pChem: a modification-centric assessment tool for the performance of chemoproteomic probes

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Chemoproteomics has emerged as a key technology to expand the functional space in complex proteomes for probing fundamental biology and for discovering new small molecule-based therapies. Here we report a modification-centric computational tool termed [pChem](#) to provide a streamlined pipeline for unbiased performance assessment of chemoproteomic probes. The pipeline starts with an experimental setting for isotopically coding probe-derived modifications (PDMs) that can be automatically recognized by pChem, with masses accurately calculated and sites precisely localized. Further, pChem exports on-demand reports by scoring the profiling efficiency, modification-homogeneity and proteome-wide residue selectivity of a tested probe. The performance and robustness of pChem were benchmarked by applying it to eighteen bioorthogonal probes. Of note, the analyses reveal that the formation of unexpected PDMs can be driven by endogenous reactive metabolites (e.g., bioactive aldehydes and glutathione). Together, pChem is a powerful and user-friendly tool that aims to facilitate the development of probes for the ever-growing field of chemoproteomics.

Chemical probe coupled with mass spectrometry (MS)-based proteomics, herein termed chemoproteomics, offers versatile tools to globally profile protein features and to systematically interrogate the mode of action of small molecules in a native biological system¹. For instance, bioorthogonal probes surrogating endogenous metabolites (e.g., sugars and lipids) enable the proteome-wide mapping of post-translational modifications (PTMs) on specific amino acid residues². In addition, various activity-based protein profiling (ABPP) probes have been developed by targeting amino acid residues including cysteine^{3,4}, lysine⁵, tyrosine⁶, methionine^{7,8}, histidine⁹, aspartate and glutamate^{10,11}, as well as their PTM forms¹²⁻¹⁴, which greatly expand the chemical space in complex proteomes for probing fundamental biology and for discovering new small molecule-based therapies.

Nonetheless, the development of an efficient and selective probe for chemoproteomics can still be challenging. It is particularly difficult to unbiasedly assess its chemoselectivity at a proteome-wide scale, since a chemical probe displaying selectivity well-characterized *in vitro* would possibly generate unexpected modifications owing to potential cross-reactivity in complex biological systems. In addition, unforeseeable probe-derived modifications (PDMs) may be yielded during sample preparation or in-source MS fragmentation, thereby causing inhomogeneous modifications on the same sites and

complicating data analysis.

Notably, the last decade has witnessed tremendous progress in the development of blind search informatic tools¹⁵⁻²³. Such tools, in combination with isotope-coding approaches for probes (**Supplementary Fig. 1**), can provide an unbiased survey of PDMs that can be distinguished from non-probe-derived modifications (e.g., unmodified or endogenously modified ones), considering that only those peptides bearing PDMs would yield an isotopic MS signature (**Fig. 1a**). For example, we have previously used this pipeline (i.e., TagRecon for blind search²³) to substantiate the performance of several newly developed chemoselective probes for proteomic mapping of cysteine redox forms^{12, 13}. The pipeline also allowed us to uncover unexpected PTMs being captured by chemoproteomic probes *in situ*^{14, 24}. Most recently, Hacker and coworkers have applied a similar pipeline (i.e., MSFragger for blind search¹⁸) to systematically investigate the proteome-wide selectivity of diverse electrophilic probes²⁵. These studies underscore the power of blind search tools, which provide an ideal means to unbiasedly assess the proteome-wide residue selectivity of a probe and to uncover new chemotypes in the proteome.

Despite these advances, a set of challenges have emerged. First, most blind search tools cannot automatically unify the localization probability (or residue selectivity) and accurate masses of PDMs, as most of them only offer identification and site localization at a PSM (peptide-spectrum match) level. Second, no available tools can automatically distinguish isotopically coded PDMs from non-probed ones, while manual evaluation of dozens to hundreds of mass shifts can be a daunting task. Last but not the least, for probe developers, even those with substantial bioinformatics expertise, managing the existing tools can be tedious as all require laborious installation and setup, and output many redundant information rather than on-demand reports.

The challenges discussed above have therefore inspired us to develop an automated, user-friendly, fit-for-purpose computational tool for the ever-growing field of chemoproteomics. Here we present pChem, a modification-centric blind search and summarization tool to provide a pipeline for rapid and unbiased assessing of the performance of ABPP and metabolic labeling probes. This pipeline starts experimentally by isotopic coding of PDMs, which can be automatically recognized, paired, and accurately reported by pChem, further allowing users to score the profiling efficiency, modification-homogeneity and proteome-wide residue selectivity of chemoproteomic probe.

Results

The design of pChem. Building upon the [pFind](#) platform¹⁵, pChem first aims to generate all possible modifications using *bona fide* blind search based on a tag-index approach (**Fig. 1b**). Specifically, for each spectrum, a number of 5-mer sequence tags are extracted with a depth-first *de novo* search. The resulting tags are then scored and up to 100 ranked tags are retained before searching against the tag-indexed protein database in order to extend to a full-length peptide sequence. During the extension procedure, all mass shifts within the given range ($\pm 1,000$ Da by default) are considered as modification candidates. Note that the default-defined common modifications (e.g., oxidation of methionines and carbamidomethylation of cysteines) are involved in the sequence tag extraction procedure to improve the sensitivity of this algorithm. Alternatively, a few highly abundant modifications can be automatically detected by the initial pFind-based open-search against Unimod²⁶, a protein modification database for mass spectrometry applications, and then be specified in the following blind search.

Similar to other blind search tools, pChem will initially generate numerous mass shifts at the PSM-level, and then the mass shifts can be calibrated with the system error computed from PSMs identified without any unknown modifications. Specifically, the system error of the precursor ion mass is obtained by

averaging each mass difference between the precursor ion mass and the theoretical mass of the peptide without any modifications or with known modifications. The calibrated mass shifts on PSMs are further grouped together and the mean value is calculated for each group in order to generate the accurate mass for modification candidates and score their residue-specific localization probability (**Fig. 1b**). To eliminate non-PDMs or other unrealistic modifications, all modification-centric mass shifts are automatically paired according to the isotopic mass difference (e.g., 6.020132 Da caused by six heavy carbons) within empirically defined tolerance (i.e., ± 0.001 Da for a fine-tuned Orbitrap instrument). Furthermore, any trace PDM (i.e., byproducts from an unwanted side reaction or rearrangement) with the percent of total PSMs less than 5% by default can be neglected and therefore not reported (See **Methods**). Meanwhile, using an unsupervised outlier-detection approach based on statistical learning, pChem automatically recognizes and reports diagnostic fragment losses (DFLs, e.g., neutral and/or charged losses), if any, which facilitates the characterization of each PDM.

Finally, the pChem results will be exported into the heatmaps showing the site localization probability of all high-confidence PDMs and into several editable files for users to explore more details. Optionally, by automatic retrieving the accurate PDMs for the final round of targeted database search (i.e., a restricted search procedure eliminating the lengthy and laborious process of re-setting and re-searching modifications deduced from numerous mass shifts), pChem can also generate a radar plot showing overall performance outcomes by scoring the profiling efficiency, modification-homogeneity, and residue selectivity (**Fig. 1b**). Note that pChem is an installation-free executable (.exe) program that can be easily used on either laptops or desktops with the step-by-step guidance for operation and troubleshooting (**Supplementary Protocol**).

Performance and robustness of pChem. To benchmark the performance of pChem, we initially generated a QTRP (quantitative thiol reactivity profiling) dataset by using a ‘clickable’ iodoacetamide probe termed IPM (**Fig. 2a**) with a well-established chemoselectivity towards cysteinyl thiols⁴. Specifically, the HEK293T cell lysates were labeled with IPM and then processed into a mixture of isotopically tagged peptides with a predefined light to heavy ratio of 1.0 via CuAAC (Cu^I-catalyzed azide-alkyne cycloaddition, one type of click chemistry reactions)-mediated biotin-conjugation. The isotopically modified peptides were captured with streptavidin and selectively released for MS analysis on a Q-Exactive Plus instrument. For each of three biological replicates, pChem reports the targeted PDM of $\Delta 252.12$ Da (# PSM counts: 7762 ± 448) on cysteine with a high localization probability ($94.7 \pm 1.2\%$, **Fig. 2b**)²⁷ and a minor one ($\Delta 268.12$ Da, # PSM counts: 713 ± 86) that is mainly attributed to the inevitable mis-matching due to an oxidized methionine adjacent to the cysteine bearing $\Delta 252.12$ (**Supplementary Fig. 2, Supplementary Table 1-2, and Supplementary Note 1**). Moreover, pChem scored high on the overall performance of the IPM-based chemoproteomic method (**Fig. 2c**).

To validate the robustness of pChem, we generated additional QTRP samples and analyzed them on eight different instruments from three vendors. Note that the *.RAW data generated by Thermo ScientificTM Orbitrap instruments can be directly imported into pChem, while the *.WIFF (SCIEX) and *.TDF (Bruker) data from Time-of-Flight (TOF) instruments need to be first transformed into *.mzML (mass spectrometry markup language) using MSconvert²⁸. As expected, the results as above could be reproduced (**Supplementary Fig. 3a and Supplementary Table 3**), demonstrating pChem as a cross-platform tool. Next, we sought to evaluate the effect of predefined light to heavy ratios on the performance of pChem. To this end, we analyzed a published QTRP dataset from light and heavy tagged samples mixed in different ratios (L/H = 1:10, 1:5, 1:2, 1:1, 2:1, 5:1 and 10:1) that has been created for

quantification accuracy evaluation⁴. Also, pChem generated the same results, no matter how one pre-mixed the light and heavy tagged samples (**Supplementary Fig. 3b** and **Supplementary Table 3**). Likewise, we found that pChem analysis was insensitive to the types of samples/species and the corresponding protein databases (**Supplementary Fig. 3c** and **Supplementary Table 2**). Taken together, the results above demonstrate pChem as a robust assessment tool for the performance of chemoproteomic probes.

Application of pChem for residue-reactive probes. To further evaluate the performance of pChem, we applied it to re-assess several residue-reactive probes in their usage for chemoproteomics.

First, we used pChem to analyze a set of newly collected datasets generated by using various covalent probes with diverse electrophilic warheads (**Fig. 3a**), including four cysteine-reactive probes (incl. ENE, NPM, PPMS and VSF) and two lysine-reactive probes (incl. NHS and STP). Notably, the analyses not only unbiasedly identified the targeted PDM corresponding to each probe (**Fig. 3b**, **Supplementary Tables 1 and 4**), but also explained the prevalence of selected warheads for cysteine or lysine bioconjugation (**Fig. 3c**). Meanwhile, pChem allowed us to uncover a previously unknown N-term cysteine modification ($\Delta 292.12$) derived from the NPM probe (**Fig. 3b** and **Supplementary Fig. 4**). Interestingly, this modification predominantly occurred at the carboxyl side of lysine or arginine on proteins, suggesting that it formed after tryptic digestion. As shown in **Fig. 3d**, a mechanism was proposed as follows: **a**) NPM rapidly forms Michael adducts with protein cysteine residues. **b**) when peptide bonds are hydrolyzed on the carboxyl side of K/R adjacent to the initially adducted cysteine residue, the newly formed primary amine attacks the succinimido ring to undergo the intramolecular aminolysis²⁹. We also generated a dataset using a ‘cocktail’ of three thiol-reactive probes, including IPM, ENE, and NPM (**Supplementary Fig. 5a**). Notably, all and only the targeted PDMs could be automatically found by pChem (**Supplementary Fig. 5b** and **Supplementary Table 4**), which further validates the sensitivity and accuracy of this tool.

Next, we asked whether pChem could unbiasedly address the long-standing issue regarding the proteome-wide reactivity of the diazirine group that has been widely used in developing photoaffinity labeling probes³⁰. Here we generated a dataset for pChem analysis from a proteome sample labeled with an alkyl diazirine probe (**Fig. 3a**). To our surprise, such a probe that has been considered lack of residue-selectivity preferentially react with glutamic acid (**Fig. 3b**, **Supplementary Fig. 6**, **Supplementary Tables 1 and 4**). Notably, during the preparation of this manuscript, a recent study also systematically uncovered the labeling preference of diazirine probes toward acidic amino acids³¹, in accordance with our observation to some extent.

Finally, we turned our attention to probes enabling enzymatic proximity labeling (**Supplementary Fig. 7a**), which has emerged as the method-of-choice to globally study protein-protein interactions in living biological systems³². For instance, phenol probes have been predominantly used in peroxidase-catalyzed proximity labeling³³. By using restricted search, phenol probes have been shown to almost solely react with tyrosine (i.e., >98% of labeling sites) via live-cell APEX peroxidase-based proximity labeling^{34, 35}. To further investigate the proteome-wide residue selectivity by peroxidase-mediated proximity labeling in a more unbiased fashion, we generated a dataset using two types of peroxidases (i.e., *in cellulo* APEX and *in vitro* HRP) with alkyne phenol (AP) as the labeling probe (**Supplementary Fig. 7b**). The pChem analysis then revealed that APEX-catalyzed probe labeling indeed predominantly occurred on tyrosine in an expected form ($\Delta 372.18$), while an unusual PDM of $\Delta 209.08$ was also mapped onto many lysine sites (**Supplementary Fig. 7c-d** and **Supplementary Tables 1 and 4**). The latter may be formed by the

reaction between lysine and an acyl radical intermediate probably generated from the APEX-catalyzed radical reaction (**Supplementary Fig. 7e**)^{36, 37}. Instead, HRP-mediated labeling sites were mapped on both tyrosine and cysteine (**Supplementary Fig. 7c**). It is worth noting that cysteine was modified by the AP probe via an *o*-quinone intermediate (**Supplementary Fig. 7d-e** and **Supplementary Tables 1 and 4**). In fact, it has been historically known that HRP-catalyzed oxidation of phenols can produce quinonoid reactive species to adduct free thiol group^{38, 39}. Regardless, this finding suggests that the selectivity of phenol probes for proximity labeling may vary in different enzymatic conditions.

Application of pChem for oxoform-specific probes. Cysteine oxoforms, such as sulfenic acid (-SOH) and sulfinic acid (-SO₂H), represent important post-translational approaches to regulate protein functions in a redox-dependent manner⁴⁰. In the last decade, the development of various cysteine oxoform-specific probes has greatly advanced the field of thiol-based redox biology⁴¹. In particular, some of such probes designed for MS-based redox proteomics allowed us to site-specifically map and quantify hundreds to thousands of cysteine oxoforms in complex proteomes (**Fig. 4a**), thereby greatly expanding the cysteine redoxome^{12, 42-45}. Nevertheless, the chemoselectivity of oxoform-specific probes has long been controversial, even for those classic ones (i.e., dimedone for SOH)^{46, 47}. To address this controversy in an unbiased fashion, we used pChem to re-analyze the redox proteomic datasets generated in our laboratory.

As expected, the dimedone-based DYn-2 and the benzothiazine-based BTd, two most used probes for detecting and/or profiling protein SOHs^{48, 49}, produced the corresponding targeted PDMs (i.e., $\Delta 333.17$ for DYn-2 and $\Delta 418.13$ for BTd) that were predominantly localized on cysteine (**Fig. 4b**, **Supplementary Tables 1 and 5**). As compared to DYn-2 and BTd, a phosphonium ylide-based SOH probe termed WYneN (**Fig. 4a**) that is reported most recently⁵⁰ exhibited a superior overall performance (**Fig. 4b-c**). Notably, the pChem analysis revealed that the targeted PDM ($\Delta 252.12$), a triphenylphosphonium (TPP)-loss thioether product of the initial conjugate from WYneN showed a localization probability of >0.95 on cysteine. In the previous report, we also described two other two WYne probes (WYneC/O, **Fig. 4a**), albeit not ones for chemoproteomics due to the lack of profiling efficiency⁵⁰. Note that WYneC/O were found to be able to produce TPP-loss PDMs as well, but they were originally assigned as thiol ester products with theoretical Δ Ms of 265.1063 (C₁₂H₁₅N₃O₄) and 267.0855 (C₁₁H₁₃N₃O₅), respectively (**Supplementary Fig. 8** and **Supplementary Tables 1 and 5**). However, such a hypothesis was proven wrong according to the accurate masses calculated by pChem ($\Delta 265.1441$ for WYneC and $\Delta 267.1213$ for WYneO, **Fig. 4b** and **Supplementary Fig. 9**). Moreover, pChem revealed an additional PDM of $\Delta 291.1207$ for WYneO, which also seemed to be lack of TPP. As such, the formation of these TPP-loss products can be explained by the plausible mechanisms as follows (**Fig. 4d**). WYneC/O first react with endogenous formaldehyde and/or methylglyoxal (MGO, a by-product of glycolysis⁵¹) via the classic Wittig reaction to form α , β -unsaturated ketones followed by the Michael addition reaction of the latter with reduced cysteines or other biological nucleophiles. The new mechanism was further supported by the evidence from an *in vitro* labeling experiment⁵⁰, in which we failed to detect any TPP-loss modifications from WYneC/O. Overall, this mechanism apparently precludes the use of WYneC/O for live-cell labeling protein sulfenic acids in any applications.

In addition, pChem confirmed that DiaAlk (**Supplementary Fig. 10a**) mostly targets protein sulfinic acids by generating the expected PDM of $\Delta 387.17$ (#PSMs: 172) on cysteine with the known neutral or charged losses (**Supplementary Fig. 10b**, **Supplementary Tables 1 and 5**)¹². The analysis also provided two minor PDMs, including $\Delta 471.23$ (#PSMs: 46) and $\Delta 369.17$ (#PSMs: 12) both on tryptophan

(**Supplementary Fig. 10b-d**), which are most likely formed from an unwanted side-reaction by DiaAlk. Specifically, the mechanism is initial attack at the electrophilic nitrogen of DiaAlk by C-2 of the tryptophan indole ring, followed by oxidation-induced ring open at C-3, resulting in an imine conjugate and its BOC (t-butyloxycarbonyl)-loss product (**Supplementary Fig. 10e**). One therefore needs to be cautious when using DiaAlk for any protein-centric analysis (*i.e.*, Western blotting), although this probe can still be applicable for site-centric chemoproteomic profiling of SO₂H.

Application of pChem for metabolite-derived probes. The advances in biorthogonal chemistry have inspired a series of ‘clickable’ metabolites as chemical reporters for various PTMs². For instance, metabolic glycan labeling with the use of unnatural monosaccharides bearing an azide or alkyne reporter has become a widespread approach for studying protein glycosylation⁵². Recently, Qin et al.,⁵³ reported a non-enzymatic reaction of an azido analog of N-azidoacetylmannosamine (Ac₄ManNAz) with cysteine, which may interfere with its glycoproteomic profiling. pChem further confirmed this unwanted S-glycosylation by searching the chemoproteomic data from the Ac₄ManNAz-labeling sample (**Supplementary Fig. 11, Supplementary Tables 1 and 6**). Encouraged by these efforts, we then extended pChem to re-analyze two public datasets that were generated from samples metabolically labeled by alkyne surrogates of endogenous lipid electrophiles, 4-hydroxy-2-nonenal (HNE) and 4-oxo-2-nonenal (ONE), which can readily modify cellular proteomes (**Fig. 5a**)⁵⁴. Notably, our analyses not only reported almost all known PDMs by these probes^{24, 55}, but also uncovered two major ones that have been overlooked in our original publications (**Fig. 5b-c, Supplementary Tables 1 and 6**).

A PDM of $\Delta 309.17$ from aHNE was pinpointed by pChem on peptide N-terminal (**Fig. 5b** and **Supplementary Fig. 12**). Of interest, almost all those peptides bearing this unusual PDM contain one unmodified cysteine residue, indicative of intramolecular rearrangement (**Fig. 5d**). Addition of aHNE to cysteinyl thiol leaves a reactive aldehyde to further react with the amino group on the corresponding peptide N-terminal, affording the carbinolamine intermediate. Followed by the retro-Michael reaction, a stable ketoamide product is formed. Hence, the cysteines on those N-term ketoamide peptide adducts could also be assigned as aHNE-modified sites. As such, ~92.0% of such cysteines were the same as those bearing the targeted PDM of $\Delta 311.18$ (Michael addition adduct).

A PDM of $\Delta 578.22$ from aONE predominantly occurred on lysine (**Fig. 5b** and **Supplementary Fig. 13**). Such a mass shift and that of aONE-derived ketoamide adduct differ by 289.08 Da, which happens to be one molecule of glutathione (GSH, a highly abundant intracellular antioxidant⁵⁶) with a loss of water. Note that a previous report demonstrates that GSH can be rapidly cross-linked to lysines on recombinant proteins by ONE via GSH-ONE-Michael addition and subsequent Paal-Knorr condensation⁵⁷. Likewise, the PDM of $\Delta 578.22$ should also be a C-glutathionylated pyrrole adduct formed in cells by the same mechanism (**Fig. 5e**). Moreover, the lysine-specificity of this PDM was also confirmed by the characteristic loss of aziridinone-based GSH-ONE-lysine conjugate (706.31 Da, **Supplementary Fig. 13**).

Taken together, the two PDMs newly uncovered by pChem allowed us to expand the aHNE/aONE-derived adductomes and to quantify their dynamics (**Supplementary Fig. 14, Supplementary Table 7 and Supplementary Note 2**).

Discussion

Unlike many other blind search engines, pChem, as a fit-for-purpose tool for chemoproteomics, is designed to produce modification-centric, constructive and easy-to-interpret outputs for probe developers.

At a cost, pChem relies on isotopic coding of PDMs that can be achieved by many well-established approaches with detailed protocols in the field of chemoproteomics (**Supplementary Fig. 1a**). For instance, many generalized isotope-coding agents, such as light and heavy clickable biotin/desthiobiotin tags are either commercially available or easy-to-synthesize (**Supplementary Fig. 1b**)^{55, 58-61}. Alternatively, a more direct way, albeit not very cost-effective, is to use isotopically labeled probes^{42, 60, 62}. In addition, pChem is an .exe program free of installation, so its setup-and-run can be easily managed by probe developers, even those with no experience in informatics.

Our benchmarking results establish pChem as a robust tool by demonstrating its compatibility with a variety of data sources and probes. The analyses require neither prior knowledge nor manual inspection, thereby offering a truly unbiased survey of any modifications derived from a tested probe in complex biological systems. This unique feature would greatly accelerate the assessment of proteome-wide selectivity of chemoproteomic probes, the major rate-limiting process during their developments. Considering the continuing needs to develop residue-selective chemistry for probing protein functions⁶³, discovering targeted covalent inhibitors⁶⁴, and advancing single-molecule protein sequencing and fingerprinting technologies⁶⁵, we consider pChem as an attractive tool for broad utilization in the field of proteomics, chemical biology and drug industry.

Moreover, together with a few previous reports²⁴, the pChem analyses reported herein further strengthen the notion that endogenous reactive metabolites (e.g., formaldehyde, MGO and GSH) may possess cross-reactivity with either a probe itself or its PDMs, which has been largely overlooked in the field. This notion necessitates the use of such endogenous small molecules to benchmark chemoselectivity of a probe being developed at early stages. On the other hand, this notion would also inspire exciting pursuits to uncover previously unknown PTMs in biological systems.

Projecting forward, potential improvements of pChem include implementing it into the [pFind](#) studio and/or a website server, supporting data generated from alternative fragmentation techniques (e.g., electron-transfer dissociation, ETD) that can increase modification-specific fragment ions for highly labile PDMs, assessing the proteome-wide reactivity of isotopically coded cross-linkers, and so on.

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Author contributions

J.X.H. performed the experiments, analyzed the data, and wrote the protocol, Z.C.F. designed and implemented pChem, analyze the data, and wrote the protocol, L.F., performed the QTRP experiments for various species, C.P.T. generated the AP and Diazir data sets, F.C.H. acquired funding, H.C. supervised the work, advised on pChem design and revised the manuscript. J.Y. conceived the project, supervised the work, advised on pChem design, analyzed data and the wrote the manuscript with inputs from others.

Competing Interests

The authors declare no competing interests.

Fig. 1. Principle and workflow of pChem. **a**, Isotope-coding of probe-derived modifications (PDMs) yields a paired MS signature (Left) that can be distinguished from non-PDMs (Right, i.e., endogenous and/or common modifications). **b**, Schematic of pChem workflow. MS data is first imported into pChem to perform *bona fide* blind search against the protein sequence database, which generates abundant modification candidates at the PSM level. Meanwhile, PDM-specific diagnostic fragment losses (DFLs) are recognized with a statistic learning approach. Note that the candidates allow modification ($\Delta M'$) incorporated from the common modifications (e.g., methionine oxidation $\Delta 16$ and cysteine carbamidomethylation $\Delta 57$) that can be manually or automatically pre-defined. After accurate mass calibration, those modification candidates with the same calculated $\Delta M'$ from multiple PSMs are unified together and paired according to the predefined isotopic mass difference. Then, several semi-empirical criteria (See **Methods**) are applied to automatically define the high-confidence PDMs and ranked based on abundance. All on-demand information for probe developers is summarized in an output table and heatmaps. Optionally, a radar plot can be generated for illustrating overall performance outcomes by scoring the profiling efficiency (via an additional round of targeted database search automatically invoked by pChem), modification-homogeneity and residue selectivity.

Fig. 2. Performance of pChem. **a**, Chemical structures of IPM and its expected PDM. **b**, Heatmaps generated by the default pChem search (n=3) showing the amino acid localization distribution of the IPM-derived modification $\Delta 252.12$. **c**, A representative radar plot showing the overall performance of IPM for chemoproteomic profiling of the cysteinome.

Fig. 4. Application of pChem for SOH probes. **a**, Chemical structures of the SOH probes, including DYn-2, BTd, and WYneC/O/N. **b**, Representative heatmaps showing the amino acid localization distribution of the pChem-defined PDMs for each indicated probe. **c**, Representative radar plots showing the overall performance of the tested nucleophilic probes for profiling cysteine sulfenic acids. **d**, Plausible mechanisms for the formation of unwanted PDMs from WYneC/O.

Fig. 5. Application of pChem for lipid electrophile-derived probes. **a**, Chemical structures of the lipid electrophile-derived probes, including aHNE and aONE. **b**, Representative heatmaps showing the amino acid localization distribution of the pChem-defined PDMs for each indicated probe. **c**, Pie charts showing the abundance distribution (i.e., number of PSMs) of PDMs for each indicated probe. **d**, Plausible mechanism for the formation of an unexpected N-term modification ($\Delta 309.17$) derived from aHNE. **e**, Plausible mechanism for the formation of GSH-aONE-lysine modification ($\Delta 578.22$).

Methods

pChem algorithm. pChem is an installation-free .exe program relying upon an efficient database search engine termed Open-pFind¹⁵. It is freely available at <http://pfind.ict.ac.cn/software/pChem/index.html>. It can work in any desktops or laptops running Windows operating system by configuring parameters in a text editor. It can directly import the RAW data from Orbitrap instruments or read mzML files transformed by MSconvertGUI²⁸ (part of ProteoWizard v.3.0.21193). Note that pChem requires DDA (data dependent acquisition)-based MS data with both MS1 and MS/MS spectra recorded in the high-resolution mode. The output files of pChem include (**Supplementary Protocol**): (1) an editable summary file containing many key characteristics of the isotopically paired PDMs, such as the number of PSMs and peptides, accurate masses, site-specific localization probability and DFLs; (2) a heat map showing the amino acid localization distribution for each PDM; (3) a radar plot showing the overall performance with the metrics of profiling efficiency, modification homogeneity and residue selectivity.

Blind search. pChem first automatically invokes Open-pFind¹⁵ to perform *bona fide* blind search against the sequence database that is provided as input. This step aims to generate abundant modification candidates at the PSM level using a tag-index strategy by which the sequence database (The FASTA canonical protein sequence databases from various species were obtained from [Uniprot](#)²⁶) is re-constructed into the *tag-indexed protein database* and peptide candidates are retrieved. Unlike the default search mode of Open-pFind¹⁵, pChem does not retrieve preset modification lists (i.e., the Unimod database). For each high-resolution MS/MS spectrum, a number of *k*-mer tags are extracted with a depth-first search, which is similar to *de novo* peptide sequencing⁶⁶. Note that the tag candidates allow modification incorporated from the pre-defined common modifications (e.g., oxidation of methionines and carbamidomethylation of cysteines). Alternatively, the MS/MS data can be initially searched by Open-pFind¹⁵ against Unimod to find the highly abundant non-probe derived modifications, which will then be specified and retrieved in the blind search step.

The tag candidates are then scored and only a few top-ranked ones are retained before searching against the *tag-indexed protein database*. Afterward, peptide candidates are generated by extending each of the matched tags to a full-length peptide sequence. During the extension procedure, all mass shifts within the given range ($\pm 1,000$ Da by default) are retained. Peptides that fit at least one flanking mass of each tag are further retrieved and scored at PSM-level via the same scoring strategy of Open-pFind¹⁵, and the non-zero mass shift (if any) on one side is considered as a potential modification. Then the mass shift is tested in turn on all sites of the peptide except those within the region matched with the tag, to generate different modified peptides. Finally, the modified peptides are scored with the matching spectra via a number of procedures (e.g., reranking, FDR control and protein inference) the same as those of Open-pFind¹⁵.

Calibrating accurate masses of modification candidates. Mass shifts found by blind search are kept to only two decimal places, enabling relatively high computing efficiency. Further, a mass error calibration algorithm is implemented to obtain the highly accurate modification masses. Specifically, the *system error* of precursor ion mass is first calculated by averaging the mass difference error between the measured mass of precursor ion and its matching peptide free of unknown modification as follows:

$$E_{system} = \frac{M_{precursor} - M_{seq} - M_{common}}{M_{seq} + M_{common}} \cdot 10^6$$

where E_{system} denotes the system error in p.p.m., $M_{precursor}$ is the neutral mass of the precursor ion,

M_{seq} is the mass of the corresponding peptide sequence and M_{common} is the summed mass of the common modifications of this peptide if exists.

For each mass shift being assigned to unknown modification at the PSM level, its accurate value can be inferred according to the precursor ion mass calibrated with the aforementioned *system error* and the corresponding peptide sequence as follows:

$$M_{mass} = \frac{10^6}{10^6 + E_{system}} \cdot M_{precursor} - M_{seq} - M_{common}$$

where M_{mass} denotes the accurate mass of unknown modification.

Finally, the accurate mass corresponding to the same modification candidate is unified by averaging the same calculated mass shifts from multiple PSMs as follows:

$$\bar{M}_{mass} = \frac{1}{N} \sum M_{mass}$$

where N denotes the total number of PSMs containing the same modification candidate and $\bar{M}_{mass}$ is the accurate mass computed for the modification candidate. As shown in **Supplementary Fig. 15**, pChem searches can achieve mass accuracy in the low p.p.m. range (i.e., from 0.02 to 19.7 p.p.m., with medium values of 2.4 and 3.8 p.p.m. for light and heavy PDMs characterized in this study, respectively).

Defining PDMs. In general, aforementioned steps will generate and output numerous modification candidates (typically from dozens to several hundreds, **Supplementary Fig. 16**), thereby complicating the result interpretation. To this end, several semi-empirical criteria are applied to automatically recognize the genuine PDMs. First, the isotope coding information is utilized to eliminate non-PDMs or other unrealistic modifications. For the six-heavy carbon coding strategy, the theoretical mass difference between a pair of light and heavy PDMs is 6.020132 Da. If such a measured mass difference is out of the range of $[6.020132 - 0.001, 6.020132 + 0.001]$ Da, the modification will be neglected. Second, less abundant modification candidates with the PSM counting number lower than a predefined threshold (i.e., 5% of total, by default. As a result, the PSMs of the *high-confidence PDMs* generally account for >85% of those of all *isotope-paired PDMs*, **Supplementary Fig. 17**) are also filtered out. Third, only the modifications with masses larger than the pre-defined threshold, e.g., 200 Da by default, are retained. By applying such criteria, typically, less than five high-confidence PDM pairs will be narrowed down and reported in the final output summary. For all probes tested herein, these high-confidence PDMs account for only 2.3% (median, ranging from 0.63% to 13.8%) of initial modification candidates by open-search (**Supplementary Fig. 16**), thereby facilitating result interpretation by users.

Localizing PDMs. Once PDMs are defined, the subsequent statistical analysis is carried out to estimate the corresponding localization probability by calculating the position distribution for each PDM as follows:

$$p_{site_i} = \frac{n_{site_i}}{n_{total}}$$

In this formula, p_{site_i} denotes the localization probability of PDM occurring at $site_i$, n_{site_i} is the number of PSMs related to each PDM that occur at specific $site_i$, and n_{total} denotes the total number of PSMs related to the same PDM. The positions of the top-ranked p_{site_i} are reported in the *pChem.summary* file. Meanwhile, for each PDM, a heat map showing p_{site_i} values on all possible sites will be generated automatically.

Note, in addition to twenty protein-coding amino acids, peptide N- and C-termini are also considered in the modification site determination. Specifically, given a modified peptide whose length is L , if the

modification located at the leftmost amino acid (position 1), it is also regarded as a potential N-terminal modification with a position identifier of 0. Similarly, if the modification located at the rightmost amino acid (position L), it is also regarded as a potential C-terminal modification with a position identifier of $L + 1$. For example, given a peptide sequence CEHVAEADK, if a PDM is identified on the first amino acid C with blind search, the modification is also considered to be located at the N-terminal of the peptide by the algorithm.

Recognizing diagnostic fragment losses (DFLs). Without prior knowledge of the fragmentation patterns, an unsupervised outlier-detection algorithm⁶⁶ can be adopted to automatically recognize potential PDM-specific DFL set ($\Delta = \{\delta_1, \dots, \delta_k\}$) by searching a collection of MS/MS spectra corresponding to each isotopic pair of PDMs. First, for each PSM related to a PDM, all theoretical b - and y -ion masses $T = \{t_1, \dots, t_n\}$ are calculated. For example, given a peptide VQSVEK and the PDM is identified on S, the masses of the theoretical ions with the PDM is referred to as $T' = \{b_3 \sim b_5, y_4 \sim y_5\}$. Let $S = \{s_1, \dots, s_m\}$ be the experimental spectrum peak set. A theoretical fragment ion mass t_i and an experimental peak s_j have an offset $x_{i,j} = t_i - s_j$, which can be considered as a discrete random variable. Since the frequencies of the offsets corresponding to the DFLs are expected to be much larger than those of the random offsets, the peaks in the empirical distribution of the offsets allow to reveal the genuine PDM-specific DFLs. The statistics of offsets over all experimental spectrum peaks and all theoretical peaks provides a reliable learning algorithm to generate DFL set Δ , whose frequency should exceed the half of top one. That is, if the isotopically paired PDMs also generate DFLs that can be paired as mentioned above and the unmatched DFLs will be filtered out as noises (**Supplementary Fig. 18a**). Instead, if the offset with the highest frequency is equal to zero, it suggests that the corresponding PDM is unlikely to generate any significant DFLs (**Supplementary Fig. 18b**). Note that DFL set Δ can be further refined by applying the isotopic mass difference cutoff value as above.

Restricted search. The characteristics of the PDMs (*i.e.*, the accurate mass and the Top1 localization site) are automatically retrieved by pChem for the last round of restricted search in pFind 3, so are the predefined common modifications. This procedure further increases the spectrum identification rate that is measured by the percentage of spectra identified by the search engine among all input spectra within the threshold of 1% FDR estimated by the target-decoy strategy⁶⁷ at the peptide level, thereby facilitating the performance scoring process (see below). Moreover, the restricted search can revise the identification when a PDM and one or more adjacent common modifications on the same peptide sequence are erroneously identified as a 'new' PDM by summing up masses of them.

Probe performance scoring. For tool users such as probe developers, pChem provides a general evaluation of the performance of the tested probe by scoring values as follows:

(1) *Profiling efficiency (%)*, which indicates the capability of the PDMs being identified from the MS data and is calculated as:

$$P_{pe} = \frac{N_{PDM}}{N}$$

where N_{PDM} represents the number of all PSMs assigned to the isotopically paired PDMs (limited to top-5 PDMs by default) and N is the total number of PSMs identified in the restricted search procedure.

(2) *Modification homogeneity (%)*, which measures the modification location uniformity of a PDM and is computed as:

$$p_{mh} = \frac{N_H}{N_{PDM}}$$

where N_H denotes the number of the PSMs assigned to the isotopically paired PDM with the most identified PSMs.

(3) *Residue selectivity (%)*, which evaluates the position selectivity attribution of PDM and can be formed as:

$$p_{rs} = \frac{N_S}{N_{PDM}}$$

where N_S corresponds to the number of PSMs corresponding to the site with the highest number of identified PSMs assigned to the PDMs. Note that the latter two scores are calculated right after the blind search procedure.

Chemoproteomic data sets published previously. Raw data files previously generated by using the IPM-based QTRP method are available from ProteomeXchange (PXD027762 for evaluating quantification accuracy⁴; PXD027767 for profiling the cysteinome in *Drosophila melanogaster*⁶⁸, *Caenorhabditis elegans*⁴³, *Pseudomonas syringae*⁶⁹, and *Mus musculus*⁷⁰). Raw analyses of chemoproteomic data sets based on oxoform-specific probes (i.e., DYn-2, BTd, WYnes and DiaAlk) have been described previously and are available now from ProteomeXchange (PXD027764)^{12-13, 42, 49-50}. The aHNE- and aONE-based chemoproteomic analyses have also been reported previously^{24, 55}, and the corresponding raw data sets can be obtained from ProteomeXchange (PXD027760 and PXD007149, respectively). Raw data files for the Ac₄ManNAz-based glycoproteomic analysis were generated and kindly provided by Prof. Xing Chen and colleagues⁵⁹.

Chemoproteomic data sets newly generated in this study. Raw data files newly generated by using the IPM-based QTRP method are available from ProteomeXchange (PXD027755 for initial test; PXD027758 for data collected from different instruments; PXD027767 for profiling the cysteinome in *Arabidopsis thaliana*, *Escherichia coli*, and *Rattus norvegicus*). QTRP data files generated by using other thiol-reactive probes (i.e., ENE, NPM, PPMS, VSF, and 'cocktail') are available from ProteomeXchange (PXD027756). Raw data files from the NHS- and STP-based lysinome profiling are available from ProteomeXchange (PXD027789). The AP- and Diazir-based chemoproteomic data sets are available from ProteomeXchange (PXD027591).

Sample preparation. All probes used here contain a 'clickable' alkyne tag, one of most-used functionalized handles for chemoproteomics¹. For pChem search, a well-established quantitative chemoproteomic workflow relying on the commercially available light and heavy azido-biotin reagents with a photocleavable linker was utilized to isotopically code alkyne probe-derived modifications (**Supplementary Fig. 1b**). More details are described as follows.

Reagents. IPM, ENE and VSF were house-made as previously described⁷¹⁻⁷². NPM (cat. No. TA113) was purchased from KeraFast. PPMS (cat. No. P757300) was purchased from Toronto research chemicals. STP (cat. No. 30720) was purchased from Lumiprobe. NHS (cat. No. A171448) was purchased from Aladdin. AP (9186096) was purchased from J&K Scientific. The Diazir probe was kindly provided by Prof. Zheng-Qiu Li⁷³. Iodoacetamide (IAA, cat. No. V900335), tris[(1-benzyl-1H-1,2,3-triazol-4-yl)methyl]amine (TBTA) (cat. No. 678937), and sodium ascorbate (cat. No. A7631) were purchased from Sigma-Aldrich.

Dithiothreitol (DTT, cat. No. A620058-0025) was purchased from BBI Life Sciences; Light Azido-UV-Biotin (cat. No. EVU102), and Heavy Azido-UV-Biotin (cat. No. EVU151) were purchased from KeraFast; Sequencing grade trypsin (cat. No. V5113) was purchased from Promega. Streptavidin sepharose high performance (cat. No. 17-5113-01) was purchased from GE. All other reagents were purchased from Thermo Fisher Scientific unless otherwise noted.

Preparation of cell lysates. Human HEK293T cells and rat CRL-1444 cells were purchased from the National Infrastructure of Cell Line Resource (China), cultured in DMEM supplemented with 10% FBS (cat. No. 10099141), 1% penicillin-streptomycin (Cell World, cat. No. C0160-611) and 1% Glutagro (Corning, cat. No. 25-015-CI), maintained at 37°C in a 5% CO₂ humidified atmosphere. Cells were grown to ~80% confluency, washed with prechilled PBS, and lysed in pre-chilled lysis buffer (50 mM HEPES (pH 7.6), 150 mM NaCl, and 1% IGEPAL) supplemented with 1 x protease and phosphatase inhibitors (cat. No. A32961).

E.coli strain MG1655 was cultivated 50 mL M9 minimal media containing 0.2% casamino acids and 10 mM glucose to an optical density at 600 nm of ~0.3. 4 mL of culture was then transferred into 15 mL conical centrifuge tubes with 5x minimum inhibitory concentration (MIC) of antibiotics and incubated for additional 4h. The culture was centrifuged at 1,400 × g for 10 min. Cell pellets were harvested by centrifugation and lysed as above.

Arabidopsis protoplasts were prepared as previously described⁷⁴. In brief, well-expanded leaves from *Arabidopsis thaliana* (3-4 weeks) were sliced into small leaf strips (0.5-1 mm) and incubated in 20 mM MES (pH 5.7), 1.5% (w/v) cellulase R10, 0.4% (w/v), macerozyme R10, 0.4 M mannitol, 20 mM KCl, 10 mM CaCl₂, and 0.1% BSA. The leaf strips were vacuum-infiltrated in a desiccator for 30 min and incubated for additional 3h at RT in the dark. The digested sample was then filtrated with a 75 µm nylon mesh. The protoplasts were pelleted by centrifugation at 100g for 2 min, resuspended in W5 solution (2 mM MES, pH 5.7, 154 mM NaCl, 125 mM CaCl₂ and 5 mM KCl), pelleted again on ice for 30 min by gravity, and lysed in prechilled HEPES buffer (50 mM, pH 7.6) containing 150 mM NaCl, 1mM EDTA, 0.5% Triton X-100, and 1% IGEPAL, 1 x protease and phosphatase inhibitors.

Preparation of the probe-labeled proteomes. The cell lysates were then incubated with each probe as indicated with the conditions (i.e., concentration, time, temperature) summarized in **Supplementary Table 8**. The resulting protein samples were reduced with DTT (10 mM, 1 h, RT), and subsequently alkylated with IAA (40 mM, 1 h, RT, with light protection). Proteins were then precipitated with a methanol-chloroform system (aqueous phase/methanol/chloroform, 4:4:1 (v/v/v)). With sonication, the precipitated proteins were resuspended in 50 mM ammonium bicarbonate, and digested with trypsin at a 1:50 (enzyme/substrate) ratio overnight at 37°C. The tryptic digests were desalted with HLB extraction cartridges (Waters, cat. No. 186000383), dried under vacuum, and resuspended in a water solution containing 30% acetonitrile. CuAAC reaction was performed at RT for 2 h with rotation and light protection by subsequently adding 1 mM either light or heavy Azido-UV-biotin (1 µL of a 40 mM stock), 10 mM sodium ascorbate (4 µL of a 100 mM stock), 1 mM TBTA (1 µL of a 50 mM stock), and 10 mM CuSO₄ (4 µL of a 100 mM stock). The light and heavy isotopic tagged samples were then mixed immediately following CuAAC, cleaned with strong cation exchange (SCX, Nest group, cat. No SMM HIL-SCX) spin columns, and then enriched with streptavidin for 2 h at RT. Streptavidin beads were then washed with 50 mM NaAc (pH=4.5), 50 mM NaAc containing 2 M NaCl (pH=4.5), and deionized water twice each with end-to-end rotations, then resuspended in 25 mM ammonium bicarbonate, transferred to glass tubes

(VWR), and irradiated with UV lamp at 365 nm (2 h, RT, with magnetic stirring). The supernatant was collected, concentrated under vacuum, and desalted with HLB cartridges. The resulting peptides were evaporated to dryness and reconstituted in 0.1% formic acid for LC-MS/MS analysis.

LC-MS/MS. The LC-MS/MS data sets (incl. 44 representative files) used here for benchmarking pChem were generated on ten different LC-MS/MS instruments by three vendors from four independent laboratories (**Supplementary Table 9**). The settings for each LC-MS/MS instrument are summarized in **Supplementary Table 10**.

Data analysis. For pChem search, the running time depends on the computer configuration, the complexity of the tested probe-label peptide samples, and the types and settings of LC-MS/MS instruments. Here, all analyses were performed on a desktop PC running Microsoft Windows 10 Home (v.19041.110), with one 2.90GHz CPU Intel i7-10700 processor with 64 GB of installed RAM. In this computing system, for example, approximately 20 min would be required for the default pChem search of the data generated from a single 75-min EasyLC tandem Q-Exactive plus analysis of the IPM-based QTRP sample. The running times used for pChem search of MS data generated from other instruments are shown in **Supplementary Fig. 3a**.

For the pFind 3-based re-analyses of aHNE and aONE data sets, MS data files were searched against *Homo sapiens* Uniprot canonical database (Downloaded on May 16, 2021, 20,395 entries). Precursor ion mass and fragmentation tolerance were set as ± 10 p.p.m. and ± 20 p.p.m., respectively. The maximum number of modifications allowed per peptide was three, as was the maximum number of missed cleavages allowed. Common modifications (i.e., oxidation of methionines and carbamidomethylation of cysteines) and PDMs (refer to **Supplementary Table 1**) were all searched as variable modifications. A differential modification of 6.020132 Da on PDM was used for stable-isotopic quantification. The FDRs at spectrum, peptide, and protein level were $\leq 1\%$. The MS1-based quantification was performed using pQuant⁷⁵, which calculated $R_{H/L}$ values based on each identified MS scan with a 15 p.p.m.-level m/z tolerance window and assigned an interference score (σ) to each value from zero to one. In principle, the lower the calculated σ was, the less co-elution interference signal was observed in the extracted ion chromatograms. In this regard, the median values of probe-modified peptide ratios with σ less than or equal 0.5 were considered to calculate site-level ratios. Quantification results were obtained from three biological replicates with two LC-MS/MS runs for each.

Data Availability

The newly generated chemoproteomic data sets have been deposited to the ProteomeXchange Consortium via the PRIDE⁷⁶ partner repository with the dataset identifiers PXD027755 (Username: reviewer_pxd027755@ebi.ac.uk, Password: mub4elda), PXD027758 (Username: reviewer_pxd027758@ebi.ac.uk, Password: 46z1w0cj), PXD027767 (Username: reviewer_pxd027767@ebi.ac.uk, Password: EZL1H061), PXD027789 (Username: reviewer_pxd027789@ebi.ac.uk, Password: T9s2rIJ3), PXD027756 (Username: reviewer_pxd027756@ebi.ac.uk, Password: leSLiwOa); previously published data were also used to benchmark pChem in repositories with identifiers PXD027591 (Username: reviewer_pxd027591@ebi.ac.uk, Password: WgBqKEGi), PXD007149), PXD027764 (Username: reviewer_pxd027764@ebi.ac.uk, Password: Zje4S2sN), PXD027762 (Username: reviewer_pxd027762@ebi.ac.uk, Password: vLSe4BNQ), PXD027760 (Username:

reviewer_pxd027760@ebi.ac.uk, Password: djulY94L)

Code availability

pChem is open-source and is freely available at <https://github.com/pFindStudio/pChem> under a permissive license.

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