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Isolation and Biological Analysis of Aerial Parts of *Salvia Aethiopsis*

Erratum

2017-04-01

Isolation and Biological Analysis of Aerial Parts of *Salvia Aethiopis*

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ABSTRACT

The genus *Salvia* is the largest genus in the mint family (Lamiaceae), including nearly 1000 species which are widely distributed all over the world. The genus is reported for antioxidant, antimicrobial, and antiviral activities. Plants belonging to this genus are well known in folk medicine to treat epilepsy, aches, colds, bronchitis, and menstrual disorders. *Salvia aethiopis* is a perennial plant, known as Mediterranean or African sage. The ethanolic extract of aerial parts of *S. aethiopis* showed moderate level of inhibition in cannabinoid (CB1-37.0% displacement and CB2- 31.0% displacement) and opioid (Delta -46.3%, Kappa-45.3 % and Mu-32.9% displacement) receptors. Repeated silica gel column chromatography resulted in isolation of four compounds one new compound and three which were identified to be a sesquiterpene (spathulenol), and two sterols (β -sitosterol and β -sitosterol-3-O- β -D-glucoside). The structures of isolated compounds were determined using ¹H and ¹³C NMR and MS spectral data. The isolated compounds showed no activity towards cannabinoid and opioid receptor assay.

INTRODUCTION & BACKGROUND

It is a goal of the Sourcing, Isolation and Acquisition core to isolate and identify active components from plant material. The genus *Salvia* is the largest genus in the mint family (Lamiaceae), including nearly 1000 species which are widely distributed all over the world. Genus *Salvia* is a rich source of essential oils, flavonoids, and terpenoids. The genus is reported to have antioxidant, antimicrobial, and antiviral activities. Plants belonging to this genus are well known in folk medicine to treat epilepsy, aches, colds, bronchitis, and menstrual

disorders. *Salvia aethiopis* is a perennial plant, known as Mediterranean or African sage. In the United States it is best known as a noxious weed. It was probably introduced to North America as a contaminant of alfalfa seed but is native to Eurasia.

MATERIAL & METHODS

Plant material was collected during the flowering stages from foothill of Trans-Illi Alatau, Kazakhstan in June 2005. It was authenticated by botanist Dr. Nadezhda G. Gemejyeva, Botanist from the The Institute of Botany and

Phytointroduction, Almaty, Kazakhstan (Boucher specimen no.7290/25).

Dried aerial parts of *S. Aethiopsis* (0.17 kg) were ground in a Wiley-Mill plant grinder. Ground plant material was extracted using dichloromethane (1.9 L) at room temperature to yield 5.9 of DCM extract. The plant residue was further extracted using ethanol (1.7 L) producing 9.4 grams of ethanolic extract.

The ethanolic extract (4.1g) of aerial parts of *S. aethiopsis* was loaded on a normal phase column and eluted with methyl-chloride-methanol gradient to yield twelve fractions. Fraction 2 (15 mg eluted DCM) showed a single spot on a TLC and was identified to be spathulenol,

Compound II. Fraction 4 (1.2 g, 1% eluted MeOH- DCM) was subject to column chromatography using ethyl-acetate hexane gradient (1:9-1.1) to afford **Compound III** (15.0 mg). Fraction 8 (0.5 g eluted 5% MeOH-DCM) was purified on a silica gel column chromatography using CH₂-CL₂-MeOH gradient (1:0-9:1) to afford **Compound IV** (23.5 mg). Fraction 20 (0.75 g eluted 7% MeOH-DCM) was loaded on to a silica gel column chromatography using EtOAc-Hexane gradient (1:9-7:3) followed by DCM-MeOH (1:0-9:1) to afford **Compound I** (5mg).

RESULTS

ISOLATED COMPOUNDS:

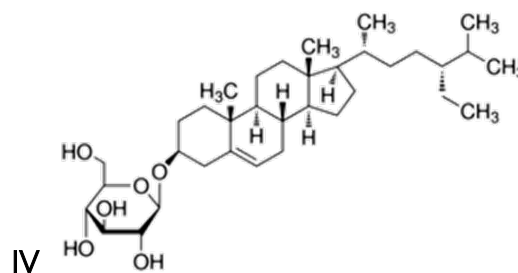
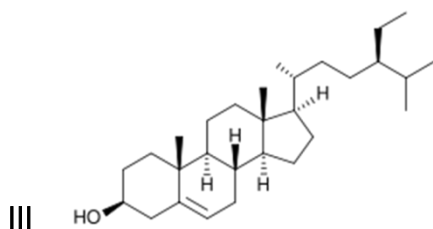
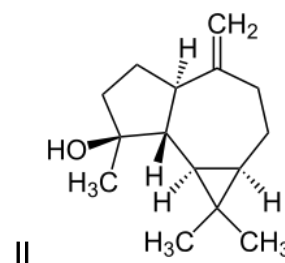
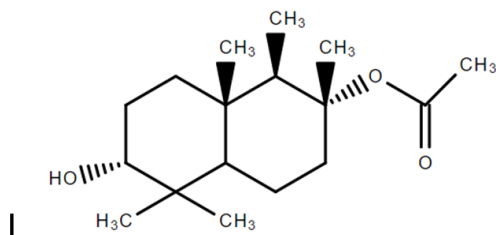


TABLE 1: ¹H, ¹³C, HMBC NMR data (500MHz, CDCL₂) for Compound 1

S.No.	¹ H NMR	¹³ C NMR (DEPT)	HMBC
1.	1.55	32.10 (T)	
2.	1.91M, 1.59M	24.99 (T)	
3.	3.35 (br,s)	75.71 (D)	
4.	-	37.42 (S)	
5.	1.53 m	48.03 (D)	
6.	2.34 m, 1.26 m	19.52 (T)	
7.	2.73 m, 1.71 m	38.84 (T)	
8.		86.29 (S)	
9.	2.27 m	55.02 (D)	C-12
10.		38.44 (S)	
11.	1.56 m, 1.28 m	29.75 (T)	
12.		178.41 (S)	
17.	1.48 (s)	19.89 (Q)	C-7,C-8, C-9
18.	0.94 (s)	28.33 (Q)	C-3, C-4,C-5, and C-19
19.	0.79 (s)	22.60 (Q)	C-3, C-4,C-5, and C-18
20.	0.82 (s)	15.65 (Q)	C-1, C-5,C-9, and C-10
<u>CH₃CO-</u>	1.87 (s)	21.95 (Q)	CH ₃ CO-
<u>CH₃CO-</u>		170.24 (S)	

DISCUSSION & CONCLUSION

The ethanolic extract of aerial parts of *S. aethiopsis* showed moderate to weak level of inhibition toward delta and kappa opioid receptors (46.3% and 45.3 %, respectively).

Bioassay guided fractionation and purification yielded four compounds (I-IV): 3 alpha-hydroxy-8alpha-acetoxy-13,14,15, 16-tetranorlabdan-12-oic acid (I, new), spathulenol (II), sitosterol (III), and sitosterol-3-O-glucoside (IV). Compound I is a new natural product.

The activity of the isolated components from *S. aethiopsis* did not show a high displacement for cannabinoid and opioid receptors. Further isolation shall continue with other species of plants of the *Salvia* genus with comparable bioassay results to find compounds active specific to these receptors that can be further studied for its effects on the body.

ACKNOWLEDGMENTS

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biological evaluation of *Salvia apiana*. Nat Prod Res

REFERENCES

- 1) Al-Aboudi AMF, Abu Zarga MH, Abu-Irmaileh BE, Awwadi FF, Khanfar MA. 2015. Three new seco-ursadiene triterpenoids from *Salvia syriaca*. Nat Prod Res. 29: 102-108.
- 2) Dennis LRJ. 1980. Gilkey's Weeds of the Pacific Northwest. Oregon State University Press, Corvallis, USA, pp. 245-246.
- 3) Gonzalez MS, San Segundo JM, Grande MC, Madarde M, Ballido IS. 1989. Sesterterpene lactones from *Salvia aethiopsis*. *Salviaethiopsisolide* and 13-*episalviaethiopsisolide*. Tetrahedron, 45: 3575-3582.
- 4) Güllüce M, Özer H, Barış Ö, Daferera D, ŞahĐn F, Polissiou M. 2006. Chemical Composition of the Essential Oil of *Salvia aethiopsis* L. Turk J Biol. 30: 231-233.
- 5) Hussain A, Adhikari A, Choudhary MI, Ayatollahi SA, Rahman A. 2016. New adduct of abietane-type diterpene from *Salvia leriifolia* Benth. Nat Prod Res. 30:1151-1156.
- 6) Jassbi AR, Eghtesadi F, Hazeri N, Ma'sumi H, Valizadeh J, Chandran JN, Schneider B, Baldwin IT. 2016. The roots of *Salvia rhytidea*: a rich source of biologically active diterpenoids. Nat Prod Res. 7:1
- 7) Srivedavyasri R, Hayes T, Ross SA. Forthcoming 2017. Phytochemical and