

Deep Interrogation of Metabolism Using a Pathway-Targeted Click-Chemistry Approach

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Cite This: <https://dx.doi.org/10.1021/jacs.0c06877>



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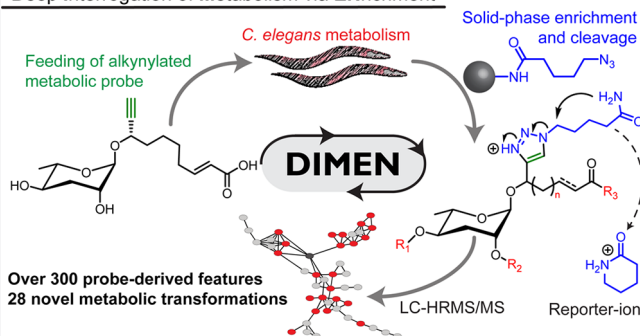
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ABSTRACT: Untargeted metabolomics indicates that the number of unidentified small-molecule metabolites may exceed the number of protein-coding genes for many organisms, including humans, by orders of magnitude. Uncovering the underlying metabolic networks is essential for elucidating the physiological and ecological significance of these biogenic small molecules. Here we develop a click-chemistry-based enrichment strategy, DIMEN (deep interrogation of metabolism via enrichment), that we apply to investigate metabolism of the ascarosides, a family of signaling molecules in the model organism *C. elegans*. Using a single alkyne-modified metabolite and a solid-phase azide resin that installs a diagnostic moiety for MS/MS-based identification, DIMEN uncovered several hundred novel compounds originating from diverse biosynthetic transformations that reveal unexpected intersection with amino acid, carbohydrate, and energy metabolism. Many of the newly discovered transformations could not be identified or detected by conventional LC-MS analyses without enrichment, demonstrating the utility of DIMEN for deeply probing biochemical networks that generate extensive yet uncharacterized structure space.

Deep Interrogation of Metabolism via ENrichment



INTRODUCTION

Small molecules represent the chemical input, biosynthetic intermediates, and endproducts of cellular activity, and thus metabolomics is essential for understanding biological processes, complementing proteomics, transcriptomics, and genomics.¹ In addition to their roles as building blocks and substrates in primary metabolism, many metabolites serve as signaling molecules, from the intra- and intercellular level, e.g., as hormones, to signaling at the interorganismal level via pheromones. Metabolomics faces challenges that do not apply to other members of the -omics family because the structural diversity of metabolites is not restricted by a limited number of templates and simple rules for assembly, as in the case of DNA, RNA, and proteins, and thus is vast and largely unpredictable. Correspondingly, LC-HRMS-based surveys of animal and plant metabolomes suggests that the number of distinct metabolites in a single species may exceed 100 000.² Future improvements of analytical instrumentation may reveal even greater compound diversity. In parallel, studies of signaling networks in *C. elegans* and other systems indicate that biogenic small molecules can have profound biological activity even at femtomolar concentrations, providing a clear rationale for systematic structural and functional characterization of low abundance metabolites.^{3–6} Given these challenges, there exists a need for methods that facilitate prioritization of detection

and structural characterization of specific compound families and biosynthetic pathways of interest.

We aimed to develop a click-chemistry-based strategy for targeted metabolomics that would enable in-depth exploration of specific metabolic pathways and compound families, beyond current detection limits, by (i) improving sensitivity through enrichment from overwhelming pools of unrelated metabolites, and (ii) facilitating recognition of relevant metabolites by means of a specific fragmentation signature in LC-MS/MS (Figure 1a). We envisioned that this could be accomplished by alkyne-labeling metabolic precursors or building blocks of interest, followed by incubation with the biological system and enrichment of labeled derivatives via click chemistry.⁷ Surprisingly, there are few examples of click chemistry-based enrichment of tagged metabolites to interrogate metabolism, whereas click chemistry has been employed extensively to enrich proteins and peptides.^{8,9}

Received: June 25, 2020

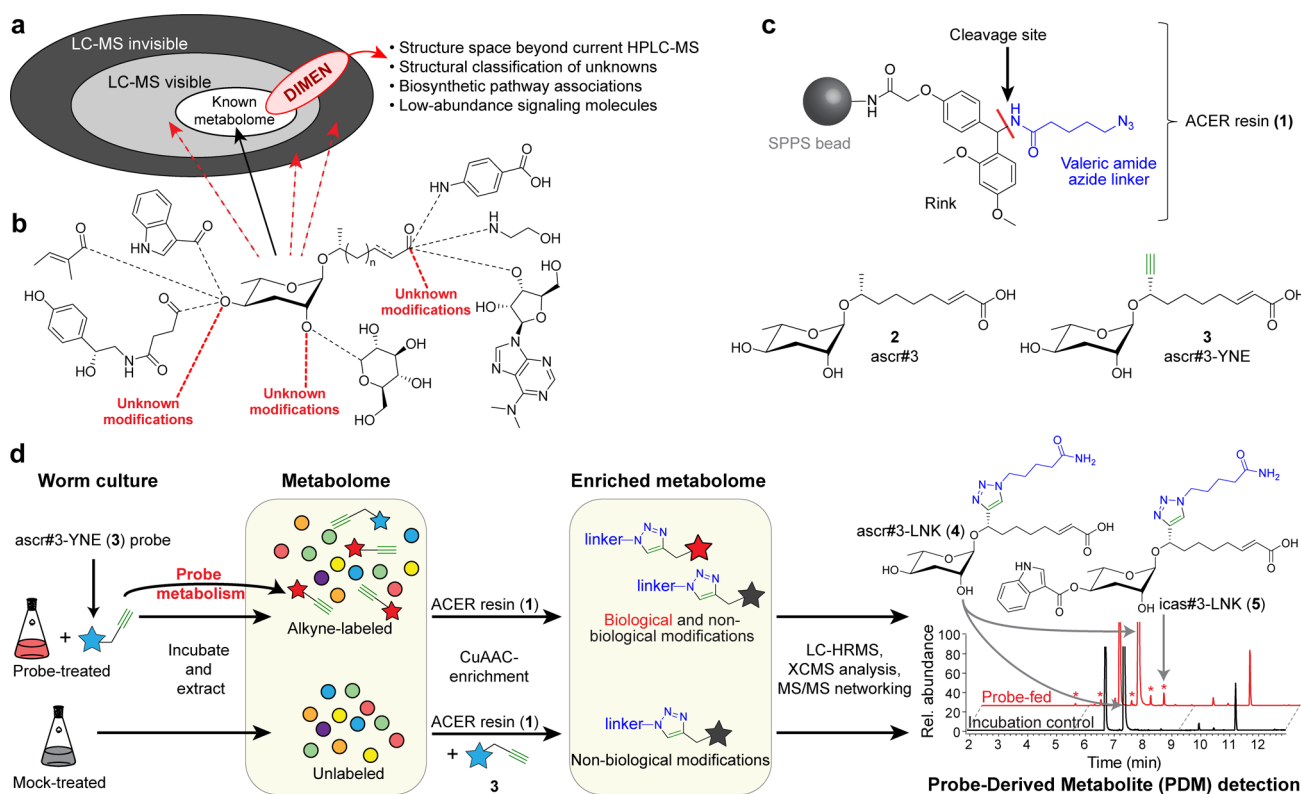


Figure 1. Development of DIMEN. (a) Schematic representation of metabolomes and purpose of DIMEN for the characterization of known, LC-MS visible (light gray) and LC-MS invisible (dark gray) molecules. (b) Glycosides of the dideoxy sugar ascarlyose are combined with diverse primary metabolic building blocks in *C. elegans*. (c) Structure of solid-support, alkyne capture and reporter-ion installation (ACER) resin (1), and alkyne probe ascr#3-YNE (3), which mimics ascr#3 (2), a precursor upstream of the biosynthesis of diverse signaling molecules. (d) Schematic overview of DIMEN analysis. *C. elegans* are grown with and without alkyne-labeled probe (blue star) and extracted. Probe is added to unlabeled samples before enrichment using the ACER resin. Enriched metabolomes are analyzed via LC-HRMS/MS to detect PDMs, e.g., ascr#3-LNK (4) and icas#3-LNK (5).

To demonstrate the utility of click chemistry-based enrichment for metabolomics, we selected a family of signaling molecules, the ascarosides, which play a central role in the life history of the model organism, *C. elegans*, and other nematodes (Figures 1b and S1a). The ascarosides form a large, structurally diverse family of compounds that is derived from modular assembly of building blocks from carbohydrate, amino acid, nucleoside, and cofactor metabolism around a core glycoside derived from the dideoxy sugar ascarlyose and a variable fatty acid-like moiety.^{10–12}

Ascarosides regulate development, aging, phenotypic plasticity, and many behaviors in *C. elegans* and other nematodes, and furthermore mediate multitrophic interactions of nematodes with fungi and plants.^{13–17} Biological activity has been observed down to femtomolar concentrations, and correspondingly ascaroside identification and metabolism has been extensively studied.^{4,12,18,19} Previous studies demonstrated uptake of synthetic ascarosides by *C. elegans*, and that introduction of an alkyne moiety in the fatty acid side chain proximal to the glycosidic linkage does not interfere with biological activity (Figure 1c).^{20–22} Feeding worms an alkyne-modified ascaroside probe may thus provide deep insight into ascaroside metabolism by enabling enrichment and recognition of derived metabolites, including derivatives of potential signaling molecules that have remained undetectable in conventional LC-HRMS.

RESULTS

Development of DIMEN. Our workflow for deep interrogation of metabolism via enrichment (DIMEN) employs the copper(I)-catalyzed azide–alkyne cycloaddition (CuAAC) to selectively bind probe-derived metabolites (PDMs) to an azide-modified solid support, using unlabeled metabolic samples with added probe as controls (Figure 1c,d). The resin-bound PDMs are extensively washed to remove unlabeled metabolites and cleaved to yield PDMs including a triazole moiety that confers a unique MS/MS signature or reporter-ion. PDMs are detected via comparative analysis of LC-HRMS data for the experimental and control samples. Finally, MS/MS fragmentation patterns are used to confirm the probe origin of the detected PDMs and further to discern structurally distinct families of PDMs via molecular networking software.²³

As the solid support we selected a Rink-amide resin typically used for solid-phase peptide synthesis (SPPS) to which we attached a simple azido-valerate linker (“ACER resin” (1), Figure 1c).²⁴ Initially we attempted to use more elaborate azide designs integrating a copper-chelating moiety in the azide linker attached to the solid support, which accelerate click chemistry and improve yields in some situations (Figure S2).^{25,26} However, in our case integration of a copper chelator in the azide linker caused extensive oxidative damage to the resin and, in some cases, also reduced stability of the linker (Figure S3). Lastly, the use of these more elaborate linkers

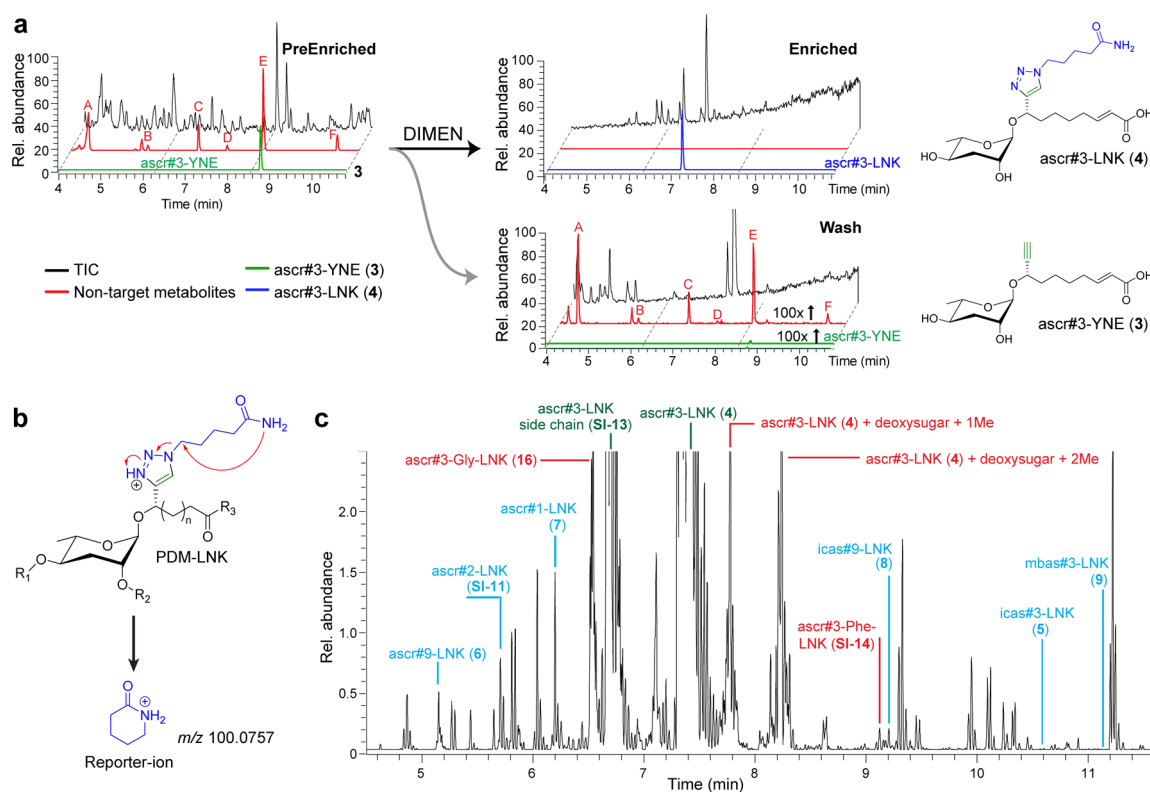


Figure 2. Enrichment of PDMs with DIMEN. (a) LC-HRMS chromatograms for a typical enrichment experiment. Peaks of highly abundant nontarget metabolites (red) are absent following enrichment and remain in the flow-through (intensities adjusted for sample volumes). (b) MS/MS of PDMs in positive mode produces a reporter-ion at m/z 100.0757. (c) Extracted-ion-chromatogram (EIC) for the reporter ion m/z 100.0757 of a DIMEN-enriched sample, showing peaks representing known metabolic transformations (annotated in blue), novel transformations (examples annotated red), and unmodified probe and nonbiological cleavage products (annotated in green). See Figure S1 for structures.

confounded MS/MS analysis by saturating spectra with linker-associated fragments.

Screening different azide designs further demonstrated that exposure to CuAAC conditions can result in unexpected side reactions. To distinguish such CuAAC side products and probe-derived artifacts from compounds that are the result of biological transformations, DIMEN includes an incubation control arm in which the alkyne probe is added to an unlabeled metabolome sample (Figure 1d). Both the probe-treated and mock-treated metabolomes were then incubated with the ACER resin (1) and subjected to CuAAC, resulting in similar artifact profiles for the control and experimental arms and facilitating recognitions of “real” PDMs via comparative LC-HRMS analysis (Figure 1d).

Optimization of enrichment conditions was accomplished by doping concentrated *C. elegans* metabolome with a trace amount of the alkyne derivative of ascr#3 (2), ascr#3-YNE (3), and monitoring the recovery of the resulting linker-modified ascr#3-LNK (4). ascr#3 is one of the major ascarosides produced by *C. elegans* and can serve as a precursor for other known ascarosides.^{4,27} Different buffers, copper chelating ligands, reagent concentrations, and types of SPPS resins were tested (Figure S4). On the basis of these experiments we selected a polyethylene glycol acrylamide (PEGA) resin as solid support for the azide linker and 2-(4-((bis((1-(*tert*-butyl)-1*H*-1,2,3-triazol-4-yl)methyl)amino)methyl)-1*H*-1,2,3-triazol-1-yl)acetic acid (BTAA) as copper chelating ligand.²⁸ Analysis of the resin supernatant following enrichment indicated that more than 99% of unlabeled metabolites had

been removed and that ~80% of added ascr#3-YNE (3) had reacted (Figure 2a).

Triazoles generally ionize well in positive-ion electrospray mode, facilitating mass spectrometric detection of PDMs (Figure S4d). Furthermore, use of the azido-valeryl modified rink resin resulted in PDMs that produce a diagnostic MS/MS fragment with a mass-to-charge of 100.0757 (Figures 2b and S5) that can be employed as a reporter-ion to automate PDM detection and corroborate that detected metabolites are in fact derived from ascr#3-YNE (Figure 2c and S6).

DIMEN Reveals Novel Metabolites. We then employed the optimized enrichment protocol to investigate metabolic transformations of ascr#3-YNE (3) in *C. elegans*. LC-HRMS/MS-based comparison of experimental and incubation control samples revealed more than 10 000 features, of which 585 were present only in the experimental samples and thus classified as likely PDMs (Figure 3a,b). Molecular networking revealed 27 clusters in which three or more nodes represent likely PDMs, most of which formed defined subclusters due to the shared reporter-ion fragment (Figure 3a).

Detailed analysis of MS/MS spectra representing likely PDMs revealed expected derivatives of known metabolites as well as a large number of features that appeared to represent novel metabolic transformations. PDMs representing known metabolites (e.g., PDMs 5–9) include the linker analogues of side chain-shortened ascarosides, e.g., the linker derivative of the dauer pheromone component ascr#1 (10), ascr#1-LNK (7) (Figures 3c and S1, Table S1).²⁹ Formation of ascr#1-LNK (4) and other chain-shortened derivatives of ascr#3-YNE (3) indicates that, similar to unmodified ascr#3 (2), the added

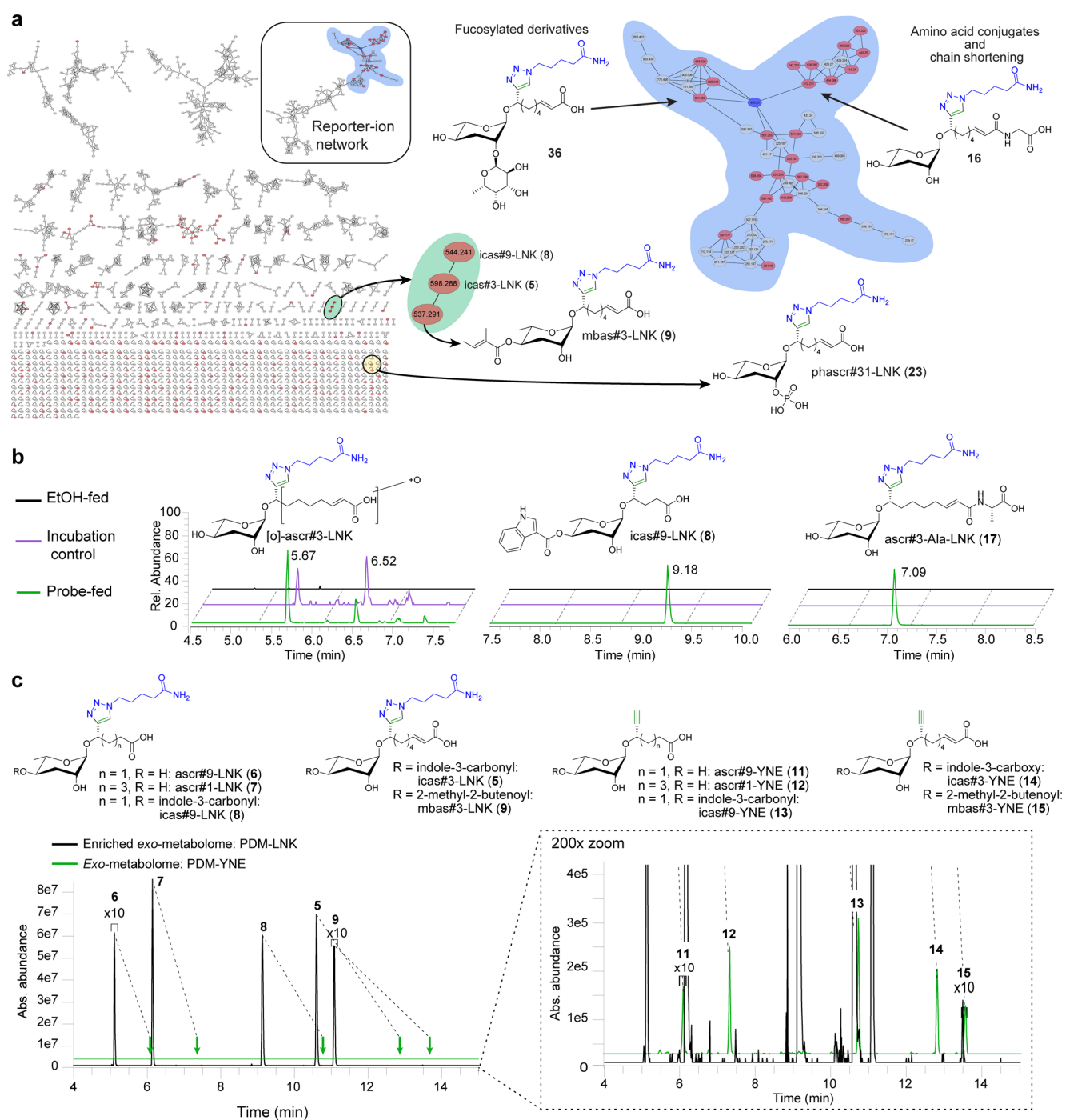


Figure 3. DIMEN of ascr#3-YNE-fed *C. elegans*. (a) Molecular network resulting from DIMEN analysis of ascr#3-YNE-treated *C. elegans* cultures. The blue node is ascr#3-LNK and the red nodes are differential between probe-fed and incubation control data sets. These include analogues of known ascarosides such as mbas#3-LNK (9) as well as several new compound families, e.g., a set of fucosylated PDMs, amino-acid ligated ascarosides, and phosphorylated ascarosides. (b) Example EICs for probe-derived artifacts detected in both the probe-fed and incubation control data sets that originate from abiological transformations (e.g., [O]-ascr#3-LNK, a product of nonenzymatic oxidation of ascr#3), PDMs only present in probe-fed data sets representing known ascaroside derivatives (e.g., icas#9-LNK (8), a derivative of the known icas#9), and PDMs only present in probe-fed data sets that represent novel metabolites (e.g., ascr#3-Ala-LNK (17)). (c) EICs filtered for alkyne-functionalized PDMs (PDM-YNE) in *exo*-metabolome (green), and linker-bound PDMs (PDM-LNK) in enriched *exo*-metabolome (black). DIMEN improves signal intensity >200-fold between pairs of PDM-YNE and PDM-LNK, due to enrichment and improvement of ionization efficiency (Figure S4d).

ascr#3-YNE (3) enters peroxisomal β -oxidation, which is known to iteratively shorten ascaroside side chains, in two-carbon steps, e.g., from 9 carbons as in ascr#3 to 7 carbons as in ascr#1.^{12,30} Another example for a PDM representing a known metabolite is icas#9-LNK (8), the linker-analogue of the aggregation pheromone icas#9 (Figure 3b and S1).^{20,31}

The detection of icas#9-LNK (8) suggests that metabolism of the probe ascr#3-YNE (3) closely mimics that of natural ascr#3 (2), which had previously been shown to undergo chain shortening and 4'-modification, e.g., attachment of an indolecarboxy moiety.⁴ Several other PDMs also represented known 4'-modified ascarosides, e.g., icas#3-LNK (5), mbas#3-

LNK (9), and others (Figures 3a and S1, Table S1). Quantitation of PDMs representing known metabolites (5–9, 11–15) in a typical DIMEN experiment indicated a signal-to-noise gain of over 2 orders of magnitude, as a result of the enrichment and generally better ionization of PDM-LNK derivatives compared to the PDM-YNE precursors (Figure 3c and S4d).

Among PDMs representing novel metabolic transformations, we prioritized identification of the three largest subclusters in the MS/MS network, which revealed previously unreported ascarosides derived from amino acid ligation and phosphorylation, as well as di- and trisaccharides derived from attachment of fucose derivatives to the ascarose. For each of these groups of PDMs, we proposed structures based on MS/MS analysis and subsequently reanalyzed the metabolome of unsupplemented *C. elegans* for candidate compounds that could constitute the corresponding naturally occurring ascaroside derivatives. To corroborate that such candidate compounds represent ascaroside derivatives, we confirmed their absence in the metabolome of *daf-22* mutants, which are known to be defective in ascaroside biosynthesis.^{12,32,33} Finally, the structures of representative members for each of the identified new families of *C. elegans* metabolites were confirmed by synthesis (see SI Methods).

Amino Acid-Ligated Ascarosides. Analysis of MS/MS spectra of one of the largest clusters of PDMs suggested that they represent amino acid conjugates of ascr#3-YNE (3) (Figure 4a). PDMs representing putative conjugates of glycine, alanine, and serine (16–18, Figures 3a,b, and 4a) were most abundant, although trace amounts of almost all proteinogenic amino acids could be detected (Figure S7). LC-HRMS analysis of the metabolome of unsupplemented *C. elegans* revealed several low abundance peaks for the expected *m/z* values that could plausibly represent amino acid conjugates of ascarosides as their production was abolished in *daf-22* mutants (19–21, Figure 4b), but peak intensities were too low to obtain MS/MS data to confirm structures. We then surveyed available metabolic data for other ascaroside biosynthetic mutants, including mutants of acyl-CoA oxidase *acox-1.1*, in which production of saturated ascarosides with a side chain of 9 carbons is greatly increased (Figure S8).^{4,34} We found that *acox-1.1* mutants produce larger amounts of compounds that have MS/MS spectra consistent with amino acid conjugates of saturated ascarosides with 9 carbon and longer side chains (19–21, Figures 4b,c and S9), e.g., ascr#10-Phe (22), whose structure was independently confirmed by synthesis (Figures 4c and S10). These data established amino acid conjugates as a new family of ascaroside derivatives which under standard laboratory conditions are produced only in very low concentrations in wild-type (Figure S9b).

Phosphorylated Ascarosides. MS/MS spectra of a second family of PDMs suggested that they represent phosphorylated ascarosides, including phascr#31-LNK (23) and phicas#31-LNK (24) (Figures 3a, 5a,b). Their structures were inferred primarily from analysis of positive ion MS/MS spectra, since negative ion MS/MS spectra were dominated by phosphate fragments (Figures 5b and S11). LC-HRMS analysis of the metabolome of unsupplemented *C. elegans* revealed a *daf-22*-dependent compound whose MS/MS spectra and retention time were consistent with a phosphorylated ascr#3, (phascr#31, 25). Subsequent targeted analysis of the *C. elegans* metabolome revealed a series of abundant compounds whose MS/MS spectra and retention times

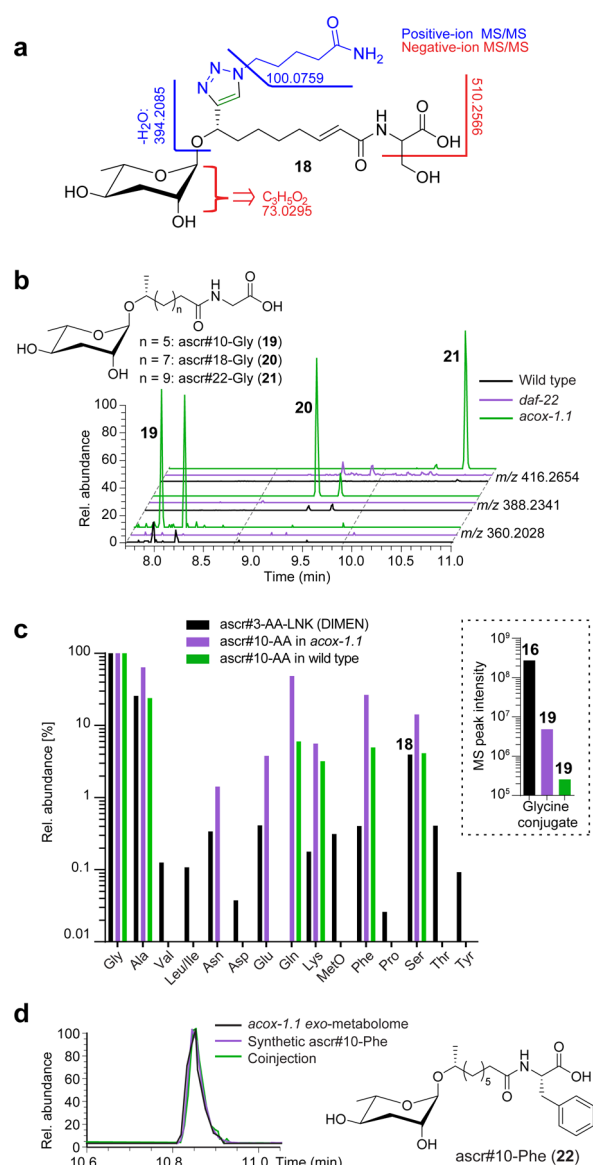


Figure 4. PDMs from amino acid ligation of ascr#3-YNE. (a) MS/MS fragmentation of ascr#3-Ser-LNK (19). (b) EICs $[M-H]^-$ for ascaroside glycine conjugates 19–22 in *acox-1.1*, wildtype, and *daf-22* exo-metabolomes. (c) Relative abundances for PDMs representing ascr#3-amino acid-LNK conjugates as detected by DIMEN (black) as well as for natural ascr#10-amino acid conjugates in *acox-1.1* mutants and wild-type worms (purple and green, respectively), normalized to the abundance of the corresponding glycine conjugate. Absolute abundances of the ascaroside-glycine conjugates are shown in the inset. (d) EICs for ascr#10-Phe (23) in *acox-1.1* exo-metabolome, synthetic ascr#10-Phe, and coinjection confirm its identification.

suggested that they represent phosphorylated ascaroside derivatives (Figures 5c–f). To confirm these assignments and determine the site of phosphorylation, we synthesized the 2'- and 4'-phosphorylated derivatives of ascr#15 (26), phascr#151 (27), and phascr#152 (28), respectively (Figures 5e and S12). Comparison of retention times of natural and synthetic samples demonstrated that natural phosphorylated ascr#15 carries the phosphate in the 2'-position and further indicated that *C. elegans* produces a series of 2'-phosphorylated derivatives (29–32) of almost all known simple ascarosides (2, 10, 33–35) (Figure 5d).

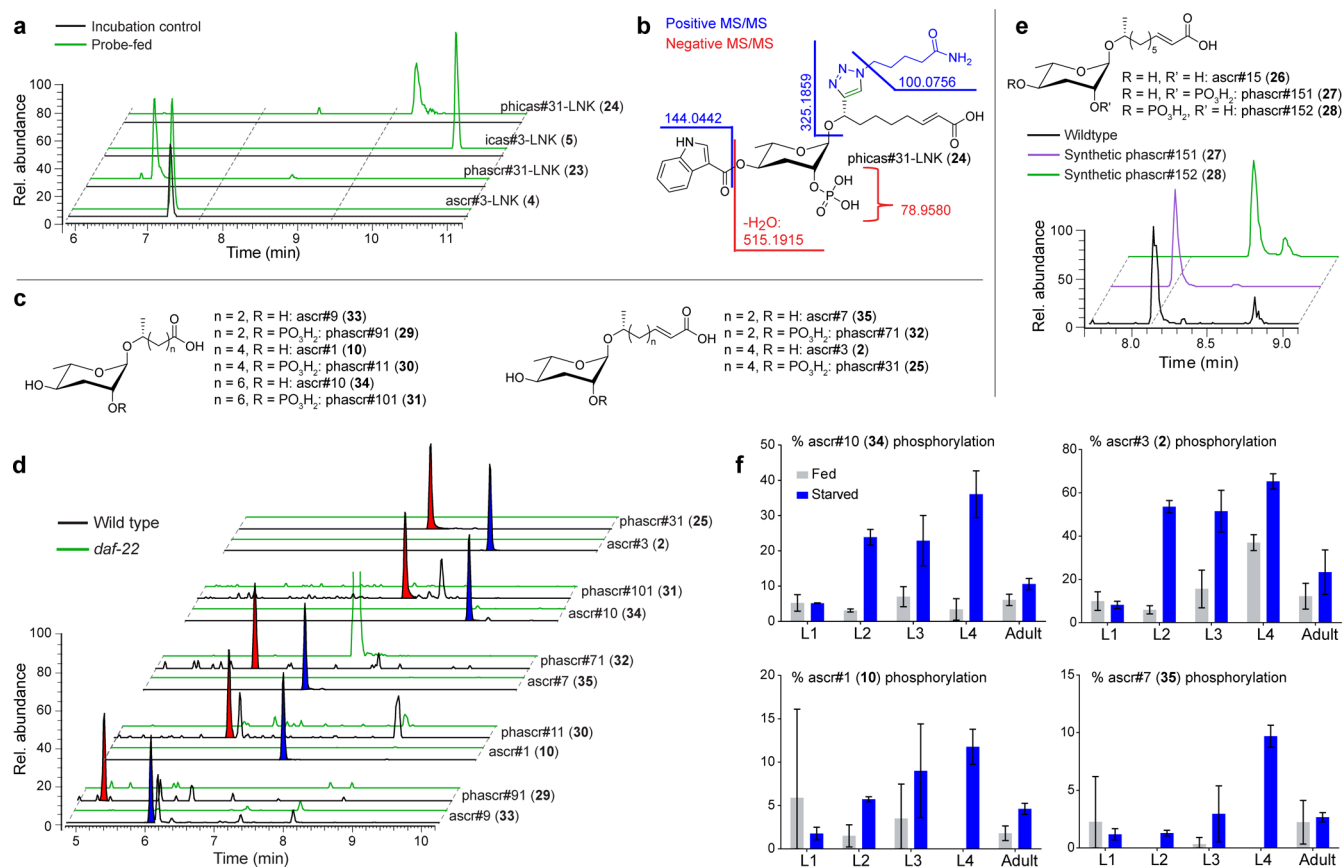


Figure 5. Identification of phosphorylated PDMs. (a) EICs for features representing phosphorylated PDMs in probe-fed and incubation control data sets. (b) MS/MS analysis of phicas#31-LNK (24). (c) Structures of simple ascarosides and their phosphorylated derivatives. (d) EICs for compounds in panel (c) in wildtype and *daf-22* *exo*-metabolomes reveal extensive phosphorylation of ascarosides. Phosphorylated and corresponding nonphosphorylated ascaroside species are highlighted with red and blue shading, respectively. (e) EICs for synthetic phascr#151 (27), synthetic phascr#152 (28), and the phosphorylated ascr#15 in wild-type *exo*-metabolome indicates phosphorylation at the 2'-position. (f) Abundances of phosphorylated ascarosides relative to corresponding nonphosphorylated ascarosides in different life stages.

To estimate the concentrations of phosphorylated ascarosides relative to their nonphosphorylated counterparts, we established standard curves for ascr#15 (26) and phascr#151 (27) (Figure S13). Quantification across different life stages and different nutritional conditions revealed that production of phosphorylated ascarosides is greatly increased in L2, L3, and L4 larvae under starved conditions (Figure 5f). In fact, over 50% of some ascarosides, e.g., ascr#3 (2), were phosphorylated in starved larvae. Levels of ascaroside phosphorylation in adults do not appear to change significantly upon starvation; however, phosphorylated ascarosides still account for a significant fraction of total ascaroside production.

Although mass features we identified here as phosphorylated ascarosides have been previously shown to be *daf-22* dependent, their uninformative MS/MS spectra prevented their recognition as an abundant class of ascarosides.¹² As pointed out above, negative-ion MS/MS spectra of phosphorylated ascarosides are dominated by phosphate fragments and lack the *m/z* 73.0295 fragment that usually is diagnostic for ascarosides (Figure S14).⁴

Fucosylated Ascarosides. A third family of PDMs was identified from the largest reporter-ion cluster for which MS/MS spectra suggested addition of one or two deoxyhexose moieties (Figure 6a,b, and S15–S17). These deoxy sugar moieties appeared to be decorated with up to two methyl groups each, resulting in a total of 11 observed combinations of

methyated carbohydrate-modified PDMs (e.g., 36a–c, 37, also see Table S1).

Similar to the amino acid conjugate PDMs, candidate metabolites corresponding to the deoxyhexosylated PDMs were detected only at very low abundances in wild-type *C. elegans*, with intensities that were too low for extensive MS/MS analysis. However, inspection of the *exo*-metabolome of *acox-1.1* mutants revealed much larger amounts of putative deoxyhexosylated ascr#10 derivatives, likely because of the very high abundance of ascr#10 (34) in this mutant (Figure S8).⁴ MS/MS analysis indicated the presence of deoxyhexosylated ascr#10 with 0, 1, and 2 additional methyl groups (38–40, Figure 6c). Additionally, we detected several series of features that appeared to represent deoxyhexosylated derivatives of other ascarosides with different side chain lengths (Figures S18–S20).

On the basis of previous reports of mono- and di-O-methylated L-fucose as a component of *C. elegans* N-glycans,³⁵ we hypothesized that the deoxyhexose in the ascr#3-YNE-derived PDMs may be fucose. To confirm the identity of the deoxyhexose, the mixture of PDMs obtained from the ascr#3-YNE (3) enrichment experiment was subjected to acidic hydrolysis to cleave all glycosidic bonds and then permethylated.^{36,37} The permethylated sample showed features with the same retention times and molecular weight as a synthetic mixture of the α - and β -isomers of permethylated fucose,

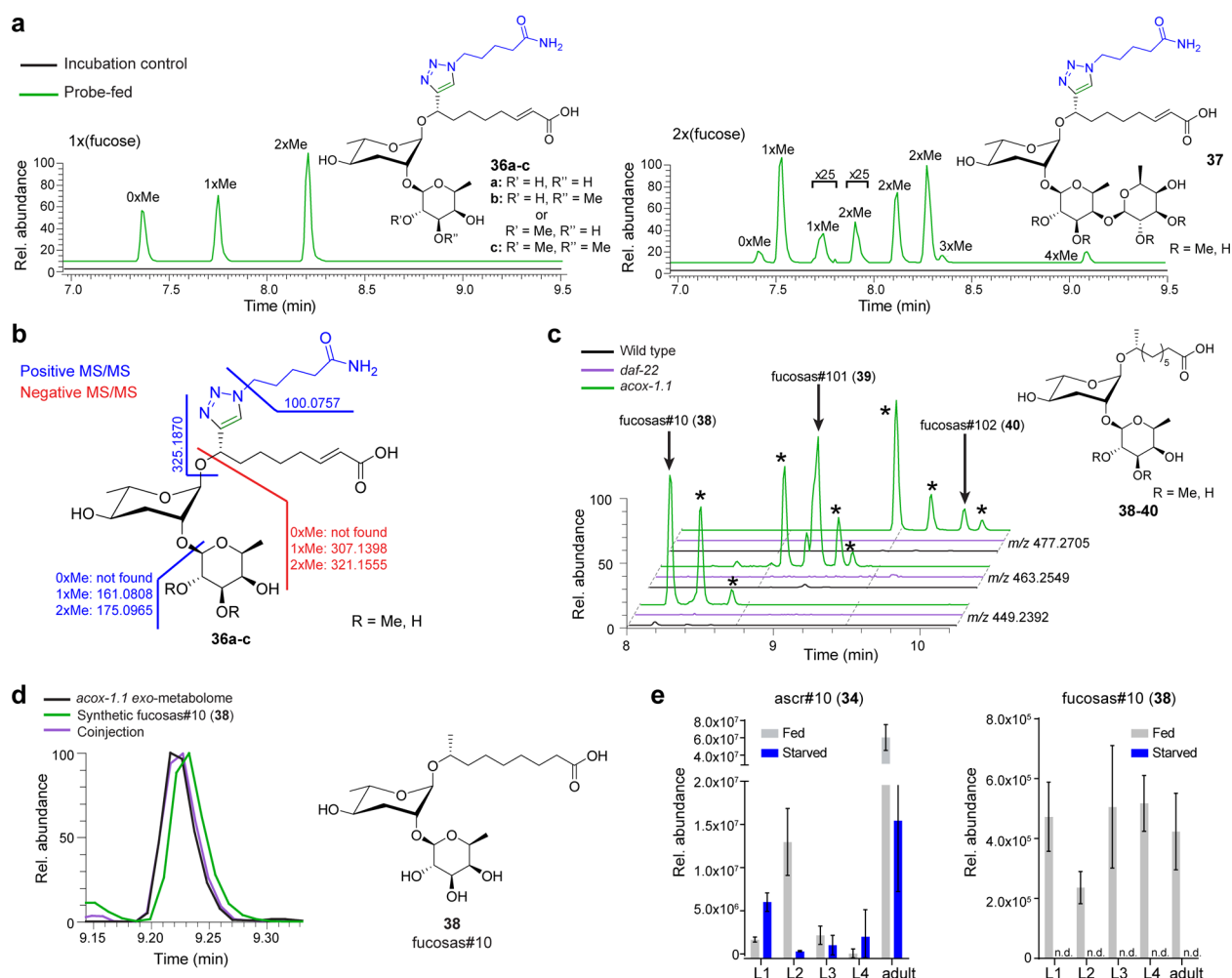


Figure 6. Identification of PDMs representing fucosylated ascarosides. (a) EICs for PDMs bearing one or two deoxyhexose moieties that are further decorated with additional methyl groups in enriched probe-fed and control data sets. (b) MS/MS analysis of fucosylated ascr#3-LNK decorated with up to two methyl groups (36a–c, also Figures S15–S17). (c) EICs for *m/z* values corresponding to fucosyl#10 (38) and derivatives with one or two additional methyl groups (fucosyl#101 (39) and fucosyl#102 (40), respectively) in *acox-1.1*, wildtype, and *daf-22* *exo*-metabolomes. Fucosylated ascr#10 derivatives are observed in wild-type *C. elegans*, and not detected in *daf-22* mutant worms. Additional isomers of fucosylated ascr#10 derivatives represent fucosyl-ascarosides with shorter side chains and mono- and di-*O*-methylation of the fucose, as suggested by MS/MS analysis (Figures S18–S20). (d) EICs for fucosyl#10 (38) in *acox-1.1* *exo*-metabolome (black), a synthetic sample (green), and in a sample where synthetic fucosyl#10 is added to *acox-1.1* *exo*-metabolome (purple) confirm identification of fucosyl#10. (e) Ascr#10 (34) and fucosyl#10 (38) were quantified in the *exo*-metabolome of wildtype worms. Production of fucosyl#10 is dependent on nutritional conditions, but not strongly dependent on life stage, unlike ascr#10.

supporting the assignment of the deoxyhexose as fucose (Figure S21). To confirm this assignment and to determine whether the fucose is attached to the 2'- or 4'-position of the ascarose, we synthesized monofucosylated ascr#10 (fucosyl#10, 38), one of the most abundant fucosylated ascarosides, using a thioglycosylation strategy (see SI Methods).³⁸ Fucosyl#10 (38) matched the retention time and MS/MS fragmentation pattern of the natural compound (Figures 6d and S22). Lastly, we quantified production of fucosylated ascarosides in the *exo*-metabolome in different life stages of wild-type *C. elegans*. Unlike ascr#10 (34), which is excreted in different amounts depending on life-stage, fucosyl#10 (38) is excreted consistently at all life stages (Figure 6e). Additionally, we found that starvation reduced production of fucosyl#10 to levels below the detection limit of LC-MS, suggesting that this compound may function in mediating starvation-associated metabolic responses (Figure 6e).

DIMEN in *E. coli* Bacteria. To test applicability of DIMEN to other biological systems, we performed a simple experiment in *E. coli* OP50 bacteria, feeding 17-octadecynoic acid (C18-YNE) as a probe, using the conditions we had developed for *C. elegans* without further optimization. Analysis of MS/MS data for the reporter-ion at *m/z* 100.0757 revealed a series of chain shortened PDMs, indicating extensive metabolism of C18-YNE via β -oxidation (Figure S23). In-depth analysis of the data obtained for C18-YNE-treated *E. coli* is beyond the scope of this study; however, the preliminary results suggest that DIMEN is broadly applicable.

DISCUSSION

Untargeted metabolic studies between different mutants, life stages, or other biological conditions result in large and often overwhelming data sets that nonetheless capture only part of the structural diversity of metabolomes, given the limitations of current MS instrumentation and data processing capabil-

ities.^{12,39–41} On the other hand, targeted studies, which often take advantage of reactive functional groups (i.e., acids, amines, thiols, etc.),^{42–44} may fail to discover novel metabolites with unexpected structural features. DIMEN enables discovery-oriented analyses in a highly sensitive, pathway-targeted manner by leveraging the advantages of both untargeted comparative metabolomics and enrichment strategies. As we demonstrated here, using a single alkyne-modified metabolite uncovered a large number of novel biosynthetic transformations that reveal unexpected intersection with amino acid metabolism, glycan chemistry, and phosphorylation.

Although click-chemistry mediated enrichment strategies have been used extensively for proteins, few studies employ click-chemistry-enabled enrichment of small molecules. Examples include a cleavable azide reactive resin (CARR) that was used to isolate azide-labeled fatty acids and phenylethylamines, as well as an azide-reactive cyclooctyne (ARCO) resin that was used to enrich azide-labeled peptides.^{8,9} In both cases, the strain-promoted azide–alkyne cycloaddition (SPAAC) click reaction was employed to enrich azide-labeled compounds.⁴⁵ These strategies have the advantage of not requiring any copper, reducing agent, or chelating ligands when compared to CuAAC. However, the requirement of a strained alkyne for enrichment results in relatively large and bulky linkers that may confer undesirable chromatographic or MS/MS properties. One of few examples of CuAAC to isolate small molecules employed an azido-coumarin loaded resin to detect alkyne-containing natural products.⁴⁶ The resulting fluorescence was then used to guide isolation of alkynated plant metabolites.

DIMEN's use of CuAAC with the ACER resin results in a relatively small and polar linker-derived modification of the enriched PDMs, which benefits ionization and limits chromatographic perturbation. However, since we observed some undesired TFA-induced cleavage of the glycosidic bond in enriched ascr#3-LNK (4), future iterations of this DIMEN could benefit from implementation of other established chemo-orthogonal cleavage strategies (Figure 2c).⁴²

Among the three newly identified compound classes, the amino acid conjugates reveal a direct connection between ascaroside signaling and amino acid abundance and thus nutritional status. Correspondingly, the previous identification of indole-carboxy- and *p*-hydroxybenzoyl-modified ascarosides (icas and hbas, respectively) connect ascaroside signaling to amino acid catabolism.¹¹ Among the identified ascaroside amino acid conjugates, the glycine derivatives are by far the most abundant (Figure 4c), although free glycine is among the less abundant amino acids in *C. elegans*.^{47,48} This suggests that, like other types of modular ascarosides, amino acid-conjugated ascarosides are products of specific biosynthetic pathways and may serve specific signaling functions.^{19,49} Under laboratory conditions, the amino acid-conjugated ascarosides are barely observable by LC-HRMS in the *exo*-metabolome of wildtype *C. elegans*, whereas they accumulate in the *acox-1.1* mutant (Figure 4b). Their identification highlights the potential of DIMEN for the discovery of low-abundance metabolites that may represent potent signaling molecules, even at low femtomolar concentrations, as shown for the case of hbas#3.⁴

In contrast, the phosphorylated ascarosides identified here are among the most abundant ascarosides in *C. elegans*, but had remained unidentified, despite the central role ascaroside signaling plays in the life history of this model organism. Although several compounds identified here as phosphorylated ascarosides had been observed previously via LC-MS based

comparative metabolomics, they remained unidentified as their ascaroside origin could not be confirmed (Figure S14).¹² DIMEN enabled structure elucidation by (i) demonstrating that these compounds must be derived from an ascaroside probe and (ii) providing robust fragmentation patterns that extended beyond phosphate fragments.

Small molecule phosphorylation regulates diverse biological processes, e.g., signal transduction through phosphorylated nucleosides, inositols, and lipid-linked inositides, and importantly, energy metabolism.^{50–52} Phosphorylation is often highly site-specific, e.g., in the case of inositol pyrophosphates.⁵³ In the case of ascarosides, phosphorylation is specific for the 2'-position. Our observation that starvation increases the percentage of excreted phosphorylated ascarosides relative to their nonphosphorylated counterparts, particularly during the larval stages of *C. elegans* (Figure 5f), suggests that phosphorylated ascarosides have distinct signaling properties.

Like the phosphorylated and amino acid conjugated PDMs, fucosylated ascarosides produced an MS/MS subcluster that enabled their quick recognition as a novel compound family (Figures 3a and 6a). Fucose is obtained by mammals from food sources via a salvage pathway and through a highly conserved 3-step transformation of GDP mannose.⁵⁴ Once acquired, fucose may be incorporated into *N*- and *O*-glycans or glycolipids through the activity of fucosyltransferases, with the most well-known example being the ABO antigens found on human blood cells.⁵⁴ Because fucose is integral for cell–cell communication and signaling, disturbances of fucosylation result in developmental disease and autoimmune disorders.⁵⁵ Fucosylation also plays an important role for chemical signaling involving bacteria, for example in host-commensal symbiosis in the human intestine.^{56,57} The *C. elegans* genome includes a partially annotated fucose biosynthetic pathway and over 30 predicted fucosyl transferases, indicating that fucose may play an important role in nematodes.^{58,59} Similar to other organisms, *C. elegans* incorporate fucose in the architecture of *N*- and *O*-glycans of glycoproteins. These fucose units can be *O*-methylated analogously to the enriched PDMs of this study.^{35,59,60} Surprisingly, there are few examples of fucosylated small molecules from other organisms, though our identification of fucosylated ascarosides and the prevalence of fucosyl transferases may suggest that fucosylated metabolites could be widespread.

In addition to the three compound classes we selected for follow up, the DIMEN MS/MS network (Figure 3a) contains many yet uncharacterized nodes that likely represent additional unknown structures and metabolic pathways. Using other alkyne-functionalized ascarosides—all data shown here were generated using a single probe—and application of DIMEN to biosynthesis mutants may reveal additional facets of ascaroside metabolism and help delineate biosynthetic pathways. Our partial analysis provides evidence for an extensive metabolic network beyond existing biochemical models in *C. elegans* and highlights the potential of click-chemistry-based enrichment and tagging approaches for uncovering hidden metabolites and biosyntheses in diverse biological systems.

■ ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/jacs.0c06877>.

DIMEN methods, synthetic procedures with NMR spectroscopic data, Supporting Figures S1–S24, and Tables S1–S7 (PDF)

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Notes

The authors declare no competing financial interest.

ACKNOWLEDGMENTS

We thank D. Kiemle (SUNY-ESF) for assistance with NMR spectroscopy, and I. O'Doherty, D. Gentile, and R. Micikas (Cornell University) for synthetic starting materials. This work was partly supported by the NIH (T32 GM008500 to J.S.H. and J.A.B., and R01 GM088290 and R35 GM131877 to F.C.S.) and the Howard Hughes Medical Institute, with which F.C.S. is a Faculty Scholar.

ABBREVIATIONS

ARCO, azide-reactive cyclooctyne; BTAA, 2-(4-((Bis((1-tert-butyl)-1H-1,2,3-triazol-4-yl)methylamino)methyl)-1H-1,2,3-triazol-1-yl)acetic acid; CARR, cleavable azide-reactive resin; CuAAC, copper catalyzed azide alkyne cycloaddition; DIMEN, deep interrogation of metabolism via enrichment;

GDP, guanosine diphosphate; LC, liquid chromatography; *m/z*, mass to charge ratio; MS, mass spectrometry; MS/MS, tandem mass spectrometry; PDM, probe-derived metabolite; PEGA, polyethylene glycol acrylamide; SPPS, solid phase peptide synthesis.

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