REVIEW

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Fatty acid isomerism: analysis and selected biological functions

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The biological functions of fatty acids and the lipids in which they are esterified are determined by their chain length, double bond position and geometry and other structural motifs such as the presence of methyl branches. Unusual isomeric features in fatty acids of human foods such as conjugated double bonds or chain branching found in dairy products, some seeds and nuts, and marine foods potentially have important effects on human health. Recent advancements in identifying fatty acids with unusual double bond positions and pinpointing the position of methyl branches have empowered the study of their biological functions. We present recent advances in fatty acid structural elucidation by mass spectrometry in comparison with the more traditional methods. The double bond position can be determined by purely instrumental methods, specifically solvent-mediated covalent adduct chemical ionization (SM-CACI) and ozone induced dissociation (OzID), with charge inversion methods showing promise. Prior derivatization using the Paternò–Büchi (PB) reaction to yield stable structures that, upon collisional activation, yield the double bond position has emerged. The chemical ionization (CI) based three ion monitoring (MRM) method has been developed to simultaneously identify and quantify low-level branched chain fatty acids (BCFAs), unattainable by electron ionization (EI) based methods. Accurate identification and quantification of unusual fatty acid isomers has led to research progress in the discovery of biomarkers for cancer, diabetes, nonalcoholic fatty liver disease (NAFLD) and atherosclerosis. Modulation of eicosanoids, weight loss and the health significance of BCFAs are also presented. This review clearly shows that the improvement of analytical capacity is critical in the study of fatty acid biological functions, and stronger coupling of the methods discussed here with fatty acid mechanistic research is promising in generating more refined outcomes.

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1. Introduction

Fatty acids are the major components of all cell membranes and serve as precursors of a wide array of signaling molecules. Although most human clinical fatty acid studies focus on 10–20 common structures,[1] a vast array of hydrocarbon fatty acid structures are found in nature, and each may have highly specific physiological effects.[2–4] Fatty acids of human foods have predominantly *cis* double bonds which are methylene-interrupted when two or more are present, and are straight chain. Unusual fatty acids are isomeric to common fatty acids, such as the well-known example of dairy products with conjugated linoleic acids and branched chain fatty acids, which are isomeric to linoleic acid and straight chain fatty acids, respectively.[2] Various edible although underutilized seeds such as melon, bitter melon and pomegranate seeds contain conjugated linolenic acids isomeric to both α-linolenic acid and γ-linolenic acid.[5] Pine and ginkgo nuts are rich in unusual polyunsaturated fatty acids (PUFAs), pinolenic acid, sciadonic acid and juniperonic acid, just to name a few.[6] The fatty acid double bond position, double bond geometry and methyl branching are major fatty acid structural motifs that determine their physiological functions. Analysis of food fatty acids can be particularly challenging, especially for foods originating from ruminants[7] or marine environments.[8]

The most common method for fatty acid analysis is by gas chromatography/mass spectrometry (GC/MS) analysis of fatty acid methyl esters (FAMEs) derived by derivatization of any lipid.[9] All fatty acyl groups esterified to any of the dozen or so lipid classes present in a typical food are converted to one

form, FAMEs, similar to the digestive process that converts to non-esterified fatty acids. Although to the analytical chemist this may seem like an undesirable structural modification and compromise, it is the most effective means to reveal the underlying details of the biology specific to mammalian fatty acid biochemistry.[10]

The analysis of fatty acid isomers, especially double bond positional isomers and branched structures, has long been an analytical challenge. Capillary GC coupled with either flame ionization detectors (FIDs) or MS is incapable of identifying fatty acid methyl esters (FAMEs) with varying double bond positions, though it can resolve *cis–trans* (*Z/E*) isomers. The conventional means for structural identification of metabolites is to obtain purified standards that can be analyzed side-by-side with unknowns to generate matched spectra. Although this approach applies to fatty acids of major abundance, conventional fragmentation methods do not enable the identification of double bond positions: in GC/MS based analysis of fatty acid methyl esters (FAMEs), electron ionization causes double bond migration; in collisional activation, fragments characteristic of double bond positions are not obtained.[11] Moreover, the number of FAME isomers for any empirical formula far exceeds the number of chemical standards that are available commercially, let alone can be practically maintained in a single laboratory. Consider for instance the number of isomers of methyl octadecenoate, represented by the chemical shorthand 18:1, where 18 is the number of carbons and 1 is the number of double bonds. This empirical formula is easily obtained from the molecular weight in MS-1. However, the double bond position can be in any of the 16 positions, from the 2–3 position to the 17–18 position. It may be *cis* (*Z*) or *trans* (*E*), and thus the number of double bond isomers is 32. Considering only single methyl chain branches, the number of chain branches extends to another 16 positions (all but COOH and CH_3), providing then for $32 \times 16 = 512$ isomers. Di- and tri-methyls and ethyl branching are also known, adding further potential isomers. Substantial numbers of these isomers are likely to be in food mixtures, especially those that have been treated by hydrogenation or by harsh processing. Similar considerations apply to FAMEs with 2–6 double bonds, common in foods, with the number of isomers rising exponentially with the number of double bonds.

This analysis clearly shows that general approaches are needed for *de novo* structural analysis of FAMEs. By "*de novo*", we refer to methods that reveal the structure of a FAME unequivocally with no need for genuine chemical standards. An ideal method will be rapid and produce strong diagnostic ions so that it can be applied to FAMEs of minor and trace abundance, as these may have important biological effects. Offline *de novo* methods have been in use since the early days of GC, first as methods such as ozone (O_3) decomposition methods with subsequent GC analysis, and later by specialized derivatives (*e.g.* picolinyl or 4,4-dimethyl oxazoline (DMOX)) that are amenable to *de novo* sequencing by GC/MS where the MS is a nominal resolution quadrupole.[11]

These methods demonstrated the key importance of fatty acid isomers but at the cost of substantial, concentrated effort, often devoted to obtaining a sufficient quantity of analytes to yield useful signals.

Because of the cumbersome and limited nature of existing methods, our laboratory developed two general *de novo* sequencing methods for FAMEs that identify almost all double bond and chain branching isomers.

Early lipidomics based on the introduction of intact, unmodified lipids typically by electrospray or atmospheric pressure chemical ionization routinely ignored fatty acid isomers, a practice that limited its applicability. Since about 2015, liquid chromatography (LC)-based lipidomics has focused on *de novo* methods for establishing structures in acyl chains, and has at this writing reached a stage where most isomers can be identified by one or the other methods[12–15] though some require offline preparation.[16,17] LC and GC methods tend to be complementary.

Here, we aim to illustrate the biological functions of some unusual fatty acids isomeric to some common fatty acids, stressing the importance of employing some state-of-the-art techniques for their accurate identification, streamlining the investigation of their biological and biochemical functions, and focusing on GC methods.

2. Analytical challenges of differentiating fatty acid isomerism

2.1 Traditional gas chromatography (GC) methods

GC-FID is the most widely used method for FAME analysis; however, structure identification relies on retention time matching of authentic standards. Thus, their accuracy is directly related to column specifications such as length/polarity and the availability of authentic standards. Polar columns such as CP-Sil 88 and BPX-70 are superior in separating FAME isomers, which provides GC a clear advantage in fatty acid analysis compared to LC. Under specially designed GC conditions, such as a 200 m ionic GC column for the analysis of *cis* and *trans* fatty acid isomers in fried foods, high separation efficiency can be observed for most fatty acid isomers.[18] However, coelution is still common for complex samples and MS analysis must be employed for accurate identifications.

GC-electron ionization (EI) MS data can be compared to comprehensive libraries, resulting in more confident identifications relative to GC-FID. The NIST EIMS library is a highly accessible and useful database which archives 0.3 million compounds. Unfortunately, GC-EIMS direct analysis of FAMEs similarly cannot differentiate double bond isomers. Notably, however, it can rule out many hypothetical structures based on the retention time, molecular weight, and common fragmentation.

Chemical ionization (CI) MS is also a traditional method for FAMEs. The most common CI method is a simple gas phase proton transfer, known as a "soft" ionization method because it transfers little excess energy to generate in-chain fragmentations, serving a high abundance of the molecular

ion. Therefore, the method is more suitable for the analysis of low-level fatty acids. Conversion of fatty acids into pentafluorobenzyl esters and analyzed by negative CI usually significantly reduces noise compared to positive CI because of suppression of all species that do not efficiently capture electrons, thereby leading to high sensitivity. Nevertheless, none of these traditional GC methods can differentiate fatty acid isomers without special derivatization treatments.

2.2 Derivatization methods for GC-EIMS analysis

Christie has reviewed methods for structural analysis of FAMEs by GC-EIMS.[19] We outline some of the more commonly used methods here.

2.2.1 DMOX. Fatty acid derivatization into DMOX esters for fatty acid structural elucidation is still a widely employed method due to its compatibility with GC-EIMS. Two main reasons drive the appeal of DMOX to many researchers. First, GC-EIMS is widely available and relatively low cost/minimum maintenance; second, the spectra of fatty acid DMOX derivatives can be matched against an EIMS comprehensive library, such as the NIST database, streamlining the fatty acid identification process.

Some of the limitations of DMOX derivatization and EIMS analysis include labor-intensive sample pretreatment and low sensitivity of diagnostic ions. In addition, fragments from DMOX derivatives do not yield unambiguous assignment of straight chain and branched chain structures. Importantly, DMOX and similar esterification regimes are challenging for minor-abundance fatty acids in complex mixtures because retention times of specialized esters change, sometimes unexpectedly compared to FAMEs, and thus matching FA-DMOX *vs.* FAME was difficult. Finally, the DMOX reaction itself, if not carefully executed, can cause double bond migration.

2.2.2 Picolinyl esters. Fatty acid picolinyl esters, upon EIMS activation, yield rich fragments including those that pinpoint double bond positions and methyl branch positions, making them more informative than the DMOX derivatization strategy. Picolinyl esters generally require column temperatures of about 50 °C higher than methyl esters, which makes the analysis of very long chain fatty acids (C ≥ 22) problematic.[19] Similar to DMOX, the picolinyl ester workflow requires complicated sample preparation and does not yield high-abundance diagnostic ions indicative of double bond positions or methyl branch positions.[20]

2.2.3 Dimethyl disulfide. Derivatizing FAME double bond (s) with dimethyl disulfide (DMDS) represents a convenient method to pinpoint double bond positions, with relatively high-abundance diagnostic ions specific to the erstwhile double bond positions. As a result, this method is more suitable for the discrimination of fatty acid double bond positional isomers in low abundance compared to DMOX or picolinyl ester approaches.

However, there are a few limitations of the DMDS method. First, it requires a very complicated sample derivatization procedure involving the use and cleanup of iodine (I$_2$). Second, the GC temperatures for DMDS analysis can be as high as
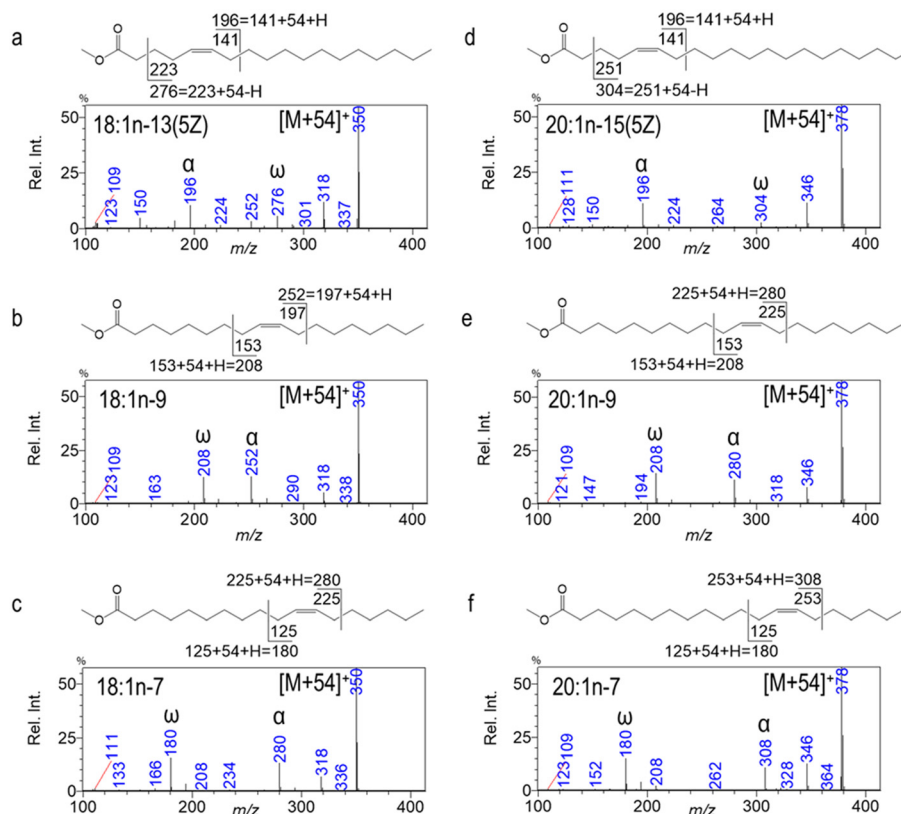
320 °C, exceeding the maximum temperature limit for most polar GC columns with high resolution. Third, the spectra of DMDS derivatives contain hundreds of ions, making the screening of diagnostic ions rather difficult especially for unusual polyunsaturated fatty acids.[21,22]

2.3 Emerging high-sensitivity, double bond derivatizing methods

2.3.1 Solvent-mediated covalent adduct chemical ionization (GC-MS). Solvent-mediated (SM) covalent adduct chemical ionization (CACI) is a state-of-the-art technique which enables structural elucidation of fatty acids in their FAME forms, and therefore greatly simplifies the sample preparation steps and is compatible with polar GC columns with a high resolving power. The technique takes advantage of a series of spontaneous reactions between acetonitrile and double bond(s) in a FAME hydrocarbon chain in the chemical ionization source, and upon collision induced dissociation (CID) yields diagnostic ions which are representative of the double bond positions.[6] The adduction reaction between the double bond in a fatty acid chain and the (1-methyleneimino)-1-ethenylium (MIE = 54; CH$_2$=C=N$^+$=CH$_2$) reactive ion is presented in Fig. 1. Depending on the number and position of double bonds, further CID activation usually results in vinylic or allylic cleavage on both sides of the erstwhile double bond position. The fragmentation rules for various FAME varieties have been worked out in several publications, with monoenes, dienes, polyenes with ≥3 homoallylic double bonds, conjugated FAMEs and polyenes with non-methylene-interrupted double bonds having nuances in spectral interpretations.[5–7,23–26]

SM-CACI enables *de novo* identification of common and unusual FAMEs without the need for authentic standards. For instance, Fig. 2 presents three isomers of 18:1 and 20:1, respectively. In this example, unusual Δ5 18:1 and 20:1 can be readily distinguished from their n-9 and n-7 counterparts in sea urchin gonads (edible parts), even without commercial standards.[23] 18:1n-13 has an α diagnostic ion at m/z 196 corresponding to an allylic cleavage at the far side of the carboxyl group. Here the number 141 is the mass of homolytic cleavage, and when the masses of the reactive ion (1-methyleneimino)-1-ethenylium (MIE = 54; CH$_2$=C=N$^+$=CH$_2$) and one hydrogen gain are accounted for, m/z 196 is observed on the SM-CACI-MS$_2$ spectrum (Fig. 2a). Similarly, a C$_2$–C$_3$ cleavage characteristic of monoenes with a Δ5 double bond leads to an ω diagnostic ion at m/z 276, having transferred a H$^•$ to the neutral leaving group during CID. Both n-9 and n-7

Fig. 1 CACI reaction forming a nitrogen containing ring and collision induced dissociation (CID) resulting in diagnostic ions pinpointing the erstwhile double bond position.

Fig. 2 CACI-MS/MS spectra of unusual Δ5 18:1 and 20:1; all [M + 54] → products. Reprinted with permission from Wang *et al.*[23] Copyright 2021 Elsevier Ltd.

isomers of 18:1 and 20:1 yield their diagnostic ion pairs allylic to both sides of the double bond, making their *de novo* identification the most straightforward (Fig. 2b, c, e and f). The SM-CACI-MS$_2$ spectra for the identification of polyunsaturated FAMEs and other unusual FAMEs by SM-CACI can be found in a series of previous publications.[5–7,23,24]

The SM-CACI technique is currently the only online derivatization mass spectrometric method coupled with a GC system, eliminating fatty acid manual derivatization. Importantly, it is the only method which allows the separation of FAMEs on the GC column before making FAME derivatives, making it highly compatible with any analytical method established on a GC-FID, and thus requiring no further offline chemistry. Briefly, the SM-CACI technique has the following advantages compared to conventional derivatization methods such as picolinyl ester: (A) no need to develop special chromatography for new esters, *i.e.*, higher temperatures. (B) Sample preparation is simple, and the preparation of FAMEs is the only procedure required. (C) Locating double bond positions of low-abundance fatty acids is possible due to its high diagnostic ion abundance and little other intrachain fragmentation.

2.3.2 Ozone induced dissociation (LC-MS). Ozone induced dissociation (OzID), first established by Blanksby and co-workers, has been extremely useful for the discrimination of phospholipid double bond positional isomers. It is also invaluable for fatty acid isomer identification when an LC system is

employed for their separation. Recently, a workflow involving the first derivatization of fatty acids into their *N*-(4-amino-methylphenyl)pyridinium (AMPP) derivatives followed by OzID activation was developed as a highly sensitive tool for fatty acid profiling and double bond position assignments.[27]

OzID generates very few intrachain fragments and high diagnostic ion abundance, enabling straightforward double bond position assignments. However, this online technique must be implemented on a customized MS instrument, compromising its easy access. An offline version was attempted by others[28] but as it does not enable mass selection before ozonolysis, it requires either pure samples or compatible mixtures. Offline ozonolysis of mixtures gives rise to extremely complicated spectra with ozonolyzed fragments originating from all fatty acids present in the original sample. LC, however, has a major advantage that sample preparation can be simpler for fatty acids, consisting of simple hydrolysis, and has no limitations for volatilization of low or high mass fatty acids. Generally, LC equipment is considerably more costly than GC, and GC chromatographic resolution can be greater.

2.3.3 Paternò–Büchi reaction (LC-MS). Paternò–Büchi reaction mass spectrometric analysis of lipid double bond positions was first developed by Xia and coworkers, serving as an alternative to OzID. It is also compatible with LC and generates very few intrachain fragments with high diagnostic ion abundance, enabling straightforward double bond position assign-
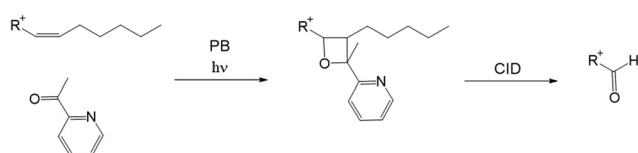
ments. The method has quickly become the most popular workflow for glycerolipid double bond position assignments. It has been adopted and adapted for various analyses, mainly due to its relatively simple derivatization apparatus – a UV lamp. It can be conveniently employed as a sample pre-treatment step or implemented into the LC-MS workflow as an online derivatization approach, as demonstrated by Xia *et al.*[29–32] and others.[33]

Fig. 3 presents the formation of the PB reaction adduct with 2-acetylpyridine as the derivatizing reagent. In the presence of UV light, a double bond on the acyl chain of a lipid molecule reacts with a ketone containing reagent, for instance, 2-acetylpyridine, acetone, *etc.*, and forms an oxygen containing ring called oxetane. Upon CID activation, site-specific cleavage at the erstwhile double bond position can be observed, and the fragments are used as the tool to pinpoint the double bond location. The resulting derivative is remarkably stable and thus can be manually admitted to the MS by infusion.

Considering its simple derivatization hardware requirements, the PB-MS/MS workflow is promising as a routine method to elucidate fatty acid isomeric features. Early work by Murphy *et al.* showed that the PB-MS/MS workflow did not result in highly interpretable spectra for polyunsaturated fatty acids, analyzed by their free form as negative ions.[34] However, further refining of the derivatization process yields highly interpretable PB-MS/MS spectra for polyenes including docosahexaenoic acid (DHA). Simply put, this was achieved either by double derivatization of the carboxyl group (*N,N*-diethyl-1,2-ethanediamine) and the double bond by PB (acetone),[35] or using a nitrogen containing PB reagent such as acetylpyridine,[33] or otherwise promoting the formation of a lithium adduct.[36] These methods allow fatty acids to be analyzed in the positive mode. While the CACI-MS/MS workflow tends to have greater sensitivity and simpler data analysis thanks to its compatibility with high resolution GC, the PB-MS/MS workflow possesses straightforward fragmentation patterns and simple mass spectra with high diagnostic ion abundances, offsetting LC's normally lower separation capacity.

The PB-MS/MS workflow was implemented for a specialized application, for quantitative analysis of deuteration levels at bis allylic positions of polyunsaturated fatty acids. In this particular case, GC-CACI-MS/MS was not possible due to the presence of companion ions (±1 Da or ±2 Da) of the CACI diagnostic ions.[17,37] This special case of analysis of a unique pharmaceutical-like species illustrates the importance of detailed mass spectral interpretation and mechanism for new analytes.
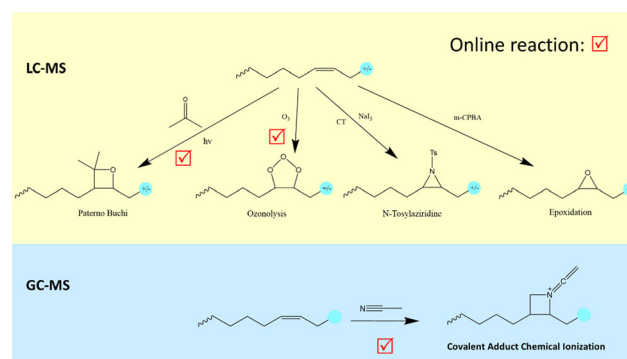
2.3.4 Other double bond derivatizing methods. There are also some other emerging derivatization methods similarly reacting with a double bond in a fatty acyl chain, yielding high-abundance diagnostic ions for convenient, highly sensitive and *de novo* identification of unsaturated fatty acid double bond variants, summarized in Fig. 4 along with the others. Among these is *N*-tosylaziridine derivatization, which replaces a double bond with a nitrogen containing cyclo-ring linked to some sulfur containing moieties. Similar to the CACI-MS/MS workflow, the positive charge remains with the nitrogen atom. During CID activation, bonds of the nitrogen containing cyclo-ring are all labile and tend to dissociate. Depending on which of the labile bonds is broken, two diagnostic ions indicating the erstwhile double bond position can be observed.[16]

Epoxidation is another double bond derivatizing strategy. Epoxidation can be induced by the *meta*-chloroperoxybenzoic acid (*m*-CPBA) reaction with double bond(s) in a fatty acyl chain at room temperature.[38] These two double bond derivatizing methods do not require any additional devices such as a UV lamp in the PB reaction or modifications to the MS instrument seen with CACI and OzID. However, they require sample offline pre-treatment, hampering their throughput compared to the online versions of CACI, PB and OzID methods. Fig. 4 summarizes the emerging double bond derivatizing MS methods and their compatible chromatographic coupling, which can be used to interrogate fatty acid structural information.

2.3.5 Charge switching chemistry. Charge switching chemistry, or ion/ion charge inversion chemistry was developed by McLuckey and coworkers, an innovative method for the release of fatty acyl chains from a glycerolipid molecule or for analysis of fatty acids themselves. Without structural modifications or adduction, fatty acids can only be analyzed in the negative mode. Previously, fatty acyl chains released from a glycerolipid molecule would be detected in the negative ion mode, which yields spectra dominated by H_2O and CO_2 neutral loss fragments with little structural information. Charge switching chemistry takes advantage of the reaction between tris-phenanthroline magnesium dications (+2 charge) and fatty acid anions (−1 charge) resulting in a cation (+1 charge), shown in

Fig. 3 Formation of a Paternò−Büchi reaction adduct with 2-acetylpyridine as the derivatizing reagent and formation of an aldehyde diagnostic ion.

Fig. 4 Emerging double bond derivatizing MS methods for fatty acid identification.

Fig. 5. This is a two-step reaction, *i.e.*, (1) ion/ion reaction yielding a complex with two phenanthroline ligands and (2) dipolar direct current (DDC) – CID removal of one of the two remaining phenanthroline ligands. Upon ion trap CID activation of the resulting fatty acid (single) phenanthroline magnesium ligand (Fig. 5), the cation yields a spectrum with rich fragmentation, at typically all carbon–carbon bond sites. A distinct difference in intensity near the double bond site or sometimes the presence of doublets at some cleavage sites indicates the location of the double bond(s). As the degree of unsaturation increases, the spectra become complicated.[39] Compared to the double bond derivatizing methods such as CACI, OzID, PB, *N*-tosylaziridine and epoxidation, the charge switching chemistry mass spectrometric method requires relatively in-depth expertise of the method for data interpretation. A more practical approach to exploit the merits of charge switching chemistry MS to identify fatty acid isomers involves the use of library matching, which has also been demonstrated.[40] Information covering most of the charge switching chemistry advancements can be found in a previous review.[41]

2.3.6 Photodissociation and electron impact excitation of ions from organics (EIEIO).

Besides various state-of-the-art derivatization and adduction methods capable of pinpointing double bond positions, there is an alternative strategy which seeks solution from innovative dissociation technologies built within the MS instrument, substituting it for CID or other conventional dissociation mechanisms. UV photodissociation utilizes short wavelength UVC to induce light-based dissociation of fatty acids and other lipids. In a pioneering study by Klein and Brodbelt, the structure of phosphatidylcholine (PC) was elucidated including the double bond positions.[42] Blanksby and coworkers adapted the principle and further demonstrated that with some suitable derivatizing reagents which are superior in absorbing the UV energy, monoene double bond positions could be revealed by manifestation of the fragmentation of an allylic bond to a double bond. Isomeric polyenes such as 18:3n-3 and 18:3n-6 could be readily distinguished by their clearly different spectra, however, straightforward localization of double bond positions is complicated by some rearrangement mechanisms around the double bond cleavages.[43]

Electron impact excitation of ions from organics (EIEIO), as an emerging ion fragmentation method, was developed recently. Its dissociation is also characterized by rich ion formation along the fatty acyl chain and a sharp decrease in ion abundance along the hydrocarbon chain indicates the presence of a double bond. This fragmentation pattern is reminiscent of the picolinyl ester-GC-EIMS method, with fragmentation conspicuously attenuated at the stronger double bond position. EIEIO was shown to elucidate structural elements, *e.g.*, double bond positions and *cis/trans* configurations of phosphatidylcholines, triglycerides and sphingomyelins.[44–46] Research is warranted to confirm that EIEIO is equally effective in determining fatty acid isomer compositions compared to methods further developed.

2.4 Identification of branched chain fatty acids (BCFAs)

2.4.1 Traditional picolinyl ester method by EIMS. BCFAs are common in certain foods,[47,48] bacteria,[49] as well as in some human tissue, notably skin.[50] Unlike for the double bond position, GC-EIMS provides information about branch points, though often the data are ambiguous.[51] Derivatization of fatty acids into picolinyl esters and analyzed by conventional GC-EIMS is the earliest method for effective identification of saturated BCFAs from their straight chain counterparts. Dating back to the 1980s, Harvey used picolinyl esters as derivatives to determine the monomethyl branch position along a fatty acid picolinyl ester chain.[20] Later Hierro *et al.* identified the anteiso-17:0 BCFA in ginkgo nuts taking advantage of the robust analytical capability of picolinyl esters.[52] Fig. 6 shows the positions of cleavage leading to high-abundance diagnostic ions, *i.e.* m/z 318 and m/z 346 for the identification of iso-17:0 and m/z 304 and m/z 332 for the identification of the anteiso-17:0 BCFA derivatized into picolinyl esters. Using anteiso-17:0 for illustration, there is no cleavage between the two respective cleavage positions due to the presence of a methyl branch at the butan-2-yl terminal position (negligible levels of m/z 318 in its spectrum).

2.4.2 Direct FAME analysis by EIMS/MS. Inspired by the work of Zirroli and Murphy,[53] Ran-Ressler *et al.* demonstrated that when tandem MS was available, a similar fragmentation pattern could be observed for anteiso-17:0 and other BCFA in their direct FAME form.[51] Fig. 7 presents the pair of high-abundance diagnostic ions m/z 227 and m/z 255 for the straightforward identification of anteiso-17:0 by GC-EIMS/MS. Identical to what is observed for the picolinyl ester analysis by GC-EIMS, the 14 Da (CH_2) interruption is missing between m/z 227 and m/z 255 (negligible levels of m/z 241) for EIMS/MS analysis.
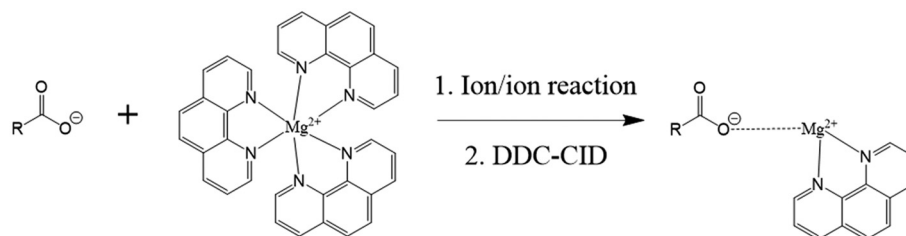

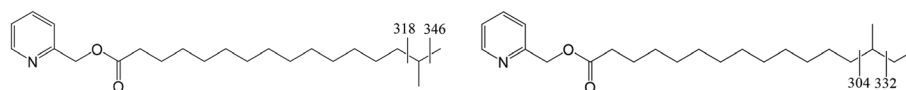**Fig. 5** Two-step formation of fatty acid phenanthroline magnesium ligands.

Fig. 6 Diagnostic ions for the identification of iso-17:0 and anteiso-17:0 BCFAs derivatized into picolinyl esters by GC–EIMS.
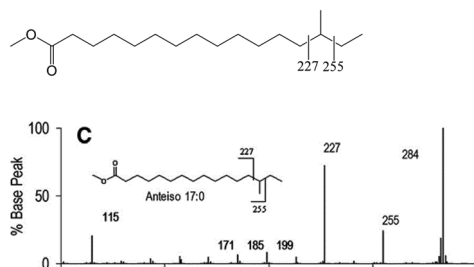
Fig. 7 EIMS/MS spectrum and diagnostic ions for the identification of the anteiso-17:0 FAME. Source: Ran-Ressler *et al.*[51]/Elsevier/CC BY 4.0. The fragmentation patterns and intensities of diagnostic ions near the methyl branch position are similar to the picolinyl ester derivatization method (*cf.* 2.4.1).

2.4.3 Three ion monitoring (MRM) of low-level BCFAs. While picolinyl ester analysis by EIMS and direct FAME analysis by EIMS/MS are both effective methods of localizing methyl branch positions, the former requires laborious derivatization efforts and special chromatography as mentioned above, and the latter has a compromised sensitivity due to strong fragmentation in both hard EI ionization and collisionally activated dissociation stages. We recently developed a chemical ionization (CI) based methodology termed "three ion monitoring" as a routine analytical workflow for samples containing BCFAs of various chain lengths. The CI method works directly towards FAMEs, and can identify and quantify trace-level BCFAs due to a sensitivity boost by a soft CI ionization method and an MRM data collecting mode.[54] Fig. 8 shows that BCFA iso form, anteiso form and straight form can be readily distinguished by the relative abundance of $[M-43]^+$ and $[M-57]^+$ ions. Thanks to the uniform response factors of branched chain FAMEs and straight chain FAMEs yielded by this CI method, quantitative analysis can be conducted without the need for rare BCFA standards.[55]
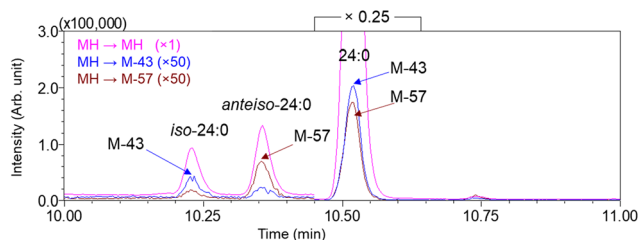
Fig. 8 Iso-24:0 is characterized by high $[M-43]^+$ (blue trace) in contrast to high $[M-57]^+$ (brown trace) in anteiso-24:0. Straight chain 24:0 has both ions at similar levels. Reprinted with permission from Wang *et al.*[54] Copyright 2020 American Chemical Society.

2.4.4 *De novo* analysis of monounsaturated BCFAs. Having introduced a couple of effective methods of saturated BCFA identification, the complete structural elucidation of monounsaturated BCFAs was a very challenging task due to the existence of two positional variants along their hydrocarbon chain, *i.e.*, the uncertain positions of a methyl branch and a double bond. Monounsaturated BCFAs are uniquely found in human surface lipids (sebaceous gland excretion) and their unambiguous identification is important for the study of human skin functionality and dermatology, such as the function of the fatty acid desaturase 2 (FADS2) gene in skin integrity and health.[50] We recently developed a two-pronged approach to address this challenge. CACI-MS/MS analysis would establish the point of unsaturation while the methyl branch structure is unresponsive to the method. Another analysis, EIMS/MS, could discriminate straight, iso and anteiso monounsaturated BCFA methyl esters (BCFAMEs) by a modified method compared to saturated BCFAMEs, due to the influence exerted by the double bond on the EIMS/MS spectra. Ratios of $[M-32]^+/[M-43]^+$ are exploited to differentiate straight and iso structures while a strong $[M-29]^+$ indicates the presence of a methyl branch at the antepenultimate position (anteiso).[24]

2.4.5 Charge switching chemistry for cyclopropane fatty acids and BCFAs. Charge switching chemistry was proven to be a useful tool to characterize cyclopropane and methyl branch structures in fatty acids and fatty acyl chains of a glycerolipid. A few subtle differences in ions could be exploited to distinguish a cyclopropane and a double bond at the same position along a hydrocarbon chain. Although it has not been confirmed, a list of expected mass values of neutral loss for varying cyclopropane positions could be used for cyclopropane fatty acid identifications.[56] With respect to BCFAs, charge switching chemistry has limited resolving capability of the anteiso BCFA from iso to straight chains, but not between iso and straight chains. However, it is still a major advancement in BCFA fingerprinting within a larger glycerolipid molecule, allowing top-down lipid structural analysis.[14]

Traditional ESI-based lipidomics with little to no fatty acyl structure elucidation has been widely applied to investigate biomarkers of health and disease. The methods described here for fatty acid and fatty acyl identification now enable full structural analysis of an ever-widening array of intact lipids as well as complex fatty acid structures. In parallel, these techniques have been applied to the discovery of novel fatty acid structures and/or unusual fatty acid isomers indicative of disease states. We briefly review recent results that depend on the unambiguous detection of unusual isomers in disease states to illustrate the power of these techniques.

3. Fatty acid isomerism in cancer

3.1 Anti-cancer effects of fatty acids

Cancer can be thought of as runaway cell replication. The synthesis of fatty acids including PUFAs is a major structural requirement for cell proliferation, though abnormalities in fatty acid biosynthetic genes can lead to unusual structures.[57] Metabolism is altered toward glycolytic production of energy (ATP) and away from the electron transport chain, a phenomenon known as the Warburg effect.[58] Various unsaturated fatty acid isomers have been studied in relation to cancer where food fatty acids modify carcinogenesis or appear to be a product of cancer cells.

3.1.1 Conjugated linoleic acids.
Conjugated linoleic acids (CLAs) are a group of fatty acids with a conjugated double bond system, with rumenic acid (9Z,11E-18:2) naturally found in dairy food. However, supplements of CLAs are synthesized by isomerization of linoleic acid which contains a mixture of various CLA isomers with 9Z,11E-18:2 and 10E,12Z-18:2 dominating. A wide range of animal and *in vitro* studies have suggested the anti-cancer effects of CLAs, including breast,[59–61] colon,[62–65] stomach,[66,67] prostate[63,68–70] and liver cancers.[71] However, there is significant controversy about which of the CLA isomers exert these anti-cancer effects. For example, 9Z,11E but not the 10E,12Z-18:2 isomer of CLA has a positive effect on prostate cancer;[70] on the other hand, another study showed that the 10E,12Z-18:2 isomer has a cytotoxic effect on rat hepatoma.[71]

Evidence from human clinical trials is scarce. One study found that the administration of CLA to breast cancer patients had some inhibitory effects on fatty acid synthesis. However, not all the measured parameters pointed to such a correlation and a mere number of 23 participants were enrolled, rendering its finding questionable.[59] Another study showed some beneficial effects of CLA supplementation on inflammatory factors and matrix metalloproteinase (MMP) enzymes in a small cohort of rectal cancer patients.[72]

As of now, there is insufficient human trial data confirming any therapeutic effects of CLAs on various cancers. Some stronger correlation of CLAs with anticarcinogenic properties, *e.g.*, against colorectal cancer was attributed to dairy intake where CLAs may only serve as a biomarker;[73] however, another study did not find any correlation between CLA rich dairy food and breast cancer incidence.[74]

3.1.2 Conjugated linolenic acids (CLnAs).
Similar to the CLA situation, the suggested anticarcinogenic properties of CLnAs are also largely based on animal and *in vitro* studies. Dietary seed oils rich in conjugated linolenic acids (CLnAs) from bitter melon and pomegranate, although with the major CLnAs being isomers, were both found to inhibit colon cancer.[75,76] Cancer cell death and tumor growth inhibition were observed for α-eleostearic acid (9Z,11E,13E-18:3)[77] and jacaric acid (8Z,10E,12Z-18:3) was found to induce leukemia and adenocarcinoma (DLD-1) cell death.[78,79] The absence of clinical trials investigating the effects of CLnAs against cancer development prevents the conclusion of its effectiveness. One

human study showed that punicic acid (9Z,11E,13Z-18:3) could be metabolically converted into rumenic acid (9Z,11E-18:3), which might partially explain the mechanisms of CLnA in suppressing cancer growth.[80]

3.1.3 Ximenynic acid.
Ximenynic acid is an 18 carbon enynic acid predominantly found in the seed oils of the *Santalaceae*, *Olacaceae*, and *Opiliaceae* families. Ximenynic acid inhibited the growth of HepG2 cells by suppressing COX-1 gene expression.[81] The health effects of ximenynic acid are very limited, and more on its positive roles in regulating insulin resistance and diabetes will be discussed in section "4.1 Diabetes".

3.2 Shift of fatty acid pathways in cancer cells

3.2.1 Isomeric 18:1(Δ11) as cancer cell markers.
The advent of precision lipidomics around the localization of lipid double bond positions has contributed to the advancement of research in various fields, including the study of cancers. In 2016, Xia's group demonstrated the ability of an emerging double bond derivatizing MS method to distinguish normal and cancerous mouse breast tissues by the abundance of 18:1 (Δ11).[30] Although the method could not show whether 18:1 (11Z) or 18:1(11E) was elevated in cancerous mouse breast tissues, it prompted a wave of studies investigating the correlation between isomeric lipid species and cancers. Human breast cancer tissues were analyzed by the same research group and showed significant differences in the concentrations of isomeric phospholipids containing 18:1(Δ9) and 18:1(Δ11) fatty acyl chains.[82] Another group of researchers used a different double bond derivatizing method to interrogate phospholipids containing 18:1(Δ9) and 18:1(Δ11) fatty acyl chains, also from human breast cancer tissues and human breast healthy tissues.[83] Although these studies did not result in completely consistent findings, probably due to very small sample sizes ($n < 10$), some isomers, *e.g.* PC 18:1–18:1 had consistently higher 18:1(Δ11) for cancerous tissues among the one mouse and two human breast cancer tissue studies.[30,82,83] A more comprehensive study then compared human breast cancer tissues, human plasma from type-2 diabetes patients and human lung cancer tissues to their healthy counterparts. The study concluded that double bond positional isomers led to the differentiation of human breast cancer tissues and human plasma from type-2 diabetes patients from their normal samples; when *sn*-isomers were also taken into account, human lung cancer tissues could also be screened.[29]

3.2.2 Sapienic acid.
Sapienic acid (16:1n-10) is almost uniquely found in human skin surface lipids. As a product of the 6-desaturase enzyme FADS2, its biosynthesis competes with n-6 and n-3 PUFA biosynthesis, especially γ-linolenic acid (18:3n-6) and stearidonic acid (18:4n-3).[84] As predicted from those *in vitro* results,[85] sapienic acid was detected in cancer cells including murine hepatocellular carcinomas (HCCs), and primary human liver and lung carcinomas, indicating that palmitic acid (16:0) was outcompeting PUFAs for membrane fatty acid biosynthesis during rapid proliferation.[86] Based on this

extraordinary finding, another research group showed that two human prostatic adenocarcinoma cell lines of different aggressiveness preferentially synthesized sapienic acid.[87] The researchers found that sapienic acid and its C18 elongated product at 6–8% and 3–6% (wt/wt of total fatty acids), respectively. As a result, sapienic acid has become a spotlight in the cancer research field, providing insights into the developmental mechanisms of cancer cells and a breakthrough point for biomarker discovery in the prognosis of various diseases. These high levels of fatty acids with unusual double bond positions highlight their prospect in all aspects of cancer research. Recently, Xia's group used the double bond derivatizing method they developed to investigate n-10 fatty acids in human breast cancer cells. Due to the excellent isomers resolving power of their method, they unambiguously demonstrated that sapienic acid and other n-10 fatty acids were significantly enriched in more than a dozen aminophospholipid species from a breast cancer cell line than a normal breast cell line, corroborating the findings by previous researchers.[88] These studies surely open up a new line of cancer research, serving as proof of concepts with preliminary data due to their limited numbers of samples ($n < 10$).

3.2.3 Fatty acid desaturase (FADS) abnormalities in cancer and fatty acid changes. The intriguing changes of fatty acid composition in cancer biomasses are correlated with the changes of activities by some genes, for example, those responsible for introducing double bonds to specific locations, *i.e.*, stearoyl-CoA desaturase (SCD) and fatty acid desaturase 2 (FADS2). The functionality of the FADS2 gene is deranged in many cancers.[57] In MCF7 human breast cancer cells, FADS2 activity is absent, suppressing the biosynthesis of the normal eicosanoid precursor arachidonic acid.[89] The absence of FADS2 unmasked the synthesis of various unusual fatty acids *via* the FADS1 directed pathways *in vitro*. *In vivo*, 3 of 9 human breast tumor samples contained sciadonic acid 20:3 (5Z,11Z,14Z), a fatty acid synthesized by FADS1 desaturation of 20:2n-6 and rarely found in any normal human tissues.[90] With the assistance of isomer-resolved lipidomics, *i.e.* OzID, Blanksby's group found a wide range of unusual fatty acids including n-8, n-10, and n-12 monounsaturates biosynthesized by FADS2 in prostate cancer cell lines and tissues. These unusual fatty acids such as 18:1n-10 could be further desaturated by FADS1 mediated pathways to form 18:2n-10 by inserting a Δ5 double bond.[91]

4. Fatty acid isomerism in non-cancerous diseases

4.1 Diabetes

The probe of diabetes lipid biomarkers has been a constant endeavor, with isomer-resolving or double bond derivatizing methods exhibiting great promise. OzID is one of the first methods developed and thus its inventors investigated this issue dating back to 2012. In the study, the researchers identified that some TAG species, especially those with n-7 and n-9

double bond positions had significantly different relative abundances between normal and diabetic plasma samples.[92] Xia's group found that the ratios of C=C isomers enjoyed less variations between individuals, and could possibly serve as a better tool than their individual abundances. They showed that one type of phosphatidylethanolamine, PE 16:0_18:1 could be used as a biomarker for type-2 diabetes when their Δ9 and Δ11 isomers were resolved and quantified.[82] In their continued work, they concluded that discriminations of plasma samples from normal and diabetic patients could be made by analyzing a dozen C=C location isomers or, alternatively, 22 *sn*-position isomers.[29]

Recently, ximenynic acid was found to be beneficial for diabetes by regulating the liver inflammation related interference to the insulin signaling pathway and intestinal microbiota in rats. Pro-inflammatory factors including IL-6, IL-1β and TNF-α, and the JNK/NF-κB inflammatory signaling pathway were all inhibited. While the PI3K/AKT insulin signaling pathway was promoted by ximenynic acid rich sandalwood seed oil.[93,94]

4.2 Nonalcoholic fatty liver disease (NAFLD)

Some studies found a link between the ratios of C=C isomers and nonalcoholic fatty liver disease (NAFLD). Specifically, the ratios of 18:1n-9/18:1n-7 and 18:3n-3/18:3n-6 could discriminate control and NAFLD mouse liver samples.[35] In contrast to serving as biomarkers for NAFLD, another study showed the preventive effects of 18:3n-6 against it owing to the fatty acid's roles in lipid metabolism and autophagy.[95]

4.3 Atherosclerosis

Research into unusual fatty acid isomer roles or their relative abundance in atherosclerosis has been mainly focused on CLA. For reference, den Hartigh extensively reviewed this area.[96] In brief, the review analyzed the findings of various animal studies and confirmed the positive efficacy of CLA against atherosclerosis. In a recent publication, the same group demonstrated the underlying mechanisms of CLA's atheroprotective effects, attributed to increasing high-density lipoprotein (HDL) particle concentration, HDL anti-inflammatory potential, and promoting beneficial effects on cholesterol efflux.[97]

5. Eicosanoid modulation

5.1 Sciadonic acid and arachidonic acid

Sciadonic acid, or 20:3(5Z,11Z,14Z), is an unusual fatty acid isomer of 20:3(8Z,11Z,14Z), which is a precursor of arachidonic acid, 20:4(5Z,8Z,11Z,14Z). Sciadonic acid has long been known to replace circulating arachidonic acid.[98] There has been interest in using sciadonic acid to modulate eicosanoid production and thus modulate many physiological responses including the immune system. Research has been focused on the interactions between sciadonic acid and prostaglandin E$_2$ (PGE$_2$). Supplementation of sciadonic acid decreased the production of a pro-inflammatory mediator, PGE$_2$, by 4–29% in mouse

macrophages, presumably due to the inhibition of inducible nitric oxide synthase (iNOS) and cyclooxygenase-2 (COX-2).[99] In a follow-up, the same authors corroborated their initial findings in a murine microglial BV-2 cell model, and extended the eicosanoid modulatory effects to two other non-methylene-interrupted fatty acids, *i.e.* 18:3(5*Z*,9*Z*,12*Z*) and 20:3 (7*Z*,11*Z*,14*Z*).[100] Using an epithelial cell line, some scholars found that during pathogenic yeast infection, sciadonic acid supplementation could inhibit the production of PGE_2 without changing the production of ω-3 fatty acid metabolites which could counteract the normal inflammatory process necessary to combat infections.[101]

6. Weight loss and conjugated fatty acids

6.1 Conjugated linoleic acids

Besides anticancer effects, another research front line of CLAs is around weight loss. Park and coworkers examined the effects of CLAs on mice body composition, and found a reduction in fat deposition and an increase in lipolysis in adipocytes.[102] Also, they tested the respective effects of common isomeric CLAs, *i.e.*, 9*Z*,11*E*-18:2 and 10*E*,12*Z*-18:2, and found that only 10*E*,12*Z*-18:2 could reduce body fat in mice. The weight loss effect was found to be linked to heparin-releasable lipoprotein lipase (HR-LPL) activity.[103] In a continued study by the same authors, they showed that only 10*E*,12*Z*-18:2 and its alcohol form could inhibit the HR-LPL activity among a wide range of structural variants.[104] Due to the scarcity of human clinical trials and even fewer reporting positive outcomes, one review by Fuke *et al.* did not explicitly indicate CLA's efficacy on human weight loss.[105] However, another review by den Hartigh suggested a positive link between 10*E*,12*Z*-18:2 and obesity, based on the evidence from preclinical trials and a few clinical trials finding either fat mass reduction or weight loss.[96]

6.2 Conjugated linolenic acids

The weight-losing effects of CLnA (18:3) may share a similar chemical and physiological mechanism with a CLA (18:2). Calendic acid (8*E*,10*E*,12*Z*-18:3) shares the same 10*E*,12*Z* structure with 10*E*,12*Z*-18:2, with an additional *trans* double bond at the C8 position. Calendic acid was found to have some weaker effects towards body fat reduction than 10*E*,12*Z*-18:2.[106] The CLnA was found to reduce perirenal and epididymal adipose tissue weight to a greater extent than the CLA.[107] However, jacaric acid (8*Z*,10*E*,12*Z*-18:3) which also contains the 10*E*,12*Z* structure, was found to have no effects on body fat reduction.[108]

7. BCFAs and health

BCFAs have been found in microbial lipids at high levels a long time ago, but are also present in other organisms including the nematode *Caenorhabditis elegans* (*C. elegans*) and mammals.[109] Ruminant fats, along with some seafood and Asian fermented foods, are BCFA-rich food sources for mammals, which have been well documented in a previous review.[110] BCFAs from the diet undergo *de novo* syntheses in organisms from bacteria to mammals and exhibit diverse health effects. Among various BCFA isomerism, the monomethyl BCFA (mmBCFA), especially long-chain ones, attracted the most attention in recent decades, due to their close involvements in metabolic regulation, anti-inflammation, anticancer, weight control, as well as neuroprotection and cardioprotection, as summarized in previous reviews.[111–113] Thus, here we aim to emphasize the significance of BCFAs in two important but not well reviewed aspects: the essential role in *C. elegans* development and the special support to human skin functions.

7.1 BCFAs are essential for *C. elegans*

The nematode *C. elegans* is a common model used in human nutritional studies for evaluating diet-related aging and longevity, obesity and the molecular basis of nutrition.[114] It was found in 2004 that mmBCFA C15iso and C17iso were essential for *C. elegans* growth and development, since suppressing their biosynthesis *via* ELO-5 and ELO-6 could arrest *C. elegans* larva development, and this situation could be reversed by giving BCFA supplements.[115] Another study showed that C17iso and its downstream metabolite glycosphingolipid could restore the *C. elegans* larva development by activating intestinal and neural mTORC1 under dietary amino acid deficiency.[116] C17iso was also found to be incorporated into phospholipids to ensure the membrane integrity of the endoplasmic reticulum for lipid droplet growth in *C. elegans*.[117] Besides, the existence of the branched side chain in mmBCFA contributes to the much lower melting point compared to the normal straight chain fatty acid with the same carbon number and saturation degree, which increases the membrane fluidity in multiple organisms. A study demonstrated decreased membrane fluidity induced by high-glucose stress in *C. elegans*, and showed that mmBCFA could rescue *C. elegans* in a high-glucose environment, suggesting the critical role of mmBCFAs in adaption to elevated glucose conditions.[118]

7.2 BCFAs support human skin function

Previous research identified BCFAs in human blood, adipose tissues, brain and skin.[112,115] Notably, BCFA isomerism showed both great richness and diversity, especially in skin, and connective tissues including skin sebaceous glands (SGs) and eyelid meibomian glands (MGs), as well as their secretion products such as *vernix caseosa* covering the newborn skin. An early paper established that the human skin fatty acid profile is unique compared to that of internal organs in the desaturation degree and fatty acid isomerism: more than 97.1% of human skin fatty acids are saturated or monounsaturated; and among them, saturated and monounsaturated BCFAs (SBCFAs and MUBCFAs) account for 5.4% and 6.6% of total fatty acids,

Table 1 Summarized functions of unusual fatty acid isomers

Unusual fatty acid isomers	Biological functions	Type of study	References
Conjugated linoleic acid (CLA)	Anticancer (breast cancer)	Clinical	McGowan et al.[59]
	Anticancer (breast cancer)	Cell lines	Rakib et al.[60]
	Anticancer (breast cancer)	Animal	Futakuchi et al.[61]
	Anticancer (colon cancer)	Animal	(1) Bassaganya-Riera et al.[62] (2) Shiraishi et al.[65]
	Anticancer (colon cancer)	Cell lines	(1) Lau et al.[63] (2) Soel et al.[64]
	Anticancer (stomach cancer)	Animal	(1) Chen et al.[66] (2) Ha et al.[67]
	Anticancer (prostate cancer)	Cell lines	(1) Lau et al.[63] (2) Kim et al.[69] (3) Song et al.[70]
	Anticancer (prostate cancer)	Animal	Cesano et al.[68]
	Anticancer (liver cancer)	Animal	Yamasaki et al.[71]
	Anticancer (rectal cancer)	Clinical	Mohammadzadeh et al.[72]
	Atheroprotective	Animal	Vaisar et al.[97]
	Atheroprotective	—	Review by den Hartigh[96]
	Weight loss or fat mass reduction	Animal	(1) Park et al.[102] (2) Park et al.[103]
	Weight loss or fat mass reduction	—	Review by den Hartigh[96]
Conjugated linolenic acid (CLnA)	Anticancer (colon cancer)	Animal	(1) Kohno et al.[76] (2) Kohno et al.[75]
	Anticancer (ferroptotic cell death)	Cell lines	Beatty et al.[77]
	Anticancer (leukemia cell death)	Cell lines	Liu et al.[78]
	Anticancer (adenocarcinoma cell death)	Cell lines	Shinohara et al.[79]
	Conversion into rumenic acid	Clinical	Yuan et al.[80]
	Weight loss or fat mass reduction	Animal	(1) Chardigny et al.[106] (2) Koba et al.[107]
Ximenynic acid	Anticancer (inhibits the growth of HepG2 cells)	Cell lines	Cai et al.[81]
	Diabetes (ameliorate insulin resistance)	Animal	(1) Gao et al.[93] (2) Zhang et al.[94]
18:1n-7 or 18:1(Δ11)	Biomarker for mouse breast cancer tissues	Animal	Ma et al.[30]
	Biomarker for human breast cancer tissues	Human tissues	(1) Zhang et al.[82] (2) Kuo et al.[83]
	Biomarker for human breast cancer tissues, human plasma from type-2 diabetes patients	Human tissues	Cao et al.[29]
	Biomarker for type-2 diabetes	Human plasma	(1) Stahlman et al.[92] (2) Zhang et al.[82]
	Biomarker for nonalcoholic fatty liver disease (NAFLD)	Animal	Xu et al.[35]
Sapienic acid or 16:1n-10	Biomarker for mouse liver cancer, human liver cancer and lung cancer	Animal and human tissues	Vriens et al.[86]
	Biomarker for human prostatic adenocarcinoma	Cell lines	Ferreri et al.[87]
	Biomarker for human breast cancer cells	Cell lines	Lin et al.[88]
	Biomarker for prostate cancer cell lines and tissues	Cell lines	Young et al.[91]
18:1n-10	Biomarker for prostate cancer cell lines and tissues	Cell lines	Young et al.[91]
Sciadonic acid or 20:3 (5Z,11Z,14Z)	Biomarker for human breast cancer	Human tissues	Park et al.[90]
	Replace circulating arachidonic acid	Cell lines	Tanaka et al.[98]
	Eicosanoid modulation	Cell lines	(1) Chen et al.[99] (2) Chen et al.[100] (3) Ells et al.[101]
γ-Linolenic acid or 18:3n-6	Inhibit nonalcoholic fatty liver disease (NAFLD)	Cell lines	Liang et al.[95]
Branched chain fatty acid (BCFA)	Essential role in C. elegans development	C. elegans culture	(1) Kniazeva et al.[115] (2) Zhu et al.[116] (3) Zhang et al.[117] (4) Vieira et al.[118]
	Biomarkers for acne	Clinical	Okoro et al.[120]
	Biomarkers for meibomian gland dysfunction (MGD)	Clinical	(1) Joffre et al.[123] (2) Joffre et al.[124]
	Intestinal health	Cell lines	Yan et al.[127]
	Intestinal health	Animal	Ran-Ressler et al.[128]

respectively, and accumulate mainly in wax esters (WEs) of skin lipids.[119] In particular, MUBCFAs with both double bond and methyl side chains were detected and presumed to assist with human sebum fluidity so as to support normal skin func-

tion, such as enhancing tolerance to extreme environmental temperatures.[50]

Several clinical studies have associated the unique skin BCFA profile with human skin function or disease pathogen-

esis. For example, a cross-sectional analytical study conducted in Nigeria in 2021 observed significantly lower levels of the anteiso-even chain BCFA with 12–18 carbons and higher levels of the normal straight chain SFA with 15–17 carbons in the facial sebum lipid profile of the acne group compared to the non-acne group.[120] Another earlier study also found the opposite change tendency of the BCFA and straight chain FA with aging: the abundance of sebum WE iso-16:1 decreased from infancy to around 20-year-old, and then increased until around 50-year-old, whereas straight chain 16:1 followed a completely reversed course.[121] A pilot study designed for applying skin sebum lipidomics in neurodegenerative disease discovery showed that both even chain and odd chain BCFAs in sebum were negatively correlated with the BMIs in both healthy and Alzheimer's disease/Parkinson's disease subjects.[122]

Meibum is a lipid-based substance synthesized in the meibomian glands of the human eyelids, excreted on to surface of the tear film to reduce evaporation. Abnormalities in meibomian gland function cause dry eye. A series of studies showed that meibum BCFA from C16–C26 in both iso- and anteiso-forms were significantly increased, while SFAs especially C16 and C18 were decreased, in meibomian gland dysfunction (MGD) patients.[123] The accumulation of BCFAs in MGD appeared to be compensation for tear film fluidity as an adaption to tear loss.[124]

Vernix caseosa, or vernix in short, is secreted by newborn SG during the 3rd trimester of gestation, covering the skin of a full-term baby during delivery, and providing as high as 29% of BCFAs for the fetus GI tract.[125,126] The function of vernix BCFAs has been summarized thoroughly in previous reviews.[111,113] In short, vernix BCFAs exhibited the ability to alleviate intestinal inflammation in both *in vitro* and *in vivo* models,[127,128] which indicated its potential in modulating a newborn immune system, though more research is warranted to expand this area. All the aforementioned functions of unusual fatty acid isomers are summarized in Table 1.

8. Conclusion

Unusual fatty acid isomers can be derived from specific human foods such as dairy, seeds, nuts and seafood. However, research on these fatty acids is relatively limited partly due to their similarity to their common isomeric counterparts and the inability to differentiate them on a *de novo* basis. The double bond position and methyl branch structure are key features for fatty acids imparting their biological functions. The interplay of advances in lipidomics and lipid functionality has led to their co-development through technological and theoretical iterations. Mass spectrometric methods capable of localizing double bond positions and methyl branch positions provide the necessary support for the functional discovery of unusual fatty acid isomers, which represent mechanism-related biomarkers for cancer, diabetes, nonalcoholic fatty liver disease (NAFLD) and atherosclerosis; modulation of eicosanoids; effects of weight loss and fat mass reduction; the essentiality of BCFAs to the growth of *C. elegans*; and BCFA's

roles in the integrity and health of skin and gut. As methods become more widely adopted, the biological effects of unusual isomeric fatty acids will emerge, with implications for foods and disease states. Adoption of these structure specific or equivalent analytical workflows will enable a new era in fatty acid and glycerolipid biology similar to the adoption of gas chromatography.

Author contributions

Donghao Wang, Zhen Wang and J. Thomas Brenna: conceptualization, design and drafting of the manuscript. Donghao Wang and Tingxiang Yang: generating figures/tables & editing. All authors approved the final version for submission.

Conflicts of interest

The authors declare no conflicts of interest.

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