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Article

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# β-Lactones: A Novel Class of $Ca^{2+}$-Independent Phospholipase $A_2$ (Group VIA $iPLA_2$) Inhibitors with Ability to Inhibit β-Cell Apoptosis

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ABSTRACT: Interest in $Ca^{2+}$-independent phospholipase $A_2$ (GVIA $iPLA_2$) has accelerated recently as it is being recognized as a participant in biological processes underlying diabetes development and autoimmune-based neurological disorders. The development of potent GVIA $iPLA_2$ inhibitors is of great importance, because only a few have been reported so far. We present a novel class of GVIA $iPLA_2$ inhibitors based on the β-lactone ring. This functionality in combination with a four-carbon chain carrying a phenyl group at position-3, and a linear propyl group at position-4 of the lactone ring confers excellent potency. *trans*-3-(4-Phenylbutyl)-4-propyloxetan-2-one (GK563) was identified as being the most potent GVIA $iPLA_2$ inhibitor ever reported ($X_I(50)$ 0.0000021, $IC_{50}$ 1 nM) and also one that is 22,000 times more active against GVIA $iPLA_2$ than GIVA $cPLA_2$. It was found to reduce β-cell apoptosis induced by pro-inflammatory cytokines, raising the possibility that it can be beneficial in countering autoimmune diseases, such as type 1 diabetes.

## INTRODUCTION

The phospholipase $A_2$ ($PLA_2$) superfamily consists of diverse enzymes, which are currently categorized into sixteen groups and many subgroups and all are able to hydrolyze the ester bond at the *sn*-2 position of glycerophospholipids.[1] A number of these enzymes do not require $Ca^{2+}$ ions either for their activity or for the translocation to membranes and are classified as $Ca^{2+}$-independent $PLA_2$s.[1-5] The initial reports of

$Ca^{2+}$-independent $PLA_2$ activity referred to a 40-kDa enzyme described as $iPLA_2$.[6,7] Subsequently, an 85-kDa $iPLA_2$ was purified and characterized from macrophages[8] and cloned from hamster, mouse, and rat[9-11] and is now designated as Group VIA (GVIA) $iPLA_2$ (also $iPLA_2\beta$). To date, the group VI $Ca^{2+}$-independent $PLA_2$ includes six different subgroups: GVIA ($iPLA_2\beta$), GVIB ($iPLA_2\gamma$), GVIC ($iPLA_2\delta$), GVID ($iPLA_2\varepsilon$), GVIE ($iPLA_2\zeta$), and GVIF ($iPLA_2\eta$).[1] Among them, GVIA is the most well studied and recognized $iPLA_2$. The human GVIA $iPLA_2$ (806 amino acids) contains seven ankyrin repeats (residues 152-382), a linker region (residues 383-474) with the eighth repeat disrupted by a 54-amino-acid insert, and a catalytic domain (residues 475-806). The active site serine of GVIA $iPLA_2$ lies within a lipase consensus sequence (Gly486-X-Ser519-X-Gly487) in the catalytic domain. Although GVIA $iPLA_2$ and the other major intracellular $PLA_2$, the calcium-dependent GIVA $cPLA_2$, both use a serine/aspartate catalytic dyad for their catalytic mechanism, GVIA $iPLA_2$ does not show arachidonic acid-selectivity while GIVA $cPLA_2$ does.[12]

$PLA_2$s have been implicated in a number of physiological and pathophysiological processes. Thus, a variety of synthetic inhibitors have been generated and studied *in vitro*, as well as *in vivo*.[1,13-15] The majority of these inhibitors have been developed to target cytosolic GIVA $cPLA_2$[16] and secreted $sPLA_2$.[17] Inhibitors of GVIA $iPLA_2$ had attracted less interest, because the enzyme's role was less well understood. The first introduced inhibitor for $iPLA_2$ was a bromoenol lactone compound (**1**, BEL,[18] Figure 1) which is an irreversible, covalent inhibitor of GVIA $iPLA_2$.[18] It has been used widely to delineate the specific role of GVIA $iPLA_2$ in variety of systems and biological processes.[1,13] Although BEL is selective against GVIA $iPLA_2$ versus other $PLA_2$s, it also inhibits other serine enzymes (i.e. magnesium-dependent phosphatidate

phosphohydrolase)[19] and therefore the data obtained from *ex vivo* and *in vivo* studies of its inhibitory activity must be carefully considered.

The GVIA iPLA$_2$ is involved in lipid signaling and pathological conditions including diabetes,[4,5,20] Barth syndrome[21] and progesterone-induced acrosome exocytosis.[22] The recent emergence of GVIA iPLA$_2$ as a contributor to pathophysiology prompted us to explore the possibility that more potent and selective GVIA iPLA$_2$ inhibitors can be developed.

In our first series of chemical synthesis, we generated polyfluoroketone-based compounds that proved to be more potent and selective than BEL, with the added feature of manifesting reversible inhibition of GVIA iPLA$_2$. These compounds contained an aromatic ring and a small aliphatic chain as a spacer between the two functional groups.[23-25] One of these first generation fluoroketones, FKGK11[23] (**2a**, Figure 1, $X_I$(50) 0.0014[24]), was used *in vivo* to demonstrate a role for GVIA iPLA$_2$ in both the onset and progression of experimental autoimmune encephalomyelitis, an animal model of multiple sclerosis.[26] Further, the combination of BEL or FKGK11 with anticancer drug paclitaxel was highly effective in blocking ovarian cancer development.[27]

Subsequent structure-activity relationship studies with the polyfluoroketones led to the generation of FKGK18[24] (**3**, Figure 1, $X_I$(50) 0.0002[24]), which contained a naphthyl ring and a trifluoromethyl group instead of a phenyl ring and a pentafluoroethyl group, respectively. FKGK18 was found to be 195 and >455 times more potent for GVIA iPLA$_2$ than for GIVA cPLA$_2$ and GV sPLA$_2$, respectively. In view of this, FKGK18 was deemed a valuable tool to explore the role of GVIA iPLA$_2$ in cells and *in vivo* models. Those studies revealed that FKGK18 was able to inhibit

β-cell apoptosis[28] and that its administration to spontaneous diabetes-prone non obese diabetic (NOD) mice significantly reduced diabetes incidence in association with reduced insulitis, improved glucose homeostasis, higher circulating insulin and β-cell preservation.[20] Subsequently, GK187[25] a more potent and selective GVIA iPLA$_2$ inhibitor (**2b**, Figure 1, $X_I$(50) 0.0001[25]) was identified. To gain a greater understanding of the enzyme-inhibitor interactions, a robust homology model was developed based on hydrogen/deuterium exchange mass specrtometry experimental data and molecular dynamics simulations.[29-31] Combining this with computational chemistry, organic synthesis and *in vitro* assays led to identification of new thioether fluoroketone inhibitors as well as a novel thioether keto-1,2,4-oxadiazole inhibitor (**4**, Figure 1, $X_I$(50) 0.0057) of GVIA iPLA$_2$.[32]

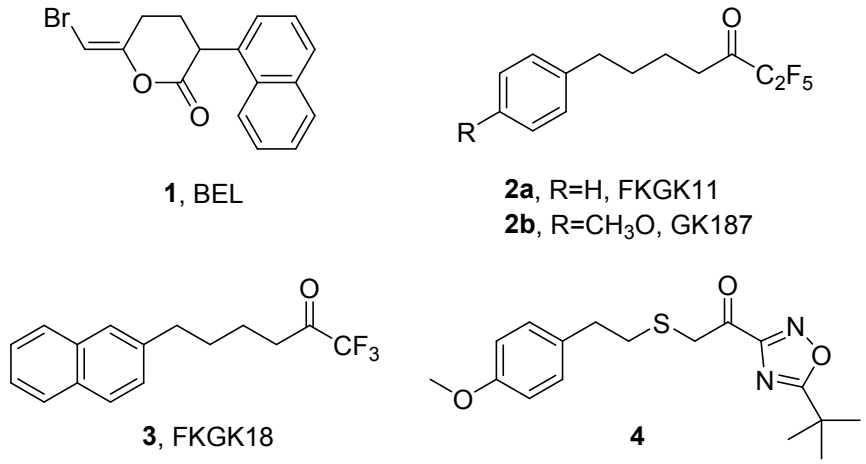



**Figure 1.** Known inhibitors of GVIA iPLA$_2$.

The need for more potent and selective inhibitors of GVIA iPLA$_2$ and the potential pharmacological limitations of fluoroketones as human therapeutics, led us to the quest of additional functional series of inhibitors. In this work, we developed a novel class of GVIA iPLA$_2$ inhibitors based on a β-lactone ring. The design and the

synthesis of a variety of these inhibitors, as well as assessment of their selectivity towards the three main human $PLA_2$s are reported. Furthermore, the ability to reduce β-cell apoptosis induced by pro-inflammatory cytokines is demonstrated.

## RESULTS AND DISCUSSION

**Design and synthesis of inhibitors.** Lipstatin (**5a**, Figure 2), a natural product isolated from *Streptomyces toxytricini*, is a potent inhibitor of pancreatic lipase[33] and the semisynthetic derivative tetrahydrolipstatin (Orlistat, **5b**, Figure 2) is an approved drug for the treatment of obesity inhibiting lipase and thus preventing the absorption of fats from the human diet. Studies on the mode of action of tetrahydrolipstatin revealed that an enzyme-inhibitor complex of an acyl-enzyme type is formed, which slowly decomposes, and that the β-lactone ring is the functional group of tetrahydrolipstatin reacting with the active site of the enzyme.[34] Several other natural products containing a β-lactone ring are enzyme inhibitors and present attractive pharmacological properties, for example marizomib (**6**, Figure 2) is a proteasome inhibitor recently approved as an orphan drug by FDA for the treatment of multiple myeloma.[35] Structural modifications of naturally occurring β-lactones have been proposed as an effective strategy for generating new drugs for treating bacterial infections, cancer, obesity and hyperlipidemia.[36]

In general, the strained β-lactone ring is expected to be attacked by the hydroxyl group of the active site serine of a serine-hydrolase. GVIA $iPLA_2$ utilizes a serine residue in its catalytic mechanism, thus, in principle it may interact with a β-lactone ring. Our previous studies on GVIA $iPLA_2$ inhibitors have shown that a potential inhibitor has to be a small non-polar molecule.[23-25] In addition, an aromatic ring

attached to a four-carbon chain seems to fit very well into the binding site of GVIA iPLA$_2$. Thus, we designed a β-lactone and chose one of the substituents to be a medium carbon chain carrying an aromatic ring in varying distances (Ar-Cn) from the lactone ring. The second substituent is a small aliphatic chain (-R) containing one to six carbon atoms. The general structure of the lactones we designed is depicted in Figure 2.

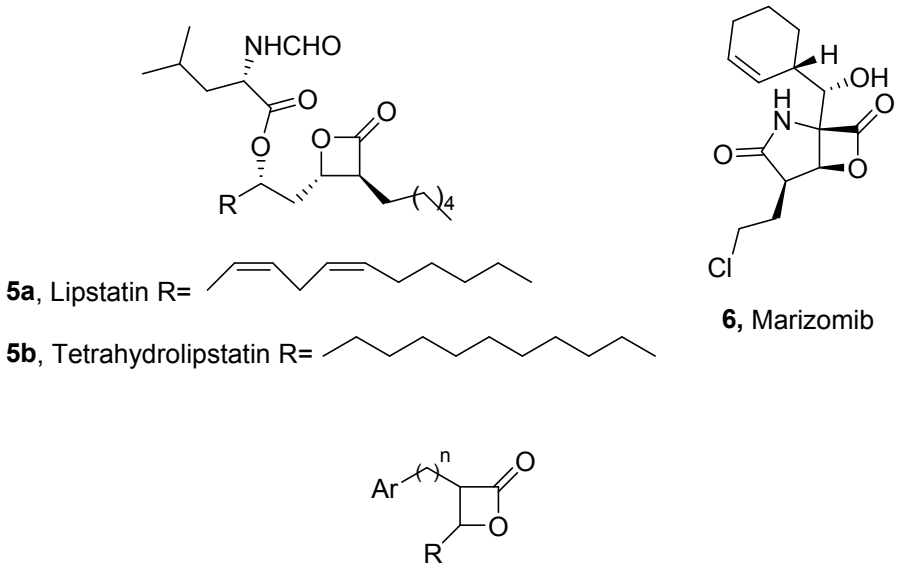
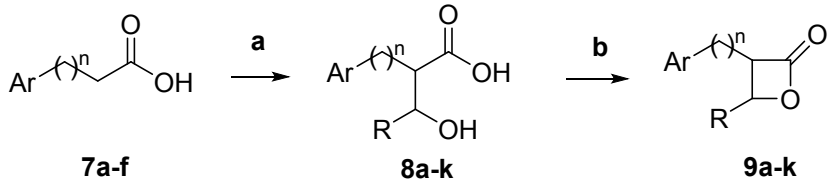
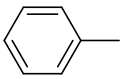
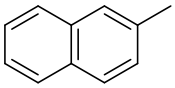
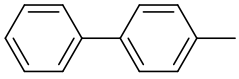
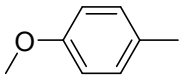
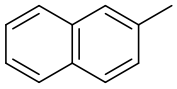
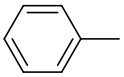


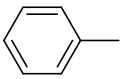
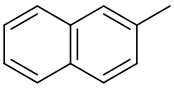
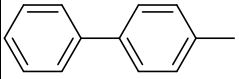
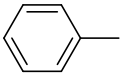
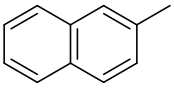
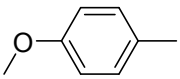
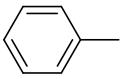
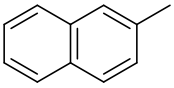
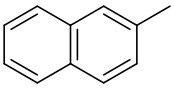
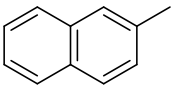
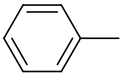

**Figure 2.** Structures of lipstatin, tetrahydrolipstatin and marizomib and general structure of β-lactones designed in this study.

The general route for the synthesis of the designed β-lactones is depicted in Scheme 1. A variety of carboxylic acids containing an aromatic group at the end of the chain were chosen as starting materials. Carboxylic acids **7a-f** were deprotonated by treatment with LDA and after reaction with commercially available aliphatic aldehydes RCHO,[37] β-hydroxy acids **8a-k** were obtained. Finally, cyclization of the intermediate α,β-substituted β-hydroxy acids **8a-k** upon treatment with p-toluenesulfonyl chloride[38] led to α,β-substituted β-lactones **9a-k** (Scheme 1).




**7a-f** **8a-k** **9a-k**

| 7 | Ar | n |
|---|---|---|
| a |  | 3 |
| b |  | 3 |
| c |  | 3 |
| d |  | 3 |
| e |  | 4 |
| f |  | 4 |

| 8, 9 | Ar | R | n |
|---|---|---|---|
| a |  | $nC_6H_{13}$ | 3 |
| b |  | $nC_6H_{13}$ | 3 |
| c |  | $nC_6H_{13}$ | 3 |
| d |  | $nC_3H_7$ | 3 |
| e |  | $nC_3H_7$ | 3 |
| f |  | $nC_3H_7$ | 3 |
| g |  | $iC_3H_7$ | 3 |
| h |  | $C_2H_5$ | 3 |
| i |  | $CH_3$ | 3 |
| j |  | $nC_3H_7$ | 4 |
| k |  | $nC_3H_7$ | 4 |

**Scheme 1.** Reagents and conditions. (a) (i) LDA (produced in situ by $(i\text{-}Pr)_2NH$ and sol. n-BuLi 1.6 M/hexane), dry THF, 0 $^oC$, 1 h, (ii) solution of RCHO in dry THF, 0 $^oC$, 1 h → r.t, o.n, (b) p-TsCl in dry pyridine, 0 $^oC$, 1 h → 4 $^oC$, 3 days.

Both β-hydroxy acids **8a-k** and β-lactones **9a-k** were obtained as mixtures of diastereomers, whose ratio was estimated by $^{1}$H NMR spectroscopy. For β-hydroxy acids **8a-k**, the ratio of the peak integrations corresponding to methinic C*H*OH signals was used to estimate the ratio of *anti*:*syn* diastereomers and varied from 7:3 to 6:4. For β-lactones, the ratio of the peak integrations corresponding to methinic proton of either C-3 or C-4 indicated a ratio of *trans*:*cis* diastereomers varying from 9:1 to 7:3. *trans* β-Lactones were obtained in excess being thermodynamically more stable than their counterparts *cis* products, and their geometry was determined by $^{1}$H NMR based on the chemical shifts reported in literature for similar compounds. In accordance to literature data, characteristic peaks for 3-CH and 4-CH are reported at 3.2 and 4.2 ppm, respectively, for *trans* β-lactones,[39] while the corresponding chemical shifts for *cis* β-lactones are reported at 3.6 and 4.5 ppm, respectively.[40] The *trans* and *cis* diastereomers of **9d, e, f, j** and **k** were separated by column chromatography. In particular for **9k**, the coupling constants between the C-3 and C-4 protons were measured to be 4.0 Hz and 6.7 Hz for the *trans* and the *cis* diastereomer (see, Supporting Information), respectively, values which are in accordance with those reported in the literature.[40] Further, in $^{13}$C NMR spectra, the chemical shifts corresponding to C-3 and C-4 are at 56.0 ppm and 77.9 ppm for the *trans* diastereomer, while at 52.6 ppm and 75.4 ppm for the *cis* diastereomer.

***In vitro* inhibition of GVIA iPLA$_2$, GIVA cPLA$_2$ and GV sPLA$_2$.** All synthesized β-lactones were tested for their *in vitro* inhibitory activity on recombinant human GVIA iPLA$_2$ using mixed micelle assays. In addition, their selectivity over human GIVA cPLA$_2$ and GV sPLA$_2$ was also studied using similar group-specific mixed micelle assays. The initial screening assays for the *in vitro* inhibition of human GVIA iPLA$_2$, GIVA cPLA$_2$ and GV sPLA$_2$ for the racemic lactones and their comparison

with previously reported fluroketones and oxadiazoles inhibitors were carried out using our previously described radioactivity-based mixed micelle assay.[41-43] For the most potent lactones, the *trans* and *cis* diastereomers were prepared and our previously described lipidomics-based mixed micelle assay was employed to determine their activities.[44,45] The inhibition results presented in Table 1 are either as percent inhibition or as $X_I(50)$ values. At first, the percent of inhibition for each $PLA_2$ enzyme at 0.091 mole fraction of each inhibitor was determined. Then, the $X_I(50)$ values were measured for compounds that displayed greater than 95% inhibition of GVIA $iPLA_2$. The $X_I(50)$ is the mole fraction of the inhibitor in the total substrate interface required to inhibit the enzyme activity by 50%. Data for inhibitors **2a**,[24] **2b**,[25] **3**[24] and **4**[32] (tested under the same radioactivity-based assay conditions) are included in Table 1 for comparison purposes.

The curves for the concentration dependence of the inhibition of GVIA $iPLA_2$ by β-lactones were fit to sigmoidal curves and those of *trans*-**9k** (GK563) and *cis*-**9k** (GK564) are presented as examples in Figure 3.

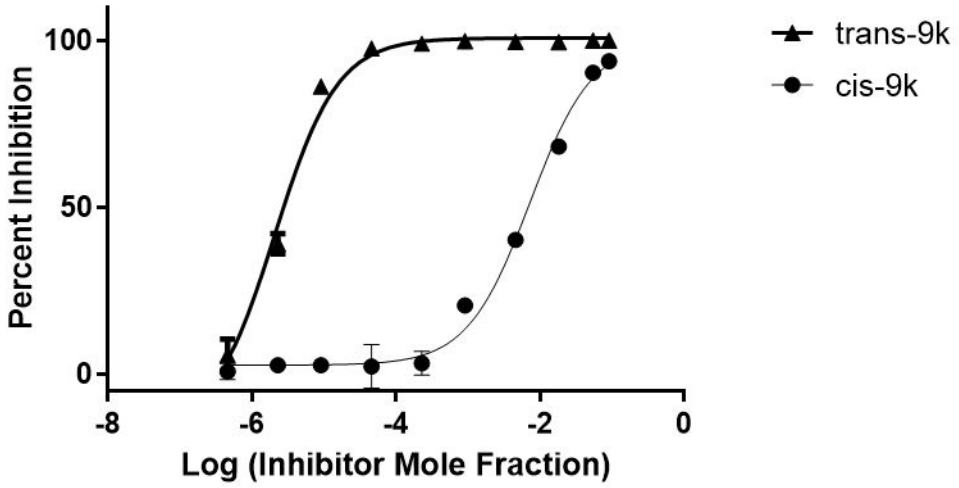


**Figure 3.** Dose-response inhibition curves for GVIA iPLA$_2$ inhibitors *trans*-**9k** and *cis*-**9k**. The curves were generated using GraphPad Prism with a nonlinear regression targeted at symmetrical sigmoidal curves based on plots of % inhibition versus log(inhibitor concentration). The reported $X_I(50)$ values were calculated from the resultant plots.

At first, β-lactones carrying an aromatic group at the end of a three-carbon atom chain and an n-hexyl chain as substituents (compounds **9a**, **9b** and **9c**) were tested. Irrespective of the nature of the aromatic group (phenyl, naphthyl, biphenyl), none of these presented significant inhibition (< 90%) of GVIA iPLA$_2$ (entries 1-3). However, it seemed that a simple phenyl (**9a**), instead of a naphthyl (**9c**) or a biphenyl (**9b**), group led to greater inhibition. Then, the hexyl chain was reduced to a shorter chain of three-carbon atoms. Interestingly, all three compounds (**9d**, **9e** and **9f**) combining an aromatic group at the end of a three-carbon atom chain and a n-propyl chain (entries 4, 7 and 10) presented significant inhibition (96-98%) of GVIA iPLA$_2$. The mixtures of these compounds were separated by column chromatography and the potencies of the corresponding *trans* and *cis* diastereomers were evaluated. Both *trans*-**9d** and *cis*-**9d** presented significant inhibition for both GVIA iPLA$_2$ and GIVA cPLA$_2$ (entries 5 and 6). However, *trans*-**9d** seemed to be more potent inhibitor of GVIA iPLA$_2$ with an $X_I(50)$ value of 0.00019 ($IC_{50}$ 95 nM, entry 5), while *cis*-**9d** more potent for GIVA cPLA$_2$ with an $X_I(50)$ value of 0.0019 ($IC_{50}$ 0.95 μM, entry 6). Among the naphthyl derivatives *trans*-**9e** and *cis*-**9e** (entries 8 and 9), *trans*-**9e** was found more potent inhibiting GVIA iPLA$_2$ with an $X_I(50)$ value of 0.00030 ($IC_{50}$ 0.15 μM, entry 8). For the para-methoxyphenyl derivatives *trans*-**9f** and *cis*-**9f** (entries 11 and 12), an interesting selectivity seems to take shape. *trans*-**9f** inhibited GVIA iPLA$_2$ with an $X_I(50)$ value equal to that estimated for *trans*-**9d** (0.00019, $IC_{50}$ 95 nM, entry

11). However, *cis*-**9f** proved to be a potent inhibitor of GIVA cPLA$_2$ ($X_I$(50) 0.00004, IC$_{50}$ 20 nM, entry 12) being almost 1000 times more potent for GIVA cPLA$_2$ than for GVIA iPLA$_2$ ($X_I$(50) 0.038, IC$_{50}$ 20 μM, entry 12).

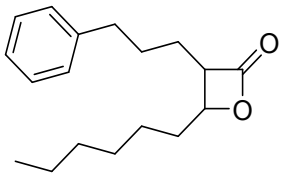
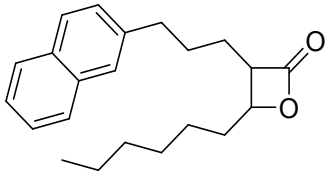
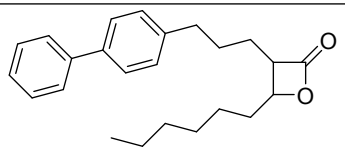
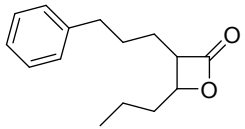
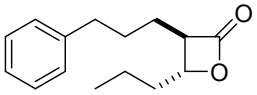
When the linear n-propyl group of **9d** was replaced by a branched isopropyl group (compound **9g**, entry 13), the inhibitory potency over GVIA iPLA$_2$ was reduced. Replacement of the n-propyl group of **9e** by an ethyl or a methyl group (compounds **9h**, **9i**, entries 14 and 15) resulted in a reduction of the inhibitory potency. Clearly, a small linear chain of three-carbon atoms led to superior inhibitory results over GVIA iPLA$_2$ in comparison to a medium chain of six-carbon atoms or a short chain of one or two carbon atoms.

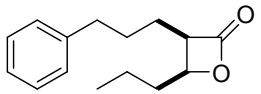
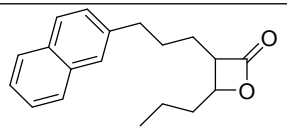
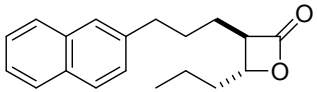
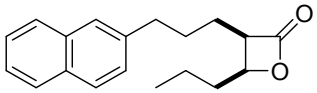
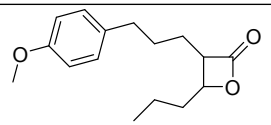
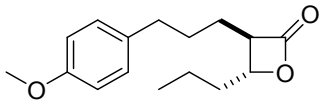
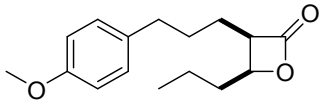
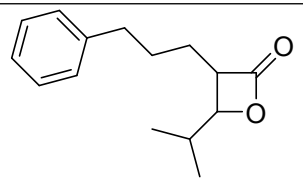
Thus, by keeping a linear three-carbon chain at position-4, the distance between the aromatic group and the lactone ring at position-3 was increased by one carbon atom resulting in compounds **9j** and **9k**, which both presented high inhibition of GVIA iPLA$_2$ (97-100%, entries 16 and 19). The diastereomers were separated by column chromatography and the potencies of both *trans* and *cis* diastereomers of **9j** and **9k** were estimated. Both the naphthyl derivatives *trans*-**9j** and *cis*-**9j** were found to inhibit GVIA iPLA$_2$ with $X_I$(50) values of 0.00009 (IC$_{50}$ 45 nM, entry17) and 0.0021 (IC$_{50}$ 1 μM, entry 18), respectively, but did not present significant inhibition of GIVA cPLA$_2$ (59% and 72% at a high concentration of 0.091 mole fraction, respectively, entries 17 and 18). Gratifyingly, the combination of a four-carbon chain carrying a phenyl group at position-3 and a linear propyl group at position-4 of the lactone ring led to the best results. The *trans* diastereomer of **9k** [*trans*-(±)-3-(4-phenylbutyl)-4-propyloxetan-2-one, GK563] was found to be a highly potent inhibitor of GVIA iPLA$_2$ with a $X_I$(50) value of 0.0000021 (IC$_{50}$ 1 nM, entry 20), while the *cis* diastereomer of **9k** [*cis*-(±)-3-(4-phenylbutyl)-4-propyloxetan-2-one, GK564] was a

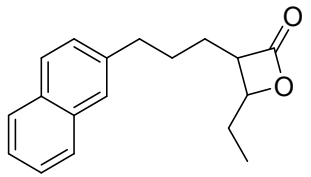
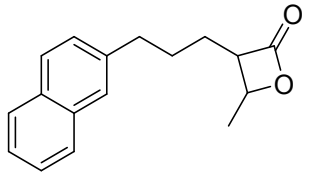
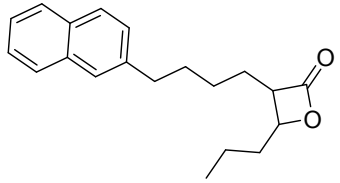
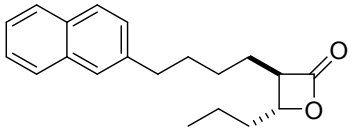
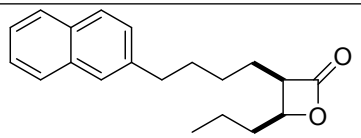
dramatically weaker inhibitor, presenting a $X_I(50)$ value of 0.007 ($IC_{50}$ 3.5 μM, entry 21). Both *trans*-**9k** and *cis*-**9k** were found to be weaker inhibitors of GIVA $cPLA_2$. In particular for inhibitor *trans*-**9k**, its $X_I(50)$ value for GIVA $cPLA_2$ was measured to be 0.042 ($IC_{50}$ 22 μM) indicating 22,000 times selectivity.

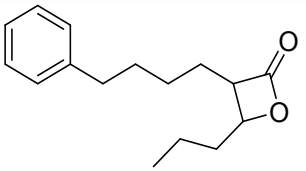
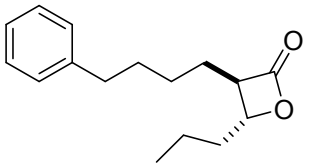
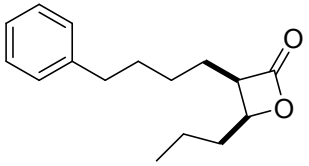
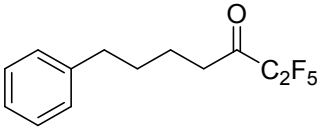
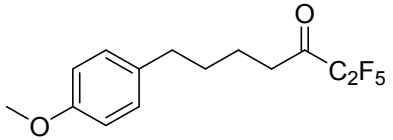
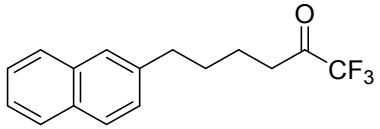
None of the β-lactones presented any appreciable inhibition of GV $sPLA_2$. The percentage inhibition of GV $sPLA_2$ did not exceed 47% (entry 1) at a high concentration of 0.091 mole fraction.

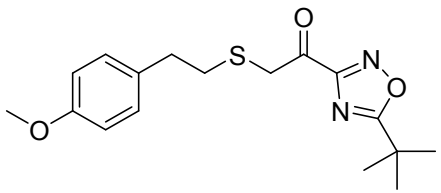
**Table 1.** *In vitro* potency and selectivity of β-lactones.

| Entry | No | Structure | GVIA iPLA$_2$ | | GIVA cPLA$_2$ | | GV sPLA$_2$ |
|---|---|---|---|---|---|---|---|
| | | | % Inhibition[a] | $X_I$(50) | % Inhibition[a] | $X_I$(50) | % Inhibition[a] |
| 1 | **9a** |  | 89 ± 2 | | 91 ± 0.5 | | 47 ± 8 |
| 2 | **9b** |  | 59 ± 7 | | 54 ± 1 | | 40 ± 8 |
| 3 | **9c** |  | 78 ± 4 | | 58 ± 3 | | 40 ± 6 |
| 4 | **9d** |  | 96 ± 1 | | 81 ± 1 | | |
| 5 | ***trans*-9d** |  | 99 ± 0 | 0.00019 ± 0.00004 | 90 ± 1 | | 25 ± 14 |

| | | | | | | | |
|---|---|---|---|---|---|---|---|
| 6 | ***cis*-9d** |  | 92 ± 1 | | 97 ± 0 | 0.0019 ± 0.0004 | N.D.[b] |
| 7 | **9e** |  | 98 ± 2 | | 77 ± 3 | | |
| 8 | ***trans*-9e** |  | 98 ± 0 | 0.00030 ± 0.00004 | 73 ± 3 | | 30 ± 3 |
| 9 | ***cis*-9e** |  | 81 ± 1 | | 89 ± 1 | | 27 ± 10 |
| 10 | **9f** |  | 96 ± 1 | | 76 ± 4 | | |
| 11 | ***trans*-9f** |  | 99 ± 1 | 0.00019 ± 0.00004 | 85 ± 2 | | 26 ± 3 |
| 12 | ***cis*-9f** |  | 85 ± 1 | 0.038 ± 0.004 | 95 ± 1 | 0.00004 ± 0.00001 | 29 ± 1 |
| 13 | **9g** |  | 93 ± 0 | | 85 ± 0.5 | | 29 ± 2 |

| | | | | | | | |
|---|---|---|---|---|---|---|---|
| 14 | **9h** |  | 93 ± 0.0 | | 85 ± 0.5 | | 29 ± 2 |
| 15 | **9i** |  | 89 ± 0.0 | | 85 ± 1 | | 26 ± 9 |
| 16 | **9j** |  | 97 ± 3 | | 71 ± 4 | | |
| 17 | ***trans*-9j** |  | 99 ± 0 | 0.00009 ± 0.00001 | 59 ± 2 | | 29 ± 5 |
| 18 | ***cis*-9j** |  | 95 ± 1 | 0.0021 ± 0.0007 | 72 ± 6 | | N.D. |

| | | | | | | | |
|---|---|---|---|---|---|---|---|
| 19 | **9k** |  | 100 ± 0 | | 84 ± 0 | | |
| 20 | ***trans*-9k** |  | 100 ± 0 | 0.0000021 ± 0.0000004 | 88 ± 3 | 0.042 ± 0.004 | 25 ± 8 |
| 21 | ***cis*-9k** |  | 99 ± 0 | 0.007 ± 0.001 | 94 ± 1 | | 34 ± 6 |
| 22 | **2a** |  | 99 ± 0[24] | 0.0014 ± 0.0001[24] | N.D.[24] | | 28 ± 1[24] |
| 23 | **2b** |  | 100[25] | 0.0001 ± 0.0000[25] | N.D.[25] | | 33[25] |
| 24 | **3** |  | 100 ± 0[24] | 0.0002 ± 0.0000[24] | 81 ± 1[24] | | 37 ± 8[24] |

| | | | | | | | |
|---|---|---|---|---|---|---|---|
| 25 | **4** |  | 98 ± 0[32] | 0.0057 ± 0.0012[32] | 70 ± 2[32] | | 38 ± 4[32] |

[a]% Inhibition at 0.091 mole fraction of each inhibitor.

[b]N.D. signifies compounds with less than 25% inhibition (or no detectable inhibition).

The results of *in vitro* inhibition clearly confirm our assumption that β-lactones inhibit the serine-based GVIA iPLA$_2$. However, a careful selection of the heterocyclic ring substituents is critical for potent inhibition. β-Lactone *trans*-**9k** stands out as the most potent inhibitor of GVIA iPLA$_2$ ever reported in literature, outperforming the potent fluoroketone FKGK18 ($X_I$(50) value of 0.0002[24], IC$_{50}$ 100 nM), which has been used successfully for *in vivo* studies.[20]

**Binding mode and interactions of 9k diastereomers.** Lactones constitute a novel class of compounds identified as potent GVIA iPLA$_2$ inhibitors. The binding mode of the most active compound in the active site of the enzyme was determined in our effort to understand its interactions with critical residues of the active site. For the docking calculations, the previously published docked structures of GIVA cPLA$_2$ and GVIA iPLA$_2$ based on our molecular dynamics simulations with two different fluoroketone compounds in the active site were used.[31,32,44] An average theoretical score of 6.0 kcal/mol was calculated for all four diastereomers of the most potent lactone GVIA iPLA$_2$ inhibitor **9k** (Table S3, Supporting Information). The binding mode of *trans*-(*S*,*S*)-**9k** in the resulting optimized docked structure showed close proximity of the carbonyl group to the oxyanion hole (Gly486/Gly487) of GVIA iPLA$_2$, while the aromatic chain was placed in the hydrophobic area of the active site, interacting with residues such as Tyr541, Met544, Val548, Phe549, Tyr643, and Leu770. The small aliphatic tail of the inhibitor was located close to Ala640 and Pro641 (Figure 4B). The lactone inhibitor *trans*-(*S*,*S*)-**9k** exhibited lower inhibition towards GIVA cPLA$_2$. The binding mode in the active site of GVIA cPLA$_2$ also showed close proximity of the carbonyl group to the oxyanion hole (Gly197/Gly198) (Figure 4A). The aromatic chain was also located in the hydrophobic area of GVIA

cPLA$_2$, but its small size does not complement the suitable aromatic interactions with the active site of GIVA cPLA$_2$.

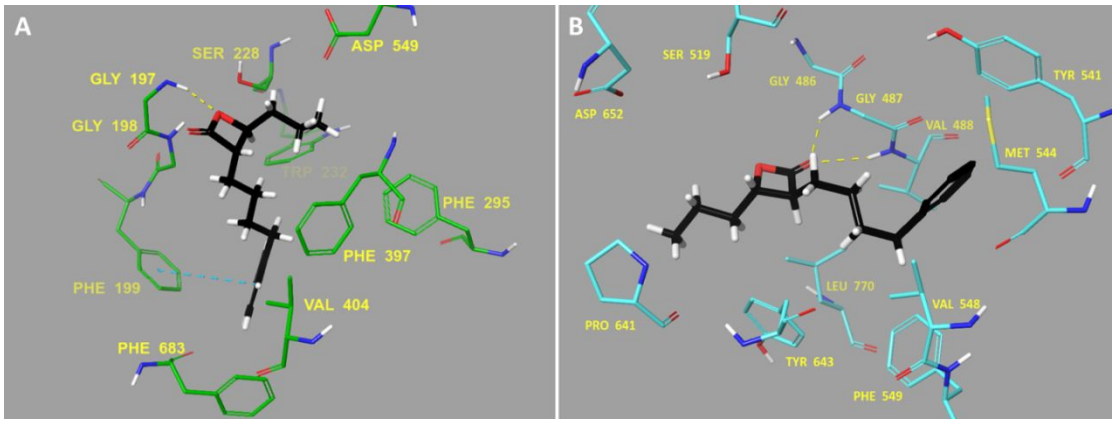
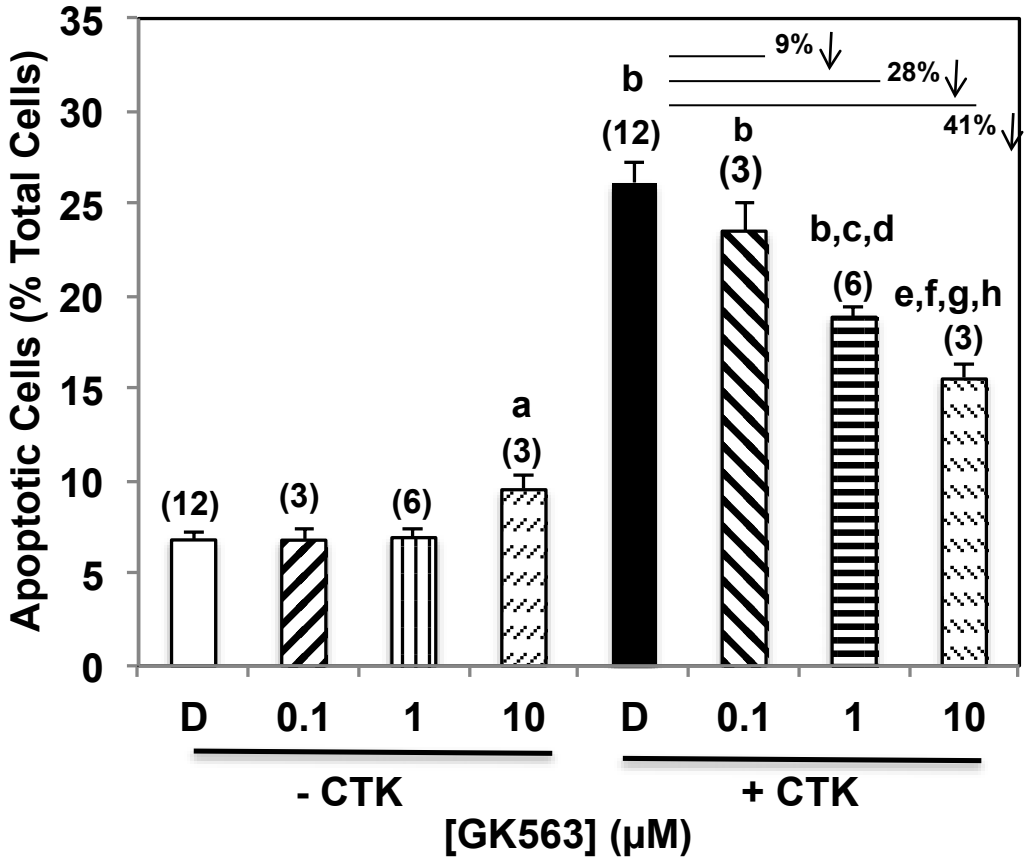



**Figure 4.** Binding mode of *trans*-(*S*,*S*)-**9k** in the active site of (A) GIVA cPLA$_2$ (PDB ID: 1CJY) and (B) GVIA iPLA$_2$ (HM based on PDB ID: 1OXW).

**Suppression of cytokine-induced β-cell apoptosis.** Type 1 diabetes (T1D) is a consequence of autoimmune destruction of islet β-cells. It is recognized that eicosanoids play important roles in promoting inflammatory responses in several diseased states, including diabetes.[46] We reported that inhibition of the GVIA iPLA$_2$ (iPLA$_2$β) mitigates β-cell death[47-52] raising the possibility that inhibitors of GVIA iPLA$_2$ may be beneficial in reducing β-cell death that leads to T1D incidence. To date, several inhibitors that can inhibit GVIA iPLA$_2$ are available, but they have limitations.[1,13-15] As we noted in introduction, recent efforts to generate more selective and potent GVIA iPLA$_2$ inhibitors identified reversible fluoroketone compounds as being selective towards GVIA.[23-25] One such inhibitor, designated FKGK18, was recently described to be more selective towards iPLA$_2$β than iPLA$_2$γ (GVIB iPLA$_2$).[28] Under *in vitro* conditions, FKGK18 inhibited insulin secretion and β-cell apoptosis.[28] Under *in vivo* conditions, it was devoid of cytotoxicity and

effective in reducing T1D incidence.[20] Thus, novel GVIA iPLA$_2$ inhibitors are very attractive as candidates for preventing β-cell apoptosis and as potential new agents for preventing T1D development.

Here, we assessed the ability of *trans*-**9k** in reducing β-cell apoptosis by treating INS-1 cells with pro-inflammatory cytokines (IL-1β + IFNγ) in the absence and presence of *trans*-**9k**. As expected, cytokine exposure resulted in a dramatic increase in β-cell apoptosis (Fig. 5). At 0.10 μM and 1.0 μM, *trans*-**9k** alone had no effect, but it promoted a slight but modest rise in cell death. Co-treatment of the cells with cytokines and *trans*-**9k** produced a concentration-dependent inhibition of β-cell apoptosis; with 0.10 μM showing minimal and non-significant effect, but significant decreases evident with 1.0 μM (28%) and 10.0 μM (41%). In comparison, these results are similar to those seen with *S*-BEL,[47] a selective inhibitor of iPLA$_2$β. However, in contrast to *S*-BEL, continuous exposure of *trans*-**9k** to cells was not cytotoxic at 0.10 or 1.0 μM, and induced only a modest rise in percent cell death (DMSO, 7.06 ± 0.36 vs. *trans*-**9k**, 9.51 ± 0.78, $p = 0.012$) at 10 μM. These findings suggest that *trans*-**9k** is another candidate inhibitor of GVIA iPLA$_2$ suitable for further studies and raise the possibility that its use *in vivo* may be beneficial in reducing β-cell death leading to T1D.




[a]Sig diff from other - CTK groups, p = 0.0065
[b]Sig diff from DMSO, $p < 10^{-10}$
[c]Sig diff from + CTK, p = 0.00045
[d]Sig diff from + CTK + 0.1 GK563, p = 0.0065
[e]Sig diff from DMSO, $p < 10^{-6}$
[f]Sig diff from + CTK, p = 0.00056
[g]Sig diff from + CTK + 1.0 GK563, p = 0.0087
[h]Sig diff from + CTK + 0.1 GK563, p = 0.0093

**Figure 5**. INS-1 cell apoptosis ± *trans*-**9k**. INS-1 cells were treated with vehicle (DMSO) or cytokines (CTK, 100 U/mL IL-1β + 300 U/mL IFNγ) for 16 h in the absence or presence of *trans*-**9k** (0.10-10 μM). The cells were then processed for TUNEL analyses and the means ± SEM (n=3-12) of percent apoptotic cells relative to total number of cells are presented.

## CONCLUSION

Herein, we describe a novel class of GVIA iPLA$_2$ inhibitors based on the β-lactone ring. This reactive functionality in combination with a four-carbon chain carrying a phenyl group at position-3, and a linear propyl group at position-4 of the lactone ring produced the best candidate inhibitor of GVIA iPLA$_2$. Inhibitor *trans*-**9k** with a $X_I(50)$ value of 0.0000021 ($IC_{50}$ 1 nM) is the most potent inhibitor of GVIA iPLA$_2$ ever reported in the literature, being a hundred times more potent than the fluoroketone inhibitor FKGK18. In addition, it is selective for GVIA iPLA$_2$, because it is 22,000 more potent for GVIA iPLA$_2$ than for GIVA cPLA$_2$. It reduces β-cell apoptosis induced by pro-inflammatory cytokines (IL-1β + IFNγ) in a concentration-dependent manner, suggesting that its use *in vivo* may be beneficial in reducing β-cell death leading to type 1 diabetes. This novel, highly potent and selective GVIA iPLA$_2$ inhibitor may be an excellent tool for the study of the role of the enzyme in cells and in animals and might help in developing novel medicinal agents.

## EXPERIMENTAL SECTION

**General.** Chromatographic purification of products was accomplished using Merck Silica Gel 60 (70-230 or 230-400 mesh). Thin-layer chromatography (TLC) was performed on Silica Gel 60 F254 aluminum plates. TLC spots were visualized with UV light and/or phosphomolybdic acid in EtOH. Melting points were determined using a Büchi 530 apparatus and were uncorrected. $^1$H and $^{13}$C NMR spectra were recorded on a Varian Mercury (200 MHz and 50 MHz, respectively) and a Bruker Avance III (600 MHz and 150 MHz, respectively) in $CDCl_3$. Chemical shifts are given in ppm, and coupling constants (*J*) in Hz. Peak multiplicities are described as

follows: s, singlet, d, doublet, t, triplet and m, multiplet. Low resolution electron spray ionization (ESI) mass spectra were recorded on a Finnigan, Surveyor MSQ Plus spectrometer, while HRMS spectra were recorded on a Bruker Maxis Impact QTOF Spectrometer. Dichloromethane was dried by standard procedures and stored over molecular sieves. All other solvents and chemicals were reagent grade and used without further purification. The purity of all compounds subjected to biological tests was determined by analytical HPLC, and was found to be ≥95%. HPLC analyses were carried out on a Shimadzu LC-2010AHT system and an ODS Hypersil (250 x 4.6 mm, 5 μm) analytical column, using $H_2O$/acetonitrile 20/80 v/v, at a flow rate of 1.0 mL/min. HPLC analyses of *trans*-**9k** and *cis*-**9k** were carried out on a Agilent 1100 system and a Daicel Chiralcel OD-H (250 × 4.6 mm, 5 μm) using hexane/i-PrOH 95/5 v/v, at a flow rate of 1.0 mL/min.

Carboxylic acids **5a** and **5f** were commercially available. Carboxylic acids **5b**,[53] **5c**,[25] **5d**[25] and **5e**[53] have been described elsewhere and their analytical data are in accordance with literature. β-Hydroxy acids **8a-k** and β-lactones **9a-k** were obtained as mixtures of diastereomers. The diastereomeric ratio (d.r.) of the mixtures was determined by $^1$H NMR spectroscopy.

**General method for the synthesis of β-hydroxy acids (8a-k).** To a stirred solution of diisopropylamine (3 mmol) in anhydrous THF (2 mL), under argon at 0 $^o$C, a solution of n-BuLi 1.6 M in hexane (3 mmol, 1.9 mL) was slowly added via syringe and the solution of LDA was stirred at 0 $^o$C for 10 minutes. The carboxylic acid **7a-f** (1 mmol) in anhydrous THF (6 mL) was then added and the solution was stirred at 0 $^o$C for 1 h. Then, aldehyde (1.3 mmol) in anhydrous THF (2 mL) was added and the solution was stirred at 0 $^o$C for 1 h and at room temperature overnight. The solvent

was removed under reduced pressure. The reaction mixture was acidified with HCl 1N to pH 2 and extracted with $Et_2O$ (3x 20 mL). The organic layers were combined, washed with brine and dried. The solvent was removed and the product was purified by column chromatography eluting with a gradient of $CHCl_3$/MeOH 95/5 to 9/1.

**3-Hydroxy-2-(3-phenylpropyl)nonanoic acid (8a).** Mixture of diastereomers (d.r. 6:4 *anti*:*syn*). Yield 51%; Oil; $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.37-7.09 (m, 5H), 3.92-3.77 (m, 0.4H), 3.76-3.61 (m, 0.6H), 2.73-2.55 (m, 2H), 2.54-2.37 (m, 1H), 1.87-1.56 (m, 4H), 1.55-1.13 (m, 10H), 0.90 (t, $J$ = 6.6 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 180.6, 141.8, 128.3, 128.2, 125.8, 72.2, 72.1, 50.9, 50.6, 35.7, 35.6, 35.2, 33.9, 31.7, 29.1, 29.0, 28.9, 28.7, 25.6, 22.6, 14.1; MS (ESI) m/z (%): 291.3 [$(M-H)^-$, 100]; HRMS: 315.1947 $(M+Na)^+$, (315.1931).

**3-Hydroxy-2-(3-(naphthalen-2-yl)propyl)nonanoic acid (8b).** Mixture of diastereomers (d.r. 7:3 *anti*:*syn*). Yield 35%; Oil; $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.86-7.65 (m, 3H), 7.55 (s, 1H), 7.48-7.21 (m, 3H), 3.93-3.75 (m, 0.3H), 3.74-3.57 (m, 0.7H), 2.85-2.65 (m, 2H), 2.56-2.29 (m, 1H), 1.91-1.56 (m, 4H), 1.56-1.03 (m, 10H), 0.84 (t, $J$ = 6.4 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 180.7, 139.3, 133.5, 131.9, 127.8, 127.5, 127.4, 127.2, 126.3, 125.8, 125.1, 72.3, 72.2, 51.1, 50.9, 36.3, 35.8, 35.4, 34.0, 31.8, 29.2, 29.0, 28.9, 28.7, 25.6, 22.6, 14.1; MS (ESI) m/z (%): 341.3 [$(M-H)^-$, 100]; HRMS: 365.2090 $(M+Na)^+$, (365.2087).

**2-(3-([1,1'-Biphenyl]-4-yl)propyl)-3-hydroxynonanoic acid (8c).** Mixture of diastereomers (d.r. 7:3 *anti*:*syn*). Yield 52%; Solid; mp 40-42 $^o$C; $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.60-7.21 (m, 9H), 3.95-3.83 (m, 0.3H), 3.79-3.63 (m, 0.7H), 2.75-2.60 (m, 3H), 1.88-1.60 (m, 4H), 1.53-1.13 (m, 10H), 0.88 (t, $J$ = 6.0 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 180.6, 141.0, 138.8, 128.8, 128.7, 127.0, 126.9, 72.3, 72.2, 51.1,

50.9, 35.5, 35.3, 33.9, 33.9, 31.8, 29.5, 29.1, 29.0, 28.7, 25.6, 22.6, 14.1; MS (ESI) m/z (%): 367.2 [(M-H)$^-$, 100].

**3-Hydroxy-2-(3-phenylpropyl)hexanoic acid (8d).** Mixture of diastereomers (d.r. 7:3 *anti*:*syn*). Yield 28%; Oil; $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.37-7.07 (m, 5H), 3.94-3.80 (m, 0.3H), 3.79-3.67 (m, 0.7H), 2.72-2.57 (m, 2H), 2.57-2.42 (m, 1H), 1.89-1.56 (m, 4H), 1.56-1.30 (m, 4H), 0.93 (t, $J$ = 6.2 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 180.6, 141.9, 128.3, 128.2, 125.8, 71.9, 71.8, 50.8, 50.7, 37.4, 36.1, 35.8, 35.6, 29.5, 29.0, 28.9, 26.3, 19.1, 18.8, 13.9; MS (ESI) m/z (%): 249.3 [(M-H)$^-$, 100]; HRMS: 273.1475 (M+Na)$^+$, (273.1461).

**3-Hydroxy-2-(3-(naphthalen-2-yl)propyl)hexanoic acid (8e).** Mixture of diastereomers (d.r. 7:3 *anti*:*syn*). Yield 20%; Solid; mp 38-40 $^o$C; $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.90-7.67 (m, 3H), 7.60 (s, 1H), 7.50-7.19 (m, 3H), 3.94-3.80 (m, 0.3H), 3.80-3.62 (m, 0.7H), 2.78 (t, $J$ = 6.0 Hz, 2H), 2.58-2.38 (m, 1H), 1.94-1.57 (m, 4H), 1.57-1.15 (m, 4H), 0.89 (t, $J$ = 6.2 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 180.4, 139.3, 133.5, 131.9, 127.9, 127.5, 127.4, 127.1, 126.4, 125.9, 125.1, 71.9, 71.8, 51.0, 50.7, 37.4, 36.0, 35.9, 35.7, 29.3, 29.0, 28.9, 26.2, 19.1, 18.8, 13.9; MS (ESI) m/z (%): 299.3 [(M-H)$^-$, 100].

**3-Hydroxy-2-(3-(4-methoxyphenyl)propyl)hexanoic acid (8f).** Mixture of diastereomers (d.r. 6:4 *anti*:*syn*). Yield 46%; Oil; $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.06 (d, $J$ = 8.0 Hz, 2H), 6.79 (d, $J$ = 8.0 Hz, 2H), 3.95-3.80 (m, 0.4H), 3.79 (s, 3H), 3.78-3.59 (m, 0.6H), 2.56 (t, $J$ = 6.8 Hz, 2H), 2.52-2.43 (m, 1H), 1.86-1.53 (m, 4H), 1.53-1.19 (m, 4H), 1.02-0.80 (m, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 180.3, 157.6, 134.0, 129.2, 113.7, 71.9. 71.8, 55.2, 50.9, 50.7, 37.4, 36.0, 34.8, 34.7, 29.7, 29.3, 28.9, 26.2, 19.1, 18.8, 13.9; MS (ESI) m/z (%): 279.2 [(M-H)$^-$, 100].

**3-Hydroxy-4-methyl-2-(3-phenylpropyl)pentanoic acid (8g).** Mixture of diastereomers (d.r. 7:3 *anti*:*syn*). Yield 28%; Solid; mp 85-90 °C; $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.46-7.03 (m, 5H), 3.64-3.45 (m, 0.3H), 3.45-3.25 (m, 0.7H), 2.72-2.44 (m, 2H and 0.7H), 2.43-2.27 (m, 0.3H), 1.89-1.40 (m, 5H), 1.05-0.72 (m, 6H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 180.9, 180.7, 141.8, 128.3, 128.2, 125.8, 77.2, 48.3, 48.0, 35.8, 35.6, 31.8, 30.9, 29.5, 29.3, 29.0, 26.1, 19.6, 19.5, 17.5, 17.4; MS (ESI) m/z (%): 268.2 [($M+NH_4^+$), 100].

**3-Hydroxy-2-(3-(naphthalen-2-yl)propyl)pentanoic acid (8h).** Mixture of diastereomers (d.r. 7:3 *anti*:*syn*). Yield 30%; Solid; mp 115-120 °C; $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.83-7.70 (m, 3H), 7.60 (s, 1H), 7.48-7.28 (m, 3H), 3.78-3.68 (m, 0.3H), 3.60-3.53 (m, 0.7H), 2.75 (t, $J$ = 6.0 Hz, 2H), 2.57-2.37 (m, 1H), 1.88-1.37 (m, 6H), 0.93 (t, $J$ = 7.0 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 179.1, 139.5, 133.4, 131.8, 127.7, 127.4, 127.2, 127.1, 126.2, 125.7, 124.9, 73.7, 73.6, 51.1, 50.6, 35.8, 35.6, 29.3, 28.9, 28.8, 27.8, 10.3, 9.8; MS (ESI) m/z (%): 304.2 [($M+NH_4^+$), 100].

**2-(1-Hydroxyethyl)-5-(naphthalen-2-yl)pentanoic acid (8i).** Mixture of diastereomers (d.r. 6:4 *anti*:*syn*). Yield 33%; Solid; mp 125-130 °C; $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.86-7.70 (m, 3H), 7.60 (s, 1H), 7.50-7.25 (m, 3H), 4.12-3.99 (m, 0.4H), 3.99-3.86 (m, 0.6H), 2.79 (t, $J$ = 6.6 Hz, 2H), 2.58-2.36 (m, 1H), 1.93-1.54 (m, 4H), 1.29-1.16 (m, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 180.0, 179.8, 139.2, 133.5, 132.0, 127.9, 127.6, 127.4, 127.1, 126.3, 125.9, 125.1, 68.3, 68.1, 52.6, 51.7, 35.9, 35.7, 29.3, 28.8, 26.7, 21.4, 20.0; MS (ESI) m/z (%): 290.4 [($M+NH_4^+$), 100].

**3-Hydroxy-2-(4-(naphthalen-2-yl)butyl)hexanoic acid (8j).** Mixture of diastereomers (d.r. 6:4 *anti*:*syn*). Yield 20%; Solid; mp 36-38 °C; $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.83-7.70 (m, 3H), 7.60 (s, 1H), 7.48-7.28 (m, 3H), 3.93-3.79 (m, 0.4H), 3.79-3.65 (m, 0.6H), 2.77 (t, $J$ = 7.2 Hz, 2H), 2.53-2.39 (m, 1H), 1.85-1.59 (m, 4H),

1.58-1.31 (m, 6H), 0.91 (t, $J$ = 6.6 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 179.8, 139.8, 133.5, 131.9, 127.8, 127.6, 127.4, 127.3, 126.3, 125.8, 125.0, 71.8, 71.7, 50.7, 50.6, 37.6, 36.1, 35.8, 31.3, 31.1, 29.3, 27.4, 27.0, 26.5, 19.1, 18.9, 13.9; MS (ESI) m/z (%): 332.2 [($M+NH_4^+$), 100].

**3-Hydroxy-2-(4-phenylbutyl)hexanoic acid (8k).** Mixture of diastereomers (d.r. 7:3 *anti*:*syn*). Yield 68%; Solid; mp 38-40 °C; $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.37-7.07 (m, 5H), 3.93-3.80 (m, 0.3H), 3.80-3.64 (m, 0.7H), 2.62 (t, $J$ = 7.6 Hz, 2H), 2.53-2.37 (m, 1H), 1.85-1.55 (m, 4H), 1.55-1.27 (m, 6H), 0.94 (t, $J$ = 7.0 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 180.5, 142.3, 128.3, 128.2, 125.7, 71.9, 71.8, 50.9, 50.7, 37.5, 36.1, 35.6, 31.4, 31.3, 29.2, 27.4, 26.9, 26.4, 19.1, 18.9, 13.9; MS (ESI) m/z (%): 282.2 [($M+NH_4^+$), 100].

**General method for the cyclization of β-hydroxy acids to β-lactones (9a-k).** To a stirred solution of β-hydroxy acid **8a-k** (1 mmol) in anhydrous pyridine (2 mL), under argon at 0 °C, p-toluenesulfonyl chloride (2 mmol) in anhydrous pyridine (1 mL) was added slowly via syringe. The solution was stirred at 0 °C for 1 h and kept at 4 °C for 3 days. Then, $Et_2O$ was added and the organic layer was washed with 10% $Na_2CO_3$, HCl 1N to pH 2 and brine. The organic layer was dried and the solvent was removed in vacuo. The product was purified by column chromatography eluting with a gradient of petroleum ether (bp 40-60 °C) /AcOEt 95/5 to 9/1.

**4-Hexyl-3-(3-phenylpropyl)oxetan-2-one (9a)**. Mixture of diastereomers (d.r. 8:2 *trans*:*cis*).Yield 75%; Oil; $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.35-7.11 (m, 5H), 4.58-4.44 (m, 0.2H), 4.27-4.13 (m, 0.8H), 3.68-3.53 (m, 0.2H), 3.26-3.11 (m, 0.8H), 2.65 (t, $J$ = 6.8 Hz, 2H), 1.93-1.66 (m, 6H), 1.40-1.22 (m, 8H), 0.89 (t, $J$ = 6.4 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 172.1, 171.4, 141.3, 128.4, 128.3, 126.0, 78.0, 75.6, 55.9,

52.4, 35.4, 34.4, 31.5, 30.1, 29.1, 28.8, 28.6, 27.3, 25.4, 24.9, 23.3, 22.5, 14.0; MS (ESI) m/z (%): 292.3 [(M+$NH_4^+$), 100]; HRMS: 297.1842 $(M+Na)^+$, (297.1825).

**4-Hexyl-3-(3-(naphthalen-2-yl)propyl)oxetan-2-one (9b).** Mixture of diastereomers (d.r. 9:1 *trans*:*cis*).Yield 44%; Oil; $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.87-7.72 (m, 3H), 7.60 (s, 1H), 7.52-7.23 (m, 3H), 4.60-4.41 (m, 0.1H), 4.27-4.10 (m, 0.9H), 3.69-3.55 (m, 0.1H), 3.27-3.10 (m, 0.9H), 2.82 (t, $J$ = 7.0 Hz, 2H), 1.97-1.63 (m, 6H), 1.45-1.16 (m, 8H), 0.87 (t, $J$ = 6.0 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 171.4, 138.8, 133.5, 132.0, 128.1, 127.6, 127.4, 127.0, 126.5, 126.0, 125.3, 78.0, 55.9, 35.6, 34.4, 31.6, 28.9, 28.5, 27.4, 25.0, 22.5, 14.0; MS (ESI) m/z (%): 342.2 [(M+$NH_4^+$), 100]; HRMS: 347.1986 $(M+Na)^+$, (347.1982).

**3-(3-([1,1'-Biphenyl]-4-yl)propyl)-4-hexyloxetan-2-one (9c).** Mixture of diastereomers (d.r. 7:3 *trans*:*cis*).Yield 57%; Solid; mp 35-38 $^o$C; $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.64-7.16 (m, 9H), 4.61-4.44 (m, 0.3H), 4.27-4.15 (m, 0.7H), 3.71-3.55 (m, 0.3H), 3.28-3.12 (m, 0.7H), 2.70 (t, $J$ = 6.8 Hz, 2H) 2.02-1.51 (m, 6H), 1.51-1.14 (m, 8H), 0.88 (t, $J$ = 6.8 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 172.1, 171.4, 140.9, 140.4, 139.0, 128.8, 128.7, 127.2, 127.1, 127.0, 78.0, 75.6, 55.9, 52.5, 35.1, 34.4, 31.6, 30.1, 29.1, 28.9, 28.6, 27.4, 25.5, 25.0, 23.6, 22.5, 14.0; MS (ESI) m/z (%): 368.3 [(M+$NH_4^+$), 100]; HRMS: 373.2141 $(M+Na)^+$, (373.2138).

**3-(3-Phenylpropyl)-4-propyloxetan-2-one (9d).** Mixture of diastereomers (d.r. 9:1 *trans*:*cis*). Yield 68%; Oil; $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.40-7.10 (m, 5H), 4.60-4.45 (m, 0.1H), 4.28-4.13 (m, 0.9H), 3.68-3.54 (m, 0.1H), 3.26-3.10 (m, 0.9H), 2.66 (t, $J$ = 7.0 Hz, 2H), 1.96-1.56 (m, 6H), 1.56-1.30 (m, 2H), 0.97 (t, $J$ = 7.2 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$) δ 172.1, 171.4, 141.3, 128.4, 128.3, 126.0, 77.9, 75.4, 56.0,

52.5, 36.4, 35.5, 32.1, 29.2, 28.7, 27.4, 23.4, 18.9, 18.4, 13.8; MS (ESI) m/z (%): 250.1 [($M+NH_4^+$), 100]; HRMS: 255.1358 $(M+Na)^+$, (255.1356).

***trans*-(±)-3-(3-Phenylpropyl)-4-propyloxetan-2-one (*trans*-9d).** $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.36-7.10 (m, 5H), 4.28-4.13 (m, 1H), 3.26-3.10 (m, 1H), 2.66 (t, $J$ = 7.0 Hz, 2H), 1.96-1.56 (m, 6H), 1.56-1.30 (m, 2H), 0.97 (t, $J$ = 7.2 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$) δ 171.4, 141.3, 128.4, 128.3, 126.0, 77.9, 56.0, 36.4, 35.5, 28.7, 27.4, 18.4, 13.8.

***cis*-(±)-3-(3-Phenylpropyl)-4-propyloxetan-2-one (*cis*-9d).** $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.36-7.10 (m, 5H), 4.60-4.45 (m, 1H), 3.68-3.54 (m, 1H), 2.66 (t, $J$ = 7.0 Hz, 2H), 1.96-1.30 (m, 8H), 0.97 (t, $J$ = 7.2 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$) δ 172.1, 141.3, 128.4, 128.3, 126.0, 75.4, 52.4, 35.4, 32.1, 29.1, 23.3, 18.8, 13.8.

**3-(3-(Naphthalen-2-yl)propyl)-4-propyloxetan-2-one (9e).** Mixture of diastereomers (d.r. 8:2 *trans*:*cis*). Yield 66%; Oil; $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.86-7.67 (m, 3H), 7.60 (s, 1H), 7.52-7.24 (m, 3H), 4.59-4.45 (m, 0.2H), 4.28-4.13 (m, 0.8H), 3.69-3.55 (m, 0.2H), 3.27-3.07 (m, 0.8H), 2.82 (t, $J$ = 6.8 Hz, 2H), 1.96-1.62 (m, 6H), 1.51-1.25 (m, 2H), 0.95 (t, $J$ = 6.8 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 172.1, 171.4, 138.8, 133.5, 132.0, 128.1, 127.6, 127.4, 127.0, 126.5, 126.0, 125.3, 77.8, 75.4, 56.0, 52.5, 36.4, 35.6, 32.1, 28.9, 28.5, 27.4, 23.4, 18.9, 18.4, 13.7; MS (ESI) m/z (%): 300.2 [($M+NH_4^+$), 100]; HRMS: 305.1516 $(M+Na)^+$, (305.1512).

***trans*-(±)-3-(3-(Naphthalen-2-yl)propyl)-4-propyloxetan-2-one (*trans*-9e).** $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.86-7.67 (m, 3H), 7.60 (s, 1H), 7.52-7.24 (m, 3H), 4.28-4.13 (m, 1H), 3.27-3.07 (m, 1H), 2.82 (t, $J$ = 6.8 Hz, 2H), 1.96-1.62 (m, 6H), 1.51-1.25 (m, 2H), 0.95 (t, $J$ = 6.8 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 171.4, 138.8,

133.5, 132.0, 128.1, 127.6, 127.4, 127.0, 126.5, 126.0, 125.3, 77.8, 56.0, 36.4, 35.6, 28.5, 27.4, 18.4, 13.8.

***cis*-(±)-3-(3-(Naphthalen-2-yl)propyl)-4-propyloxetan-2-one (*cis*-9e).** $^{1}$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.86-7.67 (m, 3H), 7.60 (s, 1H), 7.52-7.24 (m, 3H), 4.59-4.45 (m, 1H), 3.69-3.55 (m, 1H), 2.82 (t, $J$ = 6.8 Hz, 2H), 1.96-1.25 (m, 8H), 0.95 (t, $J$ = 6.8 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 172.1, 138.8, 133.5, 132.0, 128.1, 127.6, 127.4, 127.0, 126.5, 126.0, 125.3, 75.4, 52.5, 35.6, 32.1, 28.9, 23.4, 18.9, 13.8.

**3-(3-(4-Methoxyphenyl)propyl)-4-propyloxetan-2-one (9f).** Mixture of diastereomers (d.r. 8:2 *trans*:*cis*). Yield 65%; Oil; $^{1}$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.09 (d, $J$ = 8.4 Hz, 2H), 6.84 (d, $J$ = 8.4 Hz, 2H), 4.62-4.47 (m, 0.2H), 4.28-4.14 (m, 0.8H), 3.79 (s, 3H), 3.68-3.51 (m, 0.2H), 3.27-3.08 (m, 0.8H), 2.60 (t, $J$ = 7.0 Hz, 2H), 1.83-1.59 (m, 6H), 1.50-1.36 (m, 2H), 0.97 (t, $J$ = 7.4 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 172.1, 171.4, 157.8, 133.3, 129.2, 113.8, 77.9, 75.4, 55.9, 55.2, 52.5, 36.4, 34.5, 32.1, 29.3, 28.9, 27.3, 23.3, 19.0, 18.4, 13.8; MS (ESI) m/z (%): 280.3 [($M+NH_4^+$), 100]; HRMS: 285.1464 $(M+Na)^+$, (285.1461).

***trans*-(±)-3-(3-(4-Methoxyphenyl)propyl)-4-propyloxetan-2-one (*trans*-9f).** $^{1}$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.09 (d, $J$ = 8.4 Hz, 2H), 6.84 (d, $J$ = 8.4 Hz, 2H), 4.28-4.14 (m, 1H), 3.79 (s, 3H), 3.27-3.08 (m, 1H), 2.60 (t, $J$ = 7.0 Hz, 2H), 1.83-1.59 (m, 6H), 1.50-1.36 (m, 2H), 0.97 (t, $J$ = 7.4 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 171.4, 157.8, 133.3, 129.2, 113.8, 77.9, 55.9, 55.2, 36.4, 34.5, 28.9, 27.3, 18.4, 13.7.

***cis*-(±)-3-(3-(4-Methoxyphenyl)propyl)-4-propyloxetan-2-one (*cis*-9f).** $^{1}$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.09 (d, $J$ = 8.4 Hz, 2H), 6.84 (d, $J$ = 8.4 Hz, 2H), 4.62-4.47 (m, 1H), 3.79 (s, 3H), 3.68-3.51 (m, 1H), 2.60 (t, $J$ = 7.0 Hz, 2H), 1.83-1.36 (m, 8H), 0.97

(t, $J$ = 7.4 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 172.1, 157.8, 133.3, 129.2, 113.8, 75.4, 55.2, 52.5, 34.5, 32.1, 29.4, 23.3, 18.9, 13.8.

**4-Isopropyl-3-(3-phenylpropyl)oxetan-2-one (9g).** Mixture of diastereomers (d.r. 7:3 *trans*:*cis*). Yield 36%; Oil; $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.38-7.07 (m, 5H), 4.15-4.02 (m, 0.3H), 3.94-3.83 (m, 0.7H), 3.67-3.52 (m, 0.3H), 3.28-3.13 (m, 0.7H), 2.66 (t, $J$ = 6.8 Hz, 2H), 2.12-1.57 (m, 5H), 1.03 (d, $J$ = 6.6 Hz, 3H), 0.94 (d, $J$ = 6.6 Hz, 2.1H), 0.88 (d, $J$ = 6.6 Hz, 0.9H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 172.1, 171.4, 141.3, 128.4, 128.3, 126.0, 82.7, 80.2, 53.9, 51.9, 35.5, 32.3, 28.9, 28.7, 27.7, 23.7, 19.0, 18.0, 17.0; MS (ESI) m/z (%): 250.2 [($M+NH_4^+$), 100]; HRMS: 255.1357 $(M+Na)^+$, (255.1356).

**4-Ethyl-3-(3-(naphthalen-2-yl)propyl)oxetan-2-one (9h).** Mixture of diastereomers (d.r. 7:3 *trans*:*cis*).Yield 53%; Oil; $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.94-7.70 (m, 3H), 7.60 (s, 1H), 7.56-7.23 (m, 3H), 4.53-4.34 (m, 0.3H), 4.23-4.06 (m, 0.7H), 3.70-3.52 (m, 0.3H), 3.30-3.09 (m, 0.7H), 2.81 (t, $J$ = 6.4 Hz, 2H), 2.07-1.58 (m, 6H), 1.09-0.91 (m, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 172.1, 171.3, 138.7, 133.4, 131.9, 128.0, 127.5, 127.3, 127.0, 126.4, 126.0, 125.2, 79.0, 76.8, 55.4, 52.3, 35.5, 28.9, 28.5, 27.4, 23.4, 23.3, 9.8, 9.1; MS (ESI) m/z (%): 286.3 [($M+NH_4^+$), 100]; HRMS: 291.1356 $(M+Na)^+$, (291.1356).

**4-Methyl-3-(3-(naphthalen-2-yl)propyl)oxetan-2-one (9i).** Mixture of diastereomers (d.r. 7:3 *trans*:*cis*). Yield 34%; Solid; mp 35-40 $^o$C; $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.86-7.71 (m, 3H), 7.60 (s, 1H), 7.48-7.28 (m, 3H), 4.81-4.63 (m, 0.4H), 4.45-4.30 (m, 0.6H), 3.71-3.52 (m, 0.4H), 3.27-3.10 (m, 0.6H), 2.82 (t, $J$ = 6.6 Hz, 2H), 2.07-1.64 (m, 4H), 1.53 (d, $J$ = 6.2 Hz, 2.1H), 1.42 (d, $J$ = 6.2 Hz, 0.9H); $^{13}$C

NMR (50 MHz, $CDCl_3$): $\delta$ 171.7, 171.0, 138.8, 138.7, 133.5, 132.0, 128.0, 127.6, 127.4, 127.0, 126.5, 126.0, 125.3, 125.2, 74.5, 71.6, 57.4, 52.6, 35.6, 28.7, 28.4, 27.2, 23.4, 20.3, 15.6; MS (ESI) m/z (%): 272.2 [$(M+NH_4^+)$, 100]; HRMS: 277.1193 $(M+Na)^+$, (277.1199).

**3-(4-(Naphthalen-2-yl)butyl)-4-propyloxetan-2-one (9j).** Mixture of diastereomers (d.r. 7:3 *trans*:*cis*).Yield 48%; Low melting point solid; $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.87-7.69 (m, 3H), 7.60 (s, 1H), 7.52-7.23 (m, 3H), 4.59-4.42 (m, 0.3H), 4.27-4.10 (m, 0.7H), 3.66-3.50 (m, 0.3H), 3.23-3.05 (m, 0.7H), 2.79 (t, $J$ = 7.2 Hz, 2H), 1.96-1.29 (m, 10H), 0.94 (t, $J$ = 7.4 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 172.2, 171.6, 139.5, 133.5, 131.9, 127.9, 127.6, 127.3, 127.2, 126.3, 125.9, 125.1, 77.9, 75.4, 56.0, 52.5, 36.4, 35.6, 32.1, 31.0, 30.9, 27.7, 27.2, 26.5, 23.8, 18.8, 18.4, 13.7; MS (ESI) m/z (%): 314.4 [$(M+NH_4^+)$, 100]; HRMS: 319.1675 $(M+Na)^+$, (319.1669).

***trans*-(±)-3-(4-(Naphthalen-2-yl)butyl)-4-propyloxetan-2-one (*trans*-9j).** $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.87-7.69 (m, 3H), 7.60 (s, 1H), 7.52-7.23 (m, 3H), 4.27-4.10 (m, 1H), 3.23-3.05 (m, 1H), 2.79 (t, $J$ = 7.2 Hz, 2H), 1.96-1.54 (m, 6H), 1.54-1.29 (m, 4H), 0.94 (t, $J$ = 7.4 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 171.6, 139.5, 133.5, 131.9, 127.9, 127.6, 127.3, 127.2, 126.3, 125.9, 125.1, 77.9, 56.0, 36.4, 35.6, 30.9, 27.7, 26.5, 18.4, 13.7.

***cis*-(±)-3-(4-(Naphthalen-2-yl)butyl)-4-propyloxetan-2-one (*cis*-9j).** $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.87-7.69 (m, 3H), 7.60 (s, 1H), 7.52-7.23 (m, 3H), 4.59-4.42 (m, 1H), 3.66-3.50 (m, 1H), 2.79 (t, $J$ = 7.2 Hz, 2H), 1.96-1.29 (m, 10H), 0.94 (t, $J$ = 7.4 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 172.2, 139.5, 133.5, 131.9, 127.9, 127.6, 127.3, 127.2, 126.3, 125.9, 125.1, 75.4, 52.5, 35.6, 32.1, 31.0, 27.2, 23.8, 18.8, 13.8.

**3-(4-Phenylbutyl)-4-propyloxetan-2-one *(*9k).** Mixture of diastereomers (d.r. 7:3 *trans*:*cis*). Yield 68%; Oil; $^1$H NMR (200 MHz, $CDCl_3$): $\delta$ 7.38-7.10 (m, 5H), 4.59-4.47 (m, 0.3H), 4.27-4.14 (m, 0.7H), 3.65-3.55 (m, 0.3H), 3.22-3.13 (m, 0.7H), 2.65 (t, $J$ = 7.6 Hz, 2H), 1.96-1.54 (m, 6H), 1.54-1.30 (m, 4H), 0.97 (t, $J$ = 7.2 Hz, 3H); $^{13}$C NMR (50 MHz, $CDCl_3$): $\delta$ 172.2, 171.5, 142.0, 128.4, 128.3, 125.8, 77.9, 75.4, 55.9, 52.5, 36.4, 35.5, 32.1, 31.1, 31.0, 27.6, 27.1, 26.5, 23.7, 18.8, 18.3, 13.7; MS (ESI) m/z (%): 264.2 [($M+NH_4^+$), 100]; HRMS: 269.1509 $(M+Na)^+$ (269.1512).

***trans*-(±)-3-(4-Phenylbutyl)-4-propyloxetan-2-one *(trans*-9k).** The mixture of diastereomers was separated by column chromatograhpy affording the *trans* diastereomer in 64% yield. Oil; $^1$H NMR (600 MHz, $CDCl_3$): $\delta$ 7.32-7.29 (m, 2H), 7.23-7.18 (m, 3H), 4.23 (ddd, $J$ = 7.4 Hz, 6.0 Hz, 4.0 Hz, 1H), 3.18 (ddd, $J$ = 8.7 Hz, 6.7 Hz, 4.0 Hz, 1H), 2.65 (t, $J$ = 7.6 Hz, 2H), 1.92-1.81 (m, 2H), 1.80-1.74 (m, 1H), 1.73-1.66 (m, 3H), 1.55-1.36 (m, 4H), 1.00 (t, $J$ = 7.4 Hz, 3H); $^{13}$C NMR (150 MHz, $CDCl_3$): $\delta$ 171.5, 142.0, 128.3, 125.8, 77.9, 56.0, 36.4, 35.5, 31.0, 27.7, 26.5, 18.4, 13.7.

***cis*-(±)-3-(4-Phenylbutyl)-4-propyloxetan-2-one *(cis*-9k).** The mixture of diastereomers was separated by column chromatograhpy affording the *cis* diastereomer in 8% yield. Oil; $^1$H NMR (600 MHz, $CDCl_3$): $\delta$ 7.32-7.28 (m, 2H), 7.23-7.18 (m, 3H), 4.55 (ddd, $J$ = 9.8 Hz, 6.4 Hz, 3.5 Hz, 1H), 3.61 (dt, $J$ = 8.8 Hz, 6.7 Hz, 1H), 2.66 (t, $J$ = 7.6 Hz, 2H), 1.87-1.80 (m, 1H), 1.78-1.54 (m, 7H), 1.49-1.39 (m, 2H), 1.01 (t, $J$ = 7.3 Hz, 3H); $^{13}$C NMR (150 MHz, $CDCl_3$): $\delta$ 172.2, 142.1, 128.3, 125.8, 75.4, 52.6, 35.5, 32.2, 31.1, 27.2, 23.8, 18.9, 13.8.

***In vitro* $PLA_2$ activity assay.** The activities of human recombinant GVIA $iPLA_2$, GIV $cPLA_2$ and GV $sPLA_2$ were determined using a previously described radioactivity-

based group-specific mixed micelle assay.[41-43] The substrate was prepared using slightly different conditions for each enzyme to achieve optimum activity: (i) GIVA $cPLA_2$ mixed micelle substrate consisted of 400 μM Triton X-100, 95.3 μM PAPC, 1.7 μM arachidonyl-1-$^{14}$C PAPC, and 3 μM $PIP_2$ in a buffer containing 100 mM HEPES pH 7.5, 90 μM $CaCl_2$, 2 mM DTT, and 0.1 mg/ml BSA; (ii) GVIA $iPLA_2$ mixed micelle substrate consisted of 400 μM Triton X-100, 98.3 μM PAPC, and 1.7 μM arachidonyl-1-$^{14}$C PAPC in a buffer containing 100 mM HEPES pH 7.5, 2 mM ATP, and 4 mM DTT; and (iii) GV $sPLA_2$ mixed micelles substrate consisted of 400 μM Triton X-100, 98.3 μM PAPC, and 1.7 μM arachidonyl-1-$^{14}$C PAPC in a buffer containing 50 mM Tris-HCl pH 8.0, and 5 mM $CaCl_2$. The compounds were initially screened at 0.091 mole fraction (5 μL of 5 mM inhibitor in DMSO) in substrate (495 μL). $X_I(50)$ values were determined for compounds exhibiting greater than 95% inhibition. Inhibition plotting percentage of inhibition vs log (mole fraction) to calculate the reported $X_I(50)$ and its associated error.

For the *trans* and *cis* diastereomers of the most potent lactones, similar group specific $PLA_2$ assays were employed to determine the activity using a lipidomics-based mixed micelle assay as previously described.[44,45] The substrate for each enzyme consisted of 100 μM PAPC (except for GIVA $cPLA_2$ as noted), 400 μM of C12E8 surfactant, and 2.5 μM of 17:0 LPC internal standard. For GIVA $cPLA_2$, the total phospholipid concentration (100 μM) consisted of 97 μM PAPC and 3 μM of $PI(4,5)P_2$ which enhances the activity of the enzyme. A specific buffer was prepared to achieve optimum activity for each enzyme. The buffer for GIVA $cPLA_2$ contained 100 mM HEPES pH 7.5, 90 μM $CaCl_2$, and 2 mM DTT. For GVIA $iPLA_2$, the buffer consisted of 100 mM HEPES pH 7.5, 2 mM ATP, and 4 mM DTT. Finally, the buffer for GV $sPLA_2$ contained 50 mM Tris-HCl pH 8.0 and 5 mM $CaCl_2$. The enzymatic reaction

was performed in a 96 well-plate using a Benchmark Scientific H5000-H MultiTherm heating shaker for 30 min at 40 °C. Each reaction was quenched with 120 µL of methanol/acetonitrile (80/20, v/v), and the samples were analyzed using a HPLC-MS system. A blank experiment, which did not contain enzyme, was also included for each substrate to determine the non-enzymatic hydrolysis product and to detect any changes in the intensity of the 17:0 LPC internal standard.

**Docking calculations.** Enzyme structures were optimized using the PPW. The structures of the inhibitors were sketched using Maestro sketcher and they were optimized using LigPrep. Glide was used for the rigid-docking of the compounds into the enzyme active site. The grid required for the docking procedure was generated using a scaling factor of 1.0 and partial charge cutoff of 0.25, while *X*, *Y*, *Z* dimensions of the inner box were set to 12 Å. For the inhibitor docking a scaling factor of 0.8 and partial charge cutoff of 0.15 were used that allow complete flexibility of the structures. The poses were selected according to the binding mode and the XP GScore. The Glide Extra-Precision (XP) scoring function was used for the calculations.[54]

***β-Cell apoptosis* ± *inhibitor trans*-9k.** INS-1 cells were generated and cultured as previously described.[47] Briefly, the cells were cultured in RPMI 1640 medium, containing 11 mM glucose, 10% fetal calf serum, 10 mM HEPES buffer, 2 mM glutamine, 1 mM sodium pyruvate, 50 mM mercaptoethanol (BME), and 0.1% (w/v) each of penicillin and streptomycin in cell culture conditions (37 °C, 5% $CO_2$ /95% air), as described.[55] The cells were treated with vehicle (DMSO, 1 µL/mL) alone or with IL-1β (100 U/mL) + IFNγ (300 U/mL) for 16 h in the absence or presence of

*trans*-**9k** (0.10-10 μM). The cells were then processed for apoptosis, by TUNEL analyses, as described.[47,50] Apoptotic cells (green fluorescence) and total number of cells, identifed by nuclear DAPI (blue) stain, in 6 fields on each slide were counted. Each slide represented one replicate (n=3-12). Percent apoptotic cells relative to total number of cells in each field was calculated and an average of the 6 fields/replicate was generated. The replicates were then averaged to generate means ± SEM for each condition and these are presented in Fig. 5.

## Associated Content

### Supporting Information.

Code numbers of tested compounds, elemental analyses of synthesized compounds, XP GScores for the all the diastereomers of **9k**, chromatographs of **9k**, *trans*-**9k** and *cis*-**9k**, NMR spectra of *trans*-**9k** and *cis*-**9k**. (PDF)

Molecular formula strings and inhibition data (CSV)

This material is available free of charge via the Internet at http://pubs.acs.org.

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## ABBREVIATIONS

ATP, adenosine triphosphate; BSA, bovine serum albumin; DAPI, 4′,6-diamidino-2-phenylindole; DMF, N,N-dimethylformamide; DMSO, dimethyl sulfoxide; DTT, dithiothreitol; EtOAc, ethyl acetate; GIVA cPLA$_2$, Group IVA cytosolic phospholipase A$_2$; GVIA iPLA$_2$, Group VIA calcium-independent phospholipase A$_2$; GV sPLA$_2$, Group V secreted phospholipase A$_2$; HEPES, 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid; IL-1β, interleukin 1β; INFγ, interferon γ; LDA, lithium diisopropylamide; PAPC, 1-palmitoyl-2-arachidonylphosphatidylcholine; PIP$_2$, phosphatidyl inositol (4,5)-bisphosphate; PPW, protein preparation wizard; THF, tetrahydrofuran; Tris-HCl, tris(hydroxymethyl)aminomethane hydrochloride**.**

## References

1. Dennis, E. A.; Cao, J.; Hsu, Y. H.; Magrioti, V.; Kokotos, G. Phospholipase A$_2$ enzymes: Physical structure, biological function, disease implication,

chemical inhibition, and therapeutic intervention. *Chem. Rev.* **2011**, *111*, 6130-6185.

2. Winstead, M. V.; Balsinde, J.; Dennis, E. A. Calcium-independent phospholipase $A_2$: Structure and function. *Biochim. Biophys. Acta.* **2000**, *1488*, 28-39.
3. Cedars, A.; Jenkins, C. M.; Mancuso, D. J.; Gross, R. W. J. Calcium independent phospholipases in the heart: Mediators of cellular signaling, bioenergetics and ischemia-induced electrophysiologic dysfunction. *J. Cardiovasc. Pharmacol*. **2009**, *53*, 277-289.
4. Lei, X.; Barbour, S. E.; Ramanadham, S. Group VIA $Ca^{2+}$-independent phospholipase $A_2$ ($iPLA_2$beta) and its role in beta-cell programmed cell death, *Biochimie* **2010**, *92*, 627-637.
5. Ramanadham, S.; Ali, T.; Ashley, J. W.; Bone, R. N.; Hancock, W. D.; Lei, X. Calcium-independent phospholipases $A_2$ and their roles in biological processes and diseases. *J. Lipid Res.* **2015**, 56, 1643-1668.
6. Hazen, S. L.; Stuppy, R. J.; Gross, R. W. Purification and characterization of canine myocardial cytosolic phospholipase $A_2$. A calcium-independent phospholipase with absolute f1-2 regiospecificity for diradyl glycerophospholipids. *J. Biol. Chem.* **1990**, *265*, 10622-10630.
7. Ramanadham, S.; Wolf, M. J.; Jett, P. A.; Gross, R. W.; Turk, J. Characterization of an ATP-stimulatable $Ca^{2+}$-independent phospholipase $A_2$ from clonal insulin-secreting HIT cells and rat pancreatic islets: A possible molecular component of the beta-cell fuel sensor. *Biochemistry* **1994**, *33*, 7442-7452.

8. Ackermann, E. J.; Kempner, E. S.; Dennis. E. A. $Ca^{2+}$-independent cytosolic phospholipase $A_2$ from macrophage-like P388D1 cells. Isolation and characterization. *J. Biol. Chem.* **1994**, *269*, 9227-9233.
9. Tang, J.; Kriz, R. W.; Wolfman, N.; Shaffer, M.; Seehra, J.; Jones, S. S. A novel cytosolic calcium independent phospholipase $A_2$ contains eight ankyrin motifs. *J. Biol. Chem.* **1997**, *272*, 8567-8575.
10. Balboa, M. A.; Balsinde, J.; Jones, S. S.; Dennis, E. A. Identity between the $Ca^{2+}$-independent phospholipase $A_2$ enzymes from P388D1 macrophages and Chinese hamster ovary cells. *J. Biol. Chem.* **1997**, *272*, 8576-8580.
11. Ma, Z.; Ramanadham, S.; Kempe, K.; Chi, X. S.; Ladenson, J.; Turk. J. Pancreatic islets express a $Ca^{2+}$-independent phospholipase $A_2$ enzyme that contains a repeated structural motif homologous to the integral membrane protein binding domain of ankyrin. *J. Biol. Chem.* **1997**, *272*, 11118-11127.
12. Lio, Y. C.; Dennis, E. A. Interfacial activation, lysophospholipase and transacylase activity of group VI $Ca^{2+}$-independent phospholipase $A_2$. *Biochim. Biophys. Acta* **1998**, *1392*, 320-332.
13. Ong, W.-Y.; Farooqui, T.; Kokotos, G.; Farooqui, A. A. Synthetic and natural inhibitors of phospholipases $A_2$: Their importance for understanding and treatment of neurological disorders. *ACS Chem. Neurosci.* **2015**, *6*, 814-831.
14. Kokotou, M. G.; Limnios, D.; Nikolaou, A.; Psarra, A.; Kokotos, G. Inhibitors of phospholipase $A_2$ and their therapeutic potential: An update on patents (2012–2016). *Expert Opin. Ther. Pat.* **2017***, 27*, 217-225.
15. Magrioti, V.; Kokotos, G. Phospholipase $A_2$ inhibitors for the treatment of inflammatory diseases: A patent review (2010 – present). *Expert Opin. Ther. Pat.* **2013**, *23*, 333-344.

16. Leslie, C. C. Cytosolic phospholipase $A_2$: Physiological function and role in disease. *J. Lipid Res.* **2015**, *56*, 1386-1402.

17. Murakami, M.; Taketomi, Y.; Miki, Y.; Sato, H.; Yamamoto, K.; Lambeau, G. Emerging roles of secreted phospholipase $A_2$ enzymes: The 3rd edition. *Biochimie.* **2014**, *107 Pt A*, 105-113.

18. Ackermann, E. J.; Conde-Frieboes, K.; Dennis, E. A. Inhibition of macrophage $Ca^{2+}$-independent phospholipase $A_2$ by bromoenol lactone and trifluoromethyl ketones. *J. Biol. Chem.* **1995**, 270, 445-450.

19. Balsinde, J.; Dennis, E. A. Bromoenol lactone inhibits magnesium-dependent phosphatidate phosphohydrolase and blocks triacylglycerol biosynthesis in mouse $P388D_1$ macrophages. *J. Biol. Chem.* **1996**, *271*, 31937-31941.

20. Bone, R. N.; Gai, Y.; Magrioti, V.; Kokotou, M. G.; Ali, T.; Lei, X.; Tse, H. M.; Kokotos, G.; Ramanadham, S. Inhibition of $Ca^{2+}$-independent phospholipase $A_2\beta$ ($iPLA_2\beta$) ameliorates islet infiltration and incidence of diabetes in NOD mice. *Diabetes* **2015,** *64*, 541-554.

21. Malhotra, A.; Edelman-Novemsky, I.; Xu, Y.; Plesken, H.; Ma, J.; Schlame, M.; Ren, M. C., Role of calcium-independent phospholipase $A_2$ in the pathogenesis of Barth syndrome. *PNAS* **2009,** *106*, 2337-2341.

22. Abi Nahed, R.; Martinez, G.; Escoffier, J.; Yassine, S.; Karaouzène, T.; Hograindleur, J. P.; Turk, J.; Kokotos, G.; Ray, P. F.; Bottari, S.; Lambeau, G.; Hennebicq, S.; Arnoult, C. Progesterone-induced acrosome exocytosis requires sequential involvement of calcium-independent phospholipase $A_2\beta$ ($iPLA_2\beta$) and Group X secreted phospholipase $A_2$ ($sPLA_2$). *J. Biol. Chem.* **2016**, 291, 3076-3089.

23. Baskakis, C.; Magrioti, V.; Cotton, N.; Stephens, D.; Constantinou-Kokotou, V.; Dennis, E.A.; Kokotos, G. Synthesis of polyfluoro ketones for selective inhibition of human phospholipase $A_2$ enzymes. *J. Med. Chem.* **2008**, *51*, 8027-8037.

24. Kokotos, G.; Hsu, Y.-H.; Burke, J. E.; Baskakis, C.; Kokotos, C. G.; Magrioti V.; Dennis, E. A. Potent and selective fluoroketone inhibitors of group VIA calcium-independent phospholipase $A_2$. *J. Med. Chem.* **2010**, *53*, 3602-3610.

25. Magrioti, V.; Nikolaou, A.; Smyrniotou, A.; Shah, I.; Constantinou-Kokotou, V.; Dennis, E. A.; Kokotos, G. New potent and selective polyfluoroalkyl ketone inhibitors of GVIA calcium-independent phospholipase $A_2$. *Bioorg. Med. Chem.* **2013**, *21*, 5823-5829.

26. Kalyvas, A.; Baskakis, C.; Magrioti, V.; Constantinou-Kokotou, V.; Stephens, D.; Lopez-Vales, R.; Lu, J. Q.; Yong, V. W.; Dennis, E. A.; Kokotos, G.; David. S. Differing roles for members of the phospholipase $A_2$ superfamily in experimental autoimmune encephalomyelitis. *Brain* **2009**, *132*, 1221-1235.

27. Li, J.; Zhao, Z.; Antalis, C.; Zhao, Z.; Emerson, R.; Wei, G.; Zhang, S; Zhang, Z.-Y.; Xu, Y. Combination therapy of an inhibitor of group VIA phospholipase $A_2$ with paclitaxel is highly effective in blocking ovarian cancer development. *Am. J. Pathol.* **2011**, 179, 452-461.

28. Ali, T.; Kokotos, G.; Magrioti, V.; Bone, R. N.; Mobley, J. A.; Hancock, W.; Ramanadham, S. Characterization of FKGK18 as inhibitor of Group VIA $Ca^{2+}$-independent phospholipase $A_2$ (iPLA$_2\beta$): Candidate drug for preventing beta-cell apoptosis and diabetes. *PLOS ONE* **2013**, *8*: e71748.

29. Hsu, Y. H.; Bucher, D.; Cao, J.; Li, S.; Yang, S. W.; Kokotos, G.; Woods, V. L., Jr., McCammon, J. A.; Dennis, E. A. Fluoroketone inhibition of $Ca^{2+}$-

independent phospholipase $A_2$ through binding pocket association defined by hydrogen/deuterium exchange and molecular dynamics, *J. Am. Chem. Soc.* **2013**, *135*, 1330-1337.

30. Bucher, D.; Hsu, Y. H.; Mouchlis, V. D.; Dennis, E. A.; McCammon, J. A. Insertion of the $Ca^{2+}$-independent phospholipase $A_2$ into a phospholipid bilayer via coarse-grained and atomistic molecular dynamics simulations. *PLoS Comput. Biol.* **2013,** *9*, e1003156.
31. Mouchlis, V.D.; Bucher, D.; McCammon, J.A.; Dennis, E.A. Membranes serve as allosteric activators of phospholipase $A_2$, enabling it to extract, bind, and hydrolyze phospholipid substrates. *PNAS* **2015**, *112*, E516eE525.
32. Mouchlis, V. D.; Limnios, D.; Kokotou, M. G.; Barbayianni, E.; Kokotos, G.; McCammon, J. A.; Dennis, E. A. Development of potent and selective inhibitors for group VIA calcium-independent phospholipase $A_2$ guided by molecular dynamics and structure-activity relationships. *J. Med. Chem.* **2016**, 59, 4403-4414.
33. Weibel, E. K.; Hadvary, P.; Hochuli, E.; Kupfer, E.; Lengsfeld, H. Lipstatin, an inhibitor of pancreatic lipase, produced by Streptomyces Toxytricini. I. Producing organism, fermentation, isolation and biological activity. *J. Antibiot.* **1987**, *40*, 1081-1085.
34. Borgström, B. Mode of action of tetrahydrolipstatin: a derivative of the naturally occurring lipase inhibitor lipstatin. *Biochim. Biophys. Acta.* **1988**, *962*, 308-316.
35. Bauer, R. A. Covalent inhibitors in drug discovery: from accidental discoveries to avoided liabilities and designed therapies, *Drug Discov. Today* **2015**, *20*, 1061-1073.

36. Kluge, A. F.; Petter, R. C. Acylating drugs: redesigning natural covalent inhibitors, *Curr. Opin. Chem. Biol.* **2010**, *14*, 421-427.

37. Black, T. H.; DuBay III, W. J. A new synthesis of substituted spiro butyrolactones via dyotropic rearrangement. *Tetrahedron Lett.* **1987,** *28*, 4787-4788.

38. Schmidt, Y.; Breit, B. Enantioselective total synthesis and determination of absolute configuration of vittatalactone. *Org. Lett.* **2009,** *11*, 4767-4769.

39. Meier, P.; Broghammer, F.; Latendorf, K.; Rauhut, G.; Peters, R. Cooperative Al(Salen)-pyridinium catalysts for the asymmetric synthesis of *trans*-configured β-lactones by [2+2]-cyclocondensation of acylbromides and aldehydes: Investigation of pyridinium substituent effects. *Molecules* **2012**, *17*, 7121-7150.

40. Purohit, V. C.; Richardson, R. D.; Smith, J. W.; Romo, D. Practical, catalytic, asymmetric synthesis of *β*-lactones via a sequential ketene dimerization/hydrogenation process: Inhibitors of the thioesterase domain of fatty acid synthase. *J. Org. Chem.* **2006**, *71*, 4549-4558.

41. Kokotos, G.; Six, D.A.; Loukas, V.; Smith, T.; Constantinou-Kokotou, V.; Hadjipavlou-Litina, D.; Kotsovolou, S.; Chiou, A.; Beltzner, C. C.; Dennis E. A. Inhibition of group IVA cytosolic phospholipase $A_2$ by novel 2-oxoamides *in vitro*, in cells and *in vivo*. *J. Med. Chem.* **2004**, *47*, 3615-3628.

42. Stephens, D.; Barbayianni, E.; Constantinou-Kokotou, V.; Peristeraki, A.; Six, D. A.; Cooper, J.; Harkewicz, R.; Deems, R. A.; Dennis, E. A.; Kokotos, G. Differential inhibition of group IVA and group VIA phospholipases $A_2$ by 2-oxoamides. *J. Med. Chem.* **2006**, *49*, 2821-2828.

43. Six, D. A.; Barbayianni. E,; Loukas, V.; Constantinou-Kokotou, V.; Hadjipavlou-Litina, D.; Stephens, D.; Wong, A.C.; Magrioti, V.; Moutevelis-Minakakis, P.; Baker, S. F.; Dennis, E. A.; Kokotos, G. Structure-activity relationship of 2-oxoamide inhibition of group IVA cytosolic phospholipase $A_2$ and group V secreted phopholipase $A_2$. *J. Med. Chem.* **2007**, *50*, 4222-4235.

44. Mouchlis, V. D.; Chen, Y.; McCammon, J. A.; Dennis, E. A. Membrane allostery and unique hydrophobic sites promote enzyme substrate specificity. *J. Am. Chem. Soc.* **2018**, *140*, 3285-3291.

45. Mouchlis, V. D.; Armando, A.; Dennis, E. A. Substrate specific inhibition constants for phospholipase $A_2$ acting on unique phospholipid substrates in mixed micelles and membranes using lipidomics. *J. Med. Chem.* **2019**, 62, DOI: 10.1021/acs.jmedchem.8b01568.

46. Dennis, E. A.; Norris, P. C. Eicosanoid storm in infection and inflammation. *Nat. Rev. Immunol.* **2015,** *15*, 511-523.

47. Ramanadham, S.; Hsu, F. F.; Zhang, S.; Jin, C.; Bohrer, A.; Song, H.; Bao, S.; Ma, Z.; Turk, J. Apoptosis of insulin-secreting cells induced by endoplasmic reticulum stress is amplified by overexpression of group VIA calcium-independent phospholipase $A_2$ (iPLA$_2\beta$) and suppressed by inhibition of iPLA$_2\beta$. *Biochemistry* **2004,** *43*, 918-930.

48. Lei, X.; Zhang, S.; Bohrer, A.; Bao, S.; Song, H.; Ramanadham, S. The group VIA calcium-independent phospholipase $A_2$ participates in ER stress-induced INS-1 insulinoma cell apoptosis by promoting ceramide generation via hydrolysis of sphingomyelins by neutral sphingomyelinase. *Biochemistry* **2007,** *46*, 10170-10185

49. 1. Lei, X.; Zhang, S.; Bohrer, A.; Ramanadham, S. Calcium-independent phospholipase $A_2$ (iPLA$_2\beta$)-mediated ceramide generation plays a key role in the cross-talk between the endoplasmic reticulum (ER) and mitochondria during ER stress-induced insulin-secreting cell apoptosis. *J. Biol. Chem.* **2008,** *283*, 34819-34832.

50. Lei, X.; Zhang, S.; Barbour, S. E.; Bohrer, A.; Ford, E. L.; Koizumi, A.; Papa, F. R.; Ramanadham, S. Spontaneous development of endoplasmic reticulum stress that can lead to diabetes mellitus is associated with higher calcium-independent phospholipase $A_2$ expression: A role for regulation by SREBP-1. *J. Biol. Chem.* **2010,** *285*, 6693-6705.

51. Lei, X.; Bone, R. N.; Ali, T.; Wohltmann, M.; Gai, Y.; Goodwin, K. J.; Bohrer, A. E.; Turk, J.; Ramanadham, S. Genetic modulation of islet β-cell iPLA$_2$β expression provides evidence for its impact on β-cell apoptosis and autophagy. *Islets* **2013,** *5*, 29-44.

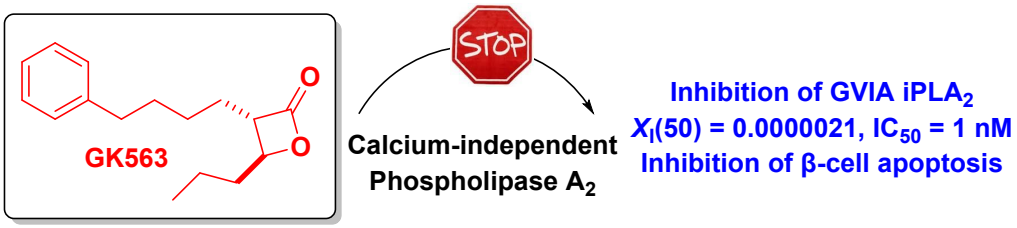
52. Lei, X.; Zhang, S.; Bohrer, A.; Barbour, S. E.; Ramanadham, S. Role of calcium-independent phospholipase $A_2$β in human pancreatic islet β-cell apoptosis. *Am. J. Physiol. Endocrinol. Metab.* **2012,** *303*, E1386-1395.

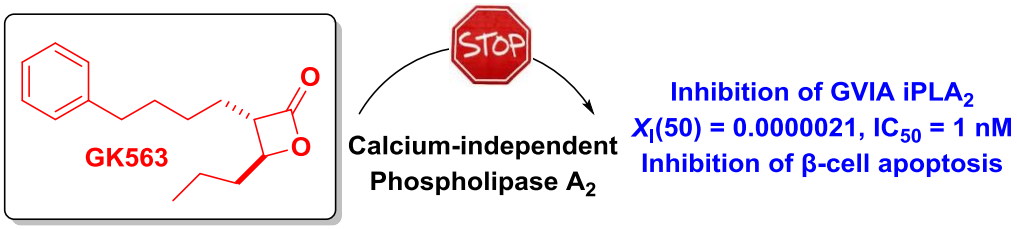
53. Huingen, R.; Rietz, U. Mittlere Ringe, XIV. Darstellung und cyclisierung der ω-[naphthyl-(2)]-fettsäuren. *Chem. Ber.* **1957,** *90*, 2768-2784.

54. Friesner, R. A.; Murphy, R. B.; Repasky, M. P.; Frye, L. L.; Greenwood, J. R.; Halgren, T. A.; Sanschagrin, P. C.; Mainz, D. T. Extra precision glide: Docking and scoring incorporating a model of hydrophobic enclosure for protein-ligand complexes. *J. Med. Chem.* **2006,** *49*, 6177-6196.

55. Ma, Z.; Zhang, S.; Turk, J.; Ramanadham, S. Stimulation of insulin secretion and associated nuclear accumulation of iPLA(2)beta in INS-1 insulinoma cells. *Am. J. Physiol. Endocrinol. Metab.* **2002**, *282*, E820-833.

## TABLE OF CONTENTS GRAPHIC




GK563
O
O
STOP
Calcium-independent
Phospholipase A2
Inhibition of GVIA iPLA2
XI(50) = 0.0000021, IC50 = 1 nM
Inhibition of β-cell apoptosis