

Mansoura University

Mansoura Journal Of Chemistry



Phytochemical and Biological Evaluation of Urospermum Picroides

Howayda I. El-Nabawy^a, Dalia M., Ayyad^a, Mamdouh S. Serag^b, Mamdouh Abdel-Mogib^{a,*} ^aDepartment of Chemistry, Faculty of Science, Mansoura University, Mansoura-35516, Egypt; ^bDepartment of Botany, Faculty of Science, Damietta University, New Damietta, Egypt.

Received 23 May 2015; accepted 2 August 2015

Keywords Abstract The chromatographic separation of the dried Urospermum picroides; Urospermum picroides aerial part extracts of (L.) phytochemical SCOP.ex.F.W. Schmidt afforded urospermal A (1) and 8,15evaluation: dihydroxygermacra-1(10),4-dien-(12,6)-olide-14-al (2).quercetin 3-O-β-D-glucopyranosid (3) and kampferol 3-O-β-Dantioxidant: antimicrobial; glucopyranoside (4). GC/MS analysis of petroleum ether cytotoxicity. fraction from aerial parts gave α -amyrine, β -amyrine, 12oleanen-3-vl olean-18-en-3β-oi, acetate. moretenol. campesterol, stigmasterol, 14B-H-pregnae and stigmast-5-en-3 β -ol, in addition to many known compounds. Inhibited of the free radicals were found to be 86.9%, 85.3%, 86.1%, and 82.4% for seeds butanol fraction, seeds ethyl acetate fraction, aerial parts ethyl acetate fraction, and aerial parts methylene chloride fraction, respectively, compared to 88.1% inhibition by ascorbic acid. The sesquiterpene-rich sub fraction containing 1 and 2 and the flavonoid-rich sub fraction containing 3 and 4 inhibited 84.7% and 83.2%, respectively, which indicated the probable synergetic effect of other constituents in their main fractions. The antimicrobial activity was found to be (68.2%, 70.8% and 69.2%), (54.5%, 50.0% and 50.0%), for aerial parts ethyl acetate fraction, and compounds 3 and 4 towards the Gram positive bacteria S. aureus, the Gram negative bacteria E. coli and the yeast C. albicans, respectively. The seeds butanol fraction and the seeds ethyl acetate fraction had activities of 68.2% and 50.50% towards Gram positive S. aureus, respectively. The seeds butanol fraction was found to be very strong cytotoxic towards MCF-7 (9.4±0.37) and strong towards HePG-2 (14.7±0.85). The aerial parts ethyl acetate fraction was found to be very strong towards MCF-(78.8±0.47) and strong towards HePG-2 (10.1±0.88).

* Corresponding Author: Dr. Mamdouh Abdel-Mogib. <u>E-mail: mamdouh_m@mans.edu.eg</u>

1. Introduction

Urospermum picroides (L.) SCOP.ex.F.W. Schmidt (Compositae) (Aufs., 1795 & Boulos, 2002) is used as a medicinal plant for treatment of many diseases such as reducing the risk of chronic inflammatory diseases (Strzelecka et al., 2005), transcription factors as target of the antiinflammatory treatment (Stalinska et al., reducing postprandial and 2005). plateletaggregation in patients with metabolic syndrome (Fragopoulou et al., 2012). U. picroidesis sold in the markets of Dalmatia (Southern Croatia) as wild vegetables (Luczaj et al., 2013).

The phytochemical investigation of *U. picroides* showed the presence of luteolin-7-glucoside, quercetin, quercetin-3-galactoside, kaempferol-3-galactoside, chlorogenic and isochlorogenic acid (Giner *et al.*, 1992) as well as sesqueiterpene lactones and glucosides (Balboul *et al.*, 1997).

This article presents the results of the phytochemical, as well as biological reinvestigation.

<u>20</u>

2. Results and Discussion

2.1. Phytochemical Evaluation

The chromatographic separation of the dried aerial part extracts of U. picroides afforded urospermal A (1) and 8,15dihydroxygermacra-1(10),4-dien-(12,6)-olide-14-al (2), (Balboul et al., 1997), as well as quercetin 3-O-B-D-glucopyranosid (3) and kampferol-3-O-β-D-glucopyranoside (4), which were previously reported from the same species (Giner et al., 1992; Islam et al., 2012 and Cai et al., 2006). Compounds 1-4 were identified by comparing their ¹H NMR spectra with those of the corresponding compounds, or literature data. GC/MS analysis of petroleum ether fraction from aerial part gave α -amyrine, β -amyrine, 12acetate, olean-18-en-3-ol, oleanen-3-yl moretenol, campesterol, stigmasterol, 14β-Hpregnane and stigmast-5-en-3-ol, in addition to many known compounds as described in the experimental section.



R = OH, quercetin 3-O- β -D-glucopyranoside (3) R = H, kampferol-3-O- β -D-glucopyranoside (4)

using ascorbic acid as a reference antioxidant material were conducted. Table 1 showed that the seeds butanol fraction had inhibited (86.9%) of the free radicals, followed by the seeds ethyl acetate fraction (85.3%) which were very near to ascorbic acid (88.1%). The aerial parts ethyl acetate fraction had inhibited 86.1% of the free radicals, followed by the aerial parts methylene chloride fraction (82.4%). The sesquiterpene-rich sub-fraction containing compounds 1 and 2 and the



R = α -methyl, urospermal A (1) R = β -methyl,8,15-dihydroxygermacra-1(10),4-dien-(12,6)-olide-14-al (2)

2.2. Biological Evaluation

2.2.1 Antioxidant activity

The antioxidant activity of petroleum ether fraction, methylene chloride fraction, ethyl acetate fraction, butanol fraction, the sesquiterpenoid mixture of compounds 1,2, the flavonoid mixture of compounds 3,4 was assessed using ABTS method (Lissi *et al.*, 1999). A control experiment and another one flavonoid-rich sub fraction containing compounds 3 and 4 inhibited 84.7% and 83.2%, respectively, which indicated the

probable synergetic effect of other constituents in their main fractions.

Table	(1)): Antioxidant ac	tivity assessment	of	fractions of	of aerial	parts and seeds.
-------	-----	-------------------	-------------------	----	--------------	-----------	------------------

	Aerial	part	Seeds part		
Fraction or compound	Absorbance Inhibition %		Absorbance	Inhibition %	
Control	0.505	0%	0.505	0%	
Ascorbic acid	0.060	88.1%	0.060	88.1%	
Petroleum ether fraction	0.321	36.4%	0.422	17.2%	
Methylene chloride fraction	0.089	82.4%	0.225	55.9%	
Ethyl acetate fraction	0.070	86.1%	0.074	85.3%	
Butanol fraction			0.067	86.9%	
Compounds 1 & 2	0.085	83.2%		40 al 4	
Compounds 3&4	0.077	84.7%			

2.2.2 Antimicrobial activity

The antimicrobial activity was assessed (Stylianakis *et al.*, 2003) using the Gram positive bacteria *Staphylococcus aureus*, and the Gram negative bacteria *Escherichia coli* and the yeast *Candida albicans*. Ampicillin and colitrimazole were used as reference antibiotics. Table 2 showed that ethyl acetate fraction of aerial parts had high antimicrobial activity 68.2%, 70.8% and 69.2%, followed by compounds 3 and 4 which had a moderate activity 54.5%, 50.0% and 50.0% towards the Gram positive bacteria S. *aureus*, and the Gram negative bacteria E. *coli* and the yeast *C*. *albicans*. Table 3 showed that the seeds butanol fraction had high activity 68.2% towards Gram positive S. *aureus* and ethyl acetate fraction had a moderate activity 50.50% towards Gram positive S. *aureus*.

Table (2): Antimicrobial activity assessment data of ethyl acetate, methylene chloride, petroleum ether fractions, and compounds (1, 2) and (3, 4) from aerial parts.

	S. aur	eus	E. co	li	C. Albicans	
Fraction or	Diameter of	% Activity	Diameter of	%	Diameter of	%
	zone (in mm)	index	zone (in mm)	index	zone (in mm)	index
Petrolum ether	4	18.2	3	12.5	5	19.2
Methylene	8	36.4	5	20.8	7	26.9
Ethyl acetate fraction	15	68.2	17	70.8	18	69.2
Compounds 1&2	7	31.8	11	45.8	8	30.8
Compounds 3&4	12	54.5	12	50.0	13	50.0
Ampicillin	22	100	24	100	NA	NA
Colitrimazole	NA	NA	NA	NA	26	100

	S. au	reus	<i>E. c</i>	oli	C. Albi	cans
Compound	Diameter of inhibition zone (in mm)	% Activity index	Diameter of inhibition zone (in mm)	% Activity index	Diameter of inhibition zone (in mm)	% activity index
Petroleum ether fraction	2	9.1	NA	NA	NA	NA
Methylene chloride	7	31.8	4	16.7	10	38.5
Ethyl acetate fraction	11	50.50	10	41.7	7	26.9
Butanol fraction	15	68.2	11	45.8	8	30.8
Ampicillin	22	100	24	100	NA	NA
Colitrimazole	NA	NA	NA	NA	26	100

Table (3): Antimicrobial activity assessment	of butanol, ethyl acetate, methylene chloride and
netroleum ether from seeds.	

2.2.3 Cytotoxic activity

Cytotoxicity was expressed as the concentration that caused 50% loss of the cell monolayer (IC₅₀). The in vitro cytotoxicity against hepatocellular carcinoma, HePG-2 (liver) and mammary gland, MCF-7 (breast) was assessed (Denizot *et al.*, 1986). 5-Fluorouracil (5-FU) was used as a standard anticancer for comparison. Table 4 showed

that butanol fraction from seeds was very strong cytotoxic towards MCF-7 (9.4 ± 0.37) and strong towards HePG-2 (14.7 ± 0.85). The activity of ethyl acetate fraction of aerial parts was found to be very strong towards MCF-(78.8 ± 0.47) and strong towards (10.1 ± 0.88) HePG-2, while the flavonoid-rich sub fraction contained compounds 3 and 4 was found to be less active.

Table (4): Cytotoxic (IC ₅₀)	values of fractions of aerial part	rts and seeds on different cell lines.
--	------------------------------------	--

	Aerial	part	Seeds p	art
Fraction or compound	In vitro Cytotoxici	ty IC ₅₀ (µg/ml)	In vitro Cytotoxicit	y IC ₅₀ (μ g/ml)
	HePG2	MCF-7	HePG2	MCF-7
5-FU	6.6±0.24	4.7±0.11	6.6±0.24	4.7±0.11
Butanol fraction			14.7±0.85	9.4±0.37
Ethyl acetate fraction	10.1±0.88	8.8±0.47	30.2±1.83	18.1±0.96
Methylene chloride	45.6±1.91	34.9±1.74	41.1±3.04	29.9±1.88
fraction				
Petroleum ether fraction	79.5±3.52	90.6±4.56	88.6±4.50	70.3±3.92
1&2	36.2±1.55	53.1±2.30		
3 & 4	13.9±1.04	22.8±1.52	· · · · · · · · · · · · · · · · · · ·	

IC₅₀ (μ g/ml): 1 – 10 (very strong). 11 – 20 (strong). 21 – 50 (moderate). 51 – 100 (weak) and above 100 (non-cytotoxic).

3. Experimental

General:

The NMR spectra were recorder on a Varian MercyryVX-300 nmr spectrometer. ¹H-NMR spectra were run at 300 MHz in

deuterated chloroform (CDCl₃) or dimethylsulphoxide (DMSO-d₆); GC/MS Analysis: **Method 1:** Aglient 6890 gas chromatograph equipped with an Aglient mass spectrometric detector, with a direct capillary interface and fused silica capillary column HP-5ms (30m x 0.32 mm x 0.25 μ m film thickness). Helium was used as carrier

gas at approximate 1.0 ml/mim, pulsed splitess mode. The solvent delay was 3 min and the injection size was 1.0 µl. The mass spectrometric detector was operated in electron impact ionization mode with an ionizing energy of 70 e.v., scanning from m/z 50 to 500. The ion source temperature was 230°C. The electron multiplier voltage (EM voltage) was maintained at 1250 V above auto tune. The instrument was manually tuned using perfluorotributyl amine (PFTBA). The GC temperature program was started at 60°C (2 min) then elevated to 280°C at a rate of 8°C/min. The detector and injector temperature were set at 300 and 280°C, respectively. Wiley and Wiley Nist mass spectral data base was used in the identification of separated peaks. (Agriculture Research Center, National Research Center, NRC, Dokki, Cairo, Egypt); Method 2: Avarian GC interfaced to Finnigan SSQ 7000 Mass Selective Detector (MSD) with ICIS V2.0 data system for MS identification of the GC components. The column used was DB-5 (J & W Scientific, Folosm, CA) cross-linked fused silica capillary column (30m long, 0.25mm interrnal diameter) coated with polydimethylsiloxane (0.5µm film thickness). The oven temperature was programmed from 50°C for 3 min., at isothermal, then heating by 7°C/min. to 250°C and isothermally for 10 min., at 250°C. Injector temperature was 200°C and the volume injected was 0.5 µl. Transition-line and ion source temperature were 250°C and 150°C respectively. The mass spectrometer had a delay of 3 min. to avoid the solvent peak and then scanned from m/z 50 to m/z 300. Ionization energy was set at 70 ev. (National Research Center (NRC), Dokki, Cairo); Solvents: petroleum ether (60-80°C), diethyl ether, petroleum ether, methylene chloride, ethyl acetate, acetone, butanol and obtained methanol were from ElgomhoriaCompany; Chemical reagents for cytotoxicity activity: RPMI-1640 medium, MTT, DMSO and 5-fluorouracil (Sigma co., St. Louis, USA), fetal bovine serum (GIBCO, UK): Thin layer chromatography and preparative (TLC) were performed on silica gel (Kieselgel 60, GF 254) of 0.25 thickness.

Plant material:

Urospermum picroides was collected on April 2014 from the garden of Faculty of Science, Damietta University, and identified by Prof. Mamdouh Salem Serag, Botany Dept., Faculty Science, of Damietta University. Plant specimens were collected for pressing on herbarium sheets and later identification of species. Identification and nomenclature were followed (Tackholm, 1974 and Boulos, 1995, 2009). The voucher plant material and herbarium specimens of species recorded have been deposited in the Herbarium of Botany Dept., Faculty of Science, and Damietta University.

Processing of plant material:

The plant material was divided into two parts; aerial parts and seeds. Each part of them was air dried in shade at room temperature and grounded to give 400 g of aerial parts and 50 g of seeds dried powder material.

The aerial parts were extracted with $CH_2Cl_2/MeOH$ (1:1) for 48 hr, after that filtrated and the solvent was evaporated. Solvent extraction by using different solvents; petroleum ether 60-80°C, methylene chloride, ethyl acetate and butanol, respectively gave four fractions; petroleum ether fraction (6g), methylene chloride fraction (0.5g), ethyl acetate fraction (1.9g) and butanol fraction (0.9g).

A sample from petroleum ether fraction was analyzed by the GC/MS technique to give 2,6,10-trimethyl neophytadiene (Rt 21.38 min, 0.36%), 2,6,10trimethyl neophytadiene isomer (Rt 21.61min, 0.63%), methyl palmitate (Rt 22.21, 0.57%), palmitic acid (Rt22.83, 0.51%), heneicosane (Rt24.20, 0.96%), phytol isomer (Rt 24.40, 2.25%), docosane (Rt 25.31, 0.43%). 26.39, tricosane (R_i 0.84%). pentacosane (Rt 28.41, 0.82%), campesterol (Rt 37.73, 1.06%), stigmasterol (Rt 38.42, 6.36%), stigmast-5-en-3-ol (Rt 39.77, 3.82%), olean-12-en-3-ol (β-amyrine) (Rt 40.61, 5.12%), urs-12-en-3-ol (α -amyrine) (R_t 41.81, 5.02%), 12-oleanen-3-yl acetate (Rt 43.51, 8.28%), olean-18-en-3-ol (R_t 43.79, 3.17%) and moretenol (R_t 44.44, 2.09%).

A sample from the methylene chloride fraction was analyzed by the GC/MS technique to give 2-butoxyethanol (R_t 5.22, 2.19%), tetrahydrothiophene1,1-dioxide (R_t 11.59, 33.47%), (E)-4-(3-hydroxybut-1-en-1yl)-3,5,5trimethylcyclohex-2-en-1-one

(R₁18.59, 8.29%), (E)-4-(4-hydroxy-2,2,6trimethyl-7-oxabicyclo[4.1.0]heptan-1-yl)but-3-en-2-one (Rt 19.16, 1.87%), eicosane (Rt 1.57%), heneicosane (Rt 24.19, 23.02, 25.31, 3.10%), 1.76%), docosane (\mathbf{R}_{t}) tricosane (Rt 26.39, 3.45%), tetracosane (Rt 27.42, 4.36%), pentacosane (Rt 28.41, 3.10%). 2.74%), hexacosane (R_t 29.36, 2.30%) and (R_t) 29.91, heptacosene heptacosane (Rt 29.91, 2.17%).

ethyl acetate fraction was The silica gel column subjected to chromatography using ethyl acetate /methanol with increasing polarity. Fractions 13-16 were combined and evaporated to give a residue (0.2g), which was re-chromatographed over silica gel column with ethyl acetate /methanol (97:3) to give a yellow solid compound (3)followed by a mixture of compounds (3) and (4) (Giner et al., 1992).

The Butanol fraction 0.9g was subjected to silica gel column chromatography using ethyl acetate /methanol solvent system with increasing polarity. The fraction obtained by ethyl acetate /methanol (9:1) as eluent gave a mixture of compound (1) and compound (2) (Balboul *et al.*, 1997).

The seeds were extracted by a sohxlet extractor using different solvents; petroleum ether 60-80°C, methylene chloride, ethyl acetate and butanol respectively to give four fractions; a sample of pet. ether extract and another one of methylene chloride extract were analyzed by the GC/MS technique.

The sample of petroleum ether extract gave 2-decenal (R_t 12.64, 0.20%), tridecane (R_t 13.29, 0.34%), 2,4-decadienal (R_t 13.70, 0.24%), tetradecane (R_t 15.13, 1.14%), pentadecane (R_t 16.85, 2.94%), hexadecane (R_t 18.48, 2.98%), 2-methylhexadecane (R_t 19.41, 1.13%), heptadecane (R_t 20.01, 1.52%), octadecane (R_t 21.45, 3.78%), nonadecane (R_t 22.83, 3.98%), eicosane (R_t

and 14β-H-pregnane 24.14, 2.28%), stigmast-5-en-3-ol (24.65,1.49%), heneicosane (R_t 24.41, 4.61%), 9,12octadecadienoic acid (Rt 26.11, 5.70%), docosane (Rt 26.60, 4.33%), tricosane (Rt 27.77, 4.71%), tetracosane (Rt 28.10, 0.20%), (E)-pentacosa-1,3-diene (Rt 29.16, 0.48%), heptacosane (Rt 29.44, 0.22%), octacosen (Rt 29.90, 3.81%), nonacosen (Rt 30.89, 3.70%), henitiacontane (Rt 44.07, 3.76%), dotriacontane (Rt 40.93, 0.77%), tritriacontane (Rt 44.07, 0.57%), (23s)-ethyl cholest-5-en-3β-ol (Rt 45.25, 0.82%).

The sample of methylene chloride 25.33, 0.46%), tetradecane (\mathbf{R}_t) gave 31.30, 0.88%), 2,6,10hexadecane (R_t) trimethyltetradecane (Rt 34.06, 0.57%), 3-0.39%). 35.93, methylheptadecane (\mathbf{R}_i) octadecane (Rt 36.69, 1.22%), pytol (Rt 37.70, 0.79%), nonadecane (Rt 39.19, 1.21%), methyl palmitate (R, 39.91, 4.39%), eicosane (Rt 41.58, 3.00%), heneicosane (Rt 43.86, 8.48%), methyl-(E)-octadec-10-enoate (R_t 43.97, 1.95%), docosane (Rt 46.06, 7.51%), tricosane (Rt 48.17, 10.73%), tetracosane (Rt 50.19, 11.77%), pentacosane (Rt 52.13, 10.89%), hexacosane (Rt 54.00, 8.56%), heptacosane (Rt 55.80, 6.44%), octacosane (Rt 57.54, 4.82%), nonacosane (Rt 59.22, 3.63%), triacontane $(R_t 60.85, 2.47\%),$ hentriacontane (\mathbf{R}_t) 62.42, 1.80%), (R_t) 64.01, 1.16%), dotriacontane tritriacontane (Rt 65.86, 0.79%) and stigmast-5-en-3-ol (Rt 66.62, 0.84%).

Biological activity

Antioxidant activity:

Antioxidant activity screening assay ABTS method. For each of the investigated fractions or compounds (2 mL) of ABTS solution (60 μ M) was added to 3 mL MnO₂ solution (25mg/mL), all prepared in (5 mL) aqueous phosphate buffer solution (pH 7, 0.1 M). The mixture was shaken, centrifuged, filtered and the absorbance of the resulting green blue solution (ABTS radical solution) at 734 nm was adjusted to approx. ca. 0.5. Then, 50 μ l of (2 mM) solution of the tested compound in spectroscopic grade MeOH/phosphate buffer (1:1) was added. The absorbance was measured and the reduction in color intensity was expressed as inhibition percentage. L-ascorbic acid was used as standard antioxidant (positive control). A blank sample was run without ABTS and using MeOH/phosphate buffer (1:1) instead of test material. Negative control was run with ABTS and MeOH/phosphate buffer (1:1) only (Lissi *et al.*, 1999). The % inhibition was calculated by the formula:

Inhibition%= $\frac{Abz(control)-Abs(test)}{Abs(control)} x100$

Antimicrobial activity:

Antimicrobial fractions or compounds were individually tested against a panel of Gram positive bacteria Staphylococcus aureus, Gram negativebacteria, Escherichia coli and the yeast Candida albicans. The fraction or compound under investigation was dissolved in DMSO (1 mg /ml). Paper discs of Whatman filter paper were prepared with standard size (5mm) and sterilized in an autoclave. The paper discs were soaked in the desired concentration of the sample solution and placed aseptically in Petri dishes containing nutrient agar media (agar 20g + beef extract 3g + peptone 5g), seeded with Staphylococcus aureus, E. coli and Candida albicans. The Petri dishes were incubated at 36°C and the inhibition zones were recorded after 24h of incubation. Each treatment was replicated three times. The antimicrobial activity of a common standard antibiotic, ampicillin and antifungal, colitrimazole was also recorded using the same procedure as above at the same concentration and solvents (Stylianakis et al., 2003). The % activity index for the test samplewas calculated by the formula:

% Activity Index = Zone of inhibition by test compound (diametre) Zone of inhibition by standard (diametre)x 100

Cytotoxicity activity:

The cell lines mentioned above were used to determine the inhibitory effects of fractions or compounds on cell growth using the MTT assay. This colorimetric assay is based on the conversion of the yellow tetrazolium bromide (MTT) to a purple formazan derivative by mitochondrial succinate dehydrogenase in viable cells. Cell lines were cultured in RPMI-1640 medium with 10% fetal bovine serum. Antibiotics added were 100 units/ml penicillin and 100µg/ml streptomycin at 37°C in a 5% CO₂ incubator. The cell lines were seeded in a 96well plate at a density of 1.0x104cells/well at 37°C for 48 h under 5% CO₂. After incubation the cells were treated with different concentrations of fractions or compounds and incubated for 24 h. After 24 h of drug treatment, 20 µl of MTT solution at 5mg/ml was added and incubated for 4 h. Dimethyl sulfoxide (DMSO) in volume of 100 µl is added into each well to dissolve the purple formazan formed. The color intensity was measured and recorded at absorbance of 570 nm using a plate reader (EXL 800) (Denizot et al., 1986). The relative cell viability in percentage was calculated as:

The relative cell viability % = (A570 of treated samples/A570 of untreated sample) X 100

Table (5):	Gas chromatography	analysis of	different	fractions	from	aerial	parts	and	seeds	parts of
Urospermu	m picroides.									

Compound name	MS data (m/z [identity] (relative abundance %))
2-butoxyethanol	$118 [M]^{+} (1.6), 100 [C_{6}H_{12}O]^{+} (3.3), 87 [C_{5}H_{11}O]^{+} (23.3), 75 [C_{3}H_{7}O_{2}]^{+}$
	$(10), 57 [C_4H_9]^+ (100).$
tetrahydrothiophene -	120 $[M]^+$ (73.3), 56 $[C_4H_8]^+$ (100).
,1,1-dioxide	
2-decenal	154 [M] ⁺ (1.6),136 (3.3), 121 (10), 110 (20), 98 (30), 83 (66.6),70 (100), 55
	(83.3).
tridecane	$184 [M]^+ (3.3), 141 [C_{10}H_{21}]^+ (1.6), 127 [C_9H_{19}]^+ (1.6), 99 [C_7H_{15}]^+ (10), 85$
	$[C_6H_{13}]^+$ (43.3), 71 $[C_5H_{15}]^+$ (66.6), 57 $[C_4H_9]^+$ (100).

2,4-decadienal	152 $[M]^+$ (6.6),123 $[C_8H_{11}O]^+$ (3.3), 109 $[C_7H_9O]^+$ (3.3), 95 $[C_6H_7O]^+$ (10), 81 $[C_7H_7O]^+$ (100) 55 $[C_7H_7O]^+$ (13.3)
tetradecane	$198 [M]^{+} (6.6), 183 [C_{13}H_{27}]^{+} (1.6), 169 [C_{12}H_{25}]^{+} (1.6), 155 [C_{11}H_{23}]^{+} (1.6), 141 [C_{10}H_{21}]^{+} (1.6), 127 [C_{9}H_{19}]^{+} (1.6), 99 [C_{7}H_{15}]^{+} (13.3), 85 [C_{11}H_{23}]^{+} (1.6), 141 [C_{10}H_{21}]^{+} (1.6), 127 [C_{9}H_{19}]^{+} (1.6), 100 [C_{12}H_{15}]^{+} (1.6), 100 [C_{12}H_{15}]^$
pentadecane	$ \begin{array}{c} [C_{6}H_{13}] & (70), 71 [C_{3}H_{15}] & (73.5), 57 [C_{4}H_{9}] & (100). \\ 212 [M]^{+} (6.6), 169 [C_{12}H_{25}]^{+} (1.6), 155 [C_{11}H_{23}]^{+} (3.3), 141 [C_{10}H_{21}]^{+} \\ (3.3), 127 [C_{9}H_{19}]^{+} (6.6), 113 [C_{8}H_{17}]^{+} (10), 99 [C_{7}H_{15}]^{+} (13.3), 85 [C_{6}H_{13}]^{+} \\ \end{array} $
hexadecane	(56.6) , 71 $[C_{5}H_{11}]$ (76.6) , 57 $[C_{4}H_{9}]$ (100) . 226 $[M]^+$ (6.6), 155 $[C_{11}H_{23}]^+$ (20), 141 $[C_{10}H_{21}]^+$ (6.6), 127 $[C_{9}H_{19}]^+$ (10), 99 $[C_{7}H_{15}]^+$ (20), 85 $[C_{6}H_{13}]^+$ (66.6), 71 $[C_{5}H_{15}]^+$ (83.3), 57 $[C_{4}H_{9}]^+$ (100).
(E)-4-(3-hydroxybut-1- en-1-yl)-	208 $[M]^+$ (1.6), 193 $[C_{12}H_{17}O_2]^+$ (1.6), 165 $[C_{11}H_{17}O]^+$ (1.6), 152 $[C_{10}H_{16}O]^+$ (13.3), 135 $[C_{10}H_{15}]^+$ (6.6), 108 $[C_7H_9O]^+$ (100), 95 $[C_6H_7O]^+$
3,5,5trimethylcyclohex- 2-en-1-one	$(10), 55 [C_4H_7]$ (1.6).
(E)-4-(4-hydroxy-2,2,6- trimethyl-7- oxabicyclo[4.1.0]heptan-	224 $[M]^+$ (1.6), 209 $[C_{12}H_{17}O_3]^+$ (1.6), 191 $[C_{12}H_{15}O_2]^+$ (1.6), 123 $[C_7H_7O_2]^+$ (100), 109 $[M-C_7H_{15}O]^+$ (6.6), 95 $[C_6H_7O]^+$ (6.6), 55 $[C_4H_7]^+$ (3.3).
1-yl)but-3-en-2-one	
2-methylhexadecane	240 $[M]^+$ (0.8), 225 $[C_{16}H_{33}]^+$ (3.3), 210 $[C_{15}H_{30}]^+$ (0.8), 197 $[C_{14}H_{29}]^+$ (20), 169 $[C_{12}H_{25}]^+$ (1.6), 155 $[C_{11}H_{23}]^+$ (1.6), 141 $[C_{10}H_{21}]^+$ (13.3), 127 $[C_{9}H_{19}]^+$ (13.3), 113 $[C_{8}H_{17}]^+$ (16.6), 99 $[C_{7}H_{15}]^+$ (26.6), 85 $[C_{6}H_{13}]^+$ (56.6), 71
heptadecane	$\begin{array}{c} [C_{3}\Pi_{13}]^{+}(0.0), 57[C_{4}\Pi_{9}]^{+}(100).\\ 240[M]^{+}(0.8), 169[C_{12}H_{25}]^{+}(13.3), 155[C_{11}H_{23}]^{+}(6.6), 141[C_{10}H_{21}]^{+}\\ (10), 127[C_{9}H_{19}]^{+}(10), 113[C_{8}H_{17}]^{+}(13.3), 99[C_{7}H_{15}]^{+}(23.3), 85[C_{6}H_{13}]^{+}\\ (66) 21[C_{9}H_{19}]^{+}(10), 113[C_{8}H_{17}]^{+}(13.3), 99[C_{7}H_{15}]^{+}(23.3), 85[C_{6}H_{13}]^{+}\\ (66) 21[C_{9}H_{19}]^{+}(10), 113[C_{8}H_{17}]^{+}(10), 113[C_{8}H_{17}]^{+}(13.3), 99[C_{7}H_{15}]^{+}(23.3), 85[C_{6}H_{13}]^{+}\\ (66) 21[C_{9}H_{19}]^{+}(10), 113[C_{8}H_{17}]^{+}(10), 113[C_{8$
2.6.10 trimethyl	$(00.0), 71 [U_5H_{15}] (80.0), 57 [U_4H_9] (100).$ 278 $[M_1^+ (12.3), 170 [U_1H_1^+ (10), 122 [U_1H_1^+ (42.2), 111 (26.6), 05]$
neonbytadiene	$[C_{H_{11}}^{+}(90) 82 [C_{H_{11}}^{+}(100) 68 [C_{H_{11}}^{+}(66.6) 57 [C_{H_{11}}^{+}(72.2)]$
octadecane	$254 \text{ [M]}^+ (6.6) 225 \text{ [C}_{4}\text{H}_{2}\text{]}^+ (0.8) 211 \text{ [C}_{4}\text{H}_{2}\text{]}^+ (1.6) 197 \text{ [C}_{4}\text{H}_{2}\text{]}^+$
oonacouno	$(3.3), 155 [C_1H_{22}]^+ (6.6), 141 [C_10H_{22}]^+ (10), 127 [C_2H_{12}]^+ (13.3), 113$
	$[C_8H_{17}]^+$ (16.6), 99 $[C_7H_{15}]^+$ (26.6), 85 $[C_6H_{13}]^+$ (73.3), 71 $[C_5H_{15}]^+$ (86.6), 57 $[C_4H_9]^+$ (100).
2,6,10-trimethyl	278 $[M]^+(16.6)$, 179 $[C_{13}H_{23}]^+(10)$, 137 $[C_{10}H_{17}]^+(13.3)$, 123 $[C_9H_{15}]^+(50)$,
neophytadiene isomer	$109 [C_8H_{13}]^+ (23.3), 95 [C_7H_{11}]^+ (90), 81 [C_6H_9]^+ (100), 68 [C_5H_8]^+ (60), 57$
methyl palmitate	$[C_4H_9]$ (66.6). 270 $[M]^+$ (13.3), 239 $[C_{16}H_{31}O]^+$ (10), 227 $[C_{14}H_{27}O_2]^+$ (13.3), 185 $[C_{14}H_2O_2]^+$ (8.5) 143 $[C_{14}H_{12}O_2]^+$ (13.2), 185
	$[C_4H_7O_2]^+(66.6), 74 [C_3H_4O_3]^+(100), 57 [C_4H_7]^+(36.6)$
palmitic acid	$268 [M]^+ (6.6), 239 [C_{17}H_{35}]^+ (1.6), 225 [C_{16}H_{33}]^+ (3.3), 197 [C_{14}H_{29}]^+$
	$(3.3), 183 [C_{13}H_{27}]^+ (6.6), 169 [C_{12}H_{25}]^+ (13.3), 155 [C_{11}H_{23}]^+ (6.6), 141$
	$[C_{10}H_{21}]^{+}$ (10), 127 $[C_{9}H_{19}]^{+}$ (13.3), 113 $[C_{8}H_{17}]^{+}$ (16.6), 99 $[C_{7}H_{15}]^{+}$ (26.6),
	85 $[C_6H_{13}]^+$ (76.6), 71 $[C_5H_{15}]^+$ (90), 57 $[C_4H_9]^+$ (100).
nonadecane	256 [M] ⁺ (23.3), 213 [C ₁₃ H ₂₅ O ₂] ⁺ (33.3), 171 [C ₁₀ H ₁₉ O ₂] ⁺ (20), 129
eicosana	$[C_7H_{13}O_2]$ (36.6), 97 $[C_6H_9O]$ (30), 73 $[C_3H_5O_2]$ (100), 57 $[C_4H_9]$ (73.3).
cicosalic	(1.6) (1.6) (1.6) (2.5) $(2.5$
	$[C_7H_{15}]^+$ (20) 85 $[C_6H_{12}]^+$ (56.6) 71 $[C_7H_{12}]^+$ (80) 57 $[C_7H_{12}]^+$ (10)
heneicosane	296 [M]^+ (3.3), 253 [C ₁₈ H ₂₂] ⁺ (1.6), 239 [C ₁₇ H ₂₆] ⁺ (1.6), 183 [C ₁₂ H ₂₇] ⁺
	$(3.3), 155[C_{11}H_{23}]^+$ (6.6), 141 $[C_{10}H_{21}]^+$ (6.6), 127 $[C_{0}H_{10}]^+$ (10), 113
	$[C_8H_{17}]^+$ (10), 99 $[C_7H_{15}]^+$ (20), 85 $[C_6H_{13}]^+$ (70), 71 $[C_5H_{11}]^+$ (86.6), 57
	[C₄H ₉] ⁺ (100).
phytol isomer	278 $[M-H_2O]^+$ (1.6), 207 $[C_{15}H_{27}]^+$ (1.6), 196 $[C_{13}H_{24}O]^+$ (1.6), 165
	$[C_{12}H_{21}]^{+}$ (1.6), 137 $[C_{10}H_{17}]^{+}$ (1.6), 123 $[C_{9}H_{15}]^{+}$ (36.6), 95 $[C_{7}H_{11}]^{+}$ (33.3),

Phytochemical and Biological Evaluation of Urospermum Picroides

$ \begin{array}{llllllllllllllllllllllllllllllllllll$		
$ \begin{array}{llllllllllllllllllllllllllllllllllll$	140.11	$\delta_{1} [C_{6} H_{9}] (33.3), 11 [C_{4} H_{7}] (33.3), 51 [C_{4} H_{7}] (33.3).$
$ \begin{array}{llllllllllllllllllllllllllllllllllll$	14p-H-pregna	288 [M] (0.8), 206 (73.3), 165 (30), 141 (20), 111 (33.3), 85 (53.3), 57 (100).
	docosane	$310 [C_{22}H_{46}]^{+} (3.3), 281 [C_{20}H_{41}]^{+} (1.6), 267 [C_{19}H_{39}]^{+} (1.6), 253 [C_{18}H_{37}]^{+}$
		$(1.6),239 [C_{17}H_{35}]^* (1.6), 169 [C_{12}H_{25}]^* (3.3), 141 [C_{10}H_{21}]^* (10), 127$
		$[C_9H_{19}]^+$ (10), 113 $[C_8H_{17}]^+$ (13.3), 99 $[C_7H_{15}]^+$ (20), 85 $[C_6H_{13}]^+$ (60),71
9,12- octadecadienoic acid tricosane 1200 [M] ⁺ (23.3), 235 [C ₁ H ₂₄ Q ₂₁ ⁺ (6.6), 165 [C ₁ P ₄₁] ⁺ (13.3), 123 (26.6), 81 (C ₆ H ₃₁ ⁺ (100), 55 [C ₄ H ₁] ⁺ (86.6). 324 [M] ⁺ (3.3), 295 [C ₁ H ₄₁] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 267 [C ₁₉ H ₃₉] ⁺ (1.6), 234 [M] ⁺ (3.3), 295 [C ₁ H ₄₁] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 285 [C ₄ H ₁₃] ⁺ (10), 127 [C ₉ H ₁₉] ⁺ (10), 13 [C ₄ H ₁₇] ⁺ (13.3), 99 [C ₇ H ₃₁] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 267 [C ₁₉ H ₃₉] ⁺ (1.6), 271 [C ₄ H ₁₇] ⁺ (10), 127 [C ₉ H ₁₉] ⁺ (10), 113 [C ₄ H ₁₇] ⁺ (13.3), 99 [C ₇ H ₁₃] ⁺ (20), 85 [C ₄ H ₄₁] ⁺ (10), 127 [C ₉ H ₁₉] ⁺ (10), 113 [C ₄ H ₁₇] ⁺ (13.3), 99 [C ₇ H ₁₃] ⁺ (20), 85 [C ₄ H ₄₁] ⁺ (10), 127 [C ₉ H ₁₉] ⁺ (1.6), 229 [C ₁₇ H ₃₁] ⁺ (1.6), 248 [C ₁₇ H ₃₁] ⁺ (1.6), 253 [C ₁₄ H ₃₂] ⁺ (6.6), 155 [C ₁₄ H ₂₁] ⁺ (6.6), 85 [C ₆ H ₁₃] ⁺ (6.6), 216 [C ₁₂ H ₂₃] ⁺ (6.5), 169 [C ₁₂ H ₂₃] ⁺ (10), 141 [C ₁₀ H ₂₁] ⁺ (6.6), 85 [C ₆ H ₁₃] ⁺ (70), 57 [C ₄ H ₃₁] ⁺ (100). (E)-pentacosa-1, 3-diene 48 [M] ⁺ (1.6), 225 [C ₁₆ H ₃₃] ⁺ (6.6), 113 [C ₈ H ₁₇] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 285 [C ₆ H ₁₃] ⁺ (6.6), 169 [C ₁₂ H ₂₃] ⁺ (6.6), 169 [C ₁₂ H ₂₃] ⁺ (10), 141 [C ₁₀ H ₂₁] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 169 [C ₁₂ H ₂₃] ⁺ (3.3), 141 [C ₁₀ H ₂₁] ⁺ (10), 127 [C ₄ H ₁₇] ⁺ (10), 113 [C ₄ H ₁₇] ⁺ (1.3), 92 [C ₄ H ₄₃] ⁺ (6.6), 169 [C ₁₂ H ₂₃] ⁺ (1.6), 128 [C ₁₀ H ₄₁] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 162 [C ₁₂ H ₂₃] ⁺ (1.6), 323 [C ₄ H ₄₁₃] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 281 [C ₄₀ H ₄₁] ⁺ (1.6), 28		$[C_{4}H_{11}]^{+}$ (80), 57 $[C_{4}H_{6}]^{+}$ (100).
octadecadienoi acid tricosane $\begin{bmatrix} C_{6}H_{3}^{-1}(100), 55 [C_{4}H_{3}^{-1}(6.6), 218 [C_{20}H_{4}]^{+}(1.6), 267 [C_{19}H_{39}]^{+}(1.6), 253 [C_{10}H_{39}]^{+}(1.6), 253 [C_{10}H_{39}]^{$	9.12-	280 [M]^{+} (23.3) 236 [CusHarOa] ⁺ (6.6) 165 [CusHarl ⁺ (13.3) 123 (26.6) 81
tricosane 124 [19] (100), 295[C ₁ ,H ₄₁] (1.6), 281 [C ₂₀ H ₄₁] (1.6), 267 [C ₁₉ H ₃₉] (1.6), 253 [C ₁₄ H ₃₇] (1.6), 239 [C ₁₇ H ₃₁] (1.6), 169 [C ₁₂ H ₂₃] (1.3), 141 [C ₁₀ H ₂₁] (1.6), 253 [C ₁₄ H ₃₇] (1.6), 239 [C ₁₇ H ₃₁] (1.6), 299 [C ₁₇ H ₃₁] (1.6), 285 [C ₄ H ₁₃] (1.6), 100, 127 [C ₄ H ₁₃] (10), 113 [C ₄ H ₁₇] (100). 338 [M1 (3.3), 232 [C ₂₃ H ₄₇] (1.6), 239 [C ₁₇ H ₃₃] (1.6), 169 [C ₁₂ H ₂₃] (2.3), 267 [C ₁₉ H ₃₉] (1.6), 253 [C ₁₄ H ₃₁] (1.6), 239 [C ₁₇ H ₃₃] (1.6), 169 [C ₁₂ H ₂₃] (2.3), 267 [C ₁₉ H ₃₉] (1.6), 253 [C ₁₄ H ₃₁] (1.6), 239 [C ₁₇ H ₃₃] (1.6), 252 [C ₁₄ H ₃₁] (1.6), 267 [C ₁₉ H ₃₉] (1.6), 253 [C ₁₄ H ₃₁] (1.6), 239 [C ₁₇ H ₃₃] (1.6), 252 [C ₁₄ H ₃₁] (1.6), 267 [C ₁₉ H ₃₉] (1.6), 253 [C ₁₄ H ₃₁] (1.6), 259 [C ₁₄ H ₃₁] (1.6), 252 [C ₁₄ H ₃₁] (1.6), 277 [C ₄ H ₃₁] (100). 288 [M1 (1.6), 225 [C ₁₄ H ₃₁] (1.6), 259 [C ₂₁ H ₄₁] (1.6), 253 [C ₁₄ H ₃₁] (1.6), 266 [M1 (1.6), 253 [C ₁₄ H ₃₁] (1.6), 253 [C ₁₄ H ₃₁] (1.6), 281 [C ₂₀ H ₄₁] (1.6), 57 [C ₄ H ₃₁] (1.6), 267 [C ₁₉ H ₃₉₁] (1.6), 253 [C ₁₄ H ₃₁] (1.6), 281 [C ₂₀ H ₄₁] (1.6), 169 [C ₁₂ H ₂₃] (3.3), 141 [C ₁₀ H ₁₁₁] (10), 127 [C ₂ H ₁₁] (10), 113 [C ₄ H ₁₇₁] (1.6), 169 [C ₁₂ H ₂₃] (3.3), 141 [C ₁₀ H ₃₁] (60), 71 [C ₄ H ₁₁] (1.6), 253 [C ₁₄ H ₃₁] (1.6), 169 [C ₁₂ H ₂₃] (3.3), 141 [C ₁₀ H ₃₁] (2.6), 155 [C ₄ H ₃₁] (1.6), 253 [C ₁₄ H ₃₁] (1.6), 267 [C ₁₄ H ₃₁] (1.6), 169 [C ₁₂ H ₂₃] (1.6), 169 [C ₁₂ H ₂₃] (1.6), 171 [C ₄ H ₁₂] (1.6), 189 [C ₄ H ₁₂] (1.6), 251 [C ₄ H ₃₁] (1.6), 251 [C ₄ H ₃₁] (1.6), 169 [C ₁₂ H ₂₃] (1.6), 113 [C ₄ H ₁₁₁] (1.6), 253 [C ₁₄ H ₃₁] (1.6), 113 [C ₄ H ₁₁₁] (1.6), 253 [C ₁₄ H ₃₁] (1.6), 257 [C ₄ H ₃₁] (1.6), 257 [C ₄ H ₃₁] (1.6), 257 [C ₄ H ₃₁] (1.6	octadecadienoic acid	$[C_{H_{a}}]^{+}(100)$ 55 $[C_{H_{a}}]^{+}(86.6)$
$ \begin{array}{llllllllllllllllllllllllllllllllllll$	tricosane	$224 [M]^{+} (2.2) 205[C_{-}U_{-}]^{+} (1.6) 281 [C_{-}U_{-}]^{+} (1.6) 267 [C_{-}U_{-}]^{+} (1.6)$
$ \begin{array}{llllllllllllllllllllllllllllllllllll$	uicosane	$524 [M] (5.5), 255 [C_2[H43] (1.0), 261 [C_{20}H41] (1.0), 207 [C_{19}H39] (1.0), 253 [C_{10}H_{10}] (1.0), 253 [C_{10}H$
$ \begin{array}{llllllllllllllllllllllllllllllllllll$		$255 [C_{18}H_{37}] (1.0), 259 [C_{17}H_{35}] (1.0), 109 [C_{12}H_{25}] (3.3), 141 [C_{10}H_{21}] (1.0), 107 [C_{11}H_{11}] (1.0), 112 [C_{11}H_{12}] (1.0), 107 [C_{11}H_{12$
		$(10), 127 [C_9H_{19}]$ $(10), 113 [C_8H_{17}]$ $(13.3), 99 [C_7H_{15}]$ $(20), 85 [C_6H_{13}]$
tetracosane 338 [M] ⁺ (3.3), 323 [C ₂₃ H ₄₃] ⁺ (1.6), 295 [C ₂₁ H ₄₃] ⁺ (1.6), 169 [C ₁₂ H ₂₃] ⁺ (3.3), 414 [C ₁₀ H ₁₃] ⁺ (10), 127 [C ₄ H ₁₉] ⁺ (10), 113 [C ₄ H ₁₇] ⁺ (1.5), 199 [C ₁ H ₄₃] ⁺ (1.6), 169 [C ₁₂ H ₂₃] ⁺ (3.3), 414 [C ₁₀ H ₄₁] ⁺ (10), 217 [C ₄ H ₁₉] ⁺ (10), 127 [C ₄ H ₁₉] ⁺ (10), 225 [C ₄ H ₄₉] ⁺ (10), 235 [C ₄ H ₄₁] ⁺ (23.3), 57 [C ₄ H ₉] ⁺ (1.6), 57 [C ₄ H ₉₁] ⁺ (1.6), 225 [C ₁₁ H ₂₃] ⁺ (6.6), 85 [C ₆ H ₁₃] ⁺ (6.3), 171 [C ₄ H ₁₁] ⁺ (80), 57 [C ₄ H ₉₁] ⁺ (100). 57 [C ₄ H ₉₁] ⁺ (100). 141 [C ₁₀ H ₂₁] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 57 [C ₄ H ₉₁] ⁺ (100). 141 [C ₁₀ H ₂₁] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 57 [C ₄ H ₉₁] ⁺ (100) a66 [C ₁₁ H ₂₁] ⁺ (1.6), 225 [C ₁₁ H ₃₃] ⁺ (1.6), 225 [C ₁₁ H ₃₃] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 57 [C ₄ H ₉₁] ⁺ (100) a53 [C ₆ H ₄₁] ⁺ (1.6), 225 [C ₁₁ H ₃₁] ⁺ (1.6), 225 [C ₁₁ H ₃₁] ⁺ (1.6), 225 [C ₁₁ H ₃₁] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 57 [C ₄ H ₉₁] ⁺ (1.6), 253 [C ₁₄ H ₃₁] ⁺ (1.6), 225 [C ₁₁ H ₃₁] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 267 [C ₁₀ H ₂₁] ⁺ (20), 81 [C ₂₁ H ₄₁] ⁺ (1.6), 225 [C ₁₁ H ₃₁] ⁺ (1.6), 253 [C ₁₁ H ₃₁] ⁺ (1.6), 251 [C ₁₁ H ₃₁] ⁺ (1.6), 253 [C ₁₁ H ₃₁] ⁺ (1.6), 251 [C ₁₁ H ₃₁] ⁺ (1.6), 267 [C ₁₀ H ₃₉] ⁺ (1.6), 253 [C ₁₁ H ₃₁] ⁺ (1.6), 257 [C ₁₀ H ₃₂] ⁺ (1.6), 253 [C ₁₁ H ₃₁] ⁺ (1.6), 257 [C ₁₀ H ₃₂] ⁺ (1.6), 253 [C ₁₁ H ₃₁] ⁺ (1.6), 257 [C ₁₀ H ₃₂] ⁺ (1.6), 253 [C ₁₂ H ₃₁] ⁺ (1.6), 257 [C ₁₁ H ₃₁] ⁺ (2.0), 113 [C ₄ H ₁₁] ⁺ (2.0), 113 [C ₄ H ₁₁] ⁺ (2.0), 257 [C ₄ H ₃₁] ⁺ (2.0), 257 [C		$(60),71 [C_{5}H_{11}]^{\circ} (80), 57 [C_{4}H_{9}]^{\circ} (100).$
$ \begin{array}{llllllllllllllllllllllllllllllllllll$	tetracosane	338 [M] ⁺ (3.3), 323 [C ₂₃ H ₄₇] ⁺ (1.6), 295[C ₂₁ H ₄₃] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6),
$ \begin{array}{llllllllllllllllllllllllllllllllllll$		$267 \left[C_{19}H_{39}\right]^{+} (1.6), 253 \left[C_{18}H_{37}\right]^{+} (1.6), 239 \left[C_{17}H_{35}\right]^{+} (1.6), 169 \left[C_{12}H_{25}\right]^{+}$
		$(3.3), 141 [C_{10}H_{21}]^{+} (10), 127 [C_{9}H_{19}]^{+} (10), 113 [C_{8}H_{17}]^{+} (13.3), 99$
$ \begin{array}{llllllllllllllllllllllllllllllllllll$		$[C_7H_{15}]^+$ (20), 85 $[C_6H_{13}]^+$ (60), 71 $[C_5H_{11}]^+$ (83.3), 57 $[C_4H_9]^+$ (100).
$ \begin{array}{llllllllllllllllllllllllllllllllllll$	pentacosane	352 [M]^{+} (3.3), 281 [C ₂₀ H ₄₁] ⁺ (10), 239 [C ₁₇ H ₃₅] ⁺ (1.6), 225 [C ₁₆ H ₃₃] ⁺ (1.6),
$ \begin{array}{llllllllllllllllllllllllllllllllllll$	P	$183 \left[C_{12}H_{22} \right]^{+} (6.6) 155 \left[C_{11}H_{22} \right]^{+} (6.6) 85 \left[C_{2}H_{12} \right]^{+} (63.3) 71 \left[C_{2}H_{11} \right]^{+} (80)$
$ \begin{array}{llllllllllllllllllllllllllllllllllll$		57 [C.H.] ⁺ (100)
	(E) mantagana 1 7 diana	$240 \text{ [N4]}^+ (1.6) - 225 \text{ [C]} U = 1^+ (6.6) - 112 \text{ [C]} U = 1^+ (22.2) 107 \text{ [C]} U = 1^+$
$ \begin{array}{llllllllllllllllllllllllllllllllllll$	(E)-pentacosa-1, 3-utene	$5^{+}6$ [$1^{+}1$] (1.0), 225 [$C_{16}1133$] (0.0), 115 [$C_{8}1177$] (25.5) 157 [$C_{14}1129$]
$\begin{array}{llllllllllllllllllllllllllllllllllll$		$(0.0), 109 [C_{12}H_{25}] (10), 141 [C_{10}H_{21}] (10.0), 85 [C_6H_{13}] (70), 57$
hexacosane 366 [M] (1.6), 323 [C ₃ H ₄₇] (1.6), 295 [C ₁ H ₄₃] (1.6), 281 [C ₂₀ H ₄₁] (1.6), 297 [C ₁ H ₂₃] ⁺ (3.3), 141 [C ₁₀ H ₂₁] ⁺ (10), 127 [C ₅ H ₁₉] ⁺ (10), 113 [C ₈ H ₁₇] ⁺ (1.6), 169 [C ₁₂ H ₂₃] ⁺ (3.3), 141 [C ₁₀ H ₂₁] ⁺ (00), 127 [C ₅ H ₁₉] ⁺ (10), 113 [C ₈ H ₁₇] ⁺ (100). 392 [M] ⁺ (0.8), 309 [C ₂₂ H ₄₃] ⁺ (3.3), 281 [C ₂₀ H ₄₁] ⁺ (6.6), 253 [C ₁₈ H ₃₇] ⁺ (6.6), 197 [C ₁₄ H ₂₉] ⁺ (6.6), 169 [C ₁₂ H ₂₅] ⁺ (10), 141 [C ₁₀ H ₂₁] ⁺ (20), 113 [C ₈ H ₁₇] ⁺ (30), 85 [C ₆ H ₁₃] ⁺ (90), 57 [C ₄ H ₉] ⁺ (100). 378 [M] ⁺ (1.6), 295 [C ₂₁ H ₄₃] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 267 [C ₁₉ H ₃₉] ⁺ (1.6), 253 [C ₁₈ H ₃₇] ⁺ (1.6), 239 [C ₁₇ H ₃₅] ⁺ (1.6), 169 [C ₁₂ H ₂₂] ⁺ (3.3), 141 [C ₁₀ H ₂₁] ⁺ (10), 127 [C ₅ H ₁₉] ⁺ (10), 113 [C ₈ H ₁₇] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 357 [C ₄ H ₉] ⁺ (1.6), 322 [C ₂₃ H ₄₇] ⁺ (1.6), 255 [C ₁₄ H ₃₃] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 267 [C ₁₉ H ₃₉] ⁺ (1.6), 253 [C ₁₈ H ₃₇] ⁺ (1.6), 251 [C ₂₃ H ₄₁] ⁺ (1.6), 285 [C ₆ H ₁₃] ⁺ (1.6), 322 [C ₂₃ H ₄₇] ⁺ (1.6), 295 [C ₂₁ H ₄₃] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 267 [C ₁₉ H ₃₉] ⁺ (1.6), 253 [C ₁₈ H ₃₇] ⁺ (1.6), 295 [C ₂₁ H ₄₃] ⁺ (1.6), 169 [C ₁₂ H ₂₅] ⁺ (3.3), 141 [C ₁₀ H ₂₁] ⁺ (10), 127 [C ₅ H ₁₉] ⁺ (10), 113 [C ₈ H ₁₇] ⁺ (13.3), 99 [C ₇ H ₁₅] ⁺ (20), 85 [C ₆ H ₁₃] ⁺ (56.6), 71 [C ₃ H ₁₁] ⁺ (80), 57 [C ₄ H ₉] ⁺ (100). 3-methylheptadecane 265 [C ₁₆ H ₃₃] ⁺ (6.6), 197 [C ₁₄ H ₂₉] ⁺ (10), 169 [C ₁₂ H ₂₅] ⁺ (10), 141 [C ₁₀ H ₂₁] ⁺ (20), 113 [C ₈ H ₁₇] ⁺ (30), 85 [C ₆ H ₁₃] ⁺ (3.3), 267 [C ₁₉ H ₃₉] ⁺ (6.6), 225 [C ₁₆ H ₃₃] ⁺ (6.6), 197 [C ₁₄ H ₂₉] ⁺ (10), 159 [C ₁₂ H ₂₅] ⁺ (10), 141 [C ₁₀ H ₂₁] ⁺ (23.3), 396 [C ₇ H ₄₅] ⁺ (13.3), 367 [C ₂₇ H ₄₃] ⁺ (6.6), 315 [C ₄ H ₁₇] ⁺ (100). 254 [M] ⁺ (2.1), 207 [C ₁₅ H ₂₇] ⁺ (1), 151 [C ₁₁ H ₁₉] ⁺ (4.3), 137 [C ₁₀ H ₁₇] ⁺ (10.8), 123 [C ₉ H ₁₃] ⁺ (58.6), 95 [C ₇ H ₁₁] ⁺ (100), 81 [C ₆ H ₉] ⁺ (6.6), 353 [C ₄ H ₁₃] ⁺ (86.9). campesterol 400 [M] ⁺ (23.3), 388 [C ₂		$[C_4H_9]$ (100)
$\begin{array}{llllllllllllllllllllllllllllllllllll$	hexacosane	$366 [M]'(1.6), 323 [C_{23}H_{47}]'(1.6), 295 [C_{21}H_{43}]'(1.6), 281 [C_{20}H_{41}]'(1.6),$
$\begin{array}{llllllllllllllllllllllllllllllllllll$		$267 [C_{19}H_{39}]^{\circ} (1.6), 253 [C_{18}H_{37}]^{\circ} (1.6), 239 [C_{17}H_{35}]^{\circ} (1.6), 169 [C_{12}H_{25}]^{\circ}$
octacosen $ \begin{bmatrix} [C_7H_{15}]^{+} (20), 85 [C_6H_{13}]^{+} (60), 71 [C_3H_{11}]^{+} (80), 57 [C_4H_9]^{+} (100). \\ 392 [M]^{+} (0.8), 309 [C_{22}H_{31}]^{+} (3.3), 281 [C_{20}H_{11}]^{+} (6.6), 253 [C_{18}H_{37}]^{+} \\ (.6.6), 197 [C_{14}H_{29}]^{+} (.6.6), 169 [C_{12}H_{25}]^{+} (10), 141 [C_{10}H_{21}]^{+} (20), 113 \\ [C_8H_{17}]^{+} (30), 85 [C_6H_{13}]^{+} (90), 57 [C_4H_9]^{+} (100). \\ 378 [M]^{+} (1.6), 295 [C_{21}H_{43}]^{+} (1.6), 281 [C_{20}H_{41}]^{+} (1.6), 267 [C_{19}H_{39}]^{+} (1.6), 253 [C_{18}H_{37}]^{+} (1.6), 295 [C_{21}H_{31}]^{+} (1.6), 399 [C_{7}H_{15}]^{+} (20), 85 [C_6H_{13}]^{+} \\ (10), 127 [C_9H_{19}]^{+} (10), 113 [C_8H_{17}]^{+} (13.3), 99 [C_7H_{15}]^{+} (20), 85 [C_6H_{13}]^{+} \\ (10), 127 [C_9H_{19}]^{+} (10), 275 [C_2H_{31}]^{+} (1.6), 281 [C_{20}H_{41}]^{+} (1.6), 367 [C_{19}H_{39}]^{+} \\ (1.6), 323 [C_{23}H_{47}]^{+} (1.6), 295 [C_{21}H_{31}]^{+} (1.6), 281 [C_{20}H_{41}]^{+} (1.6), 267 \\ [C_{19}H_{39}]^{+} (1.6), 223 [C_{13}H_{37}]^{+} (1.6), 229 [C_{17}H_{31}]^{+} (1.6), 169 [C_{12}H_{25}]^{+} (3.3), 141 [C_{10}H_{21}]^{+} \\ (20), 85 [C_6H_{13}]^{+} (56.6), 71 [C_5H_{11}]^{+} (80), 57 [C_4H_{9}]^{+} (100). \\ 1-nonacosen 406 [M]^{+} (0.8), 323 [C_{23}H_{47}]^{+} (3.3), 265 [C_{17}H_{31}]^{+} (100). \\ 406 [M]^{+} (0.8), 323 [C_{23}H_{47}]^{+} (3.3), 265 [C_{19}H_{39}]^{+} \\ (6.6), 225 [C_{16}H_{33}]^{+} (56.6), 71 [C_{3}H_{11}]^{+} (80), 57 [C_{4}H_{9}]^{+} (100). \\ 406 [M]^{+} (2.1), 225 [C_{16}H_{33}]^{+} (3.3), 265 [C_{10}H_{33}]^{+} (100). \\ 254 [M]^{+} (2.1), 225 [C_{16}H_{313}]^{+} (13), 169 [C_{12}H_{25}]^{+} (4.3), 141 [C_{10}H_{21}]^{+} (8.6), \\ 113 [C_{8}H_{17}]^{+} (100). \\ 278 [M-H_{2}O]^{+} (2.1), 207 [C_{15}H_{27}]^{+} (1), 151 [C_{11}H_{19}]^{+} (4.3), 137 [C_{10}H_{17}]^{+} \\ (10.8), 123 [C_{9}H_{13}]^{+} (58.6), 95 [C_{7}H_{11}]^{+} (100), 81 [C_{6}H_{3}]^{+} (63), 53 \\ [C_{4}H_{3}]^{+} (86.6), 207 [C_{15}H_{27}]^{+} (100), 145 [C_{11}H_{13}]^{+} \\ (23.3), 57 [C_{4}H_{9}]^{+} (23.3). 304 [C_{29}H_{46}]^{+} (13.3), 367 [C_{29}H_{43}]^{+} (66), 315 \\ [C_{22}H_{35}O]^$		$(3.3), 141 [C_{10}H_{21}]^+ (10), 127 [C_9H_{19}]^+ (10), 113 [C_8H_{17}]^+ (13.3), 99$
octacosen 392 [M] ⁺ (0.8), 309 [C ₂₂ H ₄₃] ⁺ (3.3), 281 [C ₂₀ H ₄₁] ⁺ (6.6), 253 [C ₁₈ H ₃₇] ⁺ (6.6), 197 [C ₁₄ H ₂₉] ⁺ (6.6), 159 [C ₁₂ H ₂₃] ⁺ (10), 141 [C ₁₀ H ₂₁] ⁺ (20), 113 [C ₈ H ₁₇] ⁺ (30), 85 [C ₆ H ₁₃] ⁺ (90), 57 [C ₄ H ₉] ⁺ (100). 378 [M] ⁺ (1.6), 295 [C ₂₁ H ₄₃] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 267 [C ₁₉ H ₃₉] ⁺ (1.6), 253 [C ₁₈ H ₃₇] ⁺ (1.6), 295 [C ₂₁ H ₄₃] ⁺ (1.6), 169 [C ₁₂ H ₂₅] ⁺ (3.3), 141 [C ₁₀ H ₂₁] ⁺ (10), 127 [C ₉ H ₁₉] ⁺ (10), 113 [C ₈ H ₁₇] ⁺ (13.3), 99 [C ₇ H ₁₅] ⁺ (20), 85 [C ₆ H ₁₃] ⁺ (60),71 [C ₅ H ₁₁] ⁺ (80), 57 [C ₄ H ₉] ⁺ (100). 380 [M] ⁺ (1.6), 364 [C ₂₆ H ₃₃] ⁺ (1.6), 351 [C ₂₅ H ₅₁] ⁺ (1.6), 337 [C ₂₄ H ₄₉] ⁺ (1.6), 323 [C ₂₃ H ₄₇] ⁺ (1.6), 295 [C ₂₁ H ₄₃] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 267 [C ₁₉ H ₃₉] ⁺ (1.6), 253 [C ₁₈ H ₃₇] ⁺ (1.6), 239 [C ₁₇ H ₃₅] ⁺ (1.6), 169 [C ₁₂ H ₂₅] ⁺ (3.3), 141 [C ₁₀ H ₂₁] ⁺ (10), 127 [C ₉ H ₁₉] ⁺ (10), 113 [C ₈ H ₁₇] ⁺ (1.3), 99 [C ₇ H ₁₅] ⁺ (20), 85 [C ₆ H ₁₃] ⁺ (56.6), 71 [C ₃ H ₁₁] ⁺ (80), 57 [C ₄ H ₉] ⁺ (100). 406 [M] ⁺ (0.8), 323 [C ₂₃ H ₄₇] ⁺ (1.3), 295 [C ₂₁ H ₄₃] ⁺ (1.6), 169 [C ₁₂ H ₂₅] ⁺ (10), 141 [C ₁₀ H ₂₁] ⁺ (20), 113 [C ₈ H ₁₇] ⁺ (30), 85 [C ₆ H ₁₃] ⁺ (70), 57 [C ₄ H ₉] ⁺ (100). 254 [M] ⁺ (2.1), 225 [C ₁₆ H ₃₃] ⁺ (1.5), 169 [C ₁₂ H ₂₅] ⁺ (4.3), 141 [C ₁₀ H ₂₁] ⁺ (8.6), 113 [C ₈ H ₁₇] ⁺ (100). pytol isomer 278 [M-H ₂ O] ⁺ (2.1), 207 [C ₁₅ H ₂₇] ⁺ (1), 151 [C ₁₁ H ₁₉] ⁺ (4.3), 137 [C ₁₀ H ₁₇] ⁺ (10.8), 123 [C ₉ H ₁₅] ⁺ (58.6), 95 [C ₇ H ₁₁] ⁺ (100), 81 [C ₆ H ₉] ⁺ (63), 53 [C ₄ H ₃] ⁺ (86.9). campesterol 400 [M] ⁺ (23.3), 385 [C ₂₇ H ₄₅ O] ⁺ (13.3), 367 [C ₂₇ H ₄₃] ⁺ (6.6), 315 [C ₂₂ H ₃₅ O] ⁺ (1.3)], 281 [C ₂₁ H ₂₉] ⁺ (36.6), 207 [C ₁₅ H ₂₇] ⁺ (100), 145 [C ₁₁ H ₁₃] ⁺ (23.3), 57 [C ₄ H ₉] ⁺ (23.3). 412 [M] ⁺ (83.3), 394 [C ₂₂ H ₆] ⁺ (13.3), 369 [C ₂₆ H ₄₁ O] ⁺ (20), 351 [C ₂₆ H ₃₉] ⁺ (23.3) 330 (C ₁ C ₁ H ₂₇ O) ⁺ (36.6) 271 [C ₁₀ H ₁₂₀] ⁺ (66.6) 255 [C ₁₀ H ₁₃] ⁺ (66.6)		$[C_7H_{15}]^{T}$ (20), 85 $[C_6H_{13}]^{T}$ (60),71 $[C_5H_{11}]^{T}$ (80), 57 $[C_4H_9]^{T}$ (100).
	octacosen	$392 [M]^{+} (0.8), 309 [C_{22}H_{45}]^{+} (3.3), 281 [C_{20}H_{41}]^{+} (6.6), 253 [C_{18}H_{37}]^{+}$
$ \begin{array}{llllllllllllllllllllllllllllllllllll$		$(6.6), 197 [C_{14}H_{29}]^{+} (6.6), 169 [C_{12}H_{25}]^{+} (10), 141 [C_{10}H_{21}]^{+} (20), 113$
$\begin{array}{llllllllllllllllllllllllllllllllllll$		$[C_8H_{17}]^+$ (30), 85 $[C_6H_{13}]^+$ (90), 57 $[C_4H_9]^+$ (100).
$\begin{array}{llllllllllllllllllllllllllllllllllll$	1-heptacosene	$378 [M]^+ (1.6), 295 [C_{21}H_{43}]^+ (1.6), 281 [C_{20}H_{41}]^+ (1.6), 267 [C_{19}H_{39}]^+ (1.6),$
$ \begin{array}{llllllllllllllllllllllllllllllllllll$	-	253 $[C_{18}H_{37}]^+$ (1.6),239 $[C_{17}H_{35}]^+$ (1.6), 169 $[C_{12}H_{25}]^+$ (3.3), 141 $[C_{10}H_{21}]^+$
$\begin{array}{llllllllllllllllllllllllllllllllllll$		(10), 127 $[C_0H_{10}]^+$ (10), 113 $[C_8H_{17}]^+$ (13.3), 99 $[C_7H_{15}]^+$ (20), 85 $[C_6H_{13}]^+$
heptacosane 380 [M] ⁺ (1.6), 364 [C ₂₆ H ₅₃] ⁺ (1.6), 351 [C ₂₃ H ₅₁] ⁺ (1.6), 337 [C ₂₄ H ₄₉] ⁺ (1.6), 323 [C ₂₃ H ₄₇] ⁺ (1.6), 295[C ₂₁ H ₄₃] ⁺ (1.6), 281 [C ₂₀ H ₄₁] ⁺ (1.6), 267 [C ₁₉ H ₃₉] ⁺ (1.6), 253 [C ₁₈ H ₃₇] ⁺ (1.6), 239 [C ₁₇ H ₃₅] ⁺ (1.6), 169 [C ₁₂ H ₂₅] ⁺ (3.3), 141 [C ₁₀ H ₂₁] ⁺ (10), 127 [C ₉ H ₁₉] ⁺ (1.6), 275 [C ₄ H ₉] ⁺ (100). 1-nonacosen 406 [M] ⁺ (0.8), 323 [C ₂₃ H ₄₇] ⁺ (3.3), 295 [C ₂₁ H ₄₃] ⁺ (3.3), 267 [C ₁₉ H ₃₉] ⁺ (6.6), 225 [C ₁₆ H ₃₃] ⁺ (6.6), 197 [C ₁₄ H ₂₉] ⁺ (10), 169 [C ₁₂ H ₂₅] ⁺ (10), 141 [C ₁₀ H ₂₁] ⁺ (20), 113 [C ₈ H ₁₇] ⁺ (30), 85 [C ₆ H ₁₃] ⁺ (70), 57 [C ₄ H ₉] ⁺ (100). 254 [M] ⁺ (2.1), 225 [C ₁₆ H ₃₃] ⁺ (13), 169 [C ₁₂ H ₂₅] ⁺ (4.3), 141 [C ₁₀ H ₂₁] ⁺ (8.6), 113 [C ₈ H ₁₇] ⁺ (13), 99 [C ₇ H ₁₅] ⁺ (15.2), 85[C ₆ H ₁₃] ⁺ (4.3), 137 [C ₁₀ H ₁₇] ⁺ (10.8), 123 [C ₉ H ₁₅] ⁺ (58.6), 95 [C ₇ H ₁₁] ⁺ (100), 81 [C ₆ H ₉] ⁺ (63), 53 [C ₄ H ₅] ⁺ (86.9). campesterol 400 [M] ⁺ (23.3), 385 [C ₂₇ H ₄₅ O] ⁺ (13.3), 367 [C ₂₇ H ₄₃] ⁺ (6.6), 315 [C ₂₂ H ₃₅ O] ⁺ (13.3), 281 [C ₂₁ H ₂₉] ⁺ (36.6), 207 [C ₁₅ H ₂₇] ⁺ (100), 145 [C ₁₁ H ₁₃] ⁺ (23.3), 57 [C ₄ H ₉] ⁺ (23.3). 412 [M] ⁺ (83.3), 394 [C ₂₉ H ₄₆] ⁺ (13.3), 369 [C ₂₆ H ₄₁ O] ⁺ (20), 351 [C ₂₆ H ₃₉] ⁺ (23.3) 300 [C ₁ H ₁₂ O] ⁺ (26.6) 271 [C ₁₆ H ₂₇] ⁺ (15.2) [C ₁₆ H ₄₃] ⁺ (66.6) 271 [C ₂₆ H ₃₉] ⁺		(60).71 [C ₃ H ₁₁] ⁺ (80). 57 [C ₄ H ₆] ⁺ (100).
$\begin{array}{llllllllllllllllllllllllllllllllllll$	heptacosane	380 [M]^+ (1.6) $364 \text{ [C}_{2}\text{cH}_{\text{e}1}^+$ (1.6) $351 \text{ [C}_{2}\text{cH}_{\text{e}1}^+$ (1.6) $337 \text{ [C}_{2}\text{cH}_{\text{e}0}^+$
$ \begin{array}{l} (1:0), 325 [C_{2}M_{4}]^{-} (1:0), 225 [C_{2}M_{4}]^{-} (1:0), 201^{-} (2:0), 201^{-} (1:0), 201^{-} \\ [C_{19}M_{39}]^{+} (1.6), 253 [C_{18}M_{37}]^{+} (1.6), 239 [C_{17}M_{35}]^{+} (1.6), 169 [C_{12}H_{25}]^{+} (3.3), \\ 141 [C_{10}H_{21}]^{+} (10), 127 [C_{9}H_{19}]^{+} (10), 113 [C_{8}H_{17}]^{+} (13.3), 99 [C_{7}H_{15}]^{+} \\ (20), 85 [C_{6}H_{13}]^{+} (56.6), 71 [C_{5}H_{11}]^{+} (80), 57 [C_{4}H_{9}]^{+} (100). \\ 406 [M]^{+} (0.8), 323 [C_{23}H_{47}]^{+} (3.3), 295 [C_{21}H_{43}]^{+} (3.3), 267 [C_{19}H_{39}]^{+} \\ (6.6), 225 [C_{16}H_{33}]^{+} (6.6), 197 [C_{14}H_{29}]^{+} (10), 169 [C_{12}H_{25}]^{+} (10), 141 \\ [C_{10}H_{21}]^{+} (20), 113 [C_{8}H_{17}]^{+} (30), 85 [C_{6}H_{13}]^{+} (70), 57 [C_{4}H_{9}]^{+} (100). \\ 254 [M]^{+} (2.1), 225 [C_{16}H_{33}]^{+} (13), 169 [C_{12}H_{25}]^{+} (4.3), 141 [C_{10}H_{21}]^{+} (8.6), \\ 113 [C_{8}H_{17}]^{+} (13), 99 [C_{7}H_{15}]^{+} (15.2), 85 [C_{6}H_{13}]^{+} (4.3), 137 [C_{10}H_{17}]^{+} \\ (10.8), 123 [C_{9}H_{15}]^{+} (58.6), 95 [C_{7}H_{11}]^{+} (100), 81 [C_{6}H_{9}]^{+} (63), 53 \\ [C_{22}H_{35}O]^{+} (13.3), 281 [C_{21}H_{29}]^{+} (36.6), 207 [C_{15}H_{27}]^{+} (100), 145 [C_{11}H_{13}]^{+} \\ (23.3), 57 [C_{4}H_{9}]^{+} (23.3). \\ 412 [M]^{+} (83.3), 394 [C_{29}H_{46}]^{+} (13.3), 369 [C_{26}H_{41}O]^{+} (20), 351 [C_{26}H_{39}]^{+} \\ (23.3), 300 [C_{21}H_{20}O]^{+} (36.6), 271 [C_{19}H_{27}O]^{+} (66.6) \\ 255 [C_{19}H_{29}]^{+} (66.6) \\ \end{array}$	nepuoosuno	(1.6) 323 $[C_{22}H_{23}]^{+}$ (1.6) 295 $[C_{22}H_{23}]^{+}$ (1.6) 281 $[C_{22}H_{23}]^{+}$ (1.6) 267
$ \begin{array}{llllllllllllllllllllllllllllllllllll$		$[C_{1}-H_{2}]^{+}(1.6), 253 [C_{2}-H_{2}]^{+}(1.6), 261 [C_{2}-H_{2}]^{+}(1.6), 160 [C_{2}-H_{2}]^{+}(3.3)$
$\begin{array}{llllllllllllllllllllllllllllllllllll$		$[C_{191139}]$ (1.0), 2.5 $[C_{181137}]$ (1.0), 2.5 $[C_{171135}]$ (1.0), 105 $[C_{121125}]$ (5.5),
$\begin{array}{llllllllllllllllllllllllllllllllllll$		$[141 [C_{10} n_{21}] (10), 127 [C_{9} n_{19}] (10), 115 [C_{8} n_{17}] (15.5), 99 [C_{7} n_{15}] (20), 95 [C_{11} n_{15}] (10), 115 [C_{8} n_{17}] (10), 115 [C_{8} n_{17$
1-nonacosen406 [M] (0.8) , $323 [C_{23}H_{47}]^{-}$ (3.3) , $295 [C_{21}H_{43}]^{-}$ (3.3) , $267 [C_{19}H_{39}]^{-}$ 3-methylheptadecane (6.6) , $225 [C_{16}H_{33}]^{+}$ (6.6) , $197 [C_{14}H_{29}]^{+}$ (10) , $169 [C_{12}H_{25}]^{+}$ (10) , $141 [C_{10}H_{21}]^{+}$ $254 [M]^{+}$ (2.1) , $225 [C_{16}H_{33}]^{+}$ (13) , $169 [C_{12}H_{25}]^{+}$ (4.3) , $141 [C_{10}H_{21}]^{+}$ $86 [M]^{+}$ (2.1) , $225 [C_{16}H_{33}]^{+}$ (13) , $169 [C_{12}H_{25}]^{+}$ (4.3) , $141 [C_{10}H_{21}]^{+}$ $254 [M]^{+}$ (2.1) , $225 [C_{16}H_{33}]^{+}$ (13) , $169 [C_{12}H_{25}]^{+}$ (4.3) , $141 [C_{10}H_{21}]^{+}$ $86 [M]^{+}$ (2.1) , $225 [C_{16}H_{33}]^{+}$ (13) , $169 [C_{12}H_{25}]^{+}$ (4.3) , $141 [C_{10}H_{21}]^{+}$ (6.6) , $257 [C_{4}H_{7}]^{+}$ (13) , $99 [C_{7}H_{15}]^{+}$ (15.2) , $85 [C_{6}H_{13}]^{+}$ (4.3) , $137 [C_{10}H_{17}]^{+}$ (10.8) , $123 [C_{9}H_{15}]^{+}$ (2.1) , $207 [C_{15}H_{27}]^{+}$ (1) , $151 [C_{11}H_{19}]^{+}$ (4.3) , $53 [C_{4}H_{5}]^{+}$ $(24H_{5}]^{+}$ (86.9) . $(24H_{5}]^{+}$ (23.3) , $385 [C_{27}H_{45}O]^{+}$ (13.3) , $367 [C_{27}H_{43}]^{+}$ (6.6) , $315 [C_{22}H_{35}O]^{+}$ (23.3) , $57 [C_{4}H_{9}]^{+}$ (23.3) . (23.4) , $394 [C_{29}H_{46}]^{+}$ (13.3) , $369 [C_{26}H_{41}O]^{+}$ (20) , $351 [C_{26}H_{39}]^{+}$ (23.3) , $300 [C_{31}H_{32}O]^{+}$ (36.6) , $271 [C_{19}H_{20}O]^{+}$ (66.6) , $255 [C_{10}H_{29}H_{20}]^{+}$		$(20), 85 [C_6H_{13}] (50.0), 71 [C_5H_{11}] (80), 57 [C_4H_9] (100).$
$\begin{array}{llllllllllllllllllllllllllllllllllll$	1-nonacosen	$406 [M] (0.8), 323 [C_{23}H_{47}] (3.3), 295 [C_{21}H_{43}] (3.3), 267 [C_{19}H_{39}]$
3-methylheptadecane $\begin{bmatrix} C_{10}H_{21} \\ (20), 113 \\ \begin{bmatrix} C_8H_{17} \\ (30), 85 \\ \begin{bmatrix} C_6H_{13} \\ (43), 141 \\ \begin{bmatrix} C_{10}H_{21} \end{bmatrix}^{+} (8.6), \\ 113 \\ \begin{bmatrix} C_8H_{17} \\ (2.1), 225 \\ \begin{bmatrix} C_{16}H_{33} \end{bmatrix}^{+} (13), 169 \\ \begin{bmatrix} C_{12}H_{25} \\ (4.3), 141 \\ \begin{bmatrix} C_{10}H_{21} \end{bmatrix}^{+} (8.6), \\ 113 \\ \begin{bmatrix} C_8H_{17} \\ (10) \\ (2.1), 205 \\ \begin{bmatrix} C_{15}H_{27} \end{bmatrix}^{+} (15.2), 85 \\ \begin{bmatrix} C_{6}H_{13} \\ (4.3), 137 \\ \begin{bmatrix} C_{10}H_{11} \end{bmatrix}^{+} (60), \\ 57 \\ \begin{bmatrix} C_{4}H_{7} \end{bmatrix}^{+} (100). \\ 278 \\ \begin{bmatrix} M-H_2O \end{bmatrix}^{+} (2.1), 207 \\ \begin{bmatrix} C_{15}H_{27} \end{bmatrix}^{+} (1), 151 \\ \begin{bmatrix} C_{11}H_{19} \end{bmatrix}^{+} (4.3), 137 \\ \begin{bmatrix} C_{10}H_{17} \end{bmatrix}^{+} \\ (10.8), 123 \\ \begin{bmatrix} C_{9}H_{15} \end{bmatrix}^{+} (58.6), 95 \\ \begin{bmatrix} C_{7}H_{11} \end{bmatrix}^{+} (100), 81 \\ \begin{bmatrix} C_{6}H_{9} \end{bmatrix}^{+} (63), 53 \\ \begin{bmatrix} C_{4}H_{5} \end{bmatrix}^{+} (86.9). \\ 400 \\ \begin{bmatrix} M \end{bmatrix}^{+} (23.3), 385 \\ \begin{bmatrix} C_{27}H_{45}O \end{bmatrix}^{+} (13.3), 367 \\ \begin{bmatrix} C_{27}H_{43} \end{bmatrix}^{+} (6.6), 315 \\ \begin{bmatrix} C_{22}H_{35}O \end{bmatrix}^{+} (13.3), 281 \\ \begin{bmatrix} C_{21}H_{29} \end{bmatrix}^{+} (36.6), 207 \\ \begin{bmatrix} C_{15}H_{27} \end{bmatrix}^{+} (100), 145 \\ \begin{bmatrix} C_{11}H_{13} \end{bmatrix}^{+} \\ (23.3), 57 \\ \begin{bmatrix} C_{4}H_{9} \end{bmatrix}^{+} (23.3). \\ 412 \\ \begin{bmatrix} M \end{bmatrix}^{+} (83.3), 394 \\ \begin{bmatrix} C_{29}H_{46} \end{bmatrix}^{+} (13.3), 369 \\ \begin{bmatrix} C_{26}H_{41}O \end{bmatrix}^{+} (20), 351 \\ \begin{bmatrix} C_{26}H_{39} \end{bmatrix}^{+} \\ (23.3) \\ 300 \\ \begin{bmatrix} C_{3}H_{3}O \end{bmatrix}^{+} (36.6) \\ 271 \\ \begin{bmatrix} C_{10}H_{20}O \end{bmatrix}^{+} (66.6) \\ 255 \\ \begin{bmatrix} C_{10}H_{21} \end{bmatrix}^{+$		$(6.6), 225 [C_{16}H_{33}]^{-}(6.6), 197 [C_{14}H_{29}]^{-}(10), 169 [C_{12}H_{25}]^{-}(10), 141$
3-methylheptadecane $254 [M]^{+} (2.1), 225 [C_{16}H_{33}]^{+} (13), 169 [C_{12}H_{25}]^{+} (4.3), 141 [C_{10}H_{21}]^{+} (8.6), 113 [C_{8}H_{17}]^{+} (13), 99 [C_{7}H_{15}]^{+} (15.2), 85 [C_{6}H_{13}]^{+} (43.4), 71 [C_{5}H_{11}]^{+} (60), 57 [C_{4}H_{7}]^{+} (100).$ pytol isomer $278 [M-H_2O]^{+} (2.1), 207 [C_{15}H_{27}]^{+} (1), 151 [C_{11}H_{19}]^{+} (4.3), 137 [C_{10}H_{17}]^{+} (10.8), 123 [C_{9}H_{15}]^{+} (58.6), 95 [C_{7}H_{11}]^{+} (100), 81 [C_{6}H_{9}]^{+} (63), 53 [C_{4}H_{5}]^{+} (86.9).$ campesterol $400 [M]^{+} (23.3), 385 [C_{27}H_{45}O]^{+} (13.3), 367 [C_{27}H_{43}]^{+} (6.6), 315 [C_{22}H_{35}O]^{+} (13.3), 281 [C_{21}H_{29}]^{+} (36.6), 207 [C_{15}H_{27}]^{+} (100), 145 [C_{11}H_{13}]^{+} (23.3), 57 [C_{4}H_{9}]^{+} (23.3).$ stigmasterol $412 [M]^{+} (83.3), 394 [C_{29}H_{46}]^{+} (13.3), 369 [C_{26}H_{41}O]^{+} (20), 351 [C_{26}H_{39}]^{+} (23.3).$		$[C_{10}H_{21}]'(20), 113 [C_8H_{17}]'(30), 85 [C_6H_{13}]'(70), 57 [C_4H_9]'(100).$
$\begin{array}{llllllllllllllllllllllllllllllllllll$	3-methylheptadecane	254 [M] ⁺ (2.1), 225 [C ₁₆ H ₃₃] ⁺ (13), 169 [C ₁₂ H ₂₅] ⁺ (4.3), 141 [C ₁₀ H ₂₁] ⁺ (8.6),
57 $[C_4H_7]^*$ (100).pytol isomer278 $[M-H_2O]^+$ (2.1), 207 $[C_{15}H_{27}]^+$ (1), 151 $[C_{11}H_{19}]^+$ (4.3), 137 $[C_{10}H_{17}]^+$ (10.8), 123 $[C_9H_{15}]^+$ (58.6), 95 $[C_7H_{11}]^+$ (100), 81 $[C_6H_9]^+$ (63), 53 $[C_4H_5]^+$ (86.9).400 $[M]^+$ (23.3), 385 $[C_{27}H_{45}O]^+$ (13.3), 367 $[C_{27}H_{43}]^+$ (6.6), 315 $[C_{22}H_{35}O]^+$ (13.3), 281 $[C_{21}H_{29}]^+$ (36.6), 207 $[C_{15}H_{27}]^+$ (100), 145 $[C_{11}H_{13}]^+$ stigmasterol412 $[M]^+$ (83.3), 394 $[C_{29}H_{46}]^+$ (13.3), 369 $[C_{26}H_{41}O]^+$ (20), 351 $[C_{26}H_{39}]^+$ (23.3), 300 $[C_{21}H_{22}O]^+$ (36.6), 271 $[C_{10}H_{22}O]^+$ (66.6), 255 $[C_{10}H_{21}]^+$ (66.6)		113 $[C_8H_{17}]^+$ (13), 99 $[C_7H_{15}]^+$ (15.2), 85 $[C_6H_{13}]^+$ (43.4), 71 $[C_5H_{11}]^+$ (60),
pytol isomer 278 $[M-H_2O]^+$ (2.1), 207 $[C_{15}H_{27}]^+$ (1), 151 $[C_{11}H_{19}]^+$ (4.3), 137 $[C_{10}H_{17}]^+$ (10.8), 123 $[C_9H_{15}]^+$ (58.6), 95 $[C_7H_{11}]^+$ (100), 81 $[C_6H_9]^+$ (63), 53 $[C_4H_5]^+$ (86.9). 400 $[M]^+$ (23.3), 385 $[C_{27}H_{45}O]^+$ (13.3), 367 $[C_{27}H_{43}]^+$ (6.6), 315 $[C_{22}H_{35}O]^+$ (13.3), 281 $[C_{21}H_{29}]^+$ (36.6), 207 $[C_{15}H_{27}]^+$ (100), 145 $[C_{11}H_{13}]^+$ (23.3), 57 $[C_4H_9]^+$ (23.3). stigmasterol 412 $[M]^+$ (83.3), 394 $[C_{29}H_{46}]^+$ (13.3), 369 $[C_{26}H_{41}O]^+$ (20), 351 $[C_{26}H_{39}]^+$ (23.3) 300 $[C_{21}H_{22}O]^+$ (36.6) 271 $[C_{10}H_{27}O]^+$ (66.6) 255 $[C_{10}H_{21}]^+$ (66.6)		57 $[C_4H_7]^+$ (100).
$ \begin{array}{l} \text{(10.8), 123 } [C_9H_{15}]^+ (58.6), 95 [C_7H_{11}]^+ (100), 81 [C_6H_9]^+ (63), 53 \\ [C_4H_5]^+ (86.9). \\ 400 \ [M]^+ \ (23.3), 385 \ [C_{27}H_{45}O]^+ (13.3), 367 \ [C_{27}H_{43}]^+ (6.6), 315 \\ [C_{22}H_{35}O]^+ \ (13.3), 281 \ [C_{21}H_{29}]^+ (36.6), 207 \ [C_{15}H_{27}]^+ \ (100), 145 \ [C_{11}H_{13}]^+ \\ (23.3), 57 \ [C_4H_9]^+ \ (23.3). \\ 412 \ [M]^+ \ (83.3), 394 \ [C_{29}H_{46}]^+ \ (13.3), 369 \ [C_{26}H_{41}O]^+ \ (20), 351 \ [C_{26}H_{39}]^+ \\ (23.3), 300 \ [C_{21}H_{22}O]^+ \ (36.6), 271 \ [C_{16}H_{27}O]^+ \ (66.6), 255 \ [C_{16}H_{27}]^+ \ (66.6) \\ \end{array} $	pytol isomer	278 $[M-H_2O]^+$ (2.1), 207 $[C_{15}H_{27}]^+$ (1), 151 $[C_{11}H_{19}]^+$ (4.3), 137 $[C_{10}H_{17}]^+$
campesterol $\begin{bmatrix} C_{4}H_{5} \end{bmatrix}^{+} (86.9).$ 400 [M] ⁺ (23.3), 385 [C ₂₇ H ₄₅ O] ⁺ (13.3), 367 [C ₂₇ H ₄₃] ⁺ (6.6), 315 [C ₂₂ H ₃₅ O] ⁺ (13.3), 281 [C ₂₁ H ₂₉] ⁺ (36.6), 207 [C ₁₅ H ₂₇] ⁺ (100), 145 [C ₁₁ H ₁₃] ⁺ (23.3), 57 [C ₄ H ₉] ⁺ (23.3). stigmasterol 412 [M] ⁺ (83.3), 394 [C ₂₉ H ₄₆] ⁺ (13.3), 369 [C ₂₆ H ₄₁ O] ⁺ (20), 351 [C ₂₆ H ₃₉] ⁺ (23.3), 300 [C ₂₁ H ₂₂ O] ⁺ (36.6), 271 [C ₁₆ H ₂₇ O] ⁺ (66.6), 255 [C ₁₆ H ₂₇] ⁺ (66.6)	Den en	$(10.8), 123 [C_9H_{15}]^+ (58.6), 95 [C_7H_{11}]^+ (100), 81 [C_6H_9]^+ (63), 53$
campesterol 400 $[M]^+$ (23.3), 385 $[C_{27}H_{45}O]^+$ (13.3), 367 $[C_{27}H_{43}]^+$ (6.6), 315 $[C_{22}H_{35}O]^+$ (13.3), 281 $[C_{21}H_{29}]^+$ (36.6), 207 $[C_{15}H_{27}]^+$ (100), 145 $[C_{11}H_{13}]^+$ (23.3), 57 $[C_4H_9]^+$ (23.3). stigmasterol 412 $[M]^+$ (83.3), 394 $[C_{29}H_{46}]^+$ (13.3), 369 $[C_{26}H_{41}O]^+$ (20), 351 $[C_{26}H_{39}]^+$ (23.3) 300 $[C_{21}H_{22}O]^+$ (36.6) 271 $[C_{10}H_{22}O]^+$ (66.6) 255 $[C_{10}H_{21}]^+$ (66.6)		$[C_4H_5]^+(86.9).$
stigmasterol $[C_{22}H_{35}O]^{+}(13.3), 281 [C_{21}H_{29}]^{+}(36.6), 207 [C_{15}H_{27}]^{+}(100), 145 [C_{11}H_{13}]^{+}(23.3), 57 [C_{4}H_{9}]^{+}(23.3).$ 412 [M] ⁺ (83.3), 394 [C ₂₉ H ₄₆] ⁺ (13.3), 369 [C ₂₆ H ₄₁ O] ⁺ (20), 351 [C ₂₆ H ₃₉] ⁺ (23.3), 300 [C ₂₁ H ₂₂ O] ⁺ (36.6), 271 [C ₁₉ H ₂₂ O] ⁺ (66.6), 255 [C ₁₉ H ₂₂] ⁺ (66.6)	campesterol	400 $[M]^+$ (23.3), 385 $[C_{27}H_{45}O]^+$ (13.3), 367 $[C_{27}H_{43}]^+$ (6.6), 315
stigmasterol (23.3), 57 $[C_4H_9]^+$ (23.3). 412 $[M]^+$ (83.3), 394 $[C_{29}H_{46}]^+$ (13.3), 369 $[C_{26}H_{41}O]^+$ (20), 351 $[C_{26}H_{39}]^+$ (23.3) 300 $[C_{24}H_{23}O]^+$ (36.6) 271 $[C_{16}H_{27}O]^+$ (66.6) 255 $[C_{16}H_{27}]^+$ (66.6)	-	$[C_{22}H_{35}O]^+$ (13.3), 281 $[C_{21}H_{29}]^+$ (36.6), 207 $[C_{15}H_{27}]^+$ (100), 145 $[C_{11}H_{13}]^+$
stigmasterol 412 $[M]^+$ (83.3), 394 $[C_{29}H_{46}]^+$ (13.3), 369 $[C_{26}H_{41}O]^+$ (20), 351 $[C_{26}H_{39}]^+$ (23.3), 300 $[C_{24}H_{32}O]^+$ (36.6), 271 $[C_{19}H_{32}O]^+$ (66.6), 255 $[C_{19}H_{32}]^+$ (66.6)		$(23.3), 57 [C_4H_0]^+ (23.3).$
(23.3) 300 [C ₂₁ H ₂₂ O] ⁺ (36.6) 271 [C ₁₀ H ₂₂ O] ⁺ (66.6) 255 [C ₁₀ H ₂₂] ⁺ (66.6)	stigmasterol	412 $[M]^+$ (83.3), 394 $[C_{29}H_{46}]^+$ (13.3), 369 $[C_{26}H_{41}O]^+$ (20), 351 $[C_{26}H_{20}]^+$
(23.5), 500 [0211320] (50.0), 271 [01912/0] (00.0), 255 [01917/1] (00.0).		$(23.3), 300 [C_{21}H_{32}O]^+ (36.6), 271 [C_{19}H_{27}O]^+ (66.6), 255 [C_{19}H_{27}]^+ (66.6).$

Howayda I. El-Nabawy, et al.

ationnant 5 an 2 al	159 $[C_{12}H_{15}]^+$ (56.6), 83 $[C_6H_{11}]^+$ (80), 55 $[C_4H_7]^+$ (100). 414 $[M_1^+$ (60) 381 $[C_{22}H_{12}]^+$ (23.3) 329 $[C_{22}H_{12}O]^+$ (46.6) 281 $[C_{21}H_{22}]^+$
stigmast-3-en-3-or	$(50), 207 [C_{15}H_{27}]^+ (100), 145 [C_{11}H_{13}]^+ (53.3), 55 [C_{4}H_{7}]^+ (33.3).$
olean-12-en-3-ol (β-	426 $[M]^+$ (3.3), 411 $[C_{29}H_{47}O]^+$ (1.6), 218 $[C_{16}H_{26}]^+$ (100), 203 $[C_{15}H_{23}]^+$
amyrine)	$(63.3), 203 [C_{15}H_{23}]^{+} (63.3), 135 [C_{10}H_{15}]^{+} (10), 119 [C_{9}H_{11}]^{+} (10), 95$
	$[C_6H_7O]'$ (13.3).
urs-12-en-3-ol (a-	426 $[M]^+$ (10), 411 $[C_{29}H_{47}O]^+$ (6.6), 393 $[C_{29}H_{45}]^+$ (1.6), 218 $[C_{16}H_{26}]^+$
amyrine)	$(100), 203 [C_{15}H_{23}]^{+} (26.6), 189 [C_{14}H_{21}]^{+} (26.6), 175 [C_{13}H_{19}]^{-} (13.3), 135$
	$[C_{10}H_{15}]'(30), 121 [C_{9}H_{15}]'(26.6), 95 [C_{6}H_{7}O]'(26.6).$
12-oleanen-3-yl acetate	468 $[M]^{+}(3.3)$, 393 $[C_{29}H_{45}]^{+}(1.6)$, 218 $[C_{16}H_{26}]^{+}(100)$, 189 $[C_{14}H_{21}]^{+}$
	$(16.6), 135 [C_{10}H_{15}]^{\circ} (10), 119 [C_{9}H_{11}]^{\circ} (10), 95 [C_{6}H_{7}O]^{\circ} (10).$
olean-18-en-3-ol	$426 [M]^{+}(1.6), 393 [C_{29}H_{45}]^{+}(3.3), 204 [C_{15}H_{24}]^{+}(90), 189 [C_{14}H_{21}]^{+}(100),$
	$[135 [C_{10}H_{15}]^{\circ} (26.6), 119 [C_{9}H_{11}]^{\circ} (26.6), 95 [C_{6}H_{7}O]^{\circ} (40), 55 [C_{4}H_{7}]$
	(23.3).
10-octadecenoic	$296 [M] (2.1), 264 [C_{18}H_{32}O_2] (13), 222 [C_{15}H_{26}O] (6.5), 180 [C_{12}H_{20}O]$
acid, methyl ester	$(8.6), 111 [C_8H_{15}]' (26), 69 [C_5H_9]' (73.9), 55 [C_4H_7]' (100).$
moretenol	426 $[M]^{+}$ (10), 327 $[C_{24}H_{39}]^{+}$ (1.6),207 $[C_{14}H_{23}O]^{+}$ (100), 189 $[C_{14}H_{21}]^{+}$
	(50), $175 [C_{13}H_{19}]^{+}$ (13.3), $135 [C_{10}H_{15}]^{+}$ (26.6), $121 [C_{9}H_{13}]^{+}$ (26.6), 55
	$[C_4H_7]^{-}(13.3).$
(23S)-ethyl cholest-5-	414 [M] ⁺ (70), 371 [C ₂₆ H ₄₃ O] ⁺ (1.6), 273 [C ₁₉ H ₂₉ O] ⁺ (26.6), 213 [C ₁₆ H ₂₁] ⁺
en-3β-ol	$(46.6), 145 [C_{11}H_{13}]' (50), 57 [C_4H_9]' (100).$
triacontane	422 $[M]^{+}(2.1), 379 [C_{27}H_{55}]^{+}(1), 351 [C_{25}H_{51}]^{+}(1), 309 [C_{22}H_{45}]^{+}(1), 281$
	$[C_{20}H_{41}]'(1), 253 [C_{18}H_{37}]'(1), 197 [C_{14}H_{29}]'(2.1), 169 [C_{12}H_{25}]'(4.3),$
	141 $[C_{10}H_{21}]^{+}$ (6.5), 113 $[C_{8}H_{17}]^{+}$ (13), 99 $[C_{7}H_{15}]^{+}$ (17.3), 85 $[C_{6}H_{13}]^{+}$
	$(54),71 [C_5H_{11}]^{T} (78.2), 57 [C_4H_9]^{T} (100).$
hentriacontane	436 $[M]^+$ (2.1), 309 $[C_{22}H_{45}]^+$ (1), 281 $[C_{20}H_{41}]^+$ (4.3), 253 $[C_{18}H_{37}]^+$ (2.1),
	113 $[C_8H_{17}]^+$ (13), 85 $[C_6H_{13}]^+$ (54),71 $[C_5H_{11}]^+$ (78.2), 57 $[C_4H_9]^+$ (100).
dotriacontane	450 $[M]^+(1)$, 295 $[C_{21}H_{43}]^+(1)$, 281 $[C_{20}H_{41}]^+(8.6)$, 253 $[C_{18}H_{37}]^+(4.3)$,
	225 $[C_{16}H_{33}]^{+}$ (2.1), 169 $[C_{12}H_{25}]^{+}$ (4.3), 85 $[C_{6}H_{13}]^{+}$ (54),71 $[C_{5}H_{11}]^{+}$
	$(78.2), 57 [C_4H_9]^+ (100).$
tritriacontane	$464 \ [M]^{+} (2.1),295 \ [C_{21}H_{43}]^{+} (1),281 \ [C_{20}H_{41}]^{+} (13),253 \ [C_{18}H_{37}]^{+} (6.5),$
	85 $[C_6H_{13}]^+$ (54),71 $[C_5H_{11}]^+$ (78.2), 57 $[C_4H_9]^+$ (100).

Mass spectra of some compounds



Fig. 1: Mass spectrum of β -amyrine.





Refernces

- Boulos, L.; flora of Egypt, Al hadara puplishing Cairo, Egypt, 2002, 275-267.
- Balaboul, B. A.; Ahmed, A. A.; Otsuka, H.,
 Sequiterpene lactones and glucosides from Urospermum picroides,
 Phytochemistry, 1997, 45, 369-373.
- Cai, Y. Z.; Sun, M.; Xing,J.; Luo, Q.; Corke, H., Structure –radical scavenging activity relationships of phenolic compunds from traditional Chines medicinal plants, Life sciences, 2006, 78, 2872-2888.
- Denizot, F.; Lang, R.; Immunol, J., Methods, 1986, 22, 271-277.
- Fragopoulou, E.; Detopoulou, P.; Nomikos, T.; Pliakis, E.; Panagiotakos, D. B.; Antonopoulou, S., mediterrean wild plants reduce postprandial platelet aggregation in patients with metabolic syndrome, Metabolism Journal, 2012, 61, 325-334.
- Giner, R. M.; Cuellar, M. J.; Recio, M. C.; Manez, S.; Rios, J. L., chemical constituents of *urospermum picroides*, Z.Naturforsch, **1992**, 47, 531-534.
- Islam, M.; Al-Amin, M.; Siddiqi, M. M. A.; Akter, S.; Haque, M. M.; Sultan, N.; Chowdhury, A.M.S., Isolation of quercetin-3-o-beta-d-glucopyranoside from the leaves of *azadirachtaIndica*and antimicrobial and cytotoxic screening of the crude Extracts, **2012**, 60, 11-14.

- Lissi, E.; Modak, B.; Torres, R.; Escobar, J.; Urza, A., Free Radical Res., **1999**, 30, 471-477.
- Mikania F. W. Schmidt, Samml. Phys. Okon. Aufs., 1795, 1,27.
- Stylianakis, Kolocouris, A.; Kolocouris, N.; Fytas, G.; Foscolos, G. B.; Padalko, E.; Neyts, J.; Clerq, D.; Bioorg, E., Med. Chem. Lett., 2003, 1699-1703.
- Strzelecka, M.; Bzowska, M.; Koziel, J.; Szuba, B.; Dubiel, O.; Nunez, R.; Heinrich,M.; Bereta, J., Antiinflammatory effects of extracts from some traditional Mediterranean diet plants, Journal of physiology and pharmacology, 2005, 56, 139-156.
- Stalinska, K.; Guzdek, A.; Rokicki, M.; Koj, A., Transcription factors as targets of the anti-inflammatory treatment. A cell culture study with extracts from some Mediterranean diet plants, Journal of physiology and pharmacology, 2005, 56, 157-169.
- Tuczaj, T.; Koncic, M., Z.; Milicevic, T.; Dolina, K.; Pandza, M., wild vegetables mixes sold in the markets of Dalmatia (southern Croatia), Journal of ethnobiology and ethnomedicine, 2013, 9, 2.
- Tackholm, V.; Student's flora of Egypt, Cairo university press, **1974**, 888.

التقييم الفيتوكيميائي والبيولوجي لنبات السليس

أدى فصل خلاصات الأجزاء الهوائية لنبات "يوروسبرمم بكرويدز" إلى الحصول على اثنين من نوع سسكويتربين لاكتون، وهما يوروسبرمال A (۱) و ٨، ١٥- دايهيدروكسي جيرماكرا-۱(١٠)، ٤-دايين-٦، ١٢-أولايد- ١٤-آل (٢)، واثنين من نوع فلافونويد جلايكوزايد، وهما كوارستين-٣-٥-بيتا-D-جلوكوبيرانوزايد (٣) وكامفيرول-٣-٥-بيتا-D-جلوكوبيرانوزايد (٤). تحليل عينة من خلاصة الإيثر البترولي للأجزاء الهوائية بتقنية كروماتوجرافيا الغاز المقترن بمطياف الكتلة نتج عنها تعريف ألفا أميرين، وبيتا أميرين، ١٢-أوليانين-٣-ويل أسيتات، ١٨-أولاينين-٣-بيتا-أول، موريتينول، الكتلة نتج عنها تعريف ألفا أميرين، وبيتا أميرين، ١٢-أوليانين-٣-ويل أسيتات، ١٨-أوليانين-٣-بيتا-أول، موريتينول، المنابعة.

وقد وجد أن خلاصات بيوتانول للبذور، وإيثايل أسيتات للبذور، وإيثايل أسيتات للأجزاء الهوائية، وكلوريد الميثيلين للأجزاء الهوائية تثبط الشقوق الحرة بنسب %86.9، %85.3، %86.1، %82.4 على الترتيب، مقارنة بتثبيط %88.1 بحمض الأسكوربك. بينما أدت الجزئية الغنية بالسسكويتربينات ١ و٢ والجزئية الغنية بالفلافونويدات ٣ و٤ إلى نسب تثبيط 84.7% و%83.2 على اترتيب، مما يوضح التأثير التناغمي المحتمل للمكونات الأخرى.

وجد أن الفعالية ضد الميكروبات (متمثلة في البكتريا موجبة الجرام استافيلوكوكس أوريوس والبكتريا سالبة الجرام اشرشيا كولاي واخميرة كانديدا ألبيكانز) لخلاصة إيثايل أسيتات للأجزاء الهوائية والجزئية الغنية بالفلافونويدات ٣ و ٤ هي (68.2%، 68.2% و69.2%) للأولى و(54.5%، 50.0% و50.0%) للأخيرة. كما وجد أن فعالية خلاصة بيوتانول للبذور وخلاصة إيثايل أسيتات للبذور تجاه البكتريا موجبة الجرام استافيلوكوكس أوريوس هي 68.2% و50.5% على الترتيب. كما وجد أنه لخلاصة البيوتانول للبذور سمومية قوية جداً لخلايا سرطان الثدى 7-MCF (قيمتها 70.3%) وسمومية قوية لخلايا سرطان الكبد 2-96 (قيمتها 18.5%)، ولخلاصة إيثايل أسيتات للأجزاء الهوائية والجزئية الغنية بالفلافونويدات ٣ و سرطان الثدى 7-MCF (قيمتها 18.5%) وسمومية قوية خداً لخلايا سرطان الثدى 8-10% (مح.2%) وسمومية قوية سرطان الثدى 7-MCF (قيمتها 18.5%)، والخلاصة إيثايل أسيتات للأجزاء الهوائية سمومية قوية جداً لخلايا سرطان الثدى 8-10% (مح.2%)

