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Citation

Mock, E. D., Mustafa, M., Gunduz-Cinar, O., Cinar, R., Petrie, G. N., Kantae, V., ... Stelt, M. van der. (2020). Discovery of a NAPE-PLD inhibitor that modulates emotional behavior in mice. *Nature Chemical Biology*, 16(6), 667-675. doi:10.1038/s41589-020-0528-7

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Discovery of a NAPE-PLD inhibitor that modulates emotional behavior in mice

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***N*-acylethanolamines (NAEs), which include the endocannabinoid anandamide, represent an important family of signaling lipids in the brain. The lack of chemical probes that modulate NAE biosynthesis in living systems hamper the understanding of the biological role of these lipids. Using a high-throughput screen, chemical proteomics and targeted lipidomics, we report here the discovery and characterization of LEI-401 as a CNS-active *N*-acylphosphatidylethanolamine phospholipase D (NAPE-PLD) inhibitor. LEI-401 reduced NAE levels in neuroblastoma cells and in the brain of freely moving mice, but not in NAPE-PLD KO cells and mice, respectively. LEI-401 activated the hypothalamus-pituitary-adrenal axis and impaired fear extinction, thereby emulating the effect of a cannabinoid CB₁ receptor antagonist, which could be reversed by a fatty acid amide hydrolase inhibitor. Our findings highlight the distinctive role of NAPE-PLD in NAE biosynthesis in the brain and suggest the presence of an endogenous NAE tone controlling emotional behavior.**

In recent years, lipids have come to the foreground as signaling mediators in the central nervous system (CNS)^{1,2}. While classical neurotransmitters are stored in synaptic vesicles and released on fusion with the plasma membrane of neurons, due to their lipophilic nature, lipids readily diffuse through membranes and are not stored in vesicles. It is, therefore, generally accepted that signaling lipids are produced ‘on demand’ and are rapidly metabolized to terminate their biological action³. In particular, NAEs, including *N*-palmitoylethanolamine (PEA), *N*-oleoylethanolamine (OEA) and the endocannabinoid anandamide (*N*-arachidonylethanolamine, AEA) have emerged as key lipid signaling molecules. Genetic deletion or pharmacological inhibition of the main NAE hydrolytic enzyme, fatty acid amide hydrolase (FAAH), revealed elevated anandamide, PEA and OEA levels in brain and implicated these molecules in the modulation of various physiological processes such as pain, stress, anxiety, appetite, cardiovascular function and inflammation^{4–7}. The physiological effects resulting from perturbation of the production of anandamide and other NAEs in living

systems are, however, poorly studied, partly because of a lack of pharmacological tools to modulate their biosynthetic enzymes⁸.

NAPE-PLD is generally considered a principal NAE biosynthetic enzyme^{9,10}. Biochemical and structural studies have demonstrated that NAPE-PLD is a membrane-associated, constitutively active zinc hydrolase with a metallo- β -lactamase fold¹¹. The enzyme generates a broad range of NAEs by hydrolysis of the phosphodiester bond between the phosphoglyceride and the NAE in *N*-acylphosphatidylethanolamines (NAPEs)¹². Knockout (KO) studies have shown that the Ca²⁺-dependent conversion of NAPE to NAEs bearing both saturated and polyunsaturated fatty acyl groups are fivefold reduced in brain lysates from mice that genetically lack *Napepld*¹³. In accordance, reduced levels of saturated and mono-unsaturated NAEs were observed in the brains of NAPE-PLD KO mice^{13–15}. Anandamide levels were not reduced in the transgenic model reported by Leung et al., which suggested the presence of compensatory mechanisms¹³. Indeed, multiple alternative biosynthetic pathways for anandamide have been discovered since¹⁰.

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Of note, two other NAPE-PLD KO strains have been generated, which did show a decreased brain anandamide content^{14,15}. Constitutive genetic mouse models are, however, poorly suited to study the rapid and dynamic formation of NAEs and their biological functions. Potent CNS-active inhibitors of NAPE-PLD are currently not available. Previously described NAPE-PLD inhibitors lack the potency, selectivity and/or physicochemical properties to study the role of this enzyme in *in vivo* systems in an acute setting^{16–18}. Thus, a need exists for pharmacological tools that inhibit NAE biosynthesis in the brain.

Here, we report the discovery of LEI-401 (**1**) as a brain active NAPE-PLD inhibitor. High-throughput screening (HTS) and a medicinal chemistry program led to the identification of LEI-401 as a potent and selective NAPE-PLD inhibitor. LEI-401 reduced a broad range of NAEs including anandamide in neuronal cells in a NAPE-PLD-dependent manner. Anandamide and other NAE levels were reduced in the brain of wild-type (WT) mice treated with LEI-401, but not in NAPE-PLD KO mice. The compound activated the hypothalamus–pituitary–adrenal (HPA) axis and impaired extinction of an aversive memory in mice, thereby mimicking the effects of cannabinoid CB₁ receptor antagonism. These behavioral effects were prevented by a FAAH inhibitor. Taken together, LEI-401 is a first-in-class brain active NAPE-PLD inhibitor that blocks NAE biosynthesis in the brain of freely moving mice, thereby revealing a possible endogenous tone of this lipid family in emotional behavior.

Results

New NAPE-PLD inhibitor chemotypes are identified by HTS. To identify new inhibitors for NAPE-PLD, we initiated a HTS campaign in collaboration with the European Lead Factory (Supplementary Table 1)¹⁹. A previously reported fluorescent NAPE-PLD activity assay²⁰ using the surrogate substrate *N*-((6-(2,4-dinitrophenyl)amino)hexanoyl)-2-(4,4-difluoro-5,7-dimethyl-4-bora-3a,4a-diaza-*s*-indacene-3-pentanoyl)-1-hexadecanoyl-*sn*-glycero-3-*p* phosphoethanol-amine triethylammonium salt (PED6) was converted to a HTS-compatible 1,536-well format (Fig. 1a). As an enzyme source, we used membrane fractions of human embryonic kidney 293T (HEK293T) cells that were transiently transfected with human NAPE-PLD (Supplementary Fig. 1). This resulted in a robust assay with a $Z' = 0.87$, a signal-to-background ratio of 6.7 and intraplate variability (coefficient of variability) of <3.3%. Approximately 350,000 compounds were screened at a single concentration (10 μM) in 294 plates over 3 d, affording 8,321 actives (using the cut-off value of Z score ≤ -7) (Fig. 1b,c). Hit validation at the same concentration yielded 3,885 confirmed actives. To reduce the number of compounds, the percentage effect on the enzyme activity ($\geq 40\%$) was used to obtain a list of 1,120 actives. Next, dose-response curves were generated in the presence and absence of ZnSO_4 (100 μM) to remove promiscuous Zn^{2+} -ion chelators, which yielded 352 hits. Visual inspection of the 352 compounds revealed the presence of potential assay interfering compounds that absorbed light within the visible wavelength; therefore, a second deselection assay was developed. The substrate PED6 was incubated with lysate to obtain the maximum fluorescent signal, followed by incubation with different concentrations of the confirmed actives to determine the ability of the compounds to quench the fluorescent signal under the primary assay conditions. Compounds demonstrating no effect (half-maximal inhibitory concentration with negative log (pIC₅₀) < 5) in the second deselection assay were selected (Supplementary Table 2). The compounds were checked for purity (>85%) and correct molecular weight by liquid chromatography–mass spectrometry (LC–MS) analysis. This afforded a qualified hit list of five compounds (**2–6**) with four different chemotypes (Fig. 1d).

Hit optimization provides nanomolar inhibitor LEI-401. Compound **2** (*N*-(cyclopropylmethyl)-2-(methyl(phenethyl)amino)-6-morpholinopyrimidine-4-carboxamide) displayed the most optimal biological and physicochemical characteristics (molecular weight

of 396 Da, calculated partition coefficient $c\text{-log}P = 3.84$) (Fig. 1d and Supplementary Table 2). On resynthesis, chemical analysis and retesting, the identity and submicromolar activity ($K_i = 0.30 \mu\text{M}$, 95% CI = 0.22–0.38 μM) of the hit was confirmed (Fig. 2 and Supplementary Notes). To improve the inhibitory activity of compound **2**, a medicinal chemistry program was initiated in which 112 derivatives were synthesized. A full report of the medicinal chemistry program will be described elsewhere; however, in brief, we discovered that replacing the morpholine for a (*S*)-3-hydroxypyridine (**7**) improved activity ($K_i = 0.086 \mu\text{M}$, 95% CI, 0.058–0.14 μM), while simultaneously decreasing lipophilicity ($c\text{-log}P = 3.46$) (Fig. 2a,b). The absolute configuration of the hydroxyl was not essential for binding, as the (*R*)-enantiomer **8** showed similar potency ($K_i = 0.11 \mu\text{M}$, 95% CI, 0.096–0.13 μM). Conformational restriction of the phenethylamine moiety to a (*S*)-3-phenylpiperidine (**1**) further increased the potency ($K_i = 0.027 \mu\text{M}$, 95% CI, 0.021–0.033 μM), whereas its (*R*)-enantiomer **9** was three times less active ($K_i = 0.094 \mu\text{M}$, 95% CI, 0.071–0.12 μM) (Fig. 2b). In view of the biological activity profile, we decided to further characterize compound **1** (termed LEI-401).

To profile NAPE-PLD inhibitors in mouse models, we assessed whether LEI-401 showed any species difference using recombinant mouse NAPE-PLD expressed in HEK293T cells. LEI-401 retained activity on mouse NAPE-PLD in the PED6 assay, albeit at a reduced potency ($K_i = 0.18 \mu\text{M}$, 95% CI, 0.15–0.21 μM). Next, we investigated the activity of LEI-401 on NAPE-PLD in a secondary *in vitro* assay to confirm an on-target effect of the inhibitor. To this end, we measured the NAPE-PLD enzyme activity in brain lysates of WT and NAPE-PLD KO mice by quantifying the formation of NAE (C17:0) from a synthetic C17:0-NAPE using an LC–MS-based assay²¹. LEI-401 reduced C17:0-NAE formation using brain homogenates of WT mice, but not from KO mice (Supplementary Fig. 3), thereby confirming that NAPE hydrolysis and its inhibition by LEI-401 in *ex vivo* preparations is dependent on NAPE-PLD in mouse brain. Following this, we assessed the activity profile of LEI-401 for the receptors and metabolic enzymes of the endocannabinoid system. No inhibitory activities were measured at 10 μM for the cannabinoid receptors type 1 and type 2 (Supplementary Table 3) as well as for the other enzymes involved in anandamide biosynthesis and degradation, such as phospholipase A2 group 4E (PLA2G4E) and FAAH (Supplementary Table 4), respectively. In addition, LEI-401 did not inhibit enzymes involved in the biosynthesis and degradation of the other endocannabinoid 2-arachidonoylglycerol (2-AG), including diacylglycerol lipases (DAGL α or DAGL β), monoacylglycerol lipase (MAGL) and α,β -hydrolase domain 6 (ABHD6) (Supplementary Table 4).

Photoaffinity labeling shows NAPE-PLD target engagement. To assess target engagement and selectivity of LEI-401 in cellular systems, we turned to a photoaffinity labeling approach, which was previously successfully applied for other metallo-enzymes²². Based on the structure-activity relationships of LEI-401, a photoactivatable alkyne probe (**10**) was synthesized in which the 3-phenyl-piperidine was replaced with a trifluoromethyl-diazirine-benzyl-piperazine as a photoreactive group (Fig. 3a and Supplementary Notes). In addition, the methylcyclopropyl group was substituted for a propargyl group to serve as a ligation handle to introduce reporter groups by copper(I)-catalyzed azide-alkyne cycloaddition (CuAAC, or 'click') chemistry to study target engagement. First, we confirmed that compound **10** inhibited NAPE-PLD with a K_i of 0.18 μM (95% CI, 0.11–0.27 μM) in the biochemical PED6-assay (Supplementary Fig. 4). To assess whether compound **10** was able to visualize NAPE-PLD, HEK293T cells were transfected with hNAPE-PLD-FLAG or control (mock) plasmid for 48 h, followed by *in situ* treatment with compound **10** (2 μM) for 30 min and then irradiated at 350 nm for 10 min. After lysis and clicking of a Cy5-N₃ fluorophore, the samples were resolved by SDS-polyacrylamide gel

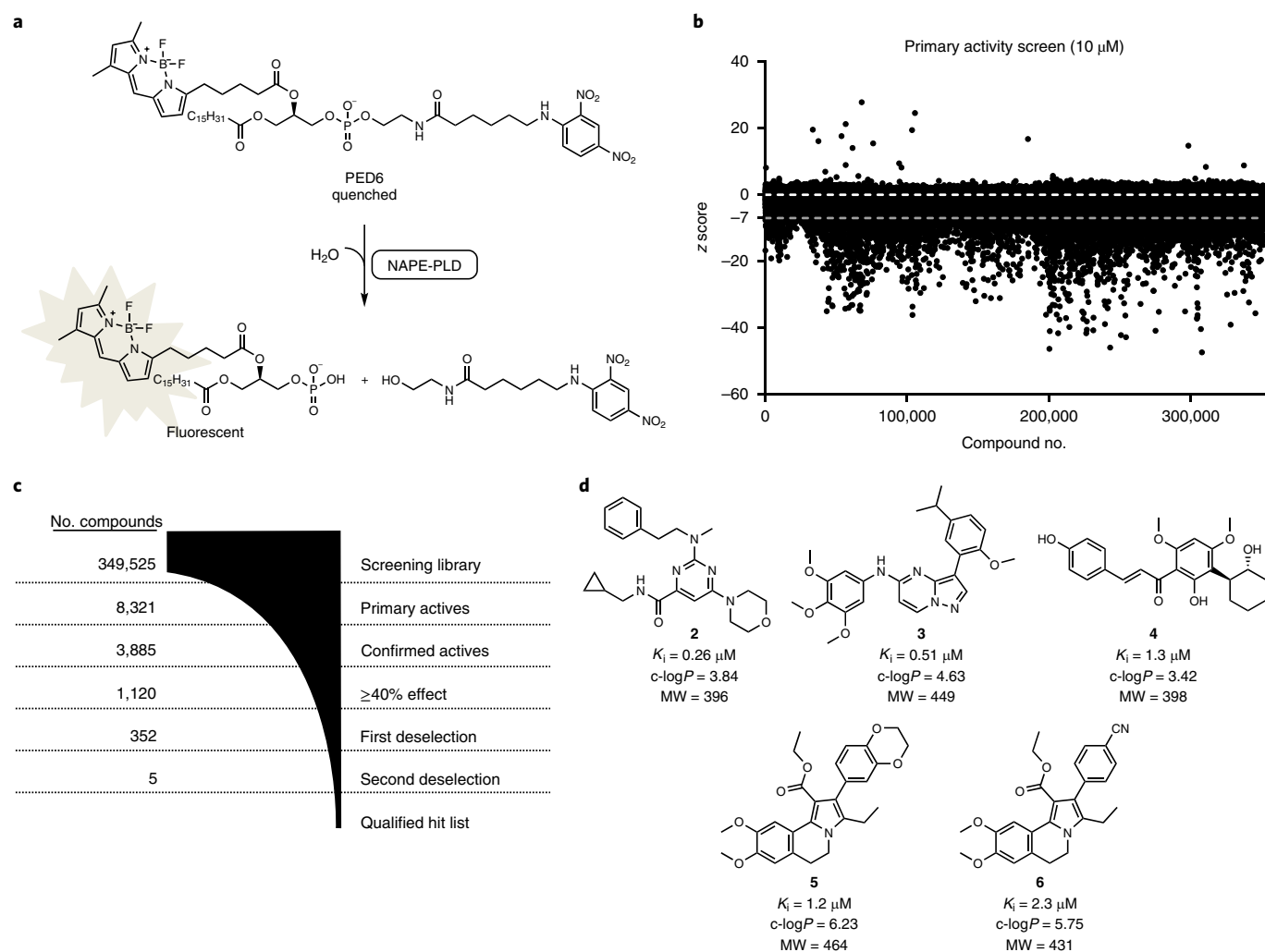


Fig. 1 | HTS of a NAPE-PLD activity assay provided potent new inhibitors 2–6. a, Cleavage of PED6 generated a fluorescent read-out of NAPE-PLD activity. **b**, Primary activity screen. **c**, Hit triage of the HTS campaign. **d**, Structures, potency data and physicochemical parameters of hit compounds (2–6). Data represent mean values \pm s.e.m. for two independent experiments with two biological replicates ($n = 4$). Dose-response curves are depicted in Supplementary Fig. 2. Complete HTS data are listed in Supplementary Tables 1 and 2.

electrophoresis (SDS-PAGE) and protein labeling was visualized by in-gel fluorescence scanning (Supplementary Fig. 5–7). An additional band at the expected molecular weight of ~ 46 kDa was apparent in the hNAPE-PLD-transfected cells but not in mock-treated cells, at the same height as the signal of an anti-FLAG antibody on western blot (Fig. 3b). The intensity of the fluorescent band at 46 kDa was reduced on coinubation by the inhibitor LEI-401 (Fig. 3b), indicating that LEI-401 bound to NAPE-PLD in living cells. Concentration-response experiments revealed that LEI-401 dose-dependently reduced the labeling of NAPE-PLD with an IC_{50} of 0.86 μM (95% CI, 0.60–1.2 μM) (Fig. 3c,d). To establish the selectivity of LEI-401, a label-free chemical proteomics experiment was performed²³. Cell lysates prepared from hNAPE-PLD-transfected HEK293T cells incubated with photoprobe **10** were clicked with biotin- N_3 , which allowed for enrichment of probe-labeled proteins using avidin agarose beads, followed by trypsin digestion and protein identification by mass spectrometry. Proteomic analysis allowed identification of 136 proteins (Fig. 3e). NAPE-PLD was enriched by probe **10** in an ultraviolet (UV)-dependent manner and its labeling could be prevented by coinubation with LEI-401 (Fig. 3f). Furthermore, LEI-401 did not compete for other protein targets of probe **10** in this analysis or in mock-transfected

HEK293T cells (Fig. 3f and Supplementary Fig. 8). In summary, the development of photoaffinity probe **10** enabled the detection of NAPE-PLD in living cells and visualization of target engagement by LEI-401 with NAPE-PLD.

LEI-401 lowers NAEs in WT but not in NAPE-PLD KO cells.

Having established that LEI-401 is a cell-permeable inhibitor and engages with NAPE-PLD, we then investigated whether LEI-401 inhibited endogenous NAPE-PLD in living cells. To this end, the mouse Neuro-2a neuroblastoma cell line was selected, since it expressed endogenous NAPE-PLD as determined by quantitative PCR (qPCR; Supplementary Table 5) and western blot using NAPE-PLD antibodies (Supplementary Fig. 9). Targeted lipidomics on lipid extracts of Neuro-2a cells allowed the quantification of eight different NAEs by LC-MS (Fig. 4a). As an appropriate negative control, two NAPE-PLD KO cell lines were generated using CRISPR/Cas9. The cell line KO-2 showed more efficient ablation of NAPE-PLD protein expression as demonstrated by western blot analysis compared to KO-1 and was therefore chosen for further analysis (Supplementary Fig. 9). Next, we incubated both WT and KO cells with LEI-401 (10 μM) for 2 h. A significant twofold reduction of anandamide was apparent in WT cells but not in NAPE-PLD KO

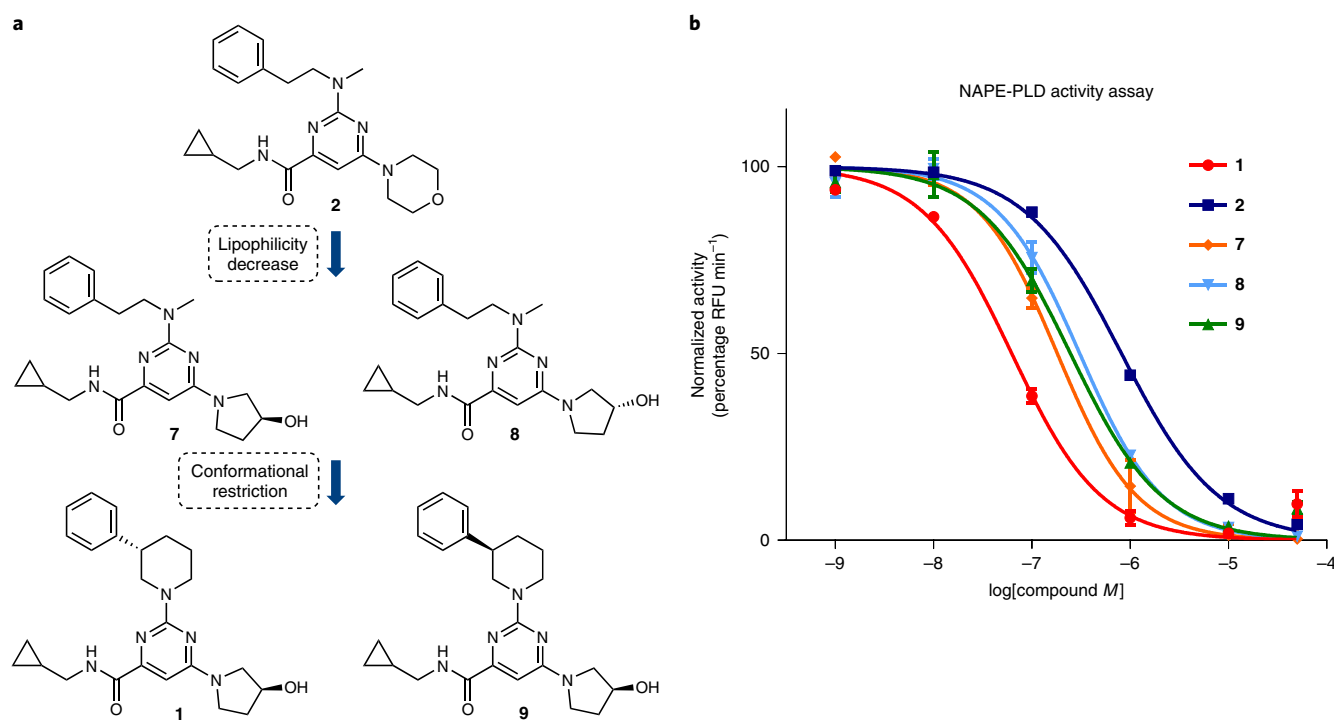


Fig. 2 | Hit optimization affords nanomolar potent inhibitor LEI-401. **a**, Respective substitution of the morpholine and *N*-methylphenethylamine of **2** for a (*S*)-3-hydroxypyrrolidine and (*S*)-3-phenylpiperidine provided LEI-401. **b**, hNAPE-PLD in vitro activity assay dose-response curves for inhibitors **1** ($K_i = 0.027 \mu\text{M}$, 95% CI 0.021–0.033 μM), **2** ($K_i = 0.30 \mu\text{M}$, 95% CI 0.22–0.38 μM), **7** ($K_i = 0.086 \mu\text{M}$, 95% CI 0.058–0.14 μM), **8** ($K_i = 0.11 \mu\text{M}$, 95% CI 0.096–0.13 μM) and **9** ($K_i = 0.094 \mu\text{M}$, 95% CI 0.071–0.12 μM). Data represent mean values \pm s.e.m. for two independent experiments with two biological replicates ($n = 4$).

cells, indicating that the LEI-401-induced reduction in anandamide levels depended on expression of NAPE-PLD in Neuro-2a cells (Fig. 4b). LEI-401-treated cells did not show any sign of reduced cell viability (Supplementary Fig. 10). It is worth noting that the smaller but significant decrease of anandamide (~25%) in NAPE-PLD KO cells compared to WT may indicate the occurrence of compensatory mechanisms in the knockout cells, as also observed by others in *Napepld*-deficient mice (Supplementary Fig. 11a)¹³. Furthermore, other saturated, mono- and (ω -6)-polyunsaturated NAEs showed a similar twofold decrease in WT cells on LEI-401 pretreatment, but not in NAPE-PLD KO cells (Fig. 4b and Supplementary Fig. 11b–f). Of note, (ω -3)-polyunsaturated *N*-eicosapentanoylethanolamine (EPEA) and *N*-docosahexanoylethanolamine (DHEA) and did not respond to LEI-401 treatment (Fig. 4b and Supplementary Fig. 11g,h). Our data are in line with previously reported biochemical substrate studies using purified recombinant NAPE-PLD¹² and indicate that endogenously expressed NAPE-PLD is responsible for biosynthesis of saturated, mono- and (ω -6)-polyunsaturated NAEs in Neuro-2a cells. The lack of effect on (ω -3)-polyunsaturated NAEs may suggest that these signaling lipids are produced via another pathway. Of note, recently it was found that dietary (ω -3)-polyunsaturated fatty acids augment NAE levels in mice via a pathway independent of NAPE-PLD²⁴.

LEI-401 is brain-penetrant and reduces brain NAE levels. To assess whether LEI-401 possesses in vivo efficacy, we first determined its in vitro absorption, distribution, metabolism and excretion (ADME) profile. LEI-401 demonstrated favorable physicochemical properties for oral bioavailability and brain penetration (molecular weight of 422 Da; $\log D = 3.3$ and topological polar surface area of 80.5 \AA^2). LEI-401 had low aqueous solubility (1.7 mg l^{-1}) and permeates well through membranes (parallel artificial membrane permeability assay (PAMPA) $P_{\text{eff}} = 0.37 \text{ nm s}^{-1}$). Clearance in human and

mouse microsomes (10 and 42 $\mu\text{l min}^{-1} \text{mg}^{-1}$ protein, respectively) as well as in human and mouse hepatocytes (6.9 and 28.8 $\mu\text{l min}^{-1}$ per 10^6 cells, respectively) was moderate, except for the low human microsomal clearance. LEI-401 exhibited high human and mouse protein binding of >99.8%. It is worth noting that it did not interact with either human or mouse PGP-transporter (efflux ratios were 1.7 and 1.8, respectively).

Pharmacokinetic analysis in male C57BL/6J mice revealed a low clearance ($\text{CL} = 5.2 \text{ ml min}^{-1} \text{kg}^{-1}$) and moderate volume of distribution ($V_{\text{ss}} = 1.21 \text{ kg}^{-1}$), resulting in a half-life of 2.7 h (Fig. 5a and Supplementary Table 6). The low in vivo clearance was unexpected based on the in vitro clearance, but could be explained by the tight protein binding. Bioavailability after oral administration reached 25% and a more favorable bioavailability was achieved via intraperitoneal (i.p.) injection ($F_{\text{ip}} = 48\%$). In both cases, good plasma exposures were obtained. The brain to plasma ratio was 0.7 at C_{max} (2 h postinjection). This led to an excellent brain exposure (>10 μM) after 30 mg kg^{-1} i.p. administration (Fig. 5b), which corresponds to approximately 70-fold the mouse NAPE-PLD K_i -value, and should be sufficient to modulate NAPE-PLD activity in mouse brain.

In view of the encouraging pharmacokinetic properties and brain exposure, we administered LEI-401 or vehicle to male C57BL/6J mice (30 mg kg^{-1} , i.p., single dose) and killed the mice after 1, 2, 4 or 8 h to determine whether NAE levels were reduced. To this end, we analyzed brain NAE levels by LC–MS. LEI-401 induced a significant and time-dependent reduction for anandamide ($P = 0.0051$), which was significant at 2 h and returned to vehicle level by 4 h (Fig. 5c). A similar trend, although not significant, was observed for OEA and DHEA (Supplementary Fig. 12a,b). Next, mice were treated with various doses of LEI-401 (3, 10 and 30 mg kg^{-1} or vehicle, i.p., single dose) and brain lipids were analyzed at 2 h. LEI-401 (30 mg kg^{-1}) significantly reduced anandamide (Fig. 5d). Other NAE lipid species showed a similar trend (Supplementary Fig. 13a–c).

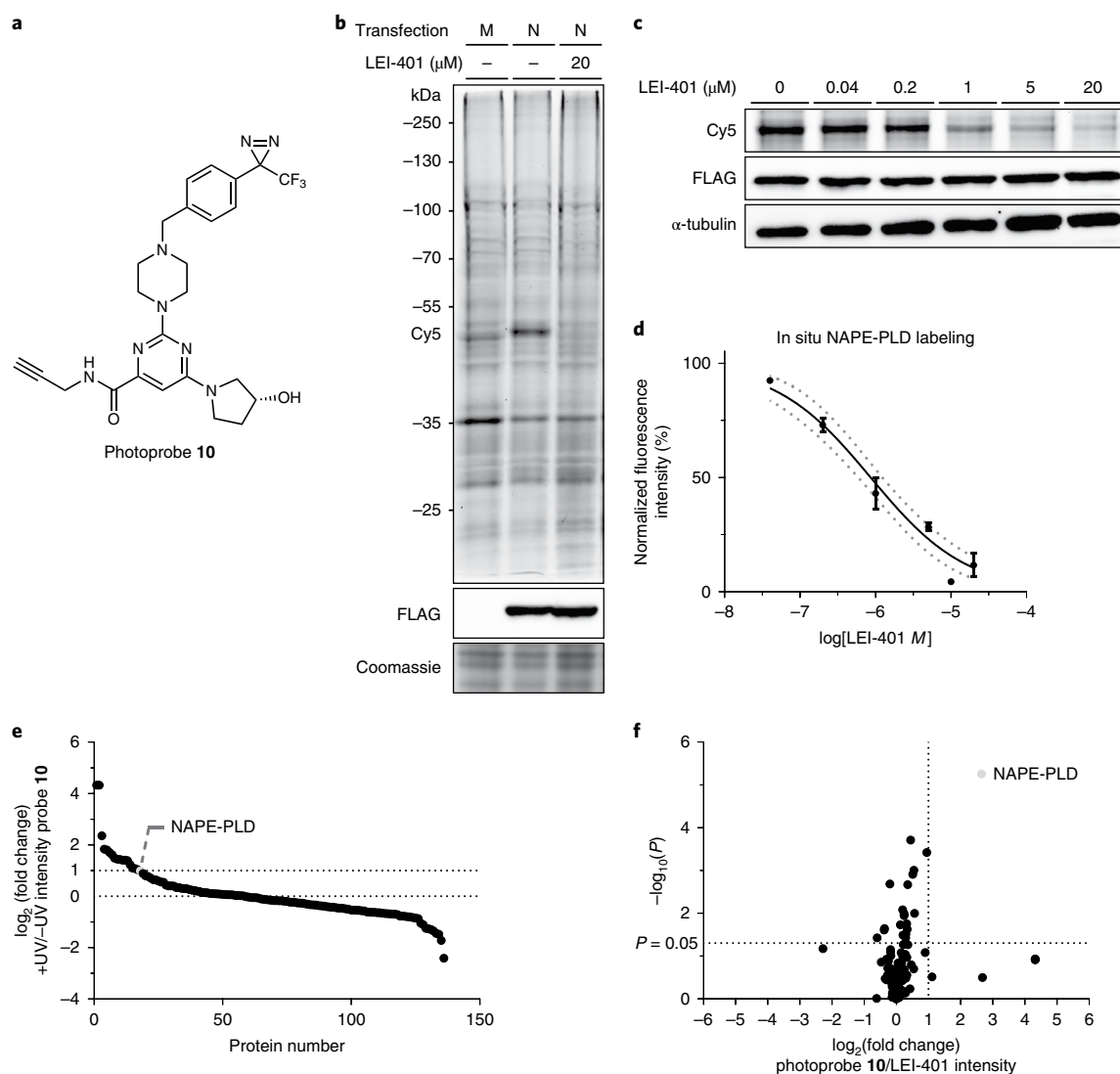


Fig. 3 | Photoprobe 10 visualizes target engagement of LEI-401 with NAPE-PLD in situ. **a**, Structure of NAPE-PLD photoprobe **10**. **b**, Fluorescent labeling of hNAPE-PLD-FLAG at ~46 kDa in overexpressing HEK293T cells (middle lane) but not in mock-transfected cells (left lane) with photoprobe **10** (2 μ M, 30 min). The NAPE-PLD band was competed out by LEI-401 (20 μ M, 30 min incubation) (right lane). Anti-FLAG western blot displays hNAPE-PLD-FLAG expression. M, mock; N, hNAPE-PLD-FLAG. Representative gel of three biologically independent experiments ($n=3$). **c**, Dose-dependent competition of photoprobe **10** (2 μ M) with LEI-401 (30 min incubation). Representative gel of three biologically independent experiments ($n=3$). For **b,c**, Coomassie or α -tubulin was used as a loading control. See Supplementary Figs. 6 and 7 for full gels and blots. **d**, Dose-response curve of LEI-401 with 95% CI (dotted lines); $IC_{50}=0.86 \mu$ M (95% CI, 0.60–1.2 μ M). Data represent mean values \pm s.e.m. for three biological replicates ($n=3$). **e**, Waterfall plot showing UV-enriched protein targets of photoprobe **10** in hNAPE-PLD-FLAG-transfected HEK293T cells (three biological replicates per condition ($n=3$)). **f**, Volcano plot displaying competed protein targets for photoprobe **10** (2 μ M) versus LEI-401 (20 μ M) in hNAPE-PLD-FLAG-transfected HEK293T cells (three biological replicates per condition ($n=3$)). Cut-off values used for **e,f**: unique peptides ≥ 2 ; $\log_2(\text{fold change}) \geq 1$ (for **e**, ratio +UV/–UV intensity probe 10 and for **f**, ratio photoprobe **10** intensity/LEI-401 intensity), $P < 0.05$ using Student's t -test (unpaired, two-tailed) and Benjamini-Hochberg correction (10% FDR).

The 2-AG levels were not significantly affected by LEI-401 treatment (30 mg kg⁻¹, i.p.) (Supplementary Figs. 12c and 13d). In line, the activity of the 2-AG producing and degrading enzymes DAGL α/β , MAGL and ABHD6 in the brain was not affected (Supplementary Fig. 14). LEI-401 did not reduce brain NAE levels in NAPE-PLD KO mice (Supplementary Fig. 15a–c,f–h), which indicated that LEI-401 is a selective NAPE-PLD inhibitor. Of note, we found that anandamide levels were not decreased by LEI-401 in WT littermates of the NAPE-PLD KO animals, which originate from a mixed 129SvJ-C57BL/6 background, while PEA and OEA levels were significantly reduced (Supplementary Fig. 15f–h). This is in line with previous genetic studies using animals from the same

background¹³. Furthermore, 2-AG levels were increased in KO mice at a dose of 60 mg kg⁻¹, whereas AA levels were decreased at this dose in animals of both genotypes (Supplementary Fig 15i,j). This indicated that at this high dose LEI-401 has an off-target that modulates 2-AG and AA levels. To determine whether acute inhibition of NAPE-PLD would affect its substrate levels, we determined NAPE and plasmalogen NAPE (pNAPE) levels in the brain of LEI-401-treated mice. No change in NAPE or pNAPE species was found for LEI-401 compared to vehicle in both WT and NAPE-PLD KO mice (Supplementary Figs. 16 and 17). Collectively, these data indicate that LEI-401 is an in vivo active NAPE-PLD inhibitor that can be used to study its physiological role.

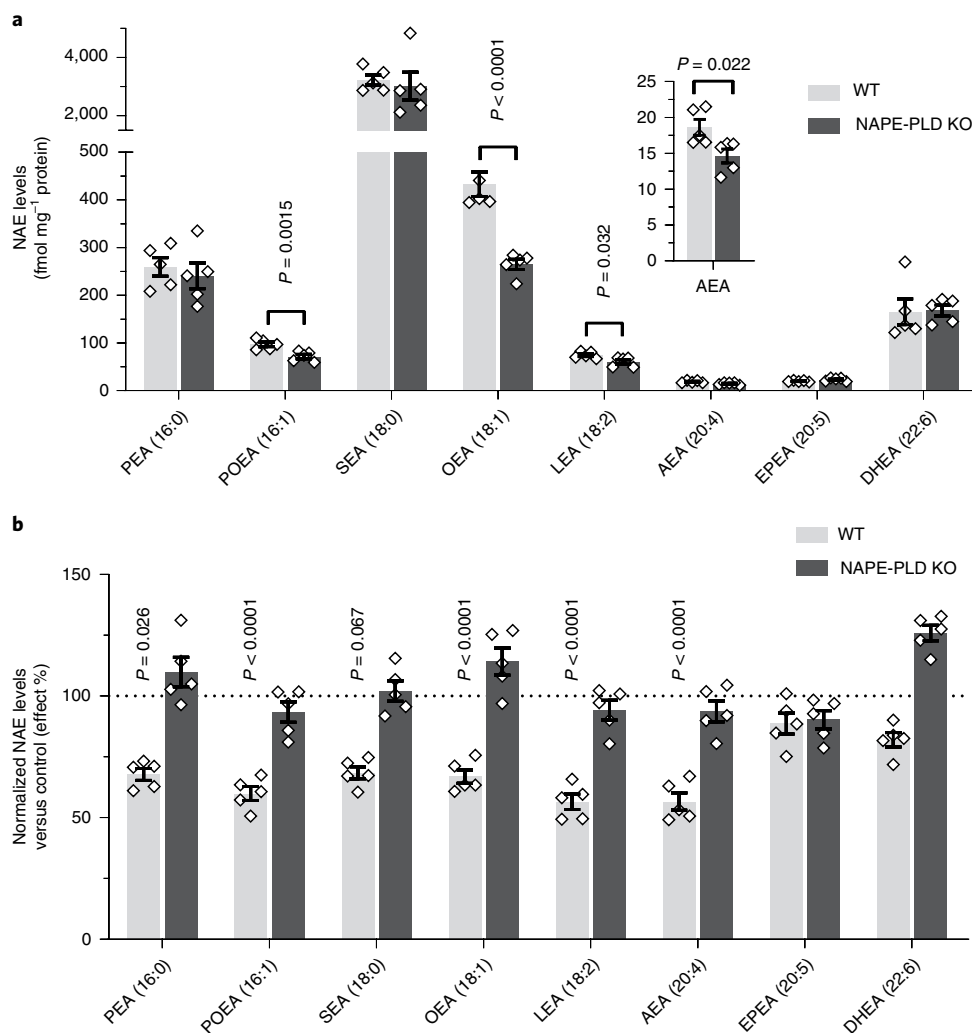


Fig. 4 | LEI-401 reduces NAE levels in Neuro-2a cells, but not in NAPE-PLD KO cells. a, Absolute NAE levels of Neuro-2a WT and NAPE-PLD KO cells. **b**, Relative NAE levels of Neuro-2a WT and NAPE-PLD KO cells treated with LEI-401 (10 μ M, 2 h), represented as effect percentage. Data were normalized against WT or NAPE-PLD KO cells treated with vehicle (DMSO). Absolute values for all NAEs are depicted in Supplementary Fig. 11. For **a, b**, data represent mean values \pm s.e.m. for five biological replicates ($n=5$). P values were calculated using one-way ANOVA with Tukey's multiple comparisons test.

LEI-401 modulates emotional behavior. NAEs are implicated in the regulation of emotional behavior, such as acute stress responses, anxiety and fear extinction^{6,25,26}. It is hypothesized that a basal tone of anandamide, acting via the cannabinoid CB₁ receptor, constrains activation of the HPA axis and controls the extinction of aversive memories⁵. This hypothesis is supported by experiments in which anandamide levels were elevated via pharmacological and genetic modulation of FAAH or by blocking the cannabinoid CB₁ receptor^{27–33}. More so, stress exposure itself is known to reduce anandamide levels in vivo and reversal of this suppression of anandamide signaling, via inhibition of FAAH, is known to counter many effects of stress such as activation of the HPA axis and alterations in anxiety^{34–36}. Direct evidence that reducing NAE levels by pharmacological tools will affect emotional behavior has, however, not been obtained yet. LEI-401 is the first chemical probe that allows us to test this hypothesis. To this end, we first measured c-Fos protein expression in the paraventricular nucleus (PVN) of the hypothalamus, a primary integratory site for the HPA axis and a nucleus that is reliably activated by stress exposure, and determined circulating corticosterone (CORT) levels in mice 2.5 h after administration of LEI-401 (30 mg kg⁻¹, i.p., single dose), a CB₁ receptor antagonist (AM251, 3 mg kg⁻¹, i.p., single dose) or vehicle. PVN brain sections

revealed that LEI-401 ($P<0.0001$) and AM251 ($P=0.011$) elevated c-Fos protein expression (Fig. 6a–d and Supplementary Fig. 18a–c). Notably, the effect of LEI-401 was reversed by cotreatment with the FAAH inhibitor URB597 (i.p., 1 mg kg⁻¹, single dose; $P=0.016$) (Fig. 6c,d). Plasma CORT levels were significantly higher in LEI-401-treated mice compared to control ($P=0.047$), but not when cotreated with a FAAH inhibitor ($P=0.0027$) (Fig. 6e). Similarly, an increase in plasma CORT was also observed for AM251-treated mice ($P=0.0042$; Supplementary Fig. 18d). These data suggest that NAPE-PLD provides a basal NAE tone, which restricts the activation of the HPA axis.

Next, to determine the consequences of depleting the endogenous NAE tone on a measure of emotional behavior, we tested LEI-401 on mouse fear extinction³⁷. First, C57BL/6J mice were trained to fear (quantified as freezing 'no movement else than breathing' levels) an auditory conditioned stimulus by repeated pairings with an unconditioned stimulus (US, footshock). One day after conditioning, mice were treated with LEI-401 (i.p., 30 mg kg⁻¹, single dose), LEI-401 cotreatment with a FAAH inhibitor (AM3506, i.p., 1 mg kg⁻¹, single dose) or vehicle 1 h and 45 min before a fear extinction training involving 50 \times conditioned stimulus presentations. A retrieval test was performed 10 d later to establish the duration of

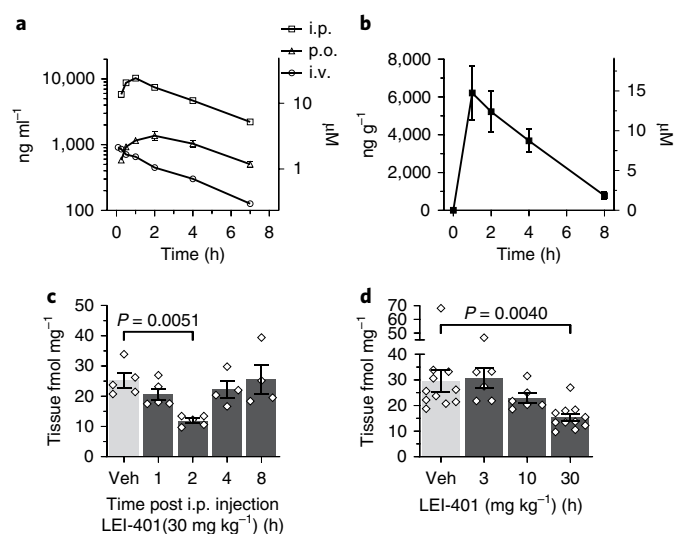


Fig. 5 | LEI-401 decreases AEA in mouse brain. **a**, In vivo pharmacokinetic profile of LEI-401 in plasma of C57BL/6J mice via intraperitoneal (i.p., 30 mg kg⁻¹), p.o. (10 mg kg⁻¹) or i.v. (1 mg kg⁻¹) administration. Data represent mean values \pm s.e.m. $n=5$ mice for time points 0.25 or 0.5 h, and $n=3$ mice for 1, 2, 4 or 7 h. **b**, Brain pharmacokinetic profile of LEI-401 in C57BL/6J mice after i.p. administration (30 mg kg⁻¹). **c**, Time-dependent effect of LEI-401 on brain AEA. C57BL/6J mice i.p. injected with LEI-401 (30 mg kg⁻¹). Veh, vehicle at 8 h. For **b** and **c**, data represent mean values \pm s.e.m. with $n=5$ mice for time points 1 or 2 h, and $n=4$ mice for 4 or 8 h. **d**, Dose-dependent effect of LEI-401 on brain AEA at 2 h after i.p. administration. Data represent mean values \pm s.e.m. with $n=5$ mice for dose groups 3 or 10 mg kg⁻¹ and $n=11$ mice for vehicle or 30 mg kg⁻¹. P values were calculated using one-way ANOVA with Dunnett's multiple comparisons test.

extinction. This entailed 5 \times conditioned stimulus presentations (Supplementary Fig. 19). LEI-401 administration produced a significant increase in freezing as compared to vehicle ($P=0.0099$), which was reversed by coadministration of a FAAH inhibitor (Fig. 6f). These results reveal a key contribution of NAPE-PLD in mediating fear extinction.

Discussion

NAEs, including anandamide, PEA and OEA, have emerged as an important class of lipid signaling molecules that regulate various physiological and behavioral functions via multiple mechanisms, including G protein-coupled receptors (cannabinoid CB₁ receptor, GPR110 and GPR119), ion channels (TRPV1 and TASK-1) and nuclear receptors (PPAR- α)³⁸. Inhibitors of NAE degradation have shed light on the role of NAEs in pain³⁹, satiety⁴⁰ and inflammation⁴¹, and emotional states such as fear⁶, stress²⁹ and anxiety²⁷. However, assessing the biological consequences of NAE reductions represent an important challenge because of a lack of selective inhibitors that block in vivo NAE production.

Here, we have addressed these important questions by developing LEI-401 as a brain active inhibitor of NAPE-PLD, an enzyme responsible for NAE biosynthesis. HTS and a medicinal chemistry program led to the discovery of LEI-401 as a potent and selective NAPE-PLD inhibitor with good pharmacokinetic properties. Acute NAPE-PLD inhibition by LEI-401 in mouse neuronal Neuro-2a cells decreased saturated, mono- and (ω -6)-polyunsaturated NAEs (including anandamide), while leaving (ω -3)-polyunsaturated NAEs unperturbed. This effect was not present in Neuro-2a cells lacking NAPE-PLD, showing the specificity of LEI-401. Notably, acute NAPE-PLD inhibition resulted in lower anandamide levels than observed in the knockout cells, thereby indicating that

long-term inactivation of NAPE-PLD in cells also may lead to activation of compensatory pathways (Supplementary Fig. 11a). LEI-401 reduced NAE formation from synthetic NAE ex vivo using brain lysates from WT mice, but not from NAPE-PLD KO mice. Administration of LEI-401 to mice resulted in a significant decrease in anandamide, PEA and OEA content in the brain. The LEI-401-mediated reductions in NAEs were absent in NAPE-PLD KO mice, thereby indicating that LEI-401 is a selective NAPE-PLD inhibitor. Of note, we found that anandamide levels were acutely reduced by LEI-401 in C57BL/6J mice, but not in animals derived from a mixed 129SvJ-C57BL/6J background. These observations are consistent with previous studies using the same mouse strains, in which genetic ablation of NAPE-PLD in C57BL/6J mice resulted in reduced anandamide levels, but not in mice with a mixed background^{13,14}. This reinforces the notion that besides NAPE-PLD alternative biosynthetic pathways exist for anandamide biosynthesis, which not only become apparent after long-term inactivation of the *Napepld* gene^{10,13}, but also appear to be functional in normal physiological conditions depending on the genetic context. While the molecular factors that govern the biosynthesis of anandamide in mice originating from a 129SvJ-C57BL/6J background remain unknown at this time, it cannot be excluded that NAPE-PLD regulates anandamide biosynthesis in specific brain regions, cell types or other tissues in these mice. Regardless of the genetic context, our data indicate that LEI-401 is an in vivo active inhibitor that can be used to study the physiological role of NAPE-PLD. However, it should be noted that when studying the behavioral effects of LEI-401 in relation to a reduction of anandamide levels it is important to take the genetic background of the animals into account.

LEI-401 provided us the opportunity to investigate the potential role of NAPE-PLD in emotional behavior. LEI-401 elicited a pronounced activation of the HPA axis, which was mimicked by a CB₁ receptor antagonist and could be blocked by coadministration of a FAAH inhibitor. These findings support the hypothesis that a basal anandamide tone controls the activation of the HPA axis via the CB₁ receptor⁵. LEI-401 also impaired fear extinction, an effect that was rescued by a FAAH inhibitor, consistent with previous findings in which genetic deletion or pharmacological blockade of the CB₁ receptor resulted in impaired fear extinction^{28,42–44}. Taken together, our data are consistent with previous studies suggesting that anandamide may set an endogenous tone that plays a role in the neural circuitries mediating stress and fear behavior and provide experimental evidence that inhibition of NAPE-PLD is sufficient to modulate stress and fear. Whether the reduction of other NAEs, besides anandamide, also contribute to the observed behavioral effects of LEI-401 cannot be excluded at the moment²⁶. Taken together, our data lend further preclinical support to test FAAH inhibitors in the clinic for stress-related disorders, such as posttraumatic stress disorder, which has been reported to exhibit reduced levels of anandamide^{45,46}.

Finally, our studies indicate that LEI-401 and photoreactive probe **10** constitute a valuable toolbox to study various aspects of NAPE-PLD and NAE biology both in animal and in cellular models. Although LEI-401 was selective in a chemical proteomics setting, did not inhibit DAGL α , DAGL β , MAGL and ABHD6, and did not affect NAE, 2-AG and AA levels in NAPE-PLD KO mice at 30 mg kg⁻¹, it modulated brain 2-AG and AA levels at a higher dose of 60 mg kg⁻¹ in both WT and KO mice. These observations indicate that LEI-401 or its metabolites may have off-target activity and highlight the valuable role of combining chemical tools with genetic models to study on-target and off-target interactions of pharmacological agents. Second generation inhibitors will benefit from the identification of the off-targets. From a translational perspective, it will be interesting to study these inhibitors in various animal models of human disease, such as metabolic syndrome⁴⁷, type II diabetes⁴⁸, neurodegeneration⁴⁹, inflammatory and neuropathic pain⁵⁰. The inhibitor reported in this study provided an opportunity

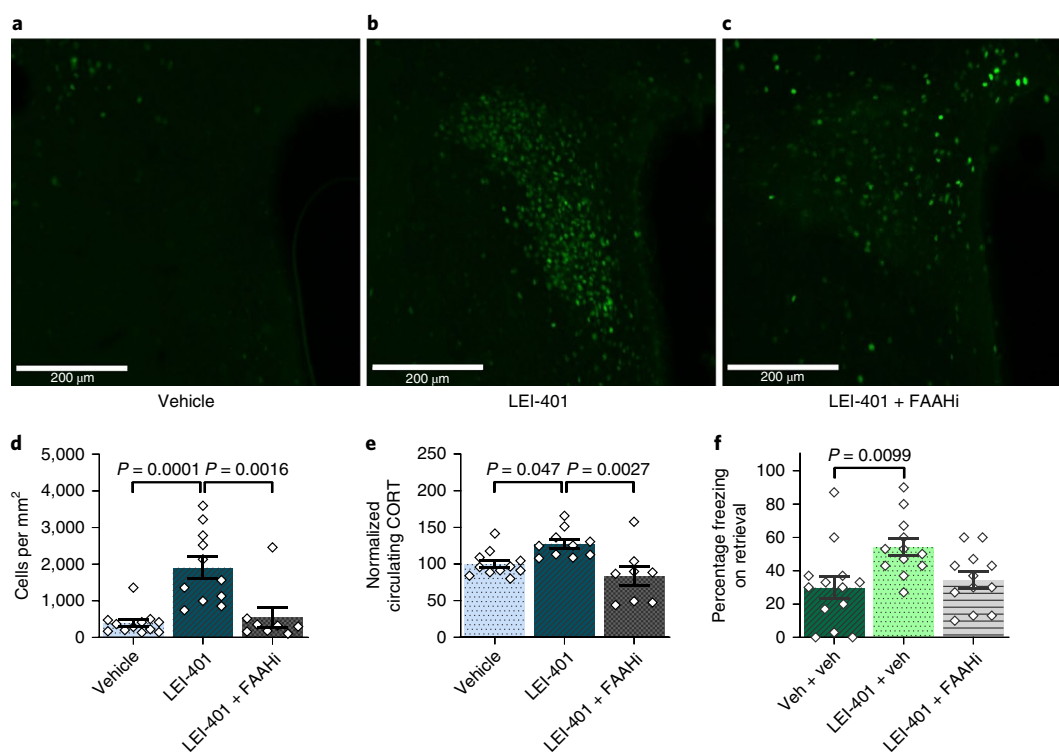


Fig. 6 | LEI-401 activates HPA axis signaling and diminishes fear extinction in vivo, which can be abolished by inhibition of FAAH. a–c, c-Fos labeling of brain sections of the PVN of the hypothalamus of mice treated with vehicle (**a**), LEI-401 (**b**) (i.p., 30 mg kg⁻¹, single dose) or LEI-401 with cotreatment of FAAH inhibitor URB597 (**c**) (i.p., 1 mg kg⁻¹, single dose) 210 min before perfusion. Scale bars, 200 μ m. **d**, Quantification of c-Fos labeled cells displayed a significant increase by LEI-401 and reversal by blockade of FAAH. For **a–d**, data represent mean values \pm s.e.m. with $n = 12$ mice for vehicle treatment, $n = 11$ mice for LEI-401 and $n = 8$ mice for LEI-401 + FAAHi. **e**, Circulating CORT levels were significantly elevated on LEI-401 treatment, but not when coadministered with the FAAH inhibitor URB597. Data represent mean values \pm s.e.m. with $n = 11$ mice for vehicle treatment, $n = 10$ mice for LEI-401 and $n = 8$ mice for LEI-401 + FAAHi. For **a–e**, P values were calculated using one-way ANOVA with Tukey's multiple comparisons test. **f**, Freezing during the fear extinction memory retrieval test 10 d after treatment with vehicle or LEI-401 (i.p., 30 mg kg⁻¹, single dose). LEI-401-treated mice exhibited increased freezing versus vehicle, but not when coadministered with FAAH inhibitor AM3506 (i.p., 1 mg kg⁻¹, single dose). Data represent mean values \pm s.e.m. with $n = 12$ mice for vehicle treatment and $n = 10$ mice for LEI-401 or LEI-401 + FAAHi. P values were calculated using one-way ANOVA with Dunnett's multiple comparisons test.

to investigate NAE biosynthesis in the brain of living animals in an acute and dynamic manner, thereby revealing a possible endogenous tone of this lipid family in emotional behavior. It is envisioned that LEI-401 may help to validate NAPE-PLD as a therapeutic target and serve as a starting point for the discovery of future drug candidates.

Online content

Any methods, additional references, Nature Research reporting summaries, source data, extended data, supplementary information, acknowledgements, peer review information; details of author contributions and competing interests; and statements of data and code availability are available at <https://doi.org/10.1038/s41589-020-0528-7>.

Received: 12 February 2019; Accepted: 27 March 2020;

Published online: 11 May 2020

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Methods

Cloning of plasmid DNA. Full length human cDNA of human or mouse *NAPEPLD* (obtained from N. Ueda⁹), *hDAGLα*, *mDagla*, *mDaglb*, *hMAGL* and *hPLA2G4E* was cloned into mammalian expression vector pcDNA3.1, containing a C-terminal FLAG-tag and genes for ampicillin and neomycin resistance. All plasmids were grown in XL-10 Z-competent cells and prepped (Maxi Prep, Qiagen). Constructs were verified by Sanger sequencing (Macrogen).

Cell culture. HEK293T or Neuro-2a cells (ATCC) were cultured at 37 °C and 7% CO₂ in DMEM (Sigma Aldrich, D6546) with GlutaMax, penicillin (100 µg ml⁻¹), streptomycin (100 µg ml⁻¹) and 10% New Born Calf serum. Cells were passaged twice a week to appropriate confluence by thorough pipetting.

Transient transfection. One day before transfection, 10⁷ cells were seeded on a 15 cm dish, and 2 h before transfection, the medium was refreshed with 13 ml medium. Transfection was performed with polyethylenimine (PEI, 60 µg per dish) in a ratio of 3:1 with plasmid DNA (20 µg per dish). The PEI and plasmid DNA were incubated in serum-free medium (2 ml per dish) at room temperature for 15 min, followed by dropwise addition to the cells. Transfection with the empty pcDNA3.1 vector was used to generate mock control samples. The medium was refreshed after 24 h and cells were collected after 48 or 72 h in cold PBS. Cells were centrifuged (10 min, 200g, 4 °C) and the supernatant was removed. The cell pellets were flash frozen in liquid N₂ and stored at -80 °C.

Cell lysate preparation. Cell pellets were resuspended in lysis buffer: 20 mM HEPES, 2 mM DTT, 0.25 M sucrose, 1 mM MgCl₂, 2.5 U ml⁻¹ benzonase and incubated 30 min on ice. The cytosolic fraction (supernatant) was separated from the membranes by ultracentrifugation (30 min, 100,000g, 4 °C). The pellet was resuspended in storage buffer comprising 20 mM HEPES and 2 mM DTT (membrane fraction). All samples were stored at -80 °C. Protein concentrations were determined using a Bradford assay (Bio-Rad).

Mouse brain lysate preparation. Mouse brains were thawed on ice, dounce homogenized in cold lysis buffer (20 mM HEPES pH 7.2, 2 mM DTT, 1 mM MgCl₂, 2.5 U ml⁻¹ benzonase) and incubated on ice for (30 min). The membrane and cytosolic fractions of cell or tissue lysates were separated by ultracentrifugation (45 min, 100,000g, 4 °C). The supernatant was collected (cytosolic fraction) and the membrane pellet was resuspended in cold storage buffer (20 mM HEPES pH 7.2, 2 mM DTT) by thorough pipetting and passage through a 1-ml insulin needle. Protein concentrations were determined using a Bradford assay (Bio-Rad). All samples were stored at -80 °C.

Western blot. Cell lysates were denatured with 4× Laemmli buffer (stock concentrations: 240 mM Tris-HCl pH 6.8, 8% w/v SDS, 40% v/v glycerol, 5% v/v β-mercaptoethanol, 0.04% v/v bromophenol blue, 30 min, room temperature) and 10–20 µg per sample was resolved on a 10% acrylamide SDS-PAGE gel (180 V, 75 min). Proteins were transferred from the gel to a 0.2-µm polyvinylidene difluoride membrane using a Trans-Blot Turbo (Bio-Rad). The membranes were washed with tris-buffered saline (TBS) (50 mM Tris-HCl pH 7.5, 150 mM NaCl) and blocked with 5% milk in TBS and Tween 20 (TBST) (50 mM Tris-HCl pH 7.5, 150 mM NaCl, 0.05% Tween 20) 1 h at room temperature or overnight at 4 °C. Primary antibodies against NAPE-PLD (Abcam, ab95397, 1:200, in TBST), FLAG-tagged proteins (Sigma Aldrich, F3165, 1:5,000 in 5% milk in TBST), α-tubulin (Genetex, GTX76511, 1:5,000 in 5% milk in TBST) or β-actin (Abcam, ab8226, 1:5,000 in 5% milk in TBST) were incubated 1 h at room temperature or overnight at 4 °C. Membranes were washed with TBST and incubated with secondary antibodies: for NAPE-PLD, goat-antirabbit-HRP (Santa Cruz, sc-2030, 1:2,000 in 5% milk in TBST); for FLAG-tagged proteins and β-actin, goat-antimouse-HRP (Santa Cruz, sc-2005, 1:5,000 in 5% milk in TBST); for α-tubulin, goat-antirat (Santa Cruz, sc-2032, 1:5,000 in 5% milk in TBST). All secondary antibodies were incubated 1 h at room temperature or overnight at 4 °C. Membranes were washed with TBST and TBS. The blot was developed in the dark using a luminol solution (10 ml, 1.4 mM luminol in Tris-HCl pH 7.5), ECL enhancer (100 µl, 6.7 mM para-hydroxycoumaric acid in DMSO) and H₂O₂ (3 µl, 30% w/w in H₂O). Chemiluminescence was visualized using a ChemiDoc Imaging System (Bio-Rad). Band intensity is normalized to α-tubulin or β-actin using ImageLab software (Bio-Rad).

qPCR. RNA isolation and cDNA synthesis was conducted as follows: total RNA from Neuro-2a cells was extracted using Trizol reagent (ThermoFisher Scientific). Once isolated, RNA concentration and purity were determined by Nanodrop (Denovix Inc.). Subsequently, cDNA synthesis was carried out with Maxima H minus first strand cDNA synthesis kit (ThermoFisher Scientific) according to the manufacturer's instructions.

qPCR analysis was conducted as follows: 10 ng of input cDNA was analyzed using SybrGreen qPCR master mix (Biotool) in combination with CFX96 optical thermal cycler (Bio-Rad). Data analysis was performed using CFX Manager software (Bio-Rad). Data are expressed in quantitation cycles (C_q) and standard deviation of three technical replicates.

Activity assays. Surrogate substrate-based fluorescence assay NAPE-PLD. The NAPE-PLD activity assay was based on a previously reported method with small alterations³⁰. The membrane fraction from transient overexpression of human or mouse NAPE-PLD in HEK293T cells was diluted to 0.4 µg µl⁻¹ in assay buffer (50 mM Tris-HCl pH 7.5, 150 mM NaCl, 0.02% v/v Triton X-100). The substrate PED6 (Invitrogen, D23739) 1 mM stock in DMSO was consecutively diluted in DMSO (10×) and in assay buffer (10×) to make a 10 µM working solution. Relevant concentrations of compound (100× working solution) were prepared in DMSO. The assay was performed in a black Greiner 96-well plate (flat bottom), final volume 100 µl. The compound or DMSO was incubated with membrane protein lysate (final concentration 0.04 µg µl⁻¹) for 30 min at 37 °C. Then, substrate PED6 was added (final concentration 1 µM) and the measurement was started immediately on a TECAN infinite M1000 pro at 37 °C (excitation, 485 nm; emission, 535 nm; gain, 100), scanning every 2 min for 1 h. Mock membrane lysate with DMSO was used for background subtraction. The slope of $t = 4$ min to $t = 14$ min was used as the enzymatic rate (relative fluorescence units (RFU per min)), which was normalized to DMSO as 100%. IC₅₀ curves were generated using Graphpad Prism v.6 (log (inhibitor) versus normalized response with variable slope). K_i values were calculated from the Cheng-Prusoff equation $K_i = IC_{50}/(1 + ((S)/K_M))$ where S = [substrate] and K_M = 0.59 µM for both mouse and human NAPE-PLD. All measurements were performed in $n = 4$ or $n = 8$ for controls, with $Z' \geq 0.6$.

HTS of NAPE-PLD activity assay. Details of the HTS of the NAPE-PLD activity assay are listed in Supplementary Table 1.

LC-MS-based endogenous NAPE-PLD activity assay. The NAPE-PLD assay to measure endogenous enzyme activity was performed as reported previously with small alterations using the synthetic substrate 1,2-dioleoyl-sn-glycero-3-phosphoethanolamine-N-heptadecanoyl (C17:0-NAPE (19), for preparation see Supplementary Notes²¹). Mouse brain membrane lysates from male and female WT or NAPE-PLD KO mice (Virginia Polytechnic Institute and State University) were used originally generated by Leung et al.¹³. In an Eppendorf tube, per sample 10 µg of mouse brain membrane lysate (5 mg ml⁻¹) was diluted in assay buffer (50 mM Tris-HCl pH 7.5 + 0.1% Triton X-100) to a final volume of 47 µl. As a negative control, a sample was denatured at 100 °C for 10 min and cooled to room temperature for 5 min. LEI-401 (0.5 µl, 100× in DMSO) or DMSO was added and incubated for 30 min at room temperature. C17:0-NAPE (10 mM stock in 9:1 CHCl₃:MeOH) was diluted in EtOH to 10 µM and then added (2.5 µl, 20×, 0.5 µM final concentration). The samples were incubated at 37 °C for 15 min. Next, 10 µl internal standard OEA-d4 (100 nM) was added, followed by 50 µl NH₄OAc buffer (0.1 M, pH 4) and 500 µl MTBE. The samples were vortexed using a Bullet Blender (4 min, Next Advance Inc.) and centrifuged for 11 min at 16,000g at 4 °C. Then, 475 ml of the supernatant was transferred to a new Eppendorf tube and evaporated to dryness. Reconstitution in 200 µl acetonitrile, vortex (3 min) and centrifugation (13,000g, 2 min). Following that, 50 µl was transferred to an LC-MS vial with a micro insert. Next, 5 µl was injected in the LC-MS/MS system. A targeted analysis of C17:0 NAE was measured using an Acquity UPLC I class binary solvent manager pump in conjugation with a Xevo-TQ-S tandem quadrupole mass spectrometer (Waters Corporation). Separation of C17:0-NAE and OEA-d4 (internal standard) was performed with an Acquity HSS-T3 column (2.1 × 100 mm², 1.8 µm particle size; Waters Corporation) maintained at 45 °C. The aqueous mobile phase A consisted of 2 mM ammonium formate and 10 mM formic acid, and the organic mobile phase B was acetonitrile. The flow rate was set to 0.6 ml min⁻¹; initial gradient conditions were 70% B held for 0.50 min and linearly ramped to 100% B over 1.5 min and held for 1 min; after 10 s the system returned to initial conditions and held for 2 min before the next injection. Electrospray ionization-MS was operated in positive mode for measurement of C17:0-NAE, and a selective multiple reaction mode was used for quantification. Data were analyzed with MassLynx v.4.1 software (Waters Corporation).

Natural substrate-based fluorescence assay DAGLα. The natural substrate assay for human and mouse DAGLα was performed as reported previously³¹. All measurements were performed in $n = 4$ or $n = 8$ for controls, with $Z' \geq 0.6$.

Surrogate substrate-based fluorescence assay DAGLβ. The biochemical mDAGLβ assay was performed as reported previously³². All measurements were performed in $n = 4$ or $n = 8$ for controls, with $Z' \geq 0.6$.

Natural substrate-based fluorescence assay MAGL. The natural substrate assay for human MAGL was performed as reported previously⁵³. All measurements were performed in $n = 4$ or $n = 8$ for controls, with $Z' \geq 0.6$.

Radioligand displacement assays CB₁ and CB₂ receptor. [³H]CP55940 displacement assays to determine the affinity for the cannabinoid CB₁ and CB₂ were performed as previously described³⁴.

Activity-based protein profiling (ABPP) for FAAH, ABHD6, ABHD12 and PLA2G4E activity. Gel-based ABPP was performed as previously described⁵². In brief, mouse brain membrane proteome or hPLA2G4E overexpressing membrane lysate

(9.5 μl and 2 $\mu\text{g}\mu\text{l}^{-1}$ or 19.5 μl and 1 $\mu\text{g}\mu\text{l}^{-1}$, respectively) was pre-incubated with vehicle or inhibitor (0.5 μl 20 \times or 0.5 μl 40 \times inhibitor stock in DMSO, respectively, 30 min, room temperature) followed by incubation with the activity-based probe MB064 (250 nM, 0.5 μl 20 \times stock in DMSO) or FP-TAMRA (500 nM, 0.5 μl 20 \times stock in DMSO) for mouse brain lysate (15 min, room temperature) or FP-TAMRA (50 nM, 0.5 μl 40 \times stock in DMSO) for PLA2G4E overexpressing lysate (5 min, room temperature). Final concentrations for the inhibitors are indicated in the main text and figure legends. Proteins were denatured with 4 \times Laemmli buffer (3.5 μl , stock concentrations: 240 mM Tris-HCl pH 6.8, 8% w/v SDS, 40% v/v glycerol, 5% v/v β -mercaptoethanol, 0.04% v/v bromophenol blue, 30 min, room temperature). The samples (10 μl per slot) were resolved by SDS-PAGE (respectively, 10 or 8% acrylamide for mouse brain or PLA2G4E lysate, 180 V, 75 min). Gels were scanned using Cy3 and Cy5 multichannel settings (605/50 and 695/55 filters, respectively) on a ChemiDoc Imaging System (Bio-Rad). Fluorescence was normalized to Coomassie staining and quantified with ImageLab (Bio-Rad).

Generation of Neuro-2a NAPE-PLD KO populations. *Single-guide RNA (sgRNA) selection.* Two sgRNAs, located in exon 2 and 3 with high efficiency and specificity as predicted by CHOPCHOP v.2 online web tool (<http://chopchop.cbu.uib.no>) were selected. Guides were cloned into the BbsI restriction site of plasmid px330-U6-Chimeric_BB-CBh-hSpCas9 (gift from F. Zhang, Addgene plasmid no. 42230). The sgRNA targeting exon 3 yielded higher knockout efficiency as determined by western blot and in the T7E1-assays. This population was used for further experiments.

Sequential transfection. Neuro-2a cells were transfected sequentially (three times within the course of 10 d) to yield populations with a high knockout efficiency. Cells were seeded at day 0, day 3 and day 6 and transfected at day 1, day 4 and day 7. Samples for T7E1 assays were collected at day 2, day 5, day 11 and after several weeks of culturing the cells.

One day before the first transfection, Neuro-2a cells were seeded to a six-well plate to reach 80–90% confluence at the time of transfection. Before transfection, culture medium was aspirated and 2 ml of fresh medium was added. A 5:1 (m/m) mixture of PEI (17.5 μg per well) and plasmid DNA (3.5 μg per well) was prepared in serum-free culture medium (250 μl each) and incubated (15 min, room temperature). Transfection was performed by dropwise addition of the PEI/DNA mixture to the cells. Then, 24 h posttransfection, the culture medium was refreshed, a small number of cells were collected for analysis by T7E1 assay and ABPP while the remainder was kept in culture under standard conditions for following transfections. After three transfection rounds, the cells were cultured according to standard protocol.

T7E1 assay. Genomic DNA was obtained by mixing 50 μl QuickExtract (Epicentre) with cell pellet (~10% of a well, from a six-well plate). The samples were incubated at 65 °C for 6 min, mixed by vortexing and then incubated at 98 °C for 2 min. Genomic DNA extracts were diluted in sterile water and directly used in PCR reactions. Genomic PCR reactions were performed on 2.5–5 μl isolated genomic DNA extract using Phusion High-Fidelity DNA Polymerase (ThermoFisher) in Phusion HF buffer in a final volume of 20 μl , for primers see Supplementary Table 7.

For the T7E1 assay, genomic PCR products were denatured and reannealed in a thermocycler using the following program: 5 min at 95 °C, 95 to 85 °C using a ramp rate of $-2\text{ }^{\circ}\text{C s}^{-1}$, 85–25 °C using a ramp rate of $-0.2\text{ }^{\circ}\text{C s}^{-1}$. To annealed PCR product (8.5 μl) was added NEB2 buffer (1 μl) and T7 endonuclease I (5 U, 0.5 μl ; New England Biolabs), followed by incubation at 37 °C for 30 min. Digested PCR products were analyzed using agarose gel electrophoresis with ethidium bromide staining.

Agarose gels were analyzed using ImageLab Software (Bio-Rad) and CRISPR gene editing efficiency was expressed as percentage T7E cleavage (volume integral of digested bands/volume integral all bands \times 100%).

Live cell photoaffinity labeling. *Gel-based photoaffinity labeling.* Here, 500,000 HEK293T cells per well were seeded in a 12-well plate 1 d before transfection. On the day of transfection, the medium was refreshed with 0.375 ml medium per well. Transfection was performed with PEI (3 μg per well) and human NAPEPLD-FLAG or mock pcDNA3.1Neo plasmids (1 μg per well). PEI and plasmids were combined in serum-free medium (0.125 ml per well) and incubated for 15 min at room temperature, then added to each well. After 24 h the medium was refreshed. Then, 48 h after transfection, the medium was removed and cells were washed with warm PBS (1 \times). This was followed by treatment with photoprobe 10 (1,000 \times in DMSO, final concentration: 2 μM) together with vehicle or competitor (600 \times in DMSO) in medium + serum (0.30 ml per well) for 30 min at 37 °C. The medium was aspirated and the cells were covered with PBS (0.15 ml per well), followed by 350 nm UV irradiation using a Caprobox at 4 °C for 10 min. The cells were collected into 1.5 ml epps with cold PBS and centrifuged for 10 min at 2,000 r.p.m. at 4 °C. The PBS was removed and the cells were flash frozen with liquid N₂ (cells can optionally be stored at $-80\text{ }^{\circ}\text{C}$). The cells were lysed with lysis buffer (30 μl , 20 mM HEPES pH 7.2, 0.25 M sucrose, 1 mM MgCl₂, benzonase 25 U ml⁻¹) followed by pipetting up and down and incubating for 30 min on ice. Protein concentrations were measured using a Bradford assay (Bio-Rad), and cell lysates were diluted to

2 $\mu\text{g}\mu\text{l}^{-1}$ with lysis buffer. Next, 18 μg cell lysate (9 μl) was then clicked with Cy5-N₃ (for molecular structure see Supplementary Fig. 2) using a click mix (1 μl per sample, final concentrations: 1 mM CuSO₄, 6 mM sodium ascorbate, 0.2 mM tris (3-hydroxypropyltriethylammonium) (THPTA), 2 μM Cy5-N₃) for 1 h at room temperature (note that it is important to separately prepare the click mix first with CuSO₄ and sodium ascorbate, until a yellow color change is observed). Samples were denatured with 4 \times Laemmli buffer (3.33 μl , stock concentrations: 240 mM Tris-HCl pH 6.8, 8% w/v SDS, 40% v/v glycerol, 5% v/v β -mercaptoethanol, 0.04% v/v bromophenol blue) and incubated for 30 min at room temperature. The samples were resolved by SDS-PAGE (10% acrylamide gel) at 180 V for 75 min, after which the gels were imaged at Cy3 and Cy5 channels (605/50 and 695/55 filters, respectively) on a ChemiDoc Imaging System (Bio-Rad). Fluorescence is normalized to Coomassie or α -tubulin by western blot using ImageLab software (Bio-Rad). IC₅₀ curves were generated using Graphpad Prism v.6.

Chemical proteomics-based photoaffinity labeling. Here, 2 \times 10⁶ HEK293T cells per well were seeded in a six-well plate 1 d before transfection. On the day of transfection, the medium was removed and refreshed with 1.125 ml fresh medium per well. Transfection was performed with PEI (9 μg per well) and human NAPEPLD-FLAG or mock pcDNA3.1Neo plasmids (3 μg per well). PEI and plasmids were combined in serum-free medium (0.375 ml per well) and incubated for 15 min at room temperature, then added to each well. After 24 h the medium was refreshed. Next, 48 h after transfection, the medium was removed and cells were washed with warm PBS (1 \times). This was followed by treatment with photoprobe 10 (1,000 \times in DMSO, final concentration of 2 μM) together with vehicle or competitor (1,000 \times in DMSO, final concentration 20 μM) in medium + serum (1 ml per well) for 30 min at 37 °C. The medium was aspirated and the cells were covered with PBS (0.5 ml per well), followed by 350 nm UV irradiation using a Caprobox at 4 °C for 10 min. The cells were collected into 1.5 ml epps with cold PBS and centrifuged for 10 min at 2,000 r.p.m. at 4 °C. The PBS was removed and the cells were flash frozen with liquid N₂ (cells can optionally be stored at $-80\text{ }^{\circ}\text{C}$). The cells were lysed with lysis buffer (30 μl , 20 mM HEPES pH 7.2, 0.25 M sucrose, 1 mM MgCl₂, benzonase 25 U ml⁻¹) followed by pipetting up and down and incubating for 30 min on ice. Protein concentrations were measured using a Bradford assay (Bio-Rad), and cell lysates were diluted to 1 $\mu\text{g}\mu\text{l}^{-1}$ with lysis buffer.

From here the label-free chemical proteomics protocol was followed as previously described³⁰. Then, 270 μg cell lysate (270 μl) was clicked with biotin-N₃ (Sigma Aldrich, 762024) using a click mix (30 μl per sample, final concentrations: 1 mM CuSO₄, 6 mM sodium ascorbate, 0.2 mM tris (3-hydroxypropyltriethylammonium) (THPTA), 4 μM biotin-N₃) for 1 h at room temperature (note that it is important to separately prepare the click mix first with CuSO₄ and sodium ascorbate, until a yellow color change is observed). Proteins were precipitated using chloroform (166 μl), methanol (666 μl) and MilliQ (366 μl H₂O). Samples were centrifuged for 10 min at 1,500g and the solvents were carefully removed. Methanol (600 μl) was added and the protein were resuspended using a probe sonicator (30% amplitude, 10 s). After centrifugation (5 min, 18,400g) the solvent was removed and the samples were resuspended again in urea buffer (250 μl , 6 M urea, 250 mM NH₄HCO₃) by pipetting up and down. DTT (2.5 μl , final concentration 10 mM) was added and the samples were incubated at 65 °C with shaking (600 r.p.m.) for 15 min. After cooling to room temperature, iodoacetamide (20 μl , final concentration 40 mM) was added and incubated in the dark at 20 °C with shaking (600 r.p.m.) for 30 min. Next, SDS (70 μl , 10%) was added and incubated at 65 °C with shaking (600 r.p.m.) for 5 min. For 18 samples, 1.8 ml of avidin agarose beads (Thermo Scientific, 20219) were divided over three 15 ml tubes and washed with PBS (3 \times 10 ml²). The beads in each tube were resuspended in PBS (6 ml) and divided over 18 tubes (1 ml each). To each tube was added the denatured sample and PBS (2 ml) and the tubes were rotated with an overhead shaker at room temperature for 3 h. After centrifugation (2 min, 2,500g) and removal of the supernatant, the beads were consecutively washed with 0.5% SDS in PBS (w/v, 6 ml) and PBS (3 \times 6 ml²), each time centrifuging (2 min, 2,500g). The beads were transferred to a 1.5-ml low binding epp (Sarstedt, 72.706.600) with on-bead digestion buffer (250 μl , 100 mM Tris pH 8.0, 100 mM NaCl, 1 mM CaCl₂, 2% v/v acetonitrile) and to each sample was added 1 μl trypsin solution (0.5 $\mu\text{g}\mu\text{l}^{-1}$ trypsin (Promega, V5111), 0.1 mM HCl). Proteins were digested at 37 °C with shaking (950 r.p.m.) overnight. Formic acid (12.5 μl) was added to each sample and the beads were filtered off using a biospin column (Bio-Rad, 7326204), the flow-through was collected in a 2-ml epp. Samples were purified using StageTips. Each StageTip was conditioned with MeOH (50 μl , centrifugation: 2 min, 300g), followed by StageTip solution B (50 μl , 80% v/v acetonitrile, 0.5% v/v formic acid in MilliQ, centrifugation for 2 min, 300g) and StageTip solution A (50 μl , 0.5% v/v formic acid in MilliQ, centrifugation for 2 min, 300g). Next, the samples were loaded, centrifuged (2 min, 600g) and the peptides on the StageTip were washed with StageTip solution A (100 μl , centrifugation for 2 min, 600g). The StageTips were transferred to a new 1.5 ml low binding epp and the peptides were eluted with StageTip solution B (100 μl , centrifugation for 2 min, 600g). The solvents were evaporated to dryness in a SpeedVac concentrator at 45 °C for 3 h. Samples were reconstituted in LC-MS solution (50 μl , 3% v/v acetonitrile, 0.1% v/v formic acid, 1 μM yeast enolase peptide digest (Waters, 186002325) in MilliQ).

The peptides were measured as described previously for the NanoACQUITY UPLC System coupled to SYNAPT G2-Si high definition mass spectrometer³⁰. A trap-elute protocol, where 5 μ l of the digest was loaded on a trap column (C18 100 \AA , 5 μ M, 180 μ M \times 20 mm, Waters) followed by elution and separation on the analytical column (HSS-T3 C18 1.8 μ M, 75 μ M \times 250 mm, Waters). The sample was brought onto this column at a flow rate of 10 μ l min⁻¹ with 99.5% solvent A for 2 min before switching to the analytical column. Peptide separation was achieved using a multistep concave gradient based on gradients previously described³⁰. The column was re-equilibrated to initial conditions after washing with 90% solvent B. The rear seals of the pump were flushed every 30 min with 10% (v/v) acetonitrile. [Glu1]-fibrinopeptide B (GluFib) was used as a lock mass compound. The auxiliary pump of the LC system was used to deliver this peptide to the reference sprayer (0.2 μ l min⁻¹). A UDMSe method was set up as previously described³⁰. Briefly, the mass range was set from 50 to 2,000 Da with a scan time of 0.6 s in positive, resolution mode. The collision energy was set to 4 V in the trap cell for low-energy MS mode. For the elevated energy scan, the transfer cell collision energy was ramped using drift-time specific collision energies. The lock mass was sampled every 30 s. For raw data processing, PLGS (v.3.0.3) was used. The MS^F identification workflow was performed with the parameters summarized in Supplementary Table 7 to search the human proteome from Uniprot (uniprot-homo-sapiens-trypsin-reviewed-2016_08_29.fasta). Protein quantification was performed using ISOQuant (v.1.5). The parameter settings used are summarized in Supplementary Table 7.

Data analysis. Protein targets of photoprobe **10** were selected based on the following cut-offs: (1) ≥ 2 -fold enrichment for UV-treated versus non-UV-treated samples; (2) unique peptides ≥ 2 ; (3) testing for significance using Student's *t*-test (unpaired, two-tailed), $P < 0.05$ was considered significant and (4) Benjamini–Hochberg correction with a false discovery rate (FDR) of 10%.

Targeted lipidomics in Neuro-2a cells. The targeted lipidomics experiments were based on previously reported methods with small alterations⁵⁵.

Sample preparation. Here, 2×10^6 Neuro-2a cells (grown at 37 °C, 7% CO₂) were seeded 1 d before treatment in 6 cm dishes. Before treatment, cells were washed twice with warm PBS and then treated with vehicle or compound (10 μ M, 0.1% DMSO) in 3 ml medium without serum (2 h at 37 °C, $n = 5$ per condition). Washing with cold PBS (1 \times) followed by gathering in 1.5 ml Eppendorf tubes and centrifugation (10 min, 1,500 r.p.m.). PBS was removed and the cell pellets were flash frozen with liquid nitrogen and stored at -80 °C. Live cell count with trypan blue was performed after compound treatment to test for cell viability. Next, 10% of each cell sample (collected during gathering) was used to determine the protein concentration using a Bradford assay (Bio-Rad) for normalization after lipid measurements.

Statistical analysis. Absolute values of lipid levels were corrected using the measured protein concentrations. Data were tested for significance with GraphPad v.6 using one-way analysis of variance (ANOVA) with Tukey correction for multiple comparisons. $P < 0.05$ was considered significant.

In vitro ADME profile. Kinetic aqueous solubility (lyophilization solubility assay). The solubility of LEI-401 in phosphate buffer at pH 6.5 from an evaporated 10 mM DMSO compound stock solution was measured. Two aliquots of the test compound were dried and dissolved in phosphate buffer at pH 6.5. The solutions were then filtered and diluted (three different dilution levels for each compound) before RapidFire MS analysis was performed. Each test compound was quantified using a six-point calibration curve prepared with the same DMSO starting solution.

Passive membrane permeability through PAMPA. PAMPA is a method that determines the permeability of substances from a donor compartment, through a lipid-infused artificial membrane into an acceptor compartment. Read-out is a permeation coefficient P_{eff} drug as well as test compound concentrations in donor, membrane and acceptor compartments. The assay was performed as described elsewhere⁵⁴.

Microsomal clearance. The microsomal clearance assay was performed as previously described⁵⁴. Pooled commercially available microsomes from male mouse microsomes (C57BL/6J, Lot 4339006) were purchased from Corning Inc.). For human, ultrapooled liver microsomes (150 mixed gender donors, BD UltraPool HLM 150, Lot 38289) were purchased to account for the biological variance in vivo from human liver tissues. For the microsomes incubations (incubation buffer 0.1 M phosphate buffer pH 7.4), 96-deep well plates were applied, which were incubated at 37 °C on a TECAN (Tecan Group Ltd) equipped with Te-Shake shakers and a warming device (Tecan Group Ltd). The reduced nicotinamide adenine dinucleotide phosphate regenerating system consisted of 30 mM glucose-6-phosphate disodium salt hydrate; 10 mM NADP; 30 mM MgCl₂ \times 6H₂O and 5 μ g μ l⁻¹ glucose-6-phosphate dehydrogenase (Roche Diagnostics) in 0.1 M potassium phosphate buffer pH 7.4. Incubations of LEI-401

at 1 μ M in microsomes incubations of 0.5 μ g μ l⁻¹ plus cofactor reduced nicotinamide adenine dinucleotide phosphate were performed in 96-well plates at 37 °C. After 1, 3, 6, 9, 15, 25, 35 and 45 min 40 μ l incubation solutions were transferred and quenched with 3:1 (v/v) acetonitrile containing internal standards. Samples were then cooled and centrifuged before analysis by LC-MS/MS. The log peak area ratios (test compound peak area/internal standard peak area) were plotted against incubation time using a linear fit. The calculated slope was used to determine the intrinsic clearance: $CL_{int} (\mu\text{l min}^{-1} \text{mg}^{-1} \text{protein}) = -\text{slope} (\text{min}^{-1}) \times 1,000 / (\text{protein concentration} (\mu\text{g} \mu\text{l}^{-1}))$.

Hepatocyte clearance. The hepatocyte clearance assay was performed as previously described⁵⁴. For animals, hepatocyte suspension cultures were either freshly prepared by liver perfusion studies or prepared from cryopreserved hepatocyte batches (pooled C57BL6 mouse hepatocytes were purchased from BioreclamationIVT). For human, commercially available, pooled (5–20 donors), cryopreserved human hepatocytes from nontransplantable liver tissues were used. For the suspension cultures, Nunc U96 PP-0.5 ml (Nunc Natural, 267245) plates were used, which were incubated in a Thermo Forma incubator from Fischer Scientific equipped with shakers from Variomag Teleshake shakers (Sterico) for maintaining cell dispersion. The cell culture medium was William's media supplemented with Glutamine, antibiotics, insulin, dexamethasone and 10% FCS. Incubations of a test compound at 1 μ M test concentration in suspension cultures of 1×10^6 cells per ml ($\sim 1 \mu\text{g} \mu\text{l}^{-1}$ protein concentration) were performed in 96-well plates and shaken at 900 r.p.m. for up to 2 h in a 5% CO₂ atmosphere at 37 °C. After 3, 6, 10, 20, 40, 60 and 120 min 100 μ l cell suspension in each well was quenched with 200 μ l methanol containing an internal standard. Samples were then cooled and centrifuged before analysis by LC-MS/MS. The log peak area ratios (test compound peak area/internal standard peak area) or concentrations were plotted against incubation time with a linear fit. The slope of the fit was used to calculate the intrinsic clearance: $CL_{int} (\mu\text{l min}^{-1} / 1 \times 10^6 \text{ cells}) = -\text{slope} (\text{min}^{-1}) \times 1,000 / (1 \times 10^6 \text{ cells})$.

Plasma protein binding. The plasma protein binding assay was performed as previously described⁵⁴. Pooled and frozen plasma from human (HMPLEDTA, lot BRH1060627) and mouse (MSEPLEDTA3-C57, lot MSE196204) were obtained from BioreclamationIVT. The Teflon equilibrium dialysis plate (96-well, 150 μ l, half-cell capacity) and cellulose membranes (12–14 kDa molecular weight cut-off) were purchased from HT-Dialysis (Gales Ferry). Both biological matrix and phosphate buffer pH were adjusted to 7.4 on the day of the experiment. The determination of unbound compound was performed using a 96-well format equilibrium dialysis device with a molecular weight cut-off membrane of 12–14 kDa. The equilibrium dialysis device itself was made of Teflon to minimize nonspecific binding of the test substance. Compounds were tested in cassettes of 2–5 with an initial total concentration of 1,000 nM, one of the cassette compounds being the positive control diazepam. Equal volumes of matrix samples containing substances and blank dialysis buffer (Soerensen buffer at pH 7.4) were loaded into the opposite compartments of each well. The dialysis block was sealed and kept for 5 h at a temperature of 37 °C and 5% CO₂ environment in an incubator. After this time, equilibrium has been reached for most small molecule compounds with a molecular weight of <600. The seal was then removed and matrix and buffer from each dialysis was prepared for analysis by LC-MS/MS. All protein binding determinations were performed in triplicate. The integrity of membranes was tested in the HT-Dialysis device by determining the unbound fraction values for the positive control diazepam in each well. At equilibrium, the unbound drug concentration in the biological matrix compartment of the equilibrium dialysis apparatus was the same as the concentration of the compound in the buffer compartment. Thus, the percentage unbound fraction (f_u) was calculated by determining the compound concentrations in the buffer and matrix compartments after dialysis as follows: $\%f_u = 100 \times \text{buffer concentration after dialysis} / \text{matrix concentration after dialysis}$. The device recovery was checked by measuring the compound concentrations in the matrix before dialysis and calculating the percent recovery (mass balance). The recovery must be within 80–120% for data acceptance.

P-glycoprotein active efflux assay. P-glycoprotein (permeability-glycoprotein, abbreviated to 'P-gp' and also known as multidrug resistance protein 1 (MDR1)) is the most-studied and best-characterized drug transporter. The P-gp assay evaluates the ability of test compounds to serve as a P-gp substrate. The assay was performed as described elsewhere⁵⁴.

logD-assay. For the determination of the octanol/water distribution coefficient (logD), the carrier-mediated distribution system-assay was used as described elsewhere⁵⁴.

In vivo pharmacokinetics and lipidomics. Subjects. Male C57BL/6J mice (Jackson Laboratory) or male and female NAPE-PLD KO mice originally from a mixed 129SvJ-C57BL/6J background (Virginia Polytechnic Institute and State University) generated by Leung et al.¹⁵ served as subjects for the in vivo pharmacokinetic and brain lipid analysis of LEI-401. Animal protocols were approved by

the Institutional Animal Care and Use Committee at National Institute on Alcohol Abuse and Alcoholism, National Institute of Health, the Bundesamt für Lebensmittelsicherheit und Veterinärwesen (BLV) der Schweizerischen Eidgenossenschaft, the Dierexperimentencommissie Universiteit Leiden and by the Institutional Animal Care and Use Committee at Virginia Commonwealth University in accordance with the National Institutes of Health Guide for the Care and Use of Laboratory Animals⁵⁶. All studies involving animals are reported in accordance with the Animal Research: Reporting of In Vivo Experiments guidelines for reporting experiments involving animals⁵⁷.

Pharmacokinetic analysis of LEI-401 (mouse plasma). Test compounds were formulated according to respective protocols either by dissolution (intravenous, i.v.) or in a glass potter until homogeneity was achieved (oral (p.o.) and i.p.). Formulations were injected i.v. using a 30G needle in the lateral tail vein mice yielding a 1 mg kg⁻¹ dose. For p.o. applications, animals were gavaged (yielding a 10 mg kg⁻¹ dose) and for i.p. applications the compounds were injected into the abdomen (yielding a 30 mg kg⁻¹ dose). At the following time points blood was drawn into EDTA: 0.083, 0.25, 0.5, 1, 2, 4, 7 and 24 h (for p.o. and i.p. the first time point was omitted). Three animals each were used for the i.v. and p.o. arm and six animals were used in the i.p. arm. Animals were distributed randomly over the time course and at each time point, a volume of 100 µl of blood was taken. Quantitative plasma measurement of the compound was performed by LC-MS/MS analysis. Pharmacokinetic analysis was conducted using Phoenix WinNonlin v.6.4 software using a noncompartmental approach consistent with the route of administration. For assessment of the exposure C_{max} , T_{max} , area under curve and bioavailability were determined from the plasma concentration profiles. Parameters (CL, V_{ss} , $t_{1/2}$) were estimated using nominal sampling times relative to the start of each administration.

Pharmacokinetic analysis of LEI-401 (mouse brain). Mice were injected with LEI-401 (30 mg kg⁻¹, i.p., single dose) dissolved in sterile DMSO (D2650 Sigma) and mixed with Tween-80 and distilled H₂O at a ratio of 1:1:8. Mice were killed after 1, 2, 4 and 8 h. Brain samples were rapidly collected, washed in ice cold saline, blotted dry, frozen in liquid N₂ and stored at -80 °C until further analysis. Brain levels of LEI-401 were quantified by LC-MS/MS using an Agilent 6410 triple quadrupole mass spectrometer (Agilent Technologies) coupled with an Agilent 1200 LC system (Agilent Technologies). Chromatographic and mass spectrometer conditions were set as described previously⁵⁸. Chromatographic separation was obtained using 2.5-µl samples injected onto a Zorbax SB-C18 rapid resolution HT column (2.1 × 50 mm², 1.8 µm, Agilent). The mass spectrometer was set for electrospray ionization in positive ion mode. The source parameters were as follows: capillary voltage, 4,000 V; gas temperature, 350 °C; drying gas, 10 l min⁻¹; nitrogen was used as the nebulizing gas with a pressure of 40 psig. Collision-induced dissociation energy was performed by using nitrogen. Levels of LEI-401 were analyzed by multiple reactions monitoring. Mass spectrometric conditions were optimized for LEI-401 with injection of synthetic standard by using MassHunter Workstation Optimizer software (Agilent Technologies). The molecular ion and fragments for LEI-401 measured were as follows: m/z 422.26 → 70.1 and 422.26 → 137 (collision-induced dissociation energy, 56 and 40 V, respectively). The amounts of LEI-401 in the samples were determined against standard curves. Values are expressed as ng g⁻¹ in wet brain tissue weight.

Lipid measurements in mouse brain. Levels of NAEs, 2-AG and arachidonic acid were measured by stable isotope dilution LC-MS/MS as described previously⁵⁹.

In vitro ABPP of LEI-401-treated mice. Male C57BL/6J mice were treated with LEI-401 (30 mg kg⁻¹, i.p., single dose) or vehicle (DMSO, Tween-80 and distilled H₂O; 1:1:8) and killed after 2 h by cervical dislocation. Brain samples were rapidly collected, frozen in liquid N₂ and stored at -80 °C until further analysis. After membrane lysate preparation (vide supra), 19.5 µl lysate (2 mg ml⁻¹) per sample was treated with 0.5 µl probe (40× stock in DMSO, final concentration 250 nM for MB064, 500 nM for fluorescence polarization-boron dipyrromethene (BODIPY) (1 µM for DH379) and incubated for 15 min at room temperature or 30 min at 37 °C for DH379. Proteins were denatured with 4× Laemmli buffer (7.5 µl, stock concentrations: 240 mM Tris-HCl pH 6.8, 8% w/v SDS, 40% v/v glycerol, 5% v/v β-mercaptoethanol, 0.04% v/v bromophenol blue, 30 min, room temperature). From the DH379-labeled samples 10 µl (14 µg protein) of quenched reaction mixture was resolved on 10% acrylamide SDS-PAGE (180 V, 75 min). The MB064- and FP-BODIPY-labeled samples were combined (1:1 v/v) and resolved together on 10% acrylamide SDS-PAGE (180 V, 75 min). Fluorescence was measured using a Bio-Rad ChemiDoc MP system (fluorescence channel filters BODIPY, Cy2 (460–490 nm); BODIPY-TMR dye, Cy3 (520–545 nm); protein ladder, Cy5 (625–650 nm)). Gels were then stained using Coomassie staining and imaged for protein loading control. Fluorescence quantification was performed using ImageLab (Bio-Rad). Intensities were normalized to the DMSO control and corrected for protein loading by Coomassie staining.

Statistical analysis. Data were tested for significance with GraphPad v.6 using one-way ANOVA with Dunnett's correction for multiple comparisons. Two-way

ANOVA was used to evaluate LEI-401 versus vehicle in NAPE-PLD KO versus WT mice. $P < 0.05$ was considered significant.

HPA axis activation. Pharmacological treatment and c-Fos labeling. All drugs were dissolved in 5% DMSO, 5% Tween-80, and 90% saline the day of each experiment. The CB₁ antagonist AM251 (3 mg kg⁻¹; Cayman Chemical), FAAH inhibitor URB597 (1 mg kg⁻¹; Cayman Chemical) and the NAPE-PLD inhibitor LEI-410 (30 mg kg⁻¹) were administered intraperitoneally (i.p., 10 ml kg⁻¹).

All animals (C57BL/6J mice) received 4 d of handling before test days. On test day, animals received drug injections 105 (AM251/vehicle) and 210 min (URB597/LEI-410/vehicle) before perfusions. Animals were deeply anesthetized using pentobarbital injection and perfused intracardially with 4% paraformaldehyde and PBS. After perfusion, the brains were extracted and fixed in 4% paraformaldehyde at 4 °C overnight and then submerged in 30% sucrose in PBS at 4 °C for 48 h. Using a mouse brain atlas, brains were mounted with optimal cutting temperature compound and sectioned (40 µm) on a cryostat. Slices containing the PVN were placed in PBS until antibody treatment.

Sections were washed three times (10 min per wash) in a PBS and 0.1% Triton X-100 (PBST) solution before incubation in PBS blocking buffer containing 10% normal donkey serum at room temperature for 1 h. Sections were incubated in primary antibody rabbit anti-c-Fos (1:300 Cell Signaling) at 4 °C overnight. Tissue was then washed in PBST solution three more times (10 min each) and then incubated in donkey antirabbit Cy3 (1:100 JacksonImmuno) at room temperature for 2 h. Sections were washed in PBST three more times, then mounted in PBS and examined using a Leica fluorescent microscope. Two slices containing the PVN were collected from each specimen. Grayscale images were taken of both hemispheres separately for each slice. All images within experiments were obtained under the same exposure and magnification (20×). Images were processed on ImageJ, where they were subjected to a brightness threshold before c-Fos labeled cells were counted based on size criteria set for each experiment. Number of labeled cells inside a defined region (24,614.47 µm²) that met the set criteria were measured and the density of c-Fos expressing cells was compared across drug groups.

CORT measurements. To measure circulating CORT levels, tail blood was taken 45 (AM251/vehicle) and 150 min (URB597/LEI-410/vehicle) after injections. Blood samples were centrifuged at 3,000 r.p.m. for 20 min at 4 °C. Serum was stored at -80 °C until analysis with an enzyme-linked immunosorbent assay kit (Arbor Assays) by following the manufacturer's instructions. Samples were tested in triplicate and diluted 1:1,000 to make sure levels fitted into the standard curve. Values were normalized to the control (vehicle) group of each experiment.

Behavioral testing. Subjects. Subjects were male 7–12-week-old C57BL/6J mice obtained from The Jackson Laboratory. Mice were housed two per cage in a temperature- and humidity-controlled vivarium under a 12 h light/dark cycle (lights on 6:00 h). Experimental procedures were performed in accordance with the National Institutes of Health Guide for Care and Use of Laboratory Animals and approved by the local Animal Care and Use Committee. The number of mice used in each experiment is indicated in the figure legends. Before testing, each mouse was handled for 2 min per day for 5 d. Mice were acclimated in a room next to test area 1 h before the experiment.

Fear conditioning. Fear conditioning was conducted in 'context A': a 30 × 25 × 21 cm³ operant chamber (Med Associates, Inc.) with metal walls and a metal rod floor. To provide an additional olfactory cue, the chamber was cleaned between subjects with a 79.5% water: 19.5% ethanol: 1% vanilla extract solution. After a 180-s stimulus-free context acclimation, there were three lots of pairings of a 30-s, 75-dB white noise cue (conditioned stimulus) and a 2-s 0.6-mA scrambled coterminating footshock (unconditioned stimulus). Between each stimulus was an increasing interpairing interval (60–90–120 s). Following the last pairing was a 120-s stimulus-free period. Stimulus presentation waps controlled by the Med Associates VideoFreeze system (Med Associates). Freezing (no visible movement except respiration) was scored every 5 s by an observer blind to condition/treatment and converted to a percentage ((freezing observations/total observations) × 100).

Fear extinction training. Fear extinction training occurred the following day in 'context B': a 27 × 27 × 14 cm³ operant chamber with transparent walls and a floor covered with wood chips, cleaned between subjects with a 99% water: 1% acetic acid solution and housed in a different room from training. One hour and forty-five minutes before extinction, mice were injected with one of four combinations of drug intervention (see the section Pharmacological treatments). After a 180-s stimulus-free baseline, there were 50 × 30-s conditioned stimulus presentations (5-s interconditioned stimulus interval).

Extinction retrieval. Extinction retrieval was tested 1 d and 10 d after extinction training in context B, via 5 × 30-s conditioned stimulus presentations (5-s interconditioned stimulus interval) beginning after a 180-s stimulus-free baseline. The mean number of freezing observations per baseline period and 5× conditioned

stimulus block was converted to a percentage ((number of freezing observations/total number of observations per period) × 100) for analysis.

Statistical analysis. The effects of treatment and trial-block on freezing during extinction training were analyzed using two-way ANOVA, with repeated measures for a trial block. The effect of drug treatment on freezing during retrieval was analyzed using ANOVA.

Pharmacological treatments. Pre-extinction treatment of NAPE-PLD inhibitor LEI-401 (30 mg kg⁻¹, i.p.), FAAH inhibitor AM3506 (1 mg kg⁻¹, i.p.) and their vehicle controls were administered 1 h 45 min before the extinction training in two consecutive injections. Every injection combination led to four groups: vehicle + vehicle, vehicle + AM3506, vehicle + LEI-401 and AM3506 + LEI-401. LEI-401 stock (30 mg ml⁻¹) was prepared in DMSO. Aliquots of LEI-401 were mixed with Tween-80 and sterile distilled H₂O (1:1:8) and AM3506 stock (10 mg ml⁻¹) was prepared in DMSO and mixed with saline (1:9) just before administration (in a volume of 10 ml kg⁻¹) before extinction training.

Reporting Summary. Further information on research design is available in the Nature Research Reporting Summary linked to this article.

Data availability

All data generated or analyzed during this study are included in this published article (and its Supplementary Information files) or are available from the corresponding author on reasonable request. The mass spectrometry proteomics data (raw data and ISOQuant output tables for proteins groups and peptides) have been deposited in the ProteomeXchange Consortium (<http://proteomecentral.proteomexchange.org>) via the PRIDE partner repository with the dataset identifier PXD017586.

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Acknowledgements

The research leading to these results has received support from the Innovative Medicines Initiative Joint Undertaking under grant agreement no. 115489, resources of which are composed of financial contribution from the European Union's Seventh Framework Programme (no. FP7/2007-2013) and EFPIA companies' in kind contribution. M.v.d.S. was supported by a VICI-grant from the Netherlands Organization for Scientific Research and funding from Oncode Institute. Leiden University, Faculty of Science 'Profiling Programme: Endocannabinoids' is also acknowledged for financial support to E.D.M., V.K., T.H. and M.v.d.S. This research was also supported by National Institutes of Health National Institute on Drug Abuse (grant nos. R01DA039942 and P30DA039340), Canadian Institutes of Health Research (grant no. FDN-143329 to M.N.H.) as well as start-up funds from the VCU School of Pharmacy to A.H.L. and the Intramural Research Program of NIAAA/NIH (to P.P., O.G.-C., L.I.C., C.M.D. and A.H.). We kindly acknowledge the Pharmaceutical Sciences division of F. Hoffman-La Roche Ltd for their technical assistance with the drug metabolism and pharmacokinetics experiments.

Author contributions

E.D.M., T.H., M.N.H., P.P., A.H.L. and M.v.d.S. conceived the project and designed the experiments. J.W., H.v.d.H. and C.A.A.v.B. performed the HTS study. E.D.M. and I.K. synthesized the compounds. E.D.M., A.C.M.v.E., A.M.F.v.d.G., A.M., T.v.d.W., M.S., M.J., T.J.W. and A.P.A.J., performed biochemical and cellular experiments. A.T.B. and B.I.F. performed the proteomics measurements. V.K. and X.D. performed the in vitro and cellular lipidomics measurements. E.D.M., M.M., R.C., G.N.P., D.O., Z.V.V., J.P., C.M., G.D., J.K.P., A.P.A.J., B.I.F., M.W., U.G. and M.N.H. performed in vivo pharmacokinetics, lipidomics and behavioral studies. O.G.-C., L.I.C., C.M.D. and A.H. designed, performed and analyzed fear extinction behavior. E.D.M., M.W., U.G., B.F.C., M.W.B., H.v.d.H., C.A.A.v.B., P.P., A.H.L. and M.v.d.S. analyzed the data and wrote the manuscript.

Competing interests

E.M., I.K. C.B. and M.v.d.S. are listed as inventors on patent application WO 2019/229250 A1 filed by Leiden University in which inhibitors of NAPE-PLD are disclosed.

Additional information

Supplementary information is available for this paper at <https://doi.org/10.1038/s41589-020-0528-7>.

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Give P values as exact values whenever suitable.
- For Bayesian analysis, information on the choice of priors and Markov chain Monte Carlo settings
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Software and code

Policy information about [availability of computer code](#)

Data collection

not applicable

Data analysis

SDS-PAGE analysis was performed with Image Lab v6 (Bio-Rad). Statistical analysis was performed with Graphpad Prism v6. Proteomic analysis was performed with ProteinLynx Global SERVER™ (PLGS, Waters) and ISOQuant v1.5 (<http://www.immunologie.uni-mainz.de/isoquant/>). qPCR experiments were analyzed with CFX Manager software (Bio-rad). For lipidomics experiments in cells and lysates data was analyzed with MassLynx 4.1 (Waters). Mouse plasma pharmacokinetic analysis of LEI-401 was performed using Phoenix WinNonlin 6.4 software (Certara). Brain pharmacokinetic experiments with LEI-401 were analyzed with MassHunter Workstation Optimizer software (Agilent Technologies).

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All data is provided in the manuscript or Supplementary information. Supplementary Table 7 describes proteomic data including UniprotID accession codes for detected proteins.

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Life sciences study design

All studies must disclose on these points even when the disclosure is negative.

Sample size	No statistical analysis was performed prior to experiments to predetermine sample sizes. Sample sizes were chosen based on prior knowledge in the respective experiments and their intrinsic variability as performed in previous studies (Ogasawara et al, PNAS, 2016 and Soethoudt et al., Nat. Commun., 2017)
Data exclusions	No data were excluded.
Replication	All the in vitro and cellular experiments were performed at least in two independent experiments (N=2) with biological replicates (n=2), showing reproducible results. The in vivo lipidomics was performed in a time-dependent and in a dose-dependent manner, producing similar effects with n= 5-11 animals. The behavioral studies were performed in two independent experiments, giving similar effects with n = 9-13 animals.
Randomization	Animals were allocated in a randomized fashion into experimental groups. The behavioral studies were conducted in accordance with the ARRIVE guidelines for reporting experiments involving animals.
Blinding	Investigators were blinded to condition and treatment when scoring freezing behavior. All other experiments are measurements of enzyme activity, protein labeling, metabolites and are not based on subjective assessment of animal behavior.

Reporting for specific materials, systems and methods

We require information from authors about some types of materials, experimental systems and methods used in many studies. Here, indicate whether each material, system or method listed is relevant to your study. If you are not sure if a list item applies to your research, read the appropriate section before selecting a response.

Materials & experimental systems

n/a	Involvement in the study
<input type="checkbox"/>	<input checked="" type="checkbox"/> Antibodies
<input type="checkbox"/>	<input checked="" type="checkbox"/> Eukaryotic cell lines
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<input checked="" type="checkbox"/>	<input type="checkbox"/> Human research participants
<input checked="" type="checkbox"/>	<input type="checkbox"/> Clinical data

Methods

n/a	Involvement in the study
<input checked="" type="checkbox"/>	<input type="checkbox"/> ChIP-seq
<input checked="" type="checkbox"/>	<input type="checkbox"/> Flow cytometry
<input checked="" type="checkbox"/>	<input type="checkbox"/> MRI-based neuroimaging

Antibodies

Antibodies used	<p>Primary antibodies:</p> <p>Anti-NAPE-PLD (Abcam, ab95397, 1:200) Anti-FLAG (Sigma Aldrich, F3165, 1:5000) Anti-alfa-tubulin (Genetex, GTX76511, 1:5000) Anti-beta-actin (Abcam, ab8226, 1:5000) Anti-cFos (Cell Signaling, #2250, 1:300)</p> <p>Secondary antibodies:</p> <p>Goat-anti-rabbit-HRP (Santa Cruz, sc-2030, 1:2000) Goat-anti-mouse-HRP (Santa Cruz, sc-2005, 1:5000) Goat-anti-rat (Santa Cruz, sc-2032, 1:5000) Donkey-anti-rabbit (JacksonImmuno, 711-165-152, 1:100)</p>
Validation	All antibodies were sourced commercially. Anti-NAPE-PLD was confirmed using human and mouse NAPE-PLD overexpressing HEK293T lysate. Other validation has been performed in previous studies (van Esbroeck et al. Front. Neurosci., 2019 and Baggelaar et al. JACS, 2015).

Eukaryotic cell lines

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Cell line source(s)	HEK293T and Neuro-2a were from ATCC.
Authentication	The cell lines used have been authenticated by the ATCC.
Mycoplasma contamination	Cell lines were regularly tested for mycoplasma contamination and were tested negative.
Commonly misidentified lines (See ICLAC register)	No misidentified cell lines were used.

Animals and other organisms

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Laboratory animals	For the in vivo PK, lipidomics and behavioral experiments C57BL/6J mice (Jackson Laboratory, Bar Harbor, Maine, USA) were used. WT and NAPE-PLD KO mice from a mixed 129SvJ-C57BL/6J background were also used (Virginia Polytechnic Institute and State University, Blacksburg, Virginia, USA). 7-12-week-old male and female mice were used in this study.
Wild animals	No wild animals were used in this study.
Field-collected samples	No field-collected samples were used in this study.
Ethics oversight	Animal protocols were approved by the Institutional Animal Care and Use Committee at Virginia Commonwealth University, the National Institute on Alcohol Abuse and Alcoholism (NIAAA), National Institute of Health (NIH) and Bundesamt für Lebensmittelsicherheit und Veterinärwesen (BLV) der Schweizerischen Eidgenossenschaft.

Note that full information on the approval of the study protocol must also be provided in the manuscript.