Synthesis of Alkyl-Glycerolipids Standards for Gas Chromatography Analysis: Application for Chimera and Shark Liver Oils

Michelle Pinault, Cyrille Guimaraes, Hélène Couthon, Jérôme Thibonnet, Delphine Fontaine, Aurélie Chantôme, Stephan Chevalier, Pierre Besson, Paul-Alain Jaffrès, and Christophe Vandier

Inserm, UMR1069, Université de Tours, 37000 Tours, France; Michelle.pinault@univ-tours.fr (M.P.); cyrille.guimaraes@gmail.com (C.G.); delphine.fontaine@etu.univ-tours.fr (D.F.); aurelie.chantome@univ-tours.fr (A.C.); stephan.chevalier@univ-tours.fr (S.C.); pierre.besson@univ-tours.fr (P.B.)

CEMCA, CNRS UMR6521, Université de Brest, IBSAM, 6 Av V. Le Gorgeu, 29238 Brest, France; helene.Couthon@univ-brest.fr (H.C.); Paul-Alain.Jaffres@univ-brest.fr (P.-A.J.)

Equipe SIMBA, Synthèse et Isolement de Molécules BioActives, EA 7502, Université de Tours, 37000 Tours, France; jerome.thibonnet@univ-tours.fr

Faculté de Sciences et Techniques, Université de Tours, 37000 Tours, France

Faculté de Pharmacie, Université de Tours, 37000 Tours, France

Correspondence: christophe.vandier@univ-tours.fr; Tel.: +(33)-247366024; Fax: +(33)-247366226

† These authors contributed equally to this work.

Received: 18 January 2018; Accepted: 20 March 2018; Published: 23 March 2018

Abstract: Natural O-alkyl-glycerolipids, also known as alkyl-ether-lipids (AEL), feature a long fatty alkyl chain linked to the glycerol unit by an ether bond. AEL are ubiquitously found in different tissues but are abundant in shark liver oil, breast milk, red blood cells, blood plasma, and bone marrow. Only a few AEL are commercially available, while many others with saturated or mono-unsaturated alkyl chains of variable length are not available. These compounds are, however, necessary as standards for analytical methods. Here, we investigated different reported procedures and we adapted some of them to prepare a series of 1-O-alkyl-glycerols featuring mainly saturated alkyl chains of various lengths (14:0, 16:0, 17:0, 19:0, 20:0, 22:0) and two monounsaturated chains (16:1, 18:1). All of these standards were fully characterized by NMR and GC-MS. Finally, we used these standards to identify the AEL subtypes in shark and chimera liver oils. The distribution of the identified AEL were: 14:0 (20–24%), 16:0 (42–54%) and 18:1 (6–16%) and, to a lesser extent, (0.2–2%) for each of the following: 16:1, 17:0, 18:0, and 20:0. These standards open the possibilities to identify AEL subtypes in tumours and compare their composition to those of non-tumour tissues.

Keywords: ether-lipids; alkyl-glycerolipids; standards for gas chromatography; fish oil

1. Introduction

Natural O-alkyl-glycerolipids, also known as alkyl-ether-lipids (AEL), have a long fatty alkyl chain linked to the glycerol unit by an ether bond [1]. The most prevalent glycerol-backboned AEL found in nature consist of an O-alkyl or O-alk-1′-enyl group attached to glycerol at position sn-1. In cells, AEL can be triglycerides (diacyl-glyceryl-ether lipids, DAGE) which possess two fatty acyl moieties at positions sn-2 and sn-3, or phospholipids (1-O-Alkyl-2-Acyl-sn-glycerylphospholipids, AAGPL) with one fatty acyl chain at the sn-2 position and one phosphodiester group (e.g., phosphocholine, phosphoethanolamine). The common feature of these two series of compounds is the 1-O-alkyl-glyceryl ether lipid (AGEL), as shown in Figure 1.
Natural AEL were found to stimulate the immune system [2] and hematopoiesis [3]. They also have anti-tumor properties [4], reduce the side effects induced by radiotherapy [5], and lead to a reduction in the growth of tumor cells and metastases [6]. Interestingly, ether-lipids have been also considered for potential use in the diagnosis of prostate cancer [7].

Some biological effects of these AEL can be explained by an interaction with their receptors, such as the platelet-activating factor (1-O-alkyl-2-acetyl-sn-glycero-3-phosphocholine) [8,9], or following their integration in the phospholipids of cell membranes [10], and by a modification of the activity of Protein Kinase C (PKC) [11–13] which has a high activity in breast tumors [14]. The integration of AEL into the membrane results in the formation of DAG ether-lipid analogs, alkyl-acyl-glycerols, which inhibit PKC activation unlike the “classical” ester-linked, PKC-activating diacyl-glycerol DAG [12,15]. In addition to a decrease in PKC activity, it has been shown that these lipids inhibit the activity of other kinases such as phosphatidylinositol-3-kinase [16] and MAP kinases [17], but also Na⁺/K⁺ ATPase [18]. AEL derivative edelfosine (1-O-octadecyl-2-O-methyl-sn-glycero-3-phosphocholine) was found to inhibit the activities of Na⁺/H⁺ exchanger [19] and K⁺ channel [20], as well as the transport of nutrients [21]. Thus, all of these effects can participate in the anti-proliferative effects of AEL. The replacement of the phosphocholine moiety by an aminoglycoside group led to a large series of compounds exhibiting anticancer properties [22,23]. Interestingly, with the replacement of the phosphocholine moiety of edelfosine by a saccharide unit to design glyco-AEL, we identified less toxic compounds whereas the modulation of ion channels was preserved [24–27]. Among disaccharide-AEL, [28] we identified ohmline (1-O-hexadecyl-2-O-methyl-rac-glycero-3-lactose) as a lead synthetic glycol-AEL that can prevent metastasis development by modifying the activity of ion channel complexes (KCa-Ca2⁺) located in cholesterol-rich nano-domain and regulating calcium signaling and migration of cancer cells [29–31]. Some aspects of the mechanism of action of ohmline were recently reported, showing that ohmline fluidified the plasma membrane, especially if this plasma membrane had a high level of cholesterol [32]. Altogether, these results pointed out the great therapeutic potential of synthetic ether lipids and contribute to questions about the role of natural ether lipids. AEL are abundant in many tissues, and especially in shark liver oil, breast milk, red blood cells, blood plasma, and bone marrow [33–35]. Among human tissues, bone marrow contained the highest percentage of alkyl-glycerols in both triglycerides and phospholipids fractions [35]. In mammals, there is a higher prevalence of alkyl-glycerols in phospholipid fractions compared to neutral lipid (triglycerides) fractions [36]. Interestingly, tumors have been found to contain larger amounts of AEL than normal tissues [36], but the physiological roles of these natural AEL have not been elucidated so far. The clinical significance of their accumulation in tumors remains to be determined. Furthermore, no quantitative analysis has been carried out in tissues to precisely determine the proportion of AEL molecular species based on their alkyl chain length and unsaturation level. If some AELs are commercially available (e.g., 1-O-hexadecyl-sn-glycerol, also known as chimyl alcohol, and 1-O-octadecyl-rac-glycerol also known as batyl alcohol), others with saturated alkyl chain of variable length or monounsaturated alkyl...
chains are, unfortunately, not commercially available. These compounds are necessary as standards for analytical methods. This led us to investigate reported synthesis procedures and to adapt some of them to prepare a series of 1-O-alkyl-glycerols with saturated chain length (14:0, 16:0, 17:0, 19:0, 20:0, 22:0) and two monounsaturated chains (16:1, 18:1). All of these standards were fully characterized by NMR and GC-MS. Finally, we used these standards to precisely characterize alkyl-glycerols subtypes in shark and chimera liver oils.

2. Results and Discussion

2.1. Alkyl-Glycerolipids Chemical Synthesis

Since our goal was to have alkyl-glycerol standards for analytical purpose, the chirality at the sn-2 position was not an issue since the targeted compounds (4a–i; Figure 2) possess only one asymmetric carbon and, therefore, on non-chiral support, both enantiomers will possess similar physico-chemical properties including retention time. As a consequence, the syntheses of the standards were achieved in racemic series. Solketal 1 is the common commercial precursor to prepare O-alkyl glycerol 4 since the free alcohol function can be deprotonated and then engaged in an alkylation reaction with a lipid alcohol activated with a mesyl functional group or directly with a halogenoalkyl. Prinz et al. [37] used KOH to deprotonate solketal in benzene and then myristoylmethanesulfonate (C14:0) or palmitoylmethanesulfonate (C16:0) was added. After alcohol deprotection with HCl in methanol, the alkylglycerol 4a (C14:0) and 4b (C16:0) were isolated in 77–85% yield (two steps). Aranda et al. [38] only changed the solvent (DMSO) to prepare compound 4a (C14:0) in 80–85% yield. Huang et al. [39] used NaH in toluene for deprotonation of solketal and a subsequent alkylation was achieved with bromoalkane (1-Bromotetradecane, 1-Bromohexadecane or 1-Bromoocotadecane), thus producing compounds 4a–b (yields were not reported). Another approach that retained our attention consisted of using lipid alcohol and racemic solketal 2 activated with a tosyl group as the leaving group. The use of this activated solketal was previously reported by Bhatia et al. [40] to produce 1-O-hexadecyl-sn-glycerol in 65% yield (including the acetal deprotection). We prepared the tosylate 2 from solketal 1 following a reported procedure [41]; it is worth noting that compound 2 is also commercially available (analytical data are available in the Supporting Material, SI1.7, 1.8). Then, we used the method of Bhatia et al. to introduce the alkylether moiety (Figure 2). First, the alkyl alcoholate was prepared by deprotonation of the alcohol with NaH and then this alcoholate reacted with the tosylated solketal 2 [42]. After work-up, the expected compounds 3a–i were isolated in 60–96% yield. Then the deprotection of the acetal function was achieved in acidic conditions (HCl 2N) in methanol at reflux and the final compounds 4a–i were isolated in 92–98% yield. If most of these compounds were reported and prepared by other methods, it must be noted that to the best of our knowledge, compounds 3/4c, 3/4d, 3/4g were not reported before.

For the analysis of the 1-O-alkyl glycerol by GC, protection of the alcohol function must be achieved. For this derivatization step, we have protected the alcohol function with acetate groups. For this purpose, two methods were employed. Method A consisted to use acetic anhydride in pyridine at 60 °C [43] and method B used acetyl chloride without solvent at 140 °C (Figure 2) [44]. Method A used mild conditions that should reduce degradation but the work-up (treatment with acid and basic solutions) induced sample losses. Method B is much faster and was efficient with a simplified work-up. However, this method requires heating at 140 °C that can induce, depending on the structure of the lipid chain, some degradation. In this series of compounds, no degradation was observed including for the mono-unsaturated compound 5c and 5f. The final compounds 5a–i were fully characterized and isolated in good yield (92–98%).
We found that both oils contain mainly alkyl-glycerols with C14:0, C16:0, and C18:1 alkyl chains and low amounts of C16:1 and C18:0 alkyl chain. Interestingly, C17:0 and C20:0 were found in Chimaera monstrosa and shark liver oils. Note, that we did not find C19:0 or C22:0 alkyl chains.

2.2. Liver Oils Alkyl-Glycerolipids Analysis

In order to precisely identify the alkyl-glycerol subtypes in shark and chimera liver oils we first extracted the total lipids of chimera and shark liver oil and derived alkyl-glycerols to alkyl-glycerol diacetates to be analyzed using gas chromatography as described in Sections 3.2 and 3.4.

Figure 3 shows an example of chromatograph of alkyl-glycerols diacetate obtained from chimera (Chimaera monstrosa) liver oil.

Using the alkyl-glycerols standards, we analysed the alkyl-glycerol subtypes in shark and chimera liver oils (Figure 4). We found that both oils contain mainly alkyl-glycerols with C14:0, C16:0, and C18:1 alkyl chains and low amounts of C16:1 and C18:0 alkyl chains. Interestingly, C17:0 and C20:0 were found in Chimaera monstrosa and shark liver oils. Note, that we did not find C19:0 or C22:0 alkyl chains.

Figure 2. Synthesis of alkyl-glycerolipids 4a–i and derivatization as diacetyl esters 5a–i. (i) TsCl, Pyridine, 1 h, 0 °C then 24 h, rt; (ii) preparation of the alcoholate: NaH, ROH, dimethoxyethane, 1 h, then addition of 2, 100 °C then 48 h, reflux; (iii) HCl 2N, Ethanol, 4 h, reflux. Method A (iv): acetic anhydride, pyridine, 60 °C, 30 min. and then 1 night at 20 °C; Method B (v): acetyl chloride, 5 min. 140 °C.
After derivatization as acetate, we optimized the temperature program for the separation and identification of alkyl-glycerolipids by GC-MS (see Supplementary Materials SI2.1, SI2.2). This method confirmed the results obtained using GC-FID.

![Figure 4](image-url)

**Figure 4.** Comparative analysis of alkyl-glycerols of chimera (Chimaera monstrosa) and shark liver (a mixture of two species: Centrophorus squamosus and Somniosus microcephalus). Bars represent mean value and error bars represent SD, n = 6. Neither C19:0 nor C22:0 alkyl chains were detected.

We found the presence of 14:0 (20–24%), 16:0 (42–54%), and 18:1 (6–16%) alkyl-glycerols and, to a lesser extent (0.2–2%), 16:1, 17:0, 18:0 and 20:0 alkyl-glycerols. It was generally reported that liver oil of certain shark species contain high levels of alkyl-glycerolipids as a mixture of subtype that varied by length and unsaturation of the alkyl chain with almost exclusively saturated and monounsaturated, the main alkyl moieties being the C16 and C18 chains in both lipid classes [45,46]. Interestingly, a high level of 18:1 was also found in the composition of alkyl chains in alkyl-glycerolipids from Greenland shark (Centrophorus squamosus) liver oil [47]. Thus, our standards confirm the presence of these alkyl-glycerolipids in shark and chimera liver oils.

### 3. Materials and Methods

#### 3.1. Chemical Synthesis

Chemicals and solvents ACS grade or higher, commercially available, were used without further purification (Carlo Erba, Val de Reuil, France; and Sigma-Aldrich, Saint-Quentin Fallavier, France). NMR spectra (Bruker, Wissembourg, France) were recorded on a Bruker DPX Avance 300 spectrometer (300 MHz for $^1$H, 75 MHz for $^{13}$C), Bruker DRX-500 (500 MHz for $^1$H, 125 MHz for $^{13}$C), and Bruker Ascend-600 AVANCE III HD equipped with TCI cryoprobe (600 MHz for $^1$H, 150 MHz for $^{13}$C). CDCl$_3$ was used as the solvent; chemical shifts are expressed in ppm relative to TMS as an internal standard. Purification of crude products was carried out using Puriflas® 430 (Interchim, Montluçon, France) equipped with an evaporative light-scattering detector (ELSD). Preparative thin-layer chromatography (TLC, LK5) separation was carried out using 20 × 20 cm, glass-backed, 1 mm or higher thickness silica gel Merck F$_{254}$ plates (VWR, Fontenay-sous-Bois, France). Control TLC was performed using Merck F$_{254}$ aluminium-backed silica gel sheets (VWR). Lipid compounds on TLC plates/sheets were revealed by 2′,7′-dichlorofluorescein.
HMRS analysis was performed on the HRMS CNRS ICOA platform at the University of Orleans.

### 3.2. Analytical Chemistry

Chemicals and solvents, ACS grade or higher, commercially available, were used without further purification (Carlo Erba).

After the extraction of total lipids by the Folch method [48] ester bonds were reduced using the strong reducing agent Vitride® (sodium hydride and aluminum) (0.5 mL Red-Al® for 10 mg total lipids) according to a described method [49], in a 80/20 v/v mixture of diethyl ether/hexane, giving ether-lipids (1-O-alkylglycerols and 1-O-alk-1-enyl-glycerols) and fatty alcohols. The reaction medium was neutralized at melting-ice temperature using ethanol/water (20/80, v/v). The reaction products were recovered from the diethyl ether phase and evaporated to dryness, then were diluted in chloroform/methanol (2/1, v/v), and finally separated by TLC on 20×20 silica gel plate (LK5) with chloroform/acetone (90/10, v/v) as the eluent. The plate was allowed to dry and was then sprayed with 2,7′dichlorofluorescein (0.2% in ethanol) to reveal the compounds under UV light. In parallel, batyl alcohol (C18:0 AGEL) was deposited on the same plate as a reference standard for collecting ether lipid samples at the same Rf = 0.18. The silica gel containing the ether lipids was scraped and transferred into a tube to be derivatized as 1-O-alkylglycerol-diacetate. Acetylation was performed with a large excess (600 µL) of ethanoyl chloride directly on the silica contained in the tube, by heating at 140 °C for 5 min. Only 1-O-alkylglycerol-diacetate were obtained since alkenyl lipids are destroyed by the presence of acid in the reaction. Reaction medium was neutralized with a saturated aqueous solution of sodium hydrogen carbonate. Diacetate of 1-O-alkylglycerols were extracted with diethyl ether, evaporated to dryness and dissolved in hexane before analysis by gas chromatography. Gas chromatographic separation of acetate-derivatized 1-O-alkylglycerolipids was performed on a Shimadzu GC-2010 Plus system (automatic liquid sample injection unit, on-column injector, flame ionization detector set at 325 °C) equipped with a Zebron 5-MS column (Phenomenex, Le Pecq, France), (30 m, 0.25 mm ID, 0.25 µm film thickness). The temperature program was as follows: 140 °C for 1 min, followed by a ramp at 7.0 °C·min⁻¹ up to 310 °C, then a 3 min hold. The transfer line temperature was 310 °C. The source temperature was 300 °C. The solvent delay was 3 min and the mass range 35–500 m/z.

### 3.3. Mass Spectrometry

#### Optimization of the Separation Method

GC-Ultra-ISQ (Thermo Fisher, Villebon sur Yvette, France) was used with a Zebron 5-MS column (Phenomenex, Le Pecq, France) having a 30 m length, 0.25 mm of diameter, and 0.25 µm film thickness. The injector temperature was 250 °C, the splitless injection volume was 1 µL, and the vector gas was hydrogen with a constant flow rate of 1.5 mL·min⁻¹. The temperature program was as follows: 140 °C for 1 min, followed by a ramp at 7.0 °C·min⁻¹ up to 310 °C, then a 3 min hold. The transfer line temperature was 310 °C. The source temperature was 300 °C. The solvent delay was 3 min and the mass range 35–500 m/z.

### 3.4. Experimentation Section

#### 3.4.1. Preparation of the (2,2-Dimethyl-1,3-dioxolan-4-yl)methyl 4-methylbenzenesulfonate (2)

To a cold (0 °C) solution of (±)-2,2-dimethyl-4-hydroxymethyl-1,3-dioxolane (4.0 g, 30 mmol), in anhydrous pyridine (6 mL), p-TsCl (7.0 g, 37 mmol) was added. After stirring for 24 h, diethyl ether (200 mL) was added and the solution was washed with aqueous HCl (1 M, 2×200 mL), H₂O (5×100 mL), dried over Na₂SO₄, and filtered. Diethyl ether was evaporated to give the desired product as a clear oil. Yield = 81%. ¹H NMR-CDCl₃, δ (ppm) 7.80 (d, J = 8.3 Hz, 2H, CH₂Ar), 7.36 (d,
1H NMR-CDCl₃, δ (ppm) 4.26 (1H, m, CH₆glycerol), 4.01 (1H, m, CH₂O-alkyl), 3.76 (1H, m, CH₂O-alkyl glycerol), 3.40–3.30 (4H, m, CH₂OC and CH₂O-glycerol), 1.70–1.50 (2H, m, CH₂CH₂O-glycerol), 1.48 (3H, s, CH₃ acetal), 1.41 (3H, s, CH₃ acetal), 1.29 (29H, m, CH₂), 0.91 (3H, t, J = 6.8 Hz, CH₃). ¹³C NMR-CDCl₃, δ (ppm) 130.5, 109.6, 74.2, 71.8, 71.3, 67.3, 31.7, 29.7, 29.2, 29.0, 26.6, 25.7, 23.2, 14.0.

4-(n-Heptadecyloxy)methyl-2,2-dimethyl-1,3-dioxolane (3d). This product was obtained from 1-heptadecanol (0.5 g, 1.9 mmol). TLC performed in chloroform/aceto-n (90:10). If = 0.68. Yield = 60% (0.46 g). ¹H NMR-CDCl₃, δ (ppm) 4.26 (1H, m, CH₆glycerol), 4.01 (1H, m, CH₂O-alkyl), 3.76 (1H, m, CH₂O-alkyl glycerol), 3.40–3.30 (4H, m, CH₂OC and CH₂O-glycerol), 1.70–1.50 (2H, m, CH₂CH₂O-glycerol), 1.48 (3H, s, CH₃ acetal), 1.41 (3H, s, CH₃ acetal), 1.29 (30H, m, CH₂), 0.91 (3H, t, J = 6.8 Hz, CH₃). ¹³C NMR-CDCl₃, δ (ppm) 109.6, 74.2, 71.8, 71.3, 67.3, 31.7, 29.7, 29.2, 29.0, 26.6, 25.7, 23.2, 14.0.

4-(n-Octadecyloxy)methyl-2,2-dimethyl-1,3-dioxolane (3e). This product was obtained from 1-octadecanol (0.5 g, 1.8 mmol). TLC performed in chloroform/aceto-n (90:10). If = 0.68. Yield = 65% (0.46 g). ¹H NMR-CDCl₃, δ (ppm) 4.26 (1H, m, CH₆glycerol), 4.01 (1H, m, CH₂O-alkyl), 3.76 (1H, m, CH₂O-alkyl glycerol), 3.40–3.30 (4H, m, CH₂OC and CH₂O-glycerol), 1.70–1.50 (2H, m, CH₂CH₂O-glycerol), 1.48 (3H, s, CH₃ acetal), 1.41 (3H, s, CH₃ acetal), 1.29 (30H, m, CH₂), 0.91 (3H, t, J = 6.8 Hz, CH₃). ¹³C NMR-CDCl₃, δ (ppm) 109.6, 74.2, 71.8, 71.3, 67.3, 31.7, 29.7, 29.2, 29.0, 26.6, 25.7, 23.2, 14.0.

4-(n-Heptadecyloxy)methyl-2,2-dimethyl-1,3-dioxolane (3d). This product was obtained from 1-octadecanol (0.5 g; 2.1 mmol). TLC performed in chloroform/aceto-n (90:10). If = 0.68. Yield = 96% (0.71 g). ¹H NMR-CDCl₃, δ (ppm) 4.26 (1H, m, CH₆glycerol), 4.01 (1H, m, CH₂O-alkyl), 3.76 (1H, m, CH₂O-alkyl glycerol), 3.40–3.30 (4H, m, CH₂OC and CH₂O-glycerol), 1.70–1.50 (2H, m, CH₂CH₂O-glycerol), 1.48 (3H, s, CH₃ acetal), 1.41 (3H, s, CH₃ acetal), 1.29 (26H, m, CH₂), 0.91 (3H, t, J = 6.8 Hz, CH₃). ¹³C NMR-CDCl₃, δ (ppm) 109.6, 74.2, 71.8, 71.3, 67.3, 31.7, 29.7, 29.2, 29.0, 26.6, 25.7, 23.2, 14.0.

4-(n-Heptadecyloxy)methyl-2,2-dimethyl-1,3-dioxolane (3c). This product was obtained from 1-hexadecanol (0.5 g, 2.1 mmol). TLC performed in chloroform/aceto-n (90:10). If = 0.68. Yield = 96% (0.52 g). ¹H NMR-CDCl₃, δ (ppm) 4.26 (1H, m, CH₆glycerol), 4.01 (1H, m, CH₂O-alkyl), 3.76 (1H, m, CH₂O-alkyl glycerol), 3.40–3.30 (4H, m, CH₂OC and CH₂O-glycerol), 1.70–1.50 (2H, m, CH₂CH₂O-glycerol), 1.48 (3H, s, CH₃ acetal), 1.41 (3H, s, CH₃ acetal), 1.29 (18H, m, CH₂), 0.91 (3H, t, J = 7.0 Hz, CH₃). ¹³C NMR-CDCl₃, δ (ppm) 130.5, 109.6, 74.2, 71.8, 71.3, 67.3, 31.7, 29.7, 29.2, 29.0, 28.8, 28.0, 26.6, 25.7, 23.2, 14.0.
20.3 (4H, m, CH\textsubscript{2}-CH=CH-CH\textsubscript{2}), 1.75–1.56 (2H, m, CH\textsubscript{2}CH\textsubscript{2}O-glycerol), 1.47 (3H, s, CH\textsubscript{3} acetate), 1.40 (3H, s, CH\textsubscript{3} acetate), 1.29 (22H, m, CH\textsubscript{2}H\textsubscript{2}), 0.91 (3H, t, J = 7 Hz, CH\textsubscript{3}). \textsuperscript{13}C NMR-CDCl\textsubscript{3}, \(\delta\) (ppm) 130.5, 109.6, 74.2, 71.8, 71.3, 67.3, 31.7, 29.7, 29.3, 29.2, 28.9, 28.8, 28.6, 26.5, 25.7, 23.2, 14.0.

4-(\textit{n}-Nonadecyloxy)methyl)-2,2-dimethyl-1,3-dioxolane (3g). This product was obtained from 1-nonadecanol (0.5 g, 1.75 mmol). TLC performed in chloroform/acetone (90:10). \(R_f = 0.68\). Yield = 79% (0.47 g).

1.48 (3H, s, CH\textsubscript{3} acetate); 1.29 (32H, m, CH\textsubscript{2}H\textsubscript{2}), 0.91 (3H, t, J = 6.8 Hz, CH\textsubscript{3}). \textsuperscript{13}C NMR-CDCl\textsubscript{3}, \(\delta\) (ppm) 109.6, 74.2, 71.8, 71.3, 67.3, 31.7, 29.7, 29.2, 29.0, 26.5, 25.7, 23.2, 14.0.

4-(\textit{n}-Eicosyloxy)methyl)-2,2-dimethyl-1,3-dioxolane (3h). This product was obtained from 1-eicosanol (0.5 g, 1.7 mmol). TLC performed in chloroform/acetone (90:10). \(R_f = 0.65\). Yield = 79% (0.49 g).

1.47 (3H, s, CH\textsubscript{3} acetate); 1.29 (38H, m, CH\textsubscript{2}H\textsubscript{2}), 0.91 (3H, t, J = 6.8 Hz, CH\textsubscript{3}). \textsuperscript{13}C NMR-CDCl\textsubscript{3}, \(\delta\) (ppm) 109.6, 74.2, 71.8, 71.3, 67.3, 31.7, 29.7, 29.0, 26.5, 25.7, 23.2, 14.0.

4-(\textit{n}-Docosyloxy)methyl)-2,2-dimethyl-1,3-dioxolane (3i). This product was obtained from 1-docosanol (0.5 g, 1.5 mmol). TLC performed in chloroform/acetone (90:10). \(R_f = 0.68\). Yield = 98% (0.52 g).

3.69–3.47 (4H, m, CH\textsubscript{2}O-glycerol), 1.62 (2H, m, CH\textsubscript{2}O-glycerol), 1.47 (3H, s, CH\textsubscript{3} acetate), 1.29 (32H, m, CH\textsubscript{2}H\textsubscript{2}), 0.91 (3H, t, J = 6.8 Hz, CH\textsubscript{3}). \textsuperscript{13}C NMR-CDCl\textsubscript{3}, \(\delta\) (ppm) 109.6, 74.2, 71.8, 71.3, 67.3, 31.7, 29.7, 29.2, 29.0, 26.5, 25.7, 23.2, 14.0.

3.4.3. Preparation of the Ether-Glycerol 4a-i by Hydrolysis of Acetal 3a-i.

To a solution of 4-alkoxymethyl-2,2-dimethyl-1,3-dioxolane (1.0 eq.) in methanol (approx. 5 mL), HCl (2 M, 20 eq.) was added and the solution was heated to reflux for 4 h. After cooling to room temperature, the mixture was poured into water, the organic layers were extracted with ether, dried with Na\textsubscript{2}SO\textsubscript{4} and the solvents were removed under vacuum. The residue was purified by flash chromatography with hexane/ethanol 3:1 v/v.

3-(\textit{n}-Tetradecyloxy)propane-1,2-diol (4a). This product was obtained from compound 3a (0.52 g, 1.58 mmol). TLC performed in hexane/ethyl acetate (75:25). \(R_f = 0.10\). Yield = 98% (0.45 g).

1.43 (3H, s, CH\textsubscript{3} acetate), 1.29 (38H, m, CH\textsubscript{2}H\textsubscript{2}), 0.91 (3H, t, J = 6.8 Hz, CH\textsubscript{3}). \textsuperscript{13}C NMR-CDCl\textsubscript{3}, \(\delta\) (ppm) 72.1, 71.8, 71.0, 63.5, 31.7, 29.7, 29.2, 29.0, 26.6, 23.2, 14.0.

3-(\textit{n}-Hexadecyloxy)propane-1,2-diol (4b). This product was obtained from compound 3b (0.71 g, 1.99 mmol). TLC performed in hexane/ethyl acetate (75:25). \(R_f = 0.10\). Yield = 92% (0.58 g).

1.47 (3H, s, CH\textsubscript{3} acetate), 1.29 (38H, m, CH\textsubscript{2}H\textsubscript{2}), 0.91 (3H, t, J = 6.8 Hz, CH\textsubscript{3}). \textsuperscript{13}C NMR-CDCl\textsubscript{3}, \(\delta\) (ppm) 72.1, 71.8, 71.0, 63.5, 31.7, 29.7, 29.2, 29.0, 26.6, 23.2, 14.0.

(7E)-3-(\textit{n}-Hexadec-7-enyloxy)propane-1,2-diol (4c). This product was obtained from compound 3c (0.56 g, 1.58 mmol). TLC performed in hexane/ethyl acetate (75:25). \(R_f = 0.10\). Yield = 98% (0.49 g).

1.43 (3H, s, CH\textsubscript{3} acetate), 1.29 (38H, m, CH\textsubscript{2}H\textsubscript{2}), 0.91 (3H, t, J = 6.8 Hz, CH\textsubscript{3}). \textsuperscript{13}C NMR-CDCl\textsubscript{3}, \(\delta\) (ppm) 315.2894, found 315.2894.
3-(n-Heptadecyloxy)propane-1,2-diol (4d). This product was obtained from compound 3d (0.43 g, 1.16 mmol). TLC performed in hexane/ethyl acetate (75:25). Rf = 0.10. Yield = 95% (0.36 g). 1H NMR-CDCl3, δ (ppm) 3.90 (1H, m, CH-glycerol), 3.80–3.81 (2H, m, CH2O-alkyl), 3.70–3.45 (4H, m, CH2OH and CH2O-glycerol), 2.62 (2H, bs, 2xOH), 1.62 (2H, m, CH2O-glycerol), 1.29 (28H, m, CH2), 0.91 (3H, t, J = 7 Hz, CH3). 13C NMR-CDCl3, δ (ppm) 72.1, 71.8, 71.0, 63.5, 31.7, 29.7, 29.2, 29.0, 26.6, 23.2, 14.; HRMS ([M+H]+) calcd for [C20H42O3H]+ 331.3207, found 331.3210.

3-(n-Octadecyloxy)propane-1,2-diol (4e). This product was obtained from compound 3e (0.46 g, 1.20 mmol). TLC performed in hexane/ethyl acetate (75:25). Rf = 0.10. Yield = 96% (0.40 g). 1H NMR-CDCl3, δ (ppm) 3.90 (1H, m, CH-glycerol), 3.80–3.81 (2H, m, CH2O-alkyl), 3.70–3.45 (4H, m, CH2OH and CH2O-glycerol), 2.62 (2H, bs, 2xOH), 1.62 (2H, m, CH2O-glycerol), 1.29 (30H, m, CH2), 0.91 (3H, t, J = 7 Hz, CH3). 13C NMR-CDCl3, δ (ppm) 72.1, 71.8, 71.0, 63.5, 31.7, 29.7, 29.2, 29.0, 26.6, 23.2, 14.

(9E)-3-(n-Octadec-9-enyloxy)propane-1,2-diol (4f). This product was obtained from compound 3f (0.56 g, 1.46 mmol). TLC performed in hexane/ethyl acetate (75:25). Rf = 0.10. Yield = 96% (0.48 g). 1H NMR-CDCl3, δ (ppm) 5.37–5.30 (m, 2H, CH=CH), 3.88–3.83 (m, 1H, CH), 3.72 (dd, J = 11.1, 7.0, 3.9 Hz, 1H, CH), 3.68–3.62 (m, 1H, CH), 3.54 (dd, J = 9.7, 3.8 Hz, 1H, CH), 3.50 (dd, J = 9.8, 6.0 Hz, 1H, CH), 3.48 (dd, J = 6.4, 9.2 Hz, 1H, CH), 3.45 (dd, J = 9.2, 6.5 Hz, 1H, CH), 2.58 (d, J = 5.1 Hz, 1H, CH), 2.15 (m, 1H, CH), 2.01 (dd, J = 12.8, 6.6 Hz, 4H, CH2=CH=CH-CH2), 1.39–1.15 (m, 24H, CH2), 0.88 (t, J = 6.9 Hz, 3H, CH3). 13C NMR-CDCl3, δ (ppm) 130.5, 72.1, 71.8, 71.0, 63.5, 31.7, 29.7, 29.2, 29.0, 28.8, 28.0, 26.6, 23.2, 14.0.

3-(n-Nonadecyloxy)propane-1,2-diol (4g). This product was obtained from compound 3g (0.49 g, 1.23 mmol). TLC performed in hexane/ethyl acetate (75:25). Rf = 0.10. Yield = 95% (0.42 g). 1H NMR-CDCl3, δ (ppm) 3.90 (1H, m, CH-glycerol), 3.80–3.81 (2H, m, CH2O-alkyl), 3.70–3.45 (4H, m, CH2OH and CH2O-glycerol), 2.62 (2H, bs, 2xOH), 1.62 (2H, m, CH2O-glycerol), 1.29 (32H, m, CH2), 0.91 (3H, t, J = 7 Hz, CH3). 13C NMR-CDCl3, δ (ppm) 72.1, 71.8, 71.0, 63.5, 31.7, 29.7, 29.2, 29.0, 26.6, 23.2, 14.; HRMS ([M+H]+) calcd for [C22H46O3H]+ 359.3520, found 359.3523.

3-(n-Eicosyloxy)propane-1,2-diol (4h). This product was obtained from compound 3h (0.47 g, 1.14 mmol). TLC performed in hexane/ethyl acetate (75:25). Rf = 0.10. Yield = 95% (0.40 g). 1H NMR-CDCl3, δ (ppm) 3.90 (1H, m, CH-glycerol), 3.80–3.81 (2H, m, CH2O-alkyl), 3.70–3.45 (4H, m, CH2OH and CH2O-glycerol), 2.62 (2H, bs, 2xOH), 1.62 (2H, m, CH2O-glycerol), 1.29 (34H, m, CH2), 0.91 (3H, t, J = 7 Hz, CH3). 13C NMR-CDCl3, δ (ppm) 72.1, 71.8, 71.0, 63.5, 31.7, 29.7, 29.2, 29.0, 26.6, 23.2, 14.

3-(n-Docosyloxy)propane-1,2-diol (4i). This product was obtained from compound 3i (0.53 g, 1.20 mmol). TLC performed in hexane/ethyl acetate (75:25). Rf = 0.10. Yield = 95% (0.46 g). 1H NMR-CDCl3, δ (ppm) 3.90 (1H, m, CH-glycerol), 3.80–3.81 (2H, m, CH2O-alkyl), 3.70–3.45 (4H, m, CH2OH and CH2O-glycerol), 2.62 (2H, bs, 2xOH), 1.62 (2H, m, CH2O-glycerol), 1.29 (38H, m, CH2), 0.91 (3H, t, J = 7 Hz, CH3). 13C NMR-CDCl3, δ (ppm) 72.1, 71.8, 71.0, 63.5, 31.7, 29.7, 29.2, 29.0, 26.6, 23.2, 14.

3.4.4. Derivation Procedure: Preparation of 3-(Alk oxy)propane-1,2-diacetate

Method A [43]: To a solution of 3-(alkoxy)propane-1,2-diol (1.0 eq.) in pyridine (4.0 eq.), acetic anhydride (5.5 eq.) was added and the solution was heated to 60 °C for 30 min and stirred at room temperature overnight. The organic layer was extracted with diethyl ether, washed with HCl (1 M), an aqueous solution of K2CO3 (10%), and a saturated solution of NaCl. The resulting organic layer was dried over Na2SO4 and the solvents were removed under vacuum. The residue was purified by flash chromatography using chloroform/acetone (90:10).

Method B [44]: In a screw cap tube, to the 3-(alkoxy)propane-1,2-diol (1.0 eq.), acetyl chloride (2.1 eq.) was added. The tube was closed under inert gas and heated at 140 °C for 5 min. After cooling at room temperature, diethyl ether was added. The excess of acetyl chloride was neutralized by dropwise
addition of a saturated aqueous solution of sodium hydrogen carbonate until no more bubbles were observed. The organic layer was filtered over Na2SO4 and evaporated to dryness to give the pure compound as a white powder.

3-(Tetradecyloxy)propane-1,2-diacetate (5a). This product was obtained from compound 4a by using method A (0.45 g, 1.56 mmol). TLC performed in chloroform/acetone (96:4). Rf = 0.77. Yield = 95% (0.55 g). 1H NMR-CDCl3, δ (ppm) 5.20 (1H, m, CH2-glycerol), 4.50 & 4.07 (2H, m, CH2-O-acetate), 3.67 & 3.46 (2H, m, CH2-O-alkyl), 5.33−3.30 (2H, m, CH2-O-glycerol), 2.07 (3H, s, CH3-acetate), 2.03 (3H, s, CH3-acetate), 1.55−1.47 (4H, m, CH2-alkyl), 1.30−1.24 (20H, m, CH2), 0.89 (3H, t, J = 7 Hz, CH3). 13C NMR-CDCl3, δ (ppm) 171.8, 170.5, 71.8, 70.6, 70.0, 63.7, 31.7, 29.7, 29.2, 29.0, 26.6, 23.2, 21.3, 20.7, 14.

3-(Hexadecyloxy)propane-1,2-diacetate (5b). This product was obtained from compound 4b by using method A (0.58 g, 1.83 mmol). TLC performed in chloroform/acetone (96:4). Rf = 0.77. Yield = 95% (0.70 g). 1H NMR-CDCl3, δ (ppm) 5.20 (1H, m, CH2-glycerol), 4.49 & 4.09 (2H, m, CH2-O-acetate), 3.67 & 3.46 (2H, m, CH2-O-alkyl), 3.34−3.30 (2H, m, CH2-O-glycerol), 2.07 (3H, s, CH3-acetate), 2.03 (3H, s, CH3-acetate), 1.55−1.47 (4H, m, CH2-alkyl), 1.30−1.24 (24H, m, CH2), 0.89 (3H, t, J = 7 Hz, CH3). 13C NMR-CDCl3, δ (ppm) 171.8, 170.5, 71.8, 70.6, 70.0, 63.7, 31.7, 29.7, 29.2, 29.0, 26.6, 23.2, 21.3, 20.7, 14.

(E)-3-(Hexadec-7-enyloxy)propane-1,2-diacetate (5c). This product was obtained from compound 4c by using method B (0.49 g, 1.56 mmol). TLC performed in chloroform/acetone (96:4). Rf = 0.77. Yield = 98% (0.61 g). 1H NMR-CDCl3, δ (ppm) 5.34−5.30 (2H, m, CH=CH), 5.20 (1H, m, CH2-glycerol), 4.50 (1H, m, CH2-O-acetate), 4.04 (1H, m, CH2-O-acetate), 3.66 (1H, m, CH2-O-alkyl), 3.44 (H, m, CH2-O-alkyl), 3.33 (2H, m, CH2-O-glycerol), 2.07−1.98 (10H, m, 2 x CH3 acacet and CH2-CH=CH-CH2), 1.55 (2H, m, CH2CH2O), 1.47 (2H, m, CH2), 1.33−1.26 (16H, m, CH2), 0.89 (3H, t, J = 7 Hz, CH3). 13C NMR-CDCl3, δ (ppm) 171.8, 170.5, 130.5, 71.8, 70.6, 70.0, 63.7, 31.7, 29.7, 29.3, 29.2, 29.0, 28.8, 28.0, 26.6, 23.2, 21.3, 20.7, 14.0.

3-(Heptadecyloxy)propane-1,2-diacetate (5d). This product was obtained from compound 4d by using method A (0.36 g, 1.08 mmol). TLC performed in chloroform/acetone (96:4). Rf = 0.77. Yield = 92% (0.42 g). 1H NMR-CDCl3, δ (ppm) 5.20 (1H, m, CH2-glycerol), 4.49 & 4.09 (2H, m, CH2-O-acetate), 3.67 & 3.46 (2H, s, CH2-O-alkyl), 3.34−3.30 (2H, m, CH2-O-glycerol), 2.07 (3H, s, CH3-acetate), 2.03 (3H, s, CH3-acetate), 1.55−1.47 (4H, m, CH2-alkyl), 1.30−1.24 (26H, m, CH2), 0.89 (3H, t, J = 7 Hz, CH3). 13C NMR-CDCl3, δ (ppm) 171.8, 170.5, 71.8, 70.6, 70.0, 63.7, 31.7, 29.7, 29.2, 29.0, 26.6, 23.2, 21.3, 20.7, 14.

3-Octadecyloxy)propane-1,2-diacetate (5e). This product was obtained from compound 4e by using method B (0.40 g, 1.16 mmol). TLC performed in chloroform/acetone (96:4). Rf = 0.77. Yield = 93% (0.46 g). 1H NMR-CDCl3, δ (ppm) 5.20 (1H, m, CH2-glycerol), 4.49 & 4.09 (2H, m, CH2-O-acetate), 3.67 & 3.46 (2H, m, CH2-O-alkyl), 3.34−3.30 (2H, m, CH2-O-glycerol), 2.07 (3H, s, CH3-acetate), 2.03 (3H, s, CH3-acetate), 1.55−1.47 (4H, m, CH2-alkyl), 1.30−1.24 (28H, m, CH2), 0.89 (3H, t, J = 7 Hz, CH3). 13C NMR-CDCl3, δ (ppm) 171.8, 170.5, 71.8, 70.6, 70.0, 63.7, 31.7, 29.7, 29.2, 29.0, 26.6, 23.2, 21.3, 20.7, 14.

(9E)-3-(Octadec-9-enyloxy)propane-1,2-diacetate (5f). This product was obtained from compound 4f by using method B (0.48 g, 1.40 mmol). TLC performed in chloroform/acetone (96:4). Rf = 0.77. Yield = 96% (0.57 g). 1H NMR-CDCl3, δ (ppm) 5.38−5.31 (m, 2H, CH=CH), 5.21−5.16 (m, 1H, CH), 4.33 (dd, J = 12.0, 3.6 Hz, 1H, CH), 4.16 (dd, J = 12.0, 6.4 Hz, 1H, CH), 3.55 (dd, J = 10.8, 5.4 Hz, 1H, CH), 3.54 (dd, J = 10.6, 5.1 Hz, 1H, CH), 3.50−3.38 (m, 2H, CH2), 2.09 (s, 3H, CH3 acetate), 2.07 (s, 3H, CH3 acetate), 2.01 (dd, J = 12.8, 6.8 Hz, 4H, CH2-CH=CH-CH2), 1.39−1.20 (m, 24H, CH2), 0.88 (t, J = 7.0 Hz, 3H, CH3). 13C NMR-CDCl3, δ (ppm) 171.8, 170.5, 130.5, 71.8, 70.6, 70.0, 63.7, 31.7, 29.7, 29.3, 29.2, 29.0, 28.8, 28.0, 26.6, 23.2, 21.3, 20.7, 14.0.

3-(Nonadecyloxy)propane-1,2-diacetate (5g). This product was obtained from compound 4g by using method A (0.42 g, 1.17 mmol). TLC performed in chloroform/acetone (96:4). Rf = 0.77. Yield = 95% (0.49 g). 1H NMR-CDCl3, δ (ppm) 5.20 (1H, m, CH2-glycerol), 4.49 & 4.09 (2H, m, CH2-O-acetate), 3.67 & 3.46 (2H, m, CH2-O-alkyl), 3.34−3.30 (2H, m, CH2-O-glycerol), 2.07 (3H, s, CH3-acetate), 2.03 (3H, s,
with diethyl ether. After evaporation, the sample was taken up with hexane.

were scraped into glass tubes with Teflon screw caps and acetylated with a large excess (600 µl.

Alkylglycerols were obtained by separation of lipid extracts by silica gel thin layer chromatography (see Table 1).

1(CH, m, CH₂CO₂) for 5 min at 140°C. Alkylglycerols were identified by comparison of their retention and their fragmentation peaks in mass spectroscopy with those of the synthesized standards (see Table 1).

3.5. Analytical Chemistry

Analytical Biochemistry

The total lipids were extracted from 2 mg chimera and shark liver oil by Folch’s method [48]. Alkylglycerols were obtained by separation of lipid extracts by silica gel thin layer chromatography with chloroform/acetone (90:10, v/v) as the developing solvent. Separated lipids were visualized under ultraviolet light after spraying with 0.2% solution of 2,7-dichlorofluorescein in ethanol. Alkylglycerols were scraped into glass tubes with Teflon screw caps and acetylated with a large excess (600 µl) of acetyl chloride for 5 min at 140°C. Then, the solution was neutralized with NaHCO₃ and extracted with diethyl ether. After evaporation, the sample was taken up with hexane.

For the chromatography procedure, the samples were analyzed in batches of 13 samples comprising 10 oil samples, 3 samples comprising two blanks and one mixture of synthesized alkyl lipids serving as a reference. The alkylglycerols were identified by comparison of their retention times and their fragmentation peaks in mass spectroscopy with those of the synthesized standards.

Table 1. Mass spectroscopy data.

| Alkyl Chain R | Retention Time (min) | Molecular Weight (g. x mol⁻¹) | Most Characteristic Peaks (m/z, %) |
|---------------|----------------------|--------------------------------|----------------------------------|
| C₁₄H₂₉        | 13.6                 | 372.54                         | 43 (base peak), 57 (60), 71 (35), 83 (30), 97 (26), 111 (13), 117 (19), 159 (119), 223 (4), 269 (6), 313 (35) |
| C₁₅H₁₄        | 15.0                 | 398.58                         | 43 (base peak), 57 (63), 69 (44), 81 (73), 93 (39), 117 (31), 159 (63), 222 (15), 281 (12), 339 (25) |
| C₁₆H₁₃        | 15.7                 | 400.59                         | 43 (base peak), 57 (62), 71 (38), 83 (28), 97 (27), 111 (14), 159 (7), 219 (&-, 251 (5), 255 (<), 297 (6), 341 (67) |
| C₁₇H₁₄        | 16.8                 | 414.62                         | 43 (base peak), 57 (62), 83 (26), 97 (27), 111 (15), 117 (22) 159 (25), 209 (<), 267 (8), 311 (12), 355 (54) |
| C₁₈H₁₅        | 17.5                 | 426.63                         | 43 (base peak), 57 (52), 67 (48), 81 (72), 95 (44), 117 (18), 157 (75), 191 (7), 219 (<), 281 (12), 341 (7), 367 (25) |
| C₁₉H₁₆        | 17.8                 | 428.65                         | 43 (base peak), 57 (70), 71 (45), 85 (34), 97 (33), 111 (18), 117 (27), 159 (27), 191 (<), 281 (10), 325 (9), 369 (37) |
| C₂₀H₁₇        | 18.7                 | 442.67                         | 43 (base peak), 57 (65), 83 (38), 71 (38), 97 (36), 111 (20), 117 (32), 159 (30), 233 (<), 267 (8), 311 (12), 355 (47) |
| C₂₁H₁₈        | 19.7                 | 456.70                         | 43 (base peak), 57 (70), 71 (42), 83 (36), 97 (36), 111 (18), 117 (32), 159 (27), 207 (23), 295 (12), 355 (32), 398 (40) |
| C₂₂H₁₉        | 21.4                 | 484.75                         | 43 (base peak), 57 (70), 71 (38), 83 (28), 97 (38), 111 (22), 117 (12), 125 (12), 159 (30), 207 (22), 281 (8), 355 (10), 381 (6), 423 (27) |
3.6. Validation of the Analytical Procedure

The Table 2 represents the Statistical data for analytical procedure for GC and show the fiability of the method used.

**Table 2.** Statistical data for analytical procedure for GC.

| Parameters Studied | Results |
|--------------------|---------|
| Selectivity        | Correct |
| Fidelity           | Correct (CVR = 3.73%) |
| Repetitability     | Correct (CVr = 3.73%) |
| Reproducibility    | Correct (no deviation from linearity, \( R^2 = 0.992 \)) |
| Linearity          | 2.9 pg (on-column injector, FID) |
| Detection limit LOD| 17.5 pg (on-column injector, FID) |
| Quantification limit LOQ | 1 year at \(-80^\circ C\) |

4. Conclusions

This work demonstrates that a series of 1-O-alkyl-glycerols featuring mainly saturated lipid chain lengths (14:0, 16:0, 17:0, 19:0, 20:0, 22:0) and two mono-unsaturated lipid chains (16:1, 18:1) were easily prepared and derivatized as diacetate compounds that were used as standards for GC analysis. These standards were used to determine the molecular species of AEL present in shark oils. It would be interesting to use these standards to also determine the AEL composition in tumors. Indeed, it is known that the ether-lipids fraction is larger in tumors than non-tumoral tissues, but the molecular species have not been precisely identified. Moreover, in a second time it would be interesting to synthesize standards that would identify alkenyl-lipids. Indeed, alkenyl-lipids were found to be highly present in the membrane phospholipids of some tissues such as the heart and brain [50]. The determination of the composition of the alkenyl-lipids in tumors will help to understand the role of these lipids in the biology of tumors cells.

**Supplementary Materials:** The following are available online at http://www.mdpi.com/1660-3397/16/4/101/s1. Figure SI1-1. \(^1\)H NMR (CDCl\(_3\)) spectrum of compound 2, Figure SI1-2. \(^1\)H NMR (CDCl\(_3\)) spectrum of compound 3f, Figure SI1-3. \(^1\)H NMR (CDCl\(_3\)) spectrum of compound 4f, Figure SI1-4. 2D NMR HSQC (CDCl\(_3\)) spectrum of compound 4f, Figure SI1-5. \(^1\)H NMR (CDCl\(_3\)) spectrum of compound 5f, Figure SI1-6. 2D NMR HSQC (CDCl\(_3\)) spectrum of compound 5f, Figure SI1-7. \(^1\)H NMR (CDCl\(_3\)) spectrum of compound 5f, Figure SI1-8. 2D NMR HSQC (CDCl\(_3\)) spectrum of compound 5f, Figure SI2-1. GC-MS chromatogram of a mixture of synthesized alkyl-glycerolipids and Figure SI2-2. Mass spectroscopy data.

**Acknowledgments:** This work was funded by University of Tours, the “Région Centre-Val de Loire” (LIPIDS project of ARD2020 Biomédicaments), “INSERM”, Canceropôle Grand Ouest and the Ion Channel and Cancer Network, the “Ligue Nationale Contre le Cancer”, the Association “CANCEN”, and Tours’ Hospital oncology association ACORT. We thank Bodimant of Phosphotec for providing chimera and shark liver oils. We also thank Yves Frangin, Ludovic Sochard, Lydie Nadal-Desbarats, Aurore Lecaille and Violetta Guérin for technical assistance, and Catherine Leroy for secretarial support.

**Author Contributions:** M.P., C.G., H.C., J.T., P.B., and P-A.J. designed the chemical synthesis. M.P. and C.G. performed the chemical synthesis and the analytical analyses. D.F., A.C., and S.C. provided intellectual input. M.P., C.G., P-A.J., and C.V. wrote the manuscript. All the authors critically reviewed the final version of the manuscript.

**Conflicts of Interest:** The authors declare no competing financial interest.

**References**

1. Magnusson, C.D.; Haraldsson, G.G. Ether lipids. *Chem. Phys. Lipids* 2011, 164, 315–340. [CrossRef] [PubMed]
2. Oh, S.Y.; Jadhav, L.S. Effects of dietary alkylglycerols in lactating rats on immune responses in pups. *Pediatr. Res.* 1994, 36, 300–305. [CrossRef] [PubMed]
3. Linman, J.W.; Bethell, F.H.; Long, M.J. The erythropoietic simulatory activity of batyl alcohol. *J. Lab. Clin. Med.* 1958, 52, 596–604. [PubMed]
4. Brohult, A.; Brohult, J.; Brohult, S. Regression of tumour growth after administration of alkoxyglycerols. *Acta Obstet. Gynecol. Scand.* 1978, 57, 79–83. [CrossRef] [PubMed]

5. Brohult, A.; Brohult, J.; Brohult, S.; Joelsson, I. Effect of alkoxyglycerols on the frequency of injuries following radiation therapy for carcinoma of the uterine cervix. *Acta Obstet. Gynecol. Scand.* 1977, 56, 441–448. [CrossRef] [PubMed]

6. Pedrono, F.; Martin, B.; Leduc, C.; Le Lan, J.; Saiag, B.; Legrand, P.; Moulinoux, J.P.; Legrand, A.B. Natural alkylglycerols restrain growth and metastasis of grafted tumors in mice. *Nutr. Cancer* 2004, 48, 64–69. [CrossRef] [PubMed]

7. Zhou, X.; Mao, J.; Ai, J.; Deng, Y.; Roth, M.R.; Pound, C.; Henegar, J.; Welti, R.; Bigler, S.A. Identification of plasma lipid biomarkers for prostate cancer by lipidomics and bioinformatics. *PLoS ONE* 2012, 7, e48889. [CrossRef] [PubMed]

8. Benveniste, J. Platelet-activating factor, a new mediator of anaphylaxis and immune complex deposition from rabbit and human basophils. *Nature* 1974, 249, 581–582. [CrossRef] [PubMed]

9. Benveniste, J.; Vargaftig, B.B. Platelet-activating factor: An ether lipid with biological activity. In *Ether Lipids: Biochemical and Biomedical Aspects*; Mangold, H.K., Paltauf, F., Eds.; Academic Press: New York, NY, USA, 1983; pp. 355–376.

10. Hichami, A.; Duroudier, V.; Leblais, V.; Vernhet, L.; Le Goffic, F.; Ninio, E.; Legrand, A. Modulation of platelet-activating-factor production by incorporation of naturally occurring 1-O-alkylglycerols in phospholipids of human leukemic monocyte-like THP-1 cells. *Eur. J. Biochem.* 1997, 250, 242–248. [CrossRef] [PubMed]

11. Clark, K.J.; Murray, A.W. Evidence that the bradykinin-induced activation of phospholipase D and of the mitogen-activated protein kinase cascade involve different protein kinase C isoforms. *J. Biol. Chem.* 1995, 270, 7097–7103. [CrossRef] [PubMed]

12. Marigny, K.; Pedrono, F.; Martin-Chouly, C.A.; Youmine, H.; Saiag, B.; Legrand, A.B. Modulation of endothelial permeability by 1-O-alkylglycerols. *Acta Physiol. Scand.* 2002, 176, 263–268. [CrossRef] [PubMed]

13. Warne, T.R.; Buchanan, F.G.; Robinson, M. Growth-dependent accumulation of monoalkylglycerol in Madin-Darby canine kidney cells. Evidence for a role in the regulation of protein kinase C. *J. Biol. Chem.* 1995, 270, 11147–11154. [CrossRef] [PubMed]

14. O’Brian, C.; Vogel, V.G.; Singletery, S.E.; Ward, N.E. Elevated protein kinase C expression in human breast tumor biopsies relative to normal breast tissue. *Cancer Res.* 1989, 49, 3215–3217. [PubMed]

15. Daniel, L.W.; Small, G.W.; Schmitt, J.D.; Marasco, C.J.; Ishaq, K.; Piantadosi, C. Alkyl-linked diglycerides inhibit protein kinase C activation by diacylglycerols. *Biochem. Biophys. Res. Commun.* 1988, 151, 291–297. [CrossRef]

16. Berggren, M.I.; Gallegos, A.; Dressler, L.A.; Modest, E.J.; Powis, G. Inhibition of the signalling enzyme phosphatidylinositol-3-kinase by antitumor ether lipid analogues. *Cancer Res.* 1993, 53, 4297–4302. [PubMed]

17. Zhou, X.; Lu, X.; Richard, C.; Xiong, W.; Litchfield, D.W.; Bittman, R.; Arthur, G. 1-O-octadecyl-2-O-methyl-glycerophosphocholine inhibits the transduction of growth signals via the MAPK cascade in cultured MCF-7 cells. *J. Clin. Investig.* 1996, 98, 937–944. [CrossRef] [PubMed]

18. Zheng, B.; Oishi, K.; Shoji, M.; Eibl, H.; Berdel, W.E.; Hajdu, J.; Vogler, W.R.; Kuo, J.F. Inhibition of protein kinase C, (sodium plus potassium)-activated adenosine triphosphatase, and sodium pump by synthetic phospholipid analogues. *Cancer Res.* 1990, 50, 3025–3031. [PubMed]

19. Besson, P.; Gore, J.; Vincent, E.; Hoinard, C.; Bougnoux, P. Inhibition of Na⁺/H⁺ exchanger activity by an alkyl-lysophospholipid analogue in a human breast cancer cell line. *Biochem. Pharmacol.* 1996, 51, 1153–1158. [CrossRef]

20. Potier, M.; Chantome, A.; Joulin, V.; Girault, A.; Roger, S.; Besson, P.; Jourdan, M.L.; LeGuennec, J.Y.; Bougnoux, P.; Vandier, C. The SK3/K(Ca)2.3 potassium channel is a new cellular target for edelfosine. *Br. J. Pharmacol.* 2011, 162, 464–479. [CrossRef] [PubMed]

21. Hoffman, D.R.; Thomas, V.L.; Snyder, F. Inhibition of cellular transport systems by alkyl phospholipid analogs in HL-60 human leukemia cells. *Biochim. Biophys. Acta* 1992, 1127, 74–80. [CrossRef]

22. Arthur, G.; Bittman, R. Glycosylated antitumor ether lipids: Activity and mechanism of action. *Anti-Cancer Agents Med. Chem.* 2014, 14, 592–606. [CrossRef]
23. Samadder, P.; Xu, Y.; Schweizer, F.; Arthur, G. Cytotoxic properties of d-glucose-, d-galactose- and d-mannose-configured 2-amino-2-deoxy-glycerolipids against epithelial cancer cell lines and BT-474 breast cancer stem cells. *Eur. J. Med. Chem.* 2014, 78, 225–235. [CrossRef] [PubMed]

24. Sevrain, C.M.; Haelters, J.P.; Chantome, A.; Couthon-Gourves, H.; Gueguinou, M.; Potier-Cartereau, M.; Vandier, C.; Jaffres, P.A. DiGalactosyl-Glycero-Ether Lipid: Synthetic approaches and evaluation as SK3 channel inhibitor. *Org. Biomol. Chem.* 2013, 11, 4479–4487. [CrossRef] [PubMed]

25. Girault, A.; Haelters, J.P.; Potier-Cartereau, M.; Chantome, A.; Jaffres, P.A.; Bougnoux, P.; Jouliv, V.; Vandier, C. Targeting SKCa Channels in Cancer: Potential New Therapeutic Approaches. *Curr. Med. Chem.* 2012, 19, 697–713. [CrossRef] [PubMed]

26. Sevrain, C.M.; Haelters, J.P.; Chantome, A.; Couthon-Gourves, H.; Girault, A.; Vandier, C.; Jaffres, P.A. Glyco-Phospho-Glycerophero Ether Lipid (GPGEI): Synthesis and evaluation as Small Conductance Ca2+-Activated K+ Channels (SK3) inhibitor. *MedChemComm* 2012, 3, 1417–1478. [CrossRef]

27. Gueguinou, M.; Harnois, T.; Crottes, D.; Uguen, A.; Deliot, N.; Gambade, A.; Chantome, A.; Haelters, J.P.; Jaffres, P.A.; Jourdan, M.L.; et al. SK3/TRPC1/Orai1 complex regulates SOCE-dependent colon cancer cell migration: A novel opportunity to modulate anti-EGFR mAb action by the alkyl-lipid Ohmline. *Oncotarget* 2016, 7, 36168–36184. [CrossRef] [PubMed]

28. Berthe, W.; Sevrain, C.M.; Chantome, A.; Bouchet, A.M.; Gueguinou, M.; Fourbon, Y.; Potier-Cartereau, M.; Haelters, J.P.; Couthon-Gourves, H.; Vandier, C.; et al. New Disaccharide-Based Ether Lipids as SK3 Ion Channel Inhibitors. *ChemMedChem* 2016, 11, 1531–1539. [CrossRef] [PubMed]

29. Chantome, A.; Potier-Cartereau, M.; Clarysse, L.; Fromont, G.; Marionneau-Lambot, S.; Gueguinou, M.; Pages, J.C.; Collin, C.; Oullier, T.; Girault, A.; et al. Pivotal role of the lipid Raft SK3-Orai1 complex in human cancer cell migration and bone metastases. *Cancer Res.* 2013, 73, 4852–4861. [CrossRef] [PubMed]

30. Girault, A.; Haelters, J.P.; Potier-Cartereau, M.; Chantome, A.; Pinault, M.; Marionneau-Lambot, S.; Oullier, T.; Simon, G.; Couthon-Gourves, H.; Jaffres, P.A.; et al. New alkyl-lipid blockers of SK3 channels reduce cancer cell migration and occurrence of metastasis. *Curr. Cancer Drug Targets* 2011, 11, 1111–1125. [CrossRef] [PubMed]

31. Jaffres, P.A.; Gajate, C.; Bouchet, A.M.; Couthon-Gourves, H.; Chantome, A.; Potier-Cartereau, M.; Besson, P.; Bougnoux, P.; Mollinedo, F.; Vandier, C. Alkyl ether lipids, ion channels and lipid raft reorganization in cancer therapy. *Pharmacol. Ther.* 2016, 165, 114–131. [CrossRef] [PubMed]

32. Herrera, F.E.; Sevrain, C.M.; Jaffrés, P.A.; Couthon, H.; Grèlard, A.; Dufourc, E.J.; Chantôme, A.; Potier-Cartereau, M.; Vandier, C.; Bouchet, A.M. Singular Interaction between an Antimetastatic Agent and the Lipid Bilayer: The Ohmline Case. *ACS Omega* 2017, 2, 6361–6370. [CrossRef]

33. Hallgren, B.; Larsson, S. The glycerol ethers in man and cow. *J. Lipid Res.* 1962, 3, 39–43.

34. Hallgren, B.; Larsson, S. The glycerol ethers in the liver oils of elasmobranch fish. *J. Lipid Res.* 1962, 3, 31–38.

35. Hallgren, B.; Niklasson, A.; Stallberg, G.; Thorin, H. On the occurrence of 1-O-alkyglycer-1-0-(2-methoxyalkyl)glycerolipids in human colostrum, human milk, cow’s milk, sheep’s milk, human red bone marrow, red cells, blood plasma and a uterine carcinoma. *Acta Chem. Scand.* 1965, 19, 4168–4175. [CrossRef] [PubMed]

36. Snyder, F.; Wood, R. Alkyl and alk-1-enyl ethers of glycerol in lipids from normal and neoplastic human tissues. *Cancer Res.* 1969, 29, 251–257. [PubMed]

37. Prinz, H.; Six, L.; Rues, K.P.; Liefländer, M.L. Stereoselktive Synthese von langkettigen 1-O-(β-D-Maltosyl)-3-O-alkyl-sn-glycerinen (Alkylglycerylether-Lysoglycolipide). *Liebig's Ann. Chem.* 1985, 217–225.

38. Aranda, G.; Lallemand, J.Y. Proposal for Modulable Photoaffinity Labelling. *Synth. Commun.* 1990, 20, 85–96. [CrossRef]

39. Huang, Z.; Szoka, F.C. Sterol-Modified Phospholipids: Cholesterol and Phospholipid Chimeras with Improved Biomembrane Properties. *J. Am. Chem. Soc.* 2008, 130, 15702–15712. [CrossRef] [PubMed]

40. Bhatia, S.K.; Hajdu, J. Stereospecific Synthesis of 2-Substituted Ether Phospholipids. *Synthesis* 1989, 1, 16–20. [CrossRef]

41. Nelson, W.L.; Wenerström, J.E.; Sankar, S.R. Absolute configuration of glycerol derivatives. 3. Synthesis and Cupra A circular dichroism spectra of some chiral 3-aryloxy-1,2-propanediols and 3-aryloxy-1-amino-2-propanols. *Circ. Res.* 1977, 42, 1006–1012.

42. Cymerman, C.; Hamon, D.P.G. Studies Directed toward the Synthesis of Plasmalogens. II. (±)-cis- and -trans-(n-Hexadec-1′-enoxy)-1,2-propanediol1. *J. Org. Chem.* 1965, 30, 4168–4175.
43. Bonner, T.G.; McNamara, P. The Pyridine-catalysed Acetylation of Phenols and Alcohols. *J. Med. Soc.* **1968**, *0*, 795–797.
44. George, B.; Jamieson, R.; Reid, E.H. The analysis of oils and fats by gas chromatography: III. Separation factors of acetates, alcohols and hydrocarbons. *J. Chromatogr.* A **1967**, *26*, 8–16.
45. Bordier, C.G.; Sellier, N.; Foucault, A.P.; Le Goffic, F. Purification and characterization of deep sea shark *Centrophorus squamosus* liver oil 1-O-alkylglycerol ether lipids. *Lipids* **1996**, *31*, 521–528. [CrossRef] [PubMed]
46. Spener, F.; Mangold, H.K. The alkyl moieties in wax esters and alkyl diacyl glycerols of sharks. *J. Lipid Res.* **1971**, *12*, 12–16. [PubMed]
47. Deniau, A.L.; Mosset, P.; Pedrono, F.; Mitre, R.; Le Bot, D.; Legrand, A.B. Multiple beneficial health effects of natural alkylglycerols from shark liver oil. *Mar. Drugs* **2010**, *8*, 2175–2184. [CrossRef] [PubMed]
48. Folch, J.; Lees, M.; Sloane Stanley, G.H. A simple method for the isolation and purification of total lipides from animal tissues. *J. Biol. Chem.* **1957**, *226*, 497–509. [PubMed]
49. Kuksis, A. Fatty Acids and Glycerides. *Handb. Lipid Res.* **2012**, *3*, 126.
50. Rapport, M.M. The discovery of plasmalogen structure. *J. Lipid Res.* **1984**, *25*, 1522–1527. [PubMed]

© 2018 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (http://creativecommons.org/licenses/by/4.0/).