Complementary Benzophenone Cross-Linking/Mass Spectrometry Photochemistry

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ABSTRACT: Use of a heterobifunctional photoactivatable cross-linker, sulfo-SDA (diazirine), has yielded high-density data that facilitated structure modeling of individual proteins. We expand the photoactivatable chemistry toolbox here with a second reagent, sulfo-SBP (benzophenone). This further increases the density of photo-cross-linking to a factor of 20× over conventional cross-linking. Importantly, the two different photoactivatable groups display orthogonal directionality, enabling access to different protein regions, unreachable with a single cross-linker.

Cross-linking/mass spectrometry (CLMS) is a widespread method for investigating protein structure and interactions. Selective cross-linking chemistry is one factor that limits the current resolution of this approach. In CLMS, new covalent bonds are introduced between atoms near in space but not necessarily close in the sequence of the protein. These linkages are then detected by mass spectrometry and function as distance constraints when modeling the structure. Currently, data-density of identified distance constraints is curtailed by the restricted specificity of homobifunctional N-hydroxysuccinimide (NHS) ester cross-linkers predominantly used in CLMS analysis, consequently limiting the level of protein structure detail obtainable. Alternative cross-linking chemistry, including cysteine specific cross-linking, zero-length carbodiimide coupling, acidic specific cross-linking and photo-cross-linking has failed to demonstrate significant increases in CLMS data amount, either alone or in combination. The recent exception is the heterobifunctional, photoactivatable diazirine containing cross-linker sulfo-succinimidyl 4,4′-azipentanoate (sulfo-SDA), which greatly increases cross-linked residue pair number identification in human serum albumin (HSA) from an average of 0.07 links per residue using NHS-ester based cross-linking to 2.5 links per residue using sulfo-SDA. This high-density data allows the computation of HSA domain models from samples of purified protein as well as from crude human blood serum (RMSD to crystal structure of 2.53 and 3.38 Å, respectively).

We aimed to assess whether cross-linker coverage over a protein could be improved upon by expanding our methods of cross-linking to include an additional photoactivatable moiety, the benzophenone group. Benzophenones have previously been used to capture protein interactions in the context of amino acid analogue incorporation, but this has typically involved targeting single incorporation sites, limiting the extent of identified links. A previous study aimed at comparing exogenous bifunctional cross-linkers containing different photoactivatable groups, including diazirine and benzophenone. However, resulting data from using carboxy-benzophenone-succinimide was sparse, with only 3 links observed and a total lack of identified links using a diazirine derivative cross-linker. An earlier study, having synthesized a benzophenone derivative cross-linker and its deuterated equivalent, also identified 3 linked residue pairs. In contrast, when using the diazirine derivative sulfo-SDA in a previous study, we identified 500 cross-linked residue pairs in purified HSA at 5% FDR. We aimed to establish whether the use of a benzophenone based cross-linker could result in a similarly high number of cross-links and whether cross-linking site selection would be influenced by the different chemical nature of the two photoactivatable groups.

Methods

Chemicals and Materials. The cross-linking reagent sulfo-SDA was purchased from Thermo Fisher Scientific (Rockford, IL). HSA (A8763) was purchased from Sigma-Aldrich (St. Louis, MO). Sulfo-SDA (diazirine) was synthesized as follows.

Supporting Information
and N-hydroxysulfo succinimide (50 mg, 0.23 mmol) were stirred in 1 mL of DMF and to this was added EDC-HCl (40 mg, 0.21 mmol) and the mixture stirred overnight. Solvents were removed under reduced pressure and the residue purified using semipreparative RP-HPLC (Agilent 1100 prep-HPLC system equipped with a preparative autosampler (G2260A), preparative scale pumps (G1361A), a fraction collector (G1364B-prep), and a multiwavelength UV detector (G13658)) with preparative flow cell). Material was separated at a flow rate of 20 mL/min on an Agilent RP-C-18 column (21.2 mm × 150 mm, 10 μm particle size), using the following method: 0% B for 5 min, 0% to 80% B in 40 min, 80% B for 5 min, 80% to 0% B in 5 min, 0% B for 5 min, with mobile phase A consisting of water and 0.1% trifluoroacetic acid and mobile phase B consisting of acetonitrile and 0.1% trifluoroacetic acid) to yield the title product as a white solid (20 mg, 0.05 mmol, yield: 84%).

To prepare the gel and the proteins reduced with 20 mM DTT, alkylated with a gradient: 1 min linear increase from 1% B to 9% B; linear increase to 35% B in 169 min; 5 min increase to 85% B. Eluted peptides were sprayed directly into a hybrid linear ion trap-Orbitrap mass spectrometer (LTQ-Orbitrap Velos, Thermo Fisher Scientific). Peptides were analyzed using a “high/high” acquisition strategy, detecting at high resolution in the Orbitrap and analyzing the subsequent fragments also in the Orbitrap. Survey scan (MS) spectra were recorded in the Orbitrap at 100 000 resolution. The eight most intense signals in the survey scan for each acquisition cycle were isolated with an m/z window of 2 Th and fragmented with collision-induced dissociation (CID) in the ion trap. 1+ and 2+ ions were excluded from fragmentation. Fragmentation (MS2) spectra were acquired in the Orbitrap at 7500 resolution. Dynamic exclusion was enabled with 90 s exclusion time and repeat count equal to 1. Mass spectrometric raw files were processed into peak lists using MaxQuant version 1.3.0.5 using default parameters except the setting for “Top MS/MS peaks per 100 Da” being set to 100.

Peak lists were subsequently searched against an HSA sequence database (UniProt P02768) using Xi31 (https://github.com/Rappsilber-Laboratory/XiSearch) for identification of cross-linked peptides. Search parameters for sulfo-SDA search were MS accuracy, 6 ppm; MS/MS accuracy, 20 ppm; enzyme, trypsin; specifity, fully tryptic; allowed number of missed cleavages, four; cross-linker, SDA; fixed modifications, none; variable modifications, carbamidomethylation on cysteine, oxidation on methionine, SDA-loop (SDA cross-link within a peptide that is also cross-linked to a separate peptide). Search parameters for sulfo-SBP were the same except: cross-linker, BENZO (C14H8O2); fixed modifications, none; variable modifications, carbamidomethylation on cysteine and oxidation on methionine. The linkage specificity for sulfo-SDA and sulfo-SBP was assumed to be at lysine, serine, threonine, tyrosine, and protein N-termini at one end, with the other end having specificity for any amino acid residue. False discovery rates (FDR) were estimated following a modified target-decoy search strategy.27,32,33 Cross-link results from the FDR analysis with scores at the peptide-spectrum match level and confidence values at cross-link level are also available in Tables S1 and S2. Cross-links were visualized in the crystal structure of HSA (PDB 1AO6)34 using PyMOL (http://www.pymol.org).

To compare the fragmentation behavior of peptides linked with either sulfo-SDA or sulfo-SBP, we extracted 49 common PSMs that were identified with both sulfo-SBP and sulfo-SDA. A common PSM is defined through the sequences of the two peptides in the cross-link, the cross-link sites and the charge state of the precursor. Next, we performed a two-sided Mann-Whitney-U tests to check for differences in the number of b- and y-type fragments (ignoring loss of H2O and NH3). The tests were highly significant (n = 49, p-value ≤ 0.01 for the b-ions, and ≤ 0.001 for the y-ions, respectively). The mean difference between the number of b- and y-fragments between sulfo-SDA and sulfo-SBP cross-linked peptides is 4. Next, the fragmentation preferences in terms of fragment ion intensities were compared with an adjusted cosine similarity. Instead of computing the cosine similarity of the complete spectrum only the relative intensities of shared fragment ions were considered. The two distributions are clearly distinct indicating conservation of the preferred bond cleavage during fragmentation.

**RESULTS AND DISCUSSION**

Herein we report the synthesis and use of a photoactivatable benzophenone derivative cross-linker, 4-
(sulfosuccinimidylcarboxy)benzophenone (sulfo-SBP), and compare its behavior in the reactivity with proteins to that of sulfo-SDA (Figure 1). While fewer links were detected when using sulfo-SBP compared to sulfo-SDA, complementary photochemistry allowed access to additional parts of protein structure, previously inaccessible using the diazirine-based cross-linker only.

We aimed to benchmark sulfo-SBP against the performance of sulfo-SDA. Sulfo-SBP was first synthesized in a one-step coupling procedure from the corresponding acid (Figure 1B). Purified HSA was cross-linked in triplicate using either sulfo-SDA or sulfo-SBP. Purified HSA, with either no (−) cross-linker, reacted with (+) sulfo-SBP or with (+) sulfo-SDA. Unique cross-linked residue pairs identified using only either sulfo-SBP or sulfo-SDA and those common to both.

Using sulfo-SDA, 792 unique residue pairs were identified (supported by 1874 MS spectra) at 5% FDR (using a score cutoff of 5). In contrast, using sulfo-SBP a total of 173 unique residue pairs were identified (supported by 356 MS spectra) at 5% FDR (score cutoff = 5) (Figure 1E), of which 66 were also found using sulfo-SDA and 107 were unique to sulfo-SBP. For comparison, previous work cross-linking acidic residues and primary amines using 5 different cross-linkers, was based on an average of 52 links per protein, detected in separate studies.

Unambiguous linkage site assignment is dependent on adequate flanking fragmentation events (Figure 2). Manual validation of peptide spectra matches revealed this to be the case for 313/792 (40%) sulfo-SDA cross-linked residue pairs.

![Figure 1. Comparison of sulfo-SDA and sulfo-SBP reaction with HSA. (A) Chemical structures of sulfo-SDA and sulfo-SBP. (B) Synthesis of sulfo-SBP. (C) Experimental design. HSA was cross-linked in triplicate using either sulfo-SDA or sulfo-SBP. (D) Purified HSA, with either no (−) cross-linker, reacted with (+) sulfo-SBP or with (+) sulfo-SDA. (E) Unique cross-linked residue pairs identified using only either sulfo-SBP or sulfo-SDA and those common to both.](image1)

![Figure 2. Fragmentation analysis. (A and B) LC/ESI-MS/MS spectra of the same peptide pair cross-linked with either (A) sulfo-SBP or (B) sulfo-SDA. (C and D) Comparison of b- and y-ion fragmentation of common sulfo-SDA and sulfo-SBP cross-linked peptide pairs. (C) A Mann–Whitney U test indicates highly significant changes in the number of fragments (n = 49, **p-value ≤ 0.01 for b-ions, and ***p-value ≤ 0.001 for y-ions, respectively). (D) Fragmentation spectra similarity computed as a simplified cosine similarity and compared with a random reference distribution.](image2)
and 55/173 (32%) sulfo-SBP cross-linked residue pairs. Looking further at cross-linked peptide pairs common to both sulfo-SDA and sulfo-SBP revealed differences in the numbers of fragments between cross-linkers (Figure 2C); however, cosine similarity was high, suggesting the impact was minimal (Figure 2D).

Cross-link coverage is even across the structure of HSA for both cross-linkers (Figure 3). Very few links have Ca-Ca distances in excess of 25 Å, 5.6% for sulfo-SDA (44/776 residue pairs) and 2.9% for sulfo-SBP (5/172 residue pairs). Although sulfo-SDA has a shorter calculated backbone distance compared to sulfo-SBP (3.9 Å compared to 5.7 Å as predicted for sulfo-SBP), the data suggests that sulfo-SDA has a shorter upper limit distance constraint than sulfo-SDA. This is also demonstrated by 94% of sulfo-SBP linked residue pairs having Ca-Ca distance less than 20 Å, compared to 86% of sulfo-SDA residue pairs. The number of identified linked residue pairs in excess of these proposed upper limits are in agreement with the 5% FDR applied to identifications. Sulfo-SBP data fits well to the crystal structure of HSA (PDB1AO6), corroborating with a previous study that showed benzophenone incorporation into proteins is well tolerated. While HSA does have extensive disulfide bridges, it is not incapable of changing its structure in response to changes in its surface charge. Changing the pH does result in conformational changes as does protein oxidation. The close agreement of cross-link data with the crystal structure suggests that the reduction of charged residues during the initial anchoring step of the reagent does not noticeably disrupt the native fold of the protein.

Both cross-linkers are heterobifunctional, with cross-linking occurring as a two-step reaction: as an anchoring step, the sulfo-NHS ester component of each cross-linker reacts, primarily with the amine groups of lysine residues and protein N-termini, but also with the hydroxyl groups of serine, threonine, and tyrosine. Second, following UV activation, the photoactivatable groups react by insertion into available N−H/C−H bonds at any residue (photo-cross-linking step). Reactivity can therefore be characterized in terms of both the anchoring and photoreacted sites. Distinct anchoring residue hotspots within HSA account for up to 46 unique residue pairs for a single hotspot. In total, the top eight anchoring hotspots (K161, Y162, K183, K186, K214, K223, K456, and K549) account for 26% of all identified residue pairs (Figure S1). Fewer anchoring lysine residues are observed with sulfo-SBP than sulfo-SDA at 23/59 and 45/59 lysine residues, respectively (Figure S1). However, linear peptides with sulfo-SBP and sulfo-SDA modifications (reacted at the NHS-ester) were observed for 55 and 57 (out of 59) lysine residues, respectively, suggesting that the observation of cross-links is not limited by the anchoring reaction but must depend on the photoreaction. Both cross-linkers show distinct regional preferences in the photoreaction step (Figure 3D,E). This is marked in the case of anchoring residues K223, K236, K565 and residues K160, K161, and Y162 (Figure 3D). Although all these residues have reacted with both cross-linkers, the reaction has proceeded in opposing directions. Lysine K565, in particular, was linked to 12 different residues, 5 of which by sulfo-SDA and 7 by sulfo-SBP. Sulfo-SDA linked K565 to residues in a single loop. In contrast, sulfo-SBP linked K565 to residues in 4 separate

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**Figure 3.** Sterical analysis of cross-links. (A) Observed cross-links plotted in PDB1AO6. (B) Ca-Ca distogram of observed cross-links, compared with a random distance distribution (PDB1AO6). (C) Matrix plot of residue pairs across the sequence of HSA. (D and E) Zoomed in regions of HSA in PDB1AO6 with cross-linked residue pairs involving NHS-ester reactive residue hot-spots. (D) Photoreaction exhibits a cross-linker dependent regional preference. Top left, K223. Top right, K565. Bottom left, K236. Bottom right, K160, K161, Y162; (E) Top panel, K375. Bottom panel, K183, Y185 and K186. Sulfo-SDA (red), sulfo-SBP (blue) and common to both cross-linkers (yellow). Figures were generated using PyMOL (http://www.pymol.org).
neighboring alpha helices. In the case of anchoring residue K375 and residues K183, Y185, and K186 (Figure 3E), opposing cross-linking orientation is also evident. The difference here being that there is also considerable overlap, with some degree of linker specificity.

There is an observed cross-linking bias for particular residue types, which are different for sulfo-SDA and sulfo-SBP (Figure S2). Both cross-linkers disfavor polar residues (sulfo-SDA more strongly than sulfo-SDA); however, sulfo-SDA has a preference for acidic residues (Asp, Glu), which are strongly disfavored by sulfo-SBP. Notably, we identify a bias between the benzophenone of sulfo-SBP toward methionine residues, which corroborates a previous study demonstrating a "Magnet Effect" of methionine residues. Sulfo-SBP has a stronger preference for linking hydrophobic residues than sulfo-SDA, in particular glycine and phenylalanine residues.

Photo-cross-linking has emerged as a means to greatly increase the density of cross-linking/mass spectrometry data, which allows interrogation of protein structures in very complex systems, to a level that was previously inaccessible. The mass spectrometry proteomics data have been deposited to the ProteomeXchange Consortium35 (http://proteomecentral.proteomexchange.org) via the PRIDE partner repository with the data set identifier PXD004920.

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ASSOCIATED CONTENT

1 Supporting Information
The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.analchem.6b04938.

1H and 13C NMR spectra for 4-(N-sulfosuccinimidylcarboxy) benzophenone, gel images, replica analysis and site-selection differences following photoactivation of sulfo-SDA and sulfo-SBP (PDF)

List of peptide spectrum matches for Sulfo-SBP and Sulfo-SDA (5% FDR) (ZIP)

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Notes

The authors declare no competing financial interest. The mass spectrometry proteomics data have been deposited to the ProteomeXchange Consortium35 (http://proteomecentral.proteomexchange.org) via the PRIDE partner repository with the data set identifier PXD004920.
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