New strategies for resolving oligosaccharide isomers by exploiting mechanistic and thermochemical aspects of fragment ion formation

Andres Guerrero, Carlito B. Lebrilla*  
Department of Chemistry, University of California Davis, CA 95616, United States

1. Introduction

Oligosaccharides are very common macromolecules present in nature as glycoconjugates or as free species [1–3]. These compounds can differ in size, monosaccharide composition, sequence, branching, linkage and anomeric configuration [4]. In biological samples, oligosaccharides amounts can also vary across many orders of magnitude. There has been recent significant interest in these compounds led by the increasing realization of their roles in many biological processes [5–7]. For this reason, there have been many new developments in the analysis that have made structure elucidation more facile [8]. However, despite advances in these methods, the analysis of oligosaccharides remains far from routine, and often various structural types require different analytical tools.

Mass spectrometry (MS) has emerged as arguably the most useful method for the structural elucidation of oligosaccharides [9,10]. Structural elucidation is enabled by tandem MS (MS/MS), perderivatization or enzymatic digestion of native structures [11]. MS/MS techniques have been extensively used for this purpose with variable success [12–18]. Quite common is the modification of the oligosaccharides by different chemical reactions including permethylation [11,15], peracetylation [14], reduction [17], periodate oxidation [18] or trifluoroacetolysis [16] in order to increase the signal, stabilize the oligosaccharide, eliminate the alditol form or promote a specific fragmentation.

One important biological fluid where oligosaccharides play a role is milk. Free oligosaccharides are a major component of the milk, especially in the human where they can be found in concentrations of 7–12 g/L [19]. It is well-known that human milk oligosaccharides (HMOs) have several beneficial roles during the development of the infant such as providing nutrient for beneficial bacteria [20,21], prevention of adhesion of the pathogenic bacteria on the intestinal surface [22], immunomodulation [23,24] and modification of the cell glycome [25]. Positional isomers are also abundant in HMOs and these structural differences have important biological relevance. For example, the linkage pattern of the fucosylated moiety in HMOs defines the antigens LewisA and LewisX that are known to have distinct biological functions in cell-to-cell recognition processes [26,27].

The intensities of the fragment ions in tandem MS experiments are the result of a complex combination of factors such as the relative stability of precursors and products, bond strengths, activation barriers or position of the charge [28,29]. All these factors are in turn the consequence of a specific chemical structure. Hence, even when isomers can yield the same products in tandem MS experiments, they may often be resolved by based on the relative abundances of the fragment ions.

In the present study we show the systematic examination of the pathways and product ions generated by infrared multiphoton
dissociation (IRMPD) of representative HMOs. The interpretation of
the fragmentation pattern observed for these molecules provided
an unexpected amount of information concerning its structure
and was used not only to distinguish different pure isomers
but mixtures of them. Furthermore, the experiments were per-
formed without chemical modification or previous separation of
the oligosaccharides what significantly simplified the procedure.

2. Experimental methods

Mass measurements were performed by using a MALDI-FT-ICR
(IonSpec, Irvine, CA) instrument equipped with a 7.0-T supercon-
ducting magnet. The instrument is equipped with a pulsed Nd:YAG
laser (355 nm). The details of the instrument are published else-
where [30]. The instrument was calibrated using a natural mixture
of maltooligosacharides from beer as described in previous publi-
cations [31].

The oligosaccharides used in this study include isomers LNFP I,
LNFP II, LNFP III and LNFP V, isomers LNDFH I and LNDFH II, and were
acquired from Glyko®. The compounds were of analytical grade and
were used without further purification. The MALDI matrix 2,5-DHB
and NaCl were obtained from Sigma–Aldrich®. The MALDI samples
were prepared by applying on the MALDI plate in the following
order a 0.5 μL solution of the oligosaccharide (10 μg/μL), 0.5 μL of
2,5-DHB solution (50 μg/μL) and 0.2 μL of NaCl (0.1 M), all in 50:50
AcN/H2O. The sample was dried in the vacuum in the pre-chamber
of the MALDI source.

The tandem MS IRMPD experiments were performed using a
1060 nm instrument (Parallax Tech. Inc.) with varying irradiation
times. Double resonance experiments were accomplished by apply-
ing a burst excitation pulse to the desired ion during the IRMPD
fragmentation. The voltage in each case was optimized to be the
minimum necessary to eliminate the desired fragment in order to
avoid expulsion of other informative ions.

3. Results and discussion

Three complementary procedures are described illustrating that
it is possible to resolve isomers of complicated structures employ-
ing only mass spectrometry by taking advantage of both the
mechanistic differences in the fragmentation pathways and the
thermochemical properties of the molecules. Four isomers of lacto-
N-fucopentaose and two of lacto-N-idofucohexaose (Scheme 1) from
human milk [19] were used to develop and illustrate the methods.

3.1. Distinguishing isomers by disruption of the fragmentation
pathways through double resonance

Most methods employed to identify isomers by MS/MS are based
on the assignment of production ions after the fragmentation process
relying on diagnostic peaks that are unique to specific structures.
Unfortunately many isomers do not yield unique fragments and the
tandem mass spectra are often indistinguishable. For example, the
IRMPD of LNFP I and LNFP II in Fig. 1a and c, respectively, show the
same product ions with nearly the same abundances.

Despite the close similarities in the fragment composition, it is
possible to resolve isomers by probing the differential pathways
that generate the product ions. Scheme 2 shows the potential frag-
mentation pathways of isomers LNFP I and LNFP II starting with the
sodiated parent. The diagram is based on the product ions formed
after IRMPD and shows most of the potential routes that create
them. The main difference between the two corresponding tandem
MS is the presence of the product ion m/z 714 in LNFP II, which
corresponds to the cleavage of the galactose from the branched ter-
minus. Although the presence of this very minor fragment may be
sufficient to differentiate the two isomers, it is often found in very
low abundances when present and often not at all even for LNFP II.

An examination of Scheme 2 shows a difference between the
two isomers for the formation of the ion m/z 568. Two simultaneous
pathways generate this product ion in the fragmentation of LNFP
I, the stepwise loss of the fucose and the galactose from the non-
reducing terminus and the concerted cleavage of the disaccharide
from the precursor ion. In the branched isomers LNFP II and LNFP III
the latter pathway is not possible. That is, formation of product ion
m/z 568 occurs only after the sequential losses of fucose (m/z 730)
and galactose or vice versa, loss of galactose (m/z 714) first from
the branched terminus (Scheme 2). The pathway was probed in the
FT-ICR during tandem MS with IRMPD by employing a double
resonance ejection pulse of the intermediate species involved in
the formation of product ion m/z 568. The ejection of fragment ions
m/z 730 and m/z 714 from LNFP I yields the same abundances of m/z
568 (Fig. 1b) confirming the presence of an alternative mechanism
to the sequential losses of a fucose and a galactose. However, the
ejection of the same two ions from LNFP II during IRMPD eliminates
the fragment ion m/z 568 completely suggesting that this product is
only produced through those intermediates (Fig. 1d).

In general, this method is applicable for differentiating linear
and branched structures. Another example is given in Fig. 2 with
the isomers LNDFH I and LNDFH II. Both LNDFH I and LNDFH II yield
electronically the same fragmentation products (Fig. 2a and c,
respectively). As in the previous case, the product ion m/z 568 is
generated simultaneously by two fragmentation pathways for the
linear isomer LNDFH I, while in the branched one LNDFH II the
direct cleavage of the disaccharide to form the same fragment is not
possible. Thus, the ejection of the appropriate intermediate product
causes the elimination of species m/z 568 in LNDFH II (Fig. 2d) but not in
LNDFH I (Fig. 2b).

3.2. Distinguishing isomers based on differential dopant affinities

In MALDI-MS of oligosaccharides a dopant salt, typically alkali
metals such as sodium chloride, are used to produce a single
quasimolecular ion corresponding to the coordinated metal [32].
Oligosaccharides ions formed by metal attachment present some
advantages for analytical tandem MS over the ionic species gener-
ated by proton attachment. For example, some metal-coordinated
oligosaccharide ions may give cross-ring products yielding infor-
mation regarding the linkage [33]. Furthermore, alkali metals
stabilize labile residues like sialic acid [34] thereby increasing the
sensitivity of the method. For this reason the oligosaccharide metal
coordination has been extensively studied with alkali salts, alkaline
earths and even transition metals [35].

It has been previously shown that the ionic complexes are likely
to be composed of several isostructures (metals coordinated to dif-
ferent binding sites) [36]. Consequently, each corresponding ion
signal in the MS spectrum is formed from a number of subpop-
ulations of oligosaccharides with the metal attached to different
positions. The abundance of each isostructure in the whole ionic
complex population will depend on the affinities of the different
parts of the molecule for the dopant metal.

Although the exact proportion of each isostructure for ionic
complexes is unknown, it has been shown that branched structures
have a greater affinity for the alkali metal ion than linear structures
[37]. As a consequence, after the IRMPD of molecules with branch
and linear moieties, product ions corresponding to branched frag-
ments are expected to have higher abundances compared to those
coming from linear structures. Therefore, compounds such as LNFP
II and LNFP III with branched non-reducing ends and linear reduc-
ing ends should favor ions from the non-reducing ends versus to
the reducing ends as opposed to structures such as LNFP I where
both reducing and non-reducing ends are linear.
We tested the above hypothesis by examining the IRMPD product ion abundances of the LNFP isomers varying the irradiation time. The product ion \( m/z \) 365 corresponds to the disaccharide at the reducing end Gal\( \beta(1\rightarrow4) \)Glu coordinated to sodium and is present in the fragmentation spectra of all the LNFP isomers. Fragment \( m/z \) 388 corresponds to the disaccharide at the non-reducing end Gal\( \beta(1\rightarrow4) \)GluNac in LNFP III or Gal\( \beta(1\rightarrow3) \)GluNac in LNFP I, LNFP II and LNFP V. The relative abundances of these two product ions should differ depending on the structural nature of the termini. Fig. 3 shows the ratio of the abundances of the product ions \( m/z \) 365 and \( m/z \) 388 for each isomer versus the overall degree of fragmentation which was monitored using the

---

**Fig. 1.** IRMPD fragmentation experiments of isomers LNFP I (a and b) and LNFP II (c and d). The expulsion of species \( m/z \) 730 and \( m/z \) 714 during fragmentation (b and d) supposes the disappearance of product ion \( m/z \) 568 only in LNFP II. The red boxes indicate the range of double resonance ejection during the IRMPD fragmentation. The blue boxes indicate the position of the potentially affected product ion. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)
loss of fucose from the precursor ion, ratio ($m/z$) 730/876, as a reference.

For all the isomers, the ratio ($m/z$) 365/388 reaches an asymptotic value at prolonged irradiation times. For isomers with branched non-reducing end such as LNFP II and LNFP III that ratio reaches a value of 0.1, whereas for the linear isomer LNFP I it reaches 0.25. In LNFP V the position of the fucose generates a branched-like structure at the reducing end (Scheme 1) and the ratio is in this case 0.7, as would be expected if the coordination of the sodium favors the fragment ion corresponding to the reducing end $m/z$ 365.

The influence of the position of branched moieties in the ratio ($m/z$) 365/388 was confirmed with the oligosaccharide LNDFH II. LNDFH II presents both, a branched structure at the non-reducing end similar to LNFP II or LNFP III and the fucose in position

Scheme 2. Diagram of the fragmentation pathways of isomers LNFP I and LNFP II. Red arrows correspond to internal cross-ring cleavages (CRC), blue arrows correspond to the C,Z fragmentation of the glycosidic bond and black to B,Y cleavages [32]. The $m/z$ of the species corresponds to each product ion bound to Na$^+$. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)
Fig. 2. Fragmentation experiments of isomers LNDFH I (a and b) and LNDFH II (c and d). The ejection of species $m/z$ 730 and $m/z$ 714 during IRMPD (b, d) affects the disappearance of $m/z$ 568 in LNDFH II (d). The red boxes indicate the frequency range of double resonance ejection during the IRMPD. The blue boxes indicate the position of the potentially affected product ion. (For interpretation of the references to color in this figure legend, the reader is referred to the web version of this article.)

To probe the capability of this methodology to resolve complex samples, mixtures in different proportion of LNDFH I and LNDFH II were tested. Fig. 4 shows the evolution of the ratio ($m/z$) 365/388 versus the overall degree of fragmentation for both pure and different mixtures of isomers.

Again, for both isomers, the ratio ($m/z$) 365/388 reaches an asymptotic value at prolonged irradiation times. Using the values obtained for the pure isomers as reference, the mixtures were compared and excellent agreement with the estimated theoretical average was obtained (Table 1).

3.3. Resolving isomers by monitoring competitive fragmentation pathways

During IRMPD the isomers of LFP all yield cross ring cleavages (CRC) except for LNFP V where the fucose is directly attached to the reducing end what appears to hinder this fragmentation pathway. Product ions $816/m/z$ and $756/m/z$ are CRC fragments formed by losses of $60 \mu(0.2A)$ and $120 \mu(0.4A)$, respectively, from the glucose reducing end of the precursor ion. A similar set of ions are produced when the fucose dissociates concurrently corresponding to fragments $670/m/z$ and $610/m/z$, respectively. These two processes are in competition as illustrated in Scheme 3.

The isomers that undergo the same fragmentation pathways will have different activation barriers and will yield different abundances. We monitored the loss of the fucose from the parent versus the loss of the fucose from the fragments corresponding to the CRC products. In Fig. 5 the ratio of intensities of the parent to loss of fucose ($m/z$) 730/876 was plotted versus the similar fragmentation from the CRC product ($m/z$) ($670 + 610$) / ($816 + 756$). When the fragment yield is increased, accomplished by increasing the irradiation period, the ratio of the product ion to the precursor ion increases, hence the ratio ($m/z$) 730/876 increases. Interestingly, the ratio of the fucose loss from the CRC fragments increases in linear manner. The results for the three oligosaccharides are shown. We observe that both LNFP I and LNFP II have the same slope corresponding to 1, while LNFP III has a considerably smaller value corresponding to 0.5. Thus, LNFP II and LNFP III can be finally differentiated.

Fig. 3. Evolution of the quotient ($m/z$) 365/388 during IRMPD for isomers LNFP I, LNFP II, LNFP III and LNFP V and oligosaccharide LNDFH II.

Fig. 4. Evolution of the quotient ($m/z$) 365/388 during IRMPD for isomers LNDFH I, LNDFH II and mixtures in different proportion of them.
The results provide some information regarding the kinetics of the associated fragmentation pathways. While fucose loss from either fragment or molecular ion appears to involve similar energetics for LNFP I and II, the same loss for LNFP III is markedly different.

The theme of monitoring fragmentation pathways by varying dissociation energies can also be applied to other product ions. Fragment \( m/z \) 388, present in all LNFP isomers, corresponds to either the internal fragments Gal–GlcNac and GlcNac–Gal. It is often accompanied by a homologous fragment containing an additional oxygen (\( m/z \) 406). Although the presence of these ions are the result of several complicated and competing pathways (Scheme 2), their relative abundances can be monitored and compared to the overall fragmentation and represented by the loss of fucose from the parent ion (\( m/z \) 730/876). Shown in Fig. 6 is the ratio of (\( m/z \) 388/406) as function of the ratio (\( m/z \) 730/876). The former reaches an asymptote that can be used to differentiate, LNFP II and LNFP III. This ratio is generally reproducible and can be used to determine the percentage of each isomer. Shown in Fig. 6 are mixtures of LNFP II and LNFP III with different proportions of each isomer. The value are intermediate of the pure compounds and correspond well to that expected based on the relative abundances as shown in Table 2.

**4. Conclusion**

Oligosaccharide analysis represents a considerable challenge even with the most advanced analytical tools. Analysis may require a number of tools used together to obtain comprehensive information. Mass spectrometry provides the best combination of high sensitivity with structural analysis. Furthermore, mass spectrometry provides more than just compositional analysis. By manipulating dissociation pathways and interpreting the fragmentation pattern from a mechanistic point of view, isomer differentiation is possible. The results presented in this report show that although complete structural information cannot be obtained using tandem MS, the fragmentation pathways can be used to differentiate isomers. It is possible, that the combination of these approaches may provide new and unexpected amount information that could be used to elucidate the composition of complex isomeric mixtures. The procedures described in this study appear to be quite general and may be useful to differentiate isomeric compounds. While the methods have been applied to MALDI
produced ions, they should also be applicable to ions produced by other ionization techniques such as electrospray ionization.

Acknowledgments

This manuscript is dedicated to Professor Detlef Schroeder who was a friend and colleague. A.G. wants to thank the Spanish National Research Council for the economical supply 2009 EST CSC–1536. Funding provided by NIH (GM R01HD061923) is gratefully acknowledged.

References


