Identification and Quantification of Glycoproteins Using Ion-Pairing Normal-phase Liquid Chromatography and Mass Spectrometry*§

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Glycoprotein structure determination and quantification by MS requires efficient isolation of glycopeptides from a proteolytic digest of complex protein mixtures. Here we describe that the use of acids as ion-pairing reagents in normal-phase chromatography (IP-NPLC) considerably increases the hydrophobicity differences between non-glycopeptides and glycopeptides, thereby resulting in the reproducible isolation of N-linked high mannose type and sialylated glycopeptides from the tryptic digest of a ribonuclease B and fetuin mixture. The elution order of non-glycopeptides relative to glycopeptides in IP-NPLC is predictable by their hydrophobicity values calculated using the Wimley-White water/octanol hydrophobicity scale. O-linked glycopeptides can be efficiently isolated from fetuin tryptic digests using IP-NPLC when N-glycans are first removed with PNGase. IP-NPLC recovers close to 100% of bacterial N-linked glycopeptides modified with non-sialylated heptasaccharides from tryptic digests of periplasmic protein extracts from Campylobacter jejuni 11168 and its pgID mutant. Label-free nano-flow reversed-phase LC-MS is used for quantification of differentially expressed glycopeptides from the C. jejuni wild-type and pgID mutant followed by identification of these glycoproteins using multiple stage tandem MS. This method further confirms the acetyltransferase activity of PgL and demonstrates for the first time that heptasaccharides containing monoacetylated bacillosamine are transferred to proteins in both the wild-type and mutant strains. We believe that IP-NPLC will be a useful tool for quantitative glycoproteomics. Molecular & Cellular Proteomics 8:2170–2185, 2009.

Protein glycosylation is a biologically significant and complex post-translational modification, involved in cell-cell and receptor-ligand interactions (1–4). In fact, clinical biomarkers and therapeutic targets are often glycoproteins (5–9). Comprehensive glycoprotein characterization, involving glycosylation site identification, glycan structure determination, site occupancy, and glycan isoform distribution, is a technical challenge particularly for quantitative profiling of complex protein mixtures (10–13). Both N- and O-glycans are structurally heterogeneous (i.e. a single site may have different glycans attached or be only partially occupied). Therefore, the MS1 signals from glycopeptides originating from a glycoprotein are often weaker than from non-glycopeptides. In addition, the ionization efficiency of glycopeptides is low compared with that of non-glycopeptides and is often suppressed in the presence of non-glycopeptides (11–13). When the MS signals of glycopeptides are relatively high in simple protein digests then diagnostic sugar oxonium ion fragments produced by, for example, front-end collisional activation can be used to detect them. However, when peptides and glycopeptides co-elute, parent ion scanning is required to selectively detect the glycopeptides (14). This can be problematic in terms of sensitivity, especially for detecting glycopeptides in digests of complex protein extracts.

Isolation of glycopeptides from proteolytic digests of complex protein mixtures can greatly enhance the MS signals of glycopeptides using reversed-phase LC-ESI-MS (RPLC-ESI-MS) or MALDI-MS (15–24). Hydrazide chemistry is used to isolate, identify, and quantify N-linked glycopeptides effectively, but this method involves lengthy chemical procedures and does not preserve the glycan moieties thereby losing valuable information on glycan structure and site occupancy (15–17). Capturing glycopeptides with lectins has been widely used, but restricted specificities and unspecific binding are major drawbacks of this method (18–21). Under reversed-phase LC conditions, glycopeptides from tryptic digests of gel-separated glycoproteins have been enriched using graphite powder medium (22). In this case, however, a second digestion with proteinase K is required for trimming down the peptide moieties of tryptic glycopeptides so that the glycopeptides (typically <5 amino acid residues) essentially resemble the glycans with respect to hydrophilicity for subsequent separation. Moreover, the short peptide sequences of the proteinase K digest are often inadequate for de novo sequencing of the glycopeptides.

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1 The abbreviations used are: MS, mass spectrometry; IPR, ion-pairing reagents; IP-NPLC, ion-pairing normal-phase liquid chromatography; RT, retention time; wt, wild-type; TFA, trifluoroacetic acid; BPC, base peak chromatograms; UDP, uridine diphosphate; I.D., internal diameter.
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Glycopeptide enrichment under normal-phase LC (NPLC) conditions has been demonstrated using various hydrophilic media and different capture and elution conditions (23–28). NPLC allows either direct enrichment of peptides modified by various N-linked glycan structures using a ZIC®-HILIC column (23–27) or targeting sialylated glycopeptides using a titanium dioxide micro-column (28). However, NPLC is neither effective for enriching less hydrophilic glycopeptides, e.g. the five high mannose type glycopeptides modified by 7–11 monosaccharide units from a tryptic digest of ribonuclease b (RNase B), nor for enriching O-linked glycopeptides of bovine fetuin using a ZIC-HILIC column (23). The use of Sepharose medium for enriching glycopeptides yielded only modest recovery of glycopeptides (28). In addition, binding of hydrophilic non-glycopeptides with these hydrophilic media contaminates the enriched glycopeptides (23, 28).

We have recently developed an ion-pairing normal-phase LC (IP-NPLC) method to enrich glycopeptides from complex tryptic digests using Sepharose medium and salts or bases as ion-pairing reagents (29). Though reasonably effective the technique still left room for significant improvement. For example, the method demonstrated relatively modest glycopeptide selectivity, providing only 16% recovery for high mannose type glycopeptides (29). Here we report on a new IP-NPLC method using acids as ion-pairing reagents and polyhydroxyethyl aspartamide (A) as the stationary phase for the effective isolation of tryptic glycopeptides. The method was developed and evaluated using a tryptic digest of RNase B and fetuin mixture. In addition, we demonstrate that O-linked glycopeptides can be effectively isolated from a fetuin tryptic digest by IP-NPLC after removal of the N-linked glycans by PNGase F.

The new IP-NPLC method was used to enrich N-linked glycopeptides from the tryptic digests of protein extracts of wild-type (wt) and PglD mutant strains of Campylobacter jejuni NCTC 11168. C. jejuni has a unique N-glycosylation system that glycosylates periplasmic and inner membrane proteins containing the extended N-linked sequon, D/E-X-N-X/S/T, where X is any amino acid other than proline (30–32). The N-linked glycan of C. jejuni has been previously determined to be GalNAC-α1,4-GalNAC-α1,4-[Glcβ1,3]-GalNAC-α1,4-GalNAC-α1,4-GalNAC-α1,3-Bac-β1 (BacGalNAC4,Glc residue mass: 1406 Da), where Bac is 2,4-diacetamido-2,4,6-trideoxyglucopyranose (30). In addition, the glycan structure of C. jejuni is conserved, unlike in eukaryotic systems (30–32). IP-NPLC recovered close to 100% of the bacterial N-linked glycopeptides with virtually no contamination of non-glycopeptides. Furthermore, we demonstrate for the first time that acetylation of bacillosamine is incomplete in the wt using IP-NPLC and label-free MS.

EXPERIMENTAL PROCEDURES

Materials and Reagents—Bovine RNase B, bovine fetuin, dithiothreitol, iodoacetamide, and the ion-pairing reagents were acquired from Sigma. Modified trypsin was purchased from Promega. The polyhydroxyethyl A™ Javelin® guard column (1 cm x 1 mm I.D., 5 μm) and the ZIC®-HILIC guard column (0.5 cm x 1 mm I.D., 5 μm) were purchased from Nest Group (Southborough, MA). The Luna NH2 column (1 cm x 0.3 mm I.D., 5 μm) was purchased from Advantec Industries (Aston, PA). PNGase F was purchased from Roche Applied Sciences (Mannheim, Germany).

Preparation of Periplasmic Protein Extracts of C. jejuni 11168 (wt) and the Isogenic pglD Mutant for IP-NPLC-MS Analysis—C. jejuni glycoprotein extracts were prepared from 4 liters of culture as described (30). Proteins were quantified spectrophotometrically (Nano-dropper ND-1000 Spectrophotometer; Thermo Fisher Scientific), adjusted to a concentration of 20 μg/μl using pure water (Milli-Q system, Millipore Corp.) and were either processed immediately or stored at −20 °C.

Tryptic Digestion of the Standard Glycoproteins and Complex Protein Mixtures and PNGase F Digestion of Fetuin—Bovine RNase B or fetuin at 1 mg/ml was reduced, alkylated, and digested with trypsin, as described previously (29). A 100-μl solution of periplasmic protein extracts of C. jejuni 11168 or the pglD mutant at 320 μg/μl in 50 mM NH4HCO3 was reduced with 8 mM dithiothreitol at 37 °C for 1 h and alkylated with 100 μM iodoacetamide at 37 °C for 30 min. The reagents used for reduction and alklylation were removed by centrifugal ultrafiltration (3000 MWCO) until the samples were at pH 5–6. After the addition of 60 μg of trypsin, the protein solution (120 μl with 50 mM NH4HCO3) was incubated at 37 °C for 16 h. For the O-glycopeptide isolation experiments, half a unit of PNGase F was added to 40 μl of 69 pmol/μl of a fetuin tryptic digest with 50 mM NH4HCO3 and then incubated at 37 °C for 14 h.

IP-NPLC Separation of Glycopeptides—All IP-NPLC experiments were performed either on a CapLC™ capillary LC system or nano-Acquity UPLC® system coupled to a Q-TOF-2™ hybrid quadrupole/TOF mass spectrometer (Waters). The NPLC and IP-NPLC experiments were performed as follows: 1) The sample was suspended in 7.5 μl of 80% ACN + 20% H2O (pH 4.7), if not otherwise stated, with without addition of ion-pairing reagents. All pHs in this report were measured in ACN/H2O with electrodes calibrated in water (33). 2) A gradient of 15% to 30% solvent B (100% HPLC grade H2O) for 5 min, then 30% to 50% solvent B for 5 min was used at a flow rate of 12 μl/min. The column was equilibrated at 15% solvent B for 4 min after each gradient. Solvent A is 100% ACN. The flow was split after the column so that ~400 nL/min was directed to the ESI source to obtain NPLC-ESI-MS or IP-NPLC-ESI-MS chromatograms for each sample. The remainder of the column eluate was directed to a fraction collector or discarded. 3) The peptides eluted from the column between ~1-x min are referred to below as the “non-glycopeptide fractions” whereas the peptides eluted between ~ x-13 min are referred to as the “glycopeptide fraction”. The time x refers to the elution time of the least hydrophilic glycopeptide observed from IP-NPLC-MS of the RNase B tryptic digest and was used in all IP-NPLC experiments as a reference time for determination of the starting elution time of glycopeptides. If not otherwise stated, the IP-NPLC column used in this study was polyhydroxyethyl A, which was used for months with significant deterioration of performance.

MS Analysis of Tryptic Peptide Mixtures from Complex Protein Extracts—The tryptic digest of periplasmic protein extracts of the C. jejuni 11168 and the pglD mutant (40 μg of each sample suspended in 7.5 μl of 80% ACN + 20% H2O + 1% HCl + 10 mM NH4HCO3) were repetitively subjected to IP-NPLC for off-line isolation of glycopeptides. The glycopeptide and non-glycopeptide fractions (~90 μl for each fraction) were collected and dried to ~1 μl which was then resuspended in 100 μl of 0.1% formic acid (aq) and analyzed by label-free nano-flow RPLC-ESI-MS (nanoRPLC-ESI-MS) using a nanoAcuity UPLC system coupled to a Q-TOF Ultima™ hybrid quadrupole/TOF mass spectrometer (Waters). The peptides were first
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| Method          | IPR          | pH     | BPI            |
|-----------------|--------------|--------|----------------|
| NPLC-MS         | None, pH4.7  |        | 3.0e3 ± 1026   |
| IP-NPLC-MS      | 20% CH₃COOH, pH1.1 | | 8.6e3 ± 287   |
| IP-NPLC-MS      | 0.25% HCl, pH1.1 | | 5.0e3 ± 481   |
| IP-NPLC-MS      | 1% HCl + 50 mM NH₄HCO₃, pH1.1 | | 587 ± 31     |
| IP-NPLC-MS      | 0.25% CF₃COOH, pH2.1 | | 3.5e3 ± 35    |

Relative Intensities, %

m/z
TABLE I  
List of identified non-glycopeptides, their calculated hydrophobicity values ($\Delta G_{\text{ac}}, \text{kcal/mol}$), and IP-NPLC retention times from the RNase B and fetuin trypsin digest

Sequences in italic letters are from RNase B; Glu$^0$, Asp$^0$, His$^+$, Lys$^+$, Arg$^+$, N-T NH$_2$ and C-T COO$^-$ are used for calculation of the charged non-glycopeptides (pH 1.1) using MPEXTotalizer (35, 37). A value of 4.1 kcal/mol was further added to the calculated values for peptide xiv of fetuin due to likely salt bridge formation of this peptide (36). A value of 2 kcal/mol was further added to the calculated values for peptides xxiv–xxv due to losses of NH$_3$ at C-terminal, respectively (35, 38). The tryptic digest of RNase B and fetuin mixture was injected with 0.25% HCl in 80% ACN/20% H$_2$O for IP-NPLC-MS. CamC, carboxyamidomethylcysteine.

| Peptide no. | Sequence of identified non-glycopeptides | Calculated hydrophobicity values | IP-NPLC retention time (RT) | Error of RT |
|-------------|-------------------------------------------|---------------------------------|---------------------------|------------|
| i           | (K)VWPR                                    | $-8.0$                          | 3.8                       | 42.7       |
| ii          | (K)AQFVLPSVSVFEAVAAATD$^\text{cam}$CIAK(E) | $-8.5$                          | 4.2                       | 36.4       |
| iii         | (K)FER(Q)                                 | $-8.8$                          | 5.0                       | 0.6        |
| iv          | (K)RFK(I)                                 | $-8.9$                          | 4.3                       | 3.5        |
| v           | (K)QDQQFLFTK(B)                            | $-10.1$                         | 5.1                       | 5.8        |
| vi          | (K)$^\text{cam}$CNLLAEK(Q)                | $-10.3$                         | 5.2                       | 8.1        |
| vii         | (R)$^\text{cam}$CGR(F)                    | $-10.4$                         | 5.5                       | 5.2        |
| viii        | (K)IIVACEGNYPPVHFASV(−)                   | $-10.7$                         | 5.8                       | 2.9        |
| viv         | (K)VHAVEVALATFNAESNGSYLQIEISR(A)          | $-10.7$                         | 5.4                       | 6.9        |
| x           | (K)QYGFS$^\text{cam}$CK(G)                | $-10.9$                         | 5.5                       | 1.7        |
| xi          | (K)YPN$^\text{cam}$CAK(Y)                 | $-11.4$                         | 5.7                       | 4.6        |
| xii         | (K)HLPR(G)                                | $-11.6$                         | 6.2                       | 11.5       |
| xiii        | (R)AHYDLR(H)                              | $-11.7$                         | 6.2                       | 11.5       |
| xiv         | (−)IPLPVPAGYEPA$^\text{cam}$CDDPDEQAALAADVYINK(H) | $-11.9$                         | 6.0                       | 0.6        |
| xv          | (K)ALGGEDVR(V)                            | $-12.0$                         | 5.3                       | 1.7        |
| xvi         | (K)GSIOK(A)                               | $-12.2$                         | 5.9                       | 3.5        |
| xvi         | (K)NACK(K)                                | $-12.2$                         | 6.2                       | 0.6        |
| xix         | (K)TPIVQGSPIPPPVR(V)                      | $-12.7$                         | 5.3                       | 1.7        |
| xix         | (K)$^\text{cam}$CDDPPSDAEV(K)             | $-13.3$                         | 6.7                       | 3.5        |
| x           | (K)HTLQIDSVK(V)                           | $-13.6$                         | 6.6                       | 0.6        |
| xx          | (R)ETGSSK(Y)                              | $-13.8$                         | 6.9                       | 0.6        |
| xx          | (K)TTOANK(H)                              | $-14.0$                         | 7.0                       | 1.7        |
| xxii        | (K)NGQTN$^\text{cam}$CYQYSTMSITD$^\text{cam}$CR(E) | $-14.1$                         | 7.0                       | 0.0        |
| xxii        | (K)$^\text{cam}$CVPVNTFVHESLADVQAV$^\text{cam}$CSQK(N) | $-14.9$                         | 6.5                       | 0.6        |
| xxii        | (K)$^\text{cam}$CDDPPSDAEVR(K)            | $-15.9$                         | 6.9                       | 2.9        |
| xxiv        | (K)$^\text{cam}$CDDPPSDAEVR(K)            | $-16.1$                         | 7.4                       | 0.6        |
| xxii        | (K)QQTQHAEVDG$^\text{cam}$CDHVVLK(Q)      | $-17.9$                         | 7.5                       | 0.6        |
| xxii        | (K)HTFSGASVSSSSEGAEFGVQG(T)               | $-18.4$                         | 7.1                       | 0.6        |
| xxii        | (K)OMHDSRTSASSSSY$^\text{cam}$CNQMMK(S)   | $-18.6$                         | 7.7                       | 0.6        |

loaded onto a 180 µm I.D. × 20 mm 5-µm symmetry® C18 trap (Waters), then eluted to a 100 µm I.D. × 10 cm 1.7-µm BEH130C18 column (Waters) using a linear gradient from 0% to 36% solvent B (ACN + 0.1% formic acid) in 36 min, 36–90% solvent B for 2 min. Solvent A was 0.2% formic acid in water. The peak areas (signal/noise ≥ 3) from extracted ion chromatograms of label-free nanoRPLC-ESI-MS of the glycopeptides isolated by IP-NPLC were used for differential expression analysis of periplasmic glycoproteins between the wt and mutant strains. NanoRPLC-ESI-MS/MS analyses were also performed with data-dependent analysis with a survey intensity threshold of 20 for the total digest and the glycopeptide and non-glycopeptide fractions of the periplasmic protein extract of the pglD mutant of C. jejuni 11168.

The glycopeptide fractions were then analyzed with retention time segments of peptides by nanoRPLC-ESI-MS$^2$ and MS$^3$ using an Ettn MDLC system (Amersham Biosciences AB, Uppsala, Sweden)

![Fig. 1. Isolation of glycopeptides from a RNase B and fetuin trypsin digest using IP-NPLC. Total ion mass spectrum from NPLC-ESI-MS of peptides in the glycopeptide fraction from the trypsin digest injected with the following ion-pairing reagents (IPR): a, none in 80% ACN/20% H$_2$O; (pH 4.7); b, 20% acetic acid in 80% ACN (pH 1.1); c, 0.25% HCl in 80% ACN/20% H$_2$O (pH 1.1); d, 50 mM NH$_4$HCO$_3$ + 1% HCl in 80% ACN/20% H$_2$O (pH 1.1); and e, 0.25% TFA in 85% ACN/15% H$_2$O (pH 2.1). BPI, base peak intensities. Throughout this report, the total ion mass spectrum for NPLC-ESI-MS of the glycopeptide fraction refers to the summed MS scans across the NPLC retention times between the most hydrophobic high mannose type glycopeptide (a) from RNase B (6.5–9.0 min depending on the ion-pairing reagents used [Fig. 2, 2nd top panel]) and the most hydrophilic sialylated glycopeptide from fetuin detected (earlier than 13 min). Non-glycopeptides eluted earlier than the high mannose type glycopeptide (a) of RNase B when HCl or TFA were used as ion-pairing reagents. The trypsin digest contains 14.2 pmol of RNase B tryptic digest + 13.8 pmol of fetuin tryptic digest. The solvent compositions used to dissolve the different IPR, CH$_3$COOH, HCl, and TFA with/without NH$_4$HCO$_3$ are as indicated in each panel of the figure. The pH values listed are those of the samples. All abundant ions that are not labeled in Fig. 1 are singly charged contaminants.

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coupled to an LTQ linear ion trap mass spectrometer (Thermo Fisher Scientific) to acquire MS² and MS³ mass spectra for determination of the glycan composition of the glycopeptides and for protein identification. The peptides were first loaded onto a 300 μm ID. x 5 mm C18 PepMap100™ trap (LC Packings, San Francisco, CA), then eluted off a 75 μm ID. x 5 cm C18 PicoTip™ column (New Objective, Woburn, MA) using the same gradient as described above.

Database Searching for Glycoprotein Identification—The peaklists of MS² spectra of the N-linked glycopeptides of C. jejuni proteins were generated using Xcalibur 2.0.6 and were searched against the NCBI C. jejuni database (2007.09.05) with 12283 entries using the MascotTM search engine (version 2.2.0) (Matrix Science) for protein identification. The database searches contained the following variable modifications: glycosylated asparagine by bacillosamine (2,4-diacetamido-2,4,6-trideoxyglucopyranose with a formula of C10H16N2O4) and O-linked GlcNAc residues, and each of the five high mannose type glycans attached to Asn-60 (34). Fetuin has three O-linked sites (Ser-253, Thr-262, Ser-264, and Ser-323), which have sialylated glycans attached and low O-glycosylation occupancy (12–13). Table I shows the non-glycopeptides (i–xxix) sequences of RNase B and fetuin identified from IP-NPLC-MS analysis representing 94% of the amino acid sequence of RNase B and 63% of the amino acid sequence of fetuin. The mass spectral peaks (i–xxix) labeled in Fig. 1 correspond to non-glycopeptides sequences listed in Table I. Hydrophobicity values, free energies of transfer ΔGoctw (kcal/mol) from n-octanol to water (octw), of non-glycopeptides (i–xxix) were calculated at the sample pH throughout this report using MPExTotalizer based on the Wimley-White water/octanol free energy scale for the 20 amino acids (a more negative energy value indicates a more hydrophilic peptide, Table I) (35–37). Table II lists the isolated glycopeptides (a–w) and their glycoforms from the standard tryptic digest using IP-NPLC. The mass spectral peaks (a–w) labeled in Fig. 1 correspond to the glycopeptides (a–w) listed in Table II.

When no acid was added to the sample (pH 4.7) prior to sample injection, the glycopeptides from RNase B (a–e) were not detected at all, and the fetuin glycopeptides were only observed with low MS signals from the glycopeptide fraction (e.g. the glycopeptide ion (m/z) of fetuin at m/z 1542.89(4+)),

### RESULTS

**IP-NPLC for Isolating Tryptic Glycopeptides of Standard Glycoproteins Using Acids as Ion-pairing Reagents—** A tryptic digest of an RNase B and fetuin mixture was chosen to illustrate the use of IP-NPLC for glycopeptide isolation (Fig. 1). The tryptic digestion of RNase B yields five N-linked glycopeptides: NLTK-GlcNAc2Man5–9, where Man is mannose, and GlcNAc is N-acetylgalactosamine, with each of the five high mannose type glycans attached to Asn-60 (34). Fetuin has three N-linked sites (Asn-81, Asn-138, and Asn-158) and four O-linked sites (Ser-253, Thr-262, Ser-264, and Ser-323), which have sialylated glycans attached and low O-glycosylation occupancy (12–13). Table I shows the non-glycopeptides (i–xxix) sequences of RNase B and fetuin identified from IP-NPLC-MS analysis representing 94% of the amino acid sequence of RNase B and 63% of the amino acid sequence of fetuin. The mass spectral peaks (i–xxix) labeled in Fig. 1 correspond to non-glycopeptides sequences listed in Table I. Hydrophobicity values, free energies of transfer ΔGoctw (kcal/mol) from n-octanol to water (octw), of non-glycopeptides (i–xxix) were calculated at the sample pH throughout this report using MPExTotalizer based on the Wimley-White water/octanol free energy scale for the 20 amino acids (a more negative energy value indicates a more hydrophilic peptide, Table I) (35–37). Table II lists the isolated glycopeptides (a–w) and their glycoforms from the standard tryptic digest using IP-NPLC. The mass spectral peaks (a–w) labeled in Fig. 1 correspond to the glycopeptides (a–w) listed in Table II.

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### Table II

*List of identified glycopeptides and corresponding glycoforms for the N- and O-linked oligosaccharides from the RNase B and fetuin tryptic digest*

| Protein name | Sequence of identified peptides | Oligosaccharide composition | Glycosylation site | Glycopeptide mass |
|--------------|--------------------------------|-----------------------------|--------------------|-------------------|
| a* RNase B   | (RNLT)(D)                      | GlcNAc2 Man5                | Asn-60             | 1690.70           |
| b            |                               | GlcNAc2 Man5                |                    | 1852.76           |
| c            |                               | GlcNAc2 Man7                |                    | 2014.81           |
| d            |                               | GlcNAc2 Man5                |                    | 2176.86           |
| e            |                               | Glc2 NAc2 Man5              |                    | 2338.91           |
| f Bovine fetuin | (KKLCamCPOCam CPCLAP NDSR(V) | Hex6 HexNAc5 SA3           | Asn-138            | 4728.93           |
| g            |                               | Hex6 HexNAc5 SA2            |                    | 4072.70           |
| h            |                               | Hex6 HexNAc5 SA3            |                    | 5002.02           |
| i            |                               | Hex6 HexNAc5 SA2            |                    | 3944.61           |
| j            |                               | Hex6 HexNAc5 SA2            |                    | 4600.84           |
| k            |                               | Hex6 HexNAc5 SA2            |                    | 4891.93           |
| l            | (RPRTGEYDIEIDLETTLT Cam CHVLDPPLA N Cam CBV)(R) | Hex6 HexNAc5 SA2 | Asn-81 | 5875.54 |
| m            |                               | Hex6 HexNAc5 SA2            |                    | 6531.77           |
| n            |                               | Hex6 HexNAc5 SA2            |                    | 6822.86           |
| o            | (RNVHAVEALTFAESNGSYLOLVEISR(A) | Hex6 HexNAc5 SA2 | Asn-158 | 5876.57 |
| p            |                               | Hex6 HexNAc5 SA2            |                    | 6167.66           |
| q            |                               | Hex6 HexNAc5 SA2            |                    | 6458.75           |
| r            | (RVTTCam CTFQGTQVPQGPDPQGAEAPSVAP | Hex6 HexNAc5 SA2 | T288–288 | 7326.60 |
| s            |                               | Hex6 HexNAc5 SA2            |                    | 7817.70           |
| t            |                               | Hex6 HexNAc5 SA2            |                    | 7982.64           |
| u            |                               | Hex6 HexNAc5 SA2            |                    | 8273.91           |
| v            |                               | Hex6 HexNAc5 SA2            |                    | 8639.04           |
| w            |                               | Hex6 HexNAc5 SA2            |                    | 8930.13           |

* The 23 glycopeptides detected by IP-NPLC-MS of ~300 fmol of the RNase B and fetuin tryptic digest have been assigned a letter code (a–w) in order to facilitate identification. The consensus sequence of N-glycosylation and possible O-glycosylation sites are highlighted (12, 13), Cam, carboxamidomethylcysteine.

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Ion-pairing Normal-phase LC for Isolating Glycopeptides

![Graph showing XICs of 834.12(3+), non-glycopeptide (xxiv)](image)

![Graph showing XICs of 846.44(2+), glycopeptide (a)](image)

![Graph showing XICs of glycopeptide (b) and (c)](image)

**Fig. 2.** Effect of acids as ion-pairing reagents on peptide retention in IP-NPLC. Top panel, overlaid extracted ion chromatograms (XICs) of m/z 834.12 (3+) of the hydrophilic non-glycopeptide (xxiv), CKPVNTFVHESLADVQAVCSQK, from the RNase B and fetuin tryptic digest injected with the following ion-pairing reagents (IPR): (a) none; (b) 20% CH₃COOH; (c) 0.25% HCl; (d) 1% HCl + 50 mM NH₄HCO₃; (e) 0.25% TFA; and (f) 1% TFA + 50 mM NH₄HCO₃. Second top panel, overlaid XICs of m/z 846.44 (2+) of the least hydrophilic glycopeptide (a), NLTK-GlcnAc₂Man₅ GlcNAc, from the same tryptic digest injected with the following IPRs: (B) 20% CH₃COOH; (C) 0.25% HCl; (D) 1% HCl + 50 mM NH₄HCO₃; (E) 0.25% TFA; and (F) 1% TFA + 50 mM NH₄HCO₃. Second bottom panel, (G) XIC of m/z 927.48 (2+) of the glycopeptide (b), NLTK-GlcnAc₂Man₅ GlcNAc; (H) XIC of m/z 1008.43 (2+) of the glycopeptide (c), NLTK-GlcnAc₂Man₅ GlcNAc; IPR used for G and H: 1% TFA + 50 mM NH₄HCO₃. Bottom panel, total ion chromatogram from IP-NPLC-ESI-MS of the standard tryptic digest injected with 1% TFA + 50 mM NH₄HCO₃. The magnification factors (e.g. x1) of the base peak because of the signal suppression of co-eluting non-glycopeptides (v, vii, xi, xviii-xix, xxiv, and xxv) (Fig. 1a). Upon the addition of 20% acetic acid to the sample (pH 1.1), non-glycopeptides (v, vii, xi, xviii-xix, xxiv, xxv, and xxvi) still co-eluted with the glycopeptides (Fig. 1b) and dominated the total ion mass spectrum from NPLC-ESI-MS of the glycopeptide fraction.

In contrast, when 0.25% HCl was added to the sample (pH 1.1), the non-glycopeptides (i-xxvii) were almost entirely removed, and the spectrum of the glycopeptide fraction was dominated by glycopeptide ions from the RNase B and fetuin digest (Fig. 1c and Table II). 0.25% HCl was insufficient for removal of non-glycopeptides from the glycopeptide fraction if the sample also contained 50 mM ammonium bicarbonate, a common reagent used in tryptic digestion. For example, relatively intense ion signals were observed for non-glycopeptides (v, vii, xviii-xxix) in the MS spectrum of the glycopeptide fraction when 50 mM ammonium bicarbonate was added to the sample (supplemental Fig. S1a). However, increasing the HCl concentration to 1% restored the separation of glycopeptides and non-glycopeptides (Fig. 1d).

When 0.25% trifluoroacetic acid (TFA), a common ion-pairing reagent used to increase peptide retention in RPLC, was added to the sample, a similar degree of separation between glycopeptides and non-glycopeptides was obtained when compared with 0.25% HCl (Fig. 1d). Again, when the tryptic digest contained 50 mM NH₄HCO₃, the TFA concentration needed to be increased to 1% to eliminate the interference from non-glycopeptides (supplemental Fig. S1b). A total of 23 glycopeptides with five high mannose type glycopeptides from RNase B (a-e, Asn-60; Table II), 13 sialylated N-linked glycopeptides (f–q, Asn-38, 81, and 158), and six sialylated O-linked glycopeptides from fetuin (r–w, T228–228, 288) were detected. The IP-NPLC-MS of the RNase B and fetuin tryptic digest were reproducible (relative standard deviation (RSD) < 10% on base peak ion intensities; RSD ≤ 0.2% on the retention times of base peak ions (n=3) (Fig. 1c–e and supplemental Table S1)) with recovery of high mannose type glycopeptides close to 100% (data not shown).

**Effect of Ion-pairing on Peptide Retention in IP-NPLC—The effect of different acids on retention time shift of a non-glycopeptide (e.g. xxiv) and that of a glycopeptide (e.g. a) is demonstrated (Fig. 2).** When there were no acids added to the sample, broad chromatographic peaks and peak tailing were

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observed for the non-glycopeptide (xxiv) at 12.7 min (Fig. 2, red peak a) whereas the glycopeptide (a) was not detected. When the sample was injected with 20% CH$_3$COOH (pH 1.1), the peptide peak shape was dramatically improved (Fig. 2, orange peak b) but resulted in co-elution of glycopeptide (a) and non-glycopeptide (xxiv) at 11.9 min (orange peaks B and b). In contrast, when the samples were injected with HCl or TFA with/without 50 mM NH$_4$HCO$_3$, retention time differences of 2.2 min – 5.2 min between the non-glycopeptide (xxiv) and glycopeptide (a) were observed, therefore allowing removal of non-glycopeptide (xxiv) from the glycopeptide fraction. Similar retention time shifts were found for other peptides (Table S1).

**IP-NPLC for Isolating Tryptic Glycopeptides on Different Hydrophilic Media**—IP-NPLC experiments were carried out on a ZIC-HILIC and an amino column using the same experimental conditions as for the polyhydroxyethyl A column except that a flow rate of 1.1 $\mu$L/min was used for the 0.3 mm i.D. amino column. The three stationary phases are different in that the polyhydroxyethyl A is almost neutral, the ZIC-HILIC medium is zwitterionic, and the amino column is positively charged. The addition of TFA to the sample improved isolation selectivity of all three columns. However, isolation selectivity decreased considerably as the surface charges increased (Fig. 1e and Fig. 3). The neutral polyhydroxyethyl A column was superior to the other two columns which also yielded much wider chromatographic peaks (data not shown). The results indicate that ionic interactions between peptides and the stationary phases are detrimental to glycopeptide selectivity. The polyhydroxyethyl A medium retains solutes almost solely through hydrophilic interactions, and it has only a slight positive charge on its surface under p$H$ 2. Therefore, interferences arising from ion-exchange processes in IP-NPLC using this column were absent.

**Prediction of the Elution Order of Non-glycopeptides in IP-NPLC Using Calculated Hydrophobicity Values**—We plotted the calculated hydrophobicity values of the 29 tryptic non-glycopeptides of RNase B and fetuin (Table I) against their IP-NPLC elution times (Fig. 4) with the addition of 0.25% HCl to the sample. The linearity of the plot ($r^2 = 0.84$) and the reproducibility of retention times of IP-NPLC suggest that co-elution of a non-glycopeptide with glycopeptides occurring in IP-NPLC can be estimated using their calculated hydrophobicity values. For example, the most hydrophilic non-glycopeptide (xxix), QHMDSSTSAASSSNYNQMMK, of RNase B has a hydrophobicity value of $-18.6$ kcal/mol and eluted at 7.7 min, later than all the other tryptic non-glycopeptides in the digest (i-xxviii) (Table I and supplemental Table S1). In contrast, the most hydrophobic glycopeptide (a) from RNase B, NLTG-GlcNAc$_3$Man$_9$, eluted at 8.9 min (Fig. 2, peak...
Therefore, non-glycopeptides with hydrophobicity values higher (more positive) than $\Delta G_{\text{actw}}$ = 18.6 kcal/mol are likely to elute earlier than glycopeptides modified by high mannose type heptasaccharides in IP-NPLC. The occurrence of tryptic peptides of common proteins having hydrophobicity values lower than $\Delta G_{\text{actw}}$ = 18.6 kcal/mol is low (data not shown) suggesting that IP-NPLC is capable of isolating glycopeptides cleanly from tryptic digests of complex protein mixtures.

**IP-NPLC for Isolating Sialylated O-linked Glycopeptides of Fetuin—** The O-linked glycopeptide ions from a fetuin tryptic digest (T228–288) were observed by direct IP-NPLC-MS analysis (glycopeptides r–w; Fig. 1e). However, the
MS signals of these O-linked glycopeptide ions were much lower compared with those of the N-linked glycopeptide ions, largely because of the low glycosylation occupancy (13). When the fetuin tryptic digest was treated with PNGase F to remove the N-linked glycans, the prominent ions detected in the IP-NPLC glycopeptide fraction were all from O-linked glycopeptides of fetuin (r–w) (Fig. 5). The ion at \( m/z \) 1112.3 (3+) is an unidentified glycopeptide (Fig. 5, inset).

Isolation of Glycopeptides from Complex Peptide Mixtures Using IP-NPLC for Quantitative Glycoprotein Profiling Using Label-free MS—Glycopeptides of a complex tryptic digest of total periplasmic protein extracts of the C. jejuni 11168 wt and the pgID mutant were isolated by IP-NPLC and subjected to nanoRPLC-MS analysis. The base peak chromatograms (BPCs) of the glycopeptide and non-glycopeptide IP-NPLC fractions and the total digest are presented in Fig. 6. The BPCs of the non-glycopeptide fraction and total digest (Fig. 6, b and c) were very similar whereas that of the glycopeptide fraction (Fig. 6a) was far less complex because of removal of non-glycopeptides by IP-NPLC. MS/MS experiments determined that the most prominent ions (\( m/z \) 800–1800) from the BPC of the glycopeptide fraction were all from glycopeptides (Table III) whereas non-glycopeptide ions dominated the BPCs of the non-glycopeptide fraction and total digest.

To demonstrate that IP-NPLC can isolate glycopeptides reproducibly for quantitative analysis, the wt and pgID periplasmic extracts were fractionated in triplicate (40 \( \mu \)g each), and the glycopeptide fractions were subjected to label-free nanoRPLC-MS analyses (4 \( \mu \)g each). The peak areas of extracted ion chromatograms of the identified glycopeptides...
listed in Table III were manually quantified for differential expression analysis between the wt and mutant strains. The relative standard deviation (n = 3) of the glycopeptide ions identified by this IP-NPLC label-free MS approach was between 3–45% irrespective of the peptide abundances and retention times, indicating that IP-NPLC was capable of isolating trace level glycopeptides reproducibly (Tables III and IV) (Fig. 6).

Multiple Stage Tandem MS of Differentially Expressed Tryptic Glycopeptides for Identification of Glycoproteins—The glycopeptide ions in the glycopeptide fractions of the wt and pgID mutant extracts were subjected to MS² and MS³ analysis to determine their oligosaccharide composition and their amino acid sequences. The N-linked glycan of C. jejuni has been previously determined to be GalNAc-α1,4-[GalNAc-α1,4-[Glcβ1,3]-GalNAc-α1,4-GalNAc-α1,4-GalNAc-α1,3-Bac-β1 (BacGalNAc®Glc residue mass: 1406 Da), where Bac is 2,4-diacetamido-2,4,6-trideoxyglucopyranose (30). In addition, in vitro biosynthesis experiments have identified PglD as the acetyltransferase that modifies the uridine diphosphate (UDP)-2-acetamido-4-amino-2,4,6-trideoxyglucopyranose at the C-4 position to form UDP-2,4-diacetamido-2,4,6-trideoxyglucopyranose using acetyl-coenzyme A as the acetyl group donor (39).

As an example, the MS² spectra of equivalent glycopeptides from the wt and mutant strains are presented in Fig. 7, a and b, respectively. Both MS/MS spectra are dominated by ions associated with loss of sugar residues from the glycan modification. As expected, the oligosaccharide composition of the wt glycopeptide was determined to be BacGalNAc®Glc (Fig. 7a). The MS² spectrum of the pgID glycopeptide ion (Fig. 7b) is very similar to that of the wt suggesting that oligosaccharide composition and order are alike in the two strains. However, the reducing end sugar is smaller by 42 Da in the mutant strain and has been annotated as deAcBac (2-acetamido-4-amino-2,4,6-trideoxyglucopyranose). The glycan composition and structure of this glycopeptide ion are therefore assigned as GalNAc-α1,4-GalNAc-α1,4-[Glcβ1,3]-GalNAc-α1,4-GalNAc-α1,4-GalNAc-α1,3-deAcBac-β1 (residue mass: 1364 Da).

MS³ analysis was performed on the doubly protonated MS² fragment ion at m/z 851.45, which corresponded to the unmodified peptide (Fig. 7c). Database searching matches the MS³ fragment ions to TDQNLTVAPPEFQK from the C. jejuni periplasmic protein, Cj1670c (Table III) with an ion score of 57 as defined by the MASCOT™ algorithm. A total of 9 glycopeptides corresponding to eight periplasmic glycoproteins were identified by this approach (Table III). N-glycosylation was confirmed for Cj0200c, Cj0152c, Cj1670c, and Cj0114 (30, 32), and four novel glycopeptides from Cj0168c, Cj0235c, Cj0864, and Cj0783 were identified. The MS² and MS³ spectra for these other glycopeptides are presented in supplemental Fig. S2.
Ion-pairing Normal-phase LC for Isolating Glycopeptides

**TABLE IV**

Quantitative analysis of glycoproteins from the periplasmic protein extracts of *C. jejuni* 11168 (wt) and the isogenic pgID mutant

| Gene name | Protein name | Strain | Count | Intensities in wt | % error | Intensities in mutant | % error | Ratio wt/mutant | % error |
|-----------|--------------|--------|-------|------------------|---------|----------------------|---------|----------------|---------|
| Cj0168c   | (YP_002343627) putative periplasmic protein | wt 1 | 16166 ± 503 | 3 | 1310 ± 157 | 12.0 | 12.34 ± 1.53 | 12.39 |
| pgID      | 92 ± 18     | 19 | 1496 ± 250 | 16.7 | 0.06 ± 0.02 | 25.60 |
| Cj0235c   | (NP_281430) protein export membrane protein | wt 1 | 472 ± 26 | 6 | 27 ± 3 | 11.1 | 17.49 ± 2.17 | 12.41 |
| pgID      | 70 ± 5     | 7 | 757 ± 126 | 16.6 | 0.09 ± 0.02 | 17.86 |
| Cj0200c   | (YP_002343658) putative periplasmic protein | wt 1 | 204 ± 87 | 43 | 416 ± 40 | 11.4 | 5.02 ± 2.22 | 44.33 |
| pgID      | 204 ± 12   | 6 | 2033 ± 133 | 6.5 | 0.10 ± 0.01 | 8.78 |
| Cj0152c   | (CAL34323) hypothetical protein | wt 1 | 424 ± 103 | 24 | nd | only in wt | — |
| pgID      | 242 ± 81   | 33.3 | only in mutant | — |
| Cj1670c   | (YP_002345038) putative membrane protein or CgpAa | wt 1 | 224 ± 65 | 29 | 43 ± 4 | 10.1 | 5.22 ± 1.59 | 30.51 |
| pgID      | 242 ± 81   | 33.3 | only in mutant | — |
| Cj0864    | (YP_002343574) putative periplasmic protein | wt 1 | 2186 ± 914 | 42 | 186 ± 9 | 5.1 | 11.71 ± 4.93 | 42.09 |
| pgID      | 41 ± 5     | 11 | nd | only in wt | — |
| Cj0114    | (YP_002344271) putative periplasmic protein | wt 1 | 121 ± 20 | 16 | 1393 ± 137 | 9.8 | 0.09 ± 0.02 | 18.91 |
| pgID      | 115 ± 9    | 25 | 84 ± 16 | 19.3 | 1.37 ± 0.43 | 31.37 |
| Cj0783    | (YP_002344190) Periplasmic nitrate reductase cytochrome c-type subunit | wt 1 | 527 ± 9 | 16 | nd | only in wt | — |
| pgID      | nd | nd | nd | only in both | — |

a According to Linton et al. (41).

The putative periplasmic protein, Cj0114 is identified and quantified by two glycopeptides whose detailed information can be found in Table III.

 Recovery and Selectivity of IP-NPLC for Isolating Tryptic Glycopeptides from Complex Peptide Mixtures—The nano-RPLC-MS extracted ion chromatograms for three glycopeptide ions from the pgID total digest as well as the corresponding glycopeptide and non-glycopeptide IP-NPLC fractions are presented in Fig. 8. The glycopeptide ions were detected in the total (Fig. 8, c, f, and h) and glycopeptide fractions (Fig. 8, a, d, and g) but were absent in the non-glycopeptide fraction (Fig. 8, b, e, and h). The glycopeptide ion intensities were comparable in the total and glycopeptide chromatograms. Similar results were obtained for all of the glycopeptides listed in Table III, regardless of their abundance, indicating that the recovery of these glycopeptides by IP-NPLC was close to 100%. Furthermore, only a few precursor ions from the non-glycopeptide fraction generated a fragment ion at m/z 204.09 upon CID, the majority of which were identified as non-glycopeptides (>85%), indicating that virtually no glycopeptides were being lost in the non-glycopeptide fraction (Fig. 6b, Fig. 9, and supplemental Table S2).

The glycopeptide selectivity per nanoRPLC-MS analysis (scan range: m/z 800–1800) of the glycopeptide fraction (defined as the number of base peak glycopeptides determined by the N-consensus sequence or the signature oxonium ions of glycopeptides upon CID divided by the number of the prominent peptides identified from the glycopeptide fraction) in our approach was 90% (Fig. 6 and Table III). Non-glycopeptides rich in hydrophilic amino residues were present in the glycopeptide fraction, e.g. ALDNDNQEDQTESR from flavodoxin (Cj1382c) eluting at 16.8 min (Fig. 6a). However, the occurrence of these peptides in a complex tryptic digest appeared low thus emphasizing the efficiency of our method.

**DISCUSSION**

In this report we describe a new IP-NPLC method for quantitatively isolating glycopeptides from complex tryptic digests using acids as ion-pairing reagents and polyhydroxyethyl A stationary phase. The addition of no acids or acetic acid to the sample solution can diminish the effect of HCl or TFA on the recovery of these glycopeptides (Figs. 1 and 2). In contrast, the use of HCl or TFA resulted in separations of glycopeptides from non-glycopeptides (Figs. 1 and 2, supplemental Fig. S1b, and supplemental Table S1). It is apparent that the dramatic improvement in isolation of glycopeptides using IP-NPLC over NPLC was not simply because of the pH change in the sample because only hydrochloric acid but not acetic acid led to isolation of glycopeptides under the same sample pH (Fig. 1, b and c). Other ions present in the sample solution can diminish the effect of HCl or TFA on glycopeptide isolation selectivity, e.g. NH₄HCO₃ (supplemental Fig. S1a), but this can be overcome by increasing the concentration of the two acids (Fig. 1d and supplemental Fig. S1b). The results suggest that acidic anions present in the samples significantly affect the selectivity of NPLC for the isolation of glycopeptides.

A total of 18 different sialylated glycopeptides from fetuin, including 6 O-linked and 12 N-linked glycopeptides, and 5
high mannose type N-linked glycopeptides from RNase B were isolated by IP-NPLC from a tryptic digest of a mixture of these two glycoproteins (Fig. 1e and Table II) (12, 13, 23). A typical N-linked carbohydrate derived from a eukaryotic host usually contains 7–25 monosaccharide units (O-linked carbohydrates have a similar size range but with less branching and may be as short as one monosaccharide unit) (4). Thus, NLTK-GlcNAc2Man5–9 glycopeptides from RNase B are modified with shorter high mannose type glycans and resemble non-glycopeptides in terms of hydrophilicity. Therefore, these glycopeptides have proven to be difficult to resolve from non-glycopeptides by other chromatographic methods (23, 25–26). In contrast, our IP-NPLC method was effective at resolving these less hydrophilic glycopeptides and the sialylated fetuin glycopeptides from the digest of RNase B and fetuin mixture (Fig. 1, Table I, and supplemental Table S1). The method described here is more effective than other chromatographic methods in that it works well with a wide range of glycopeptide species (23, 25, 29).

The retention time shifts of non-glycopeptides (Fig. 2 and supplemental Table S1) can be explained by ion-pairing processes that occur between the acidic anions and the counter ions of non-glycopeptides in solution (29, 43–45). The earlier elution of non-glycopeptides upon the addition of HCl or TFA to the sample rather than CH₃COOH can be attributed to formation of ion pairs –NH₃⁺CH₃COO⁻, –NH₃⁺Cl⁻, and –NH₃⁺CF₃COO⁻, which are progressively more hydrophobic (–NH₃⁺ refers to the ionized amine groups in non-glycopeptides). The higher hydrophilicity of CH₃COO⁻ or –NH₃⁺CH₃COO⁻ in IP-NPLC is likely because of the hydro-
gen bonding ability of oxygen with the stationary phase whereas chlorine cannot form strong hydrogen bonding. Covalently bound fluorine is not capable of hydrogen bonding, and it apparently dramatically weakened the hydrogen bonding ability of oxygen on CF$_3$COO$^-$ (46). As a result, NH$_3^+$CF$_3$COO$^-$ is more hydrophobic than NH$_3^+$CH$_3$COO$^-$ and NH$_3^+$Cl$^-$. The acetate anion formed relatively hydrophilic ion pairs with the non-glycopeptides allowing them to interact with the polyhydroxyethyl A medium and eluted with the glycopeptides (Fig. 2, orange peak B and b). In contrast, the ion pairs between the charged non-glycopeptides and the Cl$^-$ or CF$_3$COO$^-$ were more hydrophobic and eluted earlier from the column (Fig. 2, peak b versus peaks d–f). In addition, improvement in chromatographic peak shapes was obtained using the three acids as ion-pairing reagents over NPLC alone (Fig. 2, red peak a versus peaks b–f) (43).

The retention times of glycopeptides were effected to a much lesser extent by ion pairing because of the strong non-ionic electrostatic interactions between their glycans and the hydrophilic stationary phase (Fig. 2, peak C versus c, peak D versus d, peak E versus e, peak F versus f). As a result, non-glycopeptides eluted much earlier than glycopeptides in IP-NPLC. Our data indicate that formation of ion-pair com-

Fig. 8. Recovery and selectivity of IP-NPLC for isolating tryptic glycopeptides from complex peptide mixtures of the C. jejuni pglD mutant. Extracted ion chromatograms (XIC) of nanoRPLC-MS at m/z 1281.40 (3+) of a glycopeptide from the putative periplasmic protein, Cj0114, in the (a) Glycopeptide fraction; (b) Non-glycopeptide fraction; (c) Total digest. XICs of nanoRPLC-MS at m/z 1021.20 (3+) of a glycopeptide from protein export membrane protein, Cj0235c, in the (d) Glycopeptide fraction; (e) Non-glycopeptide fraction; (f) Total digest. XICs of nanoRPLC-MS at m/z 926.14 (3+) of a glycopeptide from the putative periplasmic protein, Cj0168c, in the (g) Glycopeptide fraction; (h) Non-glycopeptide fraction and (i) Total digest. The amino acid sequences of these glycopeptides are provided in Table III. The insets in (d), (e), and (f) show the combined mass spectra at m/z 1020.20 (3+) corresponding to the glycopeptide fraction, non-glycopeptide fraction, and total digest (23–24 min) for comparison. PA, peak area of the extracted ions; ND, not detected; Fr., fraction.
Fig. 9. NanoRPLC-MS² of the non-glycopeptide fraction of the pglD mutant using data dependent analysis: (a) Base peak intensity (BPI) chromatogram and (b) Extracted ion chromatogram (XIC) at m/z 204.09 of the non-glycopeptide fraction. The ions labeled in (b) were all identified as non-glycopeptides, except the unidentified ion at m/z 897.42 (2+), and their sequences are provided in Table S2. The magnification factors in (a) e.g. x80 for the base peak intensity (BPI) of the ion at m/z 851.51(2+) of the nanoRPLC-MS² of the non-glycopeptide fraction were relative to the most abundant ion for comparison.

The presence of predominantly monoacetylated Bac residues on glycoproteins isolated from the C. jejuni pglD mutant confirms that PglD is an acetyltransferase and that the oligosaccharyltransferase, PglB, is still capable of transferring \(^{\text{deAc}}\text{BacGalNAc}_5\text{Glc}\). However, the quantitative information on the differentially expressed N-linked glycoproteins further demonstrates the importance of Bac acetylation at C-2 for optimal PglB activity in C. jejuni (40). The quantitative proteomic data demonstrate for the first time that glycopeptides modified by \(^{\text{deAc}}\text{BacGalNAc}_5\text{Glc}\) are also present in the wt strain, albeit at low abundance, indicating that acetylation of Bac is incomplete even in the presence of an active PglD enzyme. In addition, it is apparent that there is another

plexes between acidic anions and charged non-glycopeptides can increase the retention time differences between non-glycopeptides and glycopeptides, thereby leading to fractionation of a complex tryptic digest into a glycopeptide and a non-glycopeptide fraction (Figs. 1 and 6) (supplemental Table S1).

Ion pairs \(-\text{COO}^-\text{H}^+\) between the hydronium ion (H\(_3\)O\(^+\)) and the negatively charged carboxyl group of non-glycopeptides (\(-\text{COO}^-\)) are likely to form in solution (47). In addition, Wimley et al. have determined that the neutral \(-\text{COOH}\) terminus of peptides determined by pH, are considerably more hydrophobic than \(-\text{COO}^-\) (35–37). Therefore, ion-pairing processes between hydronium ions and peptides likely contribute to the decreased retention described in Fig. 2. However, the decreased retention because of formation of hydronium ion pairs with peptides will be generally less significant compared with anions because hydrogen of an ion-pair is capable of hydrogen bonding with stationary phase whereas chlorine or covalently bound fluorine is not (46).

We demonstrate for the first time that the surface charges of the stationary phase decrease glycopeptide isolation selectivity using IP-NPLC. Furthermore, the elution order of non-glycopeptides in IP-NPLC can be predicted reasonably well using their hydrophobicity values (Fig. 4). Non-glycopeptides in IP-NPLC can be predicted reasonably well using their hydrophobicity values (Fig. 4). Non-glycopeptides in IP-NPLC can be predicted reasonably well using their hydrophobicity values (Fig. 4). Non-glycopeptides in IP-NPLC can be predicted reasonably well using their hydrophobicity values (Fig. 4). Non-glycopeptides in IP-NPLC can be predicted reasonably well using their hydrophobicity values (Fig. 4). Non-glycopeptides in IP-NPLC can be predicted reasonably well using their hydrophobicity values (Fig. 4). Non-glycopeptides in IP-NPLC can be predicted reasonably well using their hydrophobicity values (Fig. 4). Non-glycopeptides in IP-NPLC can be predicted reasonably well using their hydrophobicity values (Fig. 4). Non-glycopeptides in IP-NPLC can be predicted reasonably well using their hydrophobicity values (Fig. 4).

We applied IP-NPLC to reproducibly and quantitatively isolate glycopeptides from tryptic digests of periplasmic proteins extracted from C. jejuni 11168 wt and pgld mutant for subsequent label-free nanoRPLC-MS analysis of glycoprotein expression (Tables III and IV). We chose to re-examine the C. jejuni pgld mutant with our new technique because in our previous study (48) we detected minor amounts of protein glycosylation in this mutant but because of sensitivity issues were unable to characterize these modifications any further. Four of the nine glycopeptides identified in this study have not been observed before. For example, the most abundant glycopeptide isolated from the wt strain is derived from the putative periplasmic protein, Cjo168c. This glycoprotein has a mass of only 5,813 Da (amino acid mass only) and therefore would not have been detected by the gel-based methods used in previous studies to resolve the glycoprotein mixtures prior to MS analysis (30). Remarkably, the reproducibility of this label-free MS approach using IP-NPLC is comparable with that obtained using hydrazide chemistry and stable isotope labeling (15). In addition, IP-NPLC does not involve lengthy chemical procedures as for hydrazide chemistry and also preserves the glycan information.

The presence of predominantly monoacetylated Bac residues on glycoproteins isolated from the C. jejuni pgld mutant confirms that PglD is an acetyltransferase and that the oligosaccharyltransferase, PglB, is still capable of transferring \(^{\text{deAc}}\text{BacGalNAc}_5\text{Glc}\). However, the quantitative information on the differentially expressed N-linked glycoproteins further demonstrates the importance of Bac acetylation at C-2 for optimal PglB activity in C. jejuni (40). The quantitative proteomic data demonstrate for the first time that glycopeptides modified by \(^{\text{deAc}}\text{BacGalNAc}_5\text{Glc}\) are also present in the wt strain, albeit at low abundance, indicating that acetylation of Bac is incomplete even in the presence of an active PglD enzyme. In addition, it is apparent that there is another
acetyltransferase complementing the PgID activity as we previously predicted (48) because glycopeptides modified by BacAilNAC,Glc are also present in the mutant although at much lower abundance compared with the wt (Tables III and IV).

The glycopeptide selectivity of our approach reaches 90% based on the total number of glycopeptides and peptides identified in the C. jejuni glycopeptide fractions (Fig. 6 and Table III). Based on the retention time difference observed between the most hydrophilic non-glycopeptide (xxix) and the glycopeptides, NLTG-GlcNAC,Man 

Based on the absence of glycopeptide ion signals in the non-glycopeptide fraction (Figs. 8 and 9). This glycopeptide selectivity and recovery is comparable with that obtained with hydrazide chemistry capturing tryptic glycopeptides (16). We believe that this IP-NPLC method can be used for isolating glycopeptides at much lower levels because low abundance glycopeptides were effectively recovered from the tryptic digests of periplasmic protein extracts from C. jejuni (Fig. 6, m/z 1218.40(3+) at 25.6 min, Cj(0114).

In comparison to other chromatographic approaches for enriching glycopeptides, this new IP-NPLC method offers the best glycopeptide selectivity and recovery for complex peptide mixtures to our knowledge (19, 22, 23, 25, 28–29). Nevertheless, there is still room for improvement as some non-glycopeptides were present in the glycopeptide fraction. The non-glycopeptides abundances, the magnitude of ion-pairing equilibrium constants under a given solvent composition (49), and the hydrophobicity of non-glycopeptides are likely the important factors concerning contamination. Further optimization of experimental conditions will improve glycopeptide selectivity and may allow the isolation of glycopeptides modified by glycans as short as trisaccharides. We believe that IP-NPLC will make a significant contribution to glycoproteomics research in the future.

Acknowledgments—We thank Simon Foote and Dr. Arsalan Haqani for assistance in database searches for identification of C. jejuni glycoproteins.

* This work was supported by the National Research Council, Canada, Genomics and Health Initiative.

The on-line version of this article (available at http://www.mcp.org) contains supplemental material.

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