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Melting Proteins Confined in Nanodroplets with 10.6 μm Light Provides Clues about Early Steps of Denaturation

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Abstract

Ubiquitin confined within nanodroplets was irradiated with a variable-power CO₂ laser. Mass spectrometry analysis shows evidence for a protein “melting”-like transition within droplets prior to solvent evaporation and ion formation. Ion mobility spectrometry reveals that structures associated with early steps of denaturation are trapped because of short droplet lifetimes.

While irradiating small (~0.05 to 1.0 μm dia.) droplets containing individual ubiquitin molecules with 10.6 μm light from a CO₂ laser, we find that it is possible to induce “melting” transitions and trap early structures associated with denaturation. Mass spectrometry (MS) techniques show that upon ramping the laser power (from 0 to 20 W), the distributions shift from low charge state ([M+7H]⁷⁺ and [M+8H]⁸⁺) to high charge state ([M+9H]⁹⁺ to [M+13H]¹³⁺) species, indicating that the protein has undergone a melting-like transition within the droplet, prior to solvent evaporation and ion formation.

This transition is sensitive to solution pH, suggesting that the droplet behaves as a bulk solution at equilibrium. However, ion mobility spectrometry (IMS) measurements of conformer distributions within charge states reveal that after irradiative heating, rapid, droplet evaporation¹ traps structures associated with initial stages of denaturation, before the equilibrium of denatured states is established. In their classic 1954 paper,² “Conformation changes of proteins,” Lumry & Eyring began by stating that “[t]he term protein denaturation even in its original meaning included all those reactions destroying the solubility of native proteins and has since acquired so many other meanings as to become virtually useless.” The

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ability to denature proteins in droplets and directly examine the structures, stabilities, and dynamics of how ensembles of equilibrium melted states are formed complements traditional spectroscopic methods and provides a rare glimpse into the complex processes that are hidden within cooperative transitions. Our approach may be useful in studying conformation-specific unfolding pathways and enzyme inactivation mechanisms such as those reported by Fernandez-Lafuente and coworkers.³

We illustrate these phenomena with two examples involving ubiquitin, a small, 76-residue protein, because several populations of structures (the native β -grasp fold, extended helical A-state, and unstructured U-state) have been characterized in the condensed phase^{4,5} as well as *in vacuo*.⁶⁻⁸ First, we show that the A-state product of thermal denaturation⁹ is not produced at high laser powers for the $[M+8H]^{8+}$ ion; apparently the complex rearrangement associated with unfolding the native β -grasp structure and subsequent assembly of the hydrogen bond networks associated with the extended helices between Gln⁴⁰ – Arg⁷² that make up the A-state⁵ is impossible during the short lifetime of the droplet. Second, we show that the ubiquitin $[M+11H]^{11+}$ denatured product initially emerges in a conformation that favors a *cis*-configured Glu¹⁸–Pro¹⁹ bond; the mixture of denatured *cis*- and *trans*-configurations observed at equilibrium evolves over longer times.⁹ This is remarkable since the native structure has a *trans*-configured Glu¹⁸–Pro¹⁹ bond;⁴ thus, denaturation to form $[M+11H]^{11+}$ initiates through a *trans*→*cis* Glu¹⁸–Pro¹⁹ isomerization, a process that is often rate limiting;¹⁰ the unfolded *trans*-configured structure observed in equilibrium⁹ must involve a relatively complex mechanism, e.g., it might arise from a subsequent *cis*→*trans* Glu¹⁸–Pro¹⁹ isomerization step.

Figure 1 shows the laser-droplet activation source used in these studies. In this configuration, droplets, produced by electrospray ionization (ESI), pass through an orthogonal IR-laser beam prior to evaporation and ion formation. Changes in protein conformation upon irradiating the droplets are monitored by nested IMS-MS techniques, as described elsewhere.¹¹

Figure 2 shows equilibrium melting data (from ref. 9) obtained by thermally heating a bulk ubiquitin solution (aqueous, pH 3). When electrosprayed at 25 °C, the charge state distribution favors $[M+7H]^{7+}$. As the solution temperature is increased, a new distribution of highly charged ions centered around $[M+11H]^{11+}$ emerges. This transition is consistent with thermally unfolding the native ubiquitin protein; following earlier pioneering ESI-MS studies as a function of solution temperature,¹³ a sigmoidal two-state model fit to the weighted average charge state yields a melting temperature (T_m) of 71 ± 2 °C, in agreement with literature values.⁴

When droplets containing ubiquitin are exposed to a laser beam, mass spectra (also shown in Figure 2) are indistinguishable from those obtained from heated solutions. At 0 W (i.e., allowing droplets to pass through without laser excitation) the charge state distribution is dominated by $[M+7H]^{7+}$, as expected. When the droplets pass through a 11 W laser field, highly charged ions (centered around $[M+11H]^{11+}$) are observed, indicating that a substantial fraction of the protein molecules underwent an unfolding transition. Unfolded structures dominate the spectrum at 17 W. The weighted average charge state distribution for

these ions is also sigmoidal in shape and analysis yields a melting power, $T_p = 10.4 \pm 0.3$ W. The similar shapes of the temperature-induced and droplet-IR-irradiated charge state distribution curves were somewhat surprising to us. One might expect irradiation of droplets to induce immediate desolvation, such that solvent is removed before the structure can change – rendering only the native protein charge state in the mass spectrum. The present results require that irradiated droplets survive long enough for the protein to unfold in solution, prior to droplet evaporation and ion formation. But, does the solution environment within the droplet still influence protein stability as it does in the bulk? To explore this, we changed the pH of the solutions used to produce droplets. Figure 2 shows the weighted average charge state curves obtained for ubiquitin confined in droplets produced from pH = 2.5, 3.0, and 4.0 solutions. The midpoint associated with these transitions shifts to higher laser powers with increasing solution pH, corroborating the idea that solution environment within the droplet influences the denaturation transition.

While the MS analysis shows that our intuition – that irradiated droplets might evaporate so quickly that solution transitions might not be observed – was incorrect, IMS measurements reveal that there are differences between droplet-IR-heating and equilibrium melting studies. An ion's mobility through a buffer gas is related to its shape, or collision cross section.¹⁴ Figure 3 shows IMS cross section distributions for $[M+7H]^{7+}$ recorded at 26 °C, distributions for $[M+7H]^{7+}$, $[M+8H]^{8+}$ and $[M+11H]^{11+}$ formed at a bulk solution temperature of 96 °C, as well as these product ions formed from irradiating droplets with a laser power of 13 W, i.e., in the region where the melting transition appears to be complete. The $[M+7H]^{7+}$ species shows a single peak centered at $\Omega = 1010 \text{ \AA}^2$, consistent with highly folded structures from solutions that favor the native state.⁹ Upon increasing either the solution temperature, or the laser power used to irradiate droplets, the compact structures decrease in relative abundance. A comparison of the decreases in abundances of the $\Omega = 1010 \text{ \AA}^2$ peak, which yielded a value of $T_m = 71 \pm 2 \text{ }^\circ\text{C}$ from prior heated solution studies,⁹ with the present data obtained upon irradiating ubiquitin in droplets shows the decrease in this peak is indistinguishable between these methods. The present data shows that the loss of compact $[M+7H]^{7+}$ conformers from droplets also behaves as a melting transition.

Cross section distributions for other charge states formed from heating bulk solutions or from laser-droplet activation show marked differences. At 26 °C, the $[M+8H]^{8+}$ species exhibits a sharp peak at $\Omega = 1010 \text{ \AA}^2$. When the bulk solution of ubiquitin is heated, this peak decreases in abundance and several broad features at $\Omega \sim 1100 - 1550 \text{ \AA}^2$ increase in relative abundance (from $T \sim 45$ to $60 \text{ }^\circ\text{C}$) and then subsequently decay at temperatures above $T \sim 70 \text{ }^\circ\text{C}$. The decrease of these signals⁹ is associated with the systematic increase in the two sharp peaks at $\Omega = 1635$ and 1650 \AA^2 , previously assigned to the ubiquitin A-state,⁷ a low abundance product of melting. While the A-state is somewhat unexpected, we note that as the temperature is increased from ~ 25 to $96 \text{ }^\circ\text{C}$, the dielectric constant of water decreases from $\epsilon = 78$ to 56 , near the value of $\epsilon = 52$,¹⁵ which is similar to that of 40:60 water:methanol solution that is known to favor the A-state at $26 \text{ }^\circ\text{C}$.^{5,7}

As is the case with heating a bulk ubiquitin solution, broad features, centered at $\Omega([M+8H]^{8+}) \sim 1160$ and $\sim 1350 \text{ \AA}^2$, are also observed upon laser irradiation of droplets; however, in comparison with heated solution data, these features are shifted to lower cross sections,

and it appears that the shapes and intensities of these features are dependent upon on droplet size. When $[M+8H]^{8+}$ is formed from $\sim 0.05 \mu\text{m}$ nanodroplets, we observe two broad features at $\Omega \sim 1150$ and $\sim 1330 \text{ \AA}^2$, with the $\Omega \sim 1150 \text{ \AA}^2$ more populated than the $\Omega \sim 1330 \text{ \AA}^2$. In contrast, when the $[M+8H]^{8+}$ products are formed upon irradiation of larger $\sim 1 \mu\text{m}$ droplets, larger $\Omega \sim 1330 \text{ \AA}^2$ cross section species increase in abundance and the population of $\Omega \sim 1150 \text{ \AA}^2$ ions decreases slightly. The lifetimes of these droplets depend on their initial size, which is related to the nano-electrospray emitter diameter and solution flow rate. Although the exact droplet lifetimes are unknown, previous experiments using theta-glass emitters indicate that a change in emitter tip diameter from $1.5 \mu\text{m}$ to $0.24 \mu\text{m}$ results in a linear decrease in the solution flow rate and a reduction in the droplet lifetime from 9 to 1 μs .¹⁶ The twenty-fold difference in tip diameters used in these experiments should change the droplet lifetimes by well over an order of magnitude. Presumably the short lifetimes favor $[M+8H]^{8+}$ products in the relatively compact $\Omega \sim 1150 \text{ \AA}^2$ region. That is, these species are quenched before the $\Omega \sim 1330 \text{ \AA}^2$ species are formed. When given time to reach equilibrium (as is the case for bulk heating studies), the population of the $\Omega \sim 1330 \text{ \AA}^2$ species increases even more.⁹

An extreme case illustrating the importance of droplet size (lifetime) in producing different structures is observed in the $\Omega \sim 1625$ to 1675 \AA^2 region, where A-state structures show two characteristic peaks. Indeed, this doublet unambiguously denotes the presence of A-state species as a product of equilibrium melting in bulk solutions;⁹ however, we observe no evidence for the A-state upon laser-droplet activation. The inability to form the A-state must be due to the transient nature of a rapidly desolvating droplet, which limits the accessible unfolding products to only those that can form rapidly. The complex changes that are associated with forming the extended regions of helices between $\text{Gln}^{40} - \text{Arg}^{72}$ cannot be carried out during the short lifetime of even the largest $1.0 \mu\text{m}$ dia. droplets.

Other products of laser-induced transitions of ubiquitin confined in droplets provide additional details about the first steps of melting. Figure 3 also shows the cross section distributions for $[M+11H]^{11+}$, the dominant charge state produced at high solution temperatures and high laser power. The most notable products of high-temperature solution melting are the two sharp peaks with cross sections of $\Omega \sim 1800$ and 1875 \AA^2 . While there can be multiple conformations associated with these peaks, Pagel and von Helden used ultraviolet photodissociation of mobility-selected ions to show that the major differences arise from the orientation of $\text{Glu}^{18} - \text{Pro}^{19}$; the $\Omega \sim 1800 \text{ \AA}^2$ species has a *cis*-configured peptide bond whereas $\Omega \sim 1875 \text{ \AA}^2$ species is in the *trans* configuration.⁸ The products formed from $0.05 \mu\text{m}$ dia. nanodroplets show that the *trans* conformer is almost completely absent, as indicated by the shoulder at $\Omega \sim 1800 \text{ \AA}^2$. However, the *trans*-configured species is formed in appreciable amounts upon irradiating $1.0 \mu\text{m}$ dia. droplets. These results require that an early step in the unfolding of the native *trans*-configured $\text{Glu}^{18}\text{-Pro}^{19}$ species involves isomerization of this peptide bond. Inspection of the native structure reveals that the $\text{Glu}^{18}\text{-Pro}^{19}$ residues are in close proximity to the β -sheet formed between $\text{Met}^1\text{-Lys}^6$ and $\text{Glu}^{64}\text{-Arg}^{72}$ residues. These interactions stabilize the β -grasp fold around the hydrophobic core. A *trans* \rightarrow *cis* isomerization of $\text{Glu}^{18}\text{-Pro}^{19}$ leading to elongated structures disrupts these stabilizing interactions.

Melting of isolated molecules in electrosprayed droplets¹⁷ provides the ability to monitor structures, stabilities, and dynamics of species involved in denaturation, and provides a unique way to study structural changes that occur in confined spaces. While the droplets used here are much larger (~0.05 to 1.0 μm dia.) than the length scale of unfolded ubiquitin (10–30 \AA) larger proteins (or confinement in smaller droplets) may introduce some fascinating physical behavior associated with confinement.¹⁸ In the studies presented here, we showed that the protonation state of protein ions that underwent irradiative heating within nanodroplets reflects that of the bulk, but significant structural changes have not equilibrated, as measured using IMS. The kinetic trapping of structures provides a new view of denaturation by capturing “snapshots” of the unfolding pathway. Further characterization of structures and pathways in a range of environments using state-of-the-art condensed and *in vacuo* techniques are likely to provide detailed insights about rearrangements that occur upon protein unfolding, reconciling the notion of an “unstructured” denatured state.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

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Notes and references

1. Lee SW, Freivogel P, Schindler T, Beauchamp J. *J Am Chem Soc.* 1998; 120:11758. Silveira JA, Fort KL, Pierson NA, Clemmer DE, Russell DH. *J Am Chem Soc.* 2013; 135:19147. [PubMed: 24313458]
2. Lumry R, Eyring H. *J Phys Chem.* 1954; 58:110.
3. Rodrigues RC, Ortiz C, Berenguer-Murcia A, Torres R, Fernandez-Lafuente R. *Chem Soc Rev.* 2013; 42:6290. [PubMed: 23059445] Sanchez A, Cruz J, Rueda N, dos Santos JCS, Torres R, Ortiz C, Villalonga R, Fernández-Lafuente R. *RSC Adv.* 2016; 6:27329.
4. Lenkinski RE, Chen DM, Glickson JD, Goldstein G. *Biochim Biophys Acta.* 1977; 494:126. [PubMed: 20153] Cary PD, King DS, Crane-Robinson C, Bradbury WM, Rabbani A, Goodwin GH, Johns EW. *Eur J Biochem.* 1980; 112:577. [PubMed: 6257511] Vijay-Kumar S, Bugg CE, Cook WJ. *J Mol Biol.* 1987; 194:531. [PubMed: 3041007] Wintrode PL, Makhatadze GI, Privalov PL. *Proteins: Struct, Funct, Genet.* 1994; 18:246. [PubMed: 8202465] Briggs MS, Roder H. *Proc Natl Acad Sci U S A.* 1992; 89:2017. [PubMed: 1312711] Jackson SE. *Org Biomol Chem.* 2006; 4:1845. [PubMed: 16688326] Kony DB, Hunenberger PH, van Gunsteren WF. *Protein Sci.* 2007; 16:1101. [PubMed: 17525462]
5. Wilkinson KD, Mayer AN. *Arch Biochem Biophys.* 1986; 250:390. [PubMed: 3022649] Harding MM, Williams DH, Woolfson DN. *Biochemistry.* 1991; 30:3120. [PubMed: 1848787] Pan Y, Briggs MS. *Biochemistry.* 1992; 31:11405. [PubMed: 1332757] Stockman BJ, Euvrard A, Scahill TA. *J Biomol NMR.* 1993; 3:285. [PubMed: 8395271] Cox JPL, Evans PA, Packman LC, Williams DH, Woolfson DN. *J Mol Biol.* 1993; 234:483. [PubMed: 8230227] Brutscher B, Brüschweiler R, Ernst RR. *Biochemistry.* 1997; 36:13043. [PubMed: 9335566]
6. Jockush RA, Schnier PD, Price WD, Strittmatter EF, Demirev PA, Williams ER. *Anal Chem.* 1997; 69:1119. [PubMed: 9075403] Oh H, Breuker K, Sze KS, Carpenter BK, McLafferty FW. *Proc Natl Acad Sci USA.* 2002; 99:15863. [PubMed: 12444260] Koeniger SL, Merenbloom SI, Sevugarajan S, Clemmer DE. *J Am Chem Soc.* 2006; 128:11713. [PubMed: 16939296] Chen S-H, Russell DH. *J*

- Am Soc Mass Spectrom. 2015; 26:1433. [PubMed: 26115967] Wagner N, Kim D, Russell DH. Anal Chem. 2016; 88:5934. [PubMed: 27137645]
7. Wytenbach T, Bowers MT. J Phys Chem B. 2011; 115:12266. [PubMed: 21905704] Shi H, Clemmer DE. J Phys Chem B. 2014; 118:3498. [PubMed: 24625065]
 8. Warnke S, Baldauf C, Bowers MT, Pagel K, von Helden G. J Am Chem Soc. 2014; 136:10308. [PubMed: 25007274]
 9. El-Baba TJ, Woodall DW, Raab SA, Fuller DR, Laganowsky A, Russell DH, Clemmer DE. J Am Chem Soc. 2017; 139:6306. [PubMed: 28427262]
 10. Jackson SE, Fersht AR. Biochemistry. 1991; 30:10436. [PubMed: 1931968]
 11. Merenbloom SI, Koeniger SL, Valentine SJ, Plasencia MD, Clemmer DE. Anal Chem. 2006; 78:2802. [PubMed: 16615796] Koeniger SL, Merenbloom SI, Valentine SJ, Jarrold MF, Udseth H, Smith RS, Clemmer DE. Anal Chem. 2006; 78:4161. [PubMed: 16771547]
 12. Kudaka I, Kojima T, Saito S, Hiraoka K. Rapid Commun Mass Spectrom. 2000; 14:1588. Takamizawa A, Fujimaki S, Sunner J, Hiraoka K. J Am Soc Mass Spectrom. 2005; 16:860. [PubMed: 15907701] Nakamura M, Takamizawa A, Yamada H, Hiraoka K, Akashi S. Rapid Commun Mass Spectrom. 2007; 21:1635. [PubMed: 17465013]
 13. Chowdhury SL, Katta V, Chait BT. J Am Chem Soc. 1990; 112:9012. Loo JA, Loo RRO, Udseth HR, Edmonds CG, Smith RD. Rapid Commun Mass Spectrom. 1991; 5:101. [PubMed: 1666527] Mirza UA, Cohen SL, Chait BT. Anal Chem. 1993; 65:1. [PubMed: 8380538] Benesch JLP, Sobott F, Robinson CV. Anal Chem. 2003; 75:2208. [PubMed: 12918957] Wang G, Abzalimov RR, Kaltashov IA. Anal Chem. 2011; 83:2870. [PubMed: 21417416]
 14. Jarrold MF, Constant VA. Phys Rev Lett. 1991; 67:2994. [PubMed: 10044611] Wytenbach T, von Helden G, Bowers MT. J Am Soc Mass Spectrom. 1996; 118:8355. Mesleh MF, Hunter JM, Shvartsburg AA, Schatz GC, Jarrold MF. J Phys Chem. 1996; 100:16082. Clemmer DE, Jarrold MF. J Mass Spectrom. 1997; 32:577. Wytenbach T, von Helden G, Batka JJ, Carlat D, Bowers MT. J Am Soc Mass Spectrom. 1997; 8:275.
 15. Malmberg CG, Maryott AA. J Res Natl Bur Stand. 1956; 56:1.
 16. Mortensen DN, Williams ER. J Am Chem Soc. 2016; 138:3453. [PubMed: 26902747]
 17. Lee JK, Kim S, Nam HG, Zare RN. Proc Natl Acad Sci U S A. 2015; 112:3898. [PubMed: 25775573] Miller CF, Kulyk DS, Kim JW, Badu-Tawiah A. Analyst. 2017; 12:2152.
 18. Sakaue T, Raphaël E. Macromolecules. 2006; 39:2621.

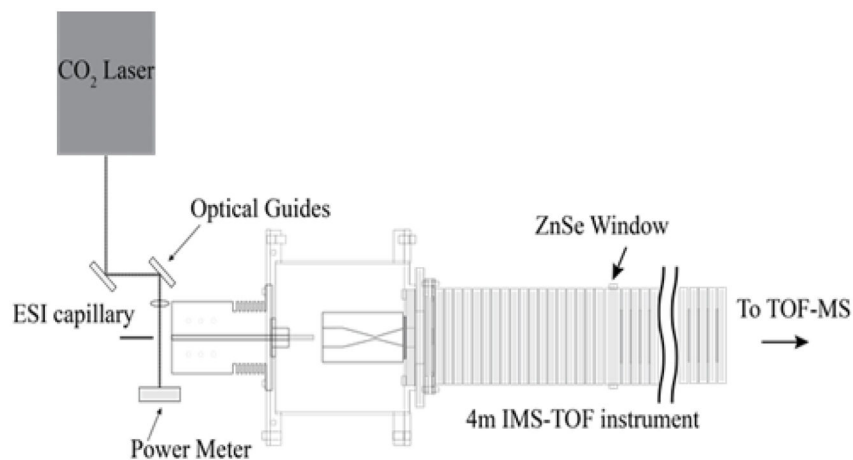


Figure 1. Schematic diagram of the instrument. Droplets diameters are estimated to be ~ 0.05 and 1.0 μm when produced from small ~ 1.0 and 20 μm dia. ESI capillary emitters (see Experimental section in the Supplementary Information). For 10 μM ubiquitin solutions, we estimate that only one in three of the droplets contains a protein molecule. After their formation by electrospray, droplets pass through a CO_2 laser beam focused at the immediate entrance to the instrument orifice. Activation in this region may induce structural changes in the protein which leads to changes in the protein charge state distribution and ion structures. This instrument is also equipped with a ZnSe window in the middle of the drift tube, which allows ions of a known mobility to be excited with 10.6 μm radiation. A series of control experiments in which the laser is focused through the drift tube shows that gaseous protein ions are not activated in the absence of solvent at the laser powers used (See Supplementary Information). This approach has similarities with an elegant “laser spray” technique, in which 10.6 μm light from an IR laser was focused into the metal-capillary tip of an ESI source to heat the bulk solution inside the capillary (see ref. 12 for details).

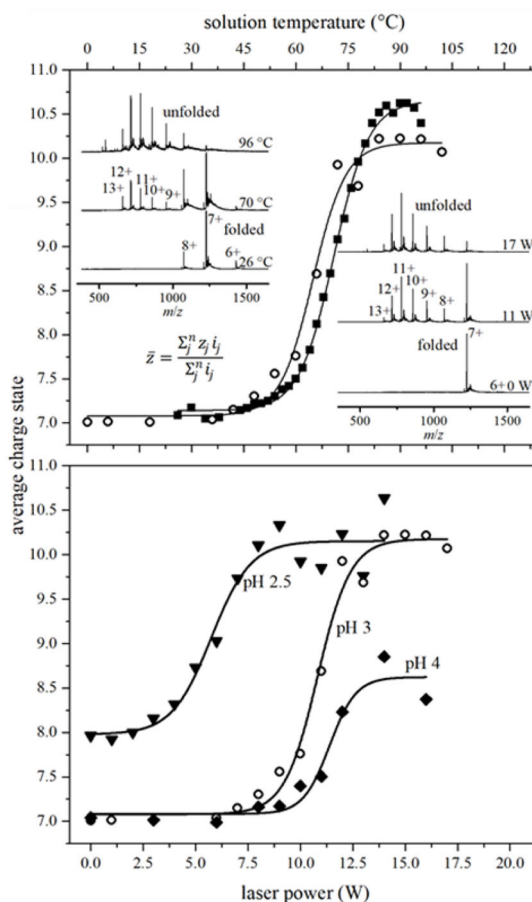


Figure 2.

(top) Average charge state as a function of solution temperature (squares) (from ref. 9) and laser power (open circles) for ubiquitin in aqueous solution at pH 3. Solid lines show the best fit of the data assuming a two-state model with $T_m = 71 \pm 2$ °C and $T_p = 10.4 \pm 0.3$ W. Insets show representative mass spectra at different solution temperatures and laser powers. Upon blocking the radiation, a melted charge state distribution immediately returns to the room temperature distribution, indicating that laser excitation heats only the droplets. The bottom plot shows ubiquitin ions heated in droplets at pH 2.5 (upside down triangles), pH 3.0 (open circles), and pH 4.0 (diamonds) with melting transitions of $T_p = 5.8 \pm 0.3$, 10.4 ± 0.3 , and 11.8 ± 0.3 W, respectively.

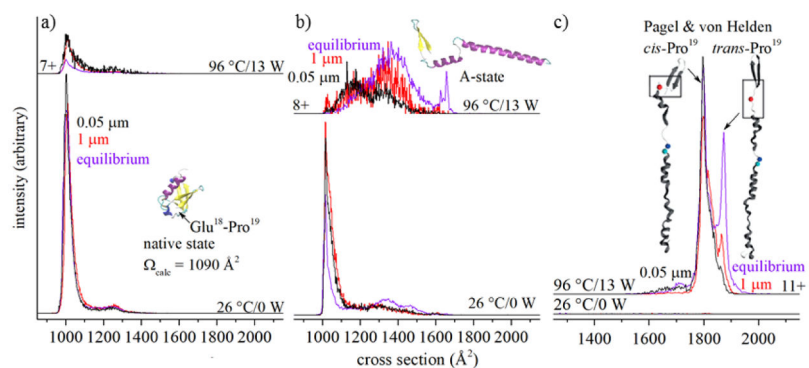


Figure 3. Cross section distributions for (a) $[M+7H]^{7+}$, (b) $[M+8H]^{8+}$, and (c) $[M+11H]^{11+}$ ions of ubiquitin at different temperatures (purple) and laser powers, with black and red lines corresponding to structures that form upon irradiating droplets formed from $\sim 1 \mu\text{m}$ and $\sim 20 \mu\text{m}$ electrospray emitters, respectively. Structures in (c) are adapted from reference 8.