Quantification of the Effect of Citrulline and Homocitrulline Residues on the Collision-Induced Fragmentation of Peptides

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ABSTRACT: Posttranslational modifications of proteins like citrullination and carbamylation are associated with several diseases. Detailed analytical characterization of citrullinated and carbamylated proteins or peptides could be difficult due to the low concentration of the analytes in complex biological samples. High structural similarity and chemical behavior of citrullinated and carbamylated residues also pose a challenge. We previously reported the “citrulline effect” phenomenon that is manifested in the generation of intense y type ions originating from Cit-Zzz amide bond scissions in collision-induced dissociation tandem mass spectra of citrullinated tryptic peptides. In this study, we created a rigorous tryptic-like model system of both citrulline and homocitrulline-containing peptides that included appropriate and well-defined controls and fragment analogues to quantify the citrulline effect and investigate whether there is an effect for homocitrulline residues as well. Our results show that citrulline residues significantly increased fragmentation at their C-terminus relatively independent of the identity of the following amino acid. In comparison, homocitrulline residues displayed inconclusive results at the same energies. However, the strength of effects was dependent on collision energy and the position of citrulline and homocitrulline in the sequences. As newer software algorithms tend to observe structure–intensity relationships during annotation, this finding increases reliable identification of modified proteins/peptides.

INTRODUCTION

Citrullination is an enzymatic posttranslational modification (PTM) of arginine residues leading to citrullines (Cit, X) in proteins.1 Homocitrulline (Hci, B) residues are formed through the nonenzymatic reaction of isocyanic acid with the amino group of lysine residues or with protein N-termini through a process known as carbamylation.2 Both citrullination and carbamylation result in a decreased basicity and thus an altered charge state, activity, or structure compared to the original species. Citrullination plays a role in several physiological functions, including epigenetic regulation3 and skin homeostasis,4 while hypercitrullination is associated with this physiological functions, including epigenetic regulation3 and skin homeostasis,4 while hypercitrullination is associated with this phenomenon that is manifested in the generation of intense y type ions originating from Cit-Zzz amide bond scissions in collision-induced dissociation tandem mass spectra of citrullinated tryptic peptides. In this study, we created a rigorous tryptic-like model system of both citrulline and homocitrulline-containing peptides that included appropriate and well-defined controls and fragment analogues to quantify the citrulline effect and investigate whether there is an effect for homocitrulline residues as well. Our results show that citrulline residues significantly increased fragmentation at their C-terminus relatively independent of the identity of the following amino acid. In comparison, homocitrulline residues displayed inconclusive results at the same energies. However, the strength of effects was dependent on collision energy and the position of citrulline and homocitrulline in the sequences. As newer software algorithms tend to observe structure–intensity relationships during annotation, this finding increases reliable identification of modified proteins/peptides.

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Another promising technique may be the use of vacuum ultraviolet photodissociation (VUVPD) to randomize the occurrence of amide bond cleavages through the whole sequence.23

We previously reported a neutral loss which was selective to citrulline residues for histone-related pentapeptides in electron-transfer dissociation (ETD).24 In addition, electron-transfer higher energy collision-induced dissociation (EThCD) was proven to be beneficial for both increasing fragmentation efficiency and the incidence of neutral losses.25 We also found that the citrulline effect, which is the enhanced cleavage probability C-terminal to citrulline residues in (higher energy) collision-induced dissociation (HCD, CID), was preserved in EThcD as well. Although several research groups18,25 examined the amide bond scission preferences of tryptic peptides at statistical levels, insufficient data are currently available regarding the effect of special amino acids including citrulline or homocitrulline.

Therefore, our aim was to characterize these effects under well-controlled circumstances, especially concentrating on—but not restricted to—the amide bonds at the C-terminus of citrulline and homocitrulline residues.

We chose the sequences AAXZAAK and AABZAAK, where X and B stand for citrulline or homocitrulline, respectively, and Z denotes any other amino acid, including some special ones like carbamidomethyl cysteine (C′) and oxidized methionine (M′) that are frequently occurring in MS-based experiments. Carbamidomethylation of Cys residues are generally carried out by iodoacetamide; however, this may negatively affect identification rates in proteomics.26 Very recently, Wiśniewski et al. reported a method involving only the reduction of Cys residues without alkylation that resulted in an improved peptide and protein identification.27 Therefore, we included both Cys and carbamidomethylated Cys residues in our experiments.

According to Hao et al., citrulline residues are converted to ornithines (Orn, O) using collision-induced dissociation.28 Lee et al. reported that carbamyllysine (Hci) residues are also prone to loss of isocyanic acid upon CID leading to lysine (Lys, K) residues.15 McGee and McLuckey observed that Orn strongly promotes amide bond cleavage at its C-terminus.29 They also proposed a neighboring group reaction mechanism in which the Orn forms a six-membered lactam. We also described a facile cleavage C-terminal to Cit, which could be explained by a loss of isocyanic acid followed by a lactam formation (Scheme 1).30 Thus, we also synthesized peptides where X or Z are substituted for ornithines (O) or lysines (K) to compare the MS/MS behavior of these residues to that of Cit and Hci.

Alanines are reported to have only a small effect on fragmentation;15 therefore, they were incorporated for elongating the peptides. This way, the peptides are relatively large to be successfully separated on C18 columns. The peptides were synthesized with a lysine C-terminus so that they model an enzymatic cleavage by trypsin or LysC, frequently used in bottom-up proteomic experiments.

We also synthesized peptides where X/B/O/K were replaced by alanine residues (AAAZAAK) as controls to examine whether the Z has a contribution in the cleavage increment or solely the X/B/O/K causes the phenomenon as several amino acids, e.g., Gly, Ser, Thr, Cys, Met, Asp, and Asn and especially Pro, have a preference to be cleaved at their N-termini.16,31,42 In addition, Leu, Ile, and Val residues also tend to produce intense peaks corresponding to the cleavage at their C-terminus.32 As Asp and Gly residues commonly occur at the +1 position to Cit residues under physiological conditions,15 alanine was selected as a control again due to its low preference to be cleaved either at its N- or C-terminus to substitute Cit residues. AAZAAK and AAZBAAK and their above-mentioned control sequences were also designed to check whether the position of X/B has a considerable impact on fragmentation. AA(X/B)ZAAK series were prepared to study the cleavage preference of the third bond, while AAZ(X/B)AAK was prepared to study that of the fourth bond. The latter may offer useful information regarding the dependence of fragmentation characteristics on the position of X residues. Altogether, 225 model peptides were synthesized.

**Scheme 1. Literature-Based Two-Step Mechanism for the CID MS/MS Fragmentation of Cit- or Hci-Containing Peptides**

**Synthesis.** Model peptides with a free carboxyl terminus were synthesized using a Multipin Peptide Synthesis Kit (Chiron Technologies Pty Ltd., Clayton VIC, Australia) on macrocrows (Mimotopes Pty Ltd., Mulgrave VIC, Australia; SynPhase L-series lantern, Wang linker, nominal loading: 15 μmol) with Fmoc/tBu strategy following the manufacturer’s instructions. Peptidemodifying Cys, Met or Trp were cleaved from the intermediates using a mixture of 82.5% trifluoroacetic acid/5% phenol/5% water/5% thioanisole/2.5% ethanedithiol, all other peptides were cleaved by a mixture consisting of 95% trifluoroacetic acid/2.5% water/2.5% triisopropylsilane for 2 h at room temperature.

Freeze-dried peptide samples were diluted to 1 μM final concentration with water prior to LC–MS analysis. To produce carbamidomethylated cysteine residues, heptapeptides containing Cys were reduced by treatment with dithiothreitol (2 μL, 200 mM in water) for 30 min at 37 °C and alkylated with iodoacetamide (2.5 μL, 200 mM in water) for 30 min at room temperature.

**UPLC–MS/MS Conditions.** An 8 min long gradient elution was used for separation of the peptides on a Waters Acquity UPLC BEH C18 1.7 μm column (2.1 × 50 mm, Wexford, Ireland). The column temperature was set to 40 °C. Eluent A was composed of 0.1% formic acid, and eluent B was composed of 80% acetonitrile and 0.1% formic acid. The following gradient was used: 0 min 2% B, 1 min 2% B, 4 min 50% B, 4.1 min 90% B, 4.5 min 90% B, 4.6 min 2% B, 8 min 2% B. The flow rate was 300 μL min⁻¹.

**Materials and Methods**
Tandem mass spectra were acquired on a Q-Exactive Focus Hybrid Quadrupole-Orbitrap instrument (Thermo Scientific, Bremen, Germany) using higher energy collision-induced dissociation (HCD). Data acquisition and processing were done with Thermo Scientific XCalibur 4.1. Positive mode was used, and a dd-MS2 (Discovery) method was applied. An isolation window of 2 m/z was employed for precursor selection. The same collision energies (10 and 15 eV) were used for the whole peptide set so that the results could be directly compared. The increase of collision energies was tested above 15 eV but was not found to be beneficial due to loss of information regarding fragments with higher m/z. The AGC target was set to the recommended 5e5 value for full MS and 1e5 for MS/MS. Resolution was set to 70000 for both full and tandem MS. The default charge state was 2, as in most proteomic applications these analytes produce the most fragment-rich spectra. Singly charged species are frequently excluded to enhance selectivity, and triply charged peptides were not present with a reasonable abundance.

**Data Evaluation.** Tandem mass spectra originating from doubly charged precursors were extracted from the chromatograms and visualized by mMass 5.5.0.35 Fragments that are usually generated by collision-induced dissociation (b- and y-ions)34 were automatically identified in the spectra using a 5 ppm mass accuracy limit. We calculated the portion of each amide bond cleavage as a relative percentage by the following formula similar to that of Kapp et al.:35

\[
\text{rel} \%_i = \frac{\sum_{k=1}^{K} b_k^{+} + y_k^{+}}{\sum_{i=1}^{N} \sum_{k=1}^{K} (b_i^{+} + y_i^{+})}
\]

where rel\%_i is the relative percentage of amide bond cleavage at the ith position, N is the total number of cleavage sites, k is the charge state of the fragment ion, K is the total number of charge states, and b^+ and y^+ are the absolute intensities of b- and y-type ions with k charges at the ith cleavage site.

Only these so-called backbone fragments were considered, and the frequently observed peaks corresponding to neutral losses were omitted. Isotope peaks and multiply charged fragments were also taken into account. Spectra were also examined manually to exclude ambiguities of annotation.

**Statistical Analysis.** As the sample and control data sets did not follow a normal distribution or sometimes had significantly differing variances, robust statistical tests that do not require the equality of population variances and less sensitive to the violation of normality were used to statistically evaluate our results. The equality of expected rank variances was tested by the Welch-type O’Brien and Levene tests.36 Robust testing of the hypothesis of stochastic equality was done with a Fligner-Policello test with Welch-like degrees of freedom,36 a Welch test on ranks, and a Brunner-Munzel test.36 In these cases, the confidence intervals for the difference of population means between the sample and control data sets could not be determined; therefore, the stochastic superiority is given instead as a measure of “strength” for the effects

\[
A_{12} = P(X_1 > X_2) + P(X_1 = X_2)/2
\]

where A_{12} is the stochastic superiority of group 1 (sample) to group 2 (control), P is the probability, X_1 are the data from group 1, and X_2 are the data from group 2. Both point and interval estimates were made for the value A_{12}. As an example, an estimated A_{12} ~ 0.8 means that by randomly selecting an X_1 and X_2 value there is a probability of ~80% that the X_1 has a higher value. One could also use the stochastic difference to characterize the probability by which a randomly selected X_1 value is higher than the X_2 value. The stochastic difference can be obtained by a scale transformation from \( \hat{A}_{12} \):

\[
\hat{\delta}_{12} = 2\hat{A}_{12} - 1
\]

By definition, \( \hat{\delta}_{12} \) can range between \([-1;1]\) or can be given in percentage \([-100\%;100\%]\). It may be important to note that these \( \hat{\delta}_{12} \) values are only representing a probability of difference and not the strength of the difference between the two population. We used the software ROPstat for the statistical analyses.39

**RESULTS AND DISCUSSION**

We synthesized 225 model peptides to quantify the effect of citrulline and homocitrulline residues on the tandem mass spectra of citrullinated and carbamylated peptides. We previously observed that Cit residues enhance the amide bond scission at their C-terminus in MS/MS.37 To the best of our knowledge, however, there are no data on Hci cleavage preferences. Therefore, in this work, we calculated all of the amide bond cleavage ratios of our new series of peptides containing Cit and Hci residues in the third and fourth position in the sequence and compared them to their variants containing alamines at these positions. We incorporated all the possible natural amino acids as well as oxidized methionine and carbamidomethylated cysteine directly preceding and following Cit, Orn, Hci, and Lys residues to assess the possible differences in cleavage preferences caused by the side chains.

Our results are in great concordance with our previous studies.34,41 As a demonstration of the citrulline effect, the comparison of the tandem mass spectrum of AAXHAAK and its control AAAHAAK can be seen in Figure 1. The intensity of y_i b_j ion corresponding to a cleavage at the Cit C-terminus is 19% higher than that of the control. It is especially interesting
that Cit-His cleavage is more favored than that of His-Ala although His residues were described to prefer fragmentation at their C-terminus.\(^3^2\)

Our results demonstrate that Cit residues in the third position substantially and statistically significantly increase the cleavage of the third amide bonds (\(\delta_{12}^\text{Cit} = 87\%\)) at 10 eV (Table 1). This effect has a very similar strength at 15 eV (\(\delta_{12}^\text{Cit} = 86\%\)) as it is also summarized in Figure 2A.

Although the increase of cleavage preference of the fourth amide bonds at Cit residues is considerable for the majority of Z (\(\delta_{12}^\text{Cit} = 92\%\)) (Table 1, Figure 3A) and statistically significant at 15 eV, there are only minor differences at 10 eV (\(\delta_{12}^\text{Cit} = 42\%\)).

**Table 1.** Stochastic Difference (\(\delta_{12}\)) of Cleavage Products (%) C-Terminal to the Respective Amino Acid Compared to the Alanine-substituted Control\(^a\)

| CE = 10 eV | Xxx in 3rd position | citrulline | ornithine | homocitrulline | lysine |
|-----------|--------------------|-----------|-----------|----------------|------|
|           |                    | 87***     | 98***     | 82***          | 32+  |
|           | Xxx in 4th position| 42**      | 99***     | -48***         | 10   |
| CE = 15 eV| Xxx in 3rd position| 86***     | 94***     | 80***          | 58***|
|           | Xxx in 4th position| 92***     | 100!      | 33*            | 71***|

\(^a\)Notation: +, \(p < 0.10\); *, \(p < 0.05\); **, \(p < 0.01\); ***, \(p < 0.001\); !, \(p\)-value cannot be determined. Xxx stands for Cit, Orn, Hci or Lys.
compared to the controls. The Cit effect was found to be less strong for the fourth position which could be explained by secondary fragmentation processes that might only be activated at higher collision energies.

Similar tendencies could be observed for Hci residues as well (Figure 2C). The values of increase are somewhat lower than that of for Cit residues but not significantly different from Cit effect when the Hci was in the third position. However, the increase was not only absent but there was a slight decrease ($\delta_{12} = -48\%$) in cleavage compared to the controls at 10 eV for Hci residues in the fourth position (Figure S2). A minor increase was observed with acceptable statistical significance if the collision energy was set to 15 eV ($\delta_{12} = 33\%$), as shown in Figure 3C, but there were several connections where the control displayed higher rel% values.

We also evaluated the data for the peptides that contain an Orn or a Lys as a substitute for Cit and Hci, respectively. These peptides could be derived by a loss of isocyanic acid from the original sequences. Orn residues displayed a statistically significantly higher increase compared to Cit residues (Figure 2B). The Orn effect was also strong when the Orn was at the fourth position (Figure 3B). The decrease of amide bond scission N-terminal to Orn was also stronger that than of the other amino acids.

On the contrary, Lys residues at the third position only produced a slightly significant and very small increase of bond scission preference ($\delta_{12} = 32\%$) and not a significant one in the fourth position at 10 eV ($\delta_{12} = 10\%$) (Table 1). This may indicate that such a low level of collision energy was not enough to efficiently fragment Lys-containing peptides. Although previous studies implicate that Orn displays a very similar or even higher proton affinity than Lys despite the higher +1 effect of the latter one, we found that while Orn residues readily fragmented even at 10 eV, Lys residues showed a lower level of fragmentation (compare Figure S1B with S1D). This may imply a higher proton affinity and, thus, a higher level of proton sequestration by Lys in accordance with both acid–base theoretical considerations and the mobile proton hypothesis of peptide fragmentation. On the other hand, a statistically significant but moderate increase was observed at 15 eV for both the third and fourth position ($\delta_{12} = 58\%$ and $\delta_{12} = 71\%$, respectively), which suggests again that a cleavage C-terminal to Lys residues became slightly favored at a higher collision energy (Table 1, Figures 2D and 3D).

The comparison of the effect of Orn and Lys residues on the fragmentation processes indicates that while the mechanism proposed by our group for the Cit effect which was explained by a Cit to Orn transition followed by Orn effect might be true, it can be conflicting to use it for homocitrulline residues. We propose that the reaction could also follow a one-step mechanism, at least in the case of homocitrulline.

We also investigated the alteration of cleavage products N-terminal to Cit/Hci/Orn/Lys residues and found that an amide bond cleavage at this position is unequivocally unfavored and suppressed compared to the control sequences (Table S1).

To highlight the effect of the Z amino acids on the cleavage preference, we calculated and plotted the difference between the Cit/Hci/Orn/Lys and their alanine-substituted variant for each Z (Figures S3 and S4). These results indicate that for Cit-Zzz connections (Cit in third position) there were no substantial differences between the Z amino acids following the Cit residue in the sequence at neither 10 nor 15 eV. These differences varied between $5\text{--}26\%$ and $2\text{--}22\%$, respectively. In the case of Cit in the fourth position, a very low positive increment ($0\text{--}2\%$) was measured for the majority of Z compared to the controls at 10 eV. For $Z = F$, $Q$, $W$, and $P$, a low decrease was observed compared to the controls. At 15 eV, all connections displayed a positive rel% difference ($2\text{--}15\%$) with the exception of $Z = H$.

On the other hand, a very high difference ($63\%$) was found between Orn-His and Orn-Ala connections at 10 eV where Orn was at the third position, while other Orn-Zzz connections were not considerably different from each other and the values ranged between 13 and 34%. At 15 eV, this discrepancy was eliminated: $5\text{--}32\%$ difference was measured with no remarkable outliers (Figure S3C,D). Orn residues preserved their effect at the fourth position (Figure S4C,D).

For Hci residues, the effect seemed to be occurring only when Hci was at the third position. The cleavage differences varied between $4\text{--}18\%$ and $4\text{--}21\%$ at 10 and 15 eV, respectively (Figure S3). With Hci at the fourth position, the effect was lacking at 10 eV (only Hci-Glu scored a positive rel% difference). At 15 eV, around half of the amino acid connections were still in the negative range (Figure S4E,F).

As for Lys residues, some connections even showed negative values when Lys was at the third position, with values ranging from $-4$ to $+6\%$ and $-5$ to $+11\%$ at 10 and 15 eV, respectively (Figure S3G,H). With Lys at the fourth positions, 6 out of 25 connections showed a positive rel% difference at 10 eV but 24 out of 25 at 15 eV (Figure S4G,H).

It should also be noted that the lowest values of rel% differences were mostly attributed to Arg and Ala residues ($X/B/O/K$ at third position). Arg is known to “sequestrate” protons and hinder fragmentation, thus giving the lower values. On the other hand, in a previous study, alanine was found to affect the MS/MS fragmentation only in a minor way. That is why we selected this amino acid as a control. Our results indicate, however, that the cleavage N-terminal to Ala residues might be rather strongly favored in at least a few cases. When $X/B/O/K$ were at the fourth position, the lowest values of rel% differences were mostly observed for His and Pro residues. For Pro residues, this can be explained by the high preference to be cleaved at its N-terminus in MS/MS. Histidine also has a similar effect that might be comparable to that of Cit and Hci in these cases.

**CONCLUSIONS**

Special amino acids or posttranslational modifications could alter the fragmentation processes of peptides in a profound way as previously demonstrated for proline residues. Some of these effects could improve or hinder the detection and identification of the peptides. We previously demonstrated an enhanced cleavage preference C-terminal to Cit residues.

Currently, there is a great effort to use the intensity values of the given fragments in the tandem mass spectra of peptides to obtain reliable structural identification and thus reduce false positive identifications. In this work, we quantified the effect of citrulline and homocitrulline residues on the higher energy collision-induced dissociation of tryptic-like model system and found that Cit residues increase the amide bond cleavage ratio at the C-terminus of Cit in a statistically and practically significant manner. The effect was virtually independent of the type of the following natural $\alpha$ amino acid (Z) as opposed to our previous study based on a proteomic data set of Lee et al. In that case, the data suggest a more pronounced effect of Cit-Zzz connections.
work, statistically strong results were only obtained for a small number of Cit-Zzz connections which were the most prevalent ones among native citrullinated peptides. Our results showed that homocitrulline residues produced a considerable effect only in the cases where Hci was in the third position in the sequence, which may indicate that the effect of Hci follows a mechanism that is less favored, in accordance with Scheme 1. Both Cit and Hci effects seemed to be stronger when Cit or Hci were farther from the C-terminus. We also found that Cit and Hci residues (as well as Orn and Lys) always suppress the cleavage at their N-terminus as well.

These results help explain the fragmentation processes of peptides containing a urea moiety and improve tandem mass spectrometry-based proteomic identification by integrating these findings into commercially available software algorithms.

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