

Structural Characterization of Monohydroxyeicosatetraenoic Acids and Dihydroxy- and Trihydroxyeicosatrienoic Acids by ESI-FTICR

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The fragmentation characteristics of monohydroxyeicosatetraenoic acids and dihydroxy- and trihydroxyeicosatrienoic acids were investigated by electrospray ionization Fourier transform ion cyclotron resonance (FTICR) mass spectrometry using sustained off-resonance irradiation collision-induced dissociation (SORI-CID) and infrared multiphoton dissociation (IRMPD). The fragmentation patterns of these compounds were associated with the number and positions of the hydroxyl substituents. The fragmentation is more complicated with increasing number of the hydroxyl groups of the compounds. In general, the major carbon-carbon cleavage of $[M - H]^-$ ions occurred at the α -position to the hydroxyl group, and the carbon-carbon cleavage occurred when there was a double-bond at the β -position to the hydroxyl group. SORI-CID and IRMPD produced some common fragmentation patterns; however, each technique provided some unique patterns that are useful for structural identification of these compounds. This study demonstrated the application of FTICR via the identification of regioisomers of trihydroxyeicosatrienoic acids in rabbit aorta samples. (J Am Soc Mass Spectrom 2008, 19, 569–585) © 2008 American Society for Mass Spectrometry

Mass spectrometry has emerged as an important tool for analysis of biomolecules. Fourier transform ion cyclotron resonance (FTICR) mass spectrometry is one of the techniques that can provide high mass accuracy and high mass resolution [1–6]. Accurate molecular weight, elemental composition, and structural information can be achieved from FTICR. Sustained off-resonance irradiation collision-induced dissociation (SORI-CID) and infrared multiphoton dissociation (IRMPD) are two different dissociation techniques often used with FTICR for MS/MS analysis. SORI-CID involves collision of the target ion slightly off its resonance frequency with a collision gas, causing the acceleration and deceleration of ions during the RF pulse. At a frequency of several kilohertz, multiple low-energy collisions occur as ions are vibrationally excited for a sustained period. Unlike SORI-CID, no collision gas is required for IRMPD. Instead, the 75 W CO₂ laser is used to irradiate the ions to form fragments. The fragments may continue to acquire some energy from the infrared laser pulse and further fragment to ions of lower masses. This study demon-

strates the utility of SORI-CID and IRMPD for the structural characterization of monohydroxyeicosatetraenoic acids (HETEs) and dihydroxy- and trihydroxyeicosatrienoic acids (DHETs and THETAs).

HETEs, DHETs, and THETAs are metabolites of arachidonic acid (AA). Different isoforms of lipoxygenases (LOX) metabolize AA to regioisomeric HETEs. 12-Hydroxy-5,8,10,14-eicosatetraenoic acid (12-HETE) and 15-HETE are important lipid mediators in inflammation, kidney, immune system, prostate diseases, and diabetes [7–14]. Cytochrome P450 epoxygenases metabolize AA to 4 regioisomeric epoxyeicosatrienoic acids (EETs), and soluble epoxide hydrolase (sEH) enzymes subsequently convert EETs to the corresponding DHETs [15–17]. EETs have various biological functions, including inhibiting the hydro-osmotic action of arginine vasopressin in the kidney, calcium mobilization, and prostaglandin formation [18, 19]. EETs stimulate relaxation in coronary rings and coronary arterioles [20–25]. A recent study showed that 14,15-DHET is a potent peroxisome proliferator-activated receptor- α (PPAR α) activator in COS-7 cells [26].

THETAs are new members of the family of endothelium-derived relaxing factors [27–31]. 11,12,15-THETA and 11,14,15-THETA were identified as endothelium-derived lipoxygenase metabolites of AA in

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the rabbit aorta. Recent studies have shown that THETAs relax rabbit small mesenteric arteries [31]. 11,12,15-THETA mediates acetylcholine-induced relaxations by activating apamin-sensitive potassium (K^+) channels in vascular smooth muscle to induce K^+ efflux, membrane hyperpolarization, and vascular relaxation [27, 28], while 11,14,15-THETA is not vasoactive [28, 29]. These studies cogently indicate that biological function is dependent upon the position of the hydroxyl groups.

Previous studies showed that HETEs and DHETs form characteristic fragments during MS^2 analysis by ion trap and triple quadrupole mass spectrometers [32–40]. The mechanisms for electrospray ionization and tandem mass spectrometry of various classes of eicosanoids have been elegantly reviewed [41]. In this study, we investigated the mass spectrometric characteristics of HETEs, DHETs, and THETAs by ESI-FTICR using SORI-CID and IRMPD. The effects of the number and the positions of the hydroxyl substituents on fragmentation patterns were characterized, and the identities of THETAs in biological samples were determined.

Experimental

Materials and Methods

11-, 12-, and 15-HETE; 11,12- and 14,15-DHET; 14,15-EET, and arachidonoyl dopamine were purchased from Cayman Chemical Co. (Ann Arbor, MI). 11,12,15-, 11,14,15-, and 13,14,15-THETA were synthesized in the laboratory of Dr. J. R. Falck [42]. Indomethacin, A23187, and L-ascorbic acid were purchased from Sigma (St. Louis, MO). C_{18} Bond Elut solid-phase extraction (SPE) columns were purchased from Varian (Harbor City,

CA). Acetonitrile was HPLC grade. Distilled, deionized water was used in all experiments.

Biological Sample Preparation

Tissue preparation and incubation. Aortas were isolated from 1- to 2-week old New Zealand White rabbits (Kuiper Rabbit Ranch, Gary, IN), placed in ice-cold HEPES buffer (in mM; 10 HEPES, 150 NaCl, 5 KCl, 2 $CaCl_2$, 1 $MgCl_2$, and 6 glucose; pH 7.4), cleaned of adhering connective tissue and fat, and cut into rings (5-mm long). Aortic rings were incubated for 10 min at 37 °C in HEPES buffer containing indomethacin (10^{-5} M). AA (10^{-4} M) was added, and the vessels were incubated for an additional 5 min. Calcium ionophore A23187 (2×10^{-5} M) was added, and the vessels were incubated for another 15 min. The reaction was stopped by the addition of ethanol to a final concentration of 15%. The incubation buffer was removed, acidified (pH < 3.5) with glacial acetic acid, and extracted on Bond Elut C-18 extraction columns as previously described [27–29]. The extracts were evaporated to dryness under a stream of N_2 and stored at -40 °C until further HPLC separation.

Separation of AA metabolites by HPLC. The extracted samples were first separated into fractions of metabolite groups by reverse-phase HPLC (Nucleosil-C18 column, 5 μm , 4.6 \times 250 mm) using water:acetonitrile mobile phase containing 0.1% glacial acetic acid. The program was a 40-min linear gradient from 50% acetonitrile in water to 100% acetonitrile [28]. The fractions (0.2 mL per fraction) corresponding to the THETAs (fractions 27–35; 5–7.5 min) were collected and extracted with

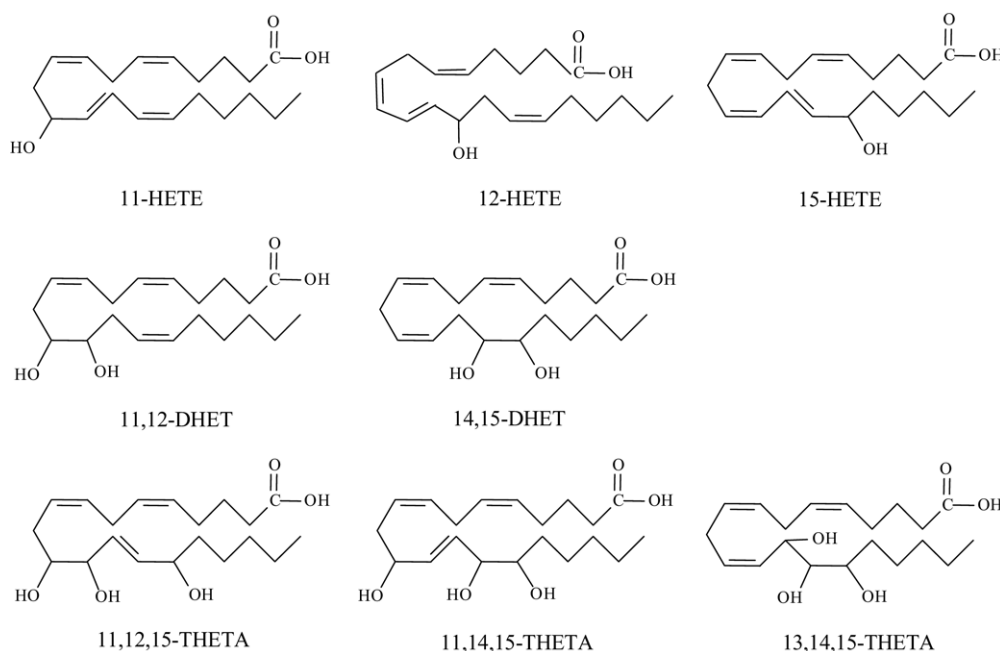


Figure 1. Structures of HETEs (11-HETE, 12-HETE and 15-HETE), DHETs (11,12-DHET and 14,15-DHET), and THETAs (11,12,15-THETA, 11,14,15-THETA and 13,14,15-THETA).

50:50 cyclohexane:ethyl acetate. The extract was dried under a stream of N_2 and redissolved in the HPLC mobile phase. The THETA fraction was rechromatographed on reverse-phase HPLC using water:acetonitrile (containing 0.1% glacial acetic acid) mobile phase. The program consisted of a 5-min isocratic phase with 35% acetonitrile in water, followed by a 35-min linear gradient to 85% acetonitrile [28]. The fractions that contained the THETAs (fractions 87–93; 17.5–18.5 min) were collected, acidified with acetic acid, and extracted with a 50:50 cyclohexane:ethyl acetate. The samples were dried under a stream of N_2 and analyzed by LC-FTICR.

Fourier Transform Ion Cyclotron Resonance Mass Spectrometry

For the standards, the experiments were performed on a 7.0 tesla FTICR (IonSpec, Lake Forest, CA) with a

Z-spray ESI source (Waters Corporation, Milford, MA) and a Model 22 syringe pump (Harvard, Holliston, MA) or a high-performance liquid chromatograph (HPLC, Agilent 1100 series, Palo Alto, CA). Standards were diluted in 50% acetonitrile:water solution. The standards were either directly infused by the syringe pump at a rate of $3 \mu\text{L}/\text{min}$ or introduced by flow injection ($1 \mu\text{L}$) through HPLC (Agilent 1100 series, Palo Alto, CA) into the electrospray source. FTICR was externally calibrated using 14,15-EET, L-ascorbic acid, and arachidonoyl dopamine. Although FTICR can detect HETEs and DHETs at 5 pg and THETAs at 15 pg, the MS/MS spectra were measured with HETEs and DHETs at 100 pg and THETAs at 500 pg. These concentrations were used to assure that the low abundant ions have adequate signals and they were accurately measured. Mass spectra were acquired in negative mode with ESI probe and source temperature of 80°C . The sample cone

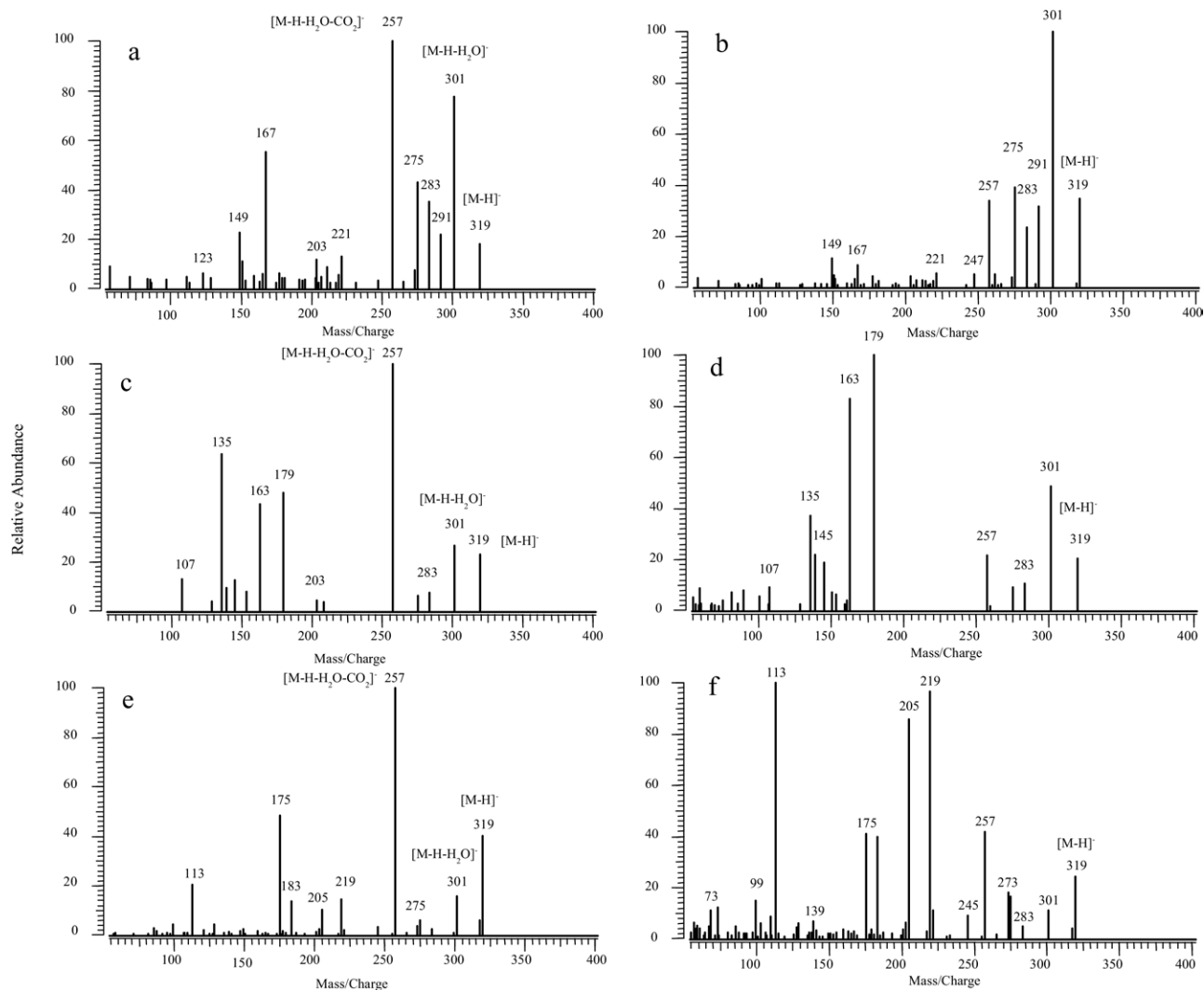


Figure 2. MS/MS spectra of $[M - H]^-$, m/z 319 for HETEs (100 pg) obtained from negative ion ESI SORI-CID and IRMPD FTICR. (a) MS/MS spectrum of 11-HETE by SORI-CID; (b) MS/MS spectrum of 11-HETE by IRMPD; (c) MS/MS spectrum of 12-HETE by SORI-CID; (d) MS/MS spectrum of 12-HETE by IRMPD; (e) MS/MS spectrum of 15-HETE by SORI-CID; (f) MS/MS spectrum of 15-HETE by IRMPD.

Table 1. High-resolution accurate mass measurements by SORI-CID FTICR for HETEs

Measured ions (m/z)	Calculated ions (m/z)	Error (ppm)	Relative abundance (%)	Elemental composition
11-HETE				
319.22872	319.22787	2.7	17.83	$C_{20}H_{31}O_3^{-1}$
301.21788	301.21730	1.9	77.16	$C_{20}H_{29}O_2^{-1}$
275.23886	275.23804	3.0	42.94	$C_{19}H_{31}O^{-1}$
257.22796	257.22747	1.9	100.00	$C_{19}H_{29}^{-1}$
207.17625	207.17544	3.9	4.95	$C_{14}H_{23}O^{-1}$
167.10801	167.10775	1.6	52.71	$C_{10}H_{15}O_2^{-1}$
149.09752	149.09719	2.2	22.75	$C_{10}H_{13}O^{-1}$
123.11851	123.11792	4.8	6.43	$C_9H_{15}^{-1}$
12-HETE				
319.22722	319.22787	-2.0	23.16	$C_{20}H_{31}O_3^{-1}$
301.21653	301.21730	-2.6	26.61	$C_{20}H_{29}O_2^{-1}$
257.22674	257.22747	-2.8	100.00	$C_{19}H_{29}^{-1}$
179.10724	179.10775	-2.8	48.37	$C_{11}H_{15}O_2^{-1}$
163.11239	163.11284	-2.8	43.48	$C_{11}H_{15}O^{-1}$
135.11756	135.11792	-2.7	63.85	$C_{10}H_{15}^{-1}$
15-HETE				
319.22738	319.22787	-1.5	42.76	$C_{20}H_{31}O_3^{-1}$
301.21563	301.21730	-5.5	17.34	$C_{20}H_{29}O_2^{-1}$
257.22673	257.22747	-2.9	100.00	$C_{19}H_{29}^{-1}$
219.13863	219.13905	-1.9	15.96	$C_{14}H_{19}O_2^{-1}$
175.14887	175.14922	-2.0	48.96	$C_{13}H_{19}^{-1}$
113.09705	113.09719	-1.2	20.82	$C_7H_{13}O^{-1}$

voltage was -45 V, the extractor cone was 10 V, Q1/Q2 RF was 70 V, and arbitrary waveform amplitude was 125 V. The probe high voltage was 3500 V. Transient signals were collected with 1024 K data points, an

analog-to-digital converter (ADC) rate of 4 MHz, and the transient length of 65.5 ms. The detection range was set at m/z 54–500. Nitrogen was used as the collision gas and cooling gas. The precursor ions were isolated in

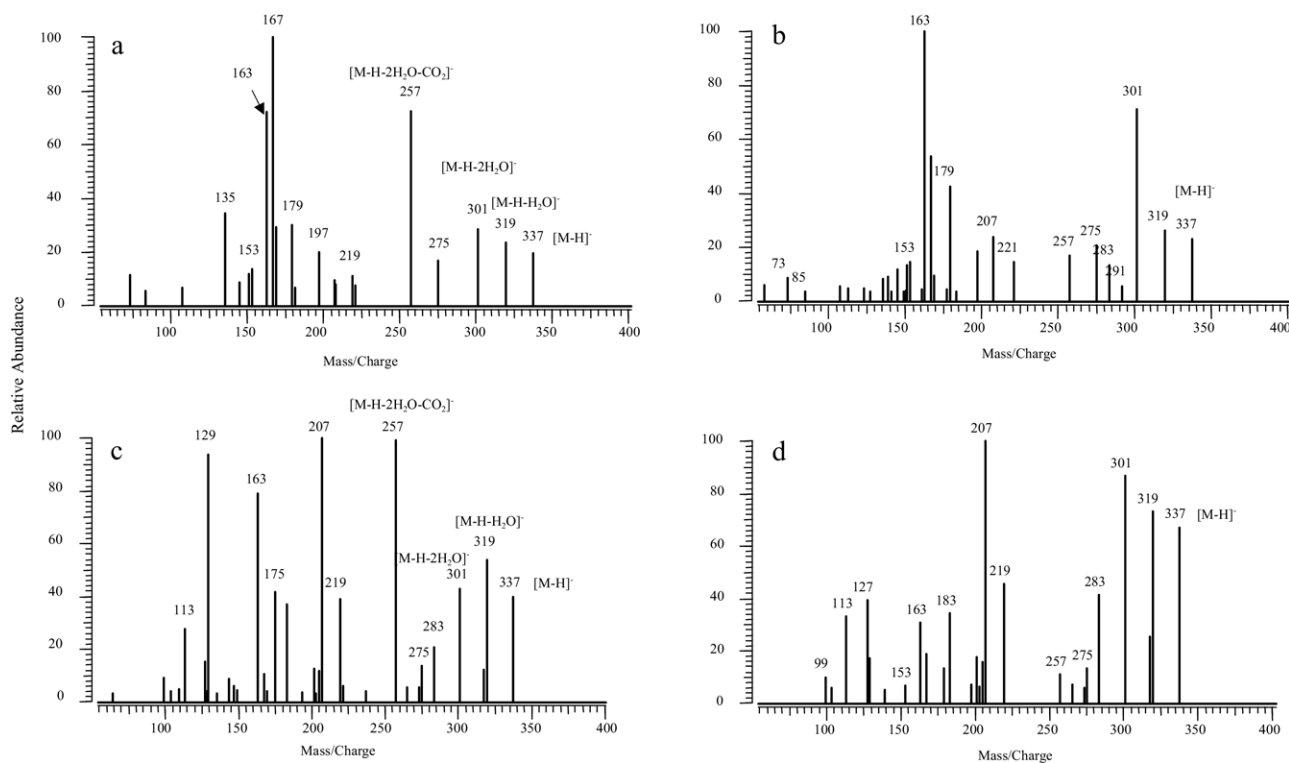


Figure 3. MS/MS spectra of $[M - H]^{-}$, m/z 337 for DHETs obtained from negative ion ESI SORI-CID and IRMPD FTICR. (a) MS/MS spectrum of 11,12-DHET by SORI-CID; (b) MS/MS spectrum of 11,12-DHET by IRMPD; (c) MS/MS spectrum of 14,15-DHET by SORI-CID; (d) MS/MS spectrum of 14,15-DHET by IRMPD.

the ICR cell and dissociated by SORI-CID or IRMPD. SORI-CID was initiated by opening a pulsed valve to admit a SORI burst with a length of 300 ms, offset frequency of 4563 Hz, amplitude of 15.5 V, and gas pulse of 15 ms. IRMPD was conducted with a 75 W continuous wave CO₂ laser at 95% power pulse, length of 300 ms, and gas pulse of 5 ms. Omega (version 8.0.201, IonSpec) software was used to control the instrument and analyze the data.

LC-FTICR Analysis of THETAs

LC-FTICR was performed with a 7.0 tesla FTICR coupled to an Agilent 1100 series liquid chromatograph. Both THETA standards and THETA fractions were analyzed by LC-FTICR. The THETA standards were diluted in acetonitrile (1000 pg/μL), and the THETA sample fraction was dissolved in 50 μL of acetonitrile. All HPLC separations were performed at ambient temperature. The samples were analyzed on a reverse-phase C₁₈ column (Kromasil 1.0 × 150 mm 5 μm; Varian) using water:acetonitrile containing 0.005% acetic acid as a mobile phase at a flow rate of 100 μL/min. The injection volume was 1 μL. The mobile phase gradient started at 20% acetonitrile in water, and linearly increased to 44% acetonitrile over 40 min, and then linearly increased to 100% in 5 min. After 25 min of run time on the LC, the FTICR was turned on to acquire data. The FTICR conditions were the same as described above for the standards.

Results and Discussion

The structures of three HETEs, two DHETs, and three THETAs are shown in Figure 1. Mass spectra of these eicosanoids exhibited carboxylate molecular ions [M – H][–] with *m/z* 319 for HETEs, *m/z* 337 for DHETs, and *m/z* 353 for THETAs as the most abundant ions. For MS/MS experiments, the molecular ions were isolated, accelerated toward the ICR cell, and dissociated. Some similar fragmentation patterns of HETEs, DHETs, and THETAs were observed, indicating the common backbone structures among these compounds. However, there were unique fragmentations reflecting the structural characteristics that can be used to identify these compounds.

Common Fragmentation Pathways for HETEs, DHETs, and THETAs

MS/MS spectra of all compounds indicated losses of H₂O and CO₂ to form a series of ions. For HETEs (Figure 2 and Table 1), which consist of one hydroxyl group, *m/z* 301 corresponding to [M – H – H₂O][–], was observed in all SORI-CID and IRMPD spectra. The *m/z* 257 was formed by a loss of CO₂ from [M – H – H₂O][–]. DHETs with two hydroxyl groups, as shown in Figure 3 and Table 2, the molecular ions (*m/z* 337) lost two H₂O molecules one after another, to form *m/z* 319 and 301, assigned as [M – H – H₂O][–] and [– H – 2H₂O][–], respectively. These ions could further lose CO₂ to form [M – H – H₂O – CO₂][–] (*m/z* 275) and [M – H – 2H₂O – CO₂][–] (*m/z* 257), respectively. The loss of H₂O and CO₂ are the major fragmentation pathways for

Table 2. High-resolution accurate mass measurements by SORI-CID FTICR for DHETs

Measured ions (<i>m/z</i>)	Calculated ions (<i>m/z</i>)	Error (ppm)	Relative abundance (%)	Elemental composition
11,12-DHET				
337.23729	337.23843	–3.4	20.04	C ₂₀ H ₃₃ O ₄ ^{–1}
319.22748	319.22787	–1.2	22.90	C ₂₀ H ₃₁ O ₃ ^{–1}
301.21718	301.21730	–0.4	27.63	C ₂₀ H ₂₉ O ₂ ^{–1}
275.23789	275.23804	–0.5	15.95	C ₁₉ H ₃₁ O ^{–1}
257.22634	257.22747	–4.4	70.41	C ₁₉ H ₂₉ ^{–1}
207.17419	207.17544	–6.0	9.09	C ₁₄ H ₂₃ O ^{–1}
197.11742	197.11832	–4.6	19.08	C ₁₁ H ₁₇ O ₃ ^{–1}
179.10725	179.10775	–2.8	30.77	C ₁₁ H ₁₅ O ₂ ^{–1}
169.12278	169.12340	–3.7	28.75	C ₁₀ H ₁₇ O ₂ ^{–1}
167.10714	167.10775	–3.7	100.00	C ₁₀ H ₁₅ O ₂ ^{–1}
163.11235	163.11284	–3.0	75.87	C ₁₁ H ₁₅ O ^{–1}
135.11769	135.11792	–1.7	36.80	C ₁₀ H ₁₅ ^{–1}
14, 15-DHET				
337.23868	337.23843	0.7	38.89	C ₂₀ H ₃₃ O ₄ ^{–1}
319.22762	319.22787	–0.8	54.05	C ₂₀ H ₃₁ O ₃ ^{–1}
301.21849	301.21730	4.0	42.51	C ₂₀ H ₂₉ O ₂ ^{–1}
275.23750	275.23804	–2.0	19.18	C ₁₉ H ₃₁ O ^{–1}
257.22729	257.22747	–0.7	98.43	C ₁₉ H ₂₉ ^{–1}
219.13885	219.13905	–0.9	38.19	C ₁₄ H ₁₉ O ₂ ^{–1}
207.13931	207.13905	1.3	100.00	C ₁₃ H ₁₉ O ₂ ^{–1}
175.14975	175.14922	3.0	40.49	C ₁₃ H ₁₉ ^{–1}
167.10810	167.10775	2.1	8.12	C ₁₀ H ₁₅ O ₂ ^{–1}
163.14947	163.14922	1.5	76.78	C ₁₂ H ₁₉ ^{–1}
129.09250	129.09210	3.1	92.71	C ₇ H ₁₃ O ₂ ^{–1}

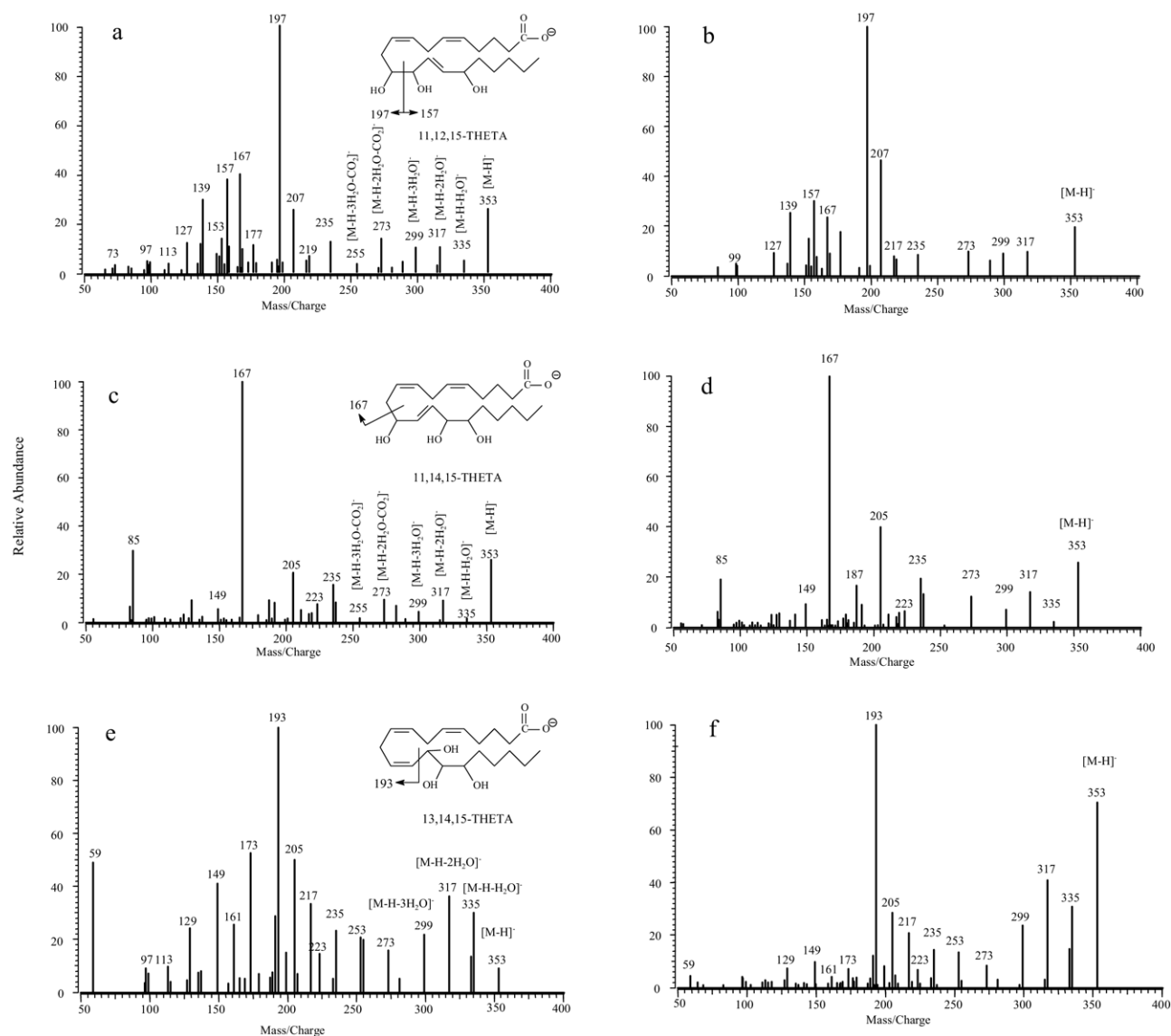


Figure 4. MS/MS spectra of $[M - H]^-$, m/z 353 for THETAs obtained from negative ion ESI SORI-CID and IRMPD FTICR. (a) MS/MS spectrum of 11,12,15-THETA by SORI-CID; (b) MS/MS spectrum of 11,12,15-THETA by IRMPD; (c) MS/MS spectrum of 11,14,15-THETA by SORI-CID; (d) MS/MS spectrum of 11,14,15-THETA by IRMPD; (e) MS/MS spectrum of 13,14,15-THETA by SORI-CID; (f) MS/MS spectrum of 13,14,15-THETA by IRMPD.

Table 3. High-resolution accurate mass measurements by SORI-CID FTICR for 11, 12, 15-THETA

Measured ions (m/z)	Calculated ions (m/z)	Error (ppm)	Relative abundance (%)	Elemental composition
353.23453	353.23335	3.3	25.57	$C_{20}H_{33}O_5^-$
335.22580	335.22278	8.9	4.53	$C_{20}H_{31}O_4^-$
317.21342	317.21222	3.8	10.20	$C_{20}H_{29}O_3^-$
299.20257	299.20165	3.1	9.90	$C_{20}H_{27}O_2^-$
273.22319	273.22239	2.9	13.59	$C_{19}H_{29}O^-$
255.21169	255.21182	-0.5	3.26	$C_{19}H_{27}^-$
235.13449	235.13397	2.2	12.17	$C_{14}H_{19}O_3^-$
207.13941	207.13905	1.7	25.31	$C_{13}H_{19}O_2^-$
197.11865	197.11832	1.7	100.00	$C_{11}H_{17}O_3^-$
167.10802	167.10775	1.6	39.96	$C_{10}H_{15}O_2^-$
157.12358	157.12340	1.1	38.32	$C_9H_{17}O_2^-$
153.12883	153.12849	2.2	14.43	$C_{10}H_{17}O^-$
139.11300	139.11284	1.2	30.20	$C_9H_{15}O^-$

Table 4. High-resolution accurate mass measurements by SORI-CID FTICR for 11, 14, 15-THETA

Measured ions (m/z)	Calculated ions (m/z)	Error (ppm)	Relative abundance (%)	Elemental composition
353.23520	353.23335	5.2	26.34	$C_{20}H_{33}O_5^{-1}$
335.22255	335.22278	-0.7	1.71	$C_{20}H_{31}O_4^{-1}$
317.21416	317.21222	6.1	9.01	$C_{20}H_{29}O_3^{-1}$
299.20354	299.20165	6.3	4.44	$C_{20}H_{27}O_2^{-1}$
273.22405	273.22239	6.1	9.48	$C_{19}H_{29}O^{-1}$
255.21371	255.21182	7.4	1.89	$C_{19}H_{27}^{-1}$
235.13527	235.13397	5.5	15.85	$C_{14}H_{29}O_3^{-1}$
223.13530	223.13397	6.0	7.50	$C_{13}H_{19}O_3^{-1}$
205.12465	205.12340	6.1	19.90	$C_{13}H_{17}O_2^{-1}$
167.10833	167.10775	3.5	100.00	$C_{10}H_{15}O_2^{-1}$
85.02952	85.02950	0.2	30.47	$C_4H_5O_2^{-1}$

HETEs and DHETs. In the SORI-CID and IRMPD spectra of m/z 353 of THETAs (Figure 4 and Tables 3, 4, and 5), which have three hydroxyl groups, the $[M - H]^-$ ions easily lost one to three H_2O molecules to give $[M - H - H_2O]^-$ (m/z 335), $[M - H - 2H_2O]^-$ (m/z 317), and $[M - H - 3H_2O]^-$ (m/z 299). After the losses of 2 or 3 H_2O , $[M - H - 2H_2O]^-$ and $[M - H - 3H_2O]^-$ could further dissociate to form $[M - H - 2H_2O - CO_2]^-$ (m/z 273) and $[M - H - 3H_2O - CO_2]^-$ (m/z 255), respectively.

Compared with HETEs and DHETs, the relative intensities of these ions (loss of H_2O and CO_2) in THETAs were lower than in HETEs and DHETs. The results suggest that the carbon-carbon bond ruptures became the major fragments of THETAs. In the comparison of three THETAs, the relative intensities of ions corresponding to the losses of H_2O and CO_2 for 13,14,15-THETA were slightly higher than for 11,14,15- and 11,12,15-THETA, suggesting that the positions of hydroxyl groups affected the losses of H_2O and CO_2 from the molecular ions. When three hydroxyl groups were adjacent, as with 13,14,15-THETA, it was easier to lose H_2O or CO_2 .

Unique Fragmentation Pathways for HETEs, DHETs, and THETAs

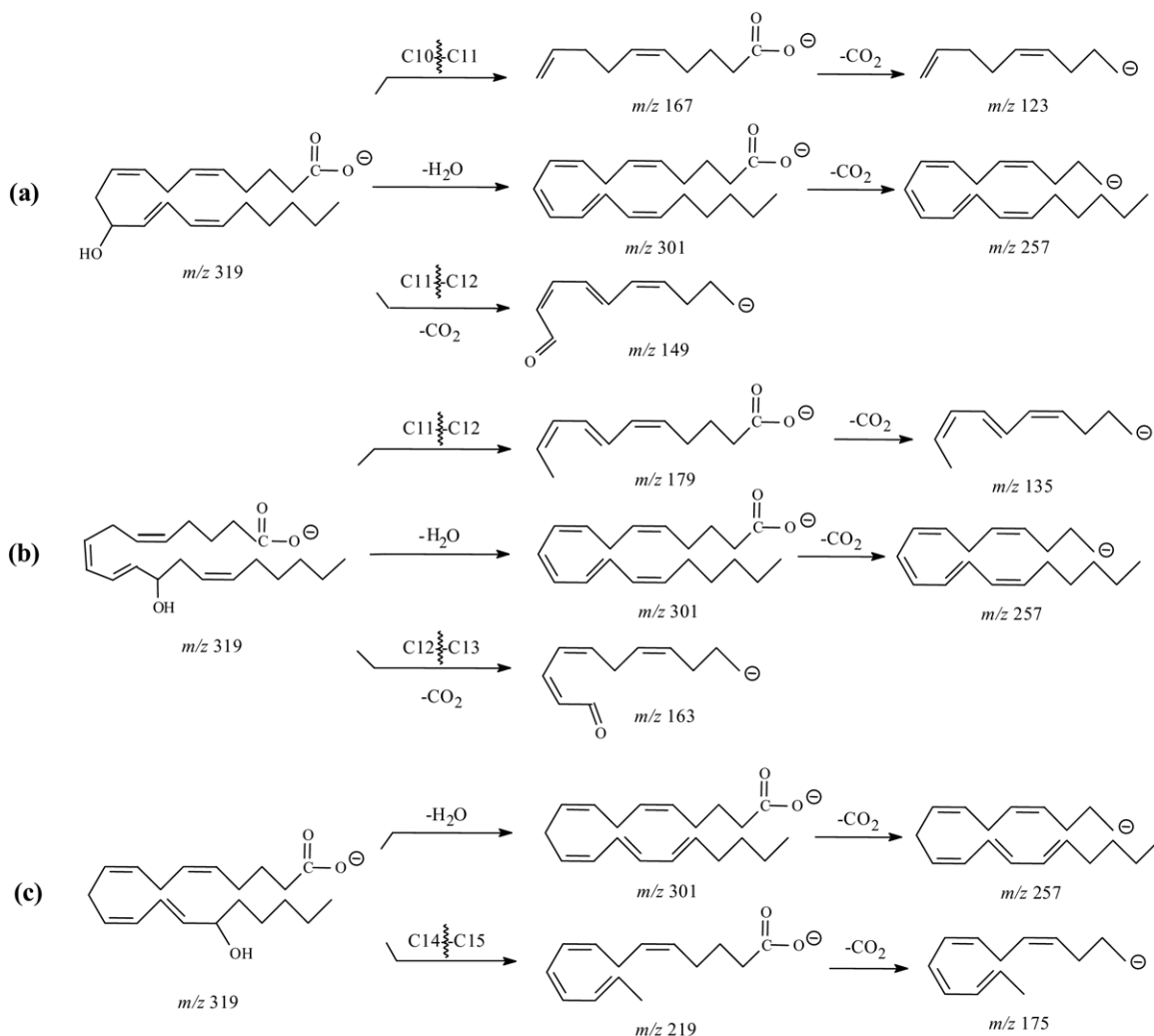
Besides the similarities of fragmentation, there were significant differences in fragmentations among HETEs, DHETs, and THETAs obtained from SORI-CID and IRMPD that indicated the characteristics of the molecular structures.

(a) MS/MS Characteristics of HETEs by SORI-CID and IRMPD.

(1) SORI-CID of HETEs. Figure 2a, c, and e show the MS/MS spectra obtained from SORI-CID FTICR for 11-, 12-, and 15-HETE, respectively. In the SORI-CID spectrum of $[M - H]^-$ for 11-HETE, the major characteristic product ions were m/z 167 and 149. The characteristic m/z 167 is similar to that reported fragmentation by triple quadrupole and ion trap mass spectrometers [33, 39, 41]. The MS/MS spectra for 11-HETE by a triple quadrupole mass spectrometer had the dominant ion of m/z 167 [33], while spectra from FTICR and ion trap [39] contained other characteristic ions. The mechanism for formation of m/z 167 was previously described [41] as the charge-remote formation of the aldehyde by proton

Table 5. High-resolution accurate mass measurements by SORI-CID FTICR for 13, 14, 15-THETA

Measured ions (m/z)	Calculated ions (m/z)	Error (ppm)	Relative abundance (%)	Elemental composition
353.23454	353.23335	3.4	9.16	$C_{20}H_{33}O_5^{-1}$
335.22447	335.22278	5.0	29.95	$C_{20}H_{31}O_4^{-1}$
317.21399	317.21222	5.6	36.52	$C_{20}H_{29}O_3^{-1}$
299.20348	299.20165	6.1	21.78	$C_{20}H_{27}O_2^{-1}$
273.22435	273.22239	7.2	15.69	$C_{19}H_{29}O^{-1}$
255.21344	255.21182	6.3	20.02	$C_{19}H_{27}^{-1}$
253.14608	253.14453	6.1	20.85	$C_{14}H_{21}O_4^{-1}$
235.13524	235.13397	5.4	23.23	$C_{14}H_{19}O_3^{-1}$
223.13486	223.13397	4.0	14.73	$C_{13}H_{19}O_3^{-1}$
217.12449	217.12340	5.0	33.61	$C_{14}H_{17}O_2^{-1}$
205.12447	205.12340	5.2	50.50	$C_{13}H_{17}O_2^{-1}$
193.12430	193.12340	4.7	100.00	$C_{12}H_{17}O_2^{-1}$
173.13440	173.13357	4.8	52.43	$C_{13}H_{17}^{-1}$
161.13442	161.13357	5.3	25.51	$C_{12}H_{17}^{-1}$
149.13423	149.13357	4.4	41.29	$C_{11}H_{17}^{-1}$
129.09271	129.09210	4.7	24.04	$C_7H_{13}O_2^{-1}$
59.01395	59.01385	1.7	49.36	$C_2H_3O_2^{-1}$



Scheme 1. Proposed major fragmentation pathways of 11-, 12- and 15-HETE by negative ion ESI-FTICR. (a) 11-HETE; (b) 12-HETE; (c) 15-HETE.

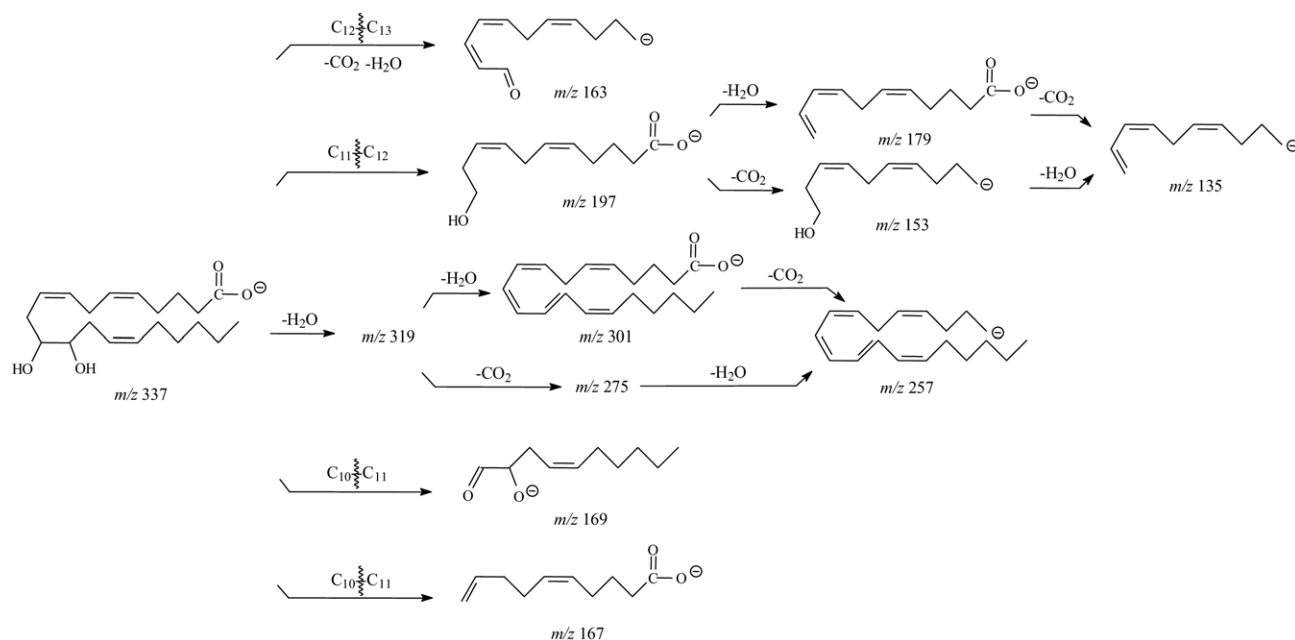
transfer and the cleavage of the C10-C11 bond. Another characteristic ion of the m/z 149 was observed and proposed as a cleavage of the C11-C12 bond and a loss of CO_2 . Another particular ion with high intensity was observed at m/z 275. It was formed by a direct loss of CO_2 from the $[\text{M} - \text{H}]^-$ ion. The losses of CO_2 and H_2O by $[\text{M} - \text{H}]^-$ ion formed the highest abundant m/z 257, although these two ions were not detected in the previous studies [33, 41] (Scheme 1a).

For 12-HETE, the major characteristic ions were m/z 179, 163, and 135 and were similar to the results from the ion trap mass spectrometer [39]. Again, the m/z 257 is the highest abundant ion. The mechanism for the m/z 179 formation was previously described [41]. We proposed that the negative charge was located at the carboxylate because the presence of the m/z 135 ion, corresponding to a loss of CO_2 from the m/z 179. The m/z 163 was presumably the result of carbon-carbon bond cleavage from C12-C13 and loss of CO_2 (Scheme 1b).

The characteristic product ions of 15-HETE were m/z 219, 175, and 113 and were similar to previous studies

[33, 39]. The m/z 219 was the product ion formed by the identical mechanism as previously described for the m/z 179 of 12-HETE [41]. The m/z 175 was the subsequent loss of CO_2 from the m/z 219 (Scheme 1c). The elemental composition of the m/z 113 indicated the fragment of $\text{C}_7\text{H}_{13}\text{O}^-$, suggesting the double-bond conjugation and cleavage of C13-C14 bond. The abundance of this ion increased in IRMPD (see below).

(2) IRMPD of HETEs. The major product ions generated from IRMPD were similar to those generated from SORI-CID, but the relative intensities of these ions were different. In IRMPD spectra of $[\text{M} - \text{H}]^-$ for all three HETEs (Figure 2b, d, and f), the relative intensity of m/z 257 decreased, while this ion was the highest abundant ion in SORI-CID spectra of HETEs. It indicated that the loss of CO_2 from the $[\text{M} - \text{H} - \text{H}_2\text{O}]^-$ ion for HETEs became less effective with IRMPD. For 12-HETE (Figure 2d), m/z 179 was the most abundant ion, indicating that the break of C11-C12 bond was the major pathway [41]. For 15-HETE (Figure 2f), m/z 219 and 113 became the major ions. The m/z 219 was formed



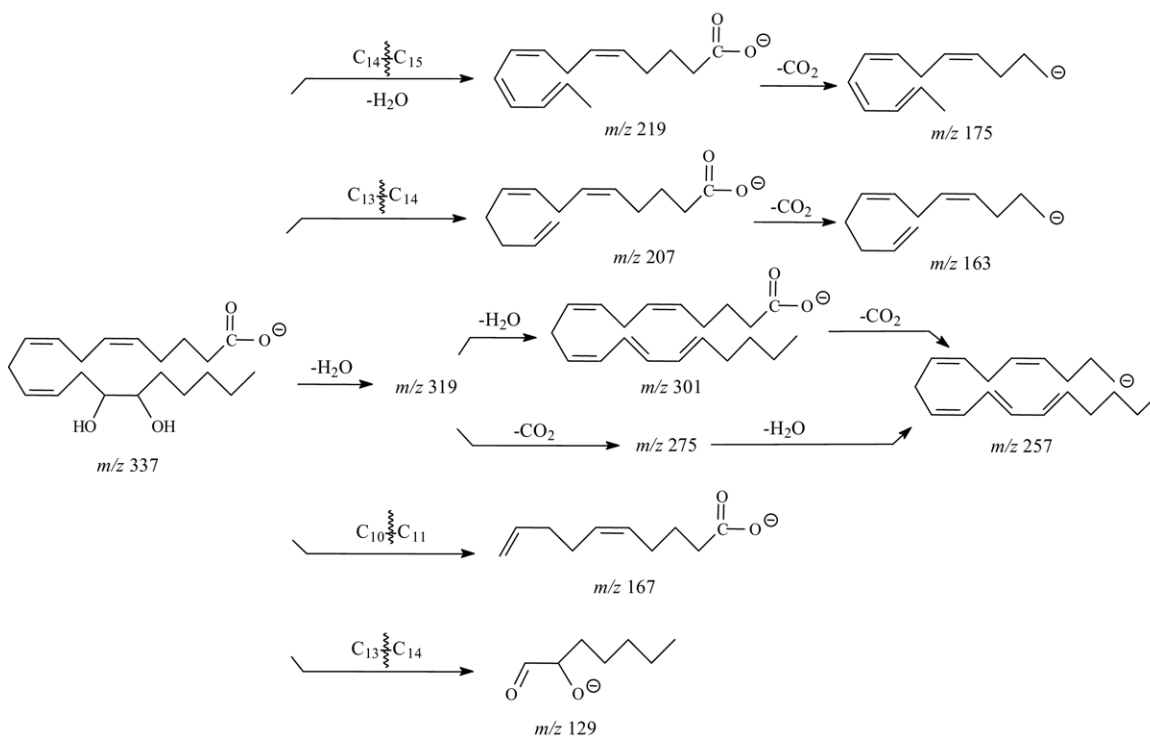
Scheme 2. Proposed major fragmentation pathways of 11,12-DHET by negative ion ESI FTICR.

by the break of C14–C15 bond and the charge was located on the carboxyl group. The formation of m/z 113 was the result of double-bond conjugation and the rupture of C13–C14 bond as in SORI-CID.

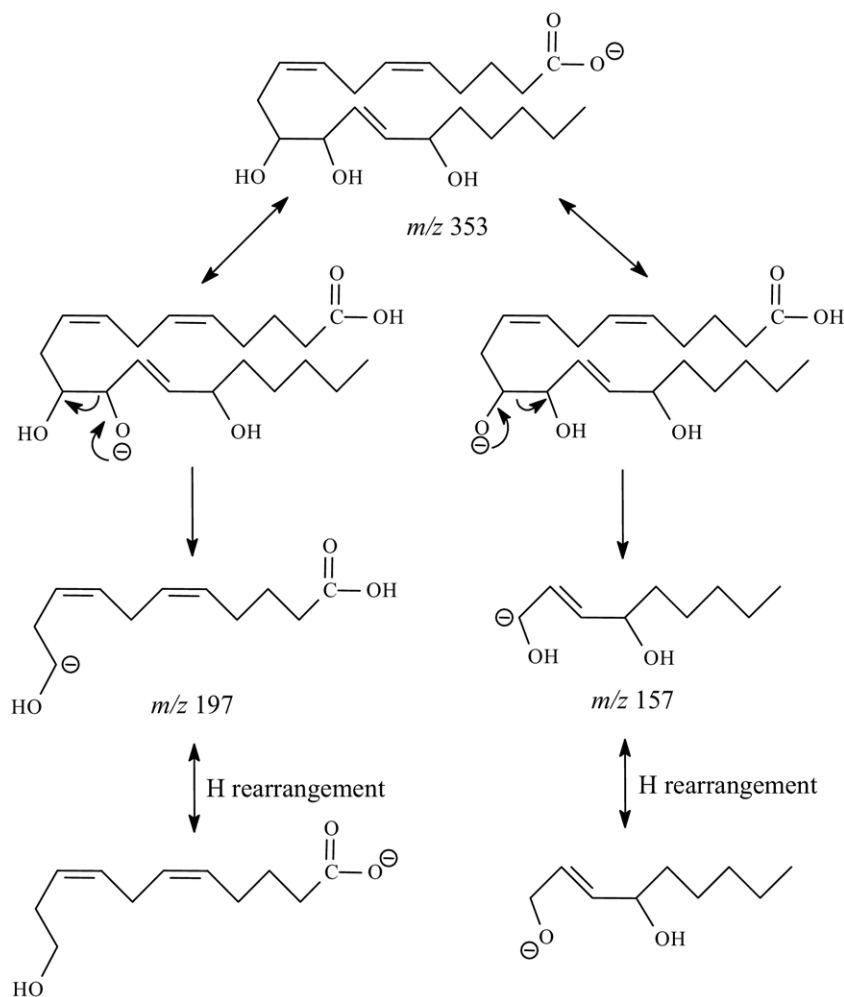
(b) MS/MS characteristics of DHETs by SORI-CID and IRMPD.

(1) SORI-CID of DHETs. In the SORI-CID spectrum of $[M - H]^-$, m/z 337, for 11,12-DHET is shown in

Figure 3a and the ions are shown in Table 2. The prominent m/z 167, 163, 135, 179, 197, 153, and 169 were observed and were similar to the results from ion trap mass spectrometer [39]. Formation of the highest abundant m/z 167 was similar to 11-HETE [41]. If the charge was relocated on the hydroxyl oxygen at C12, the m/z 169 could be formed by the same fragmentation mechanism. The m/z 197 was a result of the cleavage of



Scheme 3. Proposed major fragmentation pathways of 14,15-DHET by negative ion ESI FTICR.



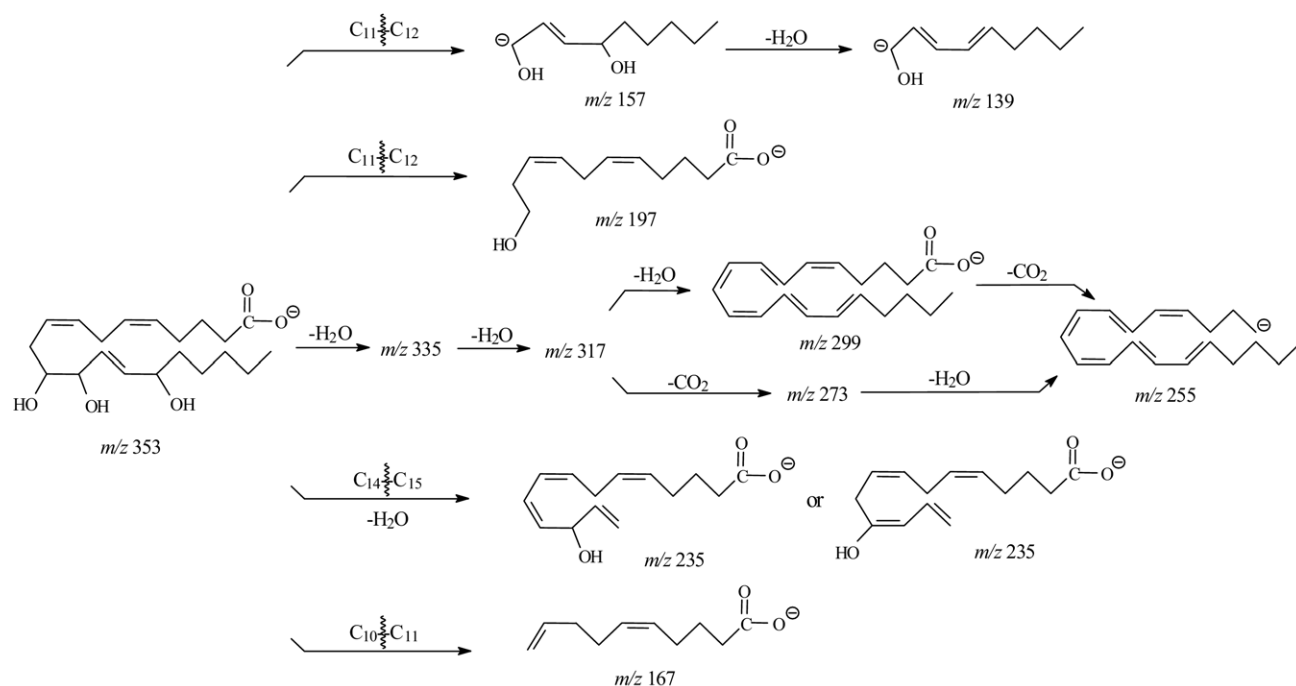
Scheme 4. Proposed mechanism of formation of the m/z 197 and m/z 157 in the negative ion SORI-CID spectrum of $[M - H]^-$ for 11,12,15-THETA.

C11–C12 bond [41]. Both m/z 179 and 153 corresponded to a loss of H_2O and CO_2 from the m/z 197, respectively. The m/z 135 was formed by a loss of CO_2 from the m/z 179 or a loss of H_2O from the m/z 153, respectively. The m/z 163 was derived from the cleavage of C12–C13 and the losses of H_2O and CO_2 . The major fragmentation pathways of 11,12-DHET were proposed in Scheme 2.

The CID spectrum of $[M - H]^-$, m/z 337, for 14,15-DHET had several characteristic ions (Table 2) that were similar to the results from the ion trap mass spectrometer [39] with the most abundant fragments of m/z 207 and 257 (Figure 3c). The m/z 257 was the result of the loss of $2H_2O$ and CO_2 from the $[M - H]^-$ ion. The m/z 207 was derived from the cleavage of the C13–C14 bond as previously described [39, 41]. In addition to m/z 207, other major ions were observed at m/z 129, 163, 175, and 219. The m/z 163 was from a loss of CO_2 from the m/z 207. If the charge was relocated on the hydroxyl oxygen at C15, m/z 129 can be formed at high abundance by the same dissociation as m/z 207. The m/z 219 resulted from a loss of H_2O and the charge-driven cleavage of the C14–C15 bond from the molecular ion as described for

15-HETE above. The subsequent loss of CO_2 by m/z 219 formed m/z 175 with high abundance. Some other low abundant ions were also observed in the spectra, such as m/z 127 derived from cleavage at the C7–C8 bond, m/z 167 derived from the cleavage at the C10–C11 bond. Scheme 3 illustrates the proposed dissociation reactions of 14,15-DHET.

(2) IRMPD of DHETs. The IRMPD spectra of $[M - H]^-$ for DHETs were similar to those formed by SORI-CID, indicating the same major fragmentation patterns. In the IRMPD spectra of $[M - H]^-$ for 11,12- and 14,15-DHETs (Figure 3b and d), the intensity of m/z 257 also decreased similar to HETEs as described above. The intensities of m/z 319 and 301, which were the results of the loss of one and two H_2O from the molecular ions, became higher, indicating easy losses of H_2O to produce $[M - H - H_2O]^-$ and $[M - H - 2H_2O]^-$. Interestingly, the m/z 163 instead of m/z 167 was observed as the high abundant ion for 11,12-DHET (Figure 3b) similar to 12-HETE. The results suggested that 11,12-DHET possibly lost H_2O at the C11 position and fragmented with the mechanism similar to 12-

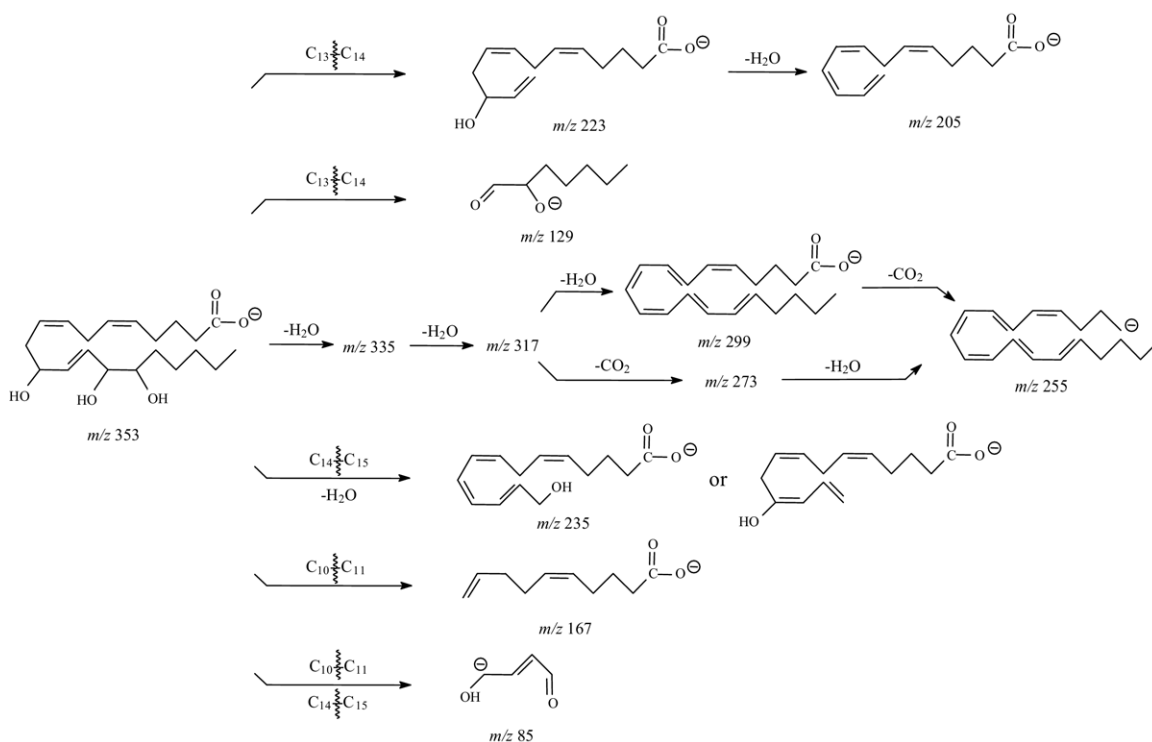


Scheme 5. Proposed fragmentation pathways of 11,12,15-THETA by negative ion ESI FTICR.

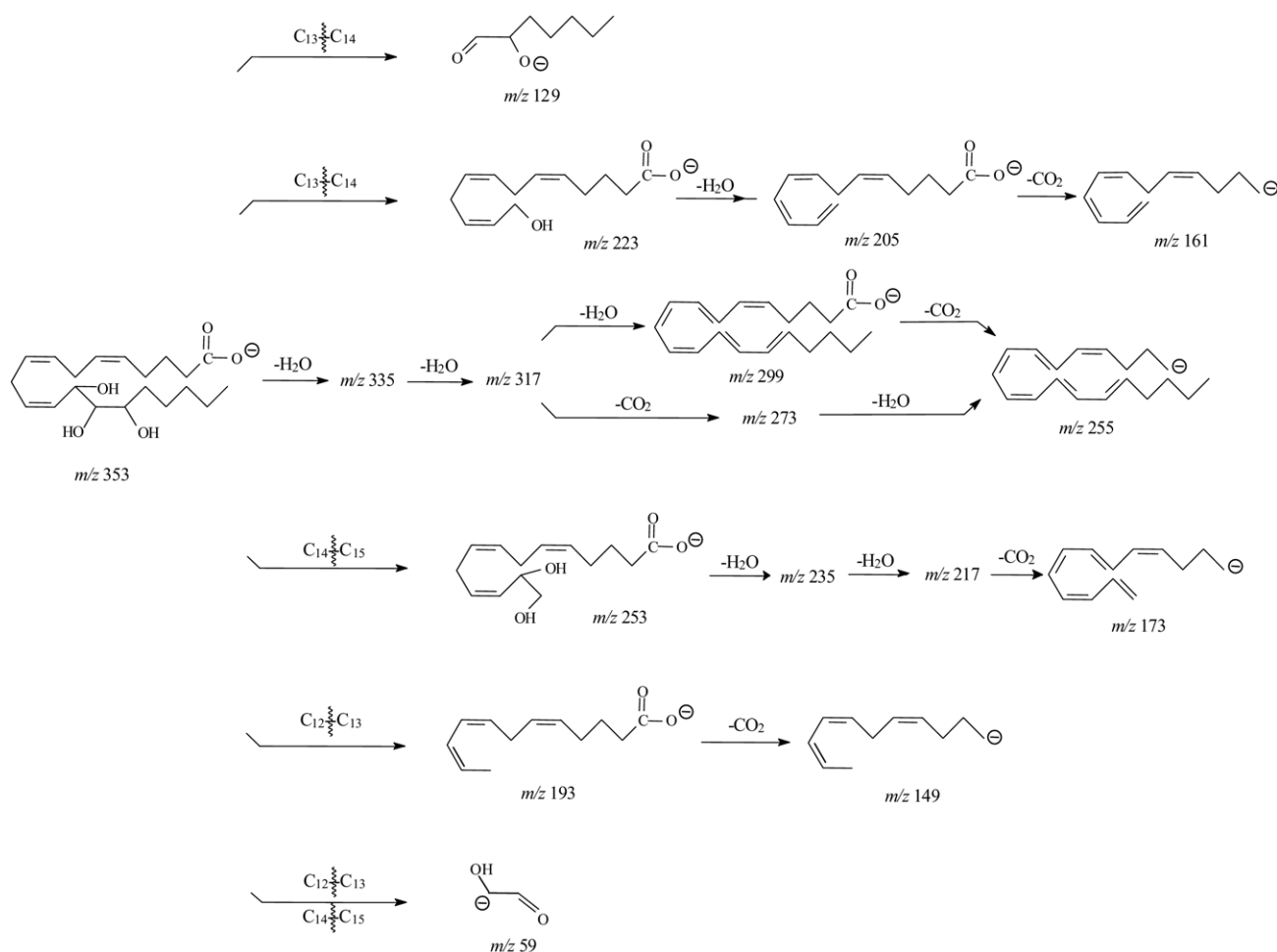
HETE. More low abundant ions were observed in the low m/z range of the IRMPD spectrum of 11,12-DHET, such as m/z 58, 123, 113, 177, and 139, indicating further fragmentation from excessive energy. For 14,15-DHET, the intensities of m/z 175, 163, and 129 became lower in the IRMPD spectrum with the m/z 207 as the highest

abundance ion (Figure 3d). This suggested the dominant C₁₃-C₁₄ cleavage.

In general, the major product ions in the CID spectra of HETEs and DHETs obtained from FTICR were similar to ions obtained from ion trap and triple quadrupole mass spectrometers [34–41]. However, more



Scheme 6. Proposed fragmentation pathways of 11,14,15-THETA by negative ion ESI FTICR.



Scheme 7. Proposed fragmentation pathway of 13,14,15-THETA by negative ion ESI FTICR.

intense and more characteristic fragmentation ions were obtained from FTICR similar to ion trap than triple quadrupole mass spectrometry. Losses of H_2O and CO_2 to form characteristic ions are more favorable in FTICR. Furthermore, the structures of these fragments can be ascertained based on the corresponding high-resolution and accurate mass data from FTICR. For example, the m/z 163 for 12-HETE and 11,12-DHET has the masses of 163.11,239 and 163.11,235, respectively. This ion has the elemental composition of $C_{11}H_{15}O^{-1}$ (Tables 1 and 2). However, the m/z 163 for 14,15-DHET has the mass of 163.14,947 with the elemental composition of $C_{12}H_{19}^{-1}$ (Table 2). These results indicated that the fragmentation mechanisms of the m/z 163 for 12-HETE/11,12-DHET, and 14,15-DHET are different as shown in Schemes 1, 2, and 3.

(c) MS/MS Characteristics of THETAs by SORI-CID and IRMPD.

(1) SORI-CID of THETAs. All THETAs exhibited identical $[M - H]^-$, m/z 353. Figure 4a, c, and e show the MS/MS spectra obtained from SORI-CID FTICR of 11,12,15-, 11,14,15-, and 13,14,15-THETA, respectively. Their high-resolution and accurate mass spectrometric data and corresponding predicted elemental composition

of their fragments are shown in Tables 3, 4, and 5, respectively. With three hydroxyl groups, the spectra of THETAs were more complex than spectra of HETEs and DHETs.

For 11,12,15-THETA, (Figure 4a and Table 3), the major ions were m/z 197, 167, 157, 139, and 127. The most abundant product ion was m/z 197. This ion was the result of the charge-driven cleavage of C11–C12 by the loss of neutral aldehyde from the enolate anion in which the charge was relocated on the hydroxyl oxygen at C12. If the charge was relocated on the hydroxyl oxygen at C11, the m/z 157 was formed by the similar fragmentation mechanism as shown in Scheme 4. The m/z 157 then lost H_2O to form m/z 139. The mechanism for formation of m/z 167 was the same as the ion of 11-HETE [41]. If the charge was relocated on the hydroxyl oxygen at C15, the m/z 235 was likely formed by the losses of H_2O and hexyl aldehyde from the cleavage of C14–C15 bond. Scheme 5 illustrates the proposed fragmentation pathways for 11,12,15-THETA.

The SORI-CID spectrum of $[M - H]^-$ for 11,14,15-THETA (Figure 4c and Table 4) was dominated by the m/z 167. The formation of the m/z 167 was the result of the double-bond conjugation and then, the break of

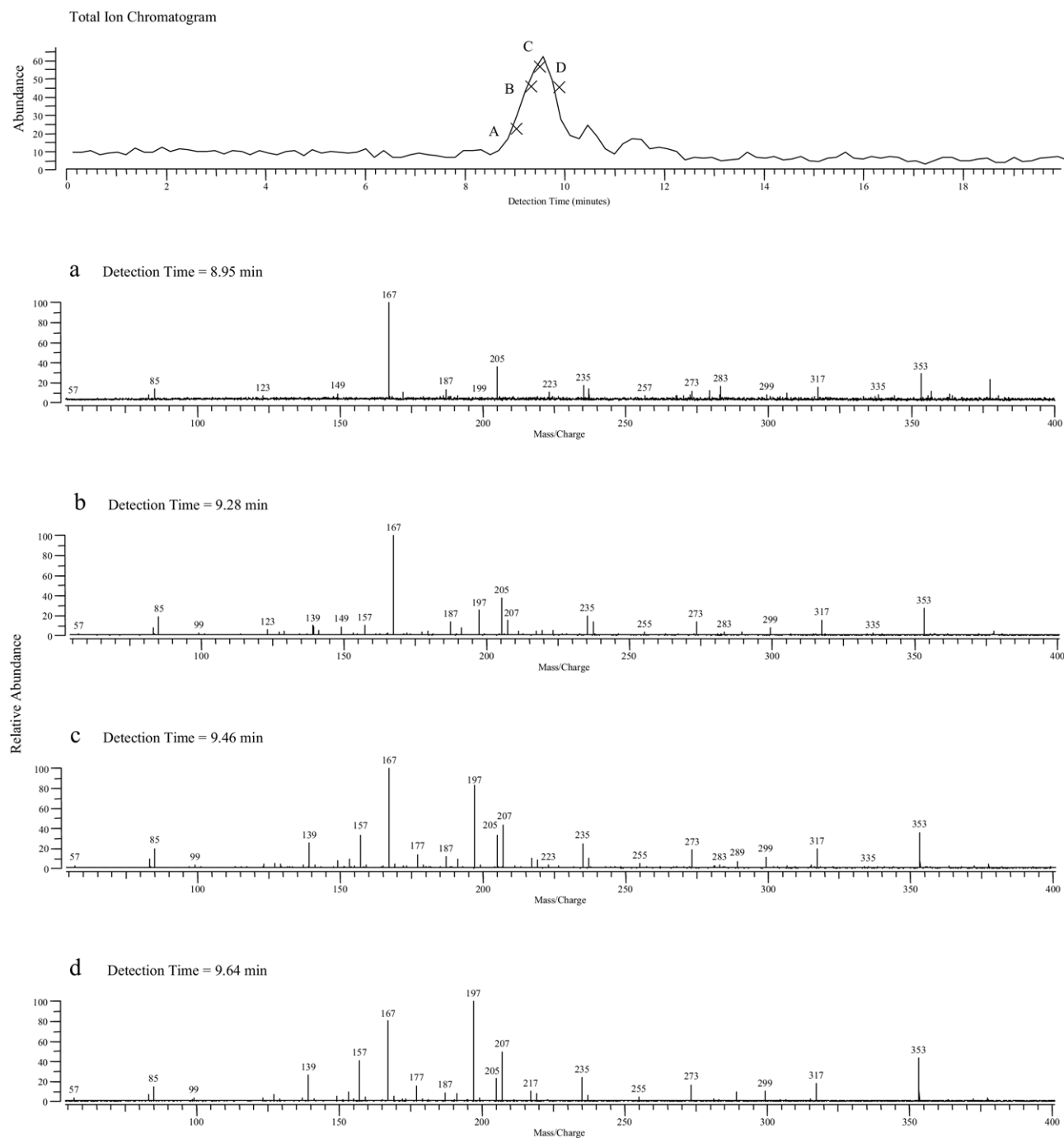


Figure 5. Total ion chromatogram (top panel) and product ion mass spectra of THETAs isolated from rabbit aorta by negative ion LC-ESI-FTICR with IRMPD. MS/MS spectrum of m/z 353 obtained from negative ion ESI FTICR at detection time of (a) 8.93 min; (b) 9.28 min; (c) 9.46 min; (d) 9.64 min.

C10–C11 bond as previously described for 11-HETE [41]. In addition, the unique characteristic m/z 85 was observed for 11,14,15-THETA. The elemental composition of this ion was assigned as $C_4H_5O_2^{-1}$. The proposed mechanism for its formation was the cleavages of C10–C11 bond and C14–C15 bond. Besides the m/z 167 and 85, the m/z 129, 205, 223, and 235 were also observed for 11,14,15-THETA. The proposed fragmen-

tation pathways for these ions were similar to DHETs as shown in Scheme 6.

When 11,12,15- and 11,14,15-THETA were compared, 11,12,15-THETA needed more collisional energy to fragment than 11,14,15-THETA. The different positions of the second hydroxyl group and the third double-bond in the structures of 11,12,15-THETA and 11,14,15-THETA resulted in a remarkable and diagnos-

Table 6. High-resolution accurate mass measurements by IRMPD FTICR for biological samples

Measured ions (<i>m/z</i>)	Calculated ions (<i>m/z</i>)	Error (ppm)	Elemental composition
353.23141	353.23335	−5.5	C ₂₀ H ₃₃ O ₅ ^{−1}
317.21112	317.21222	−3.5	C ₂₀ H ₂₉ O ₃ ^{−1}
299.20090	299.20165	−2.5	C ₂₀ H ₂₇ O ₂ ^{−1}
273.22148	273.22239	−3.5	C ₁₉ H ₂₉ O ^{−1}
235.13307	235.13397	−3.8	C ₁₄ H ₁₉ O ₃ ^{−1}
207.13821	207.13905	−4.1	C ₁₃ H ₁₉ O ₂ ^{−1}
205.12257	205.12340	−4.0	C ₁₃ H ₁₇ O ₂ ^{−1}
197.11747	197.11832	−4.3	C ₁₁ H ₁₇ O ₃ ^{−1}
167.10673	167.10775	−6.1	C ₁₀ H ₁₅ O ₂ ^{−1}
157.12265	157.12340	−4.8	C ₉ H ₁₇ O ₂ ^{−1}
139.11211	139.11284	−5.2	C ₉ H ₁₅ O ^{−1}
85.02898	85.02950	−6.1	C ₄ H ₅ O ₂ ^{−1}

tically useful difference in their major fragmentation pathways. The *m/z* 197, 207, 157, and 139 were the characteristic product ions with *m/z* 197 as the highest abundant ion for 11,12,15-THETA. The *m/z* 167, 205, and 85 were the characteristic product ions with *m/z* 167 as the most abundant ion for 11,14,15-THETA.

Figure 4e and Table 5 show the SORI CID spectrum and fragments of [M − H][−] for 13,14,15-THETA. The *m/z* 193 was observed as the most abundant ion, which was proposed to form by the double-bond conjugation and the break of C12–C13 bond similar to the formation of *m/z* 179 for 12-HETE [41]. This ion then lost CO₂ to form the *m/z* 149. Other product ions were *m/z* 253, 235, 223, 217, 205, 173, 161, and 129. If the charge was relocated at C15 oxygen group, the *m/z* 253 was formed by the charge-driven loss of neutral aldehyde from C14–C15 bond. The *m/z* 235, 217, and 173 were formed by the losses of H₂O or CO₂ from *m/z* 253. The *m/z* 223 and 129 were formed by breaking of the C13–C14 bond. The former ion lost H₂O to form the *m/z* 205, and then lost CO₂ to produce the *m/z* 161. The unique characteristic *m/z* 59 was observed for 13,14,15-THETA. The proposed formation of this ion was the cleavage of C12–C13 bond and C14–C15 bond similar to the *m/z* 85 for 11,14,15-THETA. The proposed fragmentation pathways for 13,14,15-THETA are shown in Scheme 7.

(2) IRMPD of THETAs. Figure 4b, d, and f show the product ion spectra obtained from IRMPD of 11,12,15-, 11,14,15-, and 13,14,15-THETA, respectively. The major product ions obtained from IRMPD were similar to SORI-CID, suggesting similar major fragmentation pathways of THETAs. However, there were some differences between SORI-CID and IRMPD spectra. For 11,14,15-THETA (Figure 4d), the intensities of the *m/z* 205 in the IRMPD spectrum increased 2-fold, and the abundance of *m/z* 85 decreased, compared with SORI-CID. IRMPD favored the *m/z* 205 by the cleavage of neutral aldehyde from C13–C14 bond and the subsequent loss of H₂O. This may be due to the product ion obtaining more energy from the laser, and then fragmentation to generate higher abundant *m/z* 205. For 11,12,15-THETA, the relative abundance of the *m/z* 207

increased with IRMPD. However, the abundance of the other product ions, such as *m/z* 157, 167, 177, 139, and 127, decreased. For 13,14,15-THETA, there were obvious differences between IRMPD and SORI-CID. The abundance of the *m/z* 59 and 173 decreased ten-fold in the IRMPD spectrum, and the other ions such as *m/z* 205, 149, 129, 217, 235, and 253 also decreased. These two techniques complemented each other, providing more useful structural information.

Identification of THETAs as Arachidonic Acid Metabolites in Rabbit Aorta Samples

The THETA fraction in rabbit aorta samples was analyzed by negative ion LC-FTICR to obtain structural information for identification of THETA isomers. The mass spectrum of the samples showed the most abundant ion of *m/z* 353 ([M − H][−]). The molecular weight was determined as 354 Da with the elemental composition of C₂₀H₃₄O₅.

Figure 5 (top panel) shows the total ion chromatogram of the MS/MS of *m/z* 353 by IRMPD. Figure 5a, b, c, and d show the product ions of *m/z* 353 obtained from different retention times by IRMPD. The *m/z* 335, 317, and 299 were observed, suggesting the subsequent losses of one to three H₂O from [M − H][−]. These ions indicated three OH groups were present in the structures. The low abundance of these ions also suggested that the positions of three OH groups were not adjacent. The changes in spectral patterns at different retention times (from a to d) suggested that the peak contained more than one compound. A complete separation of THETA regioisomers on a reverse phase LC column was not obtained.

These mass spectra revealed the presence of characteristic products, *m/z* 85, 167, 205, 197, 157, 139, and 207, which belonged to THETAs (Table 6). The *m/z* 167, 205, and 85 were the characteristics of 11,14,15-THETA, and the *m/z* 197, 157, 139, and 207 were the characteristics of 11,12,15-THETA. As shown in Figure 5a, the characteristics of 11,14,15-THETA, such as *m/z* 167, 205, and 85, were observed, but no *m/z* 197, 157, 139, and 207 were observed, suggesting that 11,14,15-THETA eluted first. With a longer eluting time (as shown in Figure 5b, c, and d), the *m/z* 197, 157, 139, and 207 appeared and the intensities of these ions increased, suggesting that 11,12,15-THETA eluted after 11,14,15-THETA.

Figure 6 shows the total ion chromatograms (top panel by IRMPD) and the selected ion chromatograms of *m/z* 85, 139, 157, 167, 197, 205, and 207 by both SORI-CID and IRMPD. The *m/z* 167, the major ion for 11,14,15-THETA and the minor ion for 11,12,15-THETA, exhibited a broad peak. According to the different retention times of these ions (run time of 25.00 min + detection time), the ions could be divided into two groups; a group of *m/z* 167, 205, and 85 (Figure 6c, e, g, and d, f, h) with identical detection times (9.44 min), suggesting that they were generated from one compound, 11,14,15-THETA, and another group of *m/z*

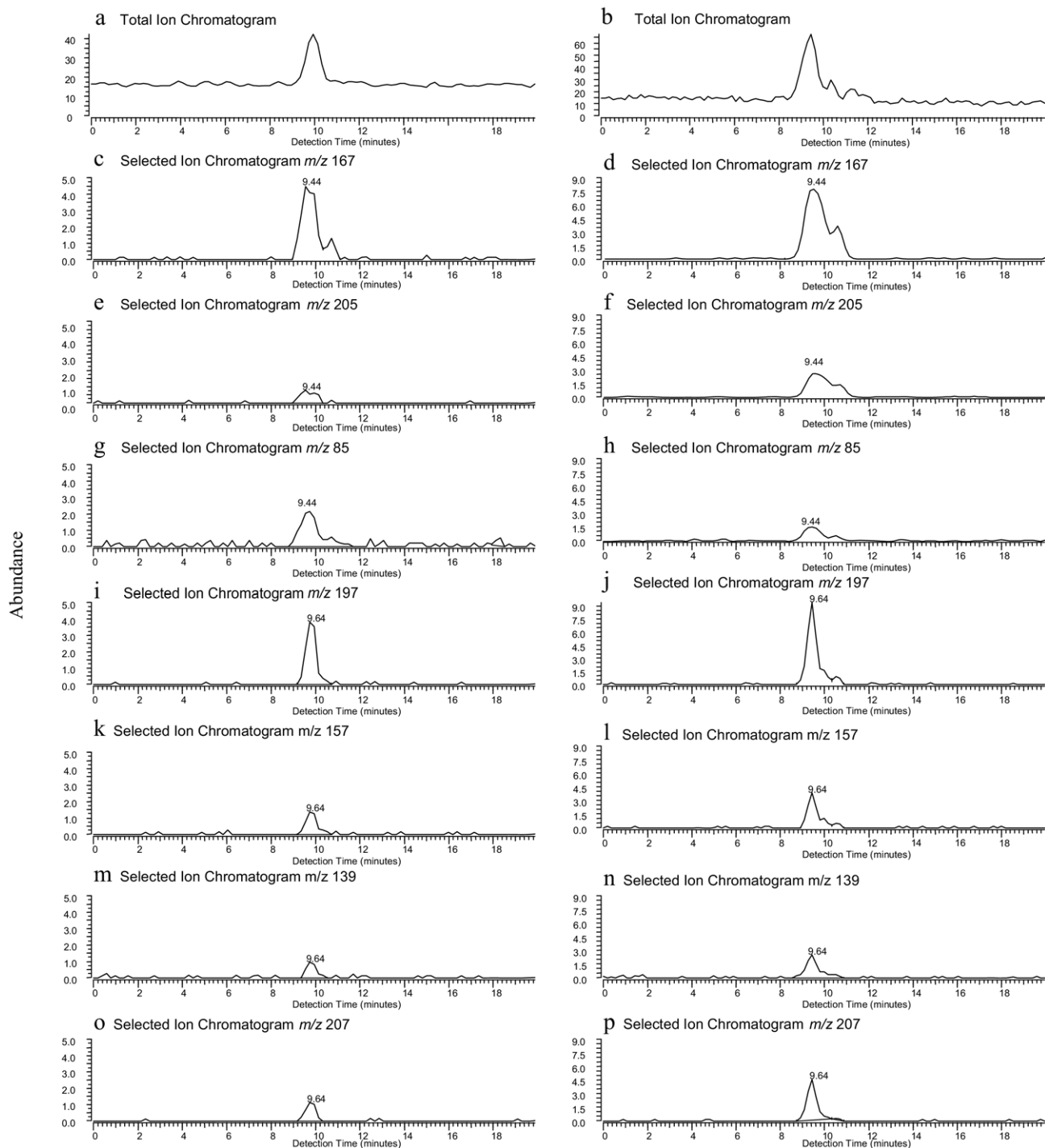


Figure 6. Total ion chromatograms and selected ion chromatograms of THETA fraction (top panels) isolated from rabbit aorta by negative ion LC-ESI-FTICR with SORI-CID and IRMPD. Total ion chromatograms by SORI-CID (a) and IRMPD (b); selected ion chromatogram of m/z 167 by SORI-CID (c) and IRMPD (d); selected ion chromatogram of m/z 205 by SORI-CID (e) and IRMPD (f); selected ion chromatogram of m/z 85 by SORI-CID (g) and IRMPD (h); selected ion chromatogram of m/z 197 by SORI-CID (i) and IRMPD (j); selected ion chromatogram of m/z 157 by SORI-CID (k) and IRMPD (l); selected ion chromatogram of m/z 139 by SORI-CID (m) and IRMPD (n); selected ion chromatogram of m/z 207 by SORI-CID (o) and IRMPD (p). Note: The indicated times were detection times (retention time = detection time + 25 min).

197,157, 13,9 and 207 (Figure 6i, k, m, o, and j, l, n, p) with identical detection times (9.64 min), suggesting that they were from 11,12,15-THETA (see also Figure 4a, b, c, and d). The m/z 205 also exhibited a small peak at a longer detection time (10.60 min). This ion is present in the spectra of both 11,14,15- and 13,14,15-THETA. However, the peak of the biological sample at 10.60 min did not have m/z 193 and the retention time did not match 13,14,15-THETA, suggesting the possibility of another THETA being present at very low concentration.

To further confirm the identity of these THETAs, a mixture of 11,12,15-THETA and 11,14,15-THETA was analyzed by LC-FTICR using the same conditions as the biological sample. The retention times and MS/MS spectra of biological THETAs were identical to the THETA standards. Comparing to the abundance of THETA standards, the concentration of THETAs in biological samples was ~ 800 pg on the column. The results indicated that both 11,12,15- and 11,14,15-THETA were produced in the rabbit aorta with 11,14,15-THETA being the major isomer.

Conclusions

Many isomers of low molecular weight, biologically active compounds such as eicosanoids can be present in biological samples. LC-MS and LC-MS/MS have been very successful in the identification and determination of eicosanoids with less complex molecular structures such as HETEs and DHETs. However, relatively newer groups of eicosanoids such as THETAs, which contain more hydroxyl groups at various positions on the arachidonic acid backbone, are more challenging. Chromatographic separation, particularly with reverse phase, of these compounds is difficult. Furthermore, the MS/MS fragmentation patterns of these compounds are very complex and not always useful for stereochemical identification.

Previously, eicosanoids were successfully identified by GC-MS. The substituents on the eicosanoids such as hydroxyl groups were typically derivatized to improve sample volatility. With increasing hydroxyl groups, the efficiency of chemical derivatization decreased. Thus, if LC-MS can accomplish the identification without the need for derivatization, it will be a powerful tool for these compounds. The SORI-CID and IRMPD in FTICR provide unique and useful spectral characteristics of highly substituted eicosanoids such as THETAs. Taking advantages of FTICR for its high mass resolution and accuracy, their fragments and molecular structures can be assigned. As demonstrated here, the regioisomers of the unresolved mixture of THETAs in biological samples were identified.

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