TFC amount increased with increasing the polarity of solvents.

The results of DPPH and ABTS with the variation of solvents are shown in Fig. 9. The best values were reached with the use of Methanol respectively. In this study, there is a correlation between total phenolic and flavonoid content and antioxidant activity. As can be noted, DPPH results were highly correlated with ABTS values in this research paper.

These results confirm that, using the best solvent (Methanol), the best extraction time (15 min.), the best amplitude 40% and with the best ultrasound mode (pulsed ultrasound) have significantly affected the extraction of TPC and TFC from *Dodonaea viscose*. Our results of TPC and TFC are commonly greater than those reported in the literature. Good results of DPPH and ABTS were obtained with using these parameters. These results indicated that the extraction compound take part in antioxidative activity.

**Solvent extraction**

Applying the pulsed ultrasound mode at 40% as amplitude for 15 min. the adoption of different solvent extraction have affected the TPC, TFC, DPPH and the ABTS of *Dodonaea viscose* (Figure 8).

The use of methanol was picked as the best solvent for extraction (The chloroform can be used) between the three extracting solvents (ethyl acetate, chloroform and methanol). The methanol has a relative polarity 0.762 compared to 0.259 for chloroform and 0.228 for ethyl acetate. This difference in polarity may be influenced the phenolic content. A lot of researches have indicated that the solubility of phenolics in the solvent is dependent on solvent polarity. Phenolic compounds are distributed in the cell according to their polarity. The extraction solvent must be elected according to the solubility.
A strong peak at 3425 cm\(^{-1}\) indicates the presence of O-H groups in phenolic compounds, and N-H groups in proteins. The peak observed at 2970 cm\(^{-1}\) is due to aliphatic C-H stretching. The peaks at (1400 and 1650) cm\(^{-1}\) indicate C=C stretching in alkenes and aromatic compounds and also C-H bending in aliphatic compounds. The peaks at (1000-1300) cm\(^{-1}\) indicate C-C and C-O stretching in the compounds like alcohols and phenols. FT-IR confirmed the presence of bio-reducing organic compounds on the surface of AgNPs which are likely to be responsible for nanoparticle synthesis and stability\(^{42}\).

**AgNPs characterization**

Green synthesis of AgNPs through *D. viscose* aqueous extract was carried out. The change of pale yellow color to dark brown color of the reaction mixture marked the synthesis of silver nanoparticles. Silver nanoparticles show a color change (pale yellow to dark brown) as a result of excitation of surface Plasmon vibrations in the particles. Characterization of synthesized nanoparticles was revealed by different methods.

AUV–Vis spectrometer was utilized to mark gradual modifications in Local Surface Plasmon Resonance of the produced silver nanoparticles. The absorption at 431 nm indicates the presence of AgNPs (Figure 10).

The FT-IR spectrum of *D. viscose* extract after synthesis of silver nanoparticles was realized to identify the water soluble organic compounds from the leaves extract which may take part in the reduction procedure of silver ions (Figure 10).

The size, size range, and distribution of the synthesized AgNPs were studied by transmission electron microscopy (TEM; JEOL-2100 made in Japan). A typical transmission electron microscope (TEM) image of the nanoparticles and their size distribution are presented in Fig. 12. The mean diameter and standard deviation of silver nanoparticles was found 30.20±19.43 (nm).

**CONCLUSION**

In the current study, ultrasound-assisted
extraction was applied to extract total phenolic and antioxidant compounds from *Dodonaea viscosa*. As far as we know, no data was reported about the extraction effect parameters on the extraction of phenolic compounds from *D. viscose* by the use of ultrasound. Results showed that TPC, TFC, DPPH and ABTS were highly affected by all the studied parameters. The optimal ultrasound extraction conditions were the solvent (Methanol), the extraction time (15 min.), the amplitude 40% and with the pulsed ultrasound mode.

In this research, we report a simple aqueous method for the environmentally friendly biosynthesis of Ag NPs. The results of this synthesis are promising.

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**Conflicts of interest**

The author declares that there are no conflicts of interest related to this article.

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