Enrichment and Site Mapping of O-Linked N-Acetylgalactosamine by a Combination of Chemical/Enzymatic Tagging, Photochemical Cleavage, and Electron Transfer Dissociation Mass Spectrometry*

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Numerous cellular processes are regulated by the reversible addition of either phosphate or O-linked β-N-acetylgalactosamine (O-GlcNAc) to nuclear and cytoplasmic proteins. Although sensitive methods exist for the enrichment and identification of protein phosphorylation sites, those for the enrichment of O-GlcNAc-containing peptides are lacking. Reported here is highly efficient methodology for the enrichment and characterization of O-GlcNAc sites from complex samples. In this method, O-GlcNAc-modified peptides are tagged with a novel biotinylating reagent, enriched by affinity chromatography, released from the solid support by photochemical cleavage, and analyzed by electron transfer dissociation mass spectrometry. Using this strategy, eight O-GlcNAc sites were mapped from a tau-enriched sample from rat brain. Sites of GlcNAcylation were characterized on important neuronal proteins such as tau, synucleins, and methyl CpG-binding protein 2. *Molecular & Cellular Proteomics* 9: 153–160, 2010.

Numerous cytoplasmic and nuclear proteins are post-translationally modified with O-linked β-N-acetylgalactosamine (O-GlcNAc).1 GlcNAcylation is involved in almost all aspects of cellular metabolism (1) and is highly dependent on the nutrient status of the cell (2). The O-GlcNAc modification rivals phosphorylation in both abundance and protein distribution. Recent studies indicate that signaling pathways can be regulated by the interplay of these two modifications at the same or proximal sites on numerous protein substrates (3).

Current understanding of the functions of O-GlcNAc and the function of O-GlcNAcylation and its relationship to phosphorylation is severely hampered by the difficulties in detecting this labile monosaccharide modification. Problems associated with the identification of O-GlcNAc sites include the following. (a) O-GlcNAc is quickly removed by hydrolases during cell lysis. (b) Like phosphorylation, O-GlcNAc is usually present in less than stoichiometric amounts at given sites on protein substrates. (c) O-GlcNAc is readily lost as an oxonium ion during conventional peptide sequence analysis by collision-activated dissociation (CAD) (supplemental Fig. 1). (d) Modified and unmodified forms of the peptide often co-elute during reverse phase HPLC (supplemental Fig. 2a), and the preferential ionization of the unmodified peptide suppresses the signal observed for the corresponding O-GlcNAc-modified peptide (supplemental Fig. 2, b and c).

Several attempts have been made to enrich samples for O-GlcNAc-modified proteins and peptides. Immunooaffinity purification of O-GlcNAc-modified peptides with an antibody (CTD 110.6) has been largely unsuccessful because of low binding avidity (4). Long, wheat germ agglutinin lectin columns (~39 ft) provide some enrichment but also bind strongly to complex glycans (5). A mutant galactosyltransferase (GalT1) has been used to label GlcNAcylated proteins with a ketone-containing galactose analog (6). Following proteolytic digestion, O-GlcNAc-modified peptides were biotinylated with hydrazine chemistry, isolated on a column packed with avidin beads, eluted with free biotin, and sequenced by ETD mass spectrometry. Failure to elute peptides with high efficiency from the avidin column and an inability to direct the fragmentation to the peptide backbone limit the usefulness of this approach. Reported here is an enrichment methodology that (a) is highly specific for O-GlcNAc-modified peptides, (b) provides for efficient release of the captured peptides from an

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1 The abbreviations used are: O-GlcNAc, O-linked β-N-acetylgalactosamine; ETD, electron transfer dissociation; CAD, collision-activated dissociation; GalT1, mutant galactosyltransferase; SCX, strong cation exchange; UDP-GalNAz, UDP-N-azidoacetylglactosamine; PC, photocleavable; MeCP2, methyl-CpG-binding protein 2; PEG, polyethylene glycol; HRP, horseradish peroxidase; LTQ, linear trap quadrupole; AGC, automatic gain control.
affinity support, and (c) facilitates complete characterization of the released peptides by ETD mass spectrometry.

**EXPERIMENTAL PROCEDURES**

Reagents and Chemical Synthesis—All chemicals were purchased from Sigma-Aldrich unless otherwise noted. PC-PEG-biotin-alkyne reagent (Fig. 1a, Reagent 1) containing a photocleavable 1,2-(nitrophenyl)ethyl moiety (PC-PEG-biotin; synthesized according to Olejnik et al. (9) or obtained from Ambergen, Watertown, MA) was prepared by treating the corresponding N-hydroxysuccinimidyl ester with a 10-fold excess of propargylamine (Sigma) in dry methanol at room temperature for 4 h in the dark. The total reaction volume was 20 μl containing 5 mg of PC-PEG-biotin (6 μmol) and 4 μl of propargylamine (60 μmol). Product (PC-PEG-biotin-alkyne) was purified by thin layer chromatography (250-μm silica gel plate, Analtech, Newark, DE) with methanol/chloroform (1:1, v/v) as the mobile phase. The product was located on the thin layer plate by brief exposure (<1 s) to 254 nm UV light and extracted by scraping the thin layer zone into dry methanol. Silica gel was removed by centrifugation. The identity of the product was confirmed by mass spectrometry (supplemental Fig. 4). The yield was greater than 98%. The purified product was stored in methanol at −20 °C until use.

Enzymatic and Chemical Derivatization of O-GlcNAc-modified Peptides, Enrichment, and Photochemical Release of Tagged Peptides—Proteins were digested with trypsin (50:1 substrate/enzyme) in 100 mM ammonium bicarbonate, pH 8 overnight at 37 °C. Trypsin was removed with a 10-kDa molecular mass cutoff membrane (Millipore, Billerica, MA). Solvent was removed under vacuum, and the residue was then resuspended in 120 μl of 10 mM HEPES, pH 7.9 containing 5 mM MnCl₂, UDP-N-acetylactyalgalactosamine (UDP-GalNAz) and 10 units of GalT1. The reaction mixture was incubated overnight at 4 °C, treated with 10 units calf intestine phosphatase (New England BioLabs, Ipswich, MA), incubated for an additional 2 h at room temperature, and then passed through a C₈ spin column (Nest Group, Southborough, MA). UDP-GalNAz and derivatized peptides were eluted in 0.1% TFA, 1% acetonitrile and 0.1% TFA, 80% acetonitrile, respectively. Solvent was removed under vacuum, and the sample was reconstituted in 20 μl solution containing 0.05 μmol of PC-PEG-biotin-alkyne, 10 mM sodium ascorbate, 1 mM tris[(1-benzyl-1H-1,2,3-triazol-4-yl) methyl]amine (in 4:1 t-butanol:DMSO), and 2 mM CuSO₄. The reaction mixture was incubated overnight at room temperature with gentle agitation. To remove excess PC-PEG-biotin-alkyne, the sample was diluted into strong cation exchange (SCX) loading buffer (5 mM KH₂PO₄, 25% acetonitrile, pH 3.0) and then passed through an SCX spin column (Nest Group). The column was washed with several column volumes of loading buffer, and the retained peptides were eluted with high salt buffer (5 mM KH₂PO₄, 400 mM KCl, 25% acetonitrile, pH 3.0). Eluant was adjusted to pH 7 with ammonium hydroxide and then allowed to interact with high capacity avidin beads (Pierce) for 2 h at room temperature. Avidin beads were washed 10 times with PBS solution and twice with 20% methanol, water and resuspended in 70% methanol, water. The suspension was transferred to a thin walled PCR tube, irradiated with 365 nm UV light (2 milliwatts/cm²) (Spectrolite ENF-240C, Westbury, NY) for 25 min at a distance of 10 cm with rotation, and the supernatant was then dried under vacuum and stored at −20 °C. The overall process is also demonstrated in Fig. 1a.

Western Blotting with HRP-conjugated Avidin—α-Crystallin (In-vitrogen) was subjected to chemoenzymatic tagging as described above but without SCX and avidin enrichment. The protein was resolved by SDS-PAGE and transferred to a nitrocellulose membrane. The membrane was blocked with 10% (w/v) bovine serum albumin in Tris-buffered saline (50 mM Tris-HCl, 150 mM NaCl, pH 7.4), incubated with HRP-streptavidin (1:20,000 dilution in TBS with 0.1% Tween 20; Pierce), and visualized by chemiluminescence.

Preparation of Tau-containing Protein Fractions from Rat Brain—Rat brain (3.4 g) was homogenized in 12.5 ml of ice-cold 1% perchloric acid (Polytron homogenizer, Glen Mills, Clifton, NJ). The resulting suspension was incubated on ice for 20 min and centrifuged at 20,000 × g for 20 min. The supernatant was passed through a 1-μm filter and concentrated with a 10-kDa molecular mass cutoff membrane with simultaneous buffer exchange to 10 mM HEPES, pH 7.5. Proteins (1.9 mg) were fractionated on a Superose 12 PC 3.2/30 gel filtration column (GE Healthcare) by using a buffer containing 20 mM HEPES and 50 mM NaCl, pH 7.5. Tau-containing fractions were combined. The total protein amount was estimated by UV absorbance at 280 nm. An aliquot of this material (about 1.3 μg) was used for O-GlcNAc site mapping.

**RESULTS AND DISCUSSION**

**Approach for Highly Specific Enrichment and Site Mapping of GlcNAcylation**—For this enrichment approach (Fig. 1a), proteins were first proteolytically digested. Following this, GlcNAcylated sites were labeled with an azido sugar using the substrate UDP-GalNAz and the galactosyltransferase GalT1.
Because the UDP product formed in this reaction potently feedback inhibits GalT1, alkaline phosphatase was added to the reaction mixture to rapidly degrade UDP and to ensure complete derivatization (supplemental Fig. 3). Peptide: N-glycosidase F, which removes N-linked glycans, was also added to the reaction mixture to ensure that N-glycans with terminal GlcNAc residues were not labeled with GalNAz. Excess UDP-GalNAz was removed using a C18 spin column. GalNAz groups were subsequently biotinylated through a copper-catalyzed 1,3-dipolar cycloaddition reaction of the free azide on GalNAz to an alkyne containing a terminal biotin group and a photocleavable linker (8). After removing excess biotinylation reagent by SCX, enrichment of the tagged O-GlcNAc-modified peptides was performed by avidin affinity chromatography. A major issue with using avidin chromatography is the difficulty in elution of biotin-containing peptides because binding of biotin to avidin is essentially irreversible ($K_a = 10^{15}$ M$^{-1}$). The elution efficiency is also low when using monomeric avidin. Therefore, harsh conditions are generally used for elution from avidin beads; however, this can easily damage O-GlcNAc. Reduction-cleavable biotin is not suitable for this application because the dipolar addition reaction requires strong reducing conditions. Acid-cleavable biotin typically requires treatment with 95% trifluoroacetic acid, which we also found was strong enough to partially hydrolyze O-GlcNAc.

Olejnik et al. (9) previously reported a photocleavable biotin reagent. Based on their findings, we synthesized a photocleavable biotin-alkyne reagent to tag GlcNAc-modified peptides (Fig. 1a, inset). This photocleavable biotin-linked alkyne contains a terminal biotin group, which facilitates enrichment of GlcNAc-modified peptides. In addition, after enrichment with avidin chromatography, the tagged GlcNAc-modified

![Fig. 1. Enrichment of O-GlcNAc-modified peptides by a combination of enzymatic labeling with UDP-GalNAz and chemical derivatization with the photocleavable reagent PC-PEG-biotin-alkyne. a, flow chart showing the overall strategy. Inset, structure of the photocleavable PC-PEG-biotin-alkyne reagent. CIP, calf intestine phosphatase; PNGase F, peptide:N-Glycosidase F; TBTA, tris[1-benzyl-1H-1,2,3-triazol-4-yl]methylamine. b, CAD mass spectrum of a tagged synthetic O-GlcNAc-modified peptide, YSPTgSPSK (gS is O-GlcNAcylated Ser), showing signature ions at m/z 300.2 and 503.1 that result from cleavage at the two sugar ketal linkages and that confirm the presence of the tagged O-GlcNAc moiety. c, ETD mass spectrum recorded on [M + 3H]$^3+$ (m/z 457.2) ions from the same peptide, YSPTgSPSK. Predicted m/z values for ions of type c’ and z’ (monoisotopic and average masses for singly and doubly charged ions, respectively) are shown above and below the peptide sequence. Observed product ions are underlined and also labeled in the spectrum. Ions in the precursor isolation window are labeled with a triangle (△). Brackets enclose ions that correspond to charge-reduced species and fragments derived from them by loss of small, neutral molecules. Product ions that result from loss of an aminomethyltriazole radical are labeled with a circle (○). M represents the tagged peptide.
peptides are efficiently released upon brief exposure to UV light (365 nm) (supplemental Fig. 4), and the released peptides carry a basic aminomethyltriazolyl acetylgalactosamine group. This overall approach is illustrated in Fig. 1a. Shown in Fig. 1b is a CAD MS/MS spectrum recorded on [M + 3H]^{3+} ions (m/z 537.10) from the same peptide, AIPVgSREEKPSSAPSS. Predicted m/z values for ions of type c^{5} and z^{12} (monoisotopic masses and average masses for singly and doubly charged ions, respectively) are shown above and below the peptide sequence. Observed product ions are underlined and also labeled in the spectrum. Ions in the precursor isolation window are labeled with a triangle (▲). Brackets enclose ions that correspond to charge-reduced species and fragments derived from them by loss of small, neutral molecules. Product ions that result from loss of an aminomethyltriazole radical are labeled with a circle (●). c, the tagging approach enables detection of O-GlcNAc-modified proteins by avidin-HRP blotting. This figure shows that tagged α-cristallin provides a strong signal after blotting with avidin-HRP blotting. Upon UV illumination, no signal is observed indicating photocleavage and loss of the biotin tag from α-cristallin.

**Detecting GlcNAcylation in α-Crystallin**—To test our enrichment strategy, we analyzed a mixture containing tryptic peptides from α-crystallin (50 fmol), which has two known O-GlcNAc sites, and bovine serum albumin (2.5 pmol). GlcNAc peptides were tagged with GalNAz followed by photocleavable biotin, enriched, subsequently released from an avidin resin, and analyzed by CAD using an LTQ-Orbitrap mass spectrometer and by ETD using an LTQ XL mass spectrometer. Shown in Fig. 2a is the CAD spectrum recorded on [M + 3H]^{3+} ions at m/z 715 from the tagged carboxyl-terminal tryptic peptide (AIPVgSREEKPSSAPSS) of α-cristallin. Note that the signals labeled as Peak 1 and Peak 2 correspond to the diagnostic oxonium ions, which are discussed above. Shown in Fig. 2b is the ETD MS/MS spectrum recorded on [M + 4H]^{4+} ions for the same peptide. Predicted fragment ions of type c and z are shown above and below the inset in Fig. 2b. Those ions that were observed are underlined. Note that the mass difference between both c^{5} and c^{4} and z^{12} and c^{11} and z^{10}.

**Fig. 2. Characterization of an O-GlcNAc-modified peptide from α-crystallin.** a, CAD mass spectrum showing signature ions characteristic of the derivatized O-GlcNAc moiety. M represents the tagged peptide. b, ETD mass spectrum recorded on [M + 4H]^{4+} ions (m/z 537.10) from the same peptide, AIPVgSREEKPSSAPSS. Predicted m/z values for ions of type c^{5} and z^{12} (monoisotopic masses and average masses for singly and doubly charged ions, respectively) are shown above and below the peptide sequence. Observed product ions are underlined and also labeled in the spectrum. Ions in the precursor isolation window are labeled with a triangle (▲). Brackets enclose ions that correspond to charge-reduced species and fragments derived from them by loss of small, neutral molecules. Product ions that result from loss of an aminomethyltriazole radical are labeled with a circle (●). c, the tagging approach enables detection of O-GlcNAc-modified proteins by avidin-HRP blotting. This figure shows that tagged α-cristallin provides a strong signal after blotting with avidin-HRP blotting. Upon UV illumination, no signal is observed indicating photocleavage and loss of the biotin tag from α-cristallin.
Novel Approach for O-GlcNAc Enrichment and Site Mapping

O-GlcNAc sites are bold with a lowercase g preceding the residue. M(ox), oxidized methionine.

**TABLE I**

| Protein                        | GI number | Peptide sequence | O-GlcNAc site(s) | Novel? |
|--------------------------------|-----------|------------------|------------------|--------|
| Tau                            | 517394    | 33VgSGDTSPFg339  | Ser'^711'        | Yes    |
| α-Synuclein                    | 149037058 | 61EOVTNGGAVgTGVTAVAQKg80 | Thr'^72'        | Yes    |
| γ-Synuclein                    | 122066261 | 59TEKQANAVgSEAVSSYNTAVKg80 | Ser'^67'        | Yes    |
| β-Synuclein*                   | 77404215  | 24QgVgTEAAEKg32  | Thr'^27'        | Yes    |
| Host cell factor C1            | 213385315 | 1147gTVAPGALER1156 | Thr'^1148'  | Yes    |
| Methyl-CpG-binding protein 2   | 149029883 | 438VAAAAGTTTTTTTTTTVAEKg455 | Thr'^443'  | Yes    |
| Spectrin β2                    | 149044856 | 251QgTVAPOMgTTTTTTTTTTTTVAEKg455 | Thr'^334' (or Thr'^444') | Yes |
|                                |           | 231QgTVAPOMgTTTTTTTTTTTTVAEKg455 | Ser'^253'  | No^3  |

* The same peptide is also in γ-synuclein.

z₁ is 589 Da rather than the 87 Da expected for an unmodified Ser residue. We concluded that the first Ser in the peptide carries the tagged O-GlcNAc moiety. Tagged EEKPAVG-TAAPK (gT is O-GlcNAcylated Thr) from the β-chain of α-crystallin was also detected (data not shown). Unmodified BSA peptides were not detected in the analysis indicating that the enrichment proceeded with high selectivity. The level of O-GlcNAc on α-crystallin is ~10% (11). Knowing this, we estimated that recovery of the two O-GlcNAc-modified peptides from α-crystallin was over 90%.

Next, we asked whether this tagging strategy could be used to detect O-GlcNAc at the intact protein level without using mass spectrometry. As shown in Fig. 2c, tagged α-crystallin was readily detectable by avidin-HRP blotting. In contrast, the signal disappeared when the tagged protein was illuminated by UV light before the sample was resolved by SDS-PAGE. We concluded that tagged proteins and proteins bound nonspecifically to the avidin beads can be readily distinguished by blotting samples before and after the photochemical cleavage.

**Detecting and Site Mapping GlcNAcylation in Complex Biological Sample**—The described protocol was used to analyze tau-containing protein fractions obtained from rat brain. These fractions were isolated by a combination of perchloric acid extraction and gel filtration chromatography. Eight O-GlcNAc sites on seven different proteins were identified (Table I and supplemental Fig. 5). Shown in Fig. 3a is the ETD mass spectrum recorded on [M + 3H]' ions from residues 709 to 717 of the tau microtubule-associated protein. Note that the mass difference between both c₃ and c₂ and z₁ and z₅ is 589 Da rather than the 87 Da expected for an unmodified Ser residue. We concluded that Thr'^711' (more commonly known as Thr'^400' according to the most common splice variant) in the carboxy-terminal region of tau is O-GlcNAc-modified. This same site can also be phosphorylated by glycogen synthase kinase-3β (12). These data support our hypothesis that one function of O-GlcNAc on tau is to prevent hyperphosphorylation in normal brain (13, 14). Because we observed limited digestion of tau with trypsin, we anticipate that additional O-GlcNAc sites will be detected when multiple proteases are used to extend the sequence coverage.

From the same tau-containing protein fractions, O-GlcNAc sites on α-, β-, and γ-synucleins (Table I) were also characterized. α-, β-, and γ-synucleins are three small proteins that are expressed primarily in neural tissues (15). Aggregated α-synuclein is a component of filamentous inclusions associated with neurodegenerative conditions, such as Alzheimer and Parkinson diseases (16). Here we report that α-synuclein is O-GlcNAc-modified on Thr'^72', a residue located within a 35-amino acid stretch that forms the protease-resistant core of α-synuclein fibrils (15). This region is prone to self-aggregation and is able to seed the formation of amyloid fibrils (17). It seems likely that GlcNAcylation on this peptide could prevent fibril formation. Interestingly, Shimura et al. (18) reported that the E3 ubiquitin protein ligase parkin binds and ubiquitiates glycrosylated α-synuclein but not the non-glycosylated form in human brain, although the identity and site of the glycosylation were unknown at that time.

Two O-GlcNAc sites were also identified on methyl-CpG-binding protein 2 (MeCP2). One site was unambiguously identified at Thr'^334' (Fig. 3b), whereas the second site was localized to either Thr'^443' or Thr'^444'. This protein is also of interest because it binds to methylated DNA and mediates transcriptional repression through interactions with a histone deacetylase and corepressor Sin3A (19). Mutations in MeCP2 cause Rett syndrome, a developmental disorder characterized by mental retardation, motor dysfunction, and autistic behavior (20). Brain-specific phosphorylation of Ser'^421' in MeCP2 by a Ca²⁺/calmodulin-dependent kinase II-dependent mechanism is thought to regulate a program of gene expression that mediates nervous system maturation (21). Whether or not GlcNAcylation on the nearby residues (Thr'^334' and either Thr'^442' or Thr'^443') alters this process will require further investigation.

In summary, we describe here an efficient enrichment protocol that is highly specific for O-GlcNAc-modified proteins and peptides and that facilitates site mapping of O-GlcNAc-modified amino acids at the low femtomole level by ETD mass spectrometry. The procedure uses a novel photocleavable biotin tag that allows for efficient release of the enriched O-GlcNAc-modified peptides from the solid affinity support.
Fig. 3. Enriched and tagged O-GlcNAc-modified peptides from tau-containing protein fractions isolated from rat brain. a, ETD MS/MS spectrum recorded on $[M + 3H]^3$ ions ($m/z$ 474.2) for the derivatized peptide VVGSDTSPR from the tau protein. b, ETD MS/MS spectrum recorded on $[M + 3H]^3$ ions ($m/z$ 748.1) for the derivatized peptide VAAAAgTTTTTTTTTTAVAEK from MeCP2. Spectra are labeled as described in Figs. 1c and 2b.
In addition, the photocleavage reaction leaves a basic aminomethyltriazole tag at the site of the O-GlcNAc modification. As a result, all modified, tryptic peptides exist in a charge state of 3+ or higher and thus fragment efficiently along the peptide backbone when subjected to ETD. For site-specific O-GlcNAc quantification, a heavy isotope-labeled photocleavable biotin alkyne is currently being synthesized.

Because the enrichment procedure is highly specific, the flow-through from the avidin chromatography can be further enriched for phosphopeptides or for peptides with other post-translational modifications. Note that in this case the use of a phosphatase during the labeling process must be omitted. A flow chart diagramming the sequential enrichment of GlcNAcylated peptides and phosphorylated peptides is shown in supplemental Fig. 6. In a recent large scale study, we applied this serial enrichment protocol (leaving out the alkaline phosphatase) to investigate the interplay of phosphorylation and GlcNAcylaton in the regulation of cytokinesis. By combining the above serial enrichment protocol with stable isotope labeling by amino acids in cell culture, we mapped, quantified, and compared relative site occupancy for over 120 specific O-GlcNAc-modified residues and over 350 phosphorylated residues by tandem mass spectrometry using only 15 μg of sample from a spindle/midbody preparation. This confirms the robustness of our protocol for analysis of complex mixtures.2 Because the sensitivity of the above approach is already on par with that used to study protein phosphorylation, we expect that the method will be a valuable tool for deciphering the “O-GlcNAcome.” Together with other methods (6, 22), we expect a significant increase of known O-GlcNAc sites in the very near future, which will greatly facilitate investigation of the function of GlcNAcylation.

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