CrossSearch, a User-friendly Search Engine for Detecting Chemically Cross-linked Peptides in Conjugated Proteins*

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Chemical cross-linking and high resolution MS have been integrated successfully to capture protein interactions and provide low resolution structural data for proteins that are refractive to analyses by NMR or crystallography. Despite the versatility of these combined techniques, the array of products that is generated from the cross-linking and proteolytic digestion of proteins is immense and generally requires the use of labeling strategies and/or data base search algorithms to distinguish actual cross-linked peptides from the many side products of cross-linking. Most strategies reported to date have focused on the analysis of small cross-linked protein complexes (<60 kDa) because the number of potential forms of covalently modified peptides increases dramatically with the number of peptides generated from the digestion of such complexes. We report herein the development of a user-friendly search engine, CrossSearch, that provides the foundation for an overarching strategy to detect cross-linked peptides from the digests of large (≥170-kDa) cross-linked proteins, i.e. conjugates. Our strategy combines the use of a low excess of cross-linker, data base searching, and Fourier transform ion cyclotron resonance MS to experimentally minimize and theoretically cull the side products of cross-linking. Using this strategy, the (αβγδ)4-phosphorylase kinase model complex was cross-linked to form with high specificity a 170-kDa βγ conjugate in which we identified residues involved in the intramolecular cross-linking of the 125-kDa β subunit between its regulatory N terminus and its C terminus. This finding provides an explanation for previously published homodimeric two-hybrid interactions of the β subunit and suggests a dynamic structural role for the regulatory N terminus of that subunit. The results offer proof of concept for the CrossSearch strategy for analyzing conjugates and are the first to reveal a tertiary structural element of either homologous α or β regulatory subunit of phosphorylase kinase.  

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PhK was purified from the psoas muscle of New Zealand rabbits. For which the number of potential products is estimated to increase exponentially with the number of peptides generated by proteolysis (6). As reactants, proteins add considerable complexity to a cross-linking reaction because they contain many reactive amino acid side chains that may be targeted by the cross-linker, leading to their modification by either cross-linking or monoderivatization (covalent modification without cross-linking) (7). In nearly all cases, the number of monoderivatization products will greatly exceed those of cross-linking, particularly for heterobifunctional cross-linkers such as the one used in this study, namely N-[γ-maleimidobutyryloxy]succinimide ester (GMBS). Because cross-linkers are usually selective as opposed to specific (8, 9), GMBS not only may react with a variety of side chains, but the resultant monoderivatization products may in turn form a variety of alternative stable secondary products as will be described herein. Further product complexity results from the fact that modifications may alter the enzyme-catalyzed hydrolysis of each protein in a conjugate by inhibiting proteolysis of amide bonds adjacent to the residue modified; this is particularly problematic for trypsin as lysine is the residue most commonly targeted by conventional chemical cross-linkers (8). The end result is that the heterogeneity of modifications possible in the formation of a conjugate and of its digestion products renders useless the direct comparison of native and cross-linked forms of proteins as a method for detecting cross-linked peptides. Labeling techniques have been developed to reduce the number of potential candidates for analysis of digests of conjugates. These include end labeling peptides with $^{18}$O (10, 11), enrichment with affinity-tagged cross-linkers (12, 13), combined isotope and affinity tagging (14), cross-linking non-labeled and $^{14}$N-labeled protein pairs (15), isotope-labeled cross-linkers (16–18), and MS-cleavable reagents (19–22). All have been used successfully in the analysis of low mass conjugates; however, the success of any labeling strategy is dependent upon many factors, especially the size of the conjugate being analyzed (6). To minimize the number of candidate ions observed in the analysis of large proteins by chemical cross-linking, we have used an alternative approach to the labeling strategies by simply using a minimal stoichiometric excess of cross-linker, which for convenience we term MSEC. The MSEC approach works particularly well with the PhK complex because it binds small organic reagents (a descriptor for most cross-linkers) with high affinity (23–25) as do many other proteins (for a review, see Ref. 26). We have shown previously that PhK specifically binds small hydrophobic cross-linkers, such as the geometric isomers of phenylenediamine (25) and GMBS (23), and that non-active structural analogs of those cross-linkers inhibit its cross-linking by the parent compound (25). Using the MSEC approach with GMBS, we demonstrate herein that this reagent cross-links the hexadecameric PhK complex to form sufficient amounts of a $\beta \gamma$ dimer for analysis and that the formation of this conjugate occurs with a tolerable amount of monoderivatized products.

Although the methods described above streamline somewhat the analysis of cross-linked proteins, predicting the chemical composition of any specific candidate ion requires considerable computational power given the large pool of products possible from the cross-linking and digestion of conjugates. A number of programs have been developed to predict the identity of cross-linked peptides; however, they either require the use of labeled reagents (6, 14–19, 27, 28) or significant user input (29–33). We developed a Web-based, user-friendly search engine, CrossSearch, that both provides a statistical approach to culling masses corresponding to side products of cross-linking and predicts the composition of cross-linked peptides from the mass output obtained for digests of cross-linked proteins. CrossSearch provides the underpinning for an overall strategy for predicting and verifying the composition of cross-linked peptides from both small and large conjugates, especially when used in conjunction with MSEC. Using this strategy in combination with FT-ICR MS, we identified residues involved in the intramolecular cross-linking of the 125-kDa regulatory β subunit of PhK in the context of the entire hexadecameric complex. This cross-linking result represents the first report of a tertiary structural element for either of the two large, homologous α and β subunits of PhK.

**EXPERIMENTAL PROCEDURES**

Proteins—PhK was purified from the psoas muscle of New Zealand White rabbits (34); dialyzed against 50 mM Hepes (pH 6.8), 0.2 mM EDTA, 10% sucrose; and stored at −80 °C. Its concentration was determined by methods described previously (34). The mAbs against the α, β, and γ subunits of PhK were described previously (35, 36), and the anti-calmodulin mAb was from Zymed Laboratories Inc. All other secondary conjugates were from Southern Biotechnology.

Cross-linking—PhK was cross-linked with GMBS essentially as described previously (23) with cross-linking initiated by addition of 125 M Tris (pH 6.8), 20% glycerol, 5% $\beta$-mercaptoethanol, 4% SDS) followed by brief vortexing. The PhK subunits were separated on 6–18% linear gradient polyacrylamide gels and stained with Coomassie Blue. Western blotting of the proteins was performed on PVDF membranes with subunit-specific mAbs as described previously (23). All cross-linking reactions were performed at least twice using different preparations of PhK.

To determine regions of cross-linking in the $\beta \gamma$ conjugate, the cross-linked PhK complex was resolved by preparative SDS-PAGE and stained with Coomassie Blue. The bands corresponding to the $\beta \gamma$ conjugate and the remaining monomeric $\alpha$ and $\gamma$ subunits were excised from the gel, sectioned, and exchanged with three aliquots (each $\times 5$ the volume of the gel slice) of 50 mM ammonium bicarbonate, 50% acetonitrile to remove SDS. The proteins were then reduced in 10 mM dithiothreitol for 1 h at 55 °C and carboxymethylated with 50 mM iodoacetic acid for 1 h in the dark. The gel pieces were washed as described above with 50 mM ammonium bicarbonate followed by several exchanges with 50 mM ammonium bicarbonate,
50% acetonitrile. After removing the last wash, the gels were dried in a SpeedVac (Savant) and treated with trypsin (Promega; 12.5 ng/µl) for 24 h at 30 °C. Peptides were extracted from the gel pieces with 50% acetonitrile, 5% formic acid.

**MS Analyses**—All samples were concentrated on a Centrivac concentrator (Labconco) to a final volume of 20 µl and pressure-loaded onto a C_{18} reverse phase nanocolumn (75-µm inner diameter fused silica packed in house with 9 cm of 100-A, 5-µm, Magic C_{18} particles, Michrom Bioresources). Following a wash with 0.1% formic acid for 15 min at 0.5 µl/min, the column was mounted on the electrospray stage of an FT-ICR mass spectrometer (LTQ FT, ThermoFinnigan, San Jose, CA), and peptides were eluted at an approximate flow rate of 0.3 µl/min over a 120-min period using a gradient of 0–90% acetonitrile (Buffer A = 0.1% formic acid; Buffer B = acetonitrile, 0.1% formic acid). The source was operated at 1.9 kV with the ion transfer temperature set to 350 °C. LC MS data were obtained in a hybrid linear ion trap FT-ICR mass spectrometer equipped with a 7-tesla magnet. The mass spectrometer was controlled using an Xcalibur software package to continuously perform mass scan analysis on the FT instrument followed by MS/MS scans on the ion trap for the six most intense ions with a dynamic exclusion of two repeat scans (30-s repeat duration and 90-s exclusion duration) of the same ion. Normalized collision energy for MS/MS was set to 35%.

**Data Analyses**—For data analyses, .dta files were created on BioWorks Browser version 3.2. The corresponding log file was used to generate a list of parent ions for which the corresponding charges and tandem mass spectra were obtained. The resulting mass list (M + H) was submitted to CrossSearch, the cross-link search engine, to generate a list of potential conjugates. The precursor ion and corresponding tandem mass spectrum matched to each cross-link assignment were visually examined and were further analyzed only if the parent ion was clearly above the noise baseline (-6x) and had an isotopic envelope that was characteristic of a peptide. An additional requirement was that the tandem mass spectrum be performed only on the monoisotopic peak with at least nine fragment ions required for assignment. The remaining tandem mass spectra were then searched by the Sequest algorithm (37) included in Bioworks 3.2 using a modified version of a contaminant protein data base supplied by the vendor (ThermoFinnigan). Data base modification included addition of the polypeptide sequences for the α (P18688), β (P12798), γ (P00518), and δ (P2593) Swiss-Prot/TrEMBL subunits of rabbit muscle PhK. Each search was performed using a 50-ppm error window for the mass of the parent ion with a fragment ion tolerance of 1.00 for peptides arising from both complete tryptic or random hydrolysis. The results of the searches were filtered using the following set of criteria for low confidence: minimum cross-correlation scores (Xcorr) of 1.5, 2.0, and 2.5 for plus 1, 2, and 3 charged ions, respectively, and a delta correlation score (Δcorr) greater than 0.08. Potential conjugates that could be matched to a protein sequence in this data base were rejected. The fragmentation patterns of the remaining candidates were analyzed for consistency with the predicted chemistry of cross-linking.

The tandem mass spectrum of each conjugate was analyzed using a combination of programs that were subsequently verified by an “in-house” spreadsheet. A list of masses from the tandem mass spectrum of each candidate parent ion was uploaded to MS2Links, part of a suite of programs (Collaboratory for MS3D) developed by Dr. Malin Young and co-workers (31) at Sandia National Laboratories, Livermore, CA. Theoretical fragmentation of the conjugate in the positive mode was accomplished using the sequences for each peptide in the conjugate and the masses for each intervening chemical cross-link and its possible fragmentation products using a mass error tolerance not exceeding 0.7 Da for each fragment ion. A modification table was generated for the GMBS cross-linker based on both predicted and observed amide fragmentations of the reagent (23). Potential matching ion assignments for each fragment mass generated by MS2Links were verified using a spreadsheet containing an array of masses predicted for the cross-linker (and fragments thereof) and also those generated for each peptide in the conjugate using MS product, a fragmentation tool in Protein Prospector developed at the University of California, San Francisco (38). Final assignments were also checked by hand for verification and accuracy.

**Cross-link Search Engine**—The CrossSearch engine was constructed with an intuitive XHTML/PHP-based “front end” that allows users to enter a peptide sample, information on sample preparation techniques, and masses identified by MS. The “back end” comprises an MySQL data base containing enzyme splicing information, tags for reactive amino acids, cross-linker mass data, and rule sets defining possible peptide reactants from digests of cross-linked proteins, the chemistry of cross-linking for the reagent chosen, and potential products and side products of the cross-linking reaction. PHP (scripting language) is used to interface the Web front end with the MySQL back end, and a combination of XHTML and cascading style sheets (CSS) languages are used to ensure compatibility with a wide range of browsers and overall ease of use.

**RESULTS**

We developed a cross-link search engine, CrossSearch, that works in combination with high resolution mass spectrometry to predict chemically cross-linked regions between, or within, proteins by generating a list of hypothetical conjugates and their corresponding masses from digests of the cross-linked proteins. A schematic of the search engine showing its basic operating principles is presented in Fig. 1. The boxes in the left half show the flow of theoretical data generated from user-entered components of the reaction (green), including the identity of the chemical cross-linker and the amino acid sequences of the proteins that are covalently linked (Pr1 and Pr2). The theoretical data are cross-referenced against the experimental data (right half), comprising mass lists obtained from MS analyses of digests of the cross-linked proteins (i.e., Pr1-Pr2 conjugate) and Pr1 and Pr2 monomers remaining after termination of the cross-linking reaction that serve as controls. Digests of the monomers help to identify, and thus eliminate, side products of the cross-linking reaction in that they are assumed to contain many of the products (and contaminants) observed in the digest of the Pr1-Pr2 conjugate that are not related directly to the cross-linking of Pr1 and Pr2.

The primary function of the search engine is to separate potential candidate cross-linked peptides from all possible side products of the cross-linking reaction. This is carried out in a series of steps, the first of which is to generate temporary data base tables (orange) of experimental masses (E2 and E3) from Pr1 and Pr2 controls and theoretical masses from an array of all possible products for comparison against the experimental mass list for the protein conjugate (E1). To calculate the array of theoretical conjugates and side products, potential reaction sites on Pr1 and Pr2 are determined by first generating corresponding cleavage maps (blue) for each protein (M1 and M2) and then selecting only those peptides that are potentially reactive through use of a rule set defined in the data base (computational and iterative steps shown in yellow).
The protein rule set defines reactive peptides (T1 and T2) as only those containing amino acids with nucleophilic side chains (His, Lys, Arg, Cys, and Tyr) with the C-terminal residue of each peptide being excluded because its modification would be assumed to inhibit cleavage by the protease (1). The user can choose other combinations of reactive amino acids to accommodate the side-chain specificity of different types of cross-linking reagents.

A cross-linker rule set is derived from the chemistries reported for the cross-linker used. As an example, possible products of the reaction between proteins and the heterobifunctional cross-linker GMBS are shown in Fig. 2, which illustrates the large number of products that may be formed. The formation of a single intermolecular cross-link between the identical residues of two proteins can actually result in conjugates having different masses based on the susceptibility of the maleimide ring to hydrolysis after cross-linking (Fig. 2A, products 3 and 5). Potentially competing side reactions are also contained in the cross-linker rule set. The preferential reactions of Cys and Lys with the succinimidyl and maleimido groups, respectively, of GMBS can form at least five relatively stable side products (products 2, 4, 6, 8, and 9) as a direct result of competing reactions between protein side chains and exogenous nucleophiles, including water, for the remaining functional group of the cross-linker. The number of both these possible side products, or monoderivatized peptides, and cross-linked conjugates is far greater than shown if one takes into account the fact that other protein side chains may react with the cross-linker to form stable products, i.e. these reagents are selective, not specific (8). The masses corre-
Fig. 2. Side products of cross-linking. A, Pr Lys ε-amines preferentially react with the N-succinimide functional group of GMBS at near-neutral pH to form product 1. Hydrolysis of the maleimide group (product 2) competes either with cross-linking reactions by Pr nucleophiles (product 3) or with exogenous nucleophiles that are present as contaminants or quench reagents to halt cross-linking at specific times (product 4). Conversion of products 3 and 4 to their corresponding maleimic derivatives occurs after hydrolysis and subsequent ring opening of the S-substituted maleimide rings (products 5 and 6). B, targeting of the maleimido functional group is generally favored by Pr Cys thiols (product 7). Hydrolysis of the succinimide group competes with cross-linking reactions by Pr nucleophiles, resulting in monodervative of the thiol (product 8). A second mass addition to the thiol may be detected after conversion of the maleimide to the corresponding maleimic derivative (product 9).

Corresponding to all these chemical modifications are loaded into tables (T4–T6) for subsequent steps of analysis. The list of monodervative peptides can also be downloaded to compare against cross-link assignments.

The next step is to identify in the experimental mass list of the digested conjugate (E1) those masses corresponding to side products generated from both theoretical and experimental (Pr1 and Pr2 controls) sources. Digests of the Pr1 and Pr2 controls (remaining Pr1 and Pr2 monomers isolated from the quenched cross-linking reaction mixture by SDS-PAGE or other separation methods) contain monodervative, incompletely digested, and non-modified peptides as well as acrylamide contaminants, the last of which are not accounted for in the theoretical list. The masses from the protein controls are loaded into tables (E2 and E3) and together with a list of common contaminants (e.g. human keratins) are compared with E1 to yield the E4 table of masses. The masses of non-modified peptides (M1 and M2) derived from the theoretical cleavage of Pr1 and Pr2 are matched with those from E4 to generate E5 because the digestion products from the Pr1 and Pr2 controls may differ from those of their corresponding counterparts in the Pr1/Pr2 conjugate (e.g. constraints could be imposed on the protease by cross-linking and/or different extents of monodervative). Inasmuch as the extent of monodervative may differ for Pr1 and Pr2 in their respective experimental control (E2 and E3) and conjugate forms (E1), a comparison of E2 and E3 with E1 may not identify all masses corresponding to monodervative peptides. To account for such differences in the monodervative peptide pools, all possible monodervative peptides from Pr1 and
Pr2 are generated theoretically (T7 and T8) by combining data tables T1, T2, and T6 with the combination subsequently identified in E5 thus annotated to yield E6. The masses in table E6 represent those masses that match either cross-linked peptides or cross-linked peptides with different extents of monoderivatization. To predict the possible composition of these products, the experimental masses from E6 are matched against all possible theoretical combinations of cross-linking (obtained in order of sequence from T1–T5, T9, and T10) and monoderivatization (T11).

Theoretical matches for each experimental mass are displayed in two different formats, the first of which is a statistical summary of the analysis showing the number of potential theoretical matches possible for seven different mass error ranges progressing in order of increasing mass accuracy from 0.1 to 0.00001-dalton error. In addition to the possible cross-linking combinations, potential theoretical matches corresponding to monoderivatized and/or non-modified peptides are also shown, providing the investigator the opportunity to either reject or select mass data for further analysis. In the second format, all possible combinations of cross-linking ± monoderivatization are listed in the final prediction table in order of increasing mass error, allowing for further analysis of all possible assignments by fragmentation, labeling, or other methods of choice. A Web-based version of the search engine using several commonly used cross-linkers is now available for beta testing.

Previously we demonstrated that GMBS is an affinity cross-linker of the \( \alpha \beta \gamma \delta \) PhK complex and that it selectively targets the \( \beta \) and \( \gamma \) subunits in activated forms of the kinase (23). In that same study, we isolated intermolecular peptide conjugates from a \( \beta \gamma \) dimer that corresponded to cross-linking between the C terminus of \( \gamma \) and the N terminus of \( \beta \); however, other peptide \( m/z \) signals were also observed that could not be eliminated as contaminants, side products of cross-linking, or intermolecular conjugates of \( \beta \) and \( \gamma \), suggesting that either one or both subunits might be intramolecularly cross-linked within the \( \beta \gamma \) dimer. To test this possibility, PhK was cross-linked with a 10-fold molar excess of GMBS over \( \alpha \beta \gamma \delta \) protomer for 2.5 min at pH 8.2 (Fig. 3A, Lane 2). As described previously, cross-linking resulted in the formation of a major conjugate corresponding to a \( \beta \gamma \) dimer by apparent mass (170 kDa) and cross-reactivity (23). The monoisotopic peak for a peptide with a signal at \( m/z \) 658.3597 (10× above background) was further subjected to a tandem MS step, and the resulting spectrum was compared against common contaminants as well as tryptic and non-enzymatic hydrolysis products of all the PhK subunits using the Sequest algorithm (37) included in Bioworks 3.2 (ThermoFinnigan). The absence of significant matches observed for the 658 \( m/z \) signal in either the Sequest data sets or mass lists obtained from digests of non-cross-linked \( \beta \) and \( \gamma \) subunit controls (A, Lane 2) together indicated the presence of a potential cross-linked peptide.
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Table I
Possible conjugates (±monoderivatization) that may correspond to a single signal (e.g., m/z 2509.3101) identified by CrossSearch from a digest of the PhK βγ dimer cross-linked with GMBS

The two separate cross-linked peptides of each conjugate are shown separated by a colon (:)...

| m/z calculated | Error (ppm) | Sequence | Cross-link | Monoderivatization |
|----------------|-------------|----------|------------|--------------------|
| 2509.3140      | 1.6         | RKVIYGHWVKKGQQQNR | 1 3       | 1 9               |
| 2509.3240      | 5.6         | DYPALR:EILGRRGSSVVR | 1 3       | 1 2 or 8          |
| 2509.3279      | 7.1         | PVTREIVIR:IIYQYRR | 1 3       | 1 9               |
| 2509.3391      | 11.6        | HFSPRGK:KVSGPHNIQLK | 1 3      | 2 or 8            |
| 2509.2705      | 15.8        | IYYQYRR:DPYALRPLR | 1 3       | 2 or 8            |
| 2509.2705      | 15.8        | IYYQYRR:DPYALRPLR | 1 3       | 2 or 8            |
| 2509.2412      | 27.4        | HFSPR:GQQQNRAALFEP | 1 3       |                  |
| 2509.4153      | 41.9        | KIRM:PVREIYRDPYALR | 1 3       |                  |
| 2509.2012      | 43.4        | ROCIK:HFSPRGKFK    | 2 5       | 2 or 8            |

* The numbers indicated in the column refer to the corresponding structure shown in Fig. 2. Side-chain substitutions will vary based on the nucleophilic chains of Arg, Cys, Lys, Tyr, or His being present in the peptides shown for each conjugate.

b Structures with the same mass are indicated as “a” or “b.”

The tandem mass spectrum of the m/z signal at 658 was first analyzed using the best match ion as a template to identify fragment masses. All major peak assignments were made within the error limits of the collision cell (supplemental Table 1). Both the total ion mass and the observed fragmentation pattern confirmed the sequence assignments for each region (21–23 and 1037–1042) of the β subunit and showed the incorporation of 1 mol of the GMBS reagent, which covalently coupled Lys-21 and Arg-1040 (Fig. 4A). Modification of these residues is consistent with the inability of trypsin to hydrolyze the amide bonds immediately following each residue (1). The chemistry of cross-linking was deduced from cleavage products resulting from disruption of the amide bonds of the intervening cross-link (Fig. 4, B and C), demonstrating that Lys-22 and Arg-1040 add, respectively, to the succinimide ester and maleimide groups to form the expected amide linkage and Michael adduct (8). This same mode of addition to each reactive group of GMBS was observed in a previous study showing cross-linking between Lys and Arg side-chain donors from the γ subunit and a peptide mimetic of the β subunit in a PhKβ peptide complex (23). Targeting of the GMBS maleimide functional group by Arg has been attributed previously to the tight initial binding of GMBS by PhK, i.e. affinity cross-linking (23), and is consistent with several reports that suggest side-chain guanidinyl nitrogens may act as nucleophiles in the formation of Arg-containing conjugates (39–41), whereas the kinetically preferred addition of Lys ε-amines over that of other protein nucleophilic side chains to succinimide esters is well characterized (8).

The second best match (~2 × the error of the first) corresponding to a monoderivatized γ267–275 peptide was also evaluated; however, only a small percentage (28%) of the fragments in the tandem mass spectrum shown in Fig. 4D could be assigned to potential fragments of this peptide, and of those few matched, almost all were assigned with overall greater mass error than fragments predicted for the β intramolecular cross-link (supplemental Table 1). The results strongly indicate that the...
The evidence reported herein for cross-linking between the N and C termini of the subunit of PhK represents the first report of a tertiary structural element for this subunit within the PhK complex (Fig. 5).

**DISCUSSION**

The computational requirements for detecting cross-linked peptides from the vast array of products that arise from the cross-linking and subsequent digestion of proteins are considerable, especially when the proteins cross-linked are not small. Several approaches have been developed to predict the identity of potential conjugates from the mass output measured for digests of cross-linked proteins (29–33); how-
ever, they either require a general understanding of the chemistry involved or the use of specific labeling techniques (6, 14–19, 27, 28). Most search strategies reported to date have focused on the analysis of products generated from the digestion of conjugates of relatively low mass (<60 kDa); however, as the mass of a conjugate increases, so does the number of potential covalent linkages it may contain (e.g. intra- and intermolecular cross-linking, monoderivatizations, and combinations of the lot). In fact, a conservative estimate suggests that the number of possible combinations may increase exponentially to the third power of the number of tryptic peptides generated (6). We report herein the development of CrossSearch, a user-friendly database search engine that provides the foundation of an overarching strategy to detect cross-linked peptides from either low or high mass (≥100 kDa) conjugates. Our strategy comprises three major steps: prediction, elimination of side products of cross-linking, and verification of cross-link assignments.

To fully test the capabilities of CrossSearch, cross-linking of the 1.3-MDa PhK complex was carried out with the heterobifunctional cross-linker GMBS followed by analysis of the resulting major conjugate, a βγ dimer. In the first step of the analysis, the extensive database of potential linkage combinations generated by CrossSearch for the 170-kDa PhK βγ conjugate virtually assured multiple assignments for each experimental mass often with several assignments falling within an error range of 5–6 ppm (Table I). Similarly multiple matches for precursor ions within this error range were also predicted for conjugates of the 176-kDa NDC complex (6). Various labeling strategies have been used successfully to reduce the pool of potential cross-links for low mass conjugates (6, 14–19, 27, 28); however, analysis of conjugates with masses over 100 kDa formed by cross-linking of the aforementioned NDC80 complex with both non-labeled and deuterated forms of a cross-linker suggests that such strategies do not adequately reduce the amount of experimental data to a level that readily facilitates manual inspection and verification of all possible cross-link candidates (6). No matter the reagent used, the study herein and those of others (6, 27) demonstrate that a rigorous assessment of the mass error for candidate m/z signals should be carried out to reduce the multiplicity of assignments generated by large conjugate data bases. Such an assessment is particularly pertinent for low intensity signals corresponding to cross-linked peptides. For large protein complexes like PhK, it is difficult to achieve high yields of cross-linking between specific sites on adjacent protein pairs (dimeric conjugates being the easiest to analyze) because dimers are generally consumed in the formation of multiple, larger low yield multimeric conjugates in the ongoing process of cross-linking (1). Our approach using non-modified peptide m/z signals with intensities bracketing those of candidate ions compensates for the dynamic range of error in mass measurements (43, 44), and it allows for removal with high confidence of those assignments with errors 2× the average mass error.

The size of the predicted data base can also be reduced significantly by eliminating the many possible side products of cross-linking, which in sum provide a significant source of potential false positives. CrossSearch provides a statistical basis for eliminating such products by mass matching using an adjustable mass error window, allowing values to be entered corresponding to the accuracy estimated for the mass spectrometer used. Choice of the correct error value is an important parameter: values that are either too small or large result correspondingly in the loss of correct assignments or the incorporation of false positive assignments. To alleviate this complication, a pool of non-modified and monoderivatized peptides is generated independently from the experimental data with the Sequest algorithm using low confidence correlation scores (see “Experimental Procedures”). The use of low correlation scores simultaneously allows for maximum capture of side products (potential false positives) and a means of comparing cross-link assignments from CrossSearch against potential side products generated from Sequest. In the event that hits from either source are observed for a single experimental m/z signal, fragmentation analyses are then carried out to verify the assignment as being correct. Although the methods described above may reduce the number of potential candidates by as much as a factor of 10, the number of remaining candidates may still be very large, depending on the mass of the conjugate and the mole ratio of cross-linker to proteins used in the cross-linking experiment.

An additional strategy used herein to limit the number of potential side reactions is to limit the mole ratio of cross-linker to protein. In screens of the PhK complex against a library of cross-linkers, GMBS was determined to be an affinity cross-linker that bound to PhK and formed significant amounts of a single conjugate at only a 10-fold excess of cross-linker over the αβγδε tetramer (23). This MSEC method used herein is an alternative approach to the use of labeled cross-linkers and has multiple advantages over labeling strategies. (i) Cross-linkers with varying chemistries and cross-linking spans may be used. (ii) It accommodates the use of zero-length cross-linkers, which in the absence of high resolution reference structures are the best reagents for determining actual contact sites between interacting proteins. (iii) Conventional cross-linkers are more economical than their labeled counterparts. The major disadvantage of MSEC is that is shared by all cross-linkers, namely the ability to selectively form a conjugate of interest.

The final step in the analysis, verification of assignments, is carried out using a combination of programs. The fragmentation patterns of candidates are first evaluated using MS2Links (31), a program that allows for rapid analysis of cross-linked peptides, ranking possible assignments for each ion fragment m/z signal in order of error. That program automatically calculates peptide backbone cleavages for the candidate peptide sequences entered; however, the user must provide mass information relating to the cross-linker, its potential cleavage products, and the amino acid side chains that are likely tar-
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ted by the reagent. In our analysis of GMBS, we loaded all possible fragments arising from cleavage of the amide bonds of the reagent (23) as well as those reported for structural analogs of GMBS (45). All assignments in this phase of the analysis are checked manually in what is the most time-consuming part of the process. Recently an algorithm has been developed that may speed up this process by directly scoring the fragmentation patterns of cross-linked peptides and by providing estimates for the probability of false positive predictions; manual inspection of the data is still required albeit on a more limited basis (6).

After eliminating probable false positives and side products from the $\beta\gamma$ conjugate data base, previously reported conjugates corresponding to intermolecular cross-linking between the C terminus of $\gamma$ and the N terminus of $\beta$ were observed (23). In addition to these intersubunit cross-links, we also observed one intramolecular $\beta\gamma$ conjugate, corresponding to cross-linking between the N and C termini of this subunit.

The low number of cross-links detected is consistent with the stoichiometry of cross-linking imposed using the MSEC approach; however, because the total sequence coverage estimated for the conjugate was $\sim \text{50\%}$ (using both Cross-Search and Sequel for verification of assignments), the presence of additional cross-links in the conjugate is certainly possible.

The cross-link reported herein between Lys-22 and Arg-1040 of the $\beta$ subunit provides the first glimpse of the tertiary structure of either of the two large, homologous, regulatory $\alpha$ and $\beta$ subunits of PhK (46). The N-terminal region containing Lys-22 is unique to the $\beta$ subunit, contains the major activating phosphorylation site of PhK (Ser-26) and a second phosphorylation site (Ser-11), and is relatively near ($\sim 12$ Å) the active site of the catalytic $\gamma$ subunit as well as its C-terminal regulatory domain (23). We previously reported that its first 31 residues mediate self-association of the $\beta$ subunit in that either their deletion or the mutation of the phosphorylatable Ser-11 and Ser-26 allows self-association to be observed in yeast two-hybrid screens (23). In those screens deletion of the C-terminal 177 residues of the 1093-residue $\beta$ subunit, which contains Arg-1040, totally eliminated self-association. The most straightforward explanation for those two-hybrid results would be that the full-length wild type $\beta$ subunit is unable to self-associate because its N terminus interacts with its C terminus, thus blocking self-association of the latter. Such a conclusion could not be reached from two-hybrid data, however, because observed homodimeric interactions could represent either intrasubunit or intersubunit interactions, which would be indistinguishable unless a single small region self-associated (47). The possibility of intersubunit interactions is of special concern with PhK because $\beta\beta$ dimers can be readily formed in activated PhK by a short cross-linker (42). Our current data corroborate and extend the previous two-hybrid results by supporting the concept of an intrasubunit interaction between the N and C termini of the $\beta$ subunit. Further the results herein considered together with those two-hybrid data suggest a structural model for PhK in which the regulatory N terminus of the $\beta$ subunit may participate in opposing interactions with the catalytic $\gamma$ subunit and with its own C terminus, depending upon its phosphorylation state. The detection and identification of a cross-linked peptide from the $\beta\gamma$ dimer demonstrates the utility of the CrossSearch strategy in the analysis of large proteins by cross-linking.

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