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A Chemical Proteomic Strategy Reveals Inhibitors of Lipoate Salvage in Bacteria and Parasites

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Abstract: The development of novel anti-infectives requires unprecedented strategies targeting pathways which are solely present in pathogens but absent in humans. Following this principle, we developed inhibitors of lipoic acid (LA) salvage, a crucial pathway for the survival of LA auxotrophic bacteria and parasites but non-essential in human cells. An LA-based probe was selectively transferred onto substrate proteins via lipoate protein ligase (LPL) in intact cells, and their binding sites were determined by mass spectrometry. Probe labeling served as a proxy of LPL activity, enabling in situ screenings for cell-permeable LPL inhibitors. Profiling a focused compound library revealed two substrate analogs (LAME and C3) as inhibitors, which were further validated by binding studies and co-crystallography. Importantly, LAME exhibited low toxicity in human cells and achieved killing of *Plasmodium falciparum* in erythrocytes with an EC₅₀ value of 15 μM, making it the most effective LPL inhibitor reported to date.

Introduction

Cofactors enhance the scope of chemical transformations in enzymes beyond the limited diversity provided by the 20 canonical amino acids.^[1] They play crucial roles in many cellular processes and are thus attractive drug targets.^[2] Recently, several chemical proteomic strategies utilized small molecule cofactor mimics, e.g. pyridoxal phosphate^[3] and heme,^[4] to infiltrate the cellular metabolic machinery and facilitate their incorporation in cofactor-dependent enzymes. A small bioorthogonal handle (alkyne or azide) is clicked to an affinity tag (e.g., biotin) for enrichment on avidin beads and subsequent identification by mass spectrometry (MS).^[5] In addition to the discovery of previously unknown cofactor-dependent enzymes, the methodology identified inhibitors against selected targets via competitive profiling.^[6]

Lipoic acid (LA) is a promising candidate for this strategy as it contributes to central metabolism in various organisms.^[7] LA is covalently bound to particular lysine residues where it catalyzes reactions such as acyl-CoA formation in dehydrogenase complexes, e.g., pyruvate dehydrogenase (oxidative decarboxylation).^[8] Bacteria and parasites exhibit a highly diverse machinery to synthesize or acquire the cofactor compared to humans. In *Escherichia coli*, the biosynthesis pathway consists of the enzymes LipB (octanoyl transferase) and LipA (lipoyl synthase). These enzymes assemble the LA cofactor directly ('on-site') on specific lysine residues of conserved domains of LA-dependent proteins via octanoylation (LipB) and subsequent introduction of the redox-active dithiolane moiety (LipA, Figure 1A).^[7a,b,d] In addition, salvage of LA is an alternate strategy by which organisms lipoylate proteins with the scavenged cofactor from the exterior environment (Figure 1A). This process is catalyzed by ATP-dependent lipoate protein ligases (LPL).^[7a,b,d,9] By contrast, no homologous LA salvage pathway is known in mammalian cells, which instead rely on LA biosynthesis in the mitochondrion.^[7a,d,10] Some bacteria and parasites, such as *Listeria monocytogenes*^[11] and *Plasmodium* spp.,^[12] respectively, are LA auxotroph organisms, i.e., they rely on the uptake and salvage of LA for survival (in case of *P. falciparum* and *P. berghei* into the mitochondrion). Thus, these organisms represent a unique opportunity for the development of selective LPL inhibitors in order to block LA-dependent central metabolism pathways and, thereby, growth.^[13] Despite this intriguing opportunity and the urgent need for novel anti-infectives, the development of such inhibitors has not been significantly

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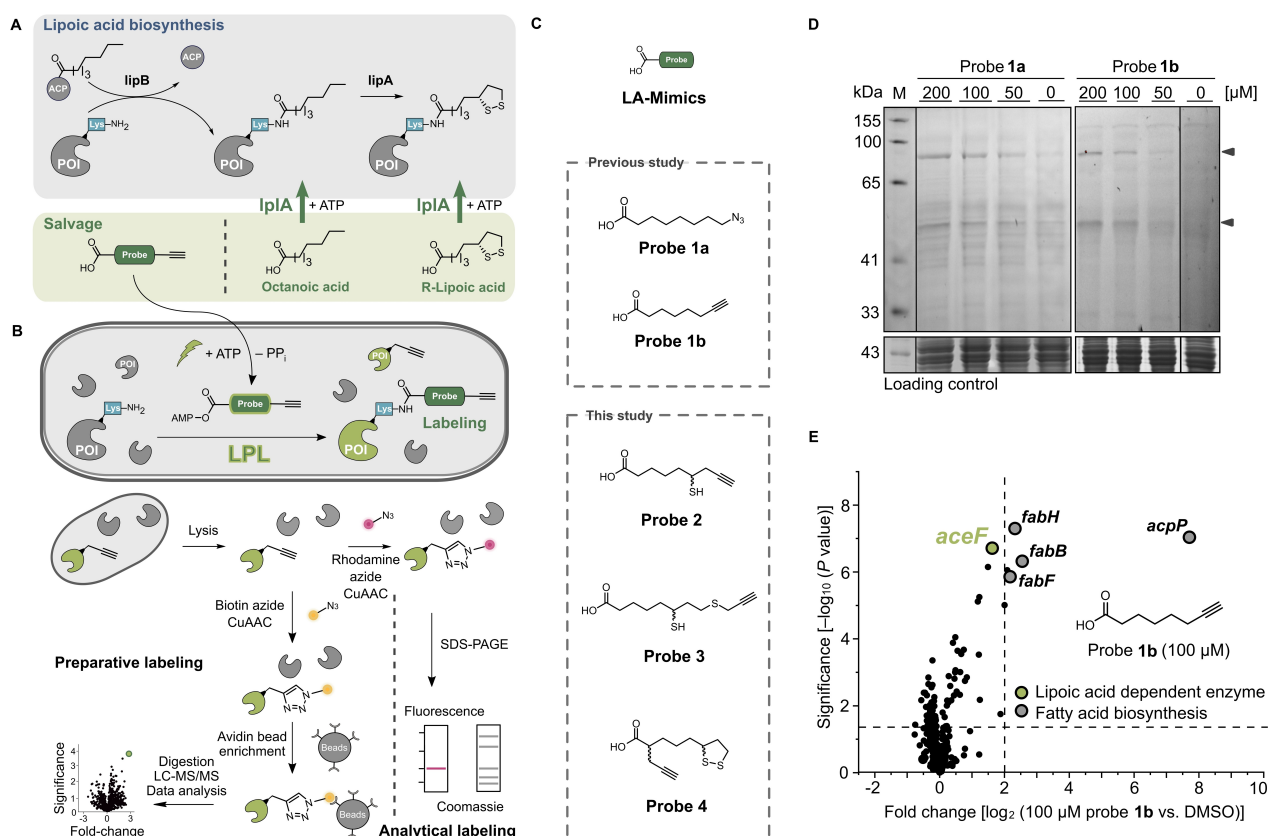


Figure 1. In situ chemical proteomic strategy to investigate lipoylation via LPL-mediated activation and transfer. A) The biosynthesis (grey box) and salvage of LA (green box) via the LPL enzyme *lplA* in *E. coli*. ACP—acyl carrier protein, POI—protein of interest. B) Intracellular probe-based infiltration strategy via ATP-dependent LPL as LA salvage enzyme with subsequent downstream analysis of labeled proteins by click-chemistry (CuAAC—copper-catalyzed azide-alkyne cycloaddition) using either gel-based rhodamine fluorescence readout by SDS-PAGE (magenta, analytical labeling) or mass spectrometry-based detection after biotin conjugation and enrichment on avidin beads (yellow, preparative labeling). C) Probe structures as LA salvage mimics used in this study. D) Analytical labeling of intact *E. coli* $\Delta lipB$ cells with probes **1a** and **1b** (2 h) in stationary phase revealed a similar pattern of concentration-dependently labeled fluorescent bands (grey arrows). The procedure with the azide-bearing C8-fatty acid probe **1a** showed higher background labeling than the alkyne-based derivative **1b**.^[20] “M” represents a marker as a protein size reference. Coomassie staining served as loading control. E) The volcano plot shows enrichment of mainly fatty acid biosynthesis proteins (grey) on a \log_2 scale after treatment of intact *E. coli* $\Delta lipB$ cells in stationary phase with the alkyne-bearing probe **1b** (100 μ M, 2 h). Of note, the LA-dependent protein product of the gene *aceF* (green) could be 3-fold enriched. The threshold lines indicate 4-fold enrichment compared to DMSO and a $-\log_{10}$ P value of 1.3 (two-sided two-sample *t*-test, $n = 4$ independent experiments per group). Gene names of proteins are shown in italic.

exploited. One example is the redox-inactive small molecule mimic of LA, 8-bromo octanoic acid (**BrO**), which is transferred via LPL to substrate proteins, but cannot further support enzymatic catalysis.^[14] **BrO** was shown to reduce growth of *P. falciparum* in erythrocytes,^[12a] *P. berghei* in HepG2 cells^[12b] and mice,^[15] and *Toxoplasma gondii* in fibroblasts,^[16] but high concentrations up to the millimolar range were needed for parasite killing.^[12,15] Thus, novel chemical tools are required to decipher more potent LPL inhibitors. Recently, two methods have been introduced to chemically modify endogenous lipoylated proteins exclusively in lysates via iodoacetamide-assisted lipoate-cyclo-octyne ligation (iLCL)^[17] or reduction of the dithiolane ring followed by thioacetalization with an alkynylated aldehyde.^[18] These methods enriched lipoylated proteins in lysates and determined their lipoylation sites on lysines. However, because they do not involve the transfer of LA via

LPL catalysis in intact cells, they are unsuitable for direct inhibitor discovery via competitive profiling.

Here, we introduce novel LA probes, which infiltrate the cellular salvage pathway in intact cells and get incorporated in cognate lipoate-dependent proteins (Figure 1B).

Chemical proteomics of intact *E. coli* and *L. monocytogenes* cells show selective enrichment of LA-dependent proteins and resolution of their modified lysine sites. Computational design, in situ profiling, and co-crystallization revealed two effective LPL inhibitors (**C3** and **LAMe**). These compounds killed *P. falciparum* in erythrocytes with EC₅₀ values of 27 μ M and 15 μ M, respectively, highlighting the value of this approach in uncovering novel anti-infectives.

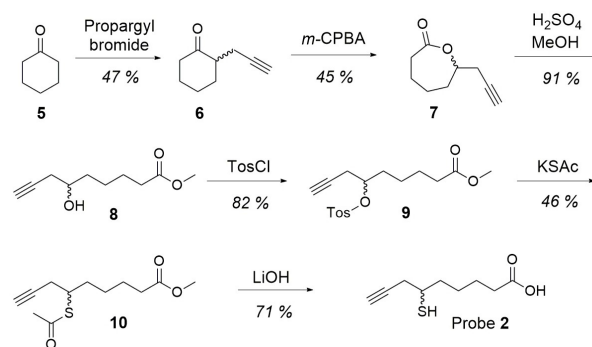
Results and Discussion

Design and Synthesis of Probes to Monitor Protein Lipoylation

As benchmarks, we started our studies with the C8-fatty acid probes **1a** (8-azido octanoic acid) and **1b** (7-octynoic acid) as previously confirmed LPL substrates (Figure 1C).^[19] First, we determined their toxicity in *E. coli* and identified up to 50 % growth reduction with probe **1a**, while probe **1b** showed no effect (Figure S1A). Next, wild-type *E. coli* and $\Delta lipB$ cells, bearing a mutation in the de novo biosynthesis pathway to maximize probe uptake, were treated with various concentrations of the non-toxic probe **1b** for 2 h in the stationary and exponential phase. Cells were lysed and click-chemistry was used to link any proteins modified by probe **1b** to rhodamine azide for fluorescent SDS-PAGE analysis (Figure 1B). Interestingly, a distinct labeling pattern was only observed in the stationary phase of $\Delta lipB$ cells at a probe concentration of 50 μM or higher (Figure 1D, S1B). A similar fluorescence pattern with higher background labeling was observed for the azide-bearing probe **1a** after linking modified proteins to rhodamine alkyne by click-chemistry.^[20] From these results, we selected optimal conditions to perform quantitative MS analysis of covalently-labeled proteins. Therefore, we treated stationary cells for 2 h with 100 μM of the non-toxic probe **1b**, followed by lysis, click to biotin azide, enrichment, digestion, and analysis via LC-MS/MS (Figure 1B). Notably, one of the three LA-dependent proteins known in *E. coli*, ODP2 (*aceF*), was 3-fold enriched (Figure 1E).^[17,18] Moreover, numerous proteins belonging to fatty acid biosynthesis were among the top hits [$-\log_{10}$ (*P* value) 6–8, \log_2 (fold-change) 2–8]. These initial results demonstrated the infiltration of bacterial LA metabolism. However, the relatively simple probe design prevented the desired specificity for LA-dependent proteins with fatty acid biosynthesis enzymes as predominant off-targets.

In order to enhance the fidelity in detection of enzyme lipoylation, we designed a suite of LA mimics as probes that are attainable by chemical synthesis (Figure 1C). Starting from probe **1b**, we incrementally increased the structural complexity to more closely mimic LA. In a first step, we placed a free thiol at position 6 of 8-nonynoic acid to obtain probe **2**. The synthesis started with propargylating cyclohexanone (**5**), followed by a Baeyer–Villiger oxidation as a key step to introduce the lactone moiety (Scheme 1).^[21] After ring opening by transesterification, the hydroxyl group at position 6 could be transformed into thioester **10** by tosylation and substitution with thioacetate. Saponification as final step generated probe **2** as a racemic mixture in six steps (5 % overall yield).

Next, the propargyl tag was linked to the thiol at position 8 of the reduced LA moiety as a thioether (probe **3**). Probe **4** presents the most similar mimic to LA after functionalization at position 2 with a propargyl handle. For probes **3** and **4**, LA was protected by esterification, followed by reduction to open the dithiolane ring. To obtain probe **3**, the thiol group at position 8 was selectively propargylated, followed by saponification to liberate the carboxylic acid as racemate. By contrast, for probe **4**, both thiols were



Scheme 1. Synthesis of probe **2**. The propargylated lactone **7** was synthesized as previously reported^[21] prior to transesterification (**8**), tosylation (**9**), and nucleophilic substitution by thioacetate. Global saponification of **10** yielded probe **2** as racemate.

protected by tritylation to install a propargyl tag at position 2, followed by global deprotection and oxidation to yield the desired probe as a mixture of all diastereoisomers (Scheme S1).

Probe 2 Labels LA-dependent Proteins in *E. coli* and *L. monocytogenes*

With a panel of probes (**2–4**) in hand, we first tested their impact on bacterial growth under labeling conditions (Figure S2A). At high concentrations of 500 μM , only probe **2** slightly reduced the growth of *E. coli* $\Delta lipB$ by 20%. Secondly, we examined their labeling performance in intact *E. coli* $\Delta lipB$ cells in both fluorescent gel-based and LC-MS/MS experiments. Labeling with probes **4** and **3** revealed high background without competition upon LA treatment (Figure S2B, C). However, probe **2** showed distinct bands on the fluorescent gel that could be competed with LA and BrO pre-treatment, indicating that LA-dependent proteins were labeled via LPL (Figure 2A). Additionally, application of probe **2** in wild-type *E. coli* cells showed no labeling indicating that it cannot compete with the de novo biosynthesis (Figure S2D). Subsequent quantitative LC-MS/MS profiling exclusively revealed all three known LA-dependent enzymes in the *E. coli* $\Delta lipB$ cells (ODP2—*aceF*, 66 kDa; ODO2—*sucB*, 44 kDa; GCSH—*gcvH*, 14 kDa) without any background (Figure S3A).^[17,18] Moreover, labeling was competed by addition of LA or BrO in 10-fold excess, demonstrating improved specificity of probe **2** (Figure 2B, S3B).

To rationalize the diverging labeling properties, we applied computational methods to predict how LA and the probe stereoisomers bind into the crystal structure of *E. coli* LPL (lplA—*lplA*, PDBID 1x2h).^[22] The structures were docked into the active site (Figure S4A) and subjected to molecular dynamics (MD) simulations to investigate their binding behavior, including binding free energies ($\Delta\Delta G_{\text{Bind}}$), root-mean-square deviation (RMSD_{Lig}), and root-mean-square fluctuation (RMSF_{Lig}). Different binding modes were observed across these compounds. The *S*-enantiomer of

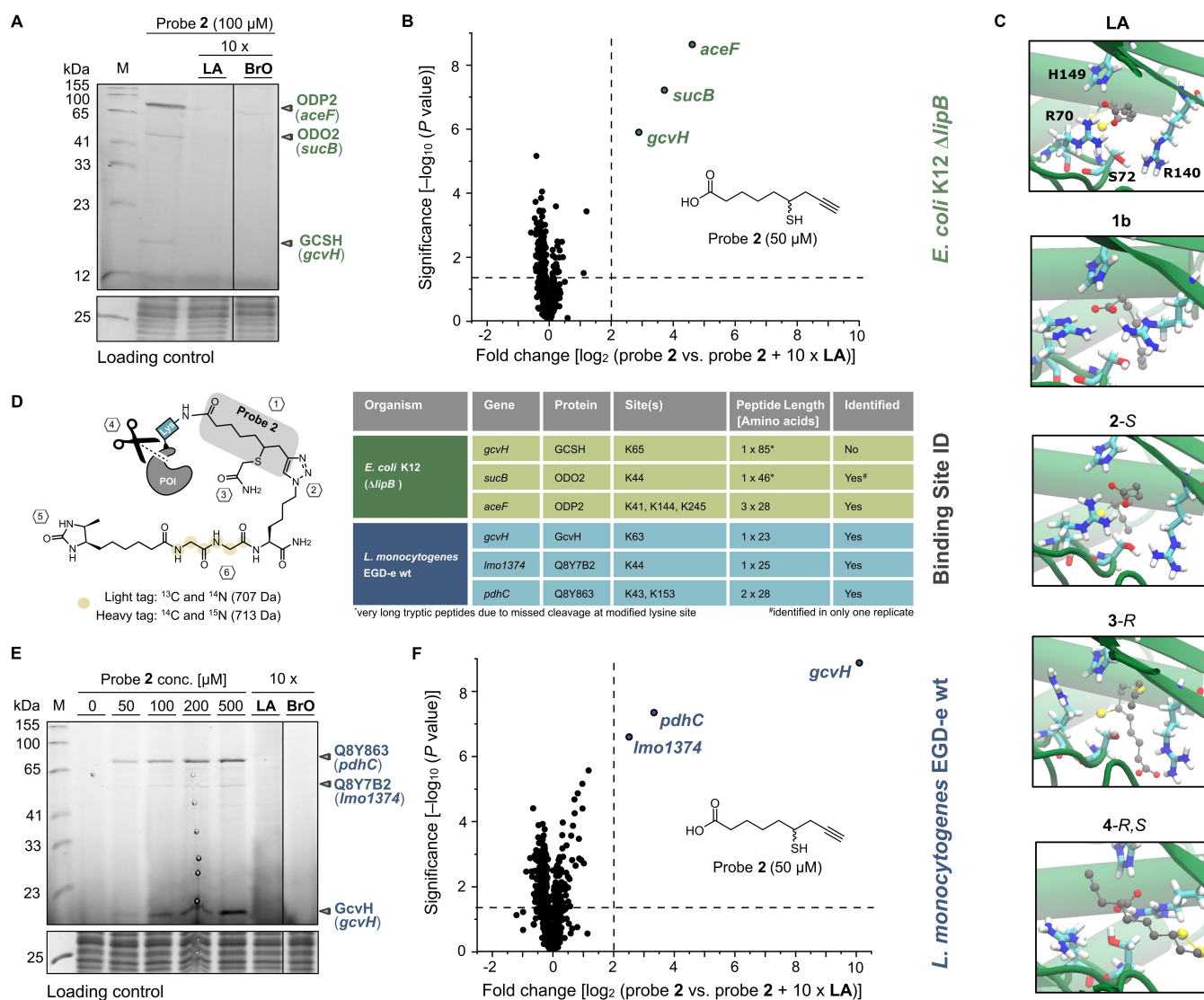


Figure 2. In situ profiling of lipoylation in *E. coli* $\Delta lipB$ (Gram-negative) and *L. monocytogenes* wild type (Gram-positive) with binding site identification. Gene names of corresponding protein products are presented in *italic*. “M” represents a marker as a protein size reference, and Coomassie staining served as loading control during analytical labeling. Of note, the protein products might show reduced migration on SDS-PAGE after rhodamine modification. The threshold lines in volcano plots indicate a $-\log_{10} P$ value of 1.3 (two-sided two-sample *t*-test, $n = 4$ independent experiments per group) and 4-fold enrichment. A) Analytical labeling of intact *E. coli* $\Delta lipB$ cells with probe 2 (100 μM , 2 h) and in competition with 10-fold excess (1 mM) of LA or BrO indicated infiltration of LA salvage via LPL. The green arrows mark the expected size of ODP2 (*aceF*, 66 kDa), ODO2 (*sucB*, 44 kDa), and GCSH (*gcvH*, 14 kDa) as known lipoylated proteins.^[17,18] B) The volcano plot presents preparative labeling results of intact *E. coli* $\Delta lipB$ cells with probe 2 (50 μM , 2 h) in comparison to probe 2 (50 μM) in competition with 10-fold excess LA (500 μM), indicating selective enrichment of LA-dependent enzymes.^[17,18] C) Representative snapshots of the native R-enantiomer of LA and computationally predicted best-performing probe candidates 1b, 2-S, 3-R and 4-R,S in *E. coli* lipA (PDBID 1x2h) are depicted at the end of molecular dynamics simulations (400 ns), highlighting comparable binding orientations of LA and 2-S in contrast to 1b, 3 and 4 (Figure S4B–D). D) The methodology of isoDTB tags^[23] was applied to identify the binding sites of probe 2 on lysine residues in *E. coli* $\Delta lipB$ and *L. monocytogenes* that corresponded to protein lipoylation sites. After labeling respective lysines in intact bacterial cells with probe 2 (1), the labeled proteome was clicked to a heavy and a light isoDTB tag as a technical replicate (2), followed by capping of free thiols with iodoacetamide (3). Proteins were digested by trypsin (4) to release the modified desthiobiotinylated peptides with one missed cleavage at the modified lysine. Tryptic peptides were enriched on avidin beads (5) and eluted to be identified by mass spectrometry. Mass shifts of 707 Da and 713 Da revealed isoDTB-tagged peptides (6). Peptides up to 28 amino acids were detected without problems in contrast to longer tryptic peptides with missed cleavage at the modified lysine, as indicated by asterisks.^[18] Peptides that were only identified in one replicate were marked with #. E) Concentration-dependent analytical labeling of intact *L. monocytogenes* wild type cells with probe 2 (0 μM –500 μM , 2 h) and competition with 10-fold excess (2 mM) of LA or BrO compared to probe 2 (200 μM) revealed infiltration of LA salvage via LPL and lipoyl transferase LipL. The blue arrows mark the expected size of Q8Y863 (*pdhC*, 58 kDa), Q8Y7B2 (*lmo1374*, 45 kDa), and GcvH (*gcvH*, 14 kDa) as known LA-dependent proteins.^[24] F) The volcano plot presents preparative labeling results of intact *L. monocytogenes* wild type cells with probe 2 (50 μM , 2 h) in comparison to probe 2 (50 μM) in competition with 10-fold excess LA (500 μM), indicating selective enrichment of LA-dependent enzymes.^[24]

probe **2** exhibited similar binding stability as **LA**, shown by RMSD_{Lig} and RMSF_{Lig} (Figure S4B), and both compounds remain in a binding pose forming hydrogen bonds with R70, H149, and S72 (Figure 2C, S4C). By contrast, probe **1b** and enantiomers and diastereomers of probes **3** and **4**, respectively, were occasionally attracted by neighboring R140 and deviated from their initial binding locations (Figure 2C, S4B–D), which may prevent their activation by ATP for subsequent transfer to **LA**-dependent proteins (Figure 1B).

With probe **2** as ideal lipoylation tool in hand, we deciphered the modified lysines in whole *E. coli* proteomes. Enrichment of the labeled peptides via tailored desthiobiotin (isoDTB) linkers^[23] and subsequent MS analysis revealed K44 of ODO2 (*sucB*) and K41/K144/K245 of ODP2 (*aceF*) as modified lipoylation sites, which is in line with literature data (Figure 2D).^[17,18]

The high fidelity of our method was also retained when we switched from the Gram-negative *E. coli* mutant to the labeling of a wild-type Gram-positive *L. monocytogenes* bacterial strain, which relies solely on an **LA** salvage pathway consisting of two LPL orthologs (lplA1–*lmo0931*; lplA2–*lmo0764*) to lipoylate GcvH. In a next step, the lipoyl group is transferred by LipL from GcvH to other **LA**-dependent proteins.^[9] Again, direct labeling of intact cells with probe **2** and competition with **LA** or **BrO** revealed three hits, comprising all known **LA**-dependent enzymes in *Listeria* (Q8Y863–*pdhC*, 58 kDa; Q8Y7B2–*lmo1374*, 45 kDa; GcvH–*gcvH*, 14 kDa; Figure 2E, F, S3C, D).^[24] Additionally, the corresponding modified lysines were identified via the isoDTB technology (Figure 2D).^[23] Of note, the binding sites—K63 of GcvH (*gcvH*), K44 of Q8Y7B2 (*lmo1374*) and K43/K153 of Q8Y863 (*pdhC*)—have been determined experimentally for the first time in *Listeria* and match theoretical predictions.^[24]

Computational Design and Screen of Novel Active Site LPL Inhibitors

The versatility of our novel labeling technology for different strains and its selectivity for the detection of lipoylated proteins represents a unique opportunity for identifying chemical compounds that lead to the manipulation of cellular lipoylation and putative therapeutic applications. Specifically, probe **2** is activated and transferred onto the cognate substrate proteins by LPL, facilitating a direct readout of its activity. The lack of an LPL-mediated salvage pathway in human cells makes this enzyme attractive for selective pathogen targeting.^[9,10,12a] Since there are no active site LPL inhibitors reported, we performed computational studies to design a suite of potential candidates for subsequent validation via our competitive profiling approach using probe **2**. We initiated the inhibitor development by a structural overlay of three representative LPL enzymes, lplA from *E. coli* (Gram-negative bacterium) and LipL1 (*lipL1*) from *P. falciparum* (parasite), for which crystal structures are available,^[22,25] and lplA1 from *L. monocytogenes* (Gram-positive bacterium) with a structural AlphaFold2 (AF2) prediction.^[26] The overlay of all

enzymes revealed a high structural similarity of the active sites as indicated by particular lysine and histidine residues (Figure S5), suggesting that inhibitors may be effective across different organisms. Three ligand binding pockets (A, B, and C, Figure S5) could be identified depending on the enzyme's catalytic state. The comparison between the **LA** bound crystal structure (*E. coli*, PDBID 1x2h, loading state)^[22] and the lipoyl-AMP (**LAQ**) bound structure (*E. coli*, PDBID 3a7r, intermediate state)^[25a] indicates that **LA** binds first to the initial loading pocket A (as demonstrated for the probe simulations, Figure S4) and is subsequently found in pocket B after adenylation by ATP, while the adenine motif of **LAQ** resides in pocket C. In the search for LPL active site inhibitors, we followed a two-tiered strategy by targeting pockets B and C. In the first approach, we virtually screened the ZINC database^[27] by a pharmacophore-guided^[28] search against the LPL active site, revealing 126 putative hits which were subsequently docked into LPL enzymes. The top 50 compounds of each ligase, selected by their docking score, were screened for overlapping compounds. The resulting 29 molecules were sorted by their averaged ranking position to identify ten commercially available compounds with the best-predicted performance (Figure S6A, B).

In a second approach, we made use of the similar catalytic activity between LPL and biotin protein ligase (BPL), for which a potent BPL inhibitor with antibacterial activity was previously reported.^[29,30] This molecule, mimicking biotin-AMP as native substrate, is composed of the signature biotin moiety linked via a triazole linker to an adenine heterocycle group (Figure 3A). Both biotin and adenine bind to distinct pockets of BPL, of which the latter adenine pocket is also present in LPL (pocket C, Figure S5). We thus tailored the molecule towards LPL to mimic **LAQ** by replacing the biotin with **LA** and the triazole with a more flexible but chemically stable amide linker while preserving the same linker length (**C3**, Figure 3A). **C3** and its cropped version, lipoic acid monomethyl amide (**LAMe**, Figure 3A), were docked into the active sites of LPL from *L. monocytogenes* and *P. falciparum* in comparison to the native ligand **LA** and **BrO** (Figure 3B, S7A) to calculate and compare their binding affinities ($\Delta\Delta G_{\text{bind}}$, Figure 3C) and stabilities (RMSD_{Lig} , RMSF_{Lig} , Figure S7B) in silico by MD simulations. Overall, the same trend of predicted binding affinities for the compounds was observed in both enzymes as follows: **C3** > **LA** \approx **LAMe** = **BrO**. Interestingly, **LA**, **LAMe**, and **BrO** were predicted to bind stronger and more stably to pocket B of *L. monocytogenes* lplA1 (approximately $-32 \text{ kcal mol}^{-1}$) compared to *P. falciparum* LipL1 (roughly $-26 \text{ kcal mol}^{-1}$), whereas **C3** showed the strongest binding free energy with around $-38 \text{ kcal mol}^{-1}$ for both ligases (Figure 3C). Remarkably, the adenine motif of **C3** in pocket C of LipL1 contributed significantly to the binding free energy ($-12 \text{ kcal mol}^{-1}$), compared to the truncated **LAMe** solely residing in pocket B. As expected, this finding underlines a significant affinity enhancement for the introduction of the adenine-linker group in addressing both pockets. Based on these promising calculations, we synthe-

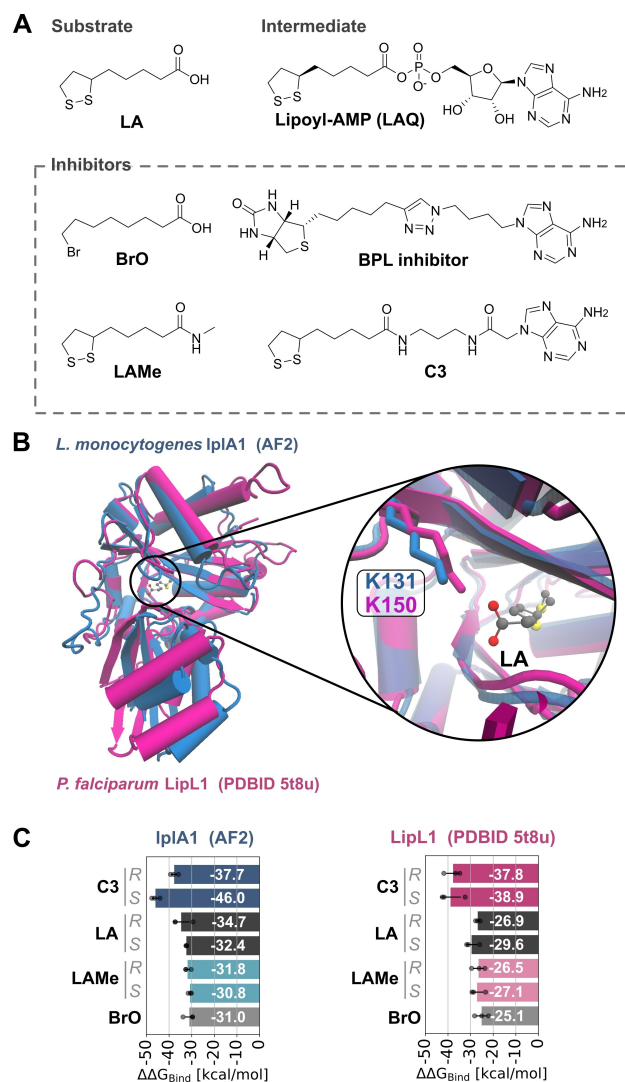


Figure 3. Docking and MD simulations of rational-designed potential inhibitors of LPL in *L. monocytogenes* (blue) and *P. falciparum* (magenta) as substrate and intermediate mimics. A) The inspiration for compound design was driven by mimicking the intermediate lipoyl-AMP (LAQ) with chemically stable amide bonds in analogy to a previously reported inhibitor of the biotin protein ligase (BPL).^[29] The simple LAMe molecule carries a monomethyl amide function that would not allow the LA moiety to be activated by ATP. C3 presents an extended version with an additional amide linker-adenine moiety. BrO was described as an inhibitor of cognate lipoylation sites as it is activated and transferred by LPL.^[14] LA as the native substrate and BrO as an established inhibitor serve as benchmarks. B) LA and all stereoisomers of potential inhibitors were docked into the active sites of *L. monocytogenes* lplA1 (AF2), and *P. falciparum* LipL1 (PDBID 5t8u), and representative binding modes of R-enantiomers are depicted exemplarily (Figure S7A). C) MD simulations in *L. monocytogenes* and *P. falciparum* predict all stereoisomers' LPL binding affinities ($\Delta\Delta G_{\text{bind}}$, mean value in white). The calculations were run in $n=3$ independent attempts, and error bars represent SEM.

sized C3 and LAMe by amide coupling for experimental comparison to BrO and LA (Scheme S2).

Biochemical and Crystallographic Studies Reveal C3 and LAMe as Active Site Inhibitors of LPL in Bacteria and Parasites

With a suite of twelve potential LPL inhibitors at hand, we tested their performance via competitive profiling with probe 2. For this, we focused on *L. monocytogenes* as a representative example of an LA-auxotrophic wild-type strain and treated cells with a 10-fold excess of competitor (1 mM) compared to the amount of probe 2 (100 μM) and analyzed the band intensity of signature protein Q8Y863 (*pdhC*, Figure 4A) via fluorescent SDS-PAGE analysis. While the ten compounds (16–25) derived from the virtual ZINC database screen largely lacked significant competition with the probe, the rationally designed C3 and LAMe compounds turned out to be potent with a gel-based EC_{50} value of about 210 nM and 330 nM, respectively (calculated for the residual in-gel modification of Q8Y863–*pdhC*; Figure 4B, S8A). This value is in the same range as competition with LA (100 nM) and 10-fold more potent than competition with the substrate analog BrO (2.3 μM). In addition, to verify that C3 is an LPL inhibitor and not metabolized to release LA after possible enzymatic amide cleavage within the cell, we demonstrated its stability in lysate over time with an HPLC-MS-based assay (Figure S8B). In line with the lack of LPL enzymes in humans,^[10a] MTT studies in HeLa cells indicate low to moderate toxicity for LAMe and C3, respectively (Figure S9A).

Accordingly, labeling with the non-toxic probe 2 (Figure S9A) in mammalian HeLa cells revealed no characteristic hits (Figure S9B). We independently confirmed a high affinity for C3 and LAMe with K_{D} values of 3.2 μM and 15 μM , respectively, which is again similar to the value of LA ($K_{\text{D}}=1.9$ μM). By contrast, the affinity of BrO was very low, and hence, binding could not be quantified (Figure 4C, S10A).

Based on these results, we aimed to better rationalize the binding mode of C3 in *L. monocytogenes* lplA1 and performed x-ray crystallography. So far, no experimental structural information of *L. monocytogenes* lplA1 is available, and we thus attempted to co-crystallize LAQ as the native intermediate. We noted that the C-terminus is flipped in the lplA1:LAQ complex (PDBID 8crj) in comparison to the predicted apo-AF2 structure (Figure S11A, B), which is in agreement with a previously co-crystallized homologous enzyme.^[25a] However, the AF2 structure perfectly aligns with the co-crystals of C3 (PDBID 8cr1) and LA (PDBID 8cr1), which confirms previous computational analysis (Figure S11B). Of note, we were able to co-crystallize LA, LAQ, and C3 solely as R-enantiomers and demonstrate inhibitor binding in the active pocket as a chemically stable LAQ analog (Figure 4D). Interestingly, C3 as an LAQ mimic could not induce the flip of the C-terminal domain (Figure S11B) as, e.g., crucial interactions with the phosphate and Lys131 are lacking, which offers a potential avenue for inhibitor refinement (Figure 4D).

Due to the structurally conserved binding pocket of *L. monocytogenes* lplA1 and *P. falciparum* LipL1 (Figure 3B, S5), we performed ITC studies with the recombinant

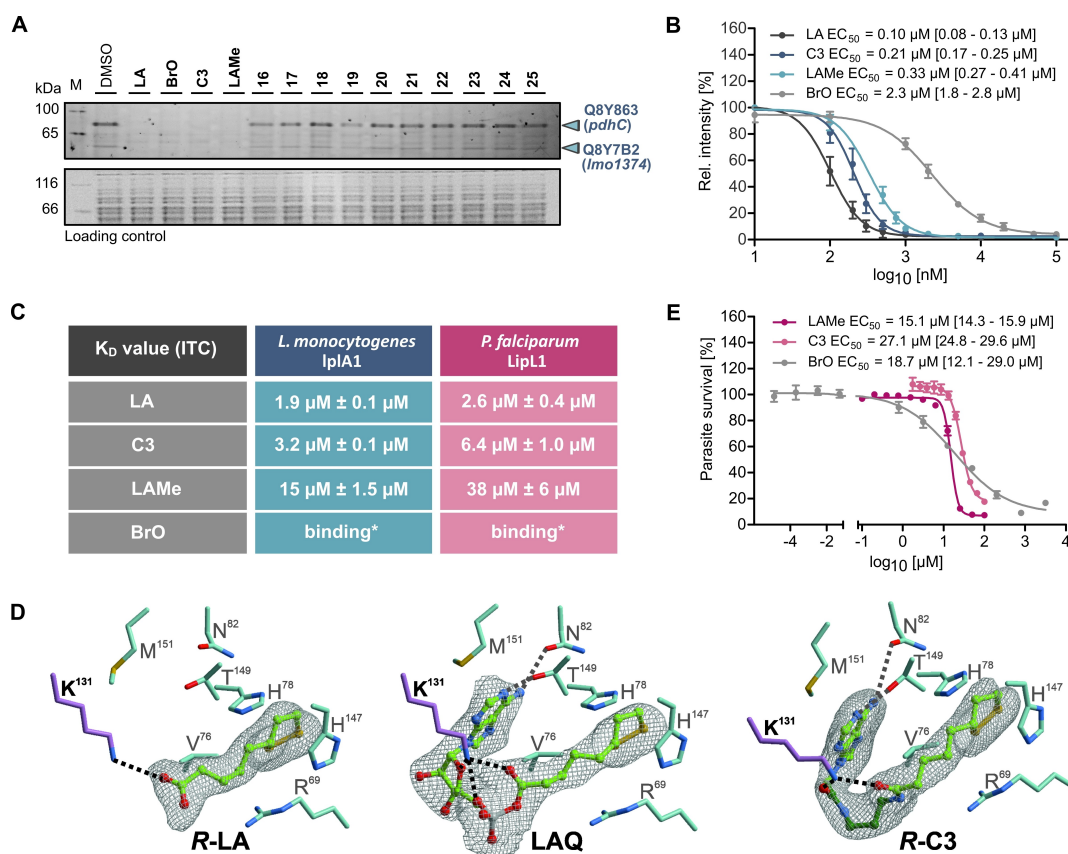


Figure 4. Identification and validation of LPL inhibitors in *L. monocytogenes* (blue) and *P. falciparum* (magenta). A) A competitive screen of the 12 selected potential inhibitors in 10-fold excess (1 mM) compared to probe **2** (100 μM, 2 h) was performed in intact *L. monocytogenes* cells to identify cell-permeable LPL inhibitors by measuring the decreased fluorescent signals of reporter proteins. SDS-PAGE with fluorescent analysis was performed, and blue arrows indicate reporter proteins Q8Y863 (*pdhC*) and Q8Y7B2 (*lmo1374*). Coomassie stain served as loading control, and “M” indicates a protein marker. B) EC₅₀ value determination of corresponding gel-based in situ competitive labeling studies in *L. monocytogenes* cells with identified inhibitors **C3** and **LAMe** versus probe **2** (100 μM, 2 h). Fluorescent signal intensities for Q8Y863 (Figure S8A) were analyzed by ImageJ. The analysis, plotted with mean values and error bars indicating SEM, was performed in *n* = 3 independent experiments, and competition with **LA** and **BrO** served as benchmarks. The error range is shown as 95% confidence interval. C) Recombinant *L. monocytogenes* lplA1 and *P. falciparum* LipL1 proteins were analyzed by ITC to separately verify the binding trends of inhibitors **C3** and **LAMe** towards LPL compared to **LA** and **BrO**. The experiments were measured at least in duplicates (*n* ≥ 2). Asterisk marks indicate qualitative ligand binding as low enthalpy change prevented quantification. D) Co-crystallization studies with *L. monocytogenes* lplA1 and ligands (**R-LA**, PDBID 8cri, left; **LAQ**, PDBID 8crj, middle; **R-C3**, PDBID 8crl, right) with respective 2F_O-F_C electron density maps (grey mesh, contoured to 1.0 σ) and amino acid interactions are depicted as a close-up of the active pocket. The analysis shows binding of **R-C3** as intermediate mimic with Lys131 (plume) as a key interaction for coordination of the lipoyl-domain and adenine unit. E) Blood-stage *P. falciparum* parasites were incubated in presence of **LAMe**, **C3**, and **BrO** concentration-dependently to measure parasite growth inhibition. Growth experiments were performed in quadruplicates (*n* = 4) in two individual experiments. Mean values were plotted, and error bars represent SEM. The error range is shown as 95% confidence interval.

ligase of the parasite, demonstrating a comparably high binding affinity of **LA** ($K_D = 2.6 \mu\text{M}$) and **C3** ($K_D = 6.4 \mu\text{M}$) followed by **LAMe** ($K_D = 38 \mu\text{M}$, Figure 4C, S10B). To evaluate the effects of **C3** and **LAMe** on the growth of **LA** auxotroph organisms, we tested the compounds against *L. monocytogenes* and *P. falciparum*. While we did not observe any effects on the growth of the bacterial strain, even in macrophage infection assays (Figure S12), **LAMe** and **C3** killed blood-stage *P. falciparum* with EC₅₀ values of 15 μM and 27 μM, respectively (Figure 4E). Although the benchmark inhibitor **BrO** achieved a similar EC₅₀ value, effective parasite eradication required millimolar concentrations, corroborating previous literature reports.^[12a] Thus,

these novel compounds represent the most potent LPL inhibitors reported to date.

Conclusion

Lipoylation is a crucial post-translational modification which enables the catalysis of essential cellular processes, such as the oxidative decarboxylation of pyruvate to acetyl-CoA (pyruvate dehydrogenase).^[8] As human and pathogenic cells differ in the salvage pathways for **LA**, selective inhibitors of pathogenic lipoylase represent an attractive strategy to develop anti-infective therapies.^[13c] However,

tools to directly monitor LPL activity in living cells have been elusive so far.

We developed a cell-permeable lipoylation-probe (probe **2**) that utilizes LPL to modify **LA**-dependent enzymes covalently and thus serves as proxy for its activity. This in situ labeling strategy exhibits several benefits over screens against recombinant LPL, including the direct readout of cell permeability and sufficient in situ target engagement. Mass spectrometry confirmed the specificity of probe **2** for intracellular lysine residues in **LA**-dependent proteins. Based on this proof-of-concept, we designed a small virtual compound library by computational methods and identified two cell-permeable and potent active site LPL inhibitors, namely **C3** and **LAMe**. Co-crystallization of **C3** provided intriguing insights into LPL inhibition as the compound forms strong bonds with Lys131 in *L. monocytogenes* lplA1. However, no antibiotic activity was observed in *L. monocytogenes*, even though previous work showed that lplA1 is essential for the growth of bacteria in macrophages.^[11a,b] We confirmed uptake and stability of **C3** in *L. monocytogenes*, raising the possibility that the lack of antibiotic activity could be due to strong competition with **LA** during salvage. Further modification of the inhibitor (e.g., mimicking the interaction of **LAQ** phosphate with Lys131 as observed in crystal structures) could improve competition with **LA**. Based on the conserved three-dimensional fold and active site pockets, we tested **C3** and **LAMe** against *P. falciparum* to see if these molecules may inhibit LPLs across species. Indeed, we saw killing of blood-stage parasites with both inhibitors, validating our approach.

Overall, we showcase LPL-dependent probes as ideal tools for identifying **LA** salvage pathway inhibitors with effective in situ activity against the malaria-causing parasite *P. falciparum*. As an outlook, iterative cycles of pre-selected computational compounds followed by intracellular LPL profiling will minimize the overall experimental effort and enhance the chances of cell-permeable hit molecules with high potency.

Author Contributions

J.N.D. and S.A.S. conceived and designed the project. J.N.D. performed synthesis, labeling, mass spectrometry, protein purifications, ITC studies, and biological assays. S.Y.C. and M.H. ran docking studies and molecular dynamics simulations. M.G. performed crystallography with help of J.N.D. M.S. conducted assays with parasites in cell culture. S.M.H. provided isoDTB tags and assisted with the evaluation of binding site identification. S.A.S., J.N.D., and S.Y.C. wrote the manuscript with input of all authors.

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Conflict of Interest

The authors declare no conflict of interest.

Data Availability Statement

The proteomics data have been deposited to the ProteomeXchange Consortium^[31] via the PRIDE^[32] partner repository, which can be accessed with the identifier PXD040359. All new crystal structures have been deposited in the RCSB Protein Data Bank (PDB) with the depository numbers 8crl, 8crj, 8crl.^[33] Analyzed proteomics data and comprehensive ZINC database^[27] screen hits are presented in supplementary files.

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