Coupling hydrophilic interaction chromatography materials with immobilized Fe$^{3+}$ for phosphopeptide and glycopeptide enrichment and separation†

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Simultaneous profiling of protein phosphorylation and glycosylation is very important to elucidate the bio-functions of these proteins. However, simultaneous enrichment of glyco- and phosphopeptides is the bottleneck in proteomics because of the low abundance of these species and ion suppression from non-modified peptides in mass spectrometry (MS). In this study, Fe$^{3+}$ immobilized hydrophilic interaction chromatography (HILIC) materials (termed polySD-SiO$_2$, recently reported in our lab) and polySD-SiO$_2$ in the HILIC mode were employed for the simultaneous enrichment and subsequent separation of glyco- and phosphopeptides. The Fe$^{3+}$ immobilized polySD-SiO$_2$ could selectively enrich glycopeptides and phosphopeptides and the co-enriched peptides were further fractionated with polySD-SiO$_2$ in the HILIC mode. With the established method, glyco- and phosphopeptides were well enriched and divided into two fractions even from tryptic digests of a-casein, fetuin and BSA at a molar ratio of 1 : 2 : 400. Application of the established method to HeLa cell lysate resulted in a total of 1903 phosphopeptides and 141 glycosylation sites. These results demonstrate that the established method could selectively and simultaneously enrich and fractionate glyco- and phosphopeptides from complex peptide mixtures.

1. Introduction

Protein phosphorylation and glycosylation are two types of most significant post-translational modification (PTM), which take part in the regulation of many vital biological processes.1 Reversible protein phosphorylation modulates cellular processes such as cell cycle and signaling.2 Protein glycosylation is related to tumor invasion and metastasis, cell adhesion, and immune response.3 The terminals of glycans in glycoproteins are often modified with sialic acids (SAs).4,5 Increasing evidence shows the increased expression of sialylated glycoproteins on the surface of tumor cells.6 Changes in the level of sialylation interplay with alternation in phosphorylation.7–9 Therefore, to further elucidate these biological processes of protein phosphorylation and glycosylation at the molecular level, it is very essential to simultaneously profile these two kinds of PTMs in biological and clinical samples.

Mass spectrometry (MS) is the method of choice for profiling protein phosphorylation and glycosylation. However, enrichment of PTM-peptides is always required prior to MS analysis owing to the low abundance of PTM-peptides in complex samples and ion suppression effect resulting from non-modified peptides in MS. Therefore, continuous efforts have been endeavored to enrich individual subset of PTM-peptides in order to reduce the complexity of samples. The commonly used methods toward phosphopeptide enrichment include immobilized metal ion affinity chromatography (IMAC),8 metal oxide affinity chromatography (representative with TiO$_2$ and ZrO$_2$),10,11 and hydrophilic interaction chromatography (HILIC).12 Meanwhile, lectin affinity chromatography,13–14 TiO$_2$ (ref. 15–17) and HILIC18,19 have been used for glycopeptide enrichment. During the last decade, HILIC materials have made tremendous progress and widely applied to enrich glycopeptides in the form of Carbon-based materials,20 polymers21 and ZIC-HILIC.22,23

In recent years, several materials have been developed and applied to simultaneous enrichment of phosphopeptides and glycopeptides. These materials include metal oxide affinity chromatography,24,25 IMAC,26,27 and HILIC.26,28,29 Even though these materials could be used to...
simultaneously capture both phosphopeptides and glycopeptides, these two PTM-peptides are co-eluted from materials, which are still very complex to be analysed with MS. To reduce the complexity of the samples, the second dimensional separation has been employed with HILIC columns or reverse phase liquid chromatography. For example, phosphopeptides and glycopeptides were enriched with TiO$_2$; after the treatment of PNGase F overnight, phosphopeptides and the deglycosylated peptides were subsequently fractionated with HILIC.$^{30,31}$ However, phosphopeptides and the deglycosylated peptides could be found in each HILIC fraction. The sever overlap of phosphopeptides and deglycosylated peptides in HILIC originates from the insufficient hydrophilicity differences between PTMs peptides. Moreover, removal of glycans is carried out under basic condition overnight, which leads to hydrolyze of co-existing phosphopeptides. Thus, it will be ideal that phosphopeptides are separated from intact glycopeptides before the deglycosylation step.

In our previous study, we synthesized bioinspired polymers Ser-Asp (SD) modified silica (denoted as polySD-SiO$_2$) to enrich glycopeptides.$^{32}$ In this article, we incorporated iron ions onto polySD-SiO$_2$ materials and used Fe$^{3+}$ immobilized polySD-SiO$_2$ to simultaneously enrich phosphopeptides and glycopeptides. Then the co-enriched PTM-peptides were further separated into phosphopeptides and intact glycopeptides fractions with polySD-SiO$_2$ under HILIC mode (Scheme 1). This method was validated with the tryptic digests of the phosphoprotein, glycoprotein and non-modified protein and even HeLa S3 cell lysate.

2. Experimental

2.1 Materials and reagents

Bovine fetuin, a-casein and bovine serum albumin (BSA), ammonium bicarbonate ($\text{NH}_4\text{HCO}_3$), glycolic acid, guanidine hydrochloride, dithiothreitol (DTT), iodoacetamide (IAA), and peptide N-Glycosidase F (PNGase F) were obtained from Sigma-Aldrich (St. Louis, MO, USA). Sequencing grade trypsin was purchased from Promega (Madison, WI, USA). Trifluoroacetic acid (TFA) was obtained from TEDIA (Fairfield, USA). Acetonitrile (ACN) was purchased from Merck (Darmstadt, Germany). Acetic acid and formic acid (FA) was obtained from Acros Organics (Geel, Belgium). Ammonium hydroxide was purchased from Fluka (Buchs, Switzerland). GE Loader tips were purchased from Eppendorf (Madison, WI). Water was prepared with a Milli-Q system (Millipore, Bedford, MA, USA).

2.2 Synthesis and characterization of Fe$^{3+}$ immobilized polySD-SiO$_2$

The previously prepared polySD-SiO$_2$ (500 mg) were suspended in water and washed twice with 1 mL 50 mM EDTA in 1 M NaCl. After centrifugation, the supernatant was removed and the remaining materials were incubated with 1 mL 0.1 M FeCl$_3$ in 0.1 M acetic acid. After incubation of 1 h, the suspension was centrifuged and the paillet was subsequently washed with 1 mL 1 M NaCl and 0.1 M acetic acid. After washing with water, the materials were dried under 60 °C. X-ray photoelectron spectroscopy (XPS) was obtained with a VG Multilab 2000. Scanning electron microscopy (SEM) spectra were recorded on a Hitachi S-4800 SEM.

2.3 Trypsin digestion of proteins

Each standard protein (1 mg) was dissolved in 100 $\mu$L of 6 M guanidine hydrochloride/50 mM ammonium bicarbonate. The denatured protein solution was reduced with 5 $\mu$L of 200 mM DTT/50 mM ammonium bicarbonate for 45 min at 56 °C. Free sulphydryl groups were alkylated with 20 $\mu$L of 200 mM IAA/25 mM ammonium bicarbonate and followed by incubation for 30 min at room temperature in the dark. The solution was diluted to 1 mL with 25 mM ammonium bicarbonate, mixed with trypsin at an enzyme/protein ratio of 1 : 20 (w/w) and incubated overnight at 37 °C. The digestion was stopped by adding 10% formic acid at the final concentration of 1%.

2.4 Enrichment of phosphopeptides and glycopeptides with Fe$^{3+}$ immobilized polySD-SiO$_2$

Fe$^{3+}$ immobilized polySD-SiO$_2$ (2.0 mg) was slurred in 40 $\mu$L of ACN and pushed into a GE Loader tip. The micro-column was equilibrated with 40 $\mu$L of loading buffer (80% CAN/5% TFA). The tryptic peptide mixture was dried, redissolved in 40 $\mu$L loading buffer and loaded onto the Fe$^{3+}$ immobilized polySD-SiO$_2$ packed micro-column. The absorbed peptides were washed twice with 40 $\mu$L of 80% ACN/5% TFA, and eluted with 40 $\mu$L of 1.0% ammonium hydroxide. The elution fraction was immediately acidified with 1% FA.

2.5 Separation of phosphopeptides and glycopeptides with PolySD-SiO$_2$

PolySD-SiO$_2$ packed micro-column was used to separate phosphopeptides from glycopeptides. Elution fraction from Fe$^{3+}$ immobilized polySD-SiO$_2$ was dried and redissolved in 40 $\mu$L of 80% ACN/5% TFA. For micro-column packing, 40 $\mu$L of ACN slurry/the PolySD-SiO$_2$ (2.0 mg) was packed into the GE Loader.

![Scheme 1](image-url) The process of simultaneous enrichment and subsequent separation of phosphopeptides and glycopeptides with Fe$^{3+}$ immobilized polySD-SiO$_2$ and polySD-SiO$_2$. 

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tip. After washing with 40 μL of 50% ACN/0.1% FA, the micro-column was equilibrated with 40 μL of 80% ACN/5% TFA. Then, the peptides were loaded onto the column and the flow-through was collected. The micro-column were rinsed twice with 40 μL of 80% ACN/5% TFA and eluted with 40 μL of 50% ACN/1% TFA. The phosphopeptides were obtained by pooling the flow-through and eluant from 80% ACN/5% TFA. The glycopeptides fraction resulting form 50% ACN/1% TFA was dried and redissolved in 40 μL of 50 mM NH₄HCO₃. After addition of 2 unit of PNGase F to glycopeptides solution, the solution was incubated overnight at 37 °C.

HeLa cells culture, protein extraction and protein digestion were carried out by using the reported methods. The phosphopeptides and glycopeptides were enriched with the above-mentioned method.

### 2.6 Enrichment and separation of phosphopeptides and glycopeptides with the reported method

For the simultaneous enrichment of phosphopeptides and glycopeptides, TiO₂ based method was applied as described previously. The eluates from TiO₂ were deglycosylated with 2 unit of PNGase F. After deglycosylation, the peptides were desalted and subsequently subjected to HILIC micro-column packed with TSKGel Amide materials (3 μm particle size, Tosoh Bioscience). The micro-column were eluted with 40 μL of 75% ACN/0.1% TFA and eluted with 40 μL of 50% ACN/0.1% TFA, respectively.

### 2.7 Validating the established method

To investigate the recovery of the established method, stable isotope dimethyl labeling protocol was followed. The standard monophosphopeptide (peptide sequence: HpSPIAPSSPSPK, p: phosphorylation site), diphosphopeptide (peptide sequence: HpSPIApSSPSPK, p: phosphorylation site) and a glycopeptide (peptide sequence: KVANKT, N: glycosylation site) were used in this experiments. Each standard peptide was dissolved in sodium acetate buffer (0.1 M, pH 6) and mixed with formaldehyde (4% in water, 1 L) and formaldehyde-D₂ (4% in water, 1 L). The mixture was vortexed again and after vortex for 5 min, the solution was then mixed immediately with freshly prepared sodium cyanoborohydride (260 mM, 1 μL). The mixture was vortexed again and then allowed to react for 5 min. After the stable isotope dimethyl labeling, the heavily labeled standard peptides were treated with the established. Before the treated samples were injected for MS analysis, the same amount of lightly labeled standard peptides was mixed with the treated samples. The recovery was calculated based on the intensity of lightly and heavily labeled standard peptides.

### 2.8 MS analysis and data searching

The peptide fractions from digests of standard samples were characterized with nano electrospray ionization (ESI)-quadrupole time-of-flight (Q-TOF) tandem mass spectrometer (Waters, Manchester, UK). Modified peptides from HeLa S3 cell lysate were analyzed were analyzed by LTQ Orbitrap Velos (Thermo, San Jose, CA) and searched with Mascot (version 2.3.0, Matrix Science, London, UK). Cysteine carboxamidomethylation was set as fixed modification and oxidation of methionine, deamidation of asparagine and phosphorylation of serine, threonine and tyrosine were set as variable modifications. Up to two missing cleavages of trypsin were allowed. Mass tolerances were 20 ppm and 0.2 Da for the parent and fragment ions, respectively. A false discovery rate of 1% was set at both the peptide and site levels.

### 3. Results and discussion

#### 3.1 Preparation of Fe³⁺ immobilized polySD-SiO₂

The Fe³⁺ immobilized polySD-SiO₂ materials were prepared by incubation of polySD-SiO₂ with FeCl₃. The prepared materials were characterized with scanning electron microscope (SEM) and XPS. SEM images indicated that the materials maintained their integrity after incorporating with Fe³⁺ (Fig. 1a and b). The XPS spectrum implies that Fe³⁺ has been modified on the polySD-SiO₂ (Fig. 1c). These results indicated that Fe³⁺ ions modified polySD-SiO₂ materials have been successfully prepared.

#### 3.2 Enrichment of phosphopeptides and glycopeptides with Fe³⁺ immobilized polySD-SiO₂

To investigate whether Fe³⁺ immobilized polySD-SiO₂ can enrich glycopeptides, tryptic digests of sialoglycoprotein bovine...
fetuin and bovine serum albumin at molar ratio of 1:100 were used as model samples. The number of glycopeptide signals is 30 (detailed information of glycopeptides in Table S1†) with the Fe³⁺ immobilized polySD-SiO₂ materials (Fig. S1†), which is comparable to that reported with polySD-SiO₂.²⁸ To further enrich both phosphopeptides and glycopeptides, tryptic digests of phosphoprotein a-casein and fetuin were used as model samples. At first, the tryptic digests of a-casein and fetuin at molar ratio of 1:3 were used to evaluate the prepared materials. Fig. 2 demonstrates that the mass spectra of peptides before and after treatment with Fe³⁺ immobilized polySD-SiO₂. Before enrichment with our materials, the mass spectrum was occupied with signals of non-modified peptides. The signals of glycopeptides could not be detected and only two phosphopeptides could be observed (Fig. 2a). The poor signals of PTM-peptides were mainly resulted from their low abundance. After enrichment with our materials, the signals of glycopeptides and phosphopeptides dominated the mass spectrum because the abundant non-PTM peptides were thoroughly removed. A total of 7 phosphopeptide signals (detailed information of phosphopeptides in Table S2†) and 20 glycopeptides could be readily found (Fig. 2b). Even 200 molar folds of BSA was mixed as interference with the tryptic digests of a-casein and fetuin at molar ratio of 1:3, our materials maintained its selectivity toward both glycopeptides signals and phosphopeptides. There are still 6 phosphopeptides and 19 glycopeptide signals could be easily found in the mass spectrum (Fig. 3a).

This result implied that Fe³⁺ immobilized polySD-SiO₂ could selectively enrich both glycopeptides and phosphopeptides.

3.3 Separation of phosphopeptides and glycopeptides with polySD-SiO₂
After glycopeptides and phosphopeptides were eluted from Fe³⁺ immobilized polySD-SiO₂, these two species were further separated with PolySD-SiO₂ under HILIC mode. In order to reduce the retention of charged peptides and enlarge the hydrophilicity difference between glycopeptides and non-glycosylated peptides, 5% TFA was added to the mobile phases.²⁹ The tryptic digests of a-casein: fetuin: BSA = 1:2:400 (molar ratio) were treated with Fe³⁺ immobilized polySD-SiO₂. After the treatment our materials, the high-abundance non-modified peptides were efficiently removed and 7 phosphopeptide signals and 16 glycopeptides were readily observed in the mass spectrum of the eluates. After subsequent separation with polySD-SiO₂ under HILIC mode, phosphopeptides and
glycopeptides were efficiently separated into two fractions. Because the further separation reduced the complexity of sample and also decreased the ion suppression effect in MS, nine phosphopeptide signals and 22 glycopeptide signals were easily detected in the mass spectrum of phosphopeptides (Fig. 3b) and glycopeptides fraction (Fig. 3c), respectively. Glycopeptides couldn’t be observed in the phosphopeptide fraction, while very low intensities of one triple phosphopeptide ([m/z = 1339.5343] were found in the glycopeptide fraction, which might result from the strong electrostatic interaction between the triple phosphopeptides and polySD-SiO₂ materials. These results demonstrated that Fe³⁺ immobilized polySD-SiO₂ enrichment coupled with PolySD-SiO₂ fractionation could efficiently separate phosphopeptides and glycopeptides from the peptide pool and further divide these two kinds of PTM-peptides into two fractions.

3.4 Enrichment and separation of phosphopeptides and glycopeptides with the reported method

We then compared the efficiency of previously reported method toward phosphopeptides and glycopeptides.⁴¹ At first, the tryptic digests of a-casein: fetuin: BSA = 1 : 2 : 400 (molar ratio) were treated with TiO₂. The phosphopeptides and glycopeptides could be enriched with TiO₂ and 7 phosphopeptide signals and 12 glycopeptide signals could be detected (Fig. 4a), while some non-PTM peptides could also be detected, such as peptide at [m/z 7874.3626(2⁺)]. Then the enriched peptides from TiO₂ was further deglycosylated with PNGase F to remove glycans. The deglycosylated peptides and phosphopeptides were further separated with TSKGel Amide micro-column under HILIC mode. Only 4 phosphopeptide signals and 4 deglycosylated peptide signals were found in the 75% ACN/0.1% TFA fraction (Fig. 4b), while there were still 2 multiple phosphopeptides and 1 deglycosylated peptide in the 50% ACN/0.1% TFA fraction. It turns out that phosphopeptides and deglycosylated peptides were co-eluted in both two fractions, which is consistent with the previous report.⁴¹ Taking together the afore-mentioned results, these data demonstrated that phosphopeptides and deglycosylated peptides overlapped in these two fractions. It is beneficial to separate phosphopeptides and glycopeptides before deglycosylation with HILIC column.

3.5 The validation of the developed strategy

To evaluate the recovery of the developed strategy, stable isotope dimethyl labeling method were employed (details in 2.7 Validating the established method).⁴⁴ The average recovery was 85.2 ± 3.2% ([n = 3]), 79.2 ± 4.1% and 84.6 ± 2.3% ([n = 3]) for standard mono-phosphopeptide, di-phosphopeptide and glycopeptide, respectively. For comparison, the recovery of the reported method was also performed, which separate phosphopeptides and glycan removed glycopeptides after TiO₂ enrichment. It turned out that the recovery of the reported method was much lower than that obtained with our developed protocol. They were 53.3 ± 1.8% and 62.9 ± 3.4% ([n = 3]) for phosphopeptides and glycopeptides, respectively. To investigate the reproducibility of the established strategy, three parallel experiments were performed to enrich and separate phosphopeptides and glycopeptides from the tryptic digests of a-casein and fetuin at molar ratio of 1 : 2 : 400 after enrichment with TiO₂ (a) and separation of phosphopeptides and deglycosylated peptides with TSKGEL-amide (b and c). (a): Phosphopeptides and glycopeptides enrichment with TiO₂; (b): peptides eluted from TSKGEL-amide with 75% ACN/0.1% TFA; (c): peptides eluted from TSKGEL-amide with 50% ACN/0.1% TFA. The phosphopeptides, glycopeptides and deglycosylated glycopeptides are marked with green asterisks, red circles and blue squares, respectively.

The developed strategy was used for identification of phosphopeptides and glycopeptides from 20 µg HeLa S3 cell lysate. The cell lysate was digested with trypsin and the tryptic peptides were treated with our established method. The phosphopeptides fraction and the deglycosylated glycopeptides fraction were analyzed by Orbitrap mass spectrometer. The selectivity of the established method for phosphopeptides and glycopeptides was 75.3%. More than 1903 phosphopeptides from 542 proteins were characterized in the phosphopeptides fraction (Table S3†) and 139 unique sites from 83 glycoproteins in glycopeptides fraction (Table S4†). No glycopeptide was detected in the
phosphopeptides fraction while 20 multiply phosphorylated peptides were characterized in the glycopeptide fraction. This overlapping ratio between phosphopeptides and glycopeptides achieved with our strategy (<5%) is much lower than that with the literature (43%). The result further demonstrated the advantages of our method for enrichment and fractionation of phosphopeptides and glycopeptides.

4. Conclusions

In summary, we have developed an efficient method for simultaneous enrichment and subsequent separation of phosphopeptides and glycopeptides, taking advantages of the strong affinity between Fe3+ immobilized polySD-SiO2 and PTM-peptides and the hydrophilicity difference between phosphopeptides and glycopeptides on polySD-SiO2. The established method has the advantages of high recovery and selectivity toward the target peptides and low overlapping between phosphopeptides and glycopeptides. We provide a powerful tool to tackle the challenge of simultaneous and selective analysis of phosphopeptides and glycopeptides. This tool may promote the discovery of more PTM-sites from limited amount of biosamples, especially clinical samples.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

We thank the National Natural Science Foundation of China (81472487 and 81773197), the Natural Science Foundation of Hubei Province (2017CFB412).

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