

**Original article:**

**ONTOGENETIC VARIATION OF VOLATILES AND ANTIOXIDANT ACTIVITY IN LEAVES OF *ASTRAGALUS COMPACTUS* LAM.  
(FABACEAE)**

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**ABSTRACT**

The genus *Astragalus* is a rich source of a variety of biologically active compounds including phenols, saponins, polysaccharides and essential oils. The present study was conducted to determine ontogenetic variation of the volatile organic compounds as well as total phenolic contents and antioxidant activity in leaves of *A. compactus*. The leaves of plant were harvested at vegetative, flowering and fructification stages and were analyzed by gas chromatography coupled with mass spectrometry (GC-MS). Total phenolic content (TPC) was determined using the Folin-Ciocalteau reagent and the antioxidant capacity was evaluated with the 1,1-diphenyl-2-picrylhydrazyl (DPPH) test. Different classes of volatile compounds were identified including alcohols, esters, hydrocarbons, sterols and terpenoides. Significant variation of these compounds was found during phenological stages of development. Sterols and hydrocarbons were the main components of essential oils at the vegetative stage. The presence of terpenoides (phytol) and alcohols (docosanol) was significant at the flowering stage. Fructification phase was characterized by the high content of sterols and hydrocarbons and absence of phytol. The antioxidant activity and phenolic content were related to the physiological stage and the highest amount detected at fructification phase. The ontogenetic variations of phenolic contents and antioxidant properties are largely contributed by climatic factors such as temperature and solar radiation.

**Keywords:** antioxidant, *Astragalus compactus*, GC-MS, phenological stages, total phenolic content, volatile compounds

**INTRODUCTION**

The genus *Astragalus* (Fabaceae) is one of the largest genera of vascular plants, comprising about 2500 species of herbs or shrubs (Lock and Simpson, 1991). *Astragalus* is mainly distributed in cool, temperate, arid and semiarid continental region of South-Western Asia, Sino Hymalayan re-

gion, Western-North and South America, Europe, North Africa and Australia (Polhill, 1981; Evans, 2002). Iran, possessing nearly 800 species, is one of the most important centres of diversity for this genus (Maassoumi, 1998).

*Astragalus* root has long been one of the most important tonic herbs used in traditional Chinese medicine and has been used

to improve resistance to infections and to aid in immunological disorders and viral infections, and as hepatoprotective (Rios and Waterman, 1997; Verotta and El-Sebakhy, 2001). It has also been proven to be effective for clinical treatment of nephritis, cardiovascular diseases, hypertension, diabetes, and cancers (Miller, 1998; Sinclair, 1998). The interesting pharmacological properties of *Astragalus* were often associated with its chemical composition. The known beneficial components of *Astragalus* are mainly flavonoid, triterpene saponins, polysaccharides and essential oils (Pistelli, 2002). Despite the fact that the phenolic compounds are among the most important antioxidants in plant (Choudhary and Swarnkar, 2011), the information on the antioxidant activity of *Astragalus* is limited. On the other hand, although a great deal of research has been carried out on phenols, saponins and polysaccharides of different *Astragalus* species (Smee et al., 1996; Nikolov and Benbassat, 1997; Krasteva and Nikolov, 2000; Gardner et al., 2001; Johnson et al., 2001), few studies have been done on the volatile compounds of the genus (Platikanov et al., 2005; Teyeb et al., 2011).

The volatile compounds consist of a complex mixture of chemicals with special chemical and physical properties, and are among the most valuable compounds produced by plants. It is shown that the volatile components of plant can be affected in a number of ways, especially by developmental and environmental factors (Circella et al., 1995; Kokkini et al., 1997; Máñez et al., 1991; Taveira et al., 2003). Knowledge of the factors that determine the chemical variability and yield for each species is important in particular for medicinal plants, to optimize the time of collection and to obtain higher yields of high-quality essential oils. As well as volatile compounds, the antioxidant capacity and phenolic content of plant species could be greatly affected by the environmental factors and growing season (Howard et al., 2002; Toor et al., 2006). Such a variation is attributed to the difference in growing temperatures and light in-

tensity. Therefore, it is hypothesized that changes in solar radiation received and temperatures in different seasons may affect the phenolic content and antioxidant components of plants.

Regarding the importance of the variability in composition of volatiles and lack of information about the effect of seasonal variation on the phenolic content and antioxidant activity of *Astragalus*, we studied ontogenetic variation of volatile and antioxidant activity in leaves of *A. compactus* Lam. *A. compactus* Lam. is one of the Asiatic species, which is famed for the production of gum tragacanth as an economically important natural product (Movafeghi et al., 2010). The aims of the presented study were: (1) to analyze the volatile composition of *A. compactus* leaves; (2) to determine the link between volatile content of plant and developmental stages; and (3) to evaluate the effect of ontogenetic variation on the antioxidant activity and phenolic contents.

## MATERIAL AND METHODS

### *Plant material*

Leaves of *A. compactus* Lam. were harvested at three phenological phases (vegetative, flowering and fructification stages) from the Payam Mountains at the east Azerbaijan province of Iran, at altitudes above 2200 m and transferred into the laboratory. Voucher specimens were deposited in the herbarium of the Tabriz University of Medical Science. The samples were dried for 10 days at room temperature and then were powdered.

### *Extraction*

A quantity of 100 grams of powdered roots was extracted with n-hexane using a Soxhlet apparatus for 8 h. The extract was concentrated to ~1 ml under reduced pressure on a rotary evaporator. The extract was stored in sealed vials at 4 °C until GC-MS analysis.

### **Analytical procedure**

Recognition of compounds was carried out by a Shimadzu GC-MS-QP 5050A gas chromatograph (Shimadzu Corporation, Kyoto, Japan). The column used for the analysis was a 60 m × 0.25 mm i.d. DB1 capillary column coated with a film of dimethylpolysiloxane (J&W Scientific, Folsom, CA, USA). An aliquot of 1 µl of the n-hexane extract was injected into the GC-MS system in the split mode (split ratio 1:42). Helium was used as the carrier gas with a flow rate of 1 ml/min. The column temperature was maintained at 100 °C for 2 min. Then, it was programmed to 300 °C at a rate of 5 °C/min and the final temperature was held for 38 min. Injector temperature and detector temperature were optimised at 270 °C and 310 °C, respectively. The MS operating parameters were as follows: ionization energy, 70 eV; ion source temperature, 280 °C; quadrupole, 100 °C; solvent delay, 3.0 min; scan speed 2000 u/s and scan range 30-600 u, EV voltage 3000 volt.

### **Identification of components**

The components were identified on the basis of comparison of their retention indices and mass spectra with those for standard compounds and by matching with the Wiley 229, Nist 107 and Nist 21 libraries.

### **DPPH radical-scavenging activity**

The DPPH quenching ability of plant methanolic extracts was measured according to Hanato et al. (1988). A quantity of 5 ml of the extract at different concentrations was added to 5 ml of a DPPH methanolic solution. The mixture was shaken vigorously and left standing at room temperature for 30 min in the dark. The absorbance of the resulting solution was then measured at 517 nm. The same procedure was applied for the quercetin as a positive control. The antiradical activity was expressed as IC<sub>50</sub> (µg/ml), the antiradical concentration required to cause a 50 % inhibition. A lower IC<sub>50</sub> value corresponds to a higher antioxidant activity of plant extracts.

The ability to scavenge the DPPH radical was calculated using the following equation:

$$\text{DPPH scavenging effect (\%)} = \frac{(A_0 - A_1)/A_0}{100}$$

where A<sub>0</sub> is the absorbance of the control at 30 min, and A<sub>1</sub> is the absorbance of the sample at 30 min.

### **Determination of TPC**

TPC was estimated using the Folin-Ciocalteau colorimetric method described previously (Meda et al., 2005) with a little modification. Briefly, 5 g of powdered samples were individually dissolved in 50 ml of aceton-water (in 4-6 ratio). After 30 min, 1 ml of these solutions were mixed with 0.2 ml of Folin-Ciocalteau reagent, and 1 ml of 2 % sodium carbonate (Na<sub>2</sub>CO<sub>3</sub>). After incubation at 40 °C for 30 min, the absorbance of the reaction mixtures were measured at 760 nm by using a spectrophotometer (Shimadzu, Kyoto, Japan). Quantification was done on the basis of the standard curve of gallic acid. Results were expressed as µg of gallic acid equivalent (GAE) per mg of dry weight (DW).

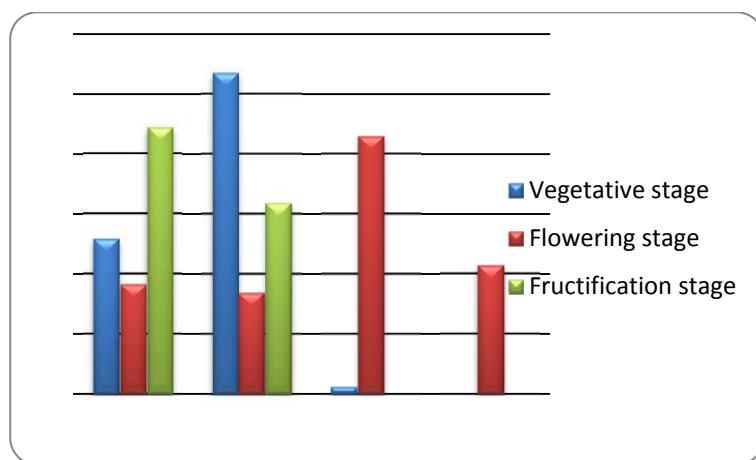
### **Statistical analysis**

The experiment was a completely randomized design with three replications. Data were subjected to analysis of variance (ANOVA) and means were separated by Duncan multiple range test at P <0.05 significant level.

## **RESULTS AND DISCUSSION**

### **GC-MS analysis**

The obtained results showed the presence of different groups of compounds including alcohols, fatty acids, hydrocarbons, sterols, terpenoids and aldehydes in *A. compactus* leaves. The identified compounds and relative quantitative distribution are shown in Table 1. Significant differences in the composition of the volatiles were found during different phenological stages (Figure 1).



**Figure 1:** Comparison of the average percentages of hydrocarbons, sterols, terpenoids and alcohols at different developmental stages

**Table 1:** Chemical composition (%) of volatiles from the leaves of *Astragalus compactus* in three developmental phases

No	Compound	K.I.	Vegetative	Flowering	Fructification
1	Octadecanal	1357	0.55	Nd	Nd
2	Pentadecanoic acid 14-methyl-methylester	1598	Nd	Nd	0.52
3	1-Heptadecene	1700	Nd	Nd	0.72
4	1-octadecene	1900	0.09	Nd	Nd
5	Phytol	1943	Nd	43.05	0.57
6	Eicosane	2000	Nd	9.72	0.54
7	Methyl linoleate	2150	Nd	Nd	1.06
8	9,12,15-octadecatrienoic acid-methylester	NA	Nd	Nd	1.68
9	Linoleic acid	2173	0.19	Nd	1.73
10	Docosane	2200	0.07	Nd	Nd
11	Tricosane	2300	0.68	8.67	1.88
12	9-Tricosene	NA	0.39	Nd	Nd
13	Tetracosane	2400	0.31	Nd	Nd
14	Docosanol	2494	Nd	21.52	Nd
15	Pentacosane	2500	Nd	Nd	1.38
16	Squalene	2790	1.32	Nd	2.25
17	Octacosane	2800	0.46	Nd	Nd
18	Triaccontane	3000	5.37	Nd	Nd
19	Triaccontan 1-bromo	NA	0.71	Nd	Nd
20	Tetracontane	NA	6.89	Nd	Nd
21	Cyclotetracosane	NA	-	Nd	3.85
22	Cyclooctacosane	NA	11.33	Nd	33.86
23	Campesterol	3355	7.41	Nd	Nd
24	Gamma-sitosterol	NA	33.28	Nd	31.92
25	Stigmasta-5,22-dien-3-ol	NA	12.95	Nd	Nd
26	Ergost-7,22-dien-9,11-epoxy-3-ol,acetate	NA	Nd	17.01	Nd
Hydrocarbons					
Steroles					
Esters					
Terpenoids					
Alcohols					
Other compounds					
Total					

KI: Kovats indices, NA: Not Available, Nd: Non detected.

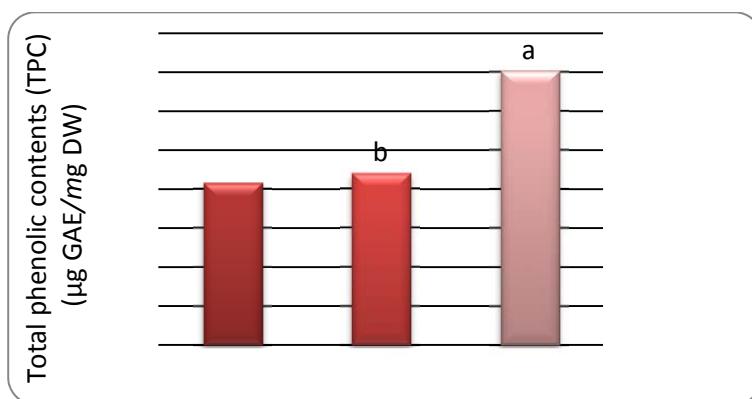
The major constituents detected in vegetative stage were sterols amounting to 53.64 %. Gamma sitosterol, campesterol and stigmasta-5,22-dien-3-ol were the predominant identified sterols. This is in agreement with the descriptions of Engvild (1995) and Ebrahimzadeh et al. (2001) that reported campesterol, gamma-sitosterol, and stigmasterol as the predominant sterols in the aerial parts and roots of the genus. Hydrocarbons (25.59 %) were the other main functional group identified in the vegetative stage. An almost complete series of n-alkanes have previously been identified in leaf materials of the different *Astragalus* species (Platikanov et al., 2005; Teyeb et al., 2011) and it is suggested that they probably function as a component part of the waxes. However, the presence of tetratetracontane and cyclooctacosane in the volatile composition of the genus has not yet been reported. Low amounts of linoleic acid and one aldehyde (octadecanal) were also detected in the vegetative stage.

The composition of the volatiles strongly changes at the stage of flowering. The main component in this stage of leaf development was phytol. Phytol protects leaves from the loss of water and parasites. This compound has also been detected in the leaf of other studied *Astragalus* species. In *A. gombiformis*, phytol is the main component of leaf volatiles at flowering stage, while in other species including *A. cicer*, *A. spruneri* and *A. glycyphyllos* hydrocarbons were the main identified volatiles at flowering (Platikanov et al., 2005; Teyeb et al., 2011). Considerable amount of docosanol was detected at flowering stage. Alcoholic compounds could have important ecological functions in plants as allelochemicals. There are reports about the role of docosanol as a male pheromone in some insects, indicating the change in the plant-insect interactions at flowering stage (Müller and Buchbauer, 2011). Docosanol also could have medicinal properties as antiviral compound. Relatively high concentration of hydrocarbons and sterols also were detected in flowering stage.

The fructification stage is characterized by the presence of high concentrations of hydrocarbons and sterols and by the reduction or absence of alcohols in leaves. Except for cyclotetracosane and cyclooctacosane, other hydrocarbons have been previously reported in other species of *Astragalus*. Gamma-sitosterol was the only identified sterol in fructification stage. The distinctive features of this stage were the disappearance of alcohols and strong reduction of phytol. Most of other studied species of *Astragalus* also show a low concentration or absence of phytol at fructification phase. The reason could be the use of this component in the synthesis of lignin and/or by further oxidation of phenol compounds catalysed by the increase of polyphenoloxidase and peroxidase activity. According to Máñez et al. (1991) the changes in the composition of the volatiles with the maturation of the organ are directly related to higher rates of cyclization and dehydration of the compounds.

#### Total phenolic content

Based on obtained results, the TPC ranged from 5.01 to 8.28 µg GAE/mg DW (Table 2). The TPC (Figure 2) was significantly higher at the fructification stage than the vegetative and flowering stage. It has been shown that the mean temperature and solar radiation are significantly higher in summer months than the other months (Toor et al., 2006). Therefore, increase of TPC during fructification could be due to an increase in the amount of UV radiation associated with an increase in solar radiation received by the plants. The increased temperature and increase in the age of plants may have also caused an accumulation of phenolics in plants. Light increases the rate of biosynthesis of phenolics in plants by increasing the activities of enzymes, especially phenylalanine ammonia-lyase (PAL) which plays an important role in converting phenylalanine into coumaric acid, which are the initial precursor molecules involved in the synthesis of phenolic components in plants (Smith, 1973).



**Figure 2:** Variation of the Total phenolic contents (TPC) in leaves of *A. compactus* during development

**Table 2:** The antioxidant activity and Total phenolic contents (TPC) in leaves of *A. compactus* during development

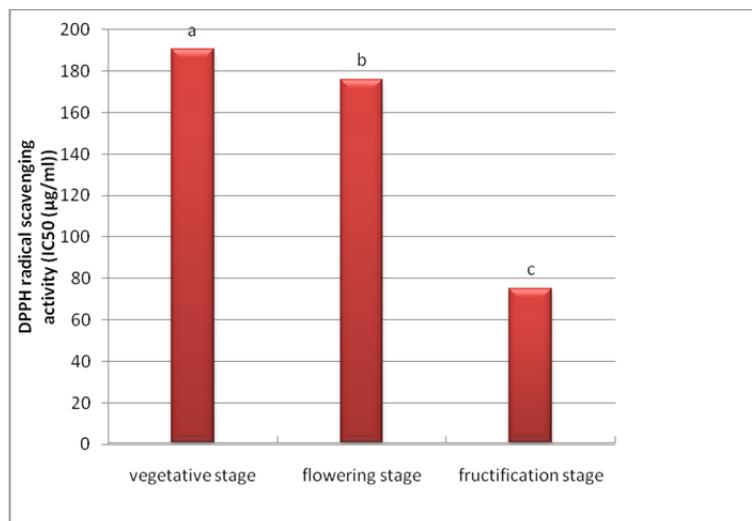
Developmental stage	Total phenolic content (TPC) ( $\mu\text{g GAE/mg DW}$ )	DPPH radical scavenging activity ( $\text{IC}_{50}$ ( $\mu\text{g/ml}$ ))
Vegetative stage	$5.01 \pm 0.09^b$	$190.4 \pm 4.89^a$
Flowering stage	$5.18 \pm 0.17^b$	$176.6 \pm 4.87^b$
Fructification stage	$8.28 \pm 0.12^a$	$75.0 \pm 3.07^c$

\* Values with different superscript letters are statistically different ( $p < 0.05$ ). Positive control (quercetin): 0.029  $\mu\text{g/ml}$

### Antioxidant activity

A developmental variation in antioxidant activity of *A. compactus* was observed with the  $\text{IC}_{50}$  ranging from 75 to 190.4  $\mu\text{g/ml}$  (Table 2). The capacity to quench free radical seemed to be related to the physiological stage. Figure 3 shows that the fructification stage exhibited the highest antioxidant activity (corresponds with the lowest  $\text{IC}_{50}$ ), whilst flowering and vegetative stages showed the less antioxidant activity (corresponds with higher  $\text{IC}_{50}$ ).

The higher antioxidant activity at fructification stage is related to the higher total phenolic content. Therefore, a positive relationship was observed between the antioxidant activity and the total phenolic content. The increase in the antioxidant activity and total phenolics at the fructification stage, which is characterized by the high temperature and a high exposure to solar radiation, leads us to conclude that *A. compactus* accumulates antioxidant compounds specially phenolic contents before and during this period probably to protect itself against solar radiations and/or grazers.



**Figure 3:** Variation of the antioxidant activity ( $\text{IC}_{50}$  ( $\mu\text{g/ml}$ )) in leaves of *A. compactus* during development

These results show that the antioxidant activity of *A. compactus* is influenced by the environmental conditions and the high solar radiation has a positive effect on the accumulation of major antioxidant components. The results suggest that *A. compactus* methanolic extracts may serve as potential sources of natural antioxidants and that the fructification phase could be considered as the best stage for the harvesting of the leaves of this plant species.

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