



(12) **DEMANDE DE BREVET CANADIEN
CANADIAN PATENT APPLICATION**

(13) **A1**

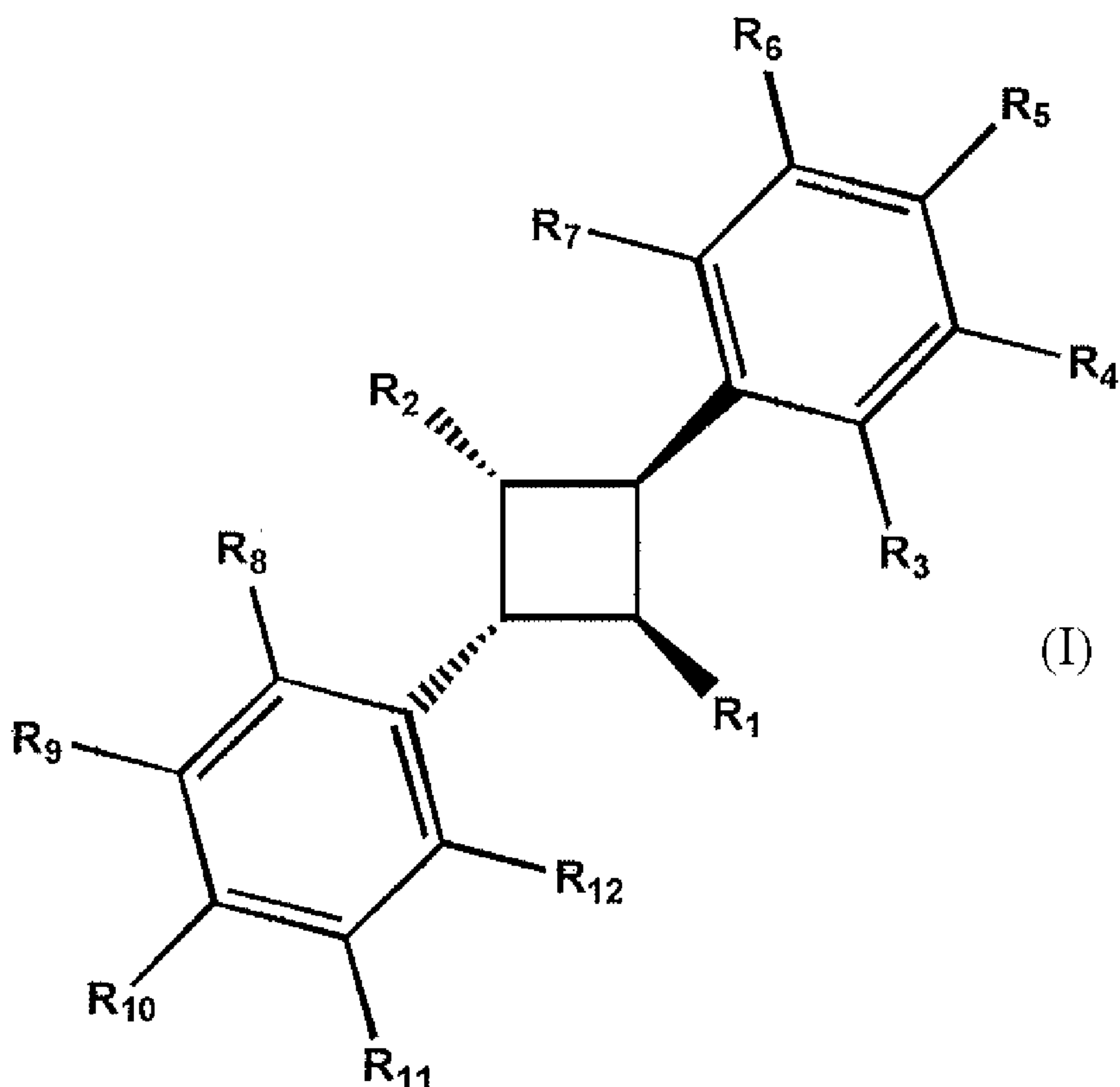
(86) Date de dépôt PCT/PCT Filing Date: 2017/03/10
(87) Date publication PCT/PCT Publication Date: 2017/09/14
(85) Entrée phase nationale/National Entry: 2018/09/11
(86) N° demande PCT/PCT Application No.: US 2017/021711
(87) N° publication PCT/PCT Publication No.: 2017/156354
(30) Priorité/Priority: 2016/03/11 (US62/307,262)

(51) Cl.Int./Int.Cl. *C07C 69/753* (2006.01),
A61K 31/167 (2006.01), *A61K 31/216* (2006.01),
A61K 31/35 (2006.01), *A61K 31/4192* (2006.01),
A61P 29/00 (2006.01), *A61P 43/00* (2006.01),
C07C 205/55 (2006.01), *C07C 233/25* (2006.01),
C07C 69/757 (2006.01), *C07D 249/06* (2006.01),
C07D 309/06 (2006.01)

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(54) Titre : DERIVES D'ACIDE α -TRUXILLIQUE ET COMPOSITIONS PHARMACEUTIQUES ASSOCIEES
(54) Title: α -TRUXILLIC ACID DERIVATIVES AND PHARMACEUTICAL COMPOSITIONS THEREOF



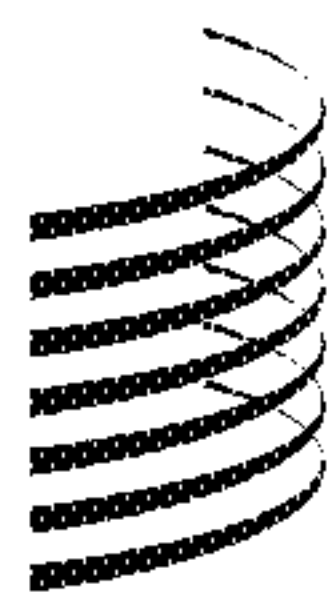
(57) Abrégé/Abstract:

The present invention provides a compound, and method of inhibiting the activity of a Fatty Acid Binding Protein (FABP) comprising contacting the FABP with a compound, said compound having the structure : Formula (I)

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(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property
Organization
International Bureau

WIPO | PCT



(10) International Publication Number

WO 2017/156354 A1

(43) International Publication Date
14 September 2017 (14.09.2017)

(51) International Patent Classification:

C07C 69/753 (2006.01) A61K 31/216 (2006.01)
 C07C 69/757 (2006.01) A61K 31/35 (2006.01)
 C07C 205/55 (2006.01) A61K 31/167 (2006.01)
 C07C 233/25 (2006.01) A61K 31/4192 (2006.01)
 C07D 309/06 (2006.01) A61P 43/00 (2006.01)
 C07D 249/06 (2006.01) A61P 29/00 (2006.01)

(21) International Application Number:

PCT/US2017/021711

(22) International Filing Date:

10 March 2017 (10.03.2017)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

62/307,262 11 March 2016 (11.03.2016) US

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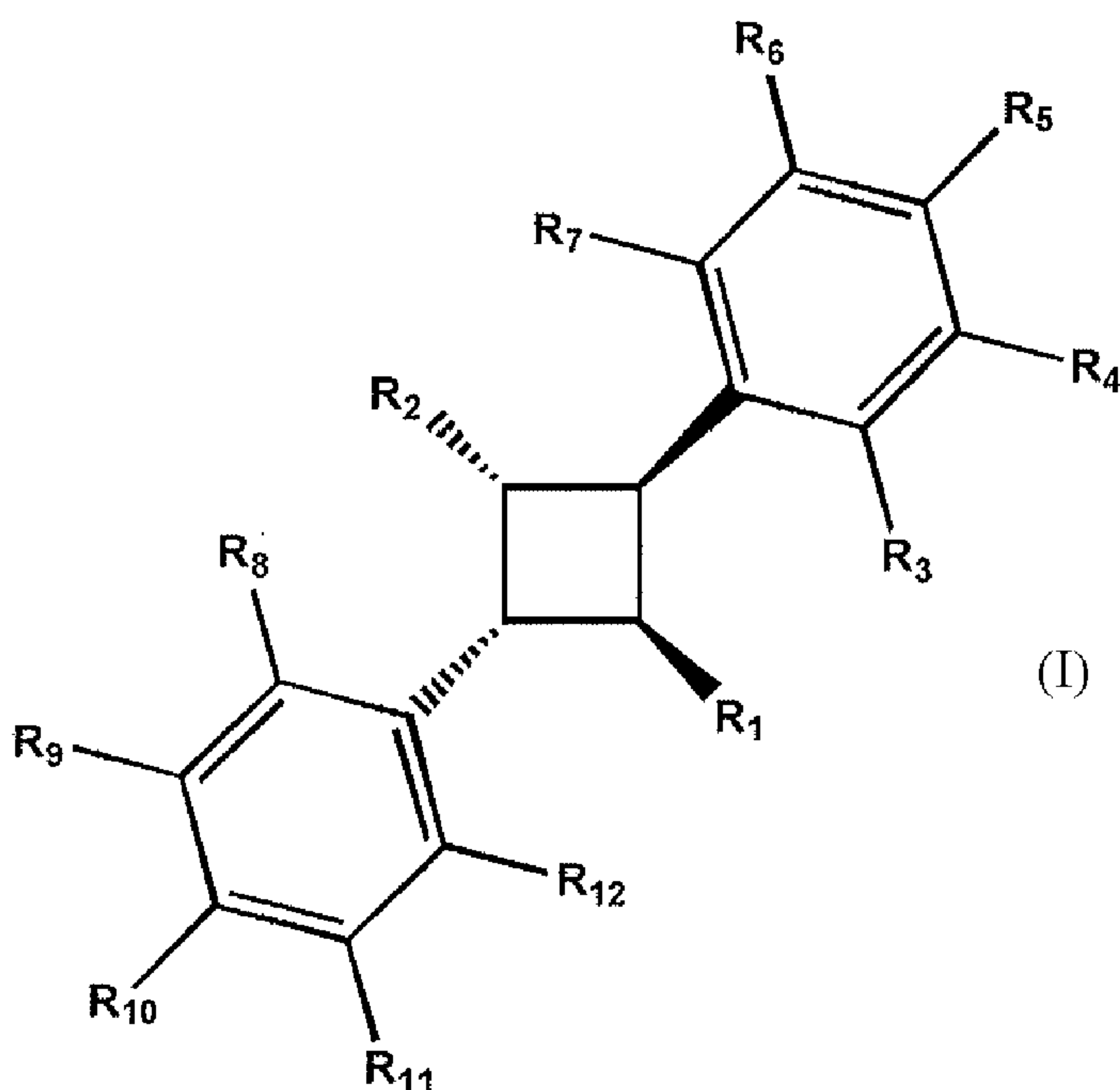
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(81) Designated States (unless otherwise indicated, for every
 kind of national protection available): AE, AG, AL, AM,
 AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY,
 BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DJ, DK, DM,
 DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT,
 HN, HR, HU, ID, IL, IN, IR, IS, JP, KE, KG, KH, KN,
 KP, KR, KW, KZ, LA, LC, LK, LR, LS, LU, LY, MA,
 MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG,
 NI, NO, NZ, OM, PA, PE, PG, PH, PL, PT, QA, RO, RS,
 RU, RW, SA, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY,
 TH, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN,
 ZA, ZM, ZW.

(84) Designated States (unless otherwise indicated, for every
 kind of regional protection available): ARIPO (BW, GH,
 GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, ST, SZ,
 TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU,

[Continued on next page]

(54) Title: α -TRUXILLIC ACID DERIVATIVES AND PHARMACEUTICAL COMPOSITIONS THEREOF

(57) Abstract: The present invention provides a com-
 pound, and method of inhibiting the activity of a Fatty
 Acid Binding Protein (FABP) comprising contacting
 the FABP with a compound, said compound having the
 structure : Formula (I)

WO 2017/156354 A1

TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK, SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, KM, ML, MR, NE, SN, TD, TG). **Published:** — *with international search report (Art. 21(3))*

**α -TRUXILLIC ACID DERIVATIVES AND PHARMACEUTICAL
COMPOSITIONS THEREOF**

5

This application claims priority of U.S. Provisional Application No. 62/307,262, filed March 11, 2016, the contents of which are hereby incorporated by reference.

10 Throughout this application, certain publications are referenced in parentheses. Full citations for these publications may be found immediately preceding the claims. The disclosures of these publications in their entireties are hereby incorporated by reference into this application in order to describe more fully the state of the art to which
15 this invention relates.

The invention was made with government support under Grant number DA032232, DA026953, and DA016419 awarded by the National Institutes of Health. The government has certain rights in the invention.

20

Background of the Invention

Lipids, owing to their water insolubility, require a variety of fatty acid binding protein (FABP) chaperones or transporters to carry them throughout cells (Furuhashi, M et al. 2008). The FABPs are part of the
25 pathway of anandamide inactivation by the fatty acid amide hydrolase (FAAH), an enzyme localized inside the cell on the endoplasmic reticulum. The endocannabinoid anandamide (arachidonoyl ethanolamide or AEA) is an uncharged neuromodulatory lipid that is inactivated through its cellular uptake by FABPs and subsequent hydrolysis by FAAH into ethanolamine and
30 arachidonic acid.

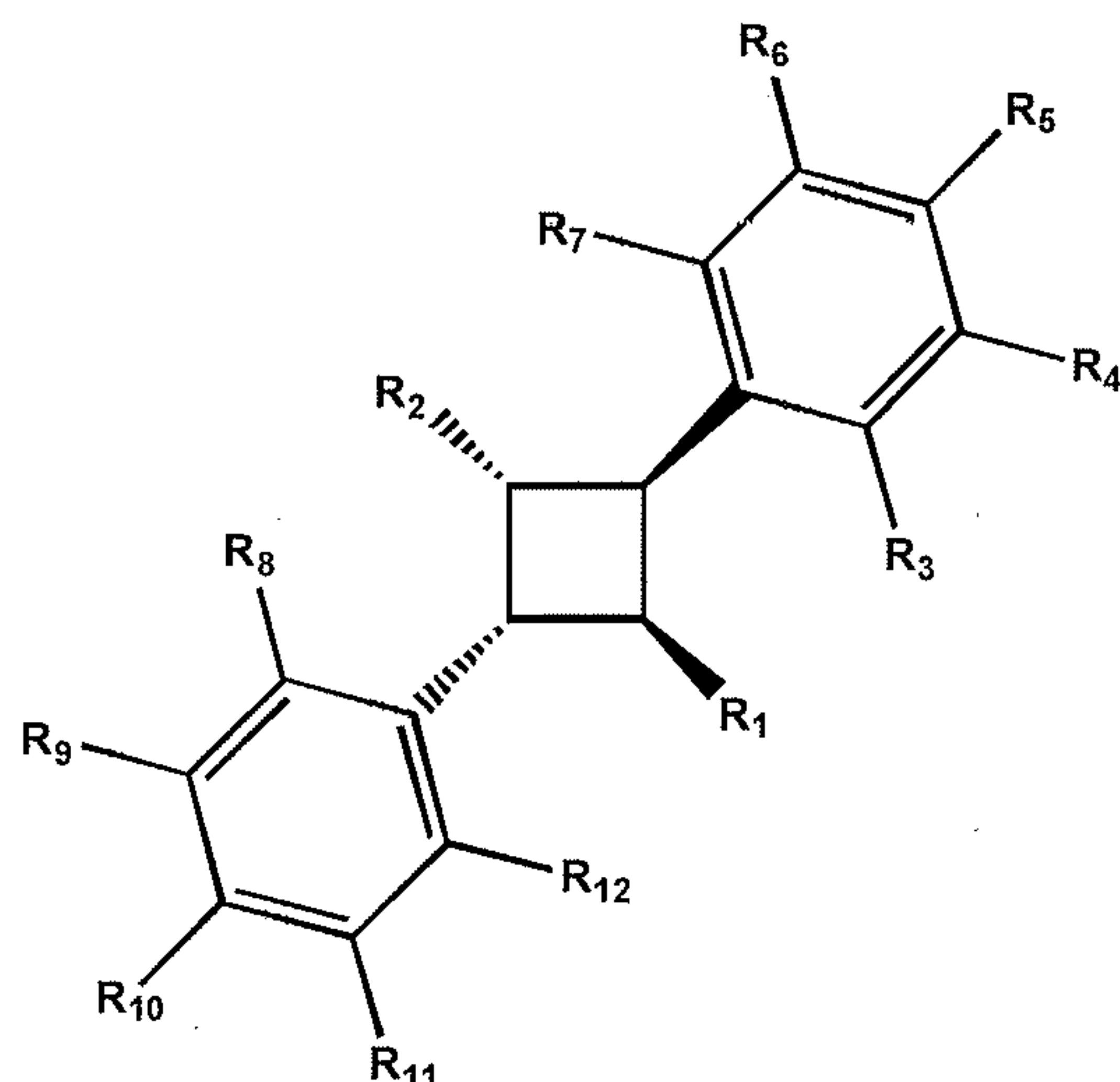
Cannabinoids such as anandamide have broad effects on the central nervous system (CNS) and influence, for example, movement, memory, nociception, endocrine regulation, thermoregulation, sensory perception, cognitive
35 functions, and mood. Similarly, genetic and pharmacological studies have revealed a broad role for endocannabinoid signaling in a variety of

physiological processes, including neuromodulator release, motor learning, synaptic plasticity, appetite, and pain sensation. Anandamide produces most of its pharmacological effects by binding and activating the cannabinoid receptor (CB-1 and CB-2) within the CNS. The increase
5 in extracellular anandamide caused by the inhibition of FABPs triggers activation of the cannabinoid receptor type 1 (CB-1) pathway leading to the relief of neurogenic and inflammatory pain.

Recently, it was shown that anandamide (an endocannabinoid) uses FABPs
10 such as FABP5 (E-FABP) and FABP7 (B-FABP) as intracellular transporters (Kaczocha, M. et al. 2009). FABPs are drug targets similar to FAAH since inhibitors of each decrease hydrolysis of anandamide and its uptake into cells, raising the levels of extracellular anandamide (Figure 1)
(Howlett, A.C. et al. 2011; Kaczocha, M. et al. 2012; Ahn, K. et al.
15 2009). Few specific FABP inhibitors have been described. There are those that were specifically designed for FABP4, such as BMS309403, which are important for the protective effects that they exert in metabolic syndrome and atherosclerosis (Barf, T. et al. 2009; Sulsky, R. et al.
2007). BMS309403 also binds other FABPs, such as FABP5 and FABP7, that
20 carry anandamide, as do other inhibitors originally designed to inhibit a putative anandamide transmembrane transporter (Kaczocha, M. et al. 2012).

Summary of the Invention

The present invention provides a compound of having the structure:



5 wherein

one of R_1 or R_2 is $-C(=O)OH$ and the other of R_1 or R_2 is $-C(=O)OR_{13}$ or $-C(=O)O\text{-alkyl-}R_{14}$,

wherein

R_{13} is cycloalkyl, aryl or heteroaryl, and

10 R_{14} is CF_3 , cycloalkyl, cycloheteroalkyl, aryl or heteroaryl; and $R_3, R_4, R_5, R_6, R_7, R_8, R_9, R_{10}, R_{11}$ and R_{12} are each independently, H $-OR_{15}$ or halogen

wherein R_{15} is H, C_{1-10} alkyl, C_{2-10} alkenyl, C_{2-10} alkynyl, aryl, or heteroaryl,

15

wherein when one of R_1 or R_2 is $-C(=O)OH$ and $R_3, R_4, R_5, R_6, R_7, R_8, R_9, R_{10}, R_{11}$ and R_{12} are each H, then the other of R_1 or R_2 is other than $-C(=O)OR_{13}$ where R_{13} is 1-naphthalene or 2-naphthalene or $-C(=O)O\text{-alkyl-}R_{14}$ where the alkyl is a branched C_2 alkyl and the R_{14} is phenyl,

20

or an enantiomer or racemate thereof;

or a pharmaceutically acceptable salt thereof.

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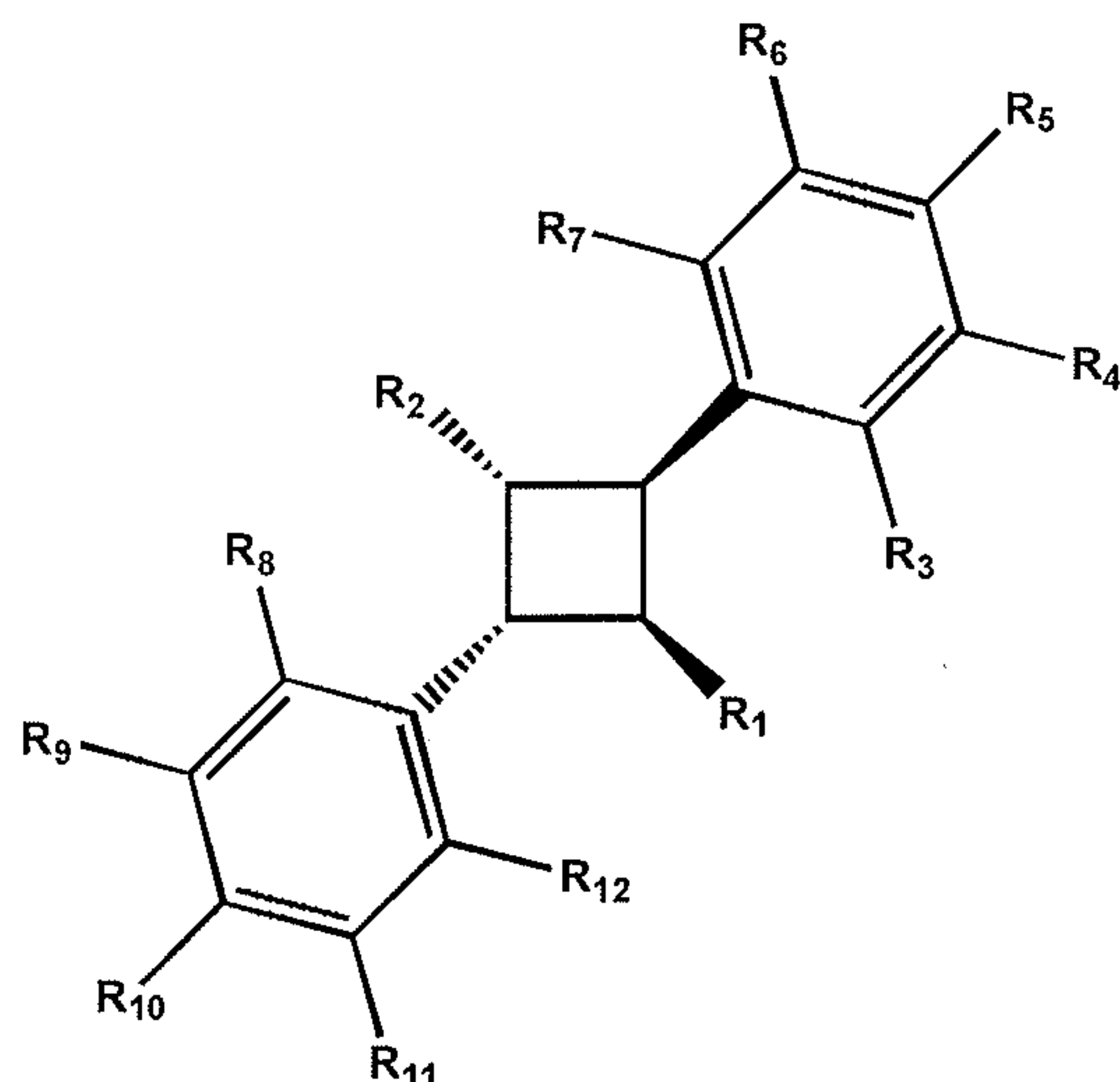
Brief Description of the Figures

Fig. 1: Scheme demonstrating anandamide inactivation and FABP drug target. Anandamide crosses the membrane by diffusion but requires FABPs for transport through the cytoplasm to the endoplasmic reticulum for breakdown by FAAH. FABP inhibitors prevent AEA from being delivered to FAAH for breakdown resulting in increased AEA levels at the receptor.

Fig. 2: Bar graphs demonstrating latency (sec) of baseline, Freund's complete adjuvant (CFA), vehicle, Compound A, Compound B, Compound C, and Compound D; and compound ($\mu\text{g/mL}$) of Compound B and Compound A. Compound A refers to Compound **3**, Compound B is a methyl ester of Compound **3**, Compound C refers to Compound **3a**, Compound D refers to Compound **4a**.

Detailed Description of the Invention

The present invention provides a compound of having the structure:



5 wherein

one of R_1 or R_2 is $-C(=O)OH$ and the other of R_1 or R_2 is $-C(=O)OR_{13}$ or $-C(=O)O\text{-alkyl-}R_{14}$,

wherein

R_{13} is cycloalkyl, aryl or heteroaryl, and

10 R_{14} is CF_3 , cycloalkyl, cycloheteroalkyl, aryl or heteroaryl; and $R_3, R_4, R_5, R_6, R_7, R_8, R_9, R_{10}, R_{11}$ and R_{12} are each independently, H $-OR_{15}$ or halogen

wherein R_{15} is H, C_{1-10} alkyl, C_{2-10} alkenyl, C_{2-10} alkynyl, aryl, or heteroaryl,

15

wherein when one of R_1 or R_2 is $-C(=O)OH$ and $R_3, R_4, R_5, R_6, R_7, R_8, R_9, R_{10}, R_{11}$ and R_{12} are each H, then the other of R_1 or R_2 is other than $-C(=O)OR_{13}$ where R_{13} is 1-naphthalene or 2-naphthalene or $-C(=O)O\text{-alkyl-}R_{14}$ where the alkyl is a branched C_2 alkyl and the R_{14} is phenyl,

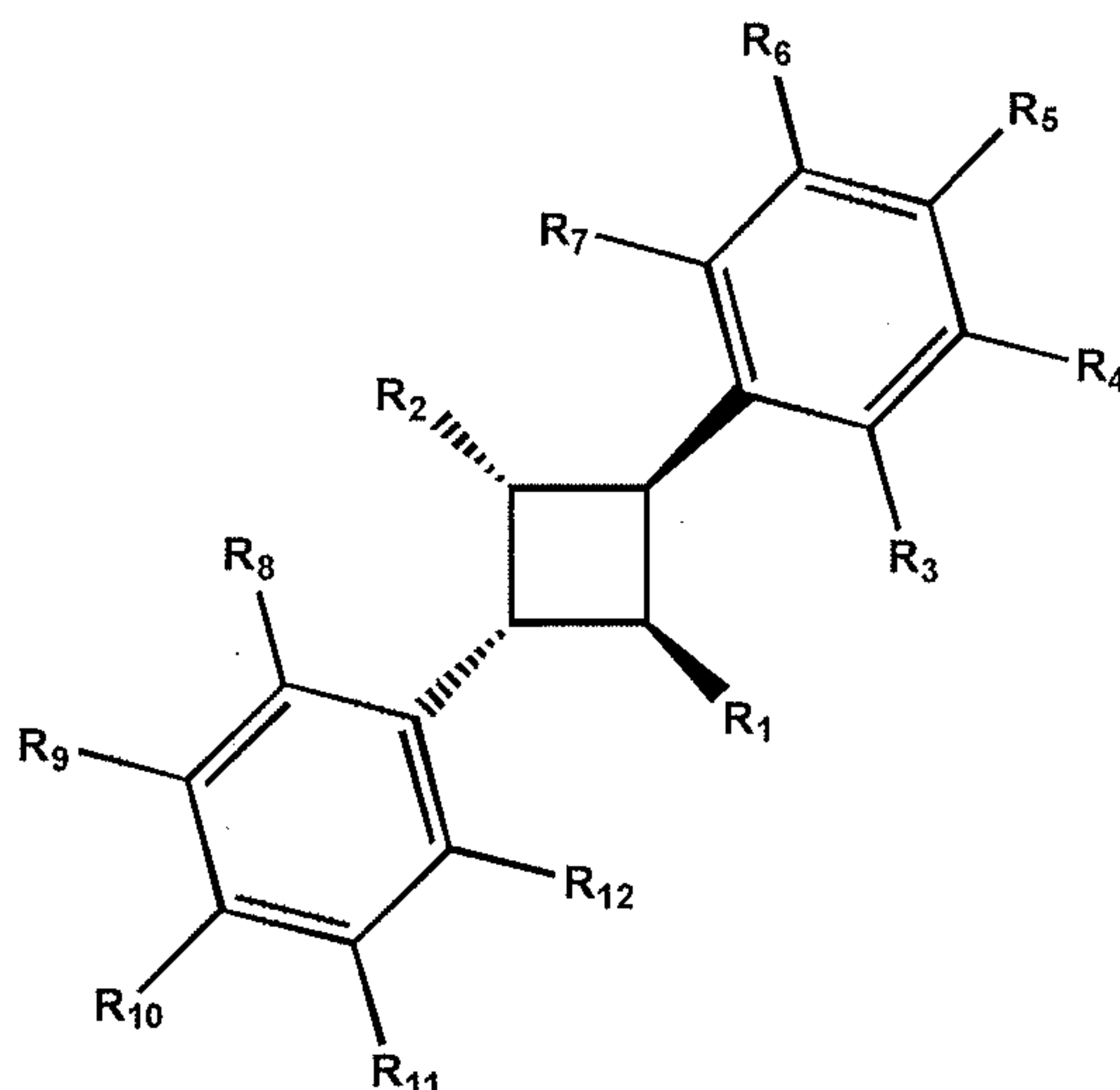
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or an enantiomer or racemate thereof;

or a pharmaceutically acceptable salt or ester thereof.

25

The present invention provides a compound of having the structure:



wherein

5 one of R_1 or R_2 is $-C(=O)OH$ and the other of R_1 or R_2 is $-C(=O)OR_{13}$ or $-C(=O)O\text{-alkyl-}R_{14}$,

wherein

R_{13} is cycloalkyl, aryl or heteroaryl, and

R_{14} is CF_3 , cycloalkyl, aryl or heteroaryl; and

10 $R_3, R_4, R_5, R_6, R_7, R_8, R_9, R_{10}, R_{11}$ and R_{12} are each independently, H $-OR_{15}$ or halogen

wherein R_{15} is H, C_{1-10} alkyl, C_{2-10} alkenyl, C_{2-10} alkynyl, aryl, or heteroaryl,

15 wherein when one of R_1 or R_2 is $-C(=O)OH$ and $R_3, R_4, R_5, R_6, R_7, R_8, R_9, R_{10}, R_{11}$ and R_{12} are each H, then the other of R_1 or R_2 is other than $-C(=O)OR_{13}$ where R_{13} is 1-naphthalene or 2-naphthalene or $-C(=O)O\text{-alkyl-}R_{14}$ where the alkyl is a branched C_2 alkyl and the R_{14} is phenyl,

20 or an enantiomer or racemate thereof;

or a pharmaceutically acceptable salt thereof.

In some embodiments, the compound wherein when one of R_1 or R_2 is $-C(=O)OH$ and $R_3, R_4, R_5, R_6, R_7, R_8, R_9, R_{10}, R_{11}$ and R_{12} are each H, then the other of R_1 or R_2 is other than $-C(=O)OR_{13}$ where R_{13} is tolyl, 1-naphthalene or 2-naphthalene or $-C(=O)O$ -alkyl- R_{14} where the alkyl is a branched C_2 alkyl and the R_{14} is phenyl.

In some embodiments, the compound wherein when one of R_1 or R_2 is $-C(=O)OH$ and the other of R_1 or R_2 is $-C(=O)OR_{13}$ where R_{13} is 1-naphthalene or 2-naphthalene, then one of $R_3, R_4, R_5, R_6, R_7, R_8, R_9, R_{10}, R_{11}$ and R_{12} is other than $-H$.

In some embodiments, the compound wherein when one of R_1 or R_2 is $-C(=O)OH$ and the other of R_1 or R_2 is $-C(=O)OR_{13}$ where R_{13} is 1-naphthalene or 2-naphthalene, then two of $R_3, R_4, R_5, R_6, R_7, R_8, R_9, R_{10}, R_{11}$ and R_{12} is other than $-H$.

In some embodiments, the compound wherein when one of R_1 or R_2 is $-C(=O)OH$ and the other of R_1 or R_2 is $-C(=O)OR_{13}$ where R_{13} is 1-naphthalene or 2-naphthalene, then four of $R_3, R_4, R_5, R_6, R_7, R_8, R_9, R_{10}, R_{11}$ and R_{12} is other than $-H$.

In some embodiments, the compound wherein

one of R_1 or R_2 is $-C(=O)R_{13}$,

wherein R_{13} is cycloalkyl, aryl or heteroaryl; and

the other of R_1 or R_2 is $-C(=O)OH$.

In some embodiments, the compound wherein

one of R_1 or R_2 is $-C(=O)O$ -alkyl- R_{14} ,

wherein R_{14} is CF_3 , cycloalkyl, aryl or heteroaryl; and

the other of R_1 or R_2 is $-C(=O)OH$.

In some embodiments, the compound wherein

one of R_1 or R_2 is $-C(=O)O$ -(C_{1-6} alkyl)- R_{14} ,

wherein R_{14} is CF_3 , cycloalkyl, aryl or heteroaryl; and

the other of R_1 or R_2 is $-C(=O)OH$.

In some embodiments, the compound wherein

one of R_1 or R_2 is $-C(=O)O-CH_3-R_{14}$,

wherein R_{14} is CF_3 , cycloalkyl, aryl or heteroaryl; and

5 the other of R_1 or R_2 is $-C(=O)OH$.

In some embodiments, the compound wherein the aryl is a substituted aryl.

10 In some embodiments, the compound wherein the aryl is substituted with a halogen, $-OH$, heteroaryl, C_2-C_6 alkynyl or $-O(alkyl)$.

In some embodiments, the compound wherein the aryl is substituted with a amide, aryl or hydroxyaryl.

15 In some embodiments, the compound wherein the aryl is substituted with a F, Cl, Br, $-OH$, triazolyl, C_2 alkylnyl or $-OCH_3$.

20 In some embodiments, the compound wherein the aryl is substituted with a F, Cl, Br, $-OH$, I, $-NHC(O)CH_3$, triazolyl, C_2 alkylnyl, phenyl, o-hydroxyphenyl or $-OCH_3$.

In some embodiments, the compound wherein the heteroaryl is a substituted heteroaryl.

25 In some embodiments, the compound wherein the heteroaryl is substituted with a halogen, $-OH$, heteroaryl, C_2-C_6 alkynyl or $-O(alkyl)$.

In some embodiments, the compound wherein the heteroaryl is substituted with an amide, aryl or hydroxyaryl.

30 In some embodiments, the compound wherein the heteroaryl is substituted with an F, Cl, Br, $-OH$, triazolyl, C_2 alkylnyl or $-OCH_3$.

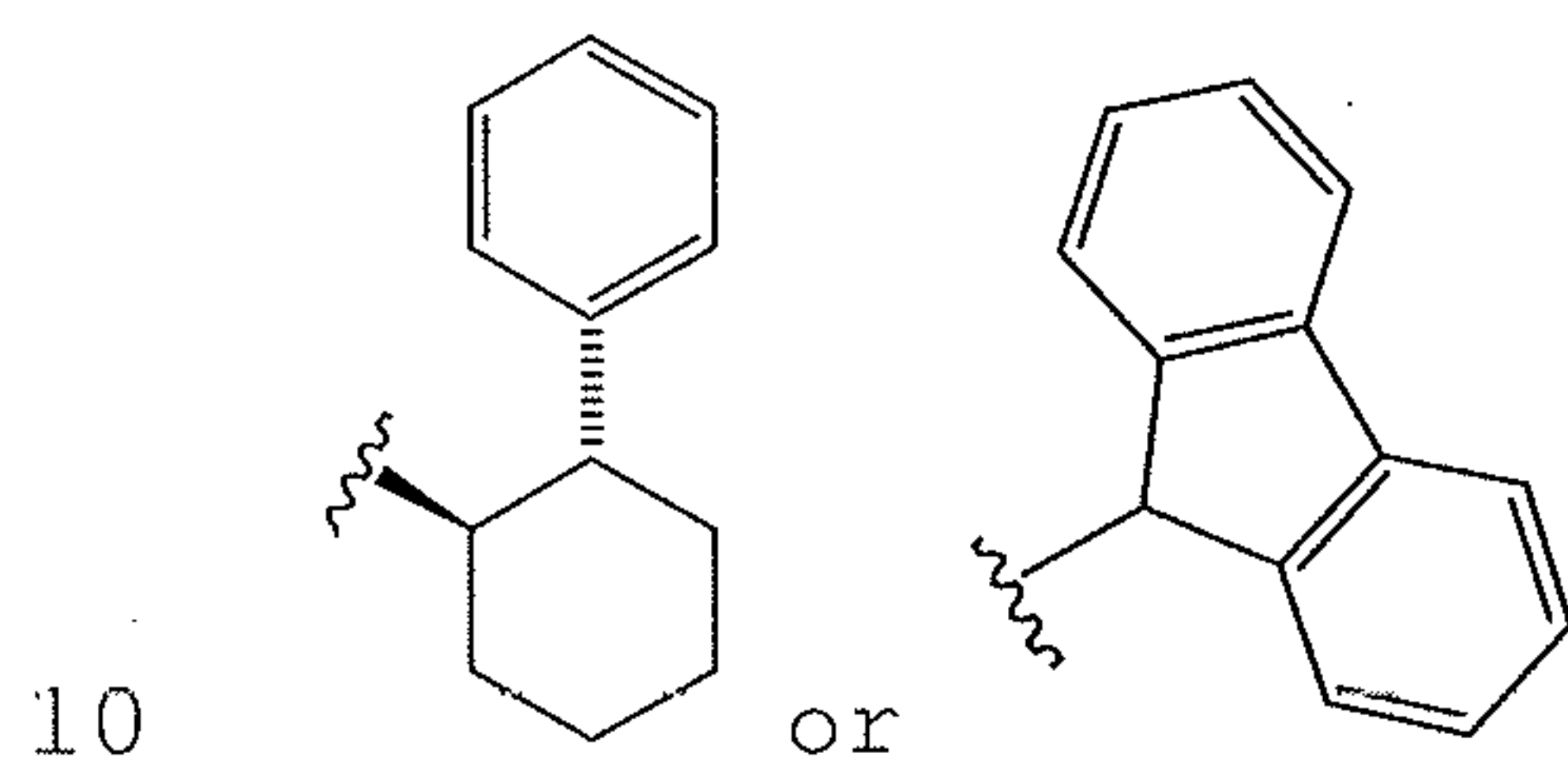
35 In some embodiments, the compound wherein the heteroaryl is substituted with a F, Cl, Br, $-OH$, I, $-NHC(O)CH_3$, triazolyl, C_2 alkylnyl, phenyl, o-hydroxyphenyl or $-OCH_3$.

In some embodiments, the compound wherein the cycloalkyl is a substituted cycloalkyl.

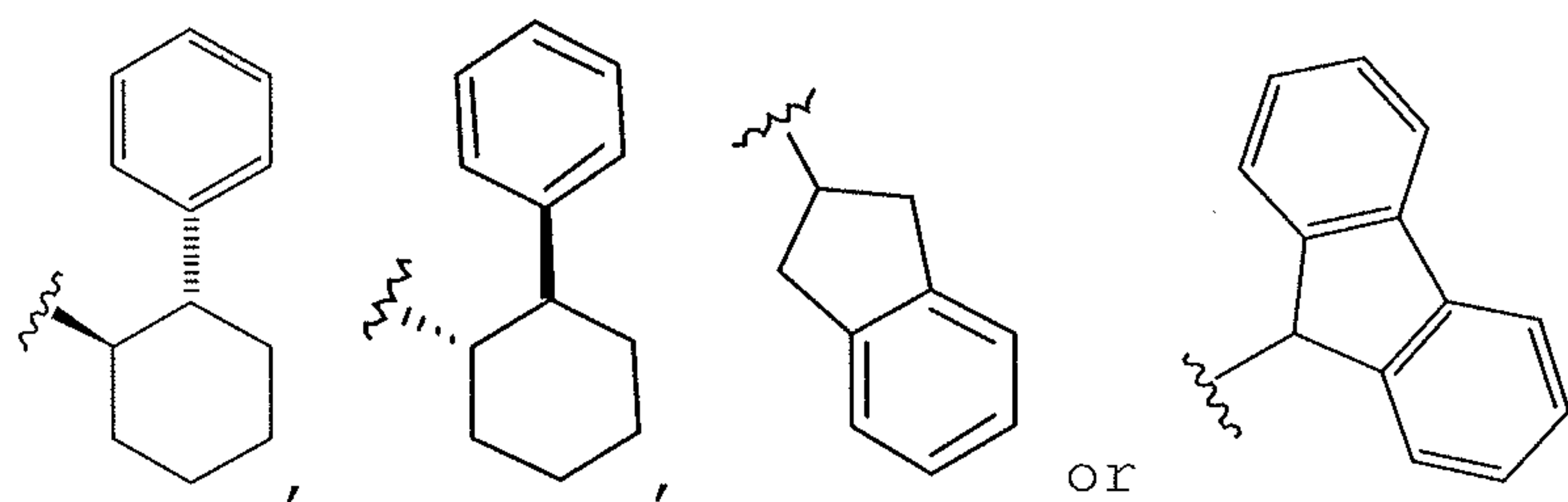
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In some embodiments, the compound wherein the cycloalkyl is substituted with an phenyl or a fused benzo group.

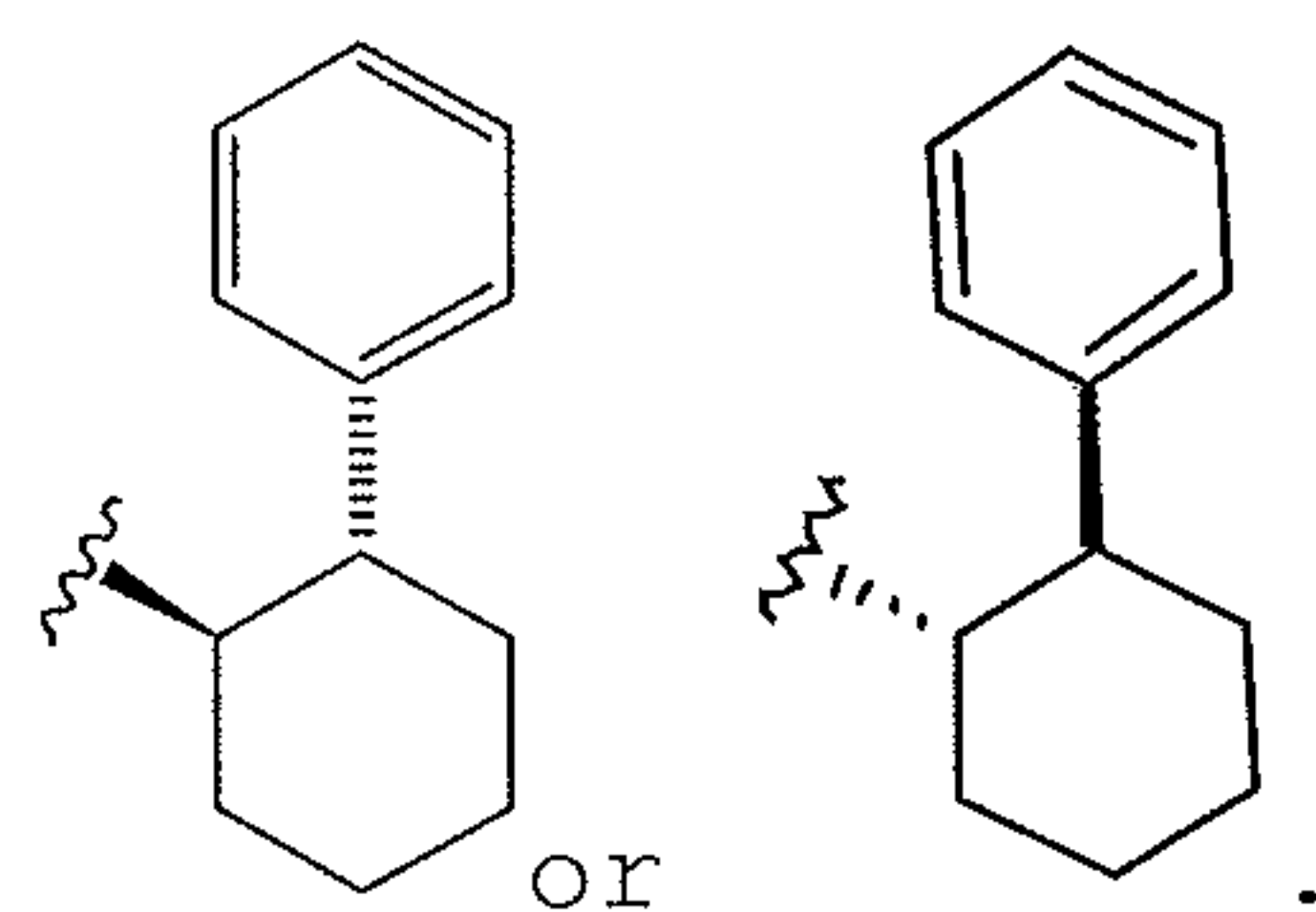
In some embodiments, the compound wherein the cycloalkyl is:



In some embodiments, the compound wherein the cycloalkyl is:

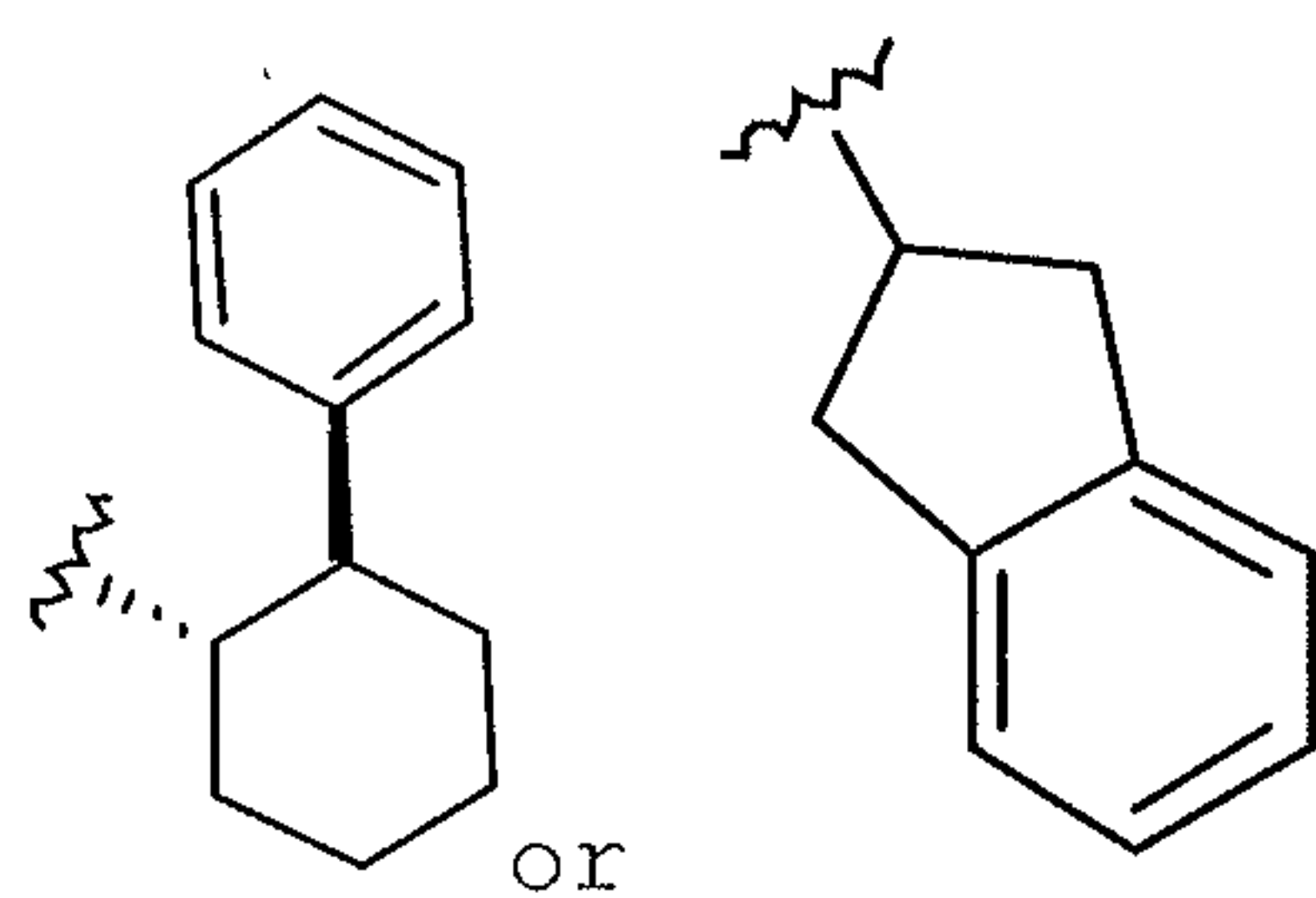


15 In some embodiments, the compound wherein the cycloalkyl is:



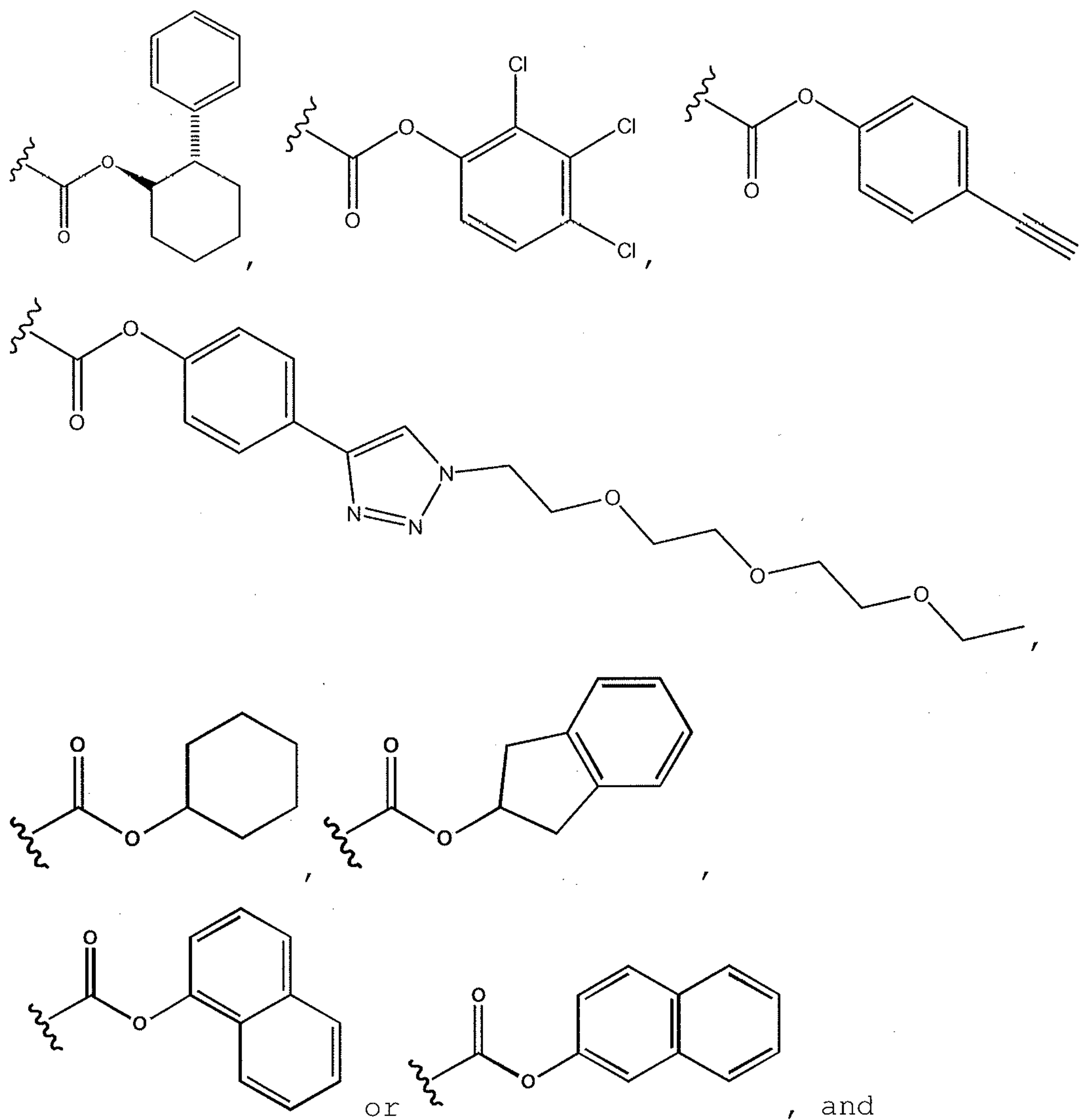
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In some embodiments, the compound wherein the cycloalkyl is:



In some embodiments, the compound wherein

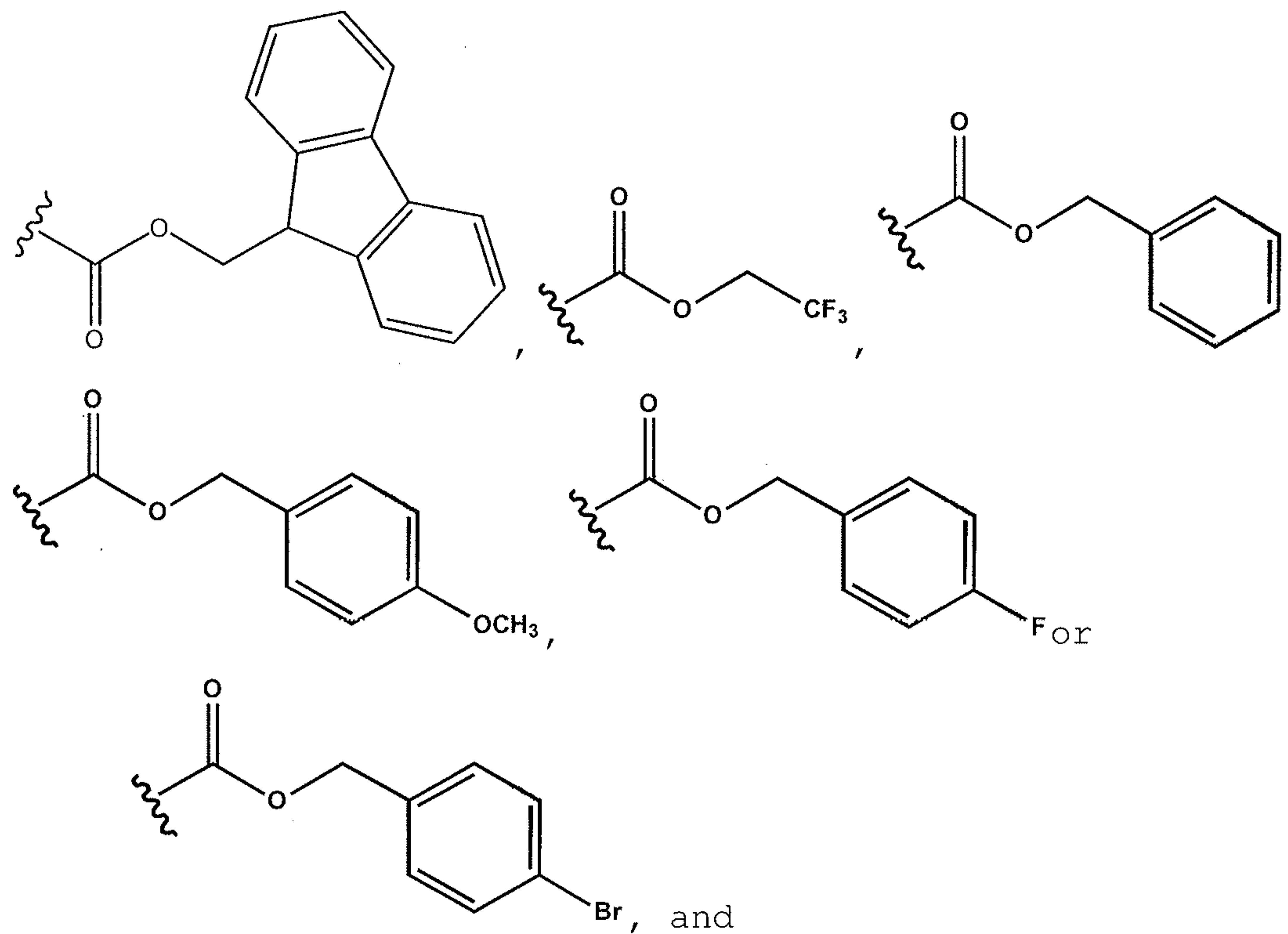
5 one of R_1 or R_2 is



the other of R_1 or R_2 is $-C(=O)OH$.

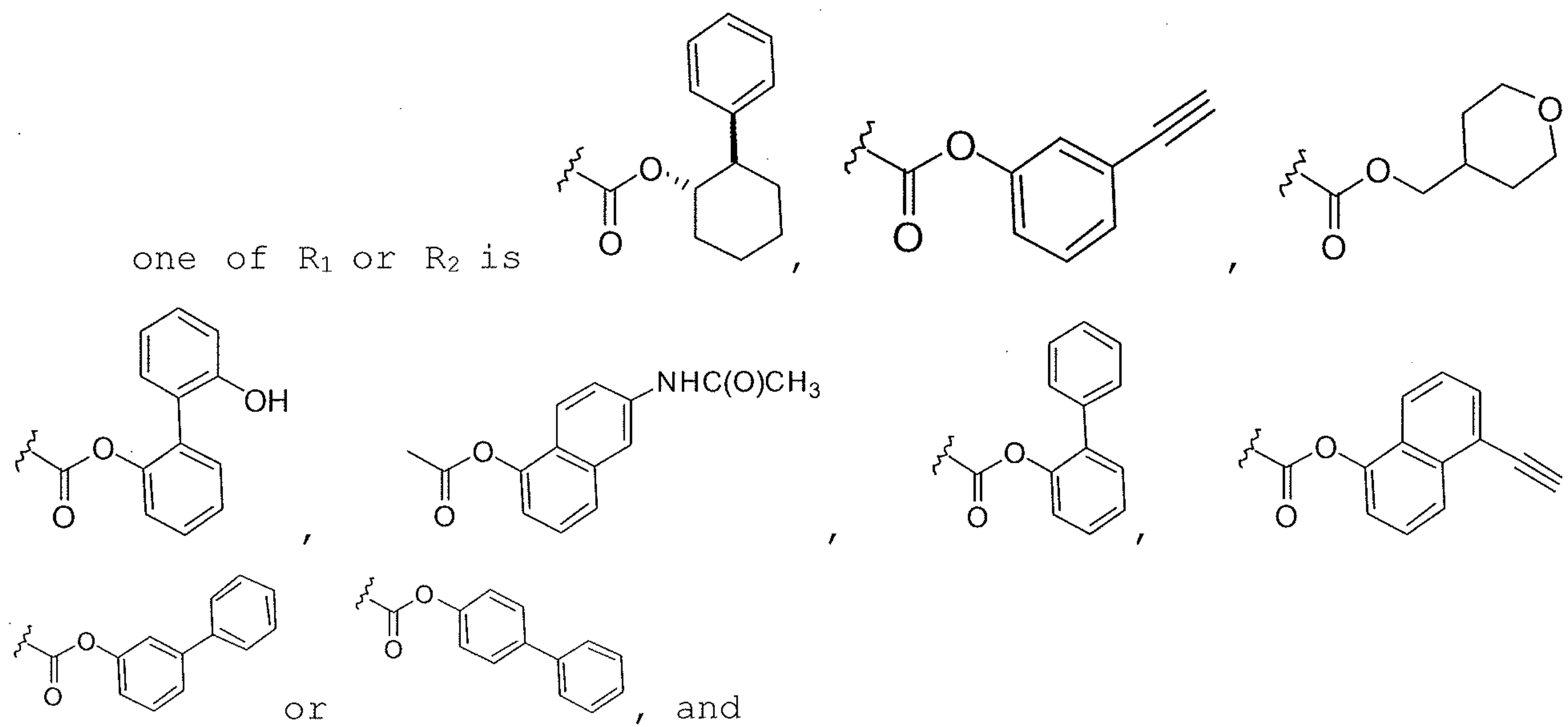
In some embodiments, the compound wherein

one of R_1 or R_2 is



the other of R_1 or R_2 is $-C(=O)OH$.

10 In some embodiments, the compound wherein



the other of R₁ or R₂ is -C(=O)OH.

In some embodiments, the compound wherein R₃, R₄, R₅, R₆, R₇, R₈, R₉, R₁₀, R₁₁ and R₁₂ are each independently, -H, halogen or -OR₁₅,

5 wherein R₁₅ is -H or C₁₋₁₀ alkyl.

In some embodiments, the compound wherein R₃, R₄, R₅, R₆, R₇, R₈, R₉, R₁₀, R₁₁ and R₁₂ are each independently, -H, F, Cl, Br or -OR₁₅,

 wherein R₁₅ is -H or C₁₋₆ alkyl.

10

In some embodiments, the compound wherein R₃, R₄, R₅, R₆, R₇, R₈, R₉, R₁₀, R₁₁ and R₁₂ are each independently, -H, Cl or -OR₁₅,

 wherein R₁₅ is -H or CH₃.

15 In some embodiments, the compound wherein R₃, R₄, R₅, R₆, R₇, R₈, R₉, R₁₀, R₁₁ and R₁₂ are each independently, -H or -NO₂.

In some embodiments, the compound wherein one of R₃, R₄, R₅, R₆, R₇, R₈, R₉, R₁₀, R₁₁ and R₁₂ is other than -H.

20

In some embodiments, the compound wherein two of R₃, R₄, R₅, R₆, R₇, R₈, R₉, R₁₀, R₁₁ and R₁₂ are other than -H.

In some embodiments, the compound wherein four of R₃, R₄, R₅, R₆, R₇, R₈, R₉, R₁₀, R₁₁ and R₁₂ are other than -H.

25

In some embodiments, the compound wherein R₃, R₄, R₆, R₇, R₈, R₉, R₁₁ and R₁₂ are each -H; and R₅ and R₁₀ are each halogen or -OR₁₅,

 wherein R₁₅ is -H, C₁₋₁₀ alkyl, C₂₋₁₀ alkenyl, C₂₋₁₀ alkynyl, aryl, or
30 heteroaryl.

In some embodiments, the compound wherein R₃, R₄, R₆, R₇, R₈, R₉, R₁₁ and R₁₂ are each -H; and R₅ and R₁₀ are each -OR₁₅,

 wherein R₁₅ is -H or C₁₋₁₀ alkyl.

35

In some embodiments, the compound wherein R_3 , R_4 , R_6 , R_7 , R_8 , R_9 , R_{11} and R_{12} are each -H; and R_5 and R_{10} are each $-OR_{15}$,

wherein R_{15} is -H or C_{1-6} alkyl.

5 In some embodiments, the compound wherein R_3 , R_4 , R_6 , R_7 , R_8 , R_9 , R_{11} and R_{12} are each -H; and R_5 and R_{10} are each $-OR_{15}$,

wherein R_{15} is -H or CH_3 .

10 In some embodiments, the compound wherein R_4 , R_5 , R_6 , R_9 , R_{10} and R_{11} are each -H; R_3 and R_{12} are each -H, halogen or $-OR_{15}$; and R_7 and R_8 are each halogen or $-OR_{15}$,

wherein R_{15} is -H, C_{1-10} alkyl, C_{2-10} alkenyl, C_{2-10} alkynyl, aryl, or heteroaryl.

15 In some embodiments, the compound wherein R_4 , R_5 , R_6 , R_9 , R_{10} and R_{11} are each -H; R_3 and R_{12} are each -H, halogen or $-OR_{15}$; and R_7 and R_8 are each halogen or $-OR_{15}$,

wherein R_{15} is -H or C_{1-10} alkyl.

20 In some embodiments, the compound wherein R_3 , R_4 , R_5 , R_6 , R_9 , R_{10} , R_{11} and R_{12} are each -H; and R_7 and R_8 are each F, Br, Cl or $-OR_{15}$,

wherein R_{15} is -H or C_{1-6} alkyl.

25 In some embodiments, the compound wherein R_3 , R_4 , R_6 , R_7 , R_8 , R_9 , R_{11} and R_{12} are each -H; and R_5 and R_{10} are each Cl or $-OR_{15}$,

wherein R_{15} is -H or CH_3 .

In some embodiments, the compound wherein R_3 , R_4 , R_7 , R_8 , R_9 and R_{12} are each -H; and R_5 , R_6 , R_{10} and R_{11} are each halogen or $-OR_{15}$,

30 wherein R_{15} is -H, C_{1-10} alkyl, C_{2-10} alkenyl, C_{2-10} alkynyl, aryl, or heteroaryl.

In some embodiments, the compound wherein R_3 , R_4 , R_7 , R_8 , R_9 and R_{12} are each -H; and R_5 , R_6 , R_{10} and R_{11} are each $-OR_{15}$,

35 wherein R_{15} is -H or C_{1-10} alkyl.

In some embodiments, the compound wherein R_3 , R_4 , R_7 , R_8 , R_9 and R_{12} are each $-H$; and R_5 , R_6 , R_{10} and R_{11} are each $-OR_{15}$,
wherein R_{15} is $-H$ or C_{1-6} alkyl.

5

In some embodiments, the compound wherein R_3 , R_4 , R_7 , R_8 , R_9 and R_{12} are each $-H$; and R_5 , R_6 , R_{10} and R_{11} are each $-OR_{15}$,
wherein R_{15} is $-H$ or CH_3 .

10 In some embodiments, the compound wherein R_3 , R_4 , R_5 , R_6 , R_8 , R_9 , R_{10} and R_{1a} are each $-H$; and R_7 and R_{12} are each NO_2 .

In some embodiments, the compound wherein
one of R_1 or R_2 is $-C(=O)OH$ and the other of R_1 or R_2 is $-C(=O)OR_{13}$ or -
15 $C(=O)O$ -alkyl- R_{14} ,

wherein

R_{13} is cycloalkyl or aryl, and

R_{14} is CF_3 or aryl; and

R_3 , R_4 , R_5 , R_6 , R_7 , R_8 , R_9 , R_{10} , R_{11} and R_{12} are each independently, H or -
20 OR_{15} ,

wherein R_{15} is H or C_{1-10} alkyl,

or an enantiomer or racemate thereof;

25 or a pharmaceutically acceptable salt thereof.

In some embodiments, the compound wherein
one of R_1 or R_2 is $-C(=O)OH$ and the other of R_1 or R_2 is $-C(=O)OR_{13}$ or -
 $C(=O)O$ -alkyl- R_{14} ,

30 wherein

R_{13} is cycloalkyl or aryl, and

R_{14} is CF_3 , cycloheteroalkyl or aryl; and

R_3 , R_4 , R_5 , R_6 , R_7 , R_8 , R_9 , R_{10} , R_{11} and R_{12} are each independently, H or -
 OR_{15} ,

35 wherein R_{15} is H or C_{1-10} alkyl,

or an enantiomer or racemate thereof;

or a pharmaceutically acceptable salt thereof.

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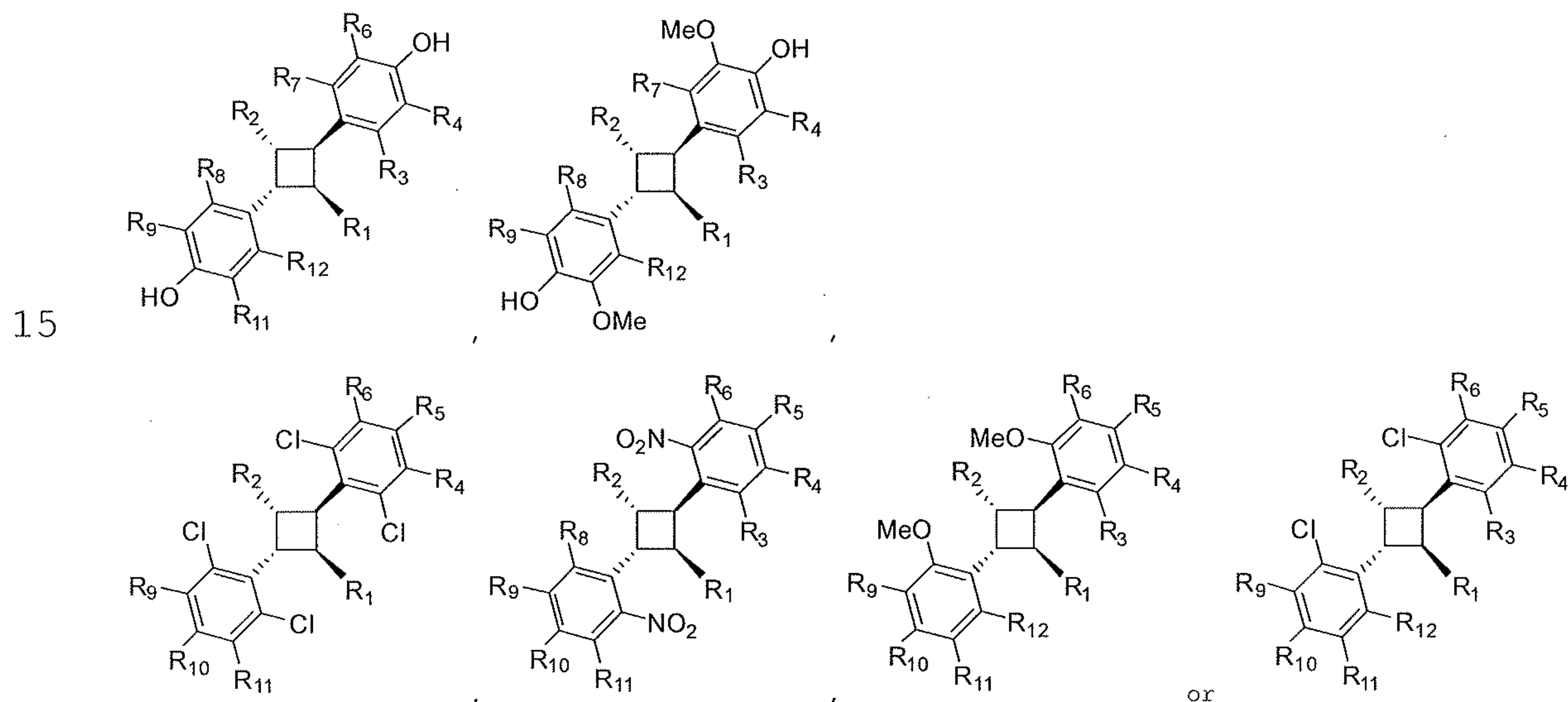
In some embodiments, the compound wherein

R₃ is identical to R₈, R₄ is identical to R₉, R₅ is identical to R₁₀,
R₆ is identical to R₁₁ and R₇ is identical to R₁₂.

10 In some embodiments, the compound wherein

R₃, R₇ and R₈ are identical to R₁₂, R₄, R₆, and R₉ are identical to R₁₁, and
R₅ is identical to R₁₀.

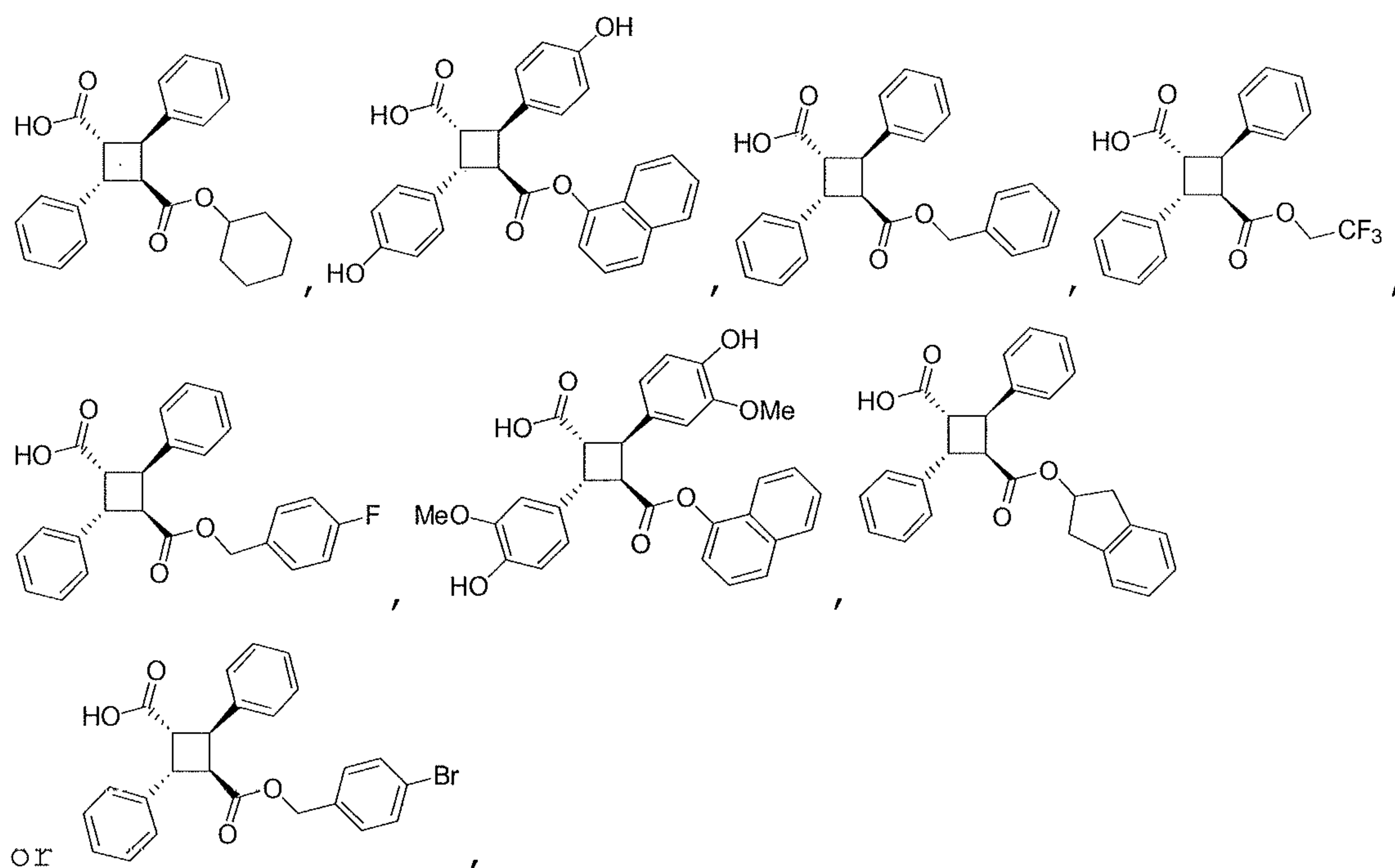
In some embodiments, the compound having the structure



20

25

In some embodiments, the compound having the structure:

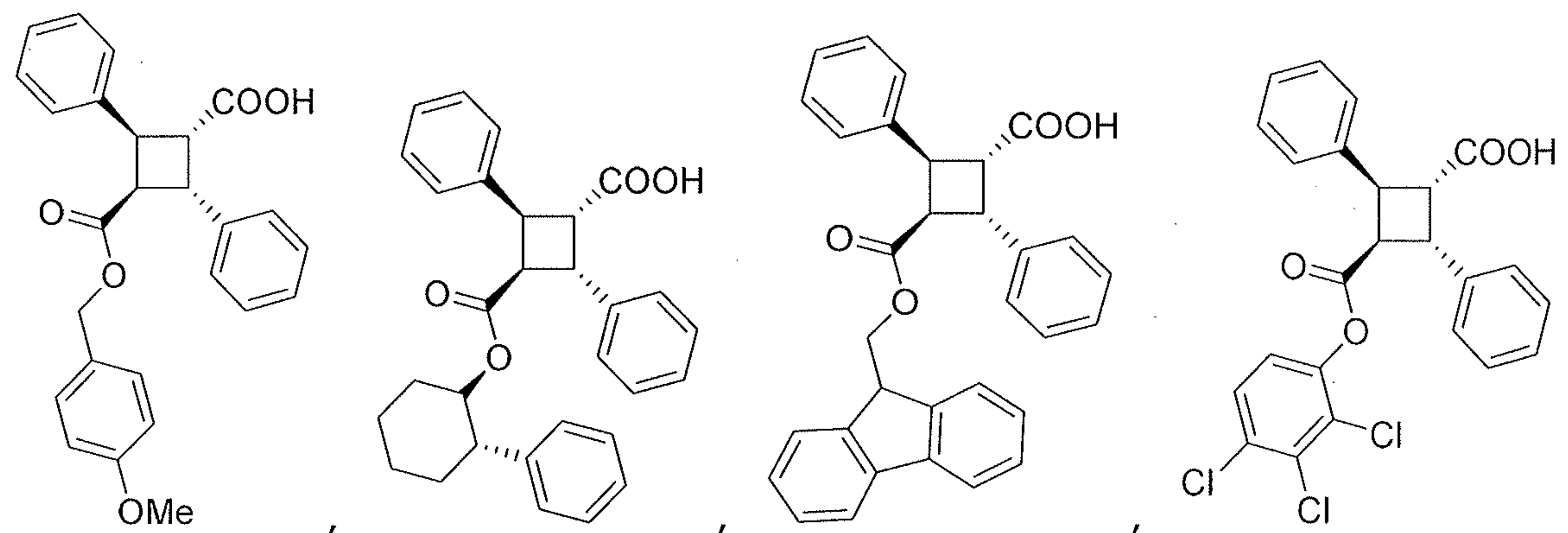


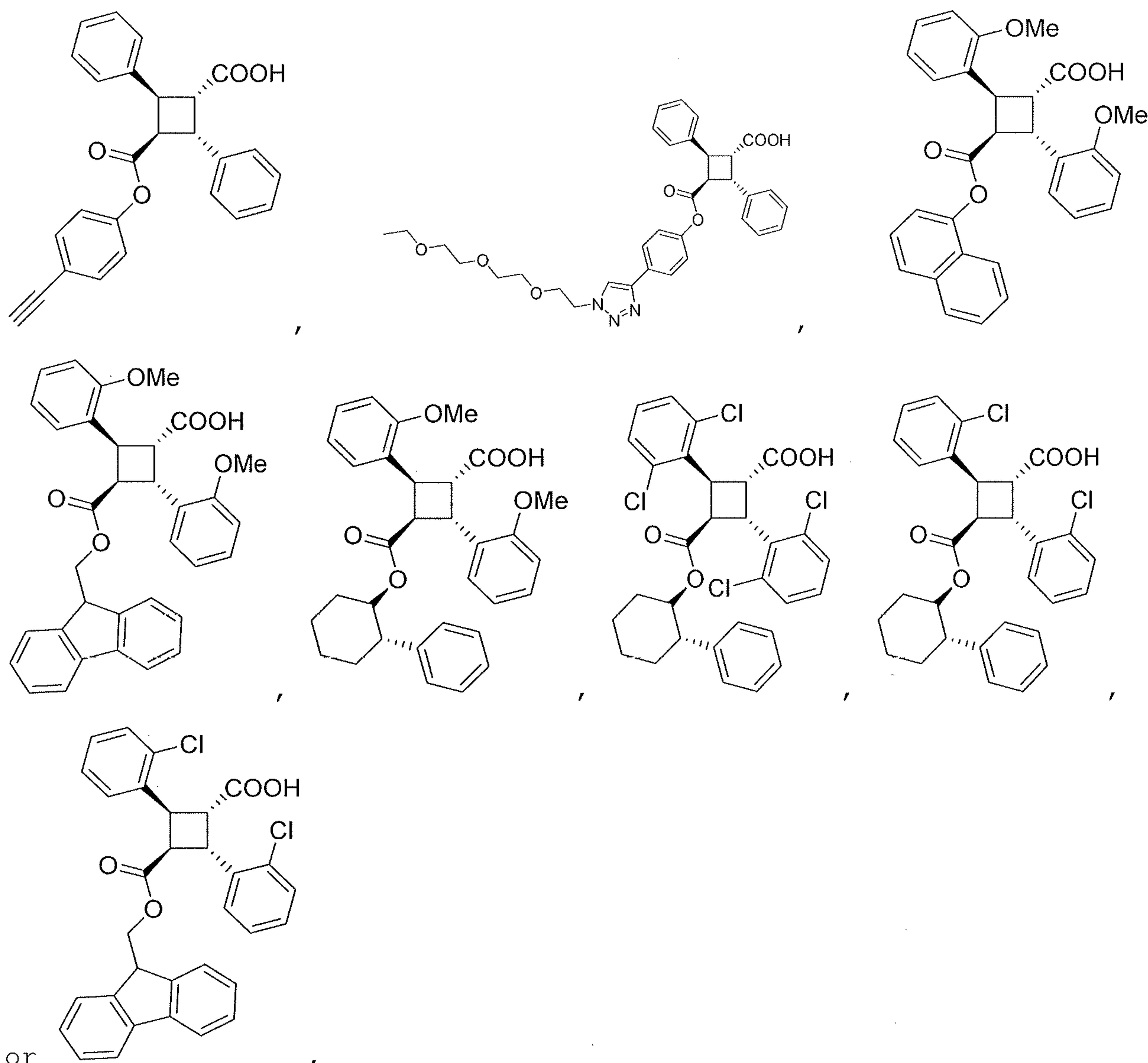
5

or an enantiomer or racemate thereof;

or a pharmaceutically acceptable salt thereof.

10 In some embodiments, the compound having the structure:

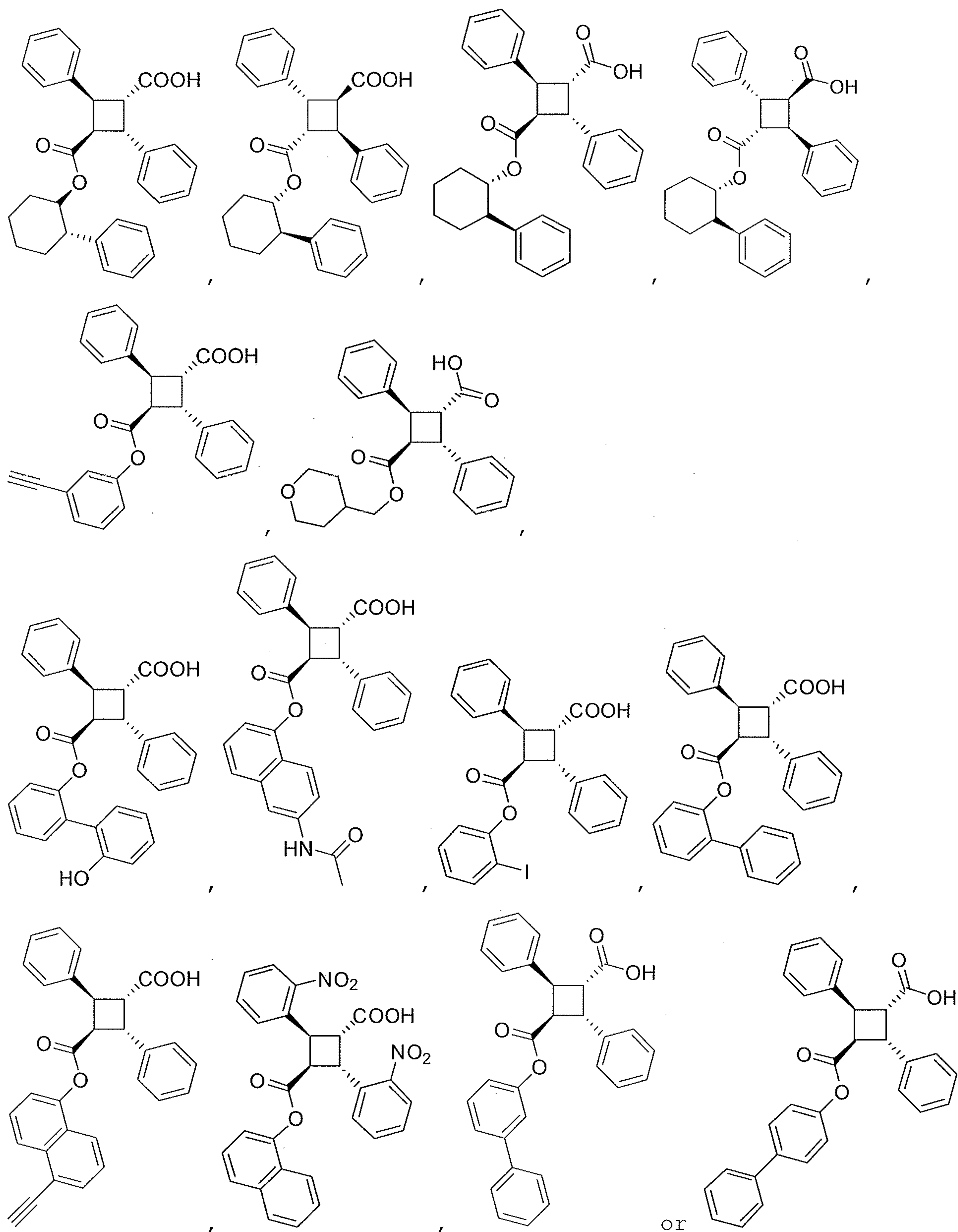




5 or enantiomer or racemate thereof;

or a pharmaceutically acceptable salt thereof.

In some embodiments, the compound having the structure:



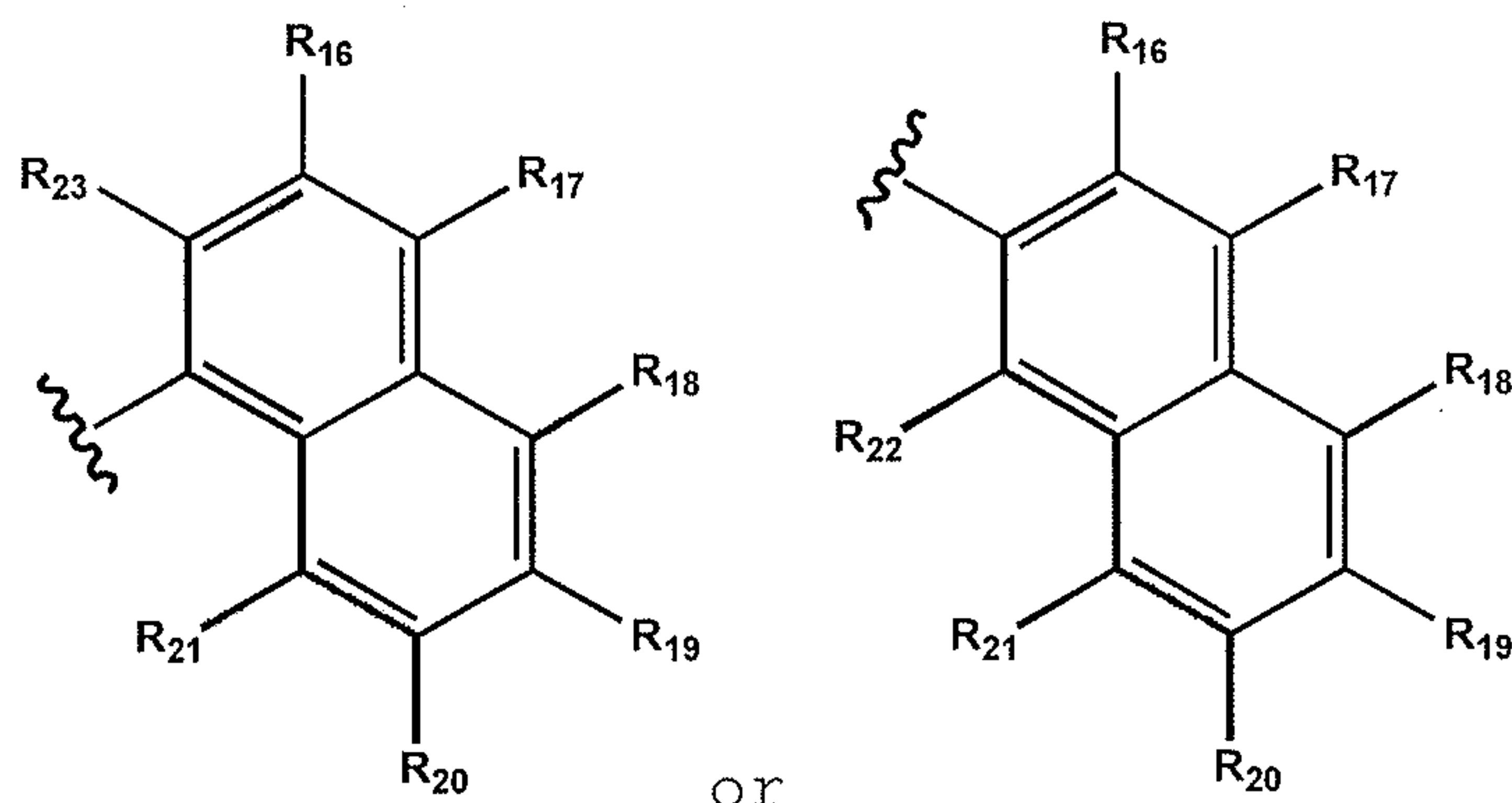
or enantiomer or racemate thereof;

or a pharmaceutically acceptable salt thereof.

In some embodiments, the compound wherein

5

R₁₃ is

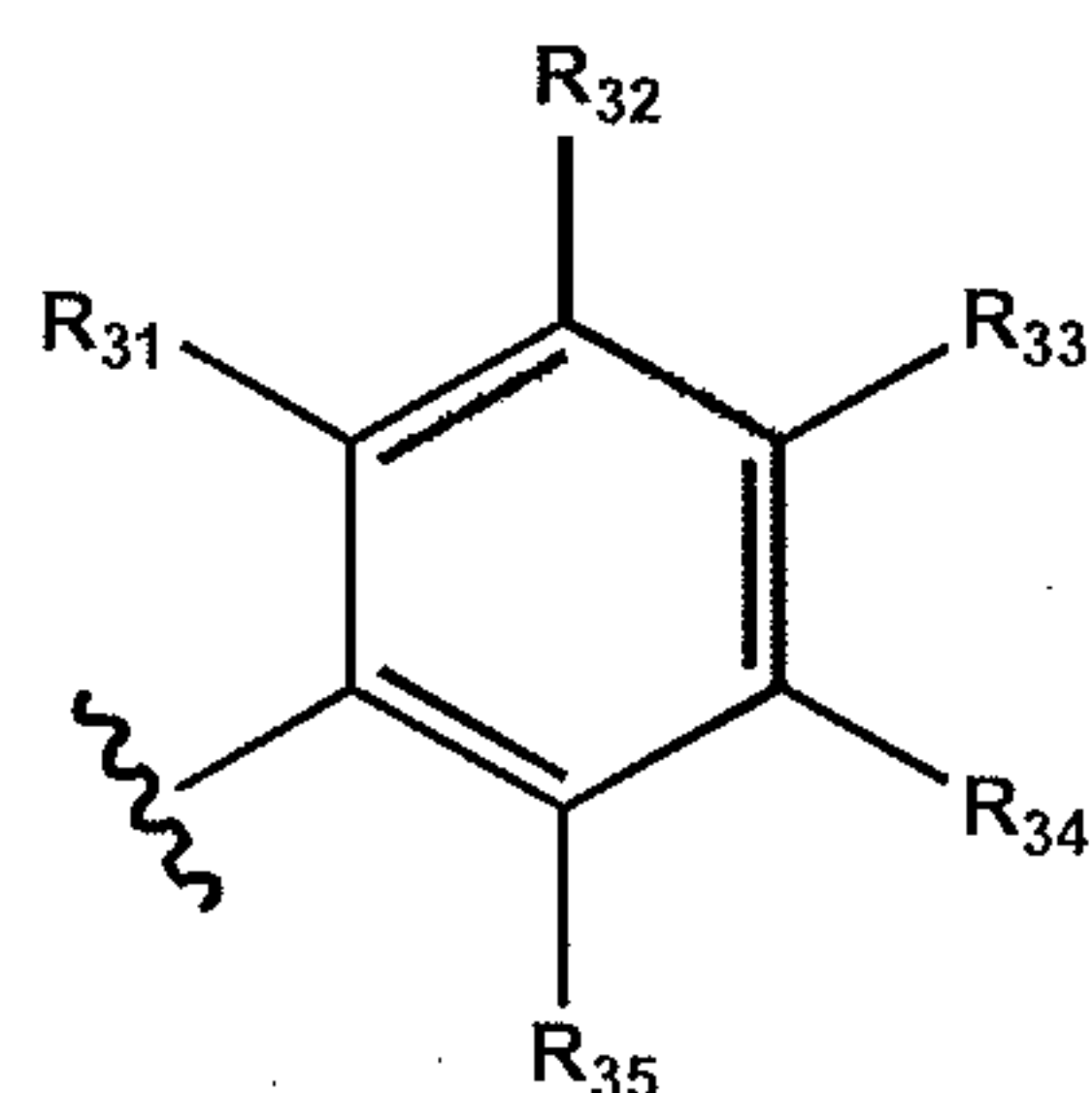


wherein each of R₁₆, R₁₇, R₁₈, R₁₉, R₂₀, R₂₁, R₂₂ and R₂₃ are each, independently, H, halogen, -NO₂, -CN, -NHR₂₄, -NR₂₄R₂₅, -SR₂₇, -SO₂R₂₈, -OR₂₉, -CO₂R₃₀, CF₃, -alkyl-NR₂₄R₂₅, -alkyl-OR₂₉, C₁₋₁₀ alkyl, C₂₋₁₀ alkenyl, C₂₋₁₀ alkynyl, aryl, heteroaryl, or heterocyclyl;

wherein R₂₄, R₂₅, R₂₆, R₂₇, R₂₈, R₂₉ and R₃₀ are each, independently, H, CF₃, C₁₋₁₀ alkyl, C₂₋₁₀ alkenyl, C₂₋₁₀ alkynyl, heteroalkyl, cycloheteroalkyl, aryl, heteroaryl, or heterocyclyl.

15 In some embodiments, the compound wherein

R₁₃ is



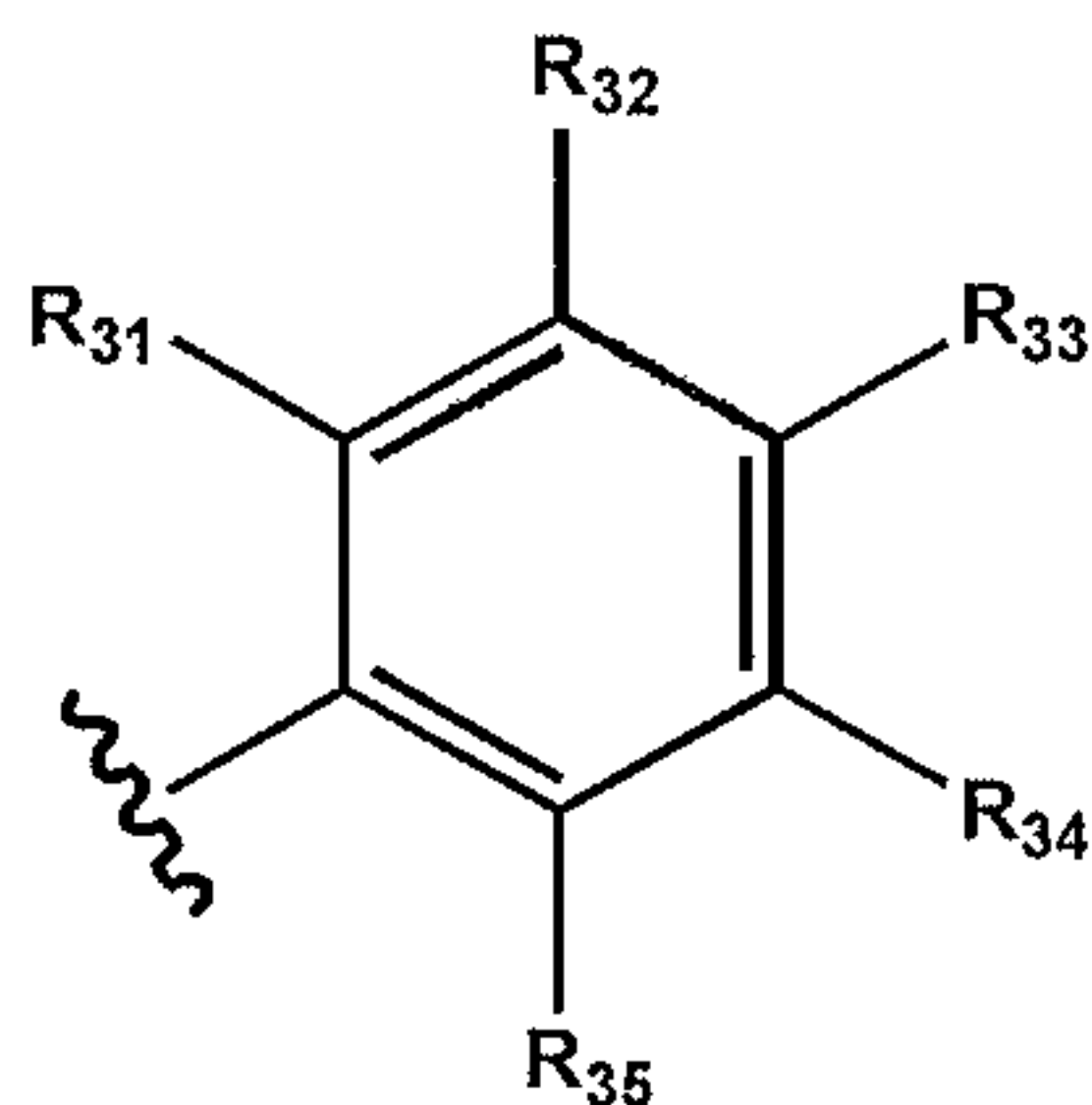
wherein each of R₃₁, R₃₂, R₃₃, R₃₄ and R₃₅ are each, independently, H, halogen, -NO₂, -CN, -NHR₂₄, -NR₂₄R₂₅, -SR₂₇, -SO₂R₂₈, -OR₂₉, -CO₂R₃₀, CF₃, -alkyl-NR₂₄R₂₅, -alkyl-OR₂₉, C₁₋₁₀ alkyl, C₂₋₁₀ alkenyl, C₂₋₁₀ alkynyl, aryl, heteroaryl, or heterocyclyl;

20

wherein R_{24} , R_{25} , R_{26} , R_{21} , R_{27} , R_{28} , R_{29} and R_{30} are each, independently, H, CF_3 , C_{1-10} alkyl, C_{2-10} alkenyl, C_{2-10} alkynyl, heteroalkyl, cycloheteroalkyl, aryl, heteroaryl, or heterocyclyl.

5 In some embodiments, the compound wherein

R_{14} is



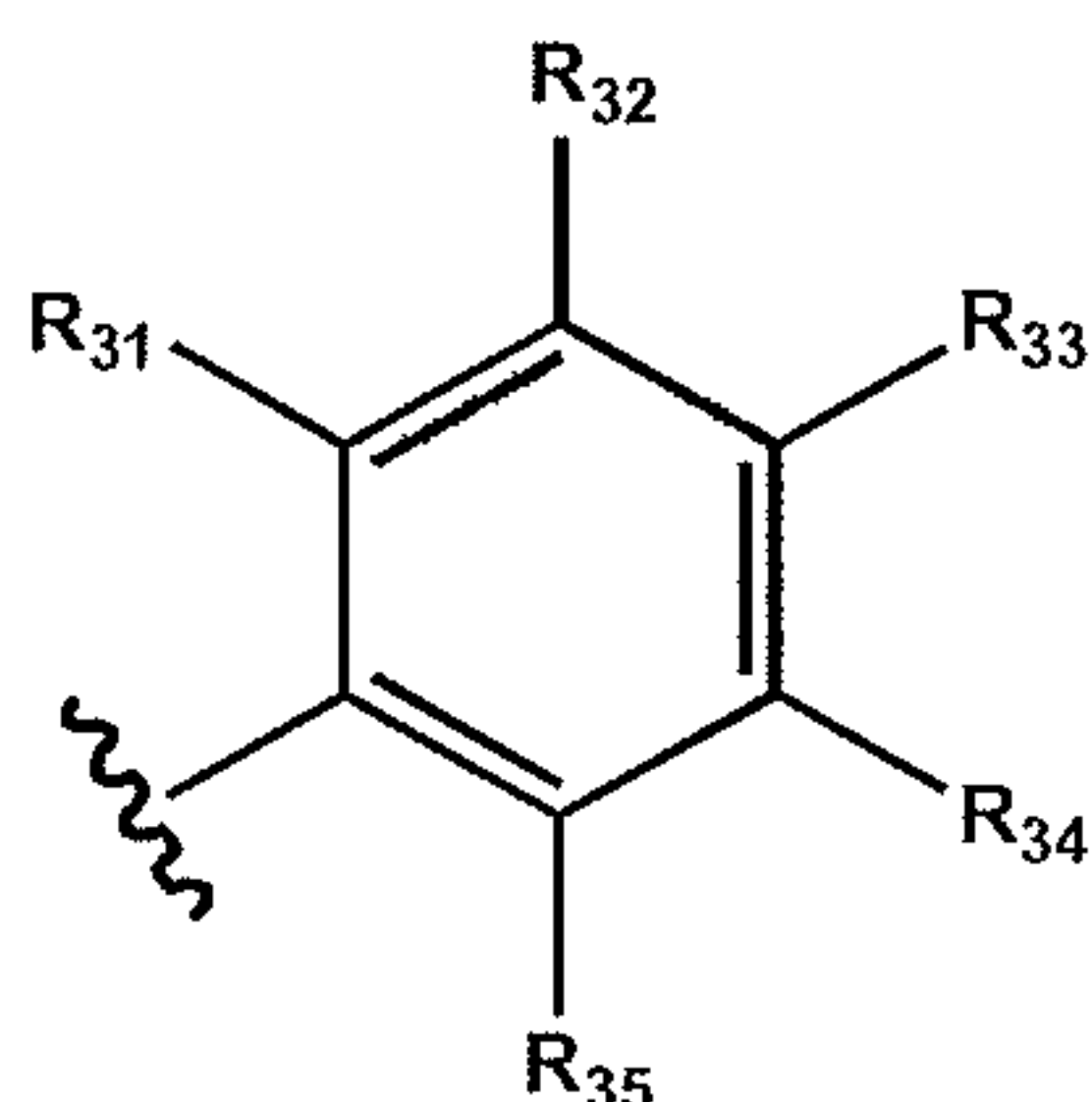
10 wherein each of R_{31} , R_{32} , R_{33} , R_{34} and R_{35} are each, independently, H, halogen, $-NO_2$, $-CN$, $-NHR_{24}$, $-NR_{24}R_{25}$, $-SR_{27}$, $-SO_2R_{28}$, $-OR_{29}$, $-CO_2R_{30}$, CF_3 , $-alkyl-NR_{24}R_{25}$, $-alkyl-OR_{29}$, C_{1-10} alkyl, C_{2-10} alkenyl, C_{2-10} alkynyl, aryl, heteroaryl, or heterocyclyl;

wherein R_{24} , R_{25} , R_{26} , R_{21} , R_{27} , R_{28} , R_{29} and R_{30} are each, independently, H, CF_3 , C_{1-10} alkyl, C_{2-10} alkenyl, C_{2-10} alkynyl, heteroalkyl, cycloheteroalkyl, aryl, heteroaryl, or heterocyclyl.

15

In some embodiments, the compound wherein

R_{14} is



20 wherein each of R_{31} , R_{32} , R_{34} and R_{35} are each, independently, H, halogen, $-NO_2$, $-CN$, $-NHR_{24}$, $-NR_{24}R_{25}$, $-SR_{27}$, $-SO_2R_{28}$, $-OR_{29}$, $-CO_2R_{30}$, CF_3 , $-alkyl-NR_{24}R_{25}$, $-alkyl-OR_{29}$, C_{1-10} alkyl, C_{2-10} alkenyl, C_{2-10} alkynyl, aryl, heteroaryl, or heterocyclyl;

25 wherein R_{24} , R_{25} , R_{26} , R_{21} , R_{27} , R_{28} , R_{29} and R_{30} are each, independently, H, CF_3 , C_{1-10} alkyl, C_{2-10} alkenyl, C_{2-10} alkynyl, heteroalkyl, cycloheteroalkyl, aryl, heteroaryl, or heterocyclyl; and

R₃₃ is halo or -OR₂₉,

wherein R₂₉ is H or C₁₋₁₀ alkyl,

In some embodiments, the triazolo is substituted with a polyether.

5

In some embodiments, R₁₃ is other than naphthyl. In some embodiments, R₁₄ is other than naphthyl. In some embodiments, R₁₃ and R₁₄ are each other than naphthyl.

10 In some embodiments, R₁₃ is cycloalkyl, monoaryl or heteroaryl.

In some embodiments, R₁₄ is CF₃, cycloalkyl, cycloheteroalkyl, monoaryl or heteroaryl.

15 In some embodiments, wherein one of R₁ or R₂ is -C(=O)O-alkyl-R₁₄,
wherein R₁₄ is CF₃, cycloalkyl, cycloheteroalkyl, aryl or
heteroaryl, and the alkyl in -C(=O)O-alkyl-R₁₄ is unbranched.

20 The present invention also provides a pharmaceutical composition
comprising the compound of the present invention and a pharmaceutically
acceptable carrier.

The present invention also provides a salt of any of the above compounds.

25 The present invention also provides an ester of any of the above
compounds.

The present invention also provides an enantiomer of any of the above
compounds.

30

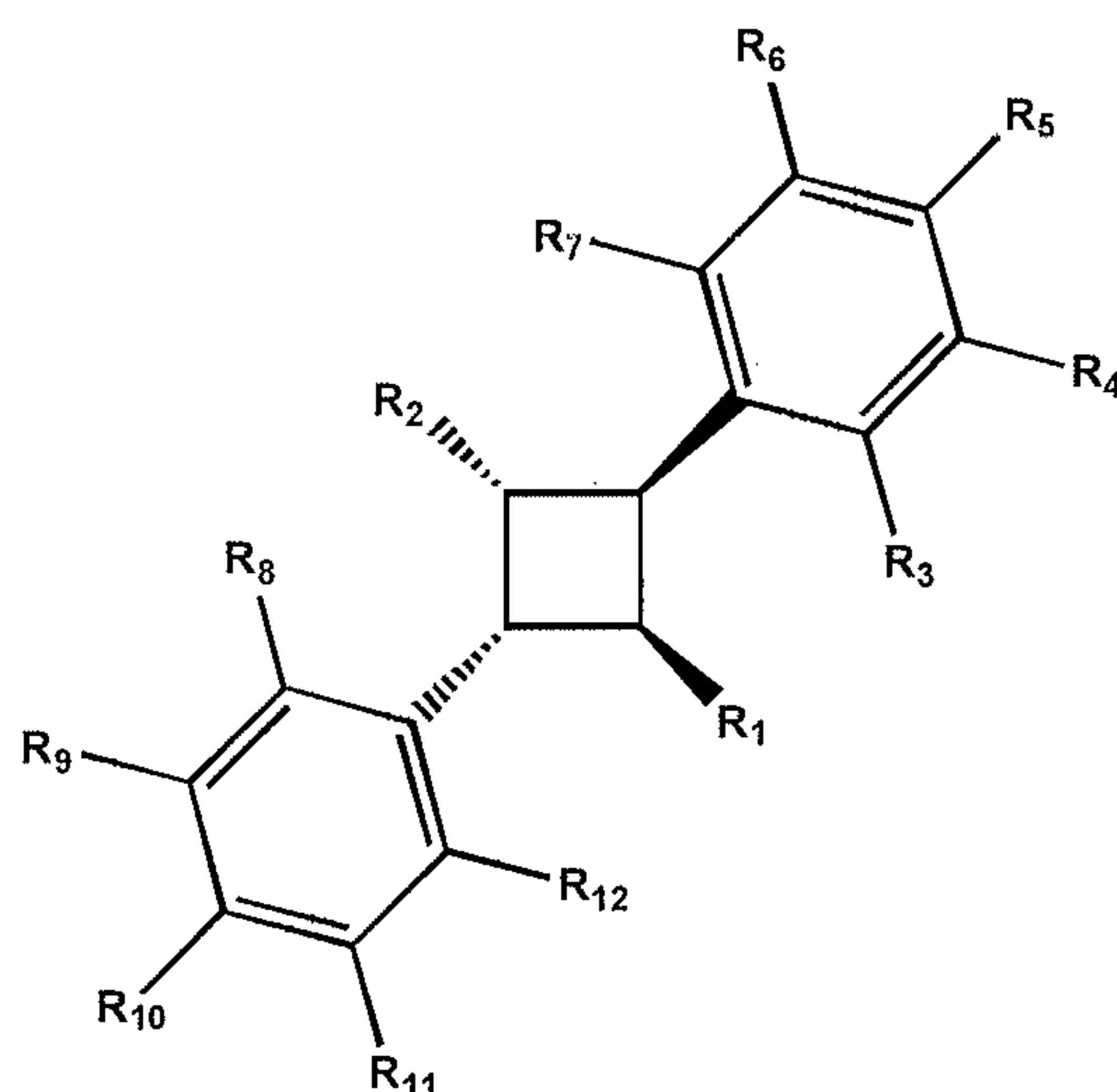
In some embodiments, a method of inhibiting the activity of a Fatty Acid
Binding Protein (FABP) comprising contacting the FABP with the compound
of the present invention.

In some embodiments, the method wherein the compound inhibits binding of an FABP ligand to the FABP.

In some embodiments, the method wherein the FABP ligand is an endocannabinoid ligand.

In some embodiments, the method wherein the FABP ligand is anandamide (AEA) or 2-arachidonoylglycerol (2-AG).

10 In some embodiments, the method of inhibiting the activity of a Fatty Acid Binding Protein (FABP) comprising contacting the FABP with a compound having the structure:



wherein

15 one of R_1 or R_2 is $-C(=O)OH$ and the other of R_1 or R_2 is $-C(=O)OR_{13}$ or $-C(=O)O\text{-alkyl-}R_{14}$,

wherein

R_{13} is H, cycloalkyl, aryl or heteroaryl, and

R_{14} is CF_3 , cycloalkyl, aryl or heteroaryl; and

20 R_3 , R_4 , R_5 , R_6 , R_7 , R_8 , R_9 , R_{10} , R_{11} and R_{12} are each independently, H, halogen or $-OR_{15}$,

wherein R_{15} is H, C_{1-10} alkyl, C_{2-10} alkenyl, C_{2-10} alkynyl, aryl, or heteroaryl,

25 or its enantiomer or racemate;

or a pharmaceutically acceptable salt thereof.

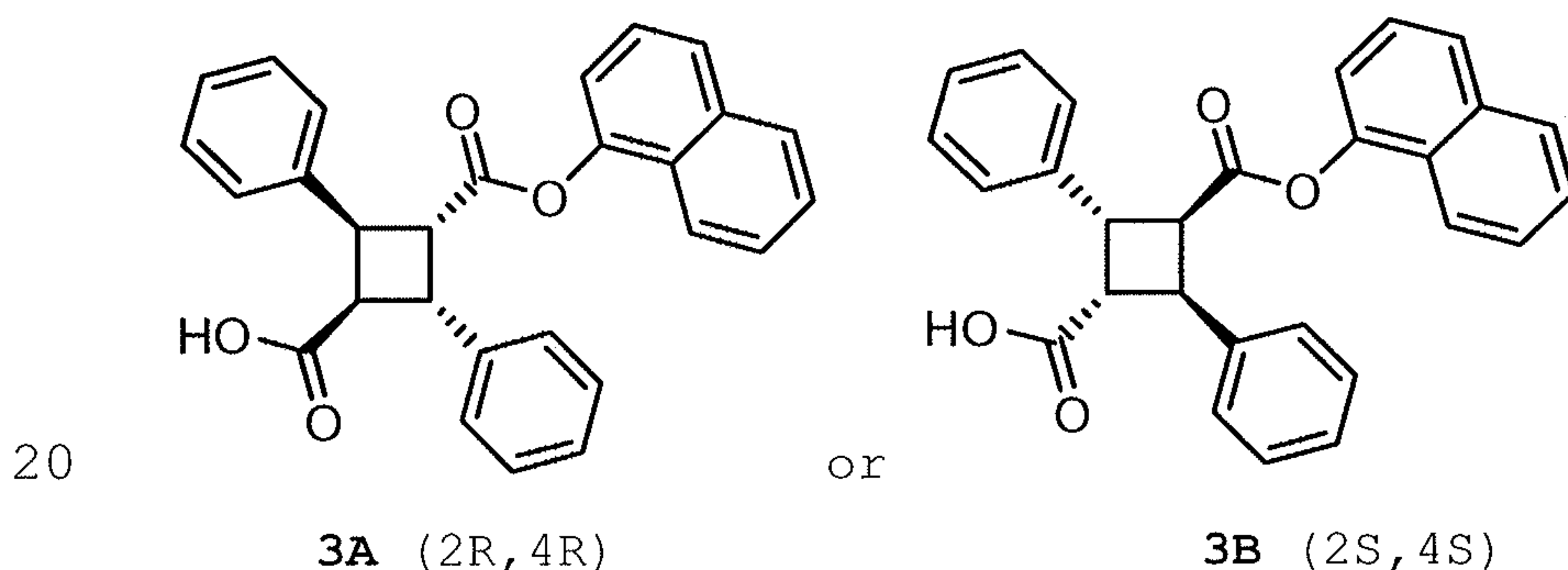
In some embodiments, the method wherein the compound inhibits binding
5 of an FABP ligand to the FABP.

In some embodiments, the method wherein the FABP ligand is an
endocannabinoid ligand.

10 In some embodiments, the method wherein the FABP ligand is anandamide
(AEA) or 2-arachidonoylglycerol (2-AG).

In some embodiments, the method wherein when one of R_1 or R_2 is $-C(=O)OH$
and $R_3, R_4, R_5, R_6, R_7, R_8, R_9, R_{10}, R_{11}$ and R_{12} are each H, then the other
15 of R_1 or R_2 is other than $-C(=O)OR_{13}$ where R_{13} is 1-naphthalene or 2-
naphthalene or $-C(=O)O$ -alkyl- R_{14} where the alkyl is a branched C_2 alkyl
and the R_{14} is phenyl.

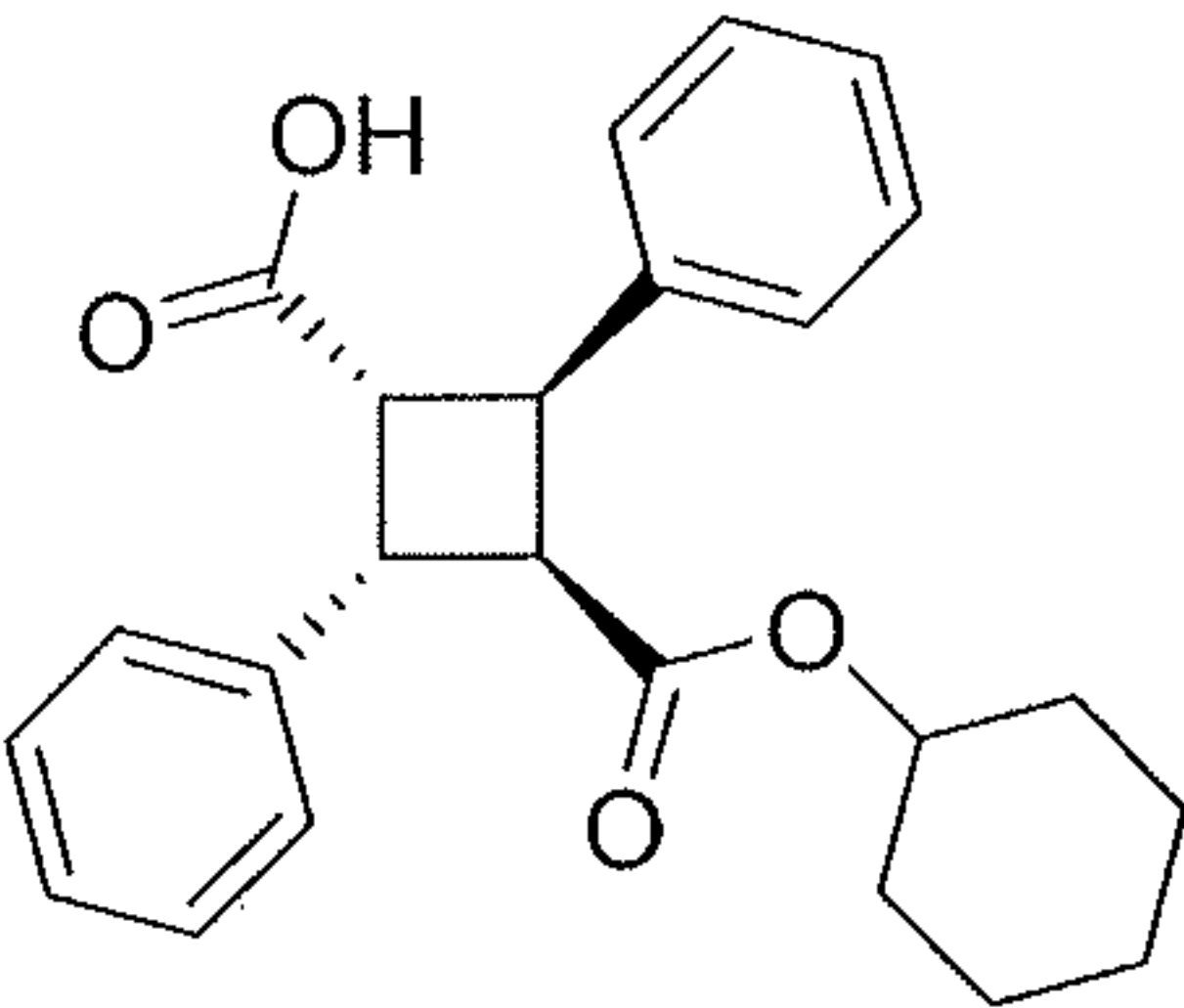
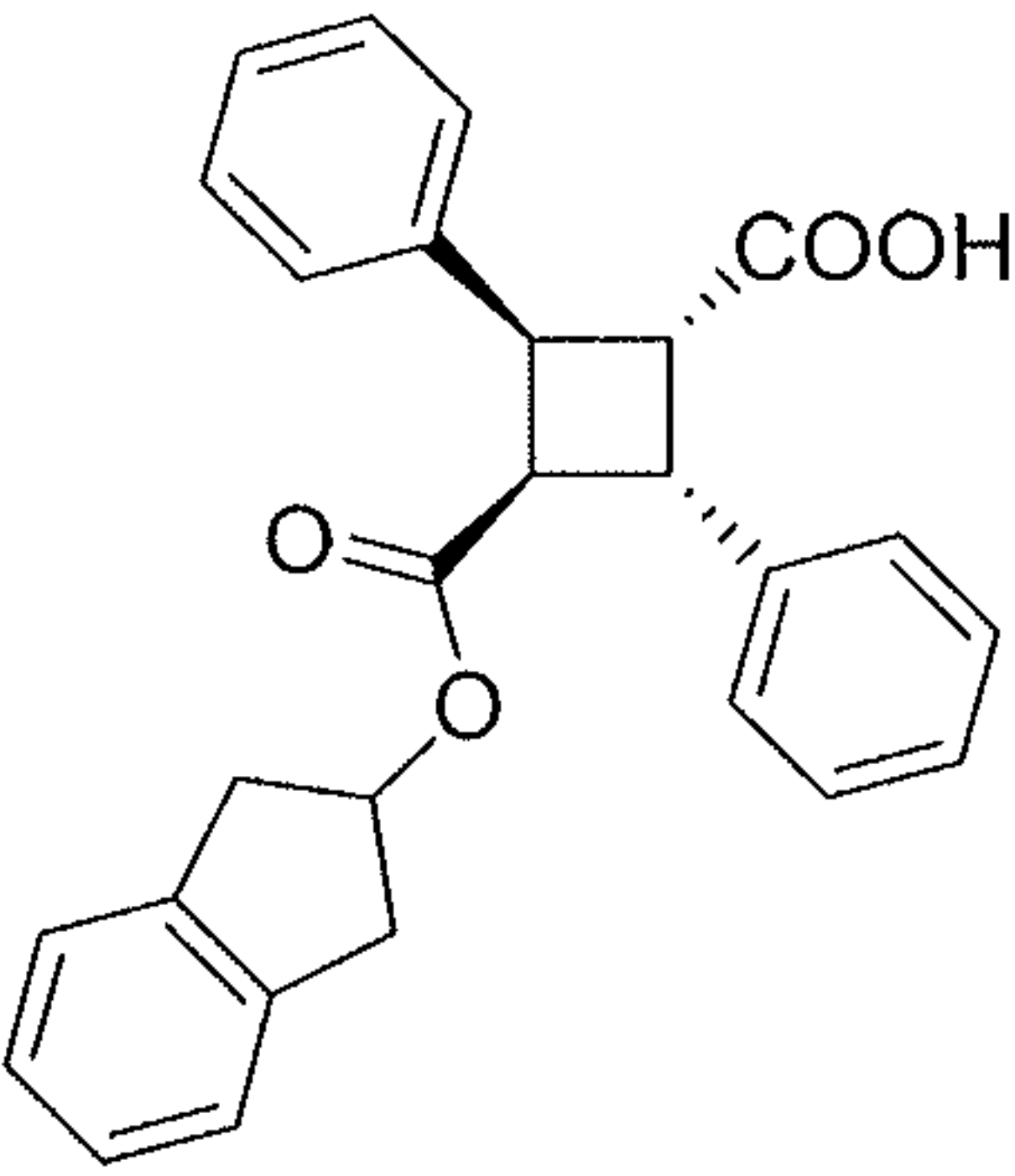
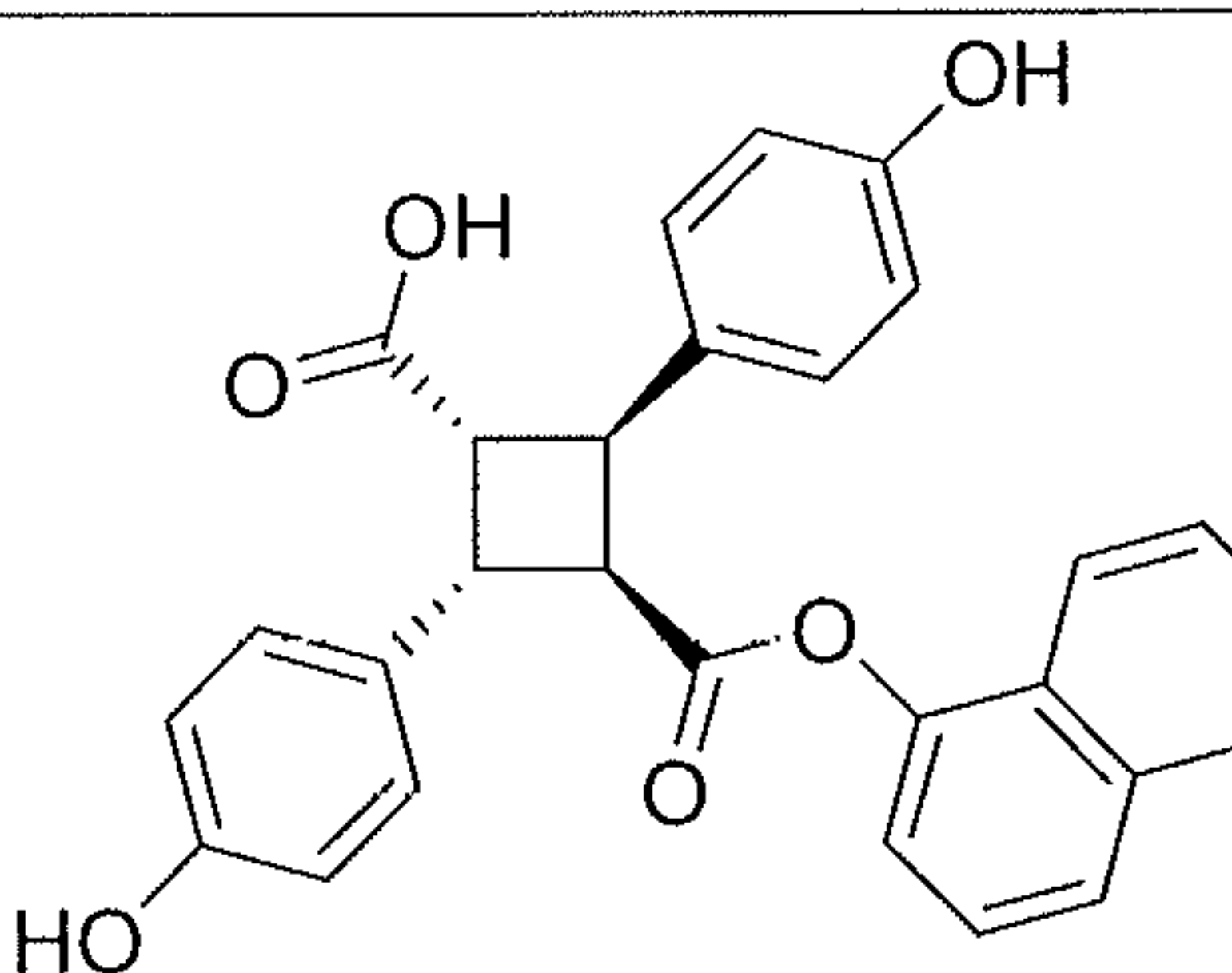
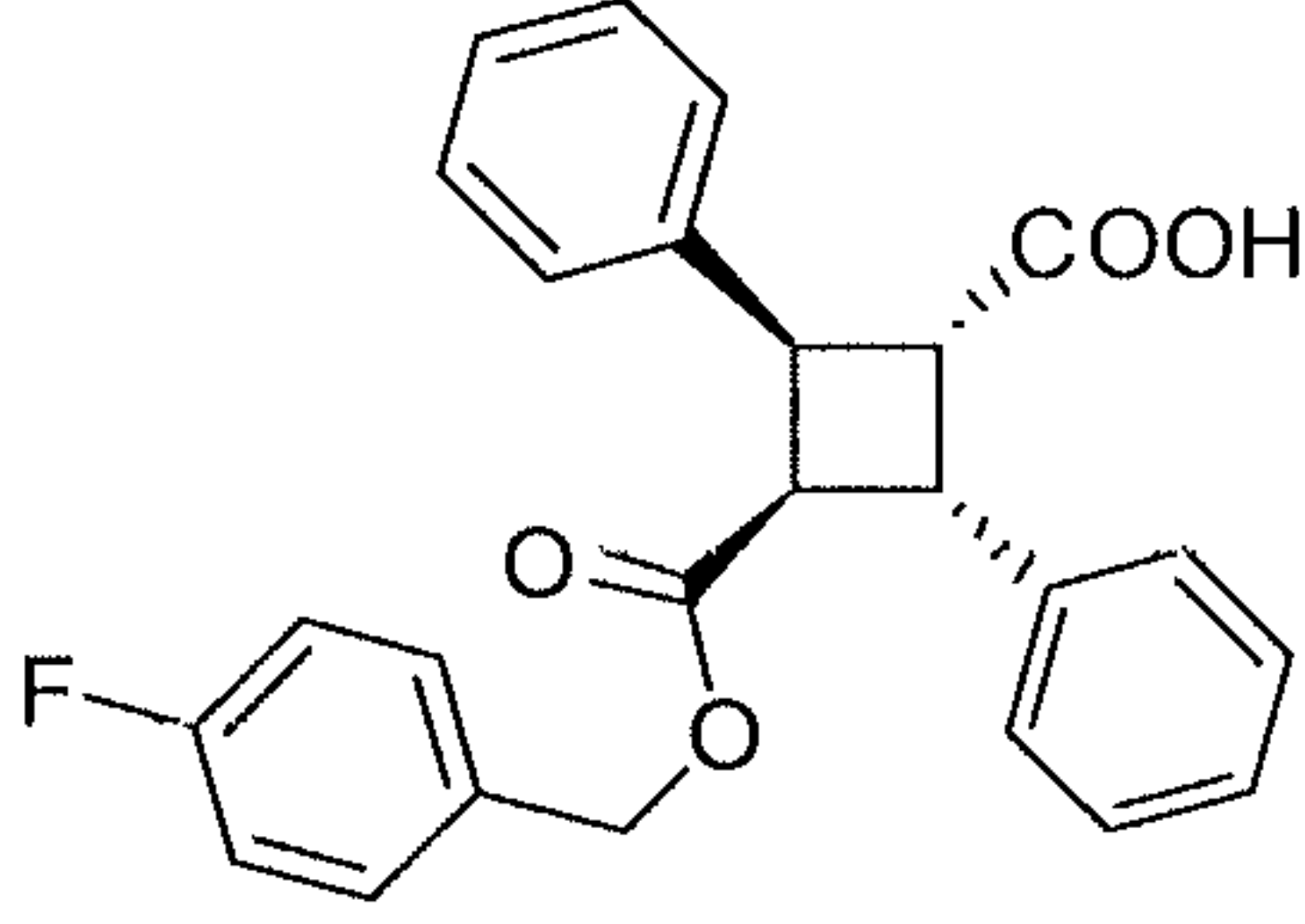
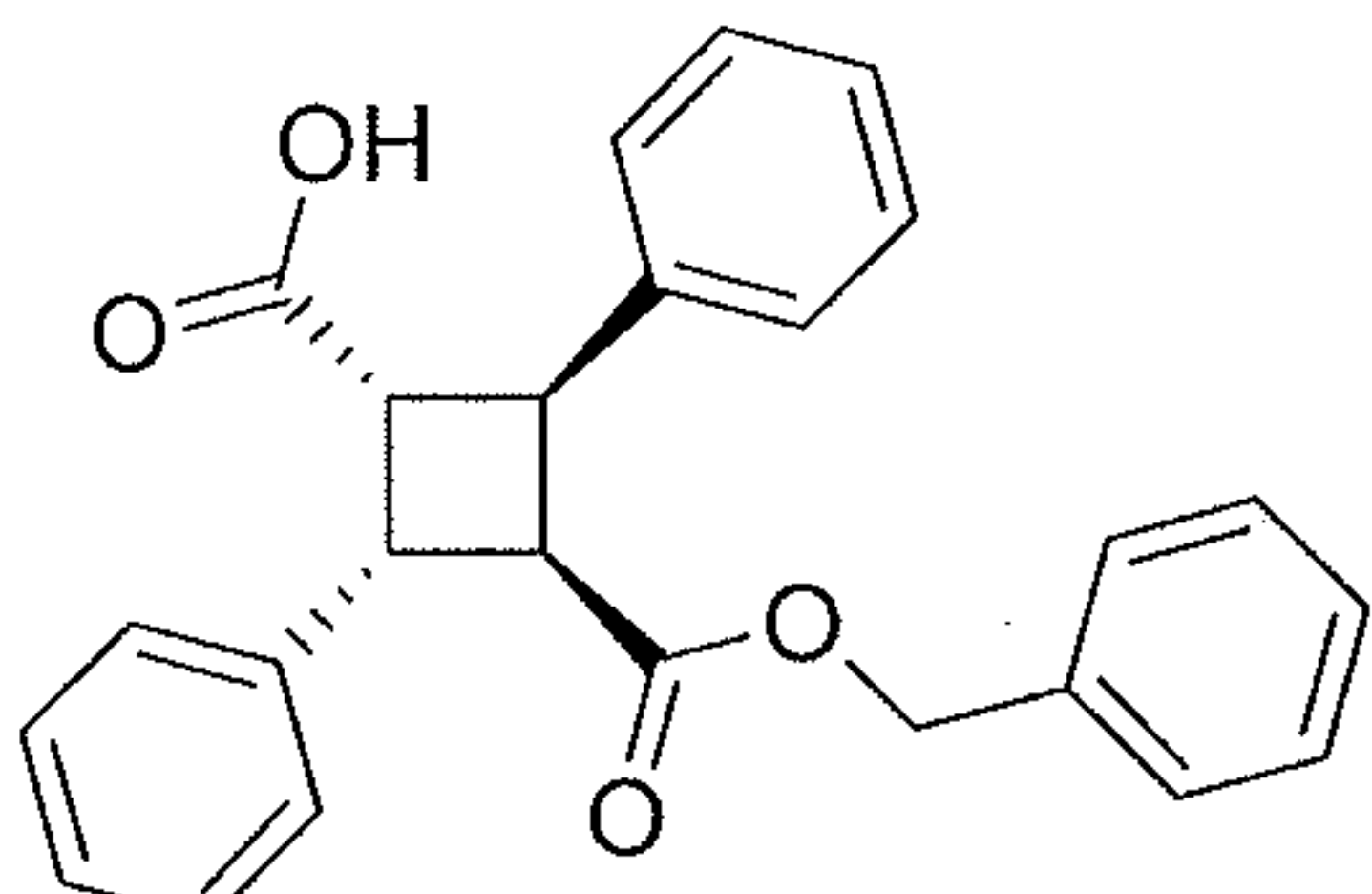
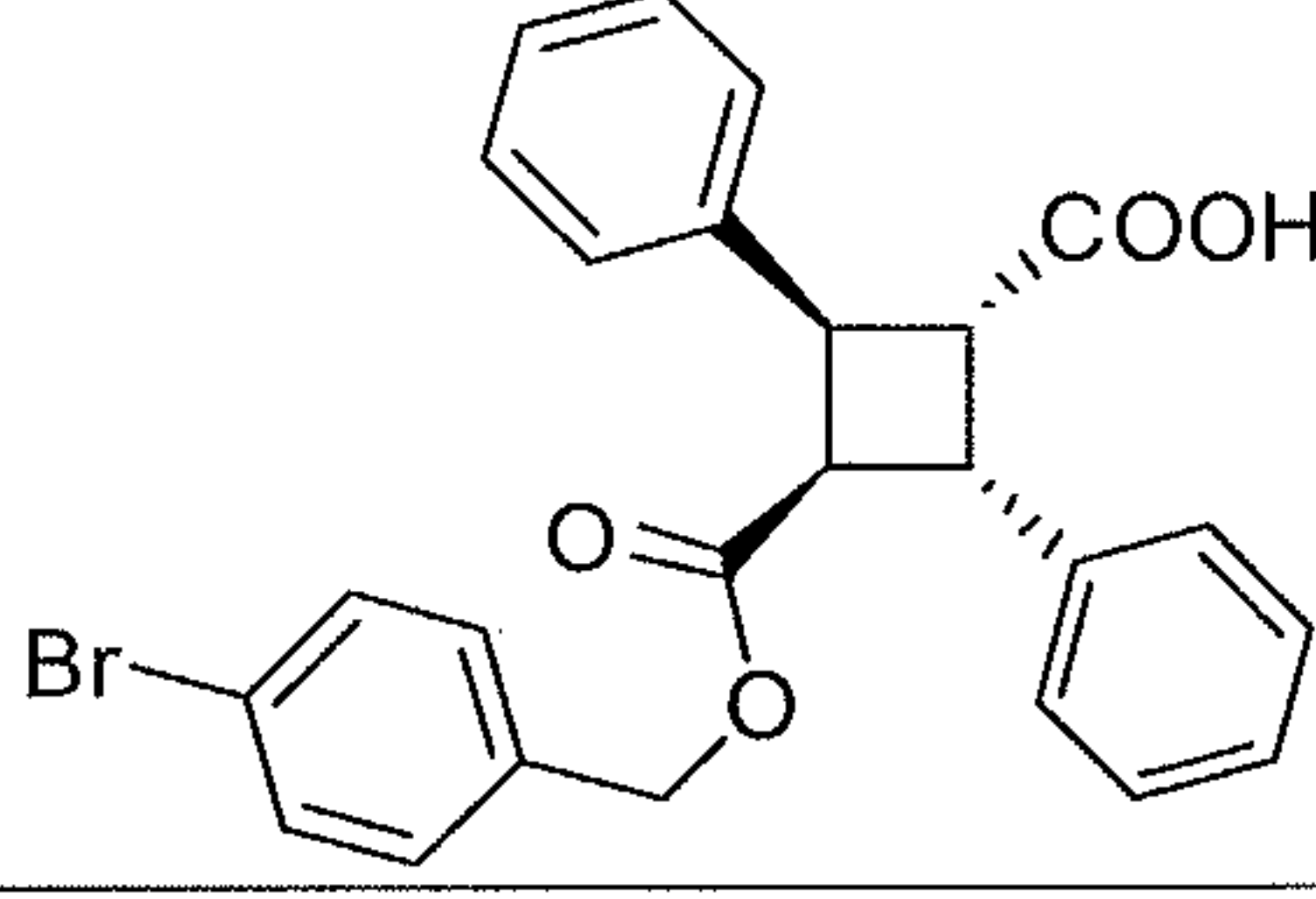
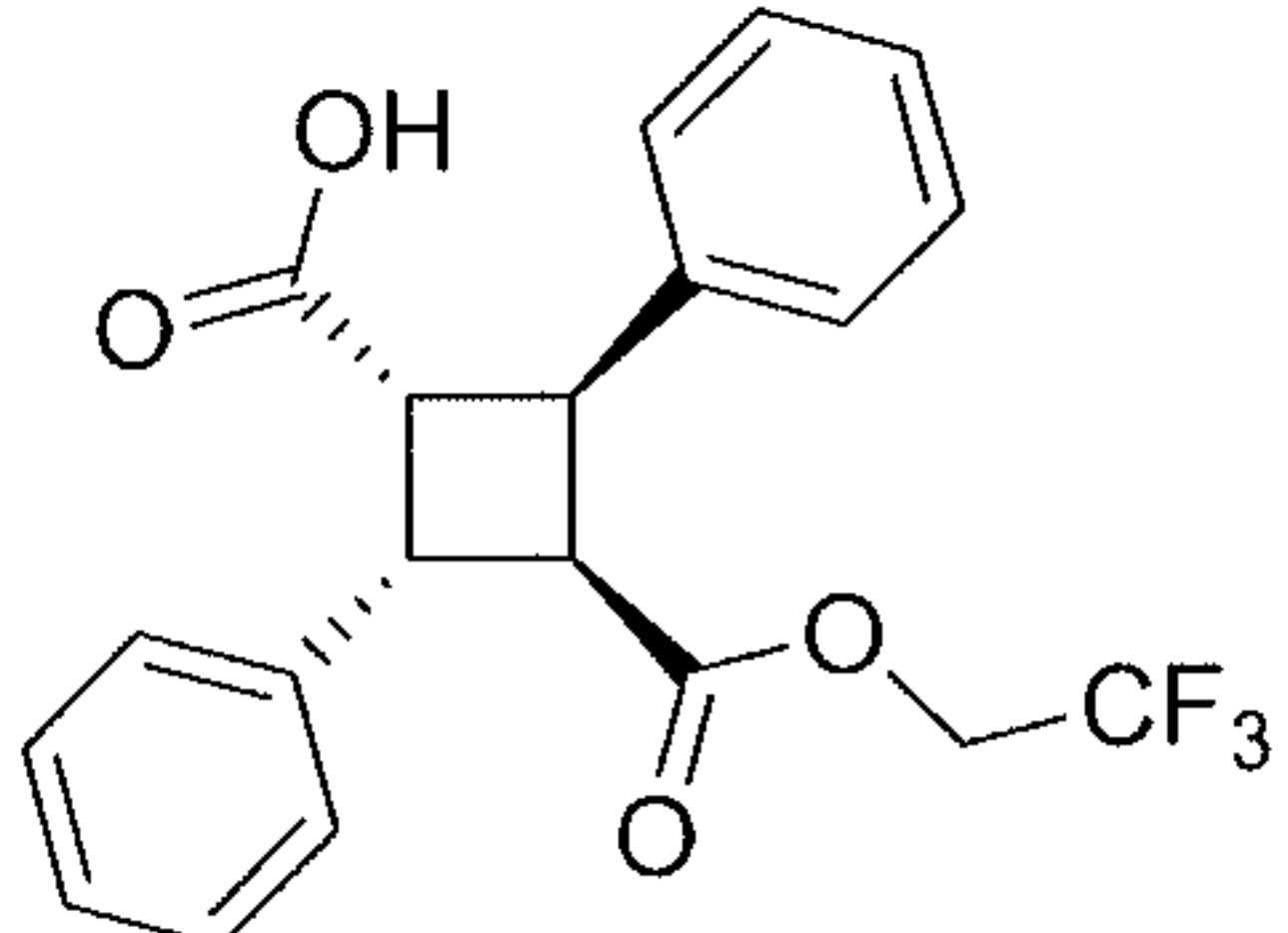
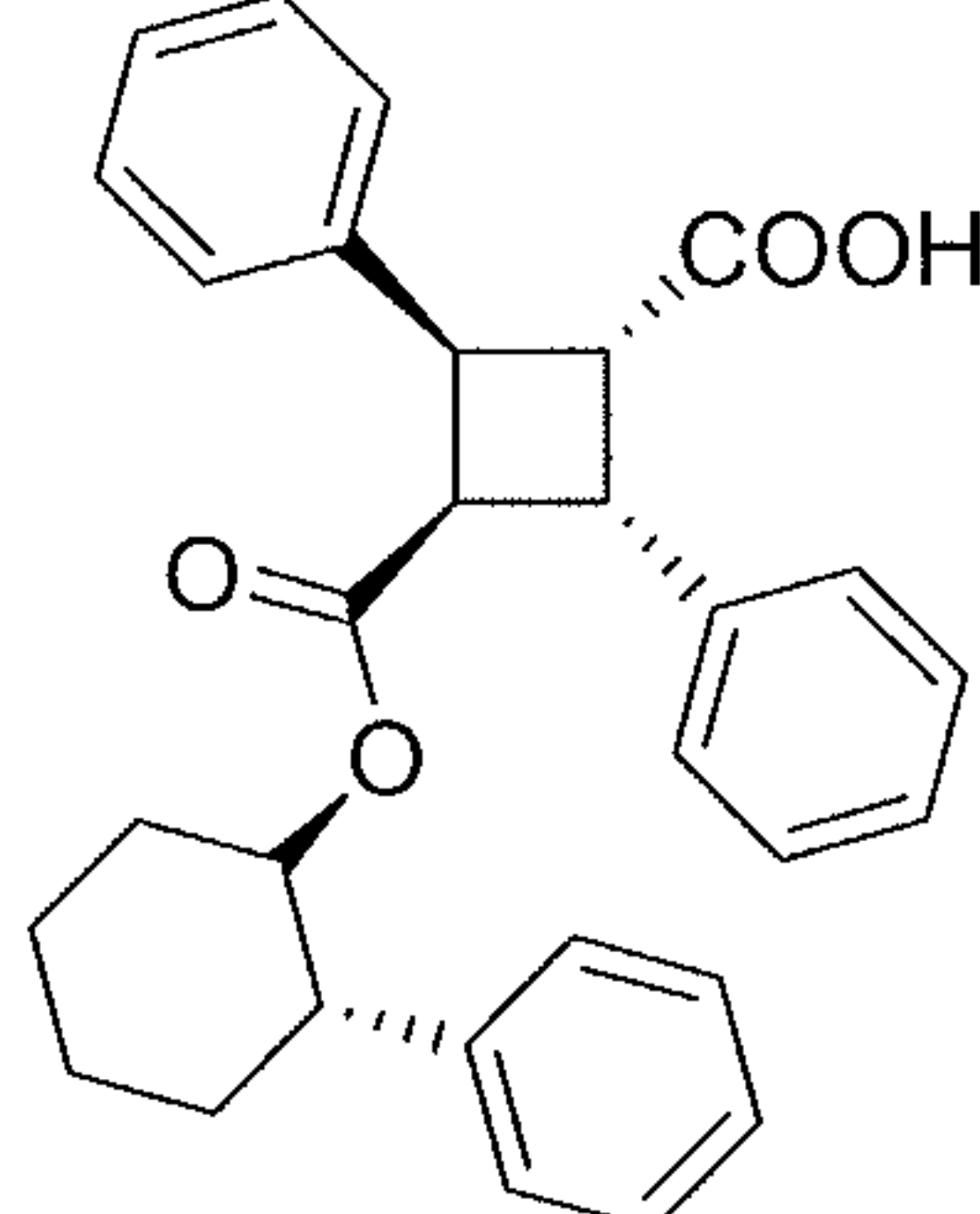
In some embodiments, the method wherein the compound has the structure:

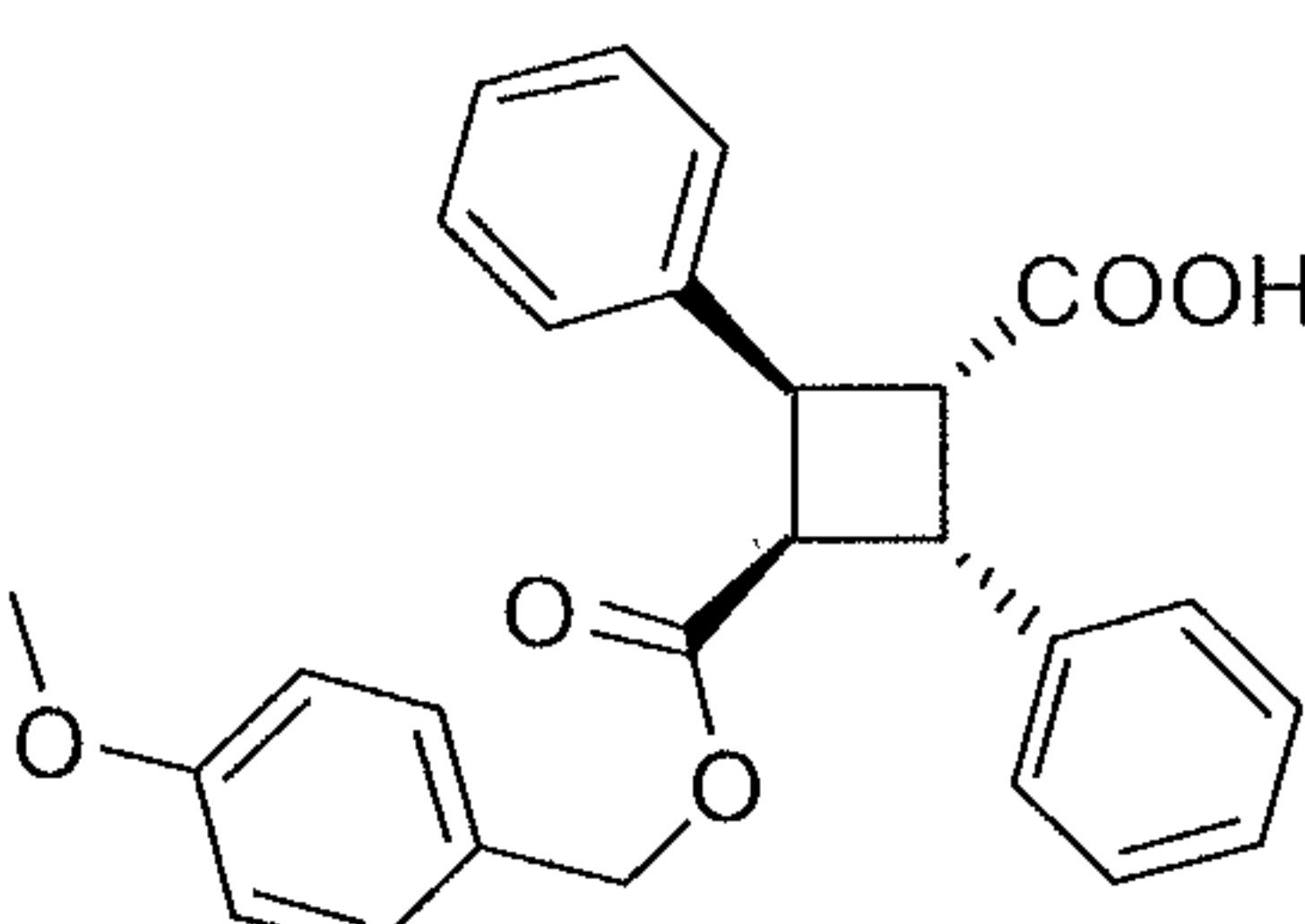
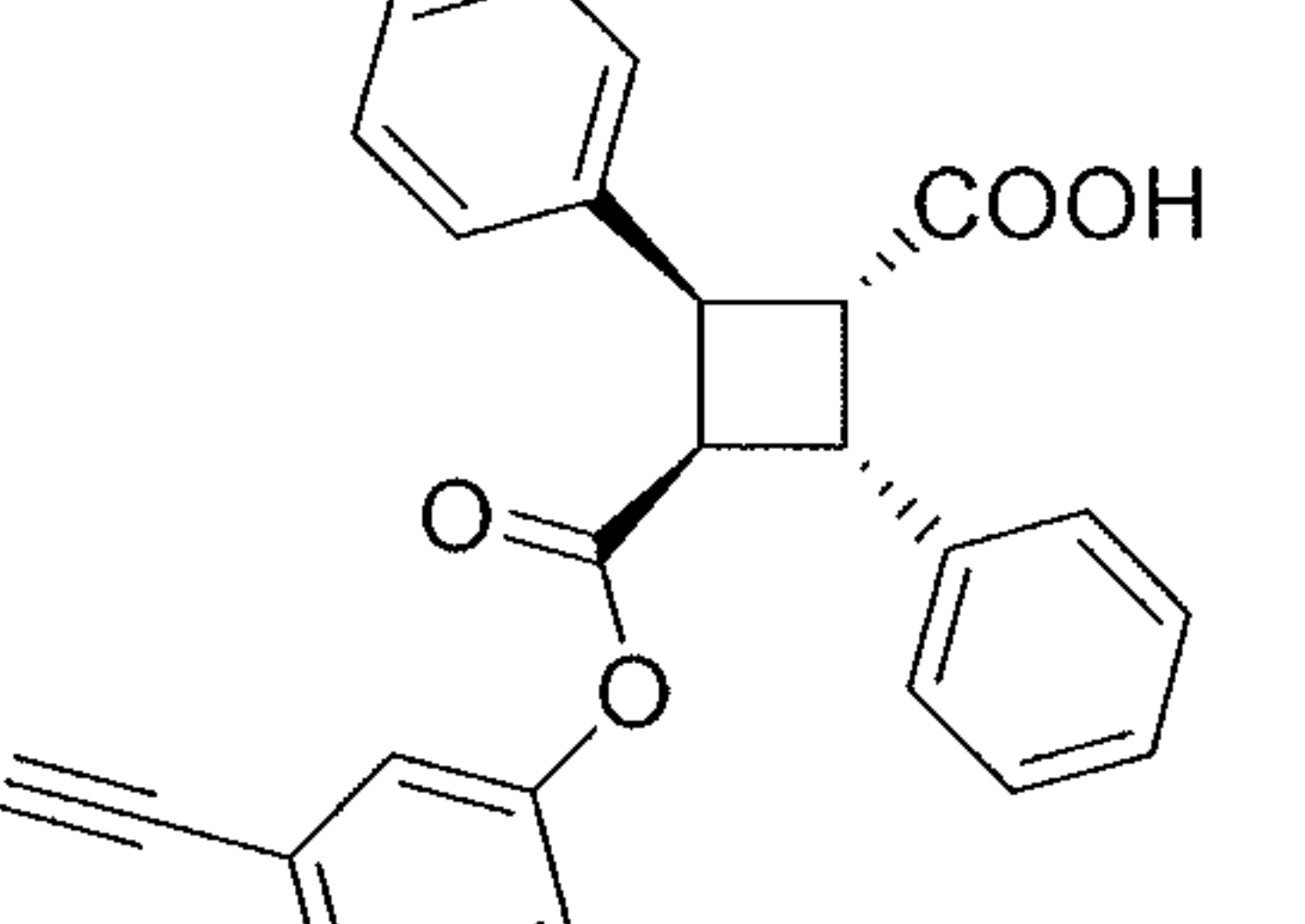
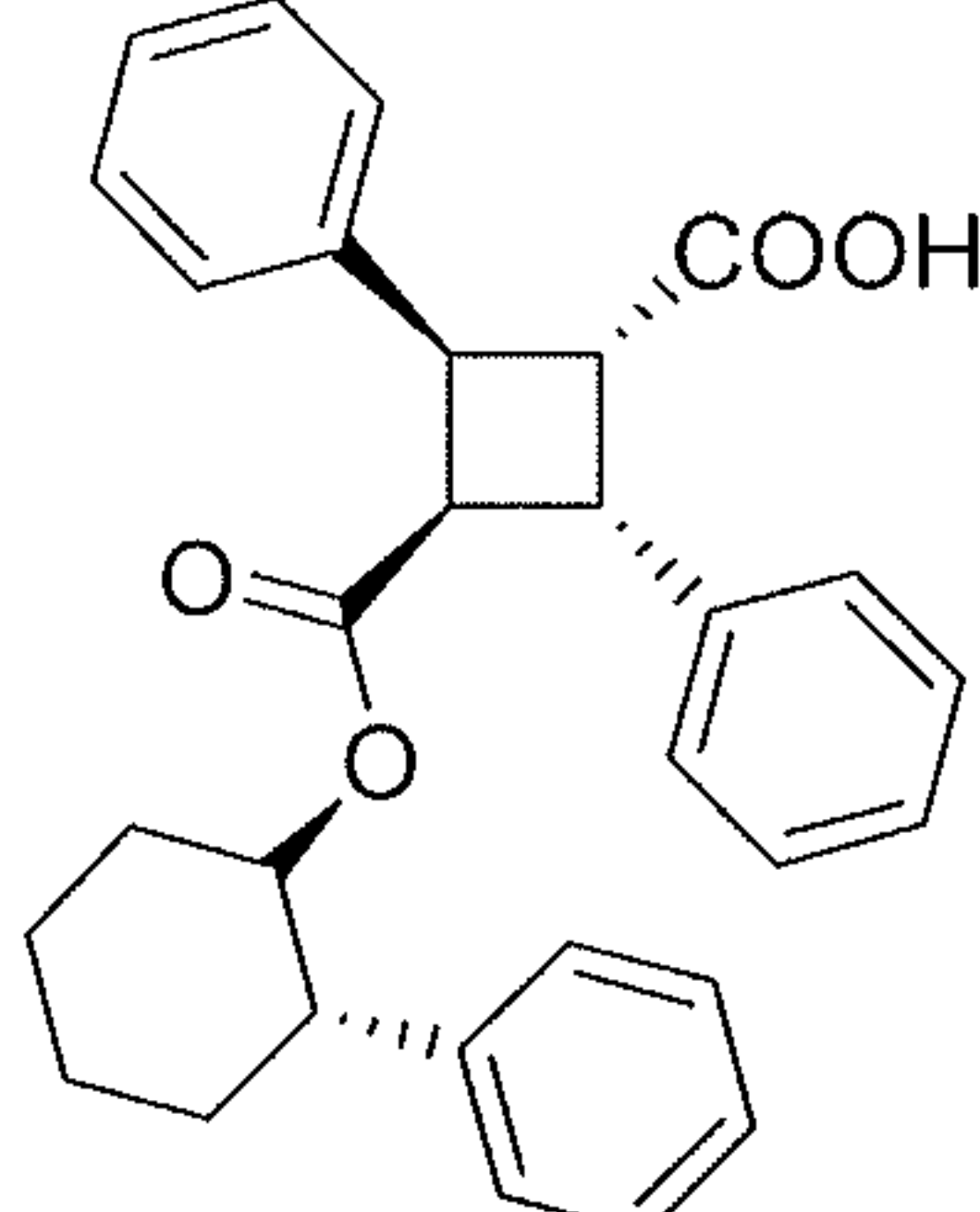
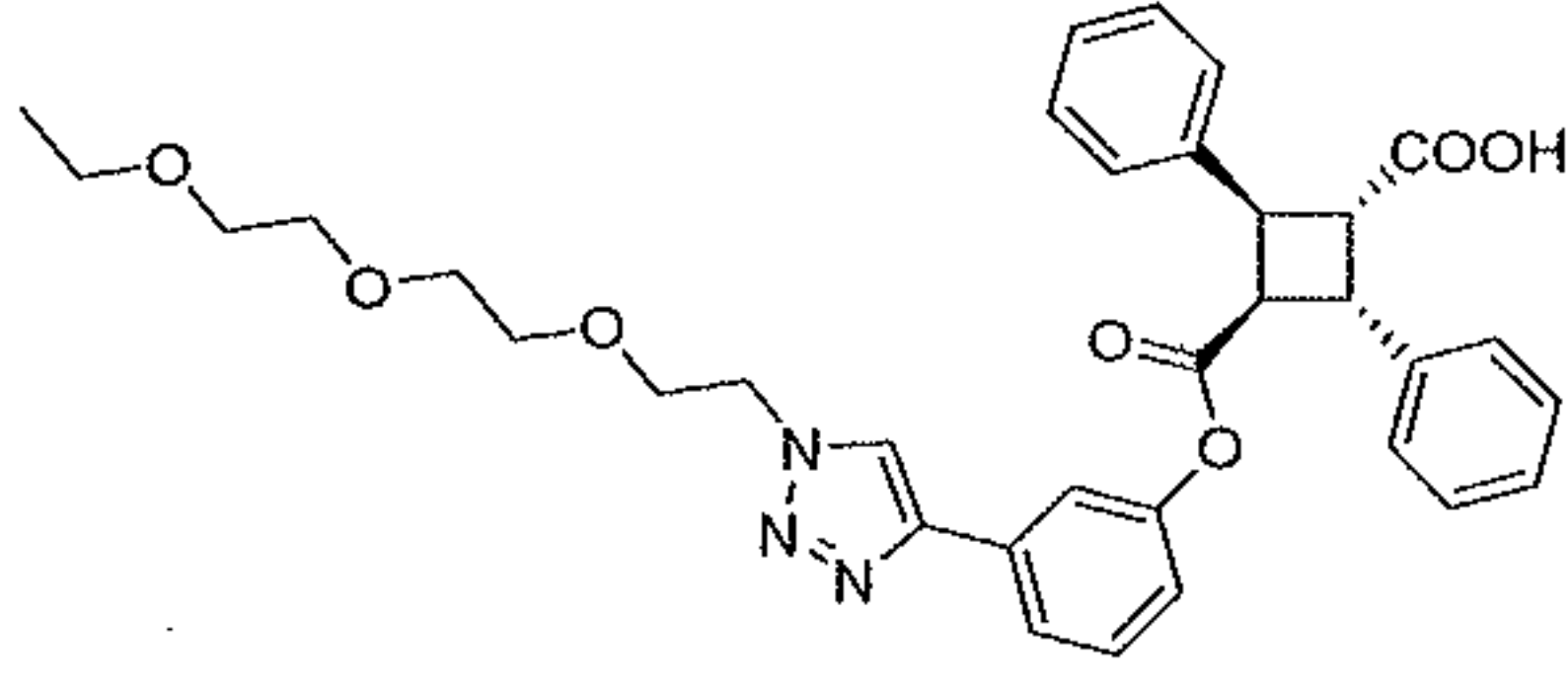
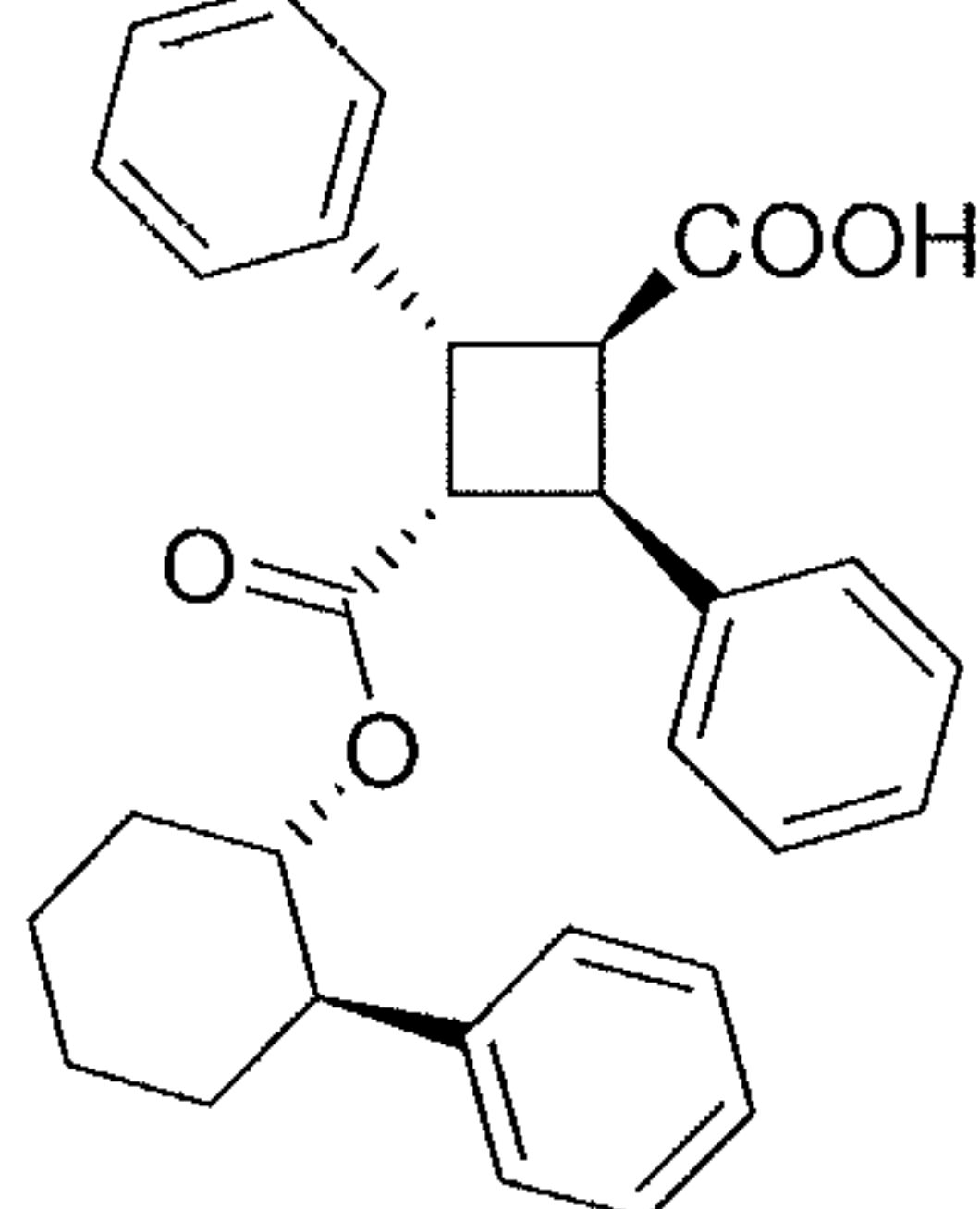
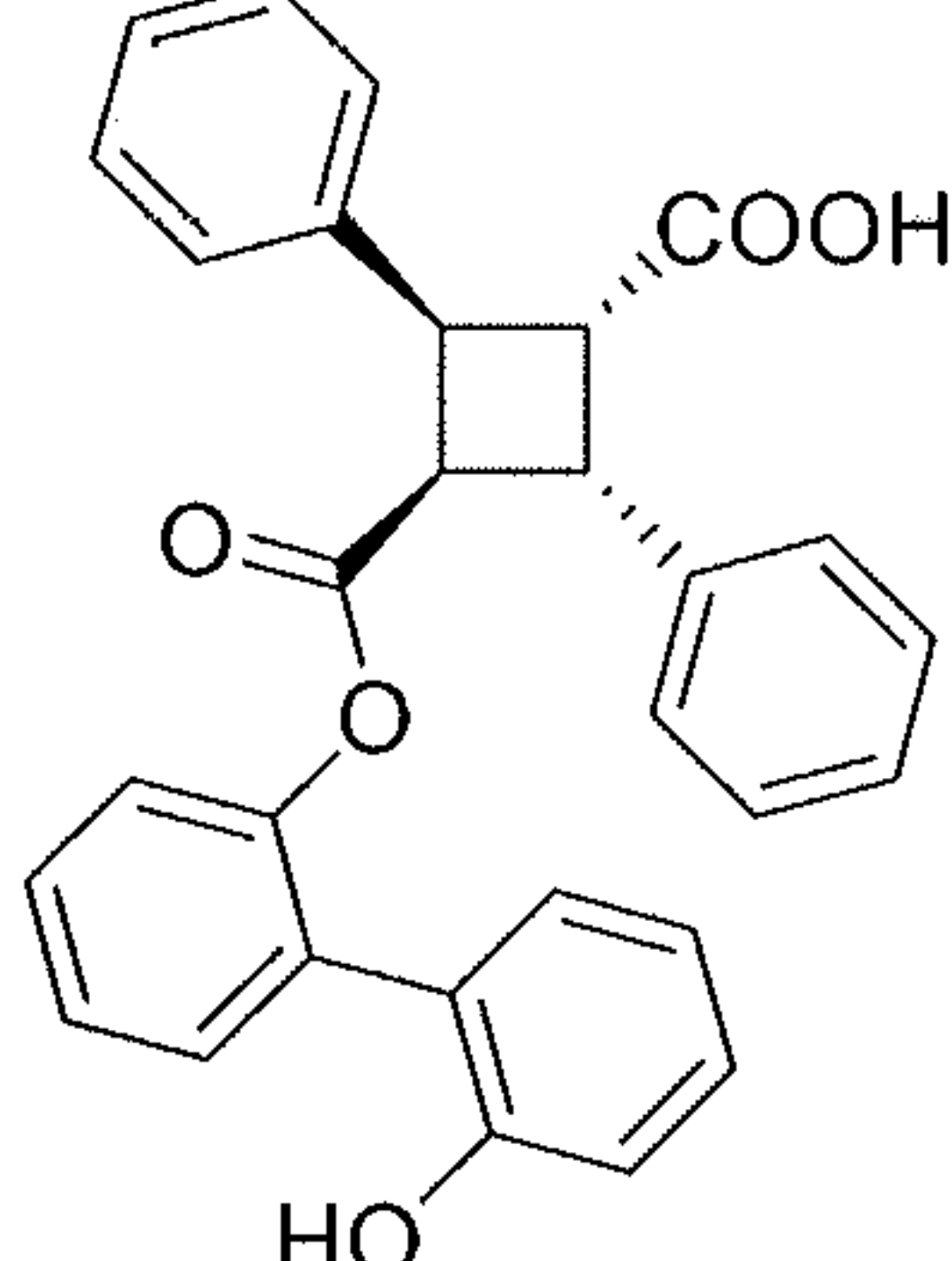
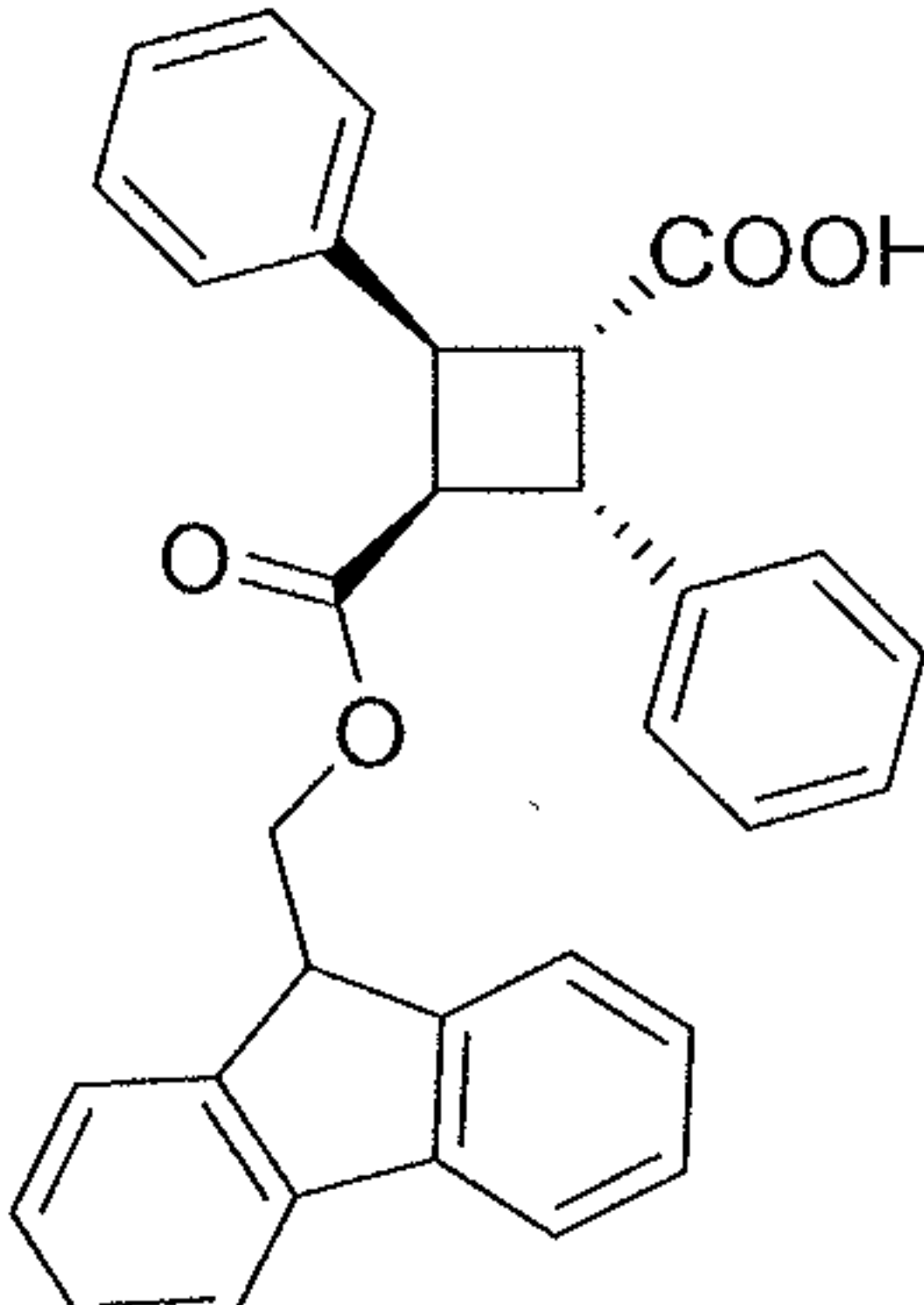
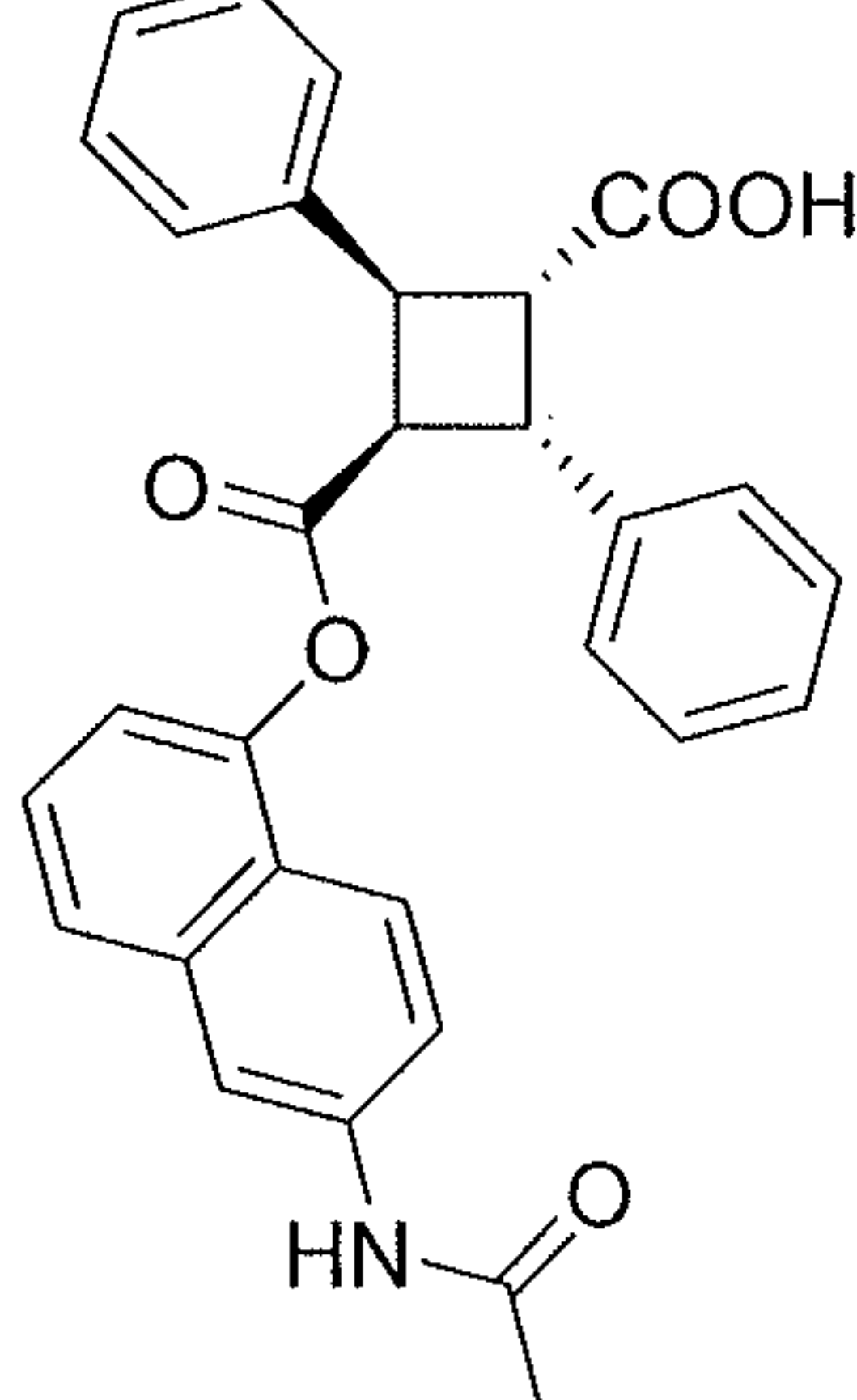


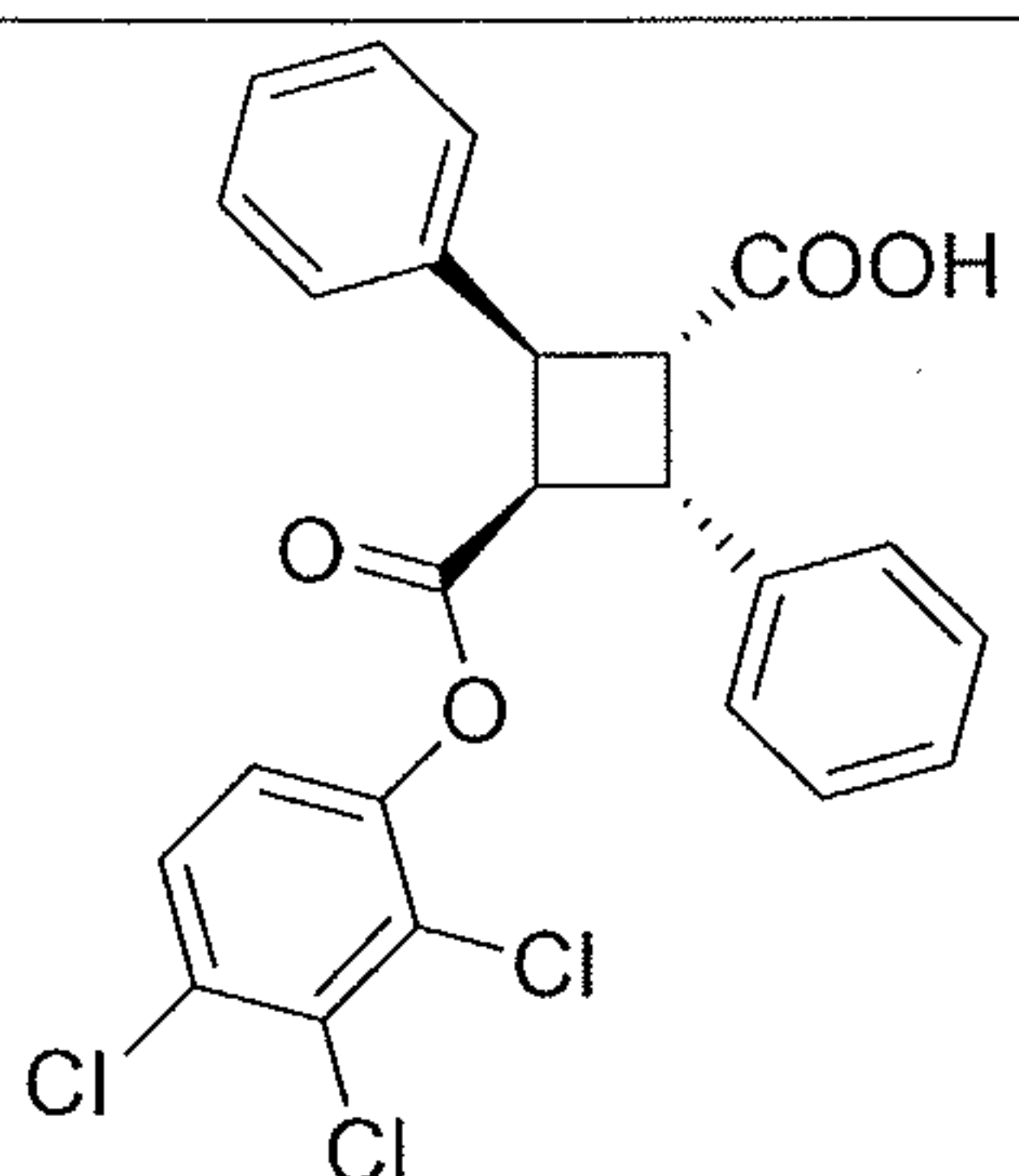
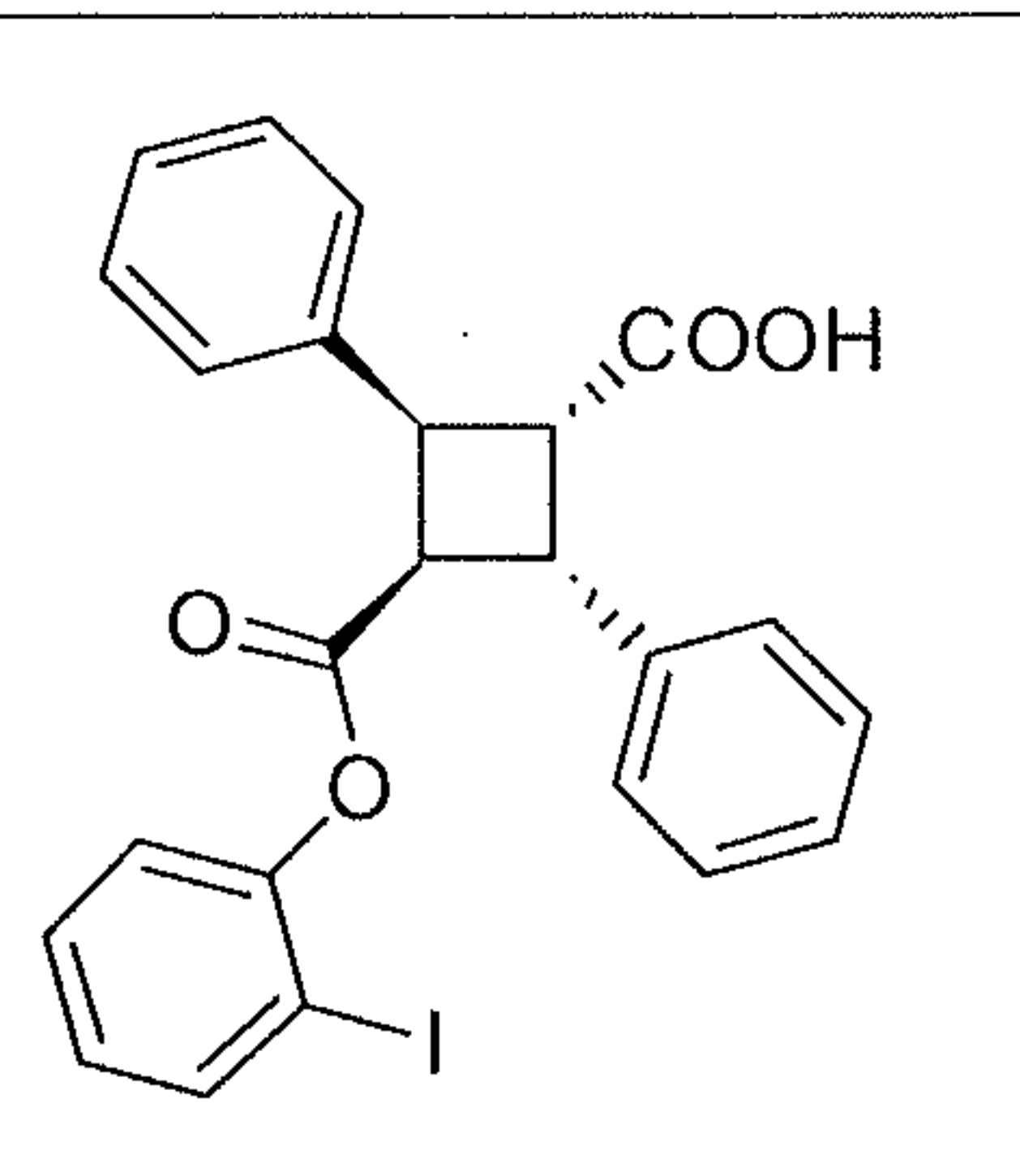
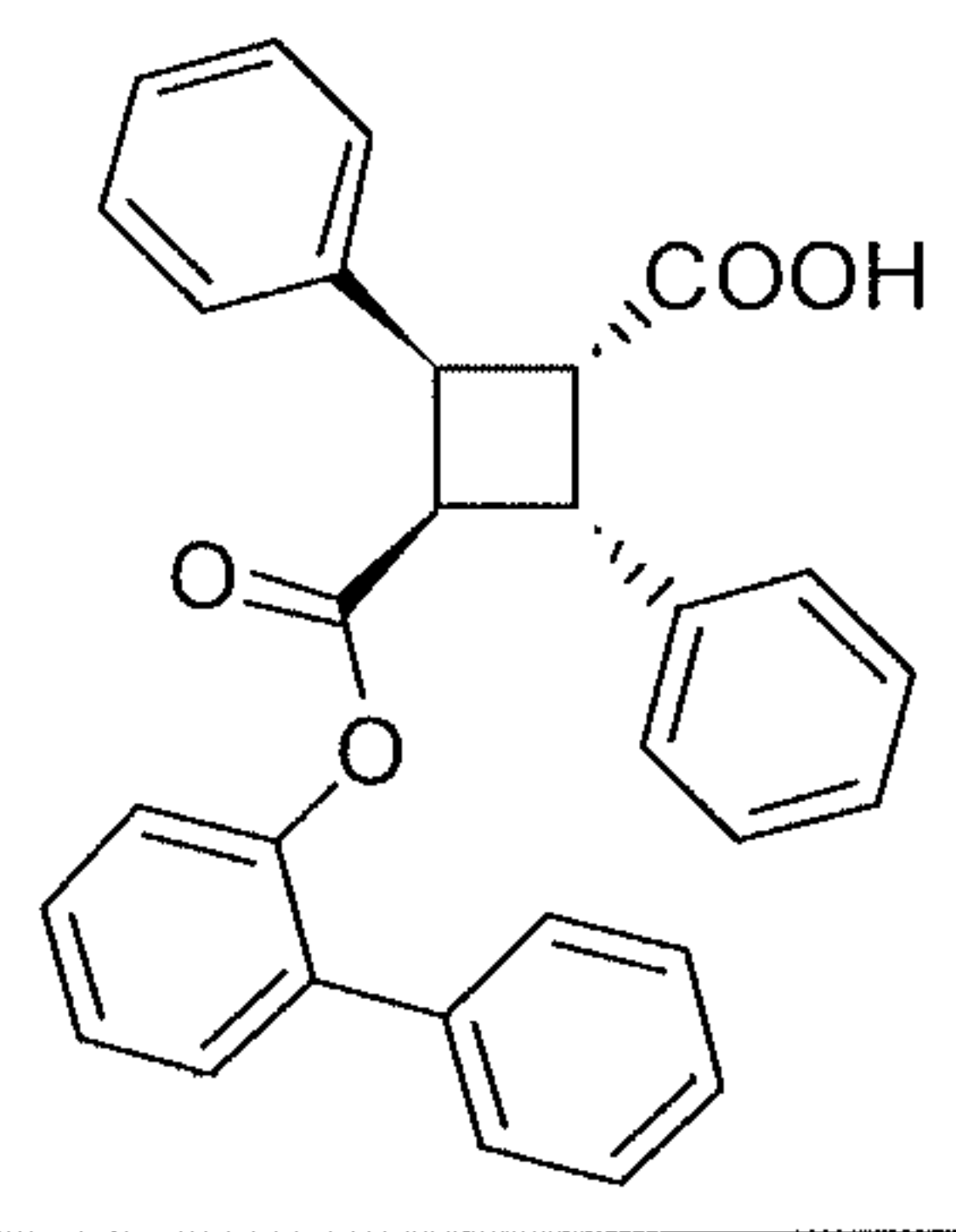
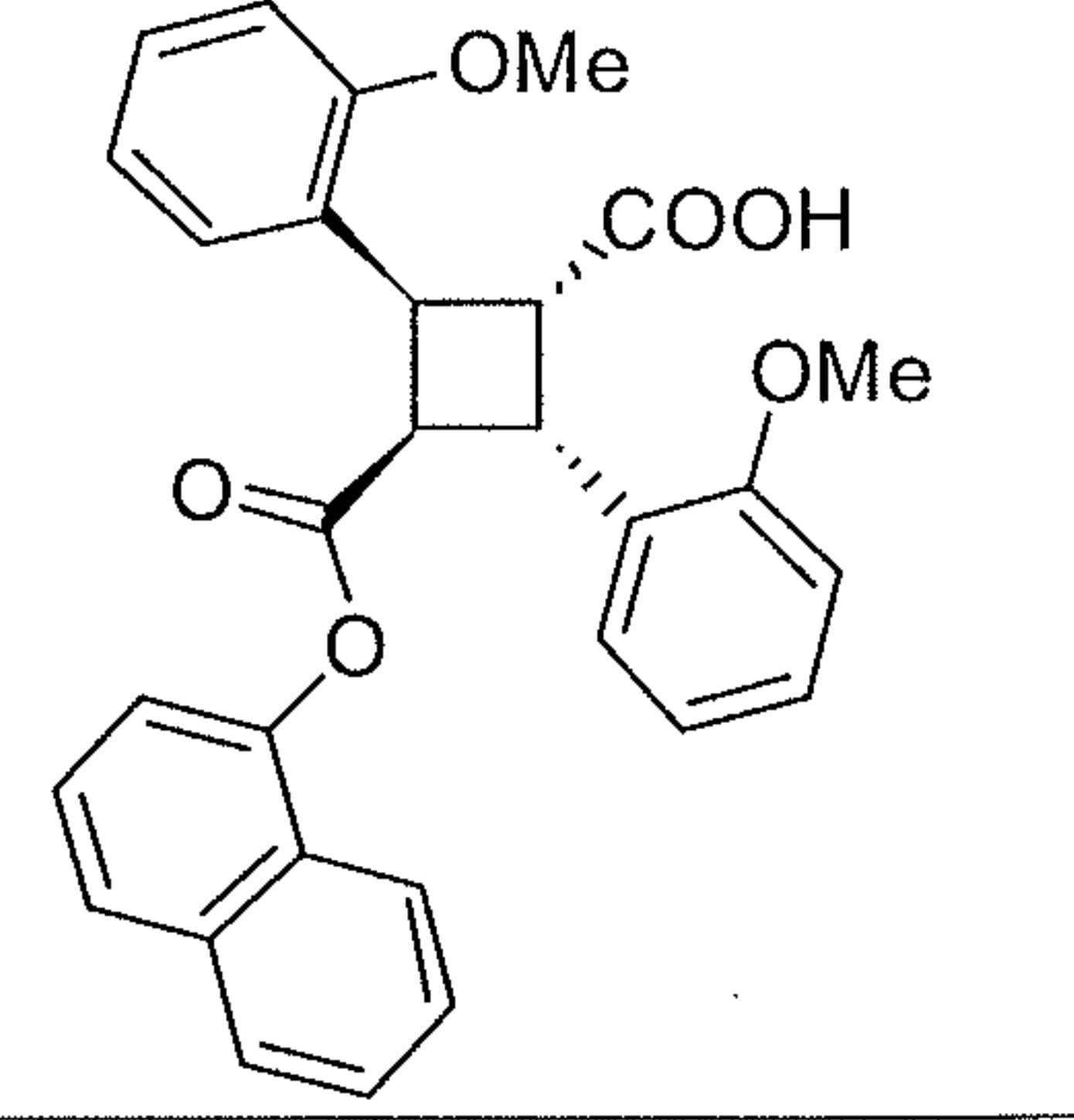
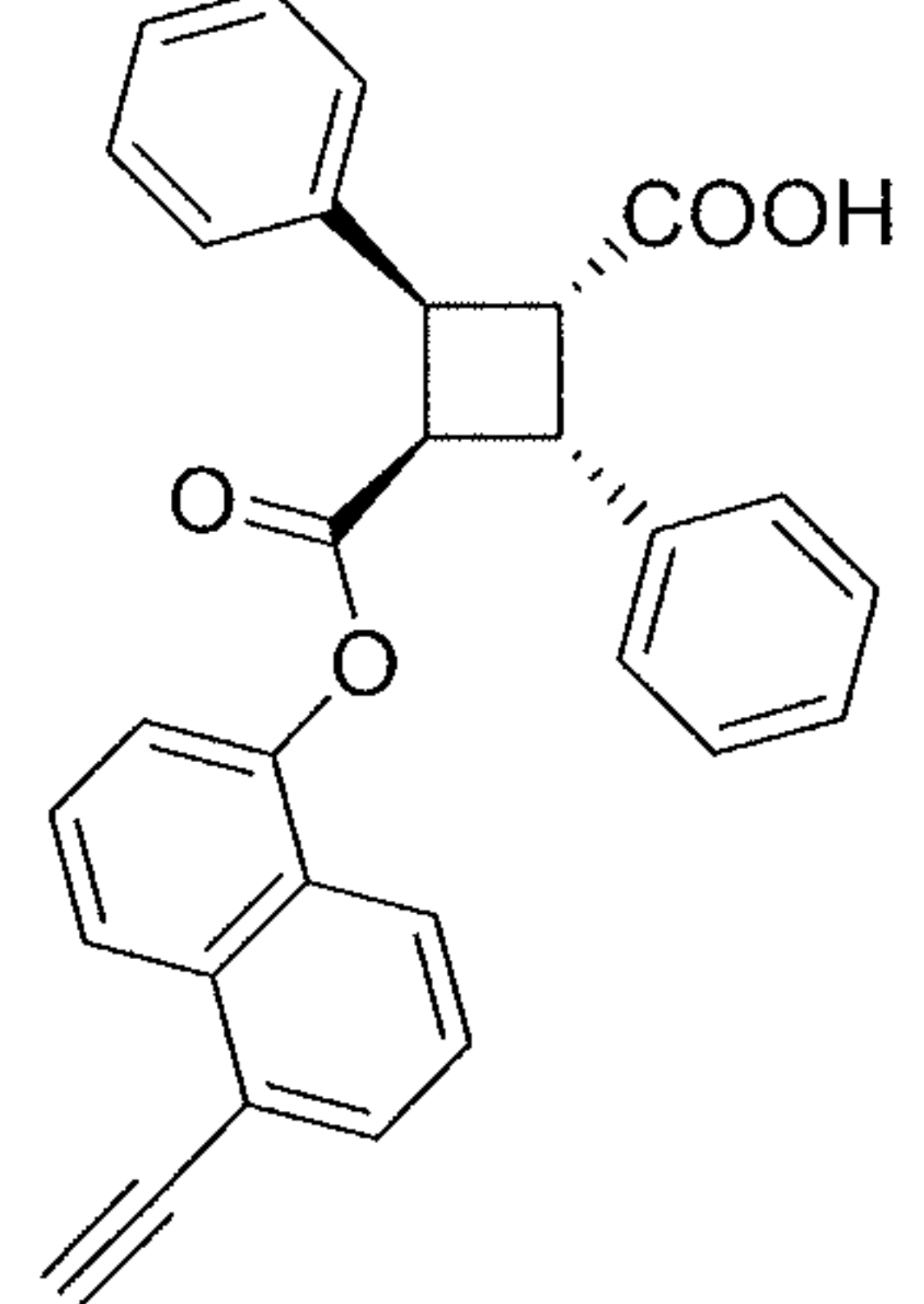
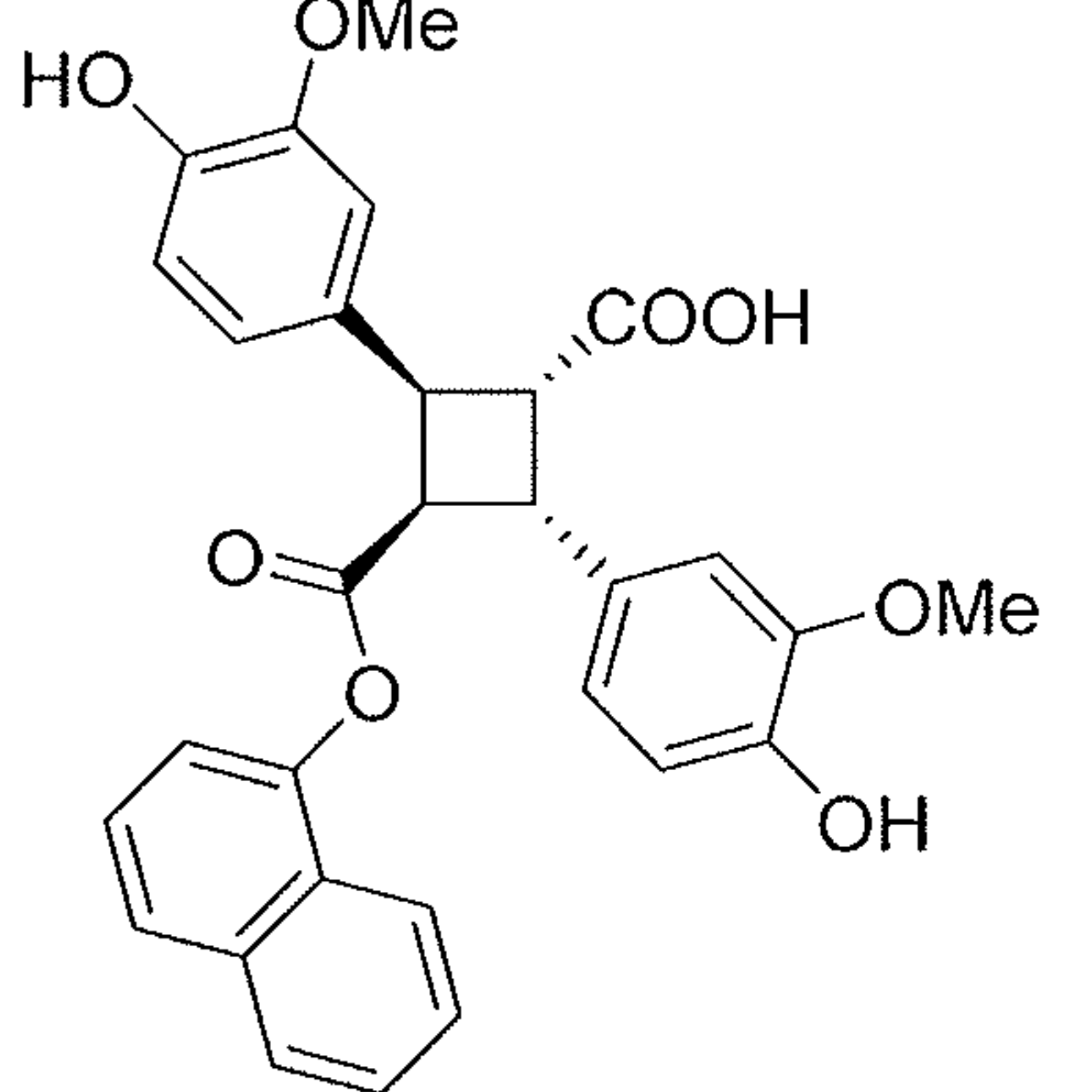
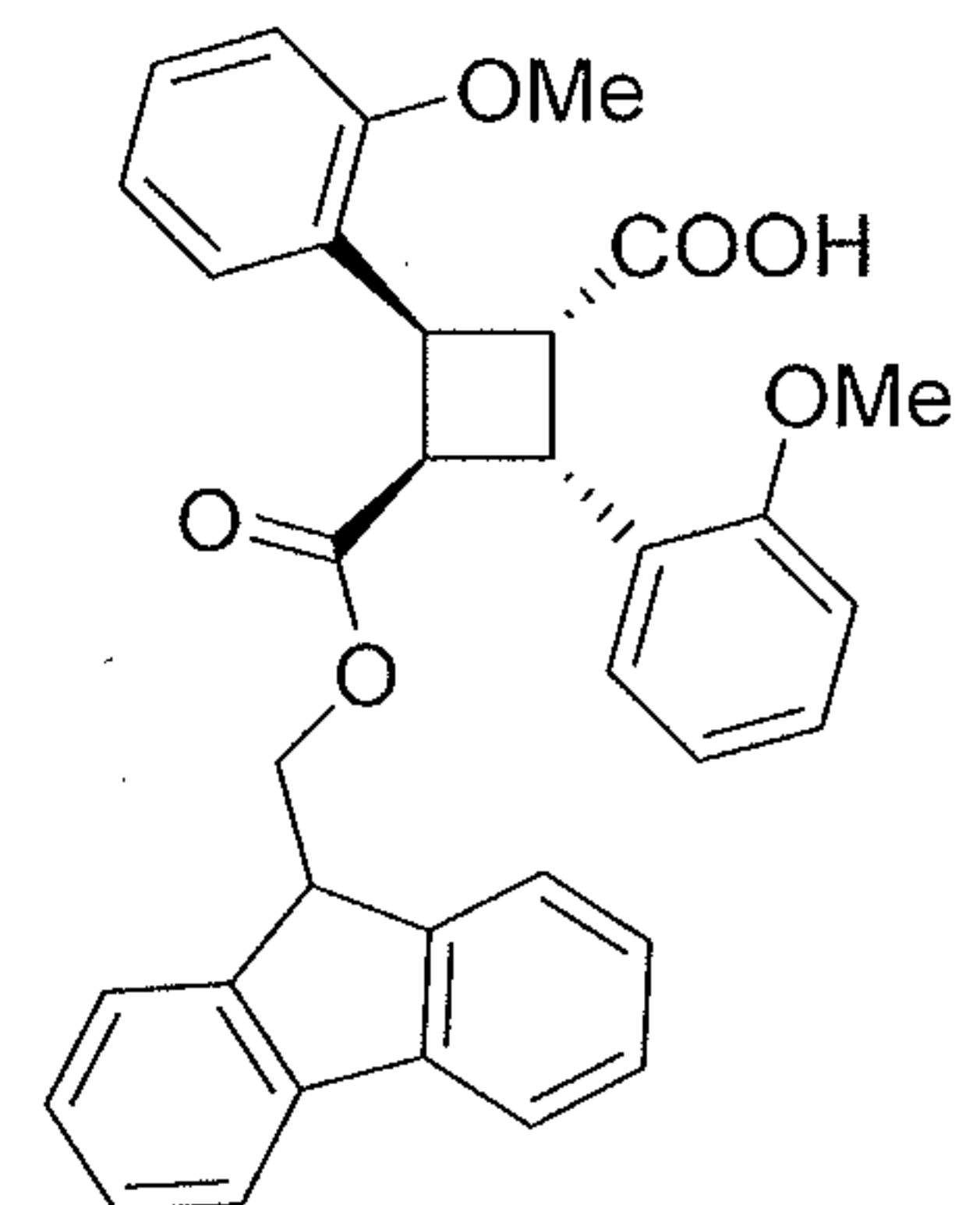
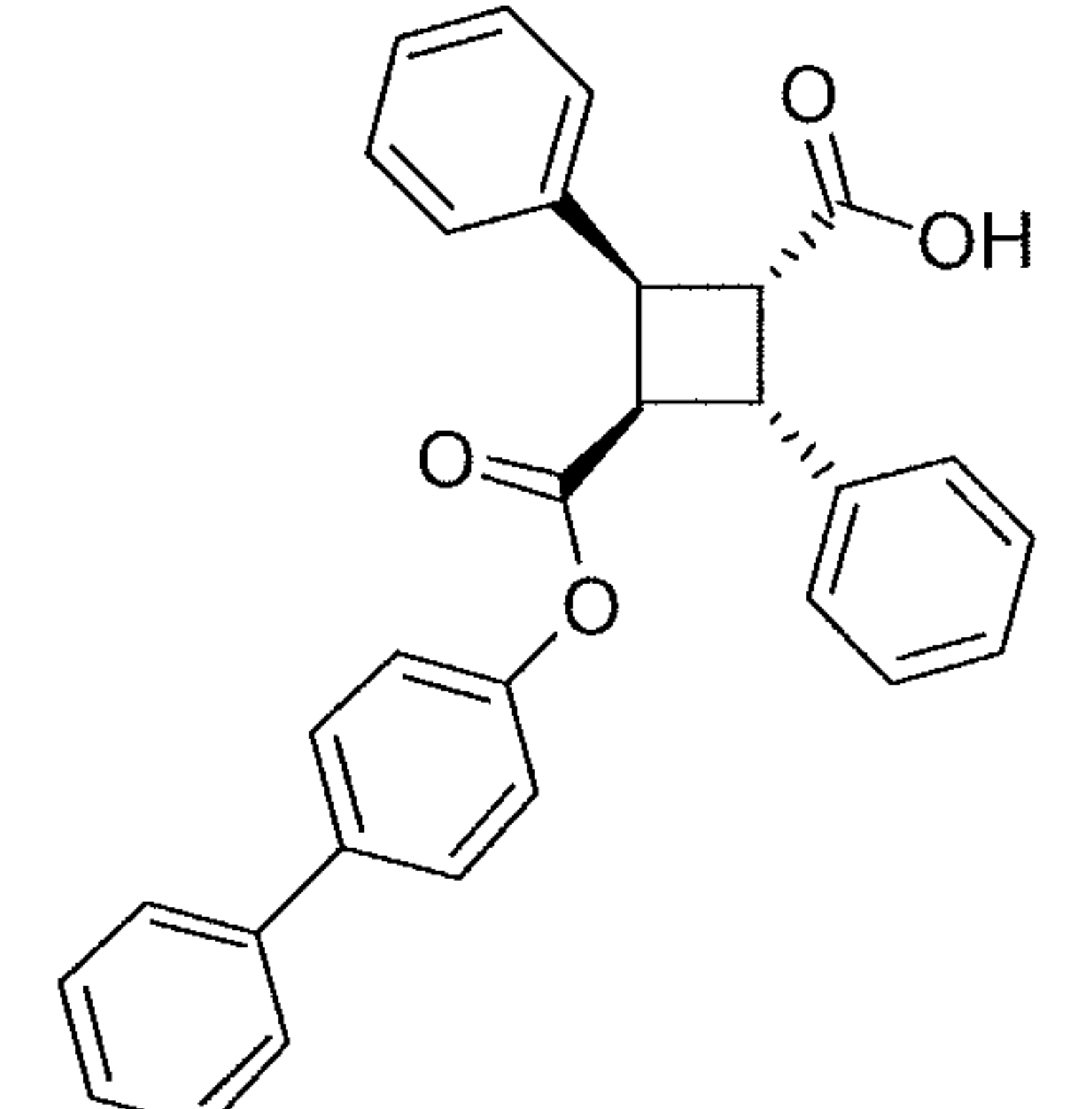
or a pharmaceutically acceptable salt thereof.

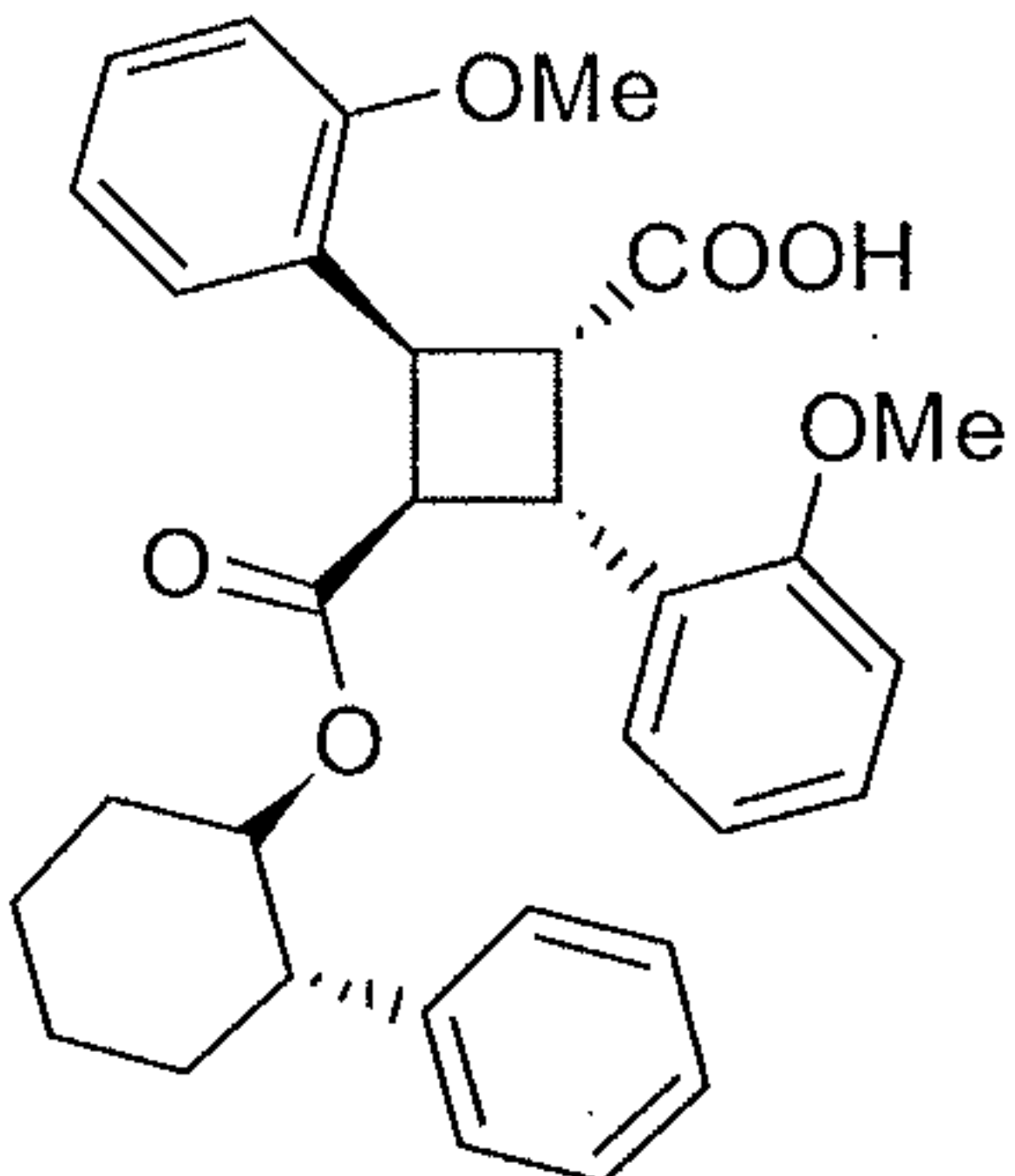
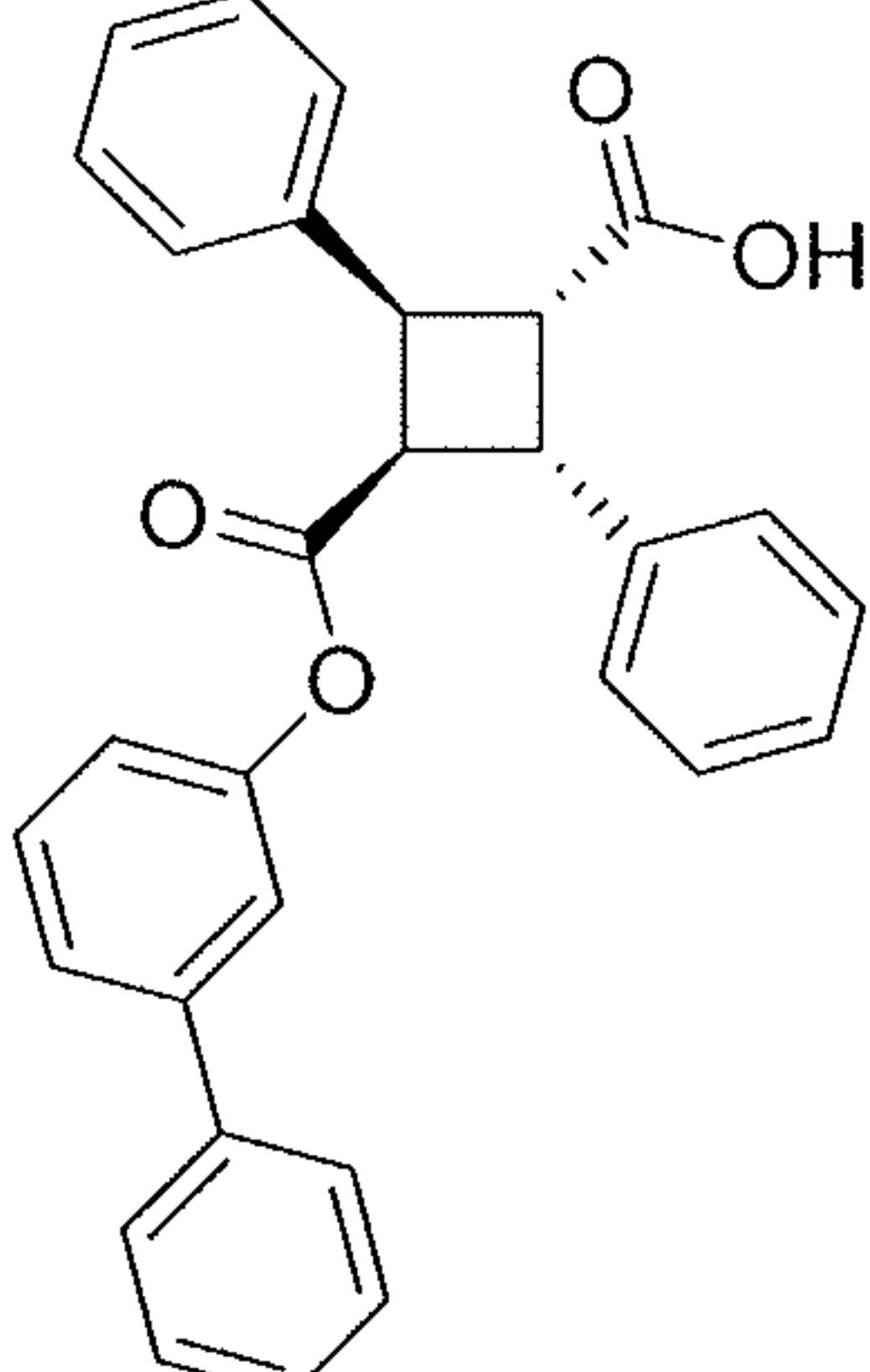
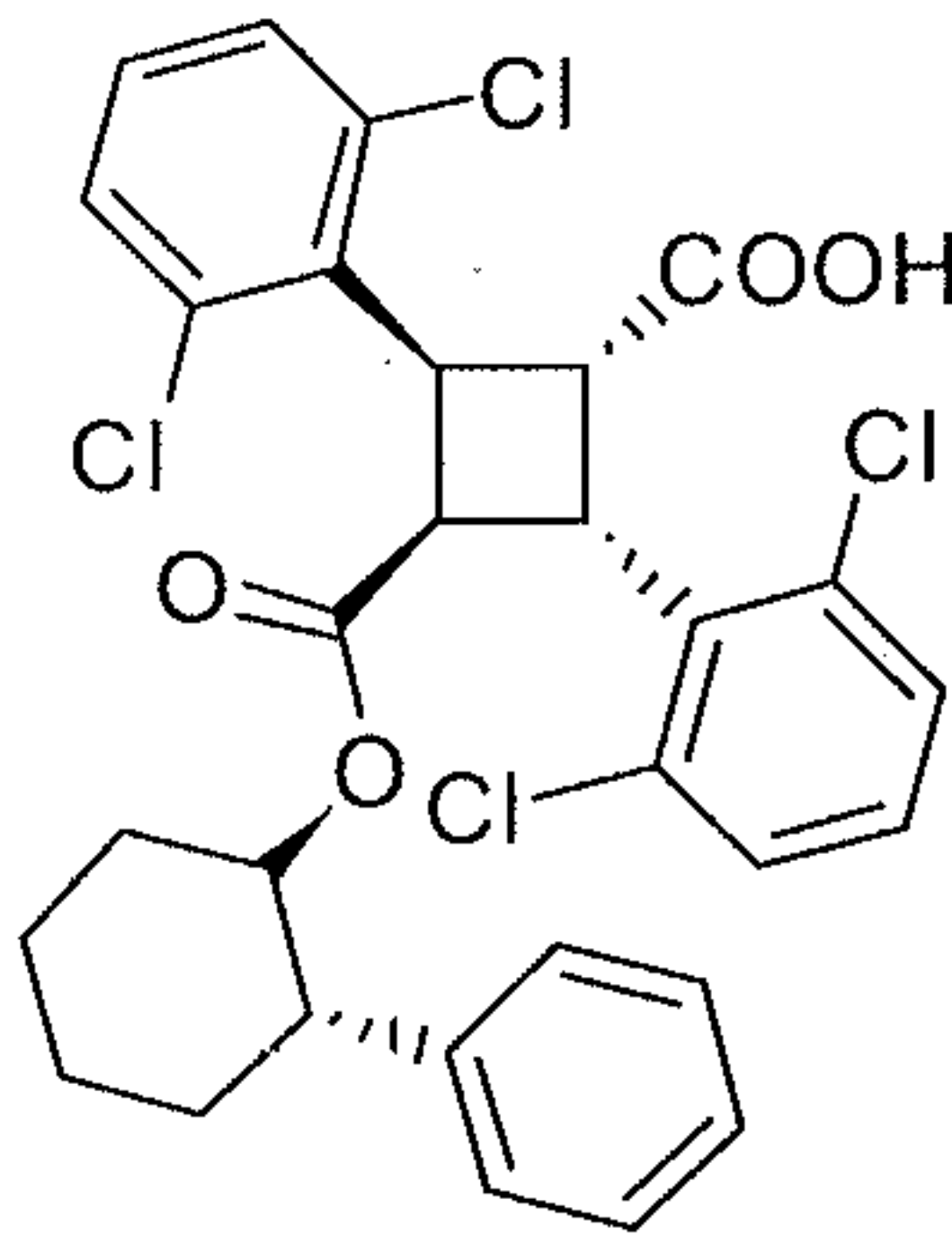
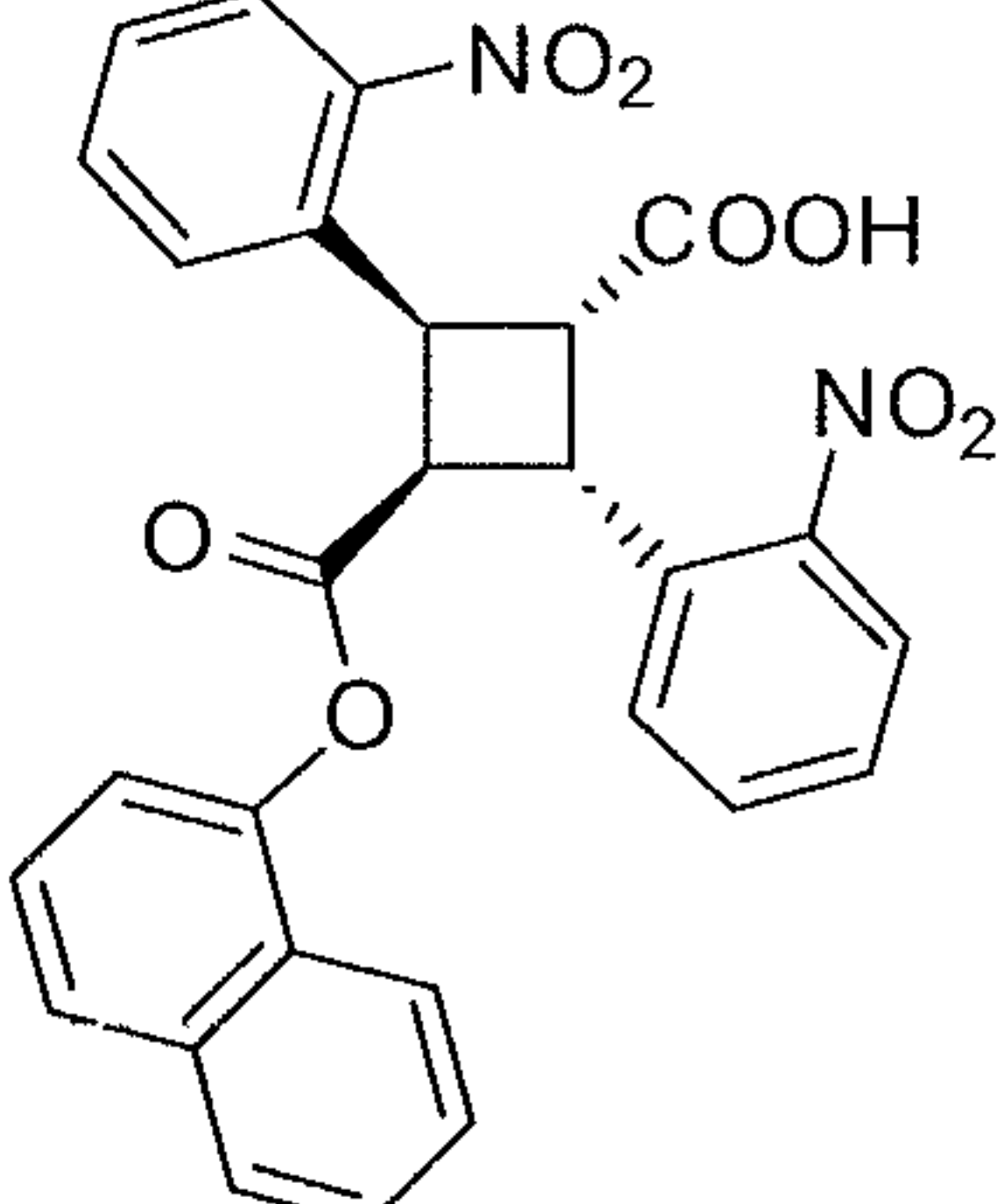
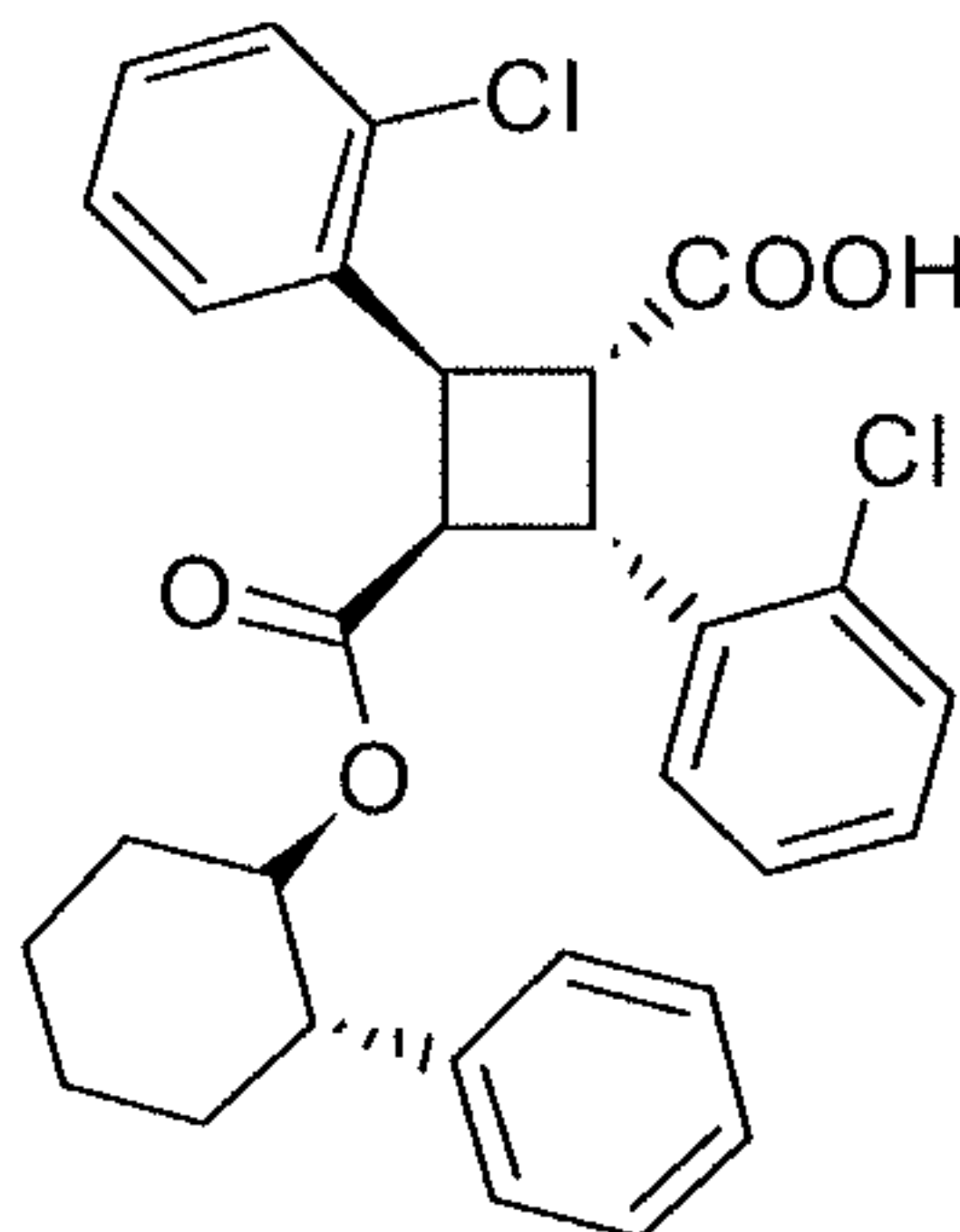
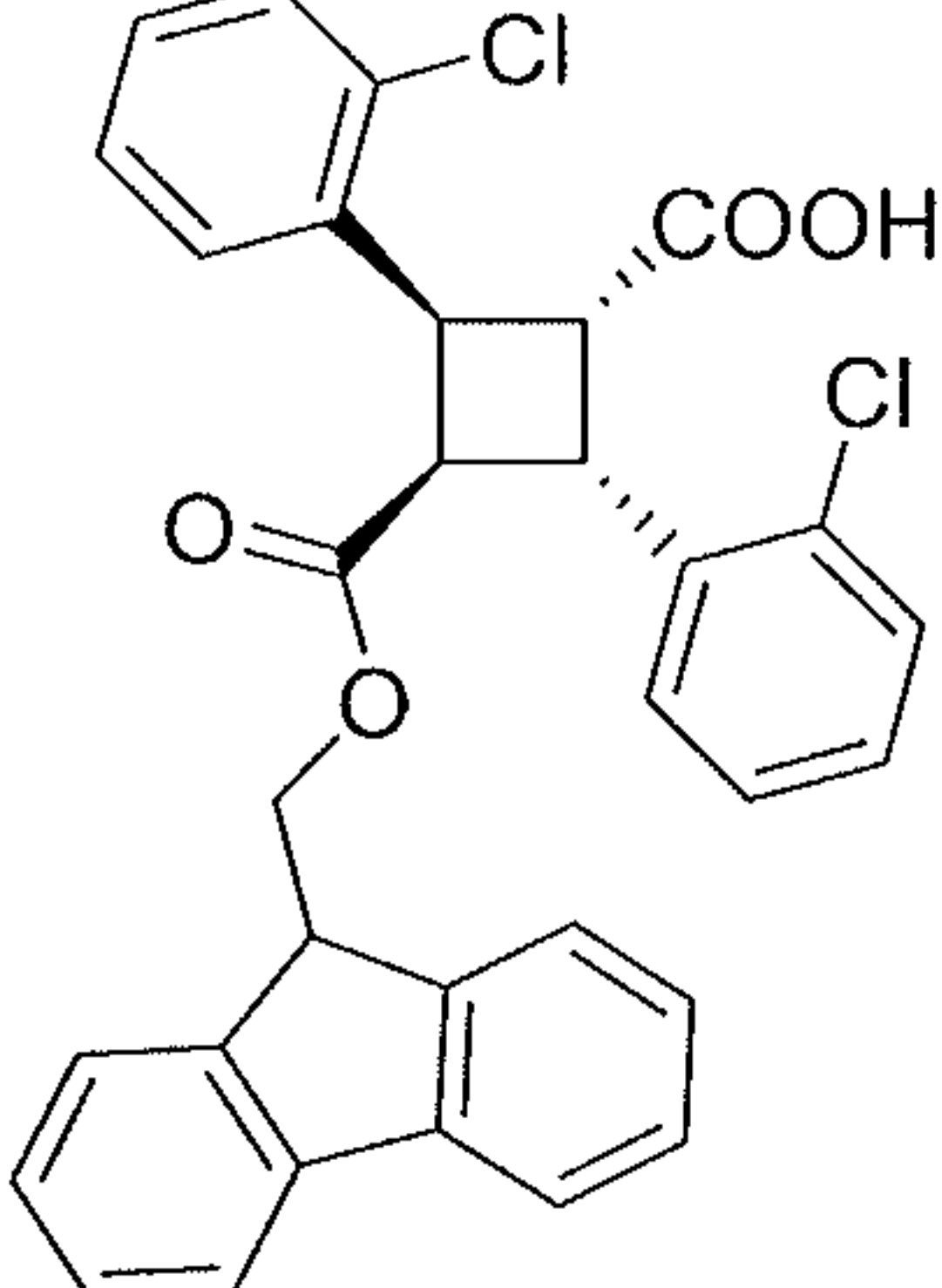
Compound **3** refers to a racemic mixture of the above compounds **3A** and **3B**.
25

Compounds of the present invention include the following:

Compound	Compound Structure	Compound	Compound Structure
3f		3m	
4d		3c	
3a		3d	
3r		3l	

3b		3f	
3l-A		3t	
3l-B		3j	
3q		3o	

3k		3e	
3g		4b	
3p		4a	
4j		3i	

4e		3h	
4g		4c	
4f		4k	

In some embodiments, the compound is the (S,S) enantiomer. In some embodiments, the compound is the (R,R) enantiomer.

5

In some embodiments, the composition comprises a mixture of enantiomers enriched in (S,S) enantiomer. In some embodiments, the composition comprises a mixture of enantiomers enriched in (R,R) enantiomer.

10 In some embodiments, the method wherein the compound is the (S,S) enantiomer. In some embodiments, the method wherein the compound is the (R,R) enantiomer.

In some embodiments of the method of inhibiting the activity of a Fatty Acid Binding Protein (FABP), wherein the compound inhibits binding of an FABP ligand to the FABP.

5

In some embodiments of the method of inhibiting the activity of a Fatty Acid Binding Protein (FABP), wherein the FABP ligand is an endocannabinoid ligand.

10 In some embodiments of the method of inhibiting the activity of a Fatty Acid Binding Protein (FABP), wherein the FABP ligand is anandamide (AEA) or 2-arachidonoylglycerol (2-AG).

15 In some embodiments, a method of treating a neurological disorder which affects at least one of movement, memory, mood, appetite, nociception, endocrine regulation, thermoregulation, sensory perception, or cognitive functions.

20 In some embodiments, a method of treating a neurological disorder associated with drug addiction, depression, compulsive behavior, neuropathic pain, or a movement disorder.

25 In some embodiments, a method of treating drug addiction, depression, compulsive behavior, neuropathic pain, inflammatory pain, or a movement disorder.

In some embodiments, a method of treating pain, neuropathic pain, or inflammatory pain.

30 In some embodiments, a method of treating a subject afflicted with a neurological disorder which affects at least one of movement, memory, mood, appetite, nociception, endocrine regulation, thermoregulation, sensory perception, or cognitive functions, comprising administering to the subject a compound of the present application.

35

In some embodiments, a method of treating a subject afflicted with a neurological disorder associated with drug addiction, depression, compulsive behavior, neuropathic pain, or a movement disorder, comprising administering to the subject a compound of the present application.

In some embodiments, a method of treating a subject afflicted with drug addiction, depression, compulsive behavior, neuropathic pain, inflammatory pain, or a movement disorder, comprising administering to the subject a compound of the present application.

In some embodiments, a method of treating a subject afflicted with pain, neuropathic pain, or inflammatory pain, comprising administering to the subject a compound of the present application.

As used herein, the term "endocannabinoid" includes any molecule that activates cannabinoid receptors. Examples of such receptors are CB1 and CB2. Examples of endocannabinoids are arachidonoyl ethanolamide (AEA) and 2-arachidonoyl glycerol (2-AG).

As used herein, the term "fatty acid binding protein" or "FABP" refers to fatty acid binding proteins (FABPs) that function as intracellular carriers that shuttle cannabinoids (and by extension fatty acid amides (FAAs)) to FAAH where cannabinoids are hydrolyzed and degraded. Further, uptake of endocannabinoids (and by extension FAAs) by the cell and the subsequent hydrolysis of endocannabinoids (and by extension FAAs) are enhanced by FABPs, and inhibiting the interaction of endocannabinoids (and by extension FAAs) with FABPs reduces endocannabinoid (and by extension FAA) uptake and hydrolysis. FABPs include, for example, fatty acid binding protein 1 (FABP 1), fatty acid binding protein 2 (FABP 2), fatty acid binding protein 3 (FABP 3), fatty acid binding protein 4 (FABP 4), fatty acid binding protein 5 (FABP 5), fatty acid binding protein 6 (FABP 6), fatty acid binding protein 7 (FABP 7), fatty acid binding protein 8 (FABP 8), fatty acid binding protein 9 (FABP 9), fatty acid binding protein 10 (FABP 10), fatty acid binding protein 11 (FABP 11),

fatty acid binding protein 5-like (FABP 5-like 1), fatty acid binding protein 5-like 2 (FABP 5-like 2), fatty acid binding protein 5-like 3 (FABP 5-like 3), fatty acid binding protein 5-like 4 (FABP 5-like 4), fatty acid binding protein 5-like 5 (FABP 5-like 5), fatty acid binding protein 5-like 6 (FABP 5-like 6), and fatty acid binding protein 5-like 7 (FABP 5-like 7) (see Chmurzynska et al. 2006 and PCT International Application Publication No. WO 2010/083532 A1, the contents of each of which are hereby incorporated by reference).

10 As used herein, the term "therapeutic agent" refers to any agent used to treat a disease or that provides a beneficial therapeutic effect to a subject.

As used herein, the phrase "inhibits the interaction" is employed herein to refer to any disruption, partial or total, of the natural effect of FABPs on the metabolism of endocannabinoids.

As used herein, the term "activity" refers to the activation, production, expression, synthesis, intercellular effect, and/or pathological or aberrant effect of the referenced molecule, either inside and/or outside of a cell. Such molecules include, but are not limited to, cytokines, enzymes, growth factors, pro-growth factors, active growth factors, and pro-enzymes. Molecules such as cytokines, enzymes, growth factors, pro-growth factors, active growth factors, and pro-enzymes may be produced, expressed, or synthesized within a cell where they may exert an effect. Such molecules may also be transported outside of the cell to the extracellular matrix where they may induce an effect on the extracellular matrix or on a neighboring cell. It is understood that activation of inactive cytokines, enzymes and pro-enzymes may occur inside and/or outside of a cell and that both inactive and active forms may be present at any point inside and/or outside of a cell. It is also understood that cells may possess basal levels of such molecules for normal function and that abnormally high or low levels of such active molecules may lead to pathological or aberrant effects that may be corrected by pharmacological intervention.

As used herein, "treating" means reducing, slowing, stopping, preventing, reversing, or in any way improving the progression of a disease or disorder or a symptom of the disease or disorder.

5

In some embodiments, the compounds of the present invention include all hydrates, solvates, and complexes of the compounds used by this invention.

10 In some embodiments, if a chiral center or another form of an isomeric center is present in a compound of the present invention, all forms of such isomer or isomers, including enantiomers and diastereomers, are intended to be covered herein.

15 In some embodiments, if a chiral center or another form of an isomeric center is present in a compound of the present invention, only enantiomeric forms are intended to be covered herein.

Compounds containing a chiral center may be used as a racemic mixture,
20 an enantiomerically enriched mixture, or the racemic mixture may be separated using well-known techniques and an individual enantiomer may be used alone. The compounds described in the present invention are in racemic form or as individual enantiomers. Example 6 provides a method by which to obtain the individual enantiomers of each of the compound
25 contained herein. A method by which to obtain the individual enantiomers is also described in WO 2014/015276, published January 23, 2014, the contents of which are hereby incorporated by reference.

As used herein, "enantiomers" are non-identical, non-superimposable
30 mirror images of each other. For any given chiral compound, only one pair of enantiomers exists. The enantiomers can be separated using known techniques, including those described in Pure and Applied Chemistry 69, 1469-1474, (1997) IUPAC.

In cases in which compounds have unsaturated carbon-carbon double bonds, both the cis (Z) and trans (E) isomers are within the scope of this invention.

5 The compounds of the subject invention may have spontaneous tautomeric forms. In cases wherein compounds may exist in tautomeric forms, such as keto-enol tautomers, each tautomeric form is contemplated as being included within this invention whether existing in equilibrium or
10 predominantly in one form.

10

In the compound structures depicted herein, hydrogen atoms are not shown for carbon atoms having less than four bonds to non-hydrogen atoms. However, it is understood that enough hydrogen atoms exist on said carbon atoms to satisfy the octet rule.

15

This invention also provides isotopic variants of the compounds disclosed herein, including wherein the isotopic atom is ^2H and/or wherein the isotopic atom ^{13}C . Accordingly, in the compounds provided herein hydrogen can be enriched in the deuterium isotope. It is to be understood that
20 the invention encompasses all such isotopic forms.

It is understood that the structures described in the embodiments of the methods hereinabove can be the same as the structures of the compounds described hereinabove.

25

It is understood that where a numerical range is recited herein, the present invention contemplates each integer between, and including, the upper and lower limits, unless otherwise stated.

30 Except where otherwise specified, if the structure of a compound of this invention includes an asymmetric carbon atom, it is understood that the compound occurs as a racemate, racemic mixture, and isolated single enantiomer. All such isomeric forms of these compounds are expressly included in this invention. Except where otherwise specified, each
35 stereogenic carbon may be of the R or S configuration. It is to be

understood accordingly that the isomers arising from such asymmetry (e.g., all enantiomers and diastereomers) are included within the scope of this invention, unless indicated otherwise. Such isomers can be obtained in substantially pure form by classical separation techniques and by stereochemically controlled synthesis, such as those described in "Enantiomers, Racemates and Resolutions" by J. Jacques, A. Collet and S. Wilen, Pub. John Wiley & Sons, NY, 1981. For example, the resolution may be carried out by preparative chromatography on a chiral column.

The subject invention is also intended to include all isotopes of atoms occurring on the compounds disclosed herein. Isotopes include those atoms having the same atomic number but different mass numbers. By way of general example and without limitation, isotopes of hydrogen include tritium and deuterium. Isotopes of carbon include C-13 and C-14.

It will be noted that any notation of a carbon in structures throughout this application, when used without further notation, are intended to represent all isotopes of carbon, such as ^{12}C , ^{13}C , or ^{14}C . Furthermore, any compounds containing ^{13}C or ^{14}C may specifically have the structure of any of the compounds disclosed herein.

It will also be noted that any notation of a hydrogen in structures throughout this application, when used without further notation, are intended to represent all isotopes of hydrogen, such as ^1H , ^2H , or ^3H . Furthermore, any compounds containing ^2H or ^3H may specifically have the structure of any of the compounds disclosed herein.

Isotopically-labeled compounds can generally be prepared by conventional techniques known to those skilled in the art using appropriate isotopically-labeled reagents in place of the non-labeled reagents employed.

In the compounds used in the method of the present invention, the substituents may be substituted or unsubstituted, unless specifically defined otherwise.

In the compounds used in the method of the present invention, alkyl, heteroalkyl, monocycle, bicycle, aryl, heteroaryl and heterocycle groups can be further substituted by replacing one or more hydrogen atoms with
5 alternative non-hydrogen groups. These include, but are not limited to, halo, hydroxy, mercapto, amino, carboxy, cyano, carbamoyl and aminocarbonyl and aminothiocarbonyl.

It is understood that substituents and substitution patterns on the
10 compounds used in the method of the present invention can be selected by one of ordinary skill in the art to provide compounds that are chemically stable and that can be readily synthesized by techniques known in the art from readily available starting materials. If a substituent is itself substituted with more than one group, it is understood that
15 these multiple groups may be on the same carbon or on different carbons, so long as a stable structure results.

In choosing the compounds used in the method of the present invention, one of ordinary skill in the art will recognize that the various
20 substituents, i.e. R_1 , R_2 , etc. are to be chosen in conformity with well-known principles of chemical structure connectivity.

As used herein, "alkyl" includes both branched and straight-chain saturated aliphatic hydrocarbon groups having the specified number of
25 carbon atoms and may be unsubstituted or substituted. Thus, C_1-C_n as in " C_1-C_n alkyl" is defined to include individual groups each having 1, 2, ..., $n-1$ or n carbons in a linear or branched arrangement. For example, C_1-C_6 , as in " C_1-C_6 alkyl" is defined to include individual groups each having 1, 2, 3, 4, 5, or 6 carbons in a linear or branched arrangement,
30 and specifically includes methyl, ethyl, n-propyl, isopropyl, n-butyl, t-butyl, pentyl, hexyl, and octyl.

As used herein, "alkenyl" refers to a non-aromatic hydrocarbon radical, straight or branched, containing at least 1 carbon to carbon double bond,
35 and up to the maximum possible number of non-aromatic carbon-carbon

double bonds may be present, and may be unsubstituted or substituted. For example, "C₂-C₆ alkenyl" means an alkenyl radical having 2, 3, 4, 5, or 6 carbon atoms, and up to 1, 2, 3, 4, or 5 carbon-carbon double bonds respectively. Alkenyl groups include ethenyl, propenyl, butenyl and cyclohexenyl.

The term "alkynyl" refers to a hydrocarbon radical straight or branched, containing at least 1 carbon to carbon triple bond, and up to the maximum possible number of non-aromatic carbon-carbon triple bonds may be present, and may be unsubstituted or substituted. Thus, "C₂-C₆ alkynyl" means an alkynyl radical having 2 or 3 carbon atoms and 1 carbon-carbon triple bond, or having 4 or 5 carbon atoms and up to 2 carbon-carbon triple bonds, or having 6 carbon atoms and up to 3 carbon-carbon triple bonds. Alkynyl groups include ethynyl, propynyl and butynyl.

"Alkylene", "alkenylene" and "alkynylene" shall mean, respectively, a divalent alkane, alkene and alkyne radical, respectively. It is understood that an alkylene, alkenylene, and alkynylene may be straight or branched. An alkylene, alkenylene, and alkynylene may be unsubstituted or substituted.

As used herein, "heteroalkyl" includes both branched and straight-chain saturated aliphatic hydrocarbon groups having the specified number of carbon atoms and at least 1 heteroatom within the chain or branch.

As used herein, "heterocycle" or "heterocyclyl" as used herein is intended to mean a 5- to 10-membered nonaromatic ring containing from 1 to 4 heteroatoms selected from the group consisting of O, N and S, and includes bicyclic groups. "Heterocyclyl" therefore includes, but is not limited to the following: imidazolyl, piperazinyl, piperidinyl, pyrrolidinyl, morpholinyl, thiomorpholinyl, tetrahydropyranyl, dihydropiperidinyl, tetrahydrothiophenyl and the like. If the heterocycle contains a nitrogen, it is understood that the corresponding N-oxides thereof are also encompassed by this definition.

As herein, "cycloalkyl" shall mean cyclic rings of alkanes of three to eight total carbon atoms, or any number within this range (i.e., cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl or cyclooctyl).

5

As used herein, "monocycle" includes any stable polyatomic carbon ring of up to 10 atoms and may be unsubstituted or substituted. Examples of such non-aromatic monocycle elements include but are not limited to: cyclobutyl, cyclopentyl, cyclohexyl, and cycloheptyl. Examples of such aromatic monocycle elements include but are not limited to: phenyl.

10

As used herein, "bicycle" includes any stable polyatomic carbon ring of up to 10 atoms that is fused to a polyatomic carbon ring of up to 10 atoms with each ring being independently unsubstituted or substituted. Examples of such non-aromatic bicycle elements include but are not limited to: decahydronaphthalene. Examples of such aromatic bicycle elements include but are not limited to: naphthalene.

15

As used herein, "aryl" is intended to mean any stable monocyclic, bicyclic or polycyclic carbon ring of up to 10 atoms in each ring, wherein at least one ring is aromatic, and may be unsubstituted or substituted. Examples of such aryl elements include phenyl, p-toluenyl (4-methylphenyl), naphthyl, tetrahydro-naphthyl, indanyl, biphenyl, phenanthryl, anthryl or acenaphthyl. In cases where the aryl substituent is bicyclic and one ring is non-aromatic, it is understood that attachment is via the aromatic ring.

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As used herein, the term "polycyclic" refers to unsaturated or partially unsaturated multiple fused ring structures, which may be unsubstituted or substituted.

30

The term "alkylryl" refers to alkyl groups as described above wherein one or more bonds to hydrogen contained therein are replaced by a bond to an aryl group as described above. It is understood that an "arylalkyl" group is connected to a core molecule through a bond from the alkyl group and that the aryl group acts as a substituent on the alkyl group. Examples

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of arylalkyl moieties include, but are not limited to, benzyl (phenylmethyl), p-trifluoromethylbenzyl (4-trifluoromethyl-phenylmethyl), 1-phenylethyl, 2-phenylethyl, 3-phenylpropyl, 2-phenylpropyl and the like.

5

The term "heteroaryl", as used herein, represents a stable monocyclic, bicyclic or polycyclic ring of up to 10 atoms in each ring, wherein at least one ring is aromatic and contains from 1 to 4 heteroatoms selected from the group consisting of O, N and S. Bicyclic aromatic heteroaryl groups include phenyl, pyridine, pyrimidine or pyridazine rings that are (a) fused to a 6-membered aromatic (unsaturated) heterocyclic ring having one nitrogen atom; (b) fused to a 5- or 6-membered aromatic (unsaturated) heterocyclic ring having two nitrogen atoms; (c) fused to a 5-membered aromatic (unsaturated) heterocyclic ring having one nitrogen atom together with either one oxygen or one sulfur atom; or (d) fused to a 5-membered aromatic (unsaturated) heterocyclic ring having one heteroatom selected from O, N or S. Heteroaryl groups within the scope of this definition include but are not limited to: benzoimidazolyl, benzofuranyl, benzofurazanyl, benzopyrazolyl, benzotriazolyl, benzothiophenyl, benzoxazolyl, carbazolyl, carbolinyl, cinnolinyl, furanyl, indolinyl, indolyl, indolazinyll, indazolyl, isobenzofuranyl, isoindolyl, isoquinolyl, isothiazolyl, isoxazolyl, naphthpyridinyl, oxadiazolyl, oxazolyl, oxazoline, isoxazoline, oxetanyl, pyranyl, pyrazinyl, pyrazolyl, pyridazinyl, pyridopyridinyl, pyridazinyl, pyridyl, pyrimidyl, pyrrolyl, quinazolinyl, quinolyl, quinoxalinyl, tetrazolyl, tetrazolopyridyl, thiadiazolyl, thiazolyl, thienyl, triazolyl, azetidinyll, aziridinyl, 1,4-dioxanyl, hexahydroazepinyl, dihydrobenzoimidazolyl, dihydrobenzofuranyl, dihydrobenzothiophenyl, dihydrobenzoxazolyl, dihydrofuranyl, dihydroimidazolyl, dihydroindolyl, dihydroisooxazolyl, dihydroisothiazolyl, dihydrooxadiazolyl, dihydrooxazolyl, dihydropyrazinyl, dihydropyrazolyl, dihydropyridinyl, dihydropyrimidinyl, dihydropyrrolyl, dihydroquinolinyl, dihydrotetrazolyl, dihydrothiadiazolyl, dihydrothiazolyl, dihydrothienyl, dihydrotriazolyl, dihydroazetidinyll, methylenedioxybenzoyl, tetrahydrofuranyl, tetrahydrothienyl, acridinyl,

carbazolyl, cinnolinyl, quinoxalinyl, pyrrazolyl, indolyl, benzotriazolyl, benzothiazolyl, benzoxazolyl, isoxazolyl, isothiazolyl, furanyl, thienyl, benzothienyl, benzofuranyl, quinolinyl, isoquinolinyl, oxazolyl, isoxazolyl, indolyl, pyrazinyl, pyridazinyl, pyridinyl, pyrimidinyl, pyrrolyl, tetra-hydroquinoline. In cases where the heteroaryl substituent is bicyclic and one ring is non-aromatic or contains no heteroatoms, it is understood that attachment is via the aromatic ring or via the heteroatom containing ring, respectively. If the heteroaryl contains nitrogen atoms, it is understood that the corresponding N-oxides thereof are also encompassed by this definition.

The term "alkylheteroaryl" refers to alkyl groups as described above wherein one or more bonds to hydrogen contained therein are replaced by a bond to an heteroaryl group as described above. It is understood that an "alkylheteroaryl" group is connected to a core molecule through a bond from the alkyl group and that the heteroaryl group acts as a substituent on the alkyl group. Examples of alkylheteroaryl moieties include, but are not limited to, $-\text{CH}_2-(\text{C}_5\text{H}_4\text{N})$, $-\text{CH}_2-\text{CH}_2-(\text{C}_5\text{H}_4\text{N})$ and the like.

The term "heterocycle" or "heterocyclyl" refers to a mono- or polycyclic ring system which can be saturated or contains one or more degrees of unsaturation and contains one or more heteroatoms. Preferred heteroatoms include N, O, and/or S, including N-oxides, sulfur oxides, and dioxides. Preferably the ring is three to ten-membered and is either saturated or has one or more degrees of unsaturation. The heterocycle may be unsubstituted or substituted, with multiple degrees of substitution being allowed. Such rings may be optionally fused to one or more of another "heterocyclic" ring(s), heteroaryl ring(s), aryl ring(s), or cycloalkyl ring(s). Examples of heterocycles include, but are not limited to, tetrahydrofuran, pyran, 1,4-dioxane, 1,3-dioxane, piperidine, piperazine, pyrrolidine, morpholine, thiomorpholine, tetrahydrothiopyran, tetrahydrothiophene, 1,3-oxathiolane, and the like.

The alkyl, alkenyl, alkynyl, aryl, heteroaryl and heterocyclyl substituents may be substituted or unsubstituted, unless specifically defined otherwise. In the compounds of the present invention, alkyl, alkenyl, alkynyl, aryl, heterocyclyl and heteroaryl groups can be further substituted by replacing one or more hydrogen atoms with alternative non-hydrogen groups. These include, but are not limited to, halo, hydroxy, mercapto, amino, carboxy, cyano and carbamoyl.

As used herein, the term "halogen" refers to F, Cl, Br, and I.

The terms "substitution", "substituted" and "substituent" refer to a functional group as described above in which one or more bonds to a hydrogen atom contained therein are replaced by a bond to non-hydrogen or non-carbon atoms, provided that normal valencies are maintained and that the substitution results in a stable compound. Substituted groups also include groups in which one or more bonds to a carbon(s) or hydrogen(s) atom are replaced by one or more bonds, including double or triple bonds, to a heteroatom. Examples of substituent groups include the functional groups described above, and halogens (i.e., F, Cl, Br, and I); alkyl groups, such as methyl, ethyl, n-propyl, isopropyl, n-butyl, tert-butyl, and trifluoromethyl; hydroxyl; alkoxy groups, such as methoxy, ethoxy, n-propoxy, and isopropoxy; aryloxy groups, such as phenoxy; arylalkyloxy, such as benzyloxy (phenylmethoxy) and p-trifluoromethylbenzyloxy (4-trifluoromethylphenylmethoxy); heteroaryloxy groups; sulfonyl groups, such as trifluoromethanesulfonyl, methanesulfonyl, and p-toluenesulfonyl; nitro, nitrosyl; mercapto; sulfanyl groups, such as methylsulfanyl, ethylsulfanyl and propylsulfanyl; cyano; amino groups, such as amino, methylamino, dimethylamino, ethylamino, and diethylamino; and carboxyl. Where multiple substituent moieties are disclosed or claimed, the substituted compound can be independently substituted by one or more of the disclosed or claimed substituent moieties, singly or pluraly. By independently substituted, it is meant that the (two or more) substituents can be the same or different.

The term "tolyl" refers to one of the three $\text{CH}_3\text{C}_6\text{H}_4$ - isomeric groups derived from toluene.

The term "naphthalene" refers to a bicyclic aromatic hydrocarbon consisting of a fused pair of benzene rings.

It is understood that substituents and substitution patterns on the compounds of the instant invention can be selected by one of ordinary skill in the art to provide compounds that are chemically stable and that can be readily synthesized by techniques known in the art, as well as those methods set forth below, from readily available starting materials. If a substituent is itself substituted with more than one group, it is understood that these multiple groups may be on the same carbon or on different carbons, so long as a stable structure results.

In choosing the compounds of the present invention, one of ordinary skill in the art will recognize that the various substituents, i.e. R_1 , R_2 , etc. are to be chosen in conformity with well-known principles of chemical structure connectivity. The various R groups attached to the aromatic rings of the compounds disclosed herein may be added to the rings by standard procedures, for example those set forth in Advanced Organic Chemistry: Part B: Reaction and Synthesis, Francis Carey and Richard Sundberg, (Springer) 5th ed. Edition. (2007), the content of which is hereby incorporated by reference.

The compounds used in the method of the present invention may be prepared by techniques well known in organic synthesis and familiar to a practitioner ordinarily skilled in the art. However, these may not be the only means by which to synthesize or obtain the desired compounds.

The compounds used in the method of the present invention may be prepared by techniques described in Vogel's Textbook of Practical Organic Chemistry, A.I. Vogel, A.R. Tatchell, B.S. Furnis, A.J. Hannaford, P.W.G. Smith, (Prentice Hall) 5th Edition (1996), March's Advanced Organic Chemistry: Reactions, Mechanisms, and Structure, Michael B. Smith, Jerry

March, (Wiley-Interscience) 5th Edition (2007), and references therein, which are incorporated by reference herein. However, these may not be the only means by which to synthesize or obtain the desired compounds.

5 Another aspect of the invention comprises a compound used in the method of the present invention as a pharmaceutical composition.

10 In some embodiments, a pharmaceutical composition comprising the compound of the present invention and a pharmaceutically acceptable carrier.

As used herein, the term "pharmaceutically active agent" means any substance or compound suitable for administration to a subject and furnishes biological activity or other direct effect in the treatment, cure, mitigation, diagnosis, or prevention of disease, or affects the structure or any function of the subject. Pharmaceutically active agents include, but are not limited to, substances and compounds described in the Physicians' Desk Reference (PDR Network, LLC; 64th edition; November 15, 2009) and "Approved Drug Products with Therapeutic Equivalence Evaluations" (U.S. Department Of Health And Human Services, 30th edition, 2010), which are hereby incorporated by reference. Pharmaceutically active agents which have pendant carboxylic acid groups may be modified in accordance with the present invention using standard esterification reactions and methods readily available and known to those having ordinary skill in the art of chemical synthesis. Where a pharmaceutically active agent does not possess a carboxylic acid group, the ordinarily skilled artisan will be able to design and incorporate a carboxylic acid group into the pharmaceutically active agent where esterification may subsequently be carried out so long as the modification does not interfere with the pharmaceutically active agent's biological activity or effect.

35 The compounds used in the method of the present invention may be in a salt form. As used herein, a "salt" is a salt of the instant compounds which has been modified by making acid or base salts of the compounds.

In the case of compounds used to treat an infection or disease caused by a pathogen, the salt is pharmaceutically acceptable. Examples of pharmaceutically acceptable salts include, but are not limited to, mineral or organic acid salts of basic residues such as amines; alkali or organic salts of acidic residues such as phenols. The salts can be made using an organic or inorganic acid. Such acid salts are chlorides, bromides, sulfates, nitrates, phosphates, sulfonates, formates, tartrates, maleates, malates, citrates, benzoates, salicylates, ascorbates, and the like. Phenolate salts are the alkaline earth metal salts, sodium, potassium or lithium. The term "pharmaceutically acceptable salt" in this respect, refers to the relatively non-toxic, inorganic and organic acid or base addition salts of compounds of the present invention. These salts can be prepared *in situ* during the final isolation and purification of the compounds of the invention, or by separately reacting a purified compound of the invention in its free base or free acid form with a suitable organic or inorganic acid or base, and isolating the salt thus formed. Representative salts include the hydrobromide, hydrochloride, sulfate, bisulfate, phosphate, nitrate, acetate, valerate, oleate, palmitate, stearate, laurate, benzoate, lactate, phosphate, tosylate, citrate, maleate, fumarate, succinate, tartrate, naphthylate, mesylate, glucoheptonate, lactobionate, and laurylsulphonate salts and the like. (See, e.g., Berge et al. (1977) "Pharmaceutical Salts", *J. Pharm. Sci.* 66:1-19).

The compounds of the present invention may also form salts with basic amino acids such as lysine, arginine, etc. and with basic sugars such as N-methylglucamine, 2-amino-2-deoxyglucose, etc. and any other physiologically non-toxic basic substance.

The compounds used in the method of the present invention may be administered in various forms, including those detailed herein. The treatment with the compound may be a component of a combination therapy or an adjunct therapy, i.e. the subject or patient in need of the drug is treated or given another drug for the disease in conjunction with one or more of the instant compounds. This combination therapy can be

sequential therapy where the patient is treated first with one drug and then the other or the two drugs are given simultaneously. These can be administered independently by the same route or by two or more different routes of administration depending on the dosage forms employed.

5

As used herein, a "pharmaceutically acceptable carrier" is a pharmaceutically acceptable solvent, suspending agent or vehicle, for delivering the instant compounds to the animal or human. The carrier may be liquid or solid and is selected with the planned manner of administration in mind. Liposomes are also a pharmaceutically acceptable carrier as are slow-release vehicles.

10

The dosage of the compounds administered in treatment will vary depending upon factors such as the pharmacodynamic characteristics of a specific chemotherapeutic agent and its mode and route of administration; the age, sex, metabolic rate, absorptive efficiency, health and weight of the recipient; the nature and extent of the symptoms; the kind of concurrent treatment being administered; the frequency of treatment with; and the desired therapeutic effect.

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A dosage unit of the compounds used in the method of the present invention may comprise a single compound or mixtures thereof with additional antitumor agents. The compounds can be administered in oral dosage forms as tablets, capsules, pills, powders, granules, elixirs, tinctures, suspensions, syrups, and emulsions. The compounds may also be administered in intravenous (bolus or infusion), intraperitoneal, subcutaneous, or intramuscular form, or introduced directly, e.g. by injection, topical application, or other methods, into or topically onto a site of disease or lesion, all using dosage forms well known to those of ordinary skill in the pharmaceutical arts.

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The compounds used in the method of the present invention can be administered in admixture with suitable pharmaceutical diluents, extenders, excipients, or in carriers such as the novel programmable sustained-release multi-compartmental nanospheres (collectively

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referred to herein as a pharmaceutically acceptable carrier) suitably selected with respect to the intended form of administration and as consistent with conventional pharmaceutical practices. The unit will be in a form suitable for oral, nasal, rectal, topical, intravenous or
5 direct injection or parenteral administration. The compounds can be administered alone or mixed with a pharmaceutically acceptable carrier. This carrier can be a solid or liquid, and the type of carrier is generally chosen based on the type of administration being used. The active agent can be co-administered in the form of a tablet or capsule,
10 liposome, as an agglomerated powder or in a liquid form. Examples of suitable solid carriers include lactose, sucrose, gelatin and agar. Capsule or tablets can be easily formulated and can be made easy to swallow or chew; other solid forms include granules, and bulk powders. Tablets may contain suitable binders, lubricants, diluents,
15 disintegrating agents, coloring agents, flavoring agents, flow-inducing agents, and melting agents. Examples of suitable liquid dosage forms include solutions or suspensions in water, pharmaceutically acceptable fats and oils, alcohols or other organic solvents, including esters, emulsions, syrups or elixirs, suspensions, solutions and/or suspensions
20 reconstituted from non-effervescent granules and effervescent preparations reconstituted from effervescent granules. Such liquid dosage forms may contain, for example, suitable solvents, preservatives, emulsifying agents, suspending agents, diluents, sweeteners, thickeners, and melting agents. Oral dosage forms optionally contain flavorants and
25 coloring agents. Parenteral and intravenous forms may also include minerals and other materials to make them compatible with the type of injection or delivery system chosen.

Techniques and compositions for making dosage forms useful in the present
30 invention are described in the following references: 7 Modern Pharmaceutics, Chapters 9 and 10 (Banker & Rhodes, Editors, 1979); Pharmaceutical Dosage Forms: Tablets (Lieberman et al., 1981); Ansel, Introduction to Pharmaceutical Dosage Forms 2nd Edition (1976); Remington's Pharmaceutical Sciences, 17th ed. (Mack Publishing Company,
35 Easton, Pa., 1985); Advances in Pharmaceutical Sciences (David

Ganderton, Trevor Jones, Eds., 1992); Advances in Pharmaceutical Sciences Vol. 7. (David Ganderton, Trevor Jones, James McGinity, Eds., 1995); Aqueous Polymeric Coatings for Pharmaceutical Dosage Forms (Drugs and the Pharmaceutical Sciences, Series 36 (James McGinity, Ed., 1989);
5 Pharmaceutical Particulate Carriers: Therapeutic Applications: Drugs and the Pharmaceutical Sciences, Vol 61 (Alain Rolland, Ed., 1993); Drug Delivery to the Gastrointestinal Tract (Ellis Horwood Books in the Biological Sciences. Series in Pharmaceutical Technology; J. G. Hardy, S. S. Davis, Clive G. Wilson, Eds.); Modern Pharmaceutics Drugs and the
10 Pharmaceutical Sciences, Vol 40 (Gilbert S. Banker, Christopher T. Rhodes, Eds.). All of the aforementioned publications are incorporated by reference herein.

Tablets may contain suitable binders, lubricants, disintegrating agents,
15 coloring agents, flavoring agents, flow-inducing agents, and melting agents. For instance, for oral administration in the dosage unit form of a tablet or capsule, the active drug component can be combined with an oral, non-toxic, pharmaceutically acceptable, inert carrier such as lactose, gelatin, agar, starch, sucrose, glucose, methyl cellulose,
20 magnesium stearate, dicalcium phosphate, calcium sulfate, mannitol, sorbitol and the like. Suitable binders include starch, gelatin, natural sugars such as glucose or beta-lactose, corn sweeteners, natural and synthetic gums such as acacia, tragacanth, or sodium alginate, carboxymethylcellulose, polyethylene glycol, waxes, and the like.
25 Lubricants used in these dosage forms include sodium oleate, sodium stearate, magnesium stearate, sodium benzoate, sodium acetate, sodium chloride, and the like. Disintegrators include, without limitation, starch, methyl cellulose, agar, bentonite, xanthan gum, and the like.

30 The compounds used in the method of the present invention may also be administered in the form of liposome delivery systems, such as small unilamellar vesicles, large unilamellar vesicles, and multilamellar vesicles. Liposomes can be formed from a variety of phospholipids such as lecithin, sphingomyelin, proteolipids, protein-encapsulated vesicles
35 or from cholesterol, stearylamine, or phosphatidylcholines. The

compounds may be administered as components of tissue-targeted emulsions.

5 The compounds used in the method of the present invention may also be coupled to soluble polymers as targetable drug carriers or as a prodrug. Such polymers include polyvinylpyrrolidone, pyran copolymer, polyhydroxylpropylmethacrylamide-phenol, polyhydroxyethylaspartamidephenol, or polyethyleneoxide-polylysine substituted with palmitoyl residues. Furthermore, the compounds may be coupled to a class of
10 biodegradable polymers useful in achieving controlled release of a drug, for example, polylactic acid, polyglycolic acid, copolymers of polylactic and polyglycolic acid, polyepsilon caprolactone, polyhydroxy butyric acid, polyorthoesters, polyacetals, polydihydropyrans, polycyanoacrylates, and crosslinked or amphipathic block copolymers of
15 hydrogels.

Gelatin capsules may contain the active ingredient compounds and powdered carriers, such as lactose, starch, cellulose derivatives, magnesium stearate, stearic acid, and the like. Similar diluents can be used to
20 make compressed tablets. Both tablets and capsules can be manufactured as immediate release products or as sustained release products to provide for continuous release of medication over a period of hours. Compressed tablets can be sugar-coated or film-coated to mask any unpleasant taste and protect the tablet from the atmosphere, or enteric coated for
25 selective disintegration in the gastrointestinal tract.

For oral administration in liquid dosage form, the oral drug components are combined with any oral, non-toxic, pharmaceutically acceptable inert carrier such as ethanol, glycerol, water, and the like. Examples of
30 suitable liquid dosage forms include solutions or suspensions in water, pharmaceutically acceptable fats and oils, alcohols or other organic solvents, including esters, emulsions, syrups or elixirs, suspensions, solutions and/or suspensions reconstituted from non-effervescent granules and effervescent preparations reconstituted from effervescent
35 granules. Such liquid dosage forms may contain, for example, suitable

solvents, preservatives, emulsifying agents, suspending agents, diluents, sweeteners, thickeners, and melting agents.

Liquid dosage forms for oral administration can contain coloring and
5 flavoring to increase patient acceptance. In general, water, asuitable
oil, saline, aqueous dextrose (glucose), and related sugar solutions and
glycols such as propylene glycol or polyethylene glycols are suitable
carriers for parenteral solutions. Solutions for parenteral
administration preferably contain a water soluble salt of the active
10 ingredient, suitable stabilizing agents, and if necessary, buffer
substances. Antioxidizing agents such as sodium bisulfite, sodium
sulfite, or ascorbic acid, either alone or combined, are suitable
stabilizing agents. Also used are citric acid and its salts and sodium
EDTA. In addition, parenteral solutions can contain preservatives, such
15 as benzalkonium chloride, methyl- or propyl-paraben, and chlorobutanol.
Suitable pharmaceutical carriers are described in Remington's
Pharmaceutical Sciences, Mack Publishing Company, a standard reference
text in this field.

20 The compounds used in the method of the present invention may also be
administered in intranasal form via use of suitable intranasal vehicles,
or via transdermal routes, using those forms of transdermal skin patches
well known to those of ordinary skill in that art. To be administered
in the form of a transdermal delivery system, the dosage administration
25 will generally be continuous rather than intermittent throughout the
dosage regimen.

Parenteral and intravenous forms may also include minerals and other
materials such as solutol and/or ethanol to make them compatible with
30 the type of injection or delivery system chosen.

The compounds and compositions of the present invention can be
administered in oral dosage forms as tablets, capsules, pills, powders,
granules, elixirs, tinctures, suspensions, syrups, and emulsions. The
35 compounds may also be administered in intravenous (bolus or infusion),

intraperitoneal, subcutaneous, or intramuscular form, or introduced directly, e.g. by topical administration, injection or other methods, to the afflicted area, such as a wound, including ulcers of the skin, all using dosage forms well known to those of ordinary skill in the pharmaceutical arts.

Specific examples of pharmaceutically acceptable carriers and excipients that may be used to formulate oral dosage forms of the present invention are described in U.S. Pat. No. 3,903,297 to Robert, issued Sept. 2, 1975.

Techniques and compositions for making dosage forms useful in the present invention are described in the following references: 7 Modern Pharmaceutics, Chapters 9 and 10 (Banker & Rhodes, Editors, 1979); Pharmaceutical Dosage Forms: Tablets (Lieberman et al., 1981); Ansel, Introduction to Pharmaceutical Dosage Forms 2nd Edition (1976); Remington's Pharmaceutical Sciences, 17th ed. (Mack Publishing Company, Easton, Pa., 1985); Advances in Pharmaceutical Sciences (David Ganderton, Trevor Jones, Eds., 1992); Advances in Pharmaceutical Sciences Vol 7. (David Ganderton, Trevor Jones, James McGinity, Eds., 1995); Aqueous Polymeric Coatings for Pharmaceutical Dosage Forms (Drugs and the Pharmaceutical Sciences, Series 36 (James McGinity, Ed., 1989); Pharmaceutical Particulate Carriers: Therapeutic Applications: Drugs and the Pharmaceutical Sciences, Vol 61 (Alain Rolland, Ed., 1993); Drug Delivery to the Gastrointestinal Tract (Ellis Horwood Books in the Biological Sciences. Series in Pharmaceutical Technology; J. G. Hardy, S. S. Davis, Clive G. Wilson, Eds.); Modern Pharmaceutics Drugs and the Pharmaceutical Sciences, Vol 40 (Gilbert S. Banker, Christopher T. Rhodes, Eds.)). All of the aforementioned publications are incorporated by reference herein.

The term "prodrug" as used herein refers to any compound that when administered to a biological system generates the compound of the invention, as a result of spontaneous chemical reaction(s), enzyme catalyzed chemical reaction(s), photolysis, and/or metabolic chemical reaction(s). A prodrug is thus a covalently modified analog or latent form of a compound of the invention.

The active ingredient can be administered orally in solid dosage forms, such as capsules, tablets, powders, and chewing gum; or in liquid dosage forms, such as elixirs, syrups, and suspensions, including, but not
5 limited to, mouthwash and toothpaste. It can also be administered parentally, in sterile liquid dosage forms.

Solid dosage forms, such as capsules and tablets, may be enteric-coated to prevent release of the active ingredient compounds before they reach
10 the small intestine. Materials that may be used as enteric coatings include, but are not limited to, sugars, fatty acids, proteinaceous substances such as gelatin, waxes, shellac, cellulose acetate phthalate (CAP), methyl acrylate-methacrylic acid copolymers, cellulose acetate succinate, hydroxy propyl methyl cellulose phthalate, hydroxy propyl
15 methyl cellulose acetate succinate (hypromellose acetate succinate), polyvinyl acetate phthalate (PVAP), and methyl methacrylate-methacrylic acid copolymers.

The compounds and compositions of the invention can be coated onto stents
20 for temporary or permanent implantation into the cardiovascular system of a subject.

Each embodiment disclosed herein is contemplated as being applicable to each of the other disclosed embodiments. Thus, all combinations of the
25 various elements described herein are within the scope of the invention.

This invention will be better understood by reference to the Experimental Details which follow, but those skilled in the art will readily appreciate that the specific experiments detailed are only illustrative
30 of the invention as described more fully in the claims which follow thereafter.

Experimental Details

Materials and Methods

Those having ordinary skill in the art of organic synthesis will appreciate that modifications to general procedures and synthetic routes contained in this application can be used to yield additional derivatives and structurally diverse compounds. Suitable organic transformations are described in March's Advanced Organic Chemistry: Reactions, Mechanisms, and Structure (Wiley-Interscience; 6th edition, 2007), the content of which is hereby incorporated by reference.

High throughput fluorescence displacement assay with NBD-stearate

FABP5 was purified and delipidated as described previously (Kaczocha, M. et al. 2012). FABP5 (30 µg), NBD-stearate (1 µM), and a competitor test compound were incubated in 30 mM Tris-HCl, 100mM NaCl buffer (pH 7.6). Competitors included arachidonic acid, BMS309403, 48 test compounds from ChemDiv library, Compound 26 and Compound 49. The initial assay was run with buffer (30mM Tris-HCl buffer), negative controls (buffer and NBD-stearate), positive controls (buffer, NBD-stearate, FABP5), and experimental wells with a variable test compound added (arachidonic acid or one of the 48 test compounds) at 10 µM. Test compounds that produced high inhibition and proved statistically significant were then added to the fluorescent assay at 10 µM and tested in triplicate to verify their success. The most effective test compound and BMS309403 were measured in increasing concentrations (0.01-50 µM), as were the Compound 26 and γ-truxillic acid 1-naphthyl ester, which were discovered following the test. The fluorescent assays were tested in the wells of Microtest 96-well Assay Plates, Optilux (BD Biosciences, Franklin Lakes, NJ) and loss of fluorescence intensity was measured with a FLUOstar OPTIMA spectrofluorometer set to excitation and emission wavelengths of 460 nm and 544 nm, respectively. For the most effective test compounds, IC₅₀ values were calculated with GraphPad Prism. GraphPad Prism was also used to determine the K_i of these select competitors from the equation $K_i = IC_{50} / (1 + ([NBD-stearate] / K_d))$. The K_d of NBD-stearate for FABP5 had been determined previously through incubating FABP5 with

increasing concentrations of NBD-stearate. One site binding analysis in GraphPad Prism indicated that the K_d of NBD-stearate for FABP5 was 0.16 μ M (Kaczocha, M. et al. 2012).

5 Patch-clamp electrophysiology in brain slices

DR neurons are visualized using an upright microscope (BX 51 WI, Olympus, Tokyo, Japan) equipped with a differential interference contrast and infrared imaging system. Somatic recordings from DR neurons are obtained with patch electrodes (3-5 $m\Omega$) back-filled with potassium gluconate based
 10 internal solution of the following composition: 120 mM potassium gluconate, 10 mM KCl, 10 mM Na_2 -phosphocreatine, 10 mM HEPES, 1 mM $MgCl_2$, 1 mM EGTA, 2 mM Na_2 -ATP, 0.25 mM Na-GTP, pH 7.3 (Adjusted with KOH; Osmolarity, 280 to 290 mOsmol/l). All the recordings were conducted in the presence of GABA_A receptor antagonist picrotoxin (100 μ M). Excitatory
 15 postsynaptic currents (EPSCs) are evoked with a single square stimulus (intensity, 1 to 10 V, duration, 100 to 200 μ s) delivered via a glass stimulating electrode. EPSCs are amplified with a Multiclamp 700B (Molecular Devices, Union City, CA, USA) and are acquired using pClamp 10 software (Molecular Devices).

20

Data Analysis

The amplitude of EPSCs is determined by measuring the average current during a 2-ms period at the peak of each EPSC and subtracted from the baseline current determined during a 5-ms time window before the
 25 stimulus. All EPSC amplitudes are normalized to the mean baseline amplitude recorded for at least 10 min before drug application. Results in the text and figures are presented as mean $\pm$ SEM. Statistical analysis is conducted using the Student's paired t-test.

30 AEA uptake

AEA uptake assays in wild-type and FABP5 knockdown HeLa cells are performed exactly as described (Kaczocha, M. et al. 2012).

FAAH enzyme assay

Enzyme assays measuring the hydrolysis of [¹⁴C]AEA in the presence of compound or the FAAH inhibitor are carried out in HeLa homogenates expressing rat FAAH as described (Kaczocha, M. et al. 2009).

5

PPAR transactivation

PPAR α and PPAR γ transactivation assays are performed in HeLa cells exactly as described (Kaczocha, M. et al. 2012). Briefly, cells are transfected with the PPAR reporter system, incubated with GW7647, 10 rosiglitazone, or compound for 6 hrs, followed by measurement of luciferase and β -galactosidase activity using a luminometer as described (Kaczocha, M. et al. 2012).

Animals

15 Male C57Bl6 mice (22-30g, Taconic Farms) are used for all experiments. The animals are group housed at room temperature and kept on a 12:12 hour light:dark cycle with *ad libitum* access to water and food. The animals are habituated to the experimental room for one week before testing. All experiments are approved by the Stony Brook University 20 Institutional Animal Care and Use Committee. The experimenter is blinded to the treatment conditions of each animal.

Carrageenan-induced paw edema and thermal hyperalgesia

Paw edema is induced by injecting 1% λ -carrageenan (20 μ l, in sterile 25 saline) into the plantar surface of the left hind paw and a control solution of saline into the right hind paw using a 27 gauge needle. Paw diameters are measured before carrageenan injection and 4 hours after injection using digital electronic calipers (Fisher) and expressed to the nearest $\pm$ 0.01 mm. The compound 20 mg/kg, i.p.) is dissolved in 30 ethanol:emulphor:saline (1:1:18), requiring sonication and gentle heating for solubilization, and is administered 45 min prior to injection of carrageenan. The cannabinoid receptor antagonists, rimonabant and SR144528 (3 mg/kg, i.p.), in ethanol:emulphor:saline (1:1:18), are injected 15 min before the FABP inhibitor. Edema is reported as the 35 change in paw diameter at 4 hr over the baseline. Changes in paw diameter

of saline-injected contralateral paws are negligible. Thermal hyperalgesia measures the latency to withdraw the paw from a focused beam of radiant heat applied to the plantar surface of the hind paw using a Hargreaves plantar apparatus (Ugo Basile) set at an intensity of 3.0.

5 For each mouse, the average latencies consisted of three trials spaced at least 5 minutes apart. The mice are habituated to the test chamber for 30 min. The cutoff time is set at 30 sec.

Formalin test

10 Mice are habituated to the observation chamber (Plexiglas box, 25 cm x 25 cm x 25 cm) for 30 min prior to formalin injection. The mice subsequently receive an injection of formalin (2.5% in saline, 20 μ l) into the plantar surface of the right hind paw using a 27 gauge needle. The animals are immediately placed back into the observation chamber and
15 nocifensive behavior (time spent licking or biting the paw) is recorded for 60 min. The formalin test consists of two phases with the first phase (0-5 min) reflecting nociceptor activation and the second phase (15-45 min) reflecting an inflammatory pain response.

20 Statistical Analyses

Behavioral data are presented as means $\pm$ S.E.M. for the vehicle and inhibitor-treated groups, each consisting of at least 6 animals. Statistical significance between vehicle and inhibitor groups are determined using unpaired t-tests or one-way ANOVA followed by Dunnett's
25 post hoc analysis. In all cases, differences of $p < 0.05$ are considered significant.

Example 1. Synthesis of α -Truxillic Acid Monoesters

30 **Analytical methods.**

NMR spectra were recorded on either a Bruker Ascend 700 spectrometer operating at 700 MHz for ^1H acquisitions and 175 MHz for ^{13}C acquisitions, a Bruker 500 Advance spectrometer operating at 500 MHz and 125 MHz for ^1H and ^{13}C acquisitions, respectively, a Bruker 400 Nanobay spectrometer

operating at 400 MHz, 100 MHz, and 376 MHz for ^1H , ^{13}C , and ^{19}F acquisitions, respectively. Chemical shifts were referenced to the residual proton solvent peaks (^1H : CDCl_3 , δ 7.26; $(\text{CD}_3)_2\text{SO}$, δ 2.50; CD_3OD , δ 3.31; CD_3CN , δ 1.94), solvent ^{13}C signals (CDCl_3 , δ 77.16; $(\text{CD}_3)_2\text{SO}$, δ 39.52; CD_3OD , δ 49.00). Signals are listed in ppm, and multiplicity identified as s = singlet, br = broad, d = doublet, t = triplet, q = quartet, m = multiplet; coupling constants in Hz; integration. High-resolution mass spectra were performed at Mass Spectrometry Services at the Univ. of Illinois at Urbana-Champaign and were obtained using Waters Q-TOF Ultima ESI mass spectrometer. Concentration under reduced pressure was performed by rotary evaporation at 25 - 30 °C at appropriate pressure.

Materials.

All air- and moisture-insensitive reactions were carried out under an ambient atmosphere, magnetically stirred, and monitored by thin layer chromatography (TLC) using Agela Technologies TLC plates pre-coated with 250 μm thickness silica gel 60 F254 plates and visualized by fluorescence quenching under UV light. Flash chromatography was performed on SiliaFlash® Silica Gel 40-63 μm 60Å particle size using a forced flow of eluent at 0.3 - 0.5 bar pressure. All air- and moisture-sensitive manipulations were performed using oven-dried glassware, including standard Schlenk and glovebox techniques under an atmosphere of nitrogen. Diethyl ether and THF were distilled from deep purple sodium benzophenone ketyl. Methylene chloride, chloroform and acetonitrile were dried over CaH_2 and distilled. Methylene chloride was degassed via three freeze-pump-thaw cycles. All other chemicals were used as received. All deuterated solvents were purchased from Cambridge Isotope Laboratories.

Chemical Synthesis

α -2,4-diphenylcyclobutene-1,3-dicarboxylic acid (1a)

E-cinnamic acid (1 g, 6.7 mmol) was placed in a pyrex dish and exposed to light at 350 nm and an intensity of 280 nW/cm² for 5 days with periodic shaking. This process was performed in the solid state and monitored by

¹H NMR. After completion of the photoreaction, the white solid was washed with diethylether (20 mL) to give α-truxillic acid.

In the same manner, other mono ester 1b to 1f were synthesized and characterized, starting from corresponding substituted (*E*)-cinnamic acid.

α-2,4-Bis(2-methoxyphenyl)cyclobutene-1,3-dicarboxylic acid (1b)

White solid; ¹H NMR (700 MHz, DMSO-d₆) δ 3.74 - 3.79 (m, 8 H), 4.42 (dd, *J* = 10.1, 7.6 Hz, 2 H), 7.20 - 7.24 (m, 2 H), 7.25 - 7.28 (m, 2 H), 11.90 (s, 2 H). ¹³C NMR (175 MHz, DMSO-d₆) δ 36.0, 44.6, 55.4, 110.5, 120.1, 127.1, 127.5, 127.9, 157.2, 173.4.

α-2,4-Bis(2-chlorophenyl)cyclobutene-1,3-dicarboxylic acid (1c)

White solid; ¹H NMR (700 MHz, DMSO-d₆) δ 3.89 (d, *J* = 6.3 Hz, 2 H), 4.63 (d, *J* = 6.3 Hz, 2 H), 7.08 (td, *J* = 7.6, 1.3 Hz, 2 H), 7.12 (t, *J* = 7.0 Hz, 2 H), 7.19 - 7.24 (m, 2 H), 7.33 (d, *J* = 7.6 Hz, 2 H), 12.58 (s, 2 H).

α-2,4-Bis(2,6-dichlorophenyl)cyclobutene-1,3-dicarboxylic acid (1d)

White solid; mp (in progress); ¹H NMR (700 MHz, dmsO-d₆) δ 3.90 (d, *J* = 6.3 Hz, 2 H), 4.64 (d, *J* = 6.3 Hz, 2 H), 7.06 - 7.06 (m, 4 H), 7.20 - 7.25 (m, 2 H), 7.33 (d, *J* = 7.6 Hz, 2 H), 12.59 (s, 2 H).

α-2,4-Bis(2-bromophenyl)cyclobutene-1,3-dicarboxylic acid (1e)

white solid; mp; ¹H NMR (700 MHz, dmsO-d₆) 3.86 (dd, *J* = 4.0, 2.4 Hz, 2H), 4.63 (d, *J* = 6.4 Hz, 2 H), 6.99 - 7.04 (m, 2 H), 7.16 (t, *J* = 7.5 Hz, 2 H), 7.33 (d, *J* = 7.7 Hz, 2 H), 7.40 (d, *J* = 7.7 Hz, 2 H), 12.58 (s, 2 H); ¹³C NMR (175 MHz, dmsO-d₆) δ 42.6, 43.9, 124.7, 127.1, 128.5, 129.1, 132.3, 137.8, 173.6; HRMS (ESI) *m/z*: calcd for C₁₈H₁₅Br₂O₄H⁺, 451.9259, found, 451.9261; (Δ = 0.44 ppm)

Benzyl α-2,4-diphenylcyclobutane-1,3-dicarboxylate (3a) and dibenzyl α-2,4-diphenylcyclobutane-1,3-dicarboxylate (5a).

To α -truxillic acid (**1a**, 200 mg, 0.66 mmol) suspended in thionyl chloride (3 mL) was added three drops of DMF and the mixture was heated to reflux for 3 h with stirring. The excess thionyl chloride and DMF were removed in *vacuo* to give the corresponding α -truxillic acid dichloride (**2a**) as light yellow solid, which was used directly in the subsequent reaction. To a solution of **2a** in THF (10 mL) were added benzyl alcohol (0.53 mmol, 0.8 eq) and pyridine (4.0 mmol), and the reaction mixture was stirred for 2 h. The reaction was quenched by adding distilled water (10 mL) with stirring for 30 min. The resulted solution was diluted with ethyl acetate (15 mL) and washed with aqueous copper sulfate (5 mL x 3) and water (5 mL x 3). The organic layer was collected, dried over MgSO_4 , and concentrated in *vacuo*. The crude mixture was purified by flash chromatography on silica gel (hexanes/AcOEt/AcOH = 79/20/1) as the eluent to give mono-ester **3a** and diester **5a**.

3a, white solid; 61% yield; ^1H NMR (300 MHz, acetone- d_6) δ 7.50 – 7.19 (m, 13H), 7.05 (s, 2H), 4.83 (d, J = 12.3 Hz, 1 H), 4.66 (d, J = 12.2 Hz, 1H), 4.55 – 4.41 (m, 2 H), 4.10 – 3.96 (m, 2H); ^{13}C NMR (101 MHz, acetone- d_6) δ 172.20, 171.56, 139.37, 139.32, 135.97, 128.32, 128.27, 128.21, 128.08, 127.85, 127.75, 127.64, 126.95, 126.81, 65.92, 46.69, 46.23, 41.76, 41.47.

5a, white solid; 13% yield; ^1H NMR (500 MHz, acetone- d_6) δ 3.99 – 4.17 (m, 2 H), 4.52 (dd, J = 10.38, 7.32 Hz, 2 H), 4.64 (d, J = 12.21 Hz, 2 H), 4.81 (d, J = 12.51 Hz, 2 H), 6.98 – 7.08 (m, 4 H), 7.23 – 7.30 (m, 7 H), 7.30 – 7.36 (m, 4 H), 7.36 – 7.45 (m, 4 H); ^{13}C NMR (126 MHz, acetone- d_6) δ 42.6, 47.5, 66.9, 128.0, 128.7, 128.8, 129.1, 129.2, 129.3, 136.9, 140.1, 172.3; HRMS (ESI) m/z calculated for $\text{C}_{32}\text{H}_{28}\text{O}_4\text{H}^+$: 477.206, found 477.2059 (Δ = 0.34 ppm)

In the same manner, other mono ester **3b** to **3r** and di-esters **5b** to **5e** were synthesized and characterized, starting from α -2,4-diphenylcyclobutene-1,3-dicarboxylic acid (**1a**).

4-Methoxybenzyl α -2,4-diphenylcyclobutane-1,3-dicarboxylate (3b) and di(4-methoxy)benzyl α -2,4-diphenylcyclobutane-1,3-dicarboxylate (5b).

5 **3b**, white solid; 40% yield; mp 99-102 °C; ^1H NMR (500 MHz, acetone- d_6) δ 3.78 (s, 3 H), 3.93 - 4.07 (m, 2 H), 4.40 - 4.50 (m, 2 H), 4.58 (d, J = 11.9 Hz, 1 H), 4.74 (d, J = 11.9 Hz, 1 H), 6.79 - 6.87 (m, 2 H), 6.93 - 7.03 (m, 2 H), 7.21 - 7.39 (m, 8 H), 7.41 (d, J = 7.32 Hz, 2 H); ^{13}C NMR (126 MHz, acetone- d_6) δ 42.4, 42.7, 47.1, 47.6, 55.6, 66.7, 114.6, 127.7,
10 127.8, 128.6, 128.7, 129.1, 129.2, 130.9, 140.3, 160.6, 172.5, 173.1; HRMS (ESI) m/z calcd for $\text{C}_{26}\text{H}_{24}\text{O}_5\text{H}^+$: 434.1962, found 434.1964 (Δ = 0.42 ppm).

5b, white solid; 12% yield; mp 145-148 °C; ^1H NMR (500 MHz, acetone- d_6)
15 δ 3.78 (s, 6 H), 3.97 - 4.08 (m, 2 H), 4.47 (dd, J = 10.38, 7.32 Hz, 2 H), 4.56 (d, J = 12.2 Hz, 2 H), 4.73 (d, J = 12.2 Hz, 2 H), 6.78 - 6.86 (m, 4 H), 6.93 - 7.01 (m, 4 H), 7.23 - 7.30 (m, 2 H), 7.30 - 7.40 (m, 8 H); ^{13}C NMR (126 MHz, acetone- d_6) δ 42.5, 47.5, 55.6, 66.7, 114.6, 127.9, 128.6, 129.3, 130.9, 140.1, 160.6, 172.3; HRMS (ESI) m/z calcd for
20 $\text{C}_{26}\text{H}_{24}\text{O}_5\text{H}^+$: 554.2536, found 544.2537 (Δ = 0.2 ppm)

4-Fluorobenzyl α -2,4-diphenylcyclobutane-1,3-dicarboxylate (3c) and di(4-fluorobenzyl) α -2,4-diphenylcyclobutane-1,3-dicarboxylate (5c).

25 **3c**: white solid; 34% yield; mp 145-148 °C; ^1H NMR (500 MHz, acetone- d_6) δ 4.03 (dd, J = 10.38, 7.32 Hz, 2 H), 4.35 - 4.58 (m, 2 H), 4.68 (d, J = 12.21 Hz, 1 H), 4.80 (d, J = 12.21 Hz, 1 H), 6.93 - 7.14 (m, 4 H), 7.21 - 7.45 (m, 10 H), 10.63 (br. s., 1 H); ^{13}C NMR (126 MHz, acetone- d_6) δ 42.4, 42.7, 47.2, 47.6, 66.1, 115.8, 116.0, 127.8, 127.9, 128.6,
30 128.7, 129.2, 129.3, 131.2, 131.3, 133.1, 140.2, 140.3, 162.4, 164.3, 172.4, 173.0; HRMS (ESI) m/z calculated for $\text{C}_{25}\text{H}_{21}\text{FO}_4$ ($\text{M}+\text{H}$) $^+$: 405.1497, found 405.1502 (Δ = 1.3 ppm).

5c: White solid; 15% yield; mp 103-105 °C; ¹H NMR (500 MHz, acetone-d₆) δ 4.06 (dd, *J* = 10.53, 7.48 Hz, 2 H), 4.51 (dd, *J* = 10.38, 7.32 Hz, 2 H), 4.66 (d, *J* = 12.36 Hz, 2 H), 4.78 (d, *J* = 12.36 Hz, 2 H), 6.93 - 7.17 (m, 8 H), 7.20 - 7.44 (m, 10 H); ¹³C NMR (126 MHz, acetone-d₆) δ 42.5, 47.4, 66.1, 115.8, 116.0, 128.0, 128.6, 129.3, 131.3, 131.3, 133.1, 133.1, 140.0, 162.4, 164.3, 172.2; HRMS (ESI) *m/z* calcd for C₃₂H₂₆F₂O₄H⁺: 513.1872, found 513.1882 (Δ = 2.0 ppm).

4-Bromobenzyl α-2,4-diphenylcyclobutane-1,3-dicarboxylate (3d).

White solid; 44% yield; mp 175-177 °C; ¹H NMR (500 MHz, acetone-d₆) δ 3.92 - 4.11 (m, 2 H), 4.40 - 4.57 (m, 2 H), 4.69 (d, *J* = 12.66 Hz, 1 H), 4.78 (d, *J* = 12.66 Hz, 1 H), 6.96 (d, *J* = 8.54 Hz, 2 H), 7.20-7.29 (m, 2 H), 7.32 (t, *J* = 7.48 Hz, 4 H), 7.35 - 7.40 (m, 2 H), 7.40-7.49 (m, 4 H); ¹³C NMR (126 MHz, acetone-d₆) δ 42.4, 42.7, 47.2, 47.5, 66.0, 122.3, 127.8, 127.9, 128.6, 128.7, 129.2, 129.3, 131.0, 132.3, 136.4, 140.2, 140.3, 172.4, 173.1; HRMS (ESI) *m/z* calcd for C₂₅H₂₁BrO₄H⁺: 465.0696, found 465.0697 (Δ = 0.27 ppm).

Di-tetrahydropyran-4-methyl α-2,4-diphenylcyclobutane-1,3-dicarboxylate (5d).

White solid; 21% yield; ¹H NMR (500 MHz, acetone-d₆) δ 0.91 - 1.10 (m, 4 H), 1.16 - 1.31 (m, 4 H), 1.38 - 1.55 (m, 2 H), 3.17 (tt, *J* = 11.71, 2.33 Hz, 4 H), 3.58 (d, *J* = 6.41 Hz, 4 H), 3.67 - 3.81 (m, 4 H), 3.90 - 4.08 (m, 2 H), 4.48 (dd, *J* = 10.38, 7.32 Hz, 2 H), 7.19 - 7.31 (m, 2 H), 7.31 - 7.46 (m, 8 H); ¹³C NMR (126 MHz, acetone-d₆) δ 35.2, 42.5, 47.7, 67.8, 67.8, 69.4, 127.9, 128.7, 129.3, 140.3, 172.3; HRMS (ESI) *m/z* calculated for C₃₀H₃₆O₆H⁺: 493.2585, found 493.2591 (Δ = 1.24 ppm).

Biphenyl -2-yl α-2,4-diphenylcyclobutane-1,3-dicarboxylate (3g).

White solid; 21% yield; mp 195-196 °C; ¹H NMR (500 MHz, acetone-d₆) δ 3.81 (dt, *J* = 11.67, 10.49 Hz, 2 H), 4.39 (t, *J* = 10.07 Hz, 1 H), 4.68 (t, *J* = 10.68 Hz, 1 H), 5.99 (dd, *J* = 8.09, 1.07 Hz, 1 H), 7.11 - 7.19 (m, 3 H), 7.20 - 7.28 (m, 2 H), 7.28 - 7.40 (m, 11 H), 7.40 - 7.46 (m,

2 H); ^{13}C NMR (126 MHz, acetone- d_6) δ 42.3, 45.4, 46.8, 47.0, 123.4, 127.1, 127.4, 128.1, 128.4, 129.2, 129.3, 129.6, 129.8, 131.3, 135.8, 138.5, 139.2, 143.2, 148.7, 170.6, 172.9; HRMS (ESI) m/z calcd for $\text{C}_{30}\text{H}_{24}\text{O}_4\text{H}^+$: 449.1747, found 449.1754 (Δ = 1.55 ppm).

5

Biphenyl-3-yl α -2,4-Diphenylcyclobutane-1,3-dicarboxylate (3h)

white solid; ^1H NMR (500 MHz, acetone- d_6) δ 4.14 (dd, J = 10.8, 7.2 Hz, 1 H), 4.29 – 4.35 (m, 1 H), 4.61 (dd, J = 10.8, 7.2 Hz, 1 H), 4.67 (dd, J = 10.8, 7.2 Hz, 1 H), 6.53 (t, J = 1.9 Hz, 1 H), 6.55 – 6.57 (m, 1 H),
10 7.29 (d, J = 7.4 Hz, 1 H), 7.33 (d, J = 7.9 Hz, 1 H), 7.36 – 7.41 (m, 4 H), 7.44 – 7.55 (m, 9 H), 7.59 (d, J = 7.5 Hz, 2 H).

Di-biphenyl-3-yl α -2,4-diphenylcyclobutane-1,3-dicarboxylate (5e).

White solid; ^1H NMR (500 MHz, acetone- d_6) δ 4.45 (dd, J = 10.8, 7.3 Hz, 2 H), 4.81 (dd, J = 10.6, 7.4 Hz, 2 H), 6.53 (m, 2 H), 6.54 – 6.58 (m, 2 H),
15 7.34 (t, J = 7.6 Hz, 2 H), 7.39 (m, 4 H), 7.43 – 7.55 (m, 14 H), 7.65 (d, J = 7.6 Hz, 4 H).

2'-HO-biphenyl-2-yl α -2,4-diphenylcyclobutane-1,3-dicarboxylate (3j)

20 2'-TIPSO-biphenyl-2-yl α -2,4-diphenylcyclobutane-1,3-dicarboxylate (3j-i) was made in the same manner of 3a. The crude product was purified by column chromatography on silica gel using ethyl acetate/hexanes (30% to 50%) as eluent to afford 3j-i as white solid (59% yield). The TIPS protecting group of 3j-i was removed by TBAF in THF to give desired
25 product 3j as a white solid (168 mg, 99%). Melting point: 55.0–57.0 °C. ^1H NMR (500 MHz, Acetone) δ 10.67 (s, 1H), 8.21 (s, 1H), 7.48 (d, J = 7.4 Hz, 2H), 7.43 (t, J = 7.5 Hz, 2H), 7.26 (ddddd, J = 17.1, 15.4, 9.2, 6.6, 2.8 Hz, 12H), 7.11 (dd, J = 7.5, 1.5 Hz, 1H), 7.03 (d, J = 8.1 Hz, 1H), 6.93 – 6.87 (m, 1H), 6.15 (dd, J = 8.0, 1.0 Hz, 1H), 4.51 (dd, J =
30 10.5, 6.8 Hz, 1H), 4.37 (dd, J = 10.4, 7.3 Hz, 1H), 4.11 (dd, J = 10.5, 6.9 Hz, 1H), 3.99 (dd, J = 10.5, 7.3 Hz, 1H). HRMS (ESI) m/z : calcd for $\text{C}_{30}\text{H}_{25}\text{O}_5\text{H}^+$, 465.1697, found, 465.1699 (Δ = 0.43 ppm).

2,4,5-Trichlorobenzyl α -2,4-diphenylcyclobutane-1,3-dicarboxylate (3k).

White solid; 70% yield; mp 191-193 °C; ¹H NMR (700 MHz, acetone-d₆) δ 4.14 (dd, *J* = 10.7, 7.3 Hz, 1 H), 4.38 (dd, *J* = 10.7, 7.1 Hz, 1 H), 4.60 (dd, *J* = 10.7, 7.1 Hz, 1 H), 4.68 (dd, *J* = 10.7, 7.3 Hz, 1 H), 6.03 (s, 1 H), 7.27 (t, *J* = 7.4 Hz, 1 H), 7.36 (t, *J* = 7.6 Hz, 2 H), 7.41 (t, *J* = 7.3 Hz, 1 H), 7.48 (dt, *J* = 15.1, 9.8 Hz, 4 H), 7.57 (d, *J* = 7.5 Hz, 2 H), 7.70 (d, *J* = 7.6 Hz, 1 H), 10.75 (s, 1 H); ¹³C NMR (175 MHz, acetone-d₆) δ 171.9, 169.2, 145.9, 139.1, 138.8, 130.9, 130.7, 129.8, 129.0, 128.8, 127.8, 127.5, 127.0, 126.3, 125.0, 46.5, 45.9, 42.0, 41.2; HRMS (ESI) *m/z* calcd for C₂₄H₁₈Cl₃O₄H⁺: 475.0265; found, 475.0264 (Δ = 0.21 ppm).

(1*R*,2*S*)-2-phenylcyclohexyl α-2,4-diphenylcyclobutane-1,3-dicarboxylate (31).

White solid; 61% yield; mp 192-193 °C; ¹H NMR (700 MHz, chloroform-d) δ 0.80 (qd, *J* = 12.8, 3.7 Hz, 1 H), 1.22 - 1.39 (m, 2 H), 1.47 - 1.59 (m, 2 H), 1.64 - 1.76 (m, 2 H), 1.90 (d, *J* = 13.4 Hz, 1 H), 2.57 (td, *J* = 12.3, 3.6 Hz, 1 H), 3.65 (ddd, *J* = 19.6, 9.9, 6.1 Hz, 2 H), 3.96 (dd, *J* = 10.0, 8.0 Hz, 1 H), 4.24 - 4.32 (m, 1 H), 4.69 (td, *J* = 10.8, 4.2 Hz, 1 H), 7.00 (d, *J* = 6.9 Hz, 2 H), 7.07 (d, *J* = 7.6 Hz, 2 H), 7.15 (d, *J* = 7.1 Hz, 2 H), 7.20 - 7.29 (m, 7 H), 7.31 (dd, *J* = 15.5, 8.0 Hz, 2 H), 10.64 (s, 1H). ¹³C NMR (175 MHz, chloroform-d) δ 24.7, 25.8, 31.4, 33.7, 40.6, 42.3, 46.1, 47.3, 49.9, 76.9, 126.7, 126.8, 127.1, 127.3, 127.6, 128.5, 128.6, 138.6, 139.2, 143.2, 171.0, 177.7; HRMS (ESI) *m/z*: calcd for C₃₀H₃₁O₄H⁺: 455.2217; found, 455.2221 (Δ 0.81 ppm).

Indan-2-yl α-2,4-diphenylcyclobutane-1,3-dicarboxylate (3m).

White solid; 34% yield; ¹H NMR (500 MHz, acetone-d₆) δ 2.05 (dt, *J* = 4.35, 2.25 Hz, 1 H), 2.10 (dd, *J* = 17.09, 2.44 Hz, 1 H), 2.74 (dd, *J* = 16.94, 2.59 Hz, 1 H), 2.87 (dd, *J* = 16.94, 6.26 Hz, 1 H), 3.11 (dd, *J* = 17.09, 6.41 Hz, 1 H), 3.90 (dd, *J* = 10.68, 6.71 Hz, 1 H), 3.97 - 4.07 (m, 1 H), 4.36 - 4.48 (m, 2 H), 5.14 - 5.21 (m, 1 H), 7.05 - 7.18 (m, 4 H), 7.19 - 7.25 (m, 1 H), 7.27 - 7.37 (m, 7 H), 7.40 (d, *J* = 7.32 Hz, 2 H); ¹³C NMR (126 MHz, acetone-d₆) δ 39.8, 40.0, 42.3, 42.6, 46.9, 47.5, 76.0, 125.3, 125.5, 127.3, 127.4, 127.7, 127.9, 128.7, 128.7, 129.1,

129.2, 140.1, 140.4, 141.4, 173.1; HRMS (ESI) m/z calcd for $C_{27}H_{24}O_4H^+$: 413.1747, found 413.1749 (Δ 0.43 ppm).

Quinolin-8-yl α -2,4-diphenylcyclobutane-1,3-dicarboxylate (3n).

5 White solid; 39% yield; ^1H NMR (700 MHz, acetone- d_6) δ 4.06 (t, J = 10.4 Hz, 1 H), 4.38 (t, J = 10.4 Hz, 1 H), 4.64 (t, J = 10.4 Hz, 1 H), 4.92 (t, J = 10.4 Hz, 1 H), 6.41 (d, J = 7.5 Hz, 1 H), 7.37 (t, J = 7.4 Hz, 1 H), 7.52 – 7.43 (m, 7 H), 7.60 (dd, J = 8.3, 4.2 Hz, 1H), 7.65 (dd, J = 7.3, 4.3 Hz, 4 H), 7.86 (d, J = 8.2 Hz, 1 H), 8.41 (dd, J = 8.2, 1.5 Hz, 1H), 8.94 (dd, J = 4.3, 1.5 Hz, 1 H), 10.75 (s, 1 H).

6-Acetamidonaphth-1-yl α -2,4-diphenylcyclobutane-1,3-dicarboxylate (3o)

^1H NMR (500 MHz, acetone- d_6) δ 2.13 (s, 3 H), 4.15 (dd, J = 10.1, 7.4 Hz, 1H), 4.54 (dd, J = 10.7, 7.2 Hz, 1 H), 4.68 (dd, J = 10.7, 7.2 Hz, 1 H), 15 6.33 (d, J = 7.5 Hz, 1 H), 7.14 (d, J = 9.0 Hz, 1 H), 7.28 (dd, J = 14.9, 7.3 Hz, 2 H), 7.36 – 7.44 (m, 4 H), 7.47 (t, J = 7.4 Hz, 2 H), 7.55 (d, J = 7.4 Hz, 2 H), 7.62 (t, J = 9.0 Hz, 3 H), 8.37 (s, 1 H), 9.34 (s, 1 H).

20 **5-Ethynylnaphth-1-yl α -2,4-diphenylcyclobutane-1,3-dicarboxylate (3p).**

White solid; 44% yield; mp 220–221 °C; ^1H NMR (500 MHz, acetone- d_6) δ 4.06 (s, 1H), 4.11 (dd, J = 10.7, 7.2 Hz, 1 H), 4.53 (dd, J = 10.7, 7.2 Hz, 1 H), 4.64 (dd, J = 10.7, 7.2 Hz, 1 H), 4.68 (dd, J = 10.7, 7.2 Hz, 1 H), 6.48 (d, J = 7.5 Hz, 1 H), 7.26 (m, 3 H), 7.34 (t, J = 7.6 Hz, 2 H), 25 H), 7.42 (m, 4 H), 7.51 (d, J = 7.5 Hz, 2 H), 7.60 (d, J = 7.2 Hz, 2 H), 7.67 (d, J = 6.8 Hz, 1 H), 8.11 (d, J = 8.4 Hz, 1 H), 10.70 (s, 1 H). ^{13}C NMR (176 MHz, acetone- d_6) δ 41.61, 42.11, 46.41, 46.77, 80.96, 83.68, 118.80, 119.77, 122.72, 123.45, 125.70, 126.49, 126.72, 126.92, 127.46, 127.85, 128.26, 128.27, 128.80, 131.41, 134.15, 139.18, 139.47, 147.18, 30 170.69, 172.12.

9-Fluorenylmethyl α -2,4-diphenylcyclobutane-1,3-dicarboxylate (3q).

White solid; 35% yield; mp 180–181 °C; ^1H NMR (700 MHz, chloroform- d) δ 3.85 (t, J = 6.7 Hz, 1 H), 4.00 (ddd, J = 18.4, 10.8, 7.4 Hz, 2 H), 4.08

(d, $J = 6.6$ Hz, 2 H), 4.37 (dd, $J = 10.8, 6.6$ Hz, 1 H), 4.47 (dd, $J = 10.8, 7.4$ Hz, 1 H), 7.25–7.39 (m, 12 H), 7.40–7.48 (m, 4 H), 7.80 (d, $J = 7.5$ Hz, 2 H); ^{13}C NMR (700 MHz, chloroform- d) δ 41.6, 41.8, 46.3, 46.7, 47.2, 66.6, 120.1, 125.0, 125.1, 127.2, 127.4, 127.5, 128.6, 128.7, 138.3, 138.4, 141.3, 141.4, 143.6, 144.1, 171.9, 177.2. HRMS (ESI) m/z : calcd for $\text{C}_{32}\text{H}_{27}\text{O}_4\text{H}^+$: 475.1904; found, 475.1907 ($\Delta = 0.63$ ppm).

2,2,2-Trifluoroethyl α -2,4-diphenylcyclobutane-1,3-dicarboxylate (3r)

white solid, 51% yield; ^1H NMR (300 MHz, acetone- d_6) δ 7.55 – 7.16 (m, 10H), 4.64 – 4.29 (m, 3H), 4.22 – 3.98 (m, 3H); ^{13}C NMR (101 MHz, Acetone) δ 172.00, 170.31, 138.93, 138.72, 128.35, 128.24, 127.69, 127.46, 127.02, 126.94, 124.64, 121.89, 60.44, 60.08, 59.71, 59.36, 46.24, 46.00, 41.71, 41.36.

Cyclohexyl α -2,4-diphenylcyclobutane-1,3-dicarboxylate (3s)

White solid; 68% yield; ^1H NMR (300 MHz, acetone- d_6): δ 0.88 (m, 1H), 1.22 (m, 5H), 1.42 (bs, 2H), 1.60 (m, 2H), 3.90–4.05 (m, 2H), 4.45 (m, 3H), 7.24–7.44 (m, 10 H); ^{13}C NMR (100 MHz, acetone- d_6) δ 23.207, 23.262, 25.081, 30.756, 31.188, 41.501, 41.660, 46.174, 46.789, 72.035, 126.753, 126.776, 127.756, 128.161, 128.184, 139.442, 139.573, 170.866, 172.183; HRMS (ESI) m/z calcd for $\text{C}_{24}\text{H}_{27}\text{O}_4\text{H}^+$: 379.1909, found 379.1909 (Δ 0 ppm).

3-[1-(3,6,9-trioxa-dodecanyl)-1,2,3-triazol-4-yl]phenyl α -2,4-diphenylcyclobutane-1,3-dicarboxylate (3t).

To a solution of **3f** in THF (8 mL) and water (1.5 mL), was added cupric sulfate pentahydrate, ascorbic acid, and 1-azido-2-(2-(2-ethoxyethoxy)ethoxy)ethane and the reaction mixture was stirred at room temperature overnight. Upon completion, the reaction mixture was diluted with water and extracted thrice with DCM. The crude mixture was purified with flash column on silica gel with 3.5% methanol in DCM as the eluent to give **3t** (180 mg, 94% yield) as white solid. mp 93–95 °C; ^1H NMR (500 MHz, acetone- d_6) δ 1.07 (t, $J = 7.02$ Hz, 6 H), 3.39 (q, $J = 7.02$ Hz, 4 H), 3.43 – 3.49 (m, 4 H), 3.49 – 3.55 (m, 4 H), 3.55 – 3.61 (m, 4 H), 3.61 – 3.71 (m, 4 H), 3.93 (t, $J = 5.19$ Hz, 4 H), 4.06 – 4.19 (m, 2 H), 4.27 – 4.36 (m, 2 H), 4.54

- 4.72 (m, 8 H), 6.28 - 6.43 (m, 2 H), 6.96 (t, $J = 1.83$ Hz, 2 H), 7.21 - 7.32 (m, 4 H), 7.33 - 7.44 (m, 6 H), 7.44 - 7.54 (m, 8 H), 7.57 (d, $J = 7.32$ Hz, 4 H), 7.69 (d, $J = 7.93$ Hz, 2 H), 8.22 (s, 2 H); ^{13}C NMR (126 MHz, acetone- d_6) δ 15.6, 42.3, 42.9, 47.0, 47.6, 51.0, 66.8, 70.1, 70.6, 71.2, 71.3, 119.5, 121.7, 122.4, 123.4, 127.8, 128.3, 128.7, 129.0, 129.2, 129.5, 130.4, 133.7, 140.1, 140.2, 146.8, 152.1, 171.3, 173.1; HRMS (ESI) m/z calculated for $\text{C}_{34}\text{H}_{37}\text{N}_3\text{O}_7\text{H}^+$: 600.2704, found 600.2705 ($\Delta = 0.13$ ppm).

10 **1-Naphthyl α -2,4-di (3-methoxy-4-triethylsiloxy) cyclobutane-1,3-dicarboxylate (4a-i).**

White solid; 26% yield; ^1H NMR (300 MHz, CDCl_3) δ ppm 0.55 - 0.77 (m, 13 H) 0.82 - 1.05 (m, 18 H) 3.75 (s, 3 H) 3.70 (s, 3 H) 3.96 (dd, $J=10.34$, 7.36 Hz, 1 H) 4.26 (dd, $J=10.24$, 7.26 Hz, 1 H) 4.37 - 4.61 (m, 2 H) 6.40 (d, $J=7.45$ Hz, 1 H) 6.63 - 6.76 (m, 1 H) 6.76 - 6.95 (m, 5 H) 7.14 - 7.46 (m, 4 H) 7.59 (d, $J=8.01$ Hz, 1 H) 7.73 (d, $J=8.01$ Hz, 1 H).

1-Naphthyl α -2,4-di (3-methoxy-4-hydroxy) cyclobutane-1,3-dicarboxylate (4a).

20 To a solution of **4a-i** (56 mg, 0.08 mmol), in 20:1 tetrahydrofuran:acetic acid (2.0 mL total), was added 1.0 M TBAF in THF (0.27 mmol) and the reaction mixture was stirred at room temperature for two hours. After completion, the reaction mixture was diluted with ethyl acetate (30 mL), washed thrice with water (15 mL), and dried over anhydrous
25 magnesium sulfate. The crude mixture was purified using flash column chromatography on silica gel with 3% methanol in dichloromethane as the eluent to give **4a** as white solid in 83% yield; ^1H NMR (400 MHz, acetone- d_6) δ ppm 3.79 (s, 3 H) 3.89 (s, 3 H) 4.02 (dd, $J=10.29$, 7.53 Hz, 1 H) 4.43 (dd, $J=10.29$, 7.53 Hz, 1 H) 4.51 - 4.67 (m, 2 H) 6.66 (d, $J=7.53$ Hz, 1 H) 6.82 (d, $J=8.03$ Hz, 1 H) 6.88 - 6.99 (m, 2 H) 7.04 (br. s., 1 H) 7.07 - 7.15 (m, 2 H) 7.20 (s, 1 H) 7.34 (d, $J=7.78$ Hz, 2 H) 7.46 (s, 1 H) 7.72 (d, $J=8.03$ Hz, 1 H) 7.85 (s, 1 H); HRMS (ESI) m/z calculated for $\text{C}_{30}\text{H}_{26}\text{O}_8\text{H}^+$: 515.1700, found 515.1743 ($\Delta = -8.34$ ppm).

1-Naphthyl α -2,4-di(2-methoxyphenyl)cyclobutane-1,3-dicarboxylate (4b).

White solid; 35% yield; mp 183.5–185.0 °C. ^1H NMR (500 MHz, acetone- d_6) δ 3.90 (s, 1 H), 4.16 – 4.21 (dd, J = 8.00, 10.50 Hz, 1 H), 4.46 – 4.49 (dd, J = 8.00, 10.50 Hz, 1 H), 4.85 – 4.88 (dd, J = 8.00, 10.50 Hz, 1 H), 4.85 – 4.88 (dd, J = 8.00, 10.50 Hz, 1 H), 4.91 – 4.95 (dd, J = 8.00, 10.50 Hz, 1 H), 6.45 – 6.49 (d, J = 7.5 Hz, 1 H), 7.01 – 7.04 (m, 2 H), 7.07 – 7.12 (m, 2 H), 7.28 – 7.31 (td, J = 4.45, 24 Hz, 1 H), 7.33 – 7.36 (t, J = 7.70 Hz, 1 H), 7.39 – 7.41 (m, 3H), 7.47 – 7.52 (m, 2 H), 7.54 – 7.56 (d, J = 7.00 Hz, 1 H), 7.72 – 7.74 (d, J = 8.00 Hz, 1 H), 7.88 – 7.89 (d, J = 8.00 Hz, 1 H). ^{13}C NMR (125 MHz, chloroform- d) δ 37.33, 37.62, 44.32, 45.46, 55.12, 55.49, 110.32, 110.48, 117.59, 120.74, 125.30, 134.41, 146.62, 157.55, 157.78, 171.30, 177.82. HRMS (ESI) m/z : calcd for $\text{C}_{30}\text{H}_{27}\text{O}_6\text{H}^+$, 483.1802; found, 483.1807 (Δ = 1.03 ppm).

1-Naphthyl α -2,4-di(2-nitrophenyl)cyclobutane-1,3-dicarboxylate (4c).

White solid; 38% yield; mp 189.0–190.0 °C. ^1H NMR (500 MHz, $\text{dmsO}-d_6$) δ 4.37 (s, 1 H), 4.68 (s, 1 H), 4.89 (s, 1 H), 5.04 (s, 1 H), 7.79 (m, 15 H), 13.23 (s, 1 H). ^{13}C NMR (125 MHz, $\text{dmsO}-d_6$) δ 40.3, 40.7, 42.2, 43.0, 118.1, 121.3, 124.5, 125.7, 126.4, 128.0, 129.6, 132.2, 132.4, 133.0, 146.3, 148.9, 171.1, 173.6.

1-Naphthyl α -2,4-di(4-tertbutyldimethylsilyloxyphenyl)cyclobutane-1,3-dicarboxylate (4d-i).

White solid; 64% yield; mp 169–170 °C; ^1H NMR (300 MHz, acetone- d_6) δ 0.24 (s, 6H), 0.27 (s, 6H), 1.01 (s, 9H), 1.04 (s, 9H), 4.04 (m, 1H), 4.45 (m, 1H), 4.60 (m, 2H), 6.50 (d, J = 7.50 Hz, 1H), 6.64 (m, 4H), 7.30 (m, 8H), 7.74 (d, J = 8.19 Hz, 1H), 7.89 (d, J = 8.19 Hz, 1 H); ^{13}C NMR (100 MHz, acetone- d_6) δ -5.2, -5.1, 17.9, 17.9, 25.2, 40.9, 41.4, 46.9, 47.1, 117.9, 119.7, 120.1, 121.4, 125.2, 125.7, 126.2, 126.3, 127.7, 129.0, 129.4, 132.0, 132.2, 134.5, 146.8, 154.7, 155.1, 170.7, 172.2.

1-Naphthyl α -2,4-di(4-tertbutyldimethylsilyloxyphenyl)cyclobutane-1,3-dicarboxylate (4d).

To a solution of compound **4d-i** (94 mg, 0.14 mmol) with THF (10 mL) and acetic acid (0.5 mL) was added 1 M TBAF (0.5 mL) and the reaction mixture was concentrated after completion. The reaction mixture was purified by flash chromatography to give **4d** as white solid (48 mg, 76 %); mp >230 °C; ¹H NMR (300 MHz, acetone-d₆) δ 4.00 (dd, *J* = 10.0, 7.5 Hz, 1H), 4.40 (dd, *J* = 10.2, 7.5 Hz, 1H), 4.62-4.51 (m, 2H), 6.61 (d, *J* = 7.5 Hz, 1H), 6.88 (q, *J* = 7.6 Hz, 4H), 7.15 (d, *J* = 8.5 Hz, 1H), 7.49-7.43 (m, 7H), 7.73 (d, *J* = 8.1 Hz, 1H), 7.87 (d, *J* = 8.2 Hz, 1H), 8.30 (s, 1H), 8.49 (s, 1H); ¹³C NMR (100 MHz, acetone-d₆) δ 40.97, 41.40, 47.03, 47.40, 115.06, 115.54, 118.05, 121.56, 125.25, 125.62, 126.14, 126.24, 126.93, 127.56, 128.99, 129.42, 129.99, 130.25, 134.47, 146.95, 156.45, 156.94, 170.85, 172.27.

(1R,2S)-2-Phenylcyclohexyl α-2,4-di(2-methoxyphenyl)cyclobutane-1,3-dicarboxylate (4e).

White solid; 48 % yield; mp 163.0-165.0 °C. ¹H NMR (500 MHz, chloroform-d) δ 0.72 (td, *J* = 12.8, 3.4 Hz, 1 H), 1.27 (ddd, *J* = 40.6, 20.3, 8.1 Hz, 2 H), 1.39 - 1.56 (m, 2 H), 1.66 (t, *J* = 13.7 Hz, 2 H), 1.82 (d, *J* = 13.2 Hz, 1 H), 2.51 - 2.40 (m, 1 H), 3.60 (s, 3 H), 3.67 (s, 4 H), 3.81 (dd, *J* = 10.5, 6.2 Hz, 1 H), 3.85 - 3.93 (m, 1 H), 4.39 - 4.29 (m, 1 H), 4.69 (dd, *J* = 10.7, 6.6 Hz, 1 H), 6.71 (d, *J* = 8.1 Hz, 1 H), 6.76 (d, *J* = 8.1 Hz, 1 H), 6.82 (t, *J* = 7.4 Hz, 1 H), 6.87 (d, *J* = 6.8 Hz, 1 H), 6.94 (t, *J* = 7.4 Hz, 1 H), 7.09 (dd, *J* = 16.6, 7.2 Hz, 3 H), 7.13 - 7.24 (m, 5 H). ¹³C NMR (125 MHz, chloroform-d) δ 24.7, 25.8, 31.4, 37.2, 44.5, 45.8, 54.9, 55.2, 76.0, 110.0, 110.1, 120.2, 126.4, 127.5, 128.0, 143.2, 157.3, 171.6, 178.5. HRMS (ESI) *m/z*: calcd for C₃₂H₃₅O₆H⁺, 515.2428, found, 515.2434 (Δ = 1.2 ppm).

(1R,2S)-2-Phenylcyclohexyl α-2,4-di(2-chlorophenyl)cyclobutane-1,3-dicarboxylate (4f).

White solid; 30% yield; ¹H NMR (500 MHz, chloroform-d) δ 1.32 - 1.6 (m, 4 H), 1.80 (d, *J* = 12.7 Hz, 1 H), 1.85 - 2.02 (m, 2 H), 2.15 - 2.33 (m, 1 H), 2.64 - 2.81 (m, 1 H), 3.52 (t, *J* = 8.4 Hz, 1 H), 3.50 - 3.77 (m, 2 H), 4.48 (dd, *J* = 16.4, 9.8 Hz, 1 H), 4.75 - 4.89 (m, 1 H), 5.16 (dt,

$J = 17.2, 7.6$ Hz, 1 H), 6.78 – 6.96 (m, 2 H), 6.99 (t, $J = 7.5$ Hz, 1 H), 7.02 – 7.11 (m, 3 H), 7.13 – 7.31 (m, 7 H). ^{13}C NMR (125 MHz, chloroform-d) δ 24.9, 25.9, 32.2, 41.5, 49.7, 49.9, 126.2, 126.6, 127.5, 127.7, 128.0, 128.5, 129.7, 134.7, 143.0, 171.0. HRMS (ESI) m/z : calcd for $\text{C}_{30}\text{H}_{29}\text{Cl}_2\text{O}_4\text{H}^+$, 523.1437, found, 523.1443 ($\Delta = 0.10$ ppm).

(1R,2S)-2-Phenylcyclohexyl α -2,4-di(2,6-dichlorophenyl)cyclobutane-1,3-dicarboxylate (4g).

White solid; 30% yield; mp 209.0–201.5 °C. ^1H NMR (700 MHz, chloroform-d) δ 1.40 (dd, $J = 14.4, 11.3$ Hz, 1 H), 1.44 – 1.61 (m, 3 H), 1.79 (d, $J = 13.2$ Hz, 1 H), 1.88 (d, $J = 10.1$ Hz, 1 H), 1.97 (d, $J = 13.4$ Hz, 1 H), 2.18 – 2.26 (m, 1 H), 2.76 (td, $J = 12.1, 3.6$ Hz, 1 H), 4.45 – 4.52 (m, 1 H), 4.70 – 4.78 (m, 1 H), 5.04 (dd, $J = 11.6, 9.3$ Hz, 1 H), 5.21 (td, $J = 10.4, 4.2$ Hz, 1 H), 5.27 (dd, $J = 11.8, 8.1$ Hz, 1 H), 6.95 (t, $J = 8.0$ Hz, 1 H), 6.99 (t, $J = 8.0$ Hz, 1 H), 7.04 (d, $J = 8.0$ Hz, 2 H), 7.12 (d, $J = 7.5$ Hz, 2 H), 7.20 (t, $J = 7.1$ Hz, 1 H), 7.30 – 7.25 (m, 5 H). ^{13}C NMR (175 MHz, chloroform-d) δ 25.0, 25.9, 32.1, 42.3, 49.8, 76.7, 126.6, 127.8, 128.1, 128.5, 129.0, 134.1, 134.3, 143.3, 171.5, 177.5. HRMS (ESI) m/z : calcd for $\text{C}_{30}\text{H}_{27}\text{Cl}_4\text{O}_4\text{H}^+$, 591.0658, found, 591.0661 ($\Delta = 0.50$ ppm).

(1R,2S)-2-Phenylcyclohexyl α -2,4-di(2-bromophenyl)cyclobutane-1,3-dicarboxylate (4h).

White solid; 29% yield; mp 136.0–138.0 °C; ^1H NMR (500 MHz, acetone- d_6) δ 1.33 – 1.44 (m, 1 H), 1.44 – 1.54 (m, 2 H), 1.62 (dd, $J = 12.9, 3.1$ Hz, 1 H), 1.76 (d, $J = 12.9$ Hz, 1 H), 1.86 (dd, $J = 28.6, 11.6$ Hz, 2 H), 2.09 (s, 1 H), 2.16 (d, $J = 9.6$ Hz, 1 H), 2.71 (td, $J = 12.2, 3.6$ Hz, 1 H), 3.61 (dd, $J = 10.1, 7.5$ Hz, 1 H), 3.74 (dd, $J = 10.0, 7.6$ Hz, 1 H), 4.44 – 4.54 (m, 1 H), 4.82 (dd, $J = 9.6, 7.7$ Hz, 1 H), 5.10 (td, $J = 10.2, 4.0$ Hz, 1 H), 7.01 (dd, $J = 12.9, 7.1$ Hz, 3 H), 7.11 (dt, $J = 15.3, 7.4$ Hz, 4 H), 7.21 (d, $J = 3.9$ Hz, 4 H), 7.34 (d, $J = 7.7$ Hz, 1 H), 7.41 (d, $J = 7.8$ Hz, 1 H). ^{13}C NMR (125 MHz, acetone- d_6) δ 25.4, 26.5, 32.9, 34.7, 43.4, 50.4, 76.9, 125.7, 125.9, 127.2, 127.9, 128.4, 129.1,

139.0, 144.1, 171.9, 173.2, HRMS (ESI) m/z : calcd for $C_{30}H_{29}Br_2O_4H^+$, 611.0427, found, 611.0424 ($\Delta = 0.49$ ppm).

(1R,2S)-2-Phenylcyclohexyl α -2,4-di(2-nitrophenyl)cyclobutane-1,3-dicarboxylate (4i).

White solid; 36% yield; mp 200.0-202.0 °C; ^1H NMR (500 MHz, acetone- d_6) δ 1.35 - 1.45 (m, 1 H), 1.49 (dd, $J = 21.7, 11.8$ Hz, 2 H), 1.64 (dd, $J = 12.9, 2.9$ Hz, 1 H), 1.77 (d, $J = 12.9$ Hz, 1 H), 1.86 (dd, $J = 25.5, 11.5$ Hz, 2 H), 1.96 (s, 1 H), 2.70 (td, $J = 12.2, 3.4$ Hz, 1 H), 3.73 (dd, $J = 10.4, 6.0$ Hz, 1 H), 3.92 (t, $J = 9.6$ Hz, 1 H), 4.76 (t, $J = 9.4$ Hz, 1 H), 4.85 (dd, $J = 10.0, 6.0$ Hz, 1 H), 5.01 - 5.13 (m, 1 H), 7.14 (d, $J = 7.6$ Hz, 2 H), 7.20 (d, $J = 4.1$ Hz, 4 H), 7.34 - 7.25 (m, 2 H), 7.37 (t, $J = 7.4$ Hz, 2 H), 7.52 (t, $J = 7.5$ Hz, 1 H), 7.68 (d, $J = 7.8$ Hz, 1 H), 7.72 (d, $J = 8.0$ Hz, 1 H). ^{13}C NMR (125 MHz, acetone- d_6) δ 25.4, 26.5, 32.8, 34.4, 40.9, 50.5, 77.3, 125.2, 125.5, 127.2, 128.5, 128.8, 129.1, 130.1, 130.2, 134.0, 144.0, 149.7, 151.0, 171.4, HRMS (ESI-TOF) (m/z): calcd for $C_{30}H_{29}N_2O_8H^+$, 545.1918, found, 545.1921 ($\Delta = 0.55$ ppm).

9-Fluorenylmethyl α -2,4-di(2-methoxyphenyl)cyclobutane-1,3-dicarboxylate (4j).

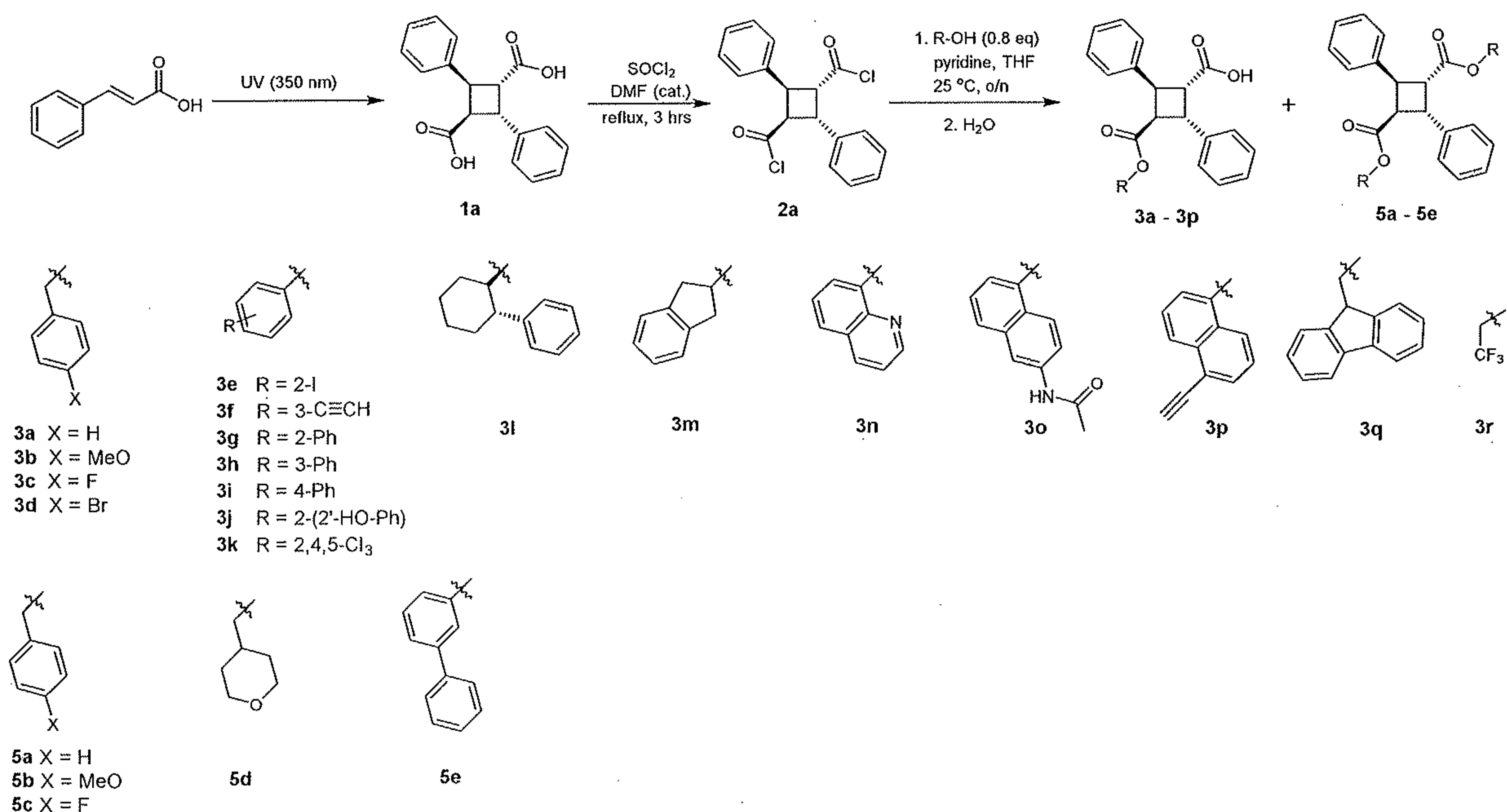
White solid; 70% yield; mp 160.0-162.0 °C; ^1H NMR (500 MHz, chloroform- d) δ 3.76 (d, $J = 11.6$ Hz, 4 H), 3.80 (s, 3 H), 3.96 (dd, $J = 10.6, 8.0$ Hz, 1 H), 4.06 (dd, $J = 10.4, 7.8$ Hz, 1 H), 4.17 - 4.10 (m, 2 H), 4.66 (dd, $J = 10.5, 7.1$ Hz, 1 H), 4.72 (dd, $J = 10.4, 7.8$ Hz, 1 H), 6.82 (d, $J = 8.1$ Hz, 1 H), 6.86 (d, $J = 8.1$ Hz, 1 H), 7.00 (q, $J = 7.1$ Hz, 2 H), 7.24 (t, $J = 7.2$ Hz, 1 H), 7.36 - 7.27 (m, 5 H), 7.41 (t, $J = 6.9$ Hz, 3 H), 7.51 (d, $J = 7.5$ Hz, 1H), 7.77 (d, $J = 7.5$ Hz, 2 H). ^{13}C NMR (125 MHz, chloroform- d) δ 37.1, 37.7, 44.6, 46.9, 55.2, 55.3, 66.4, 110.1, 110.3, 120.0, 125.1, 127.1, 127.7, 141.2, 141.3, 144.0, 144.4, 157.5, 157.6, 172.6, 178.6. HRMS (ESI) m/z : calcd for $C_{34}H_{31}O_6H^+$, 538.2208; found, 538.2228 ($\Delta = 3.72$ ppm).

9-Fluorenylmethyl α -2,4-di(2-chlorophenyl)cyclobutane-1,3-dicarboxylate (4k).

White solid; 55% yield; mp 186.5–187.5 °C. ^1H NMR (500 MHz, DMSO- d_6), δ 4.01 (dd, $J = 9.7, 7.0$ Hz, 1 H), 4.10 – 4.20 (m, 1 H), 4.23 (t, $J = 7.0$ Hz, 1 H), 4.38 (p, $J = 10.6$ Hz, 2 H), 4.60 – 4.70 (m, 1 H), 4.70 – 4.82 (m, 1 H), 7.04 – 7.20 (m, 4 H), 7.26 (dt, $J = 24.3, 8.7$ Hz, 4 H), 7.34 (d, $J = 6.2$ Hz, 1 H), 7.37 – 7.46 (m, 3H), 7.62 (t, $J = 7.5$ Hz, 2H), 7.87 (dd, $J = 7.1, 3.2$ Hz, 2 H), 12.74 (s, 1 H). ^{13}C NMR (125 MHz, DMSO- d_6) δ 41.2, 41.6, 42.5, 46.2, 66.2, 120.2, 125.2, 133.5, 140.7, 143.4, 143.6, 172.1, 173.7. HRMS (ESI) m/z : calcd for $\text{C}_{32}\text{H}_{25}\text{Cl}_2\text{O}_4\text{H}^+$, 543.1124, found, 543.1123 ($\Delta = 0.18$ ppm).

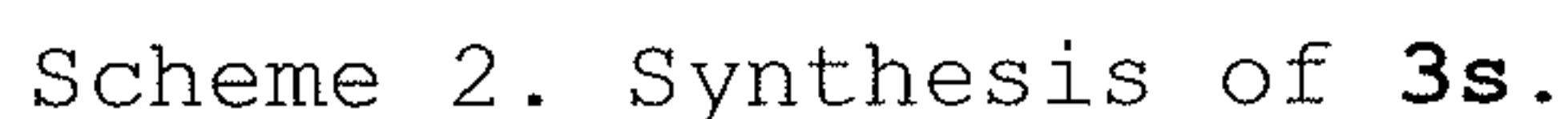
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As **Scheme 1** shows, the general procedure for the synthesis of α -truxillic acid mono-esters **3** commenced with the photochemical reaction of *trans*-cinnamic acid through irradiation in solid state at 360 nm for 3–5 days, depending on the scale. This reaction proceeded through head-to-tail [2+2] cycloaddition to give α -truxillic acid (**1a**) in 93–95% yield. Reaction of **1a** with thionyl chloride in the presence of a catalytic amount of dimethylformamide (DMF) under reflux afforded α -truxillic dichloride (**2a**), which was used directly to the subsequent reactions.

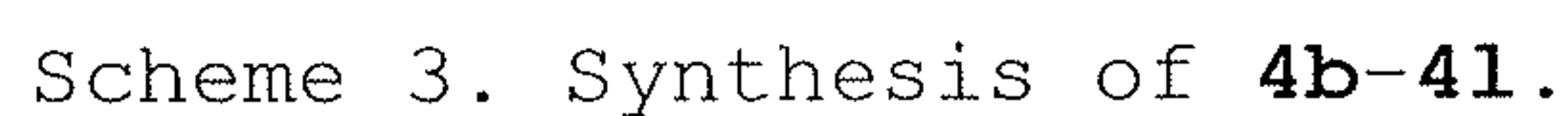


20 Scheme 1. Synthesis of α -truxillic acid mono-esters (**3**)

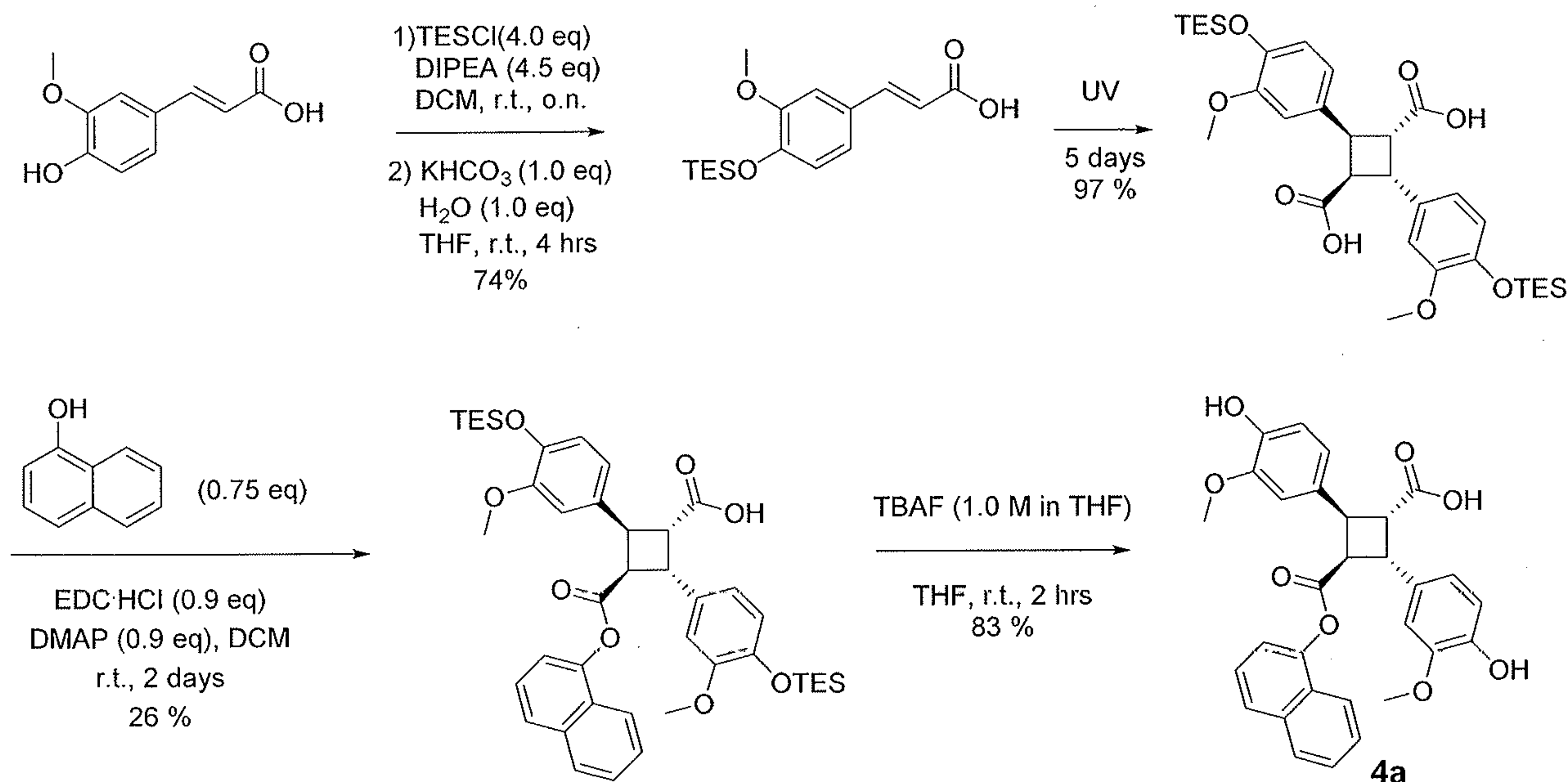
Compound **3s** was synthesized from **3f** through copper-catalyzed click reaction (Scheme 2).



10

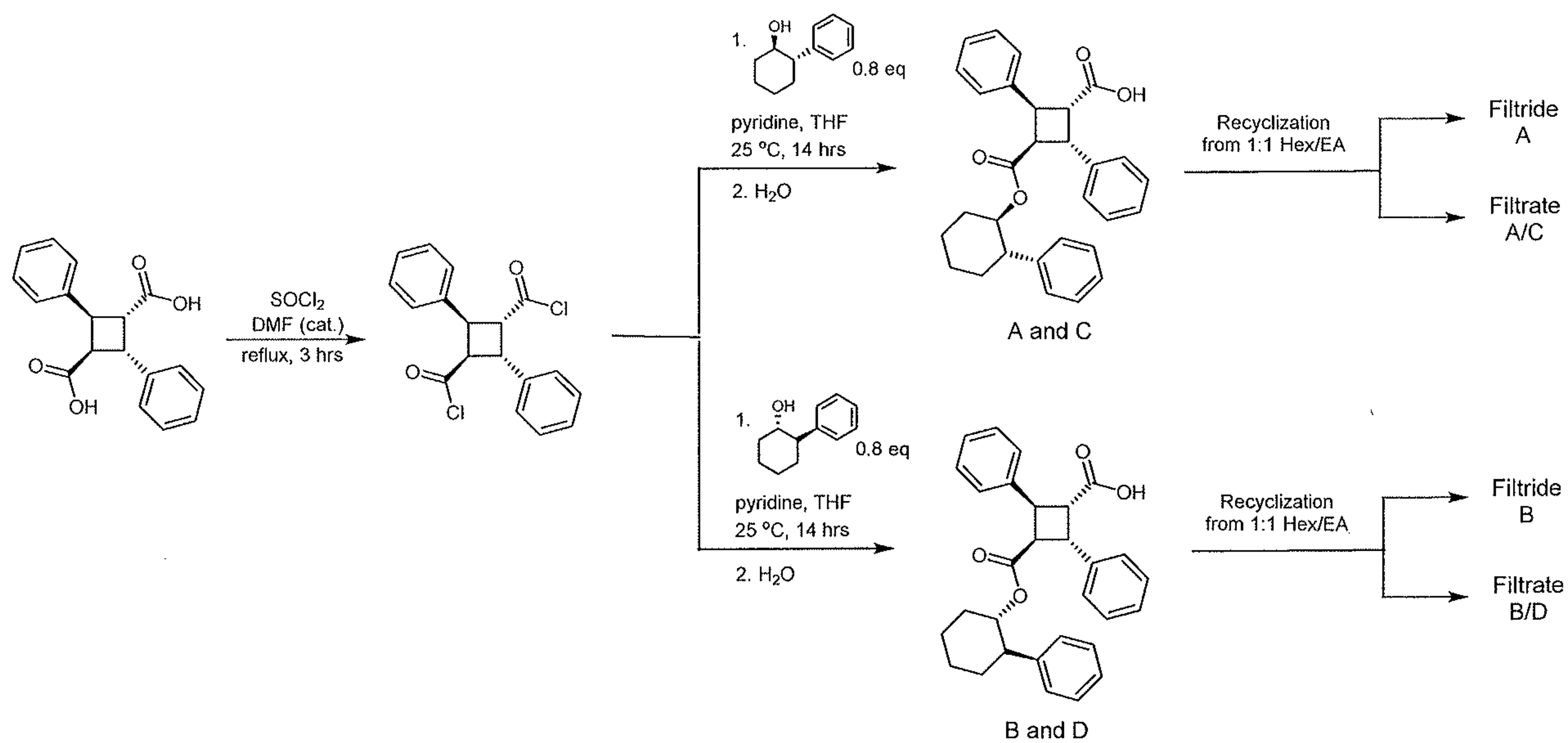
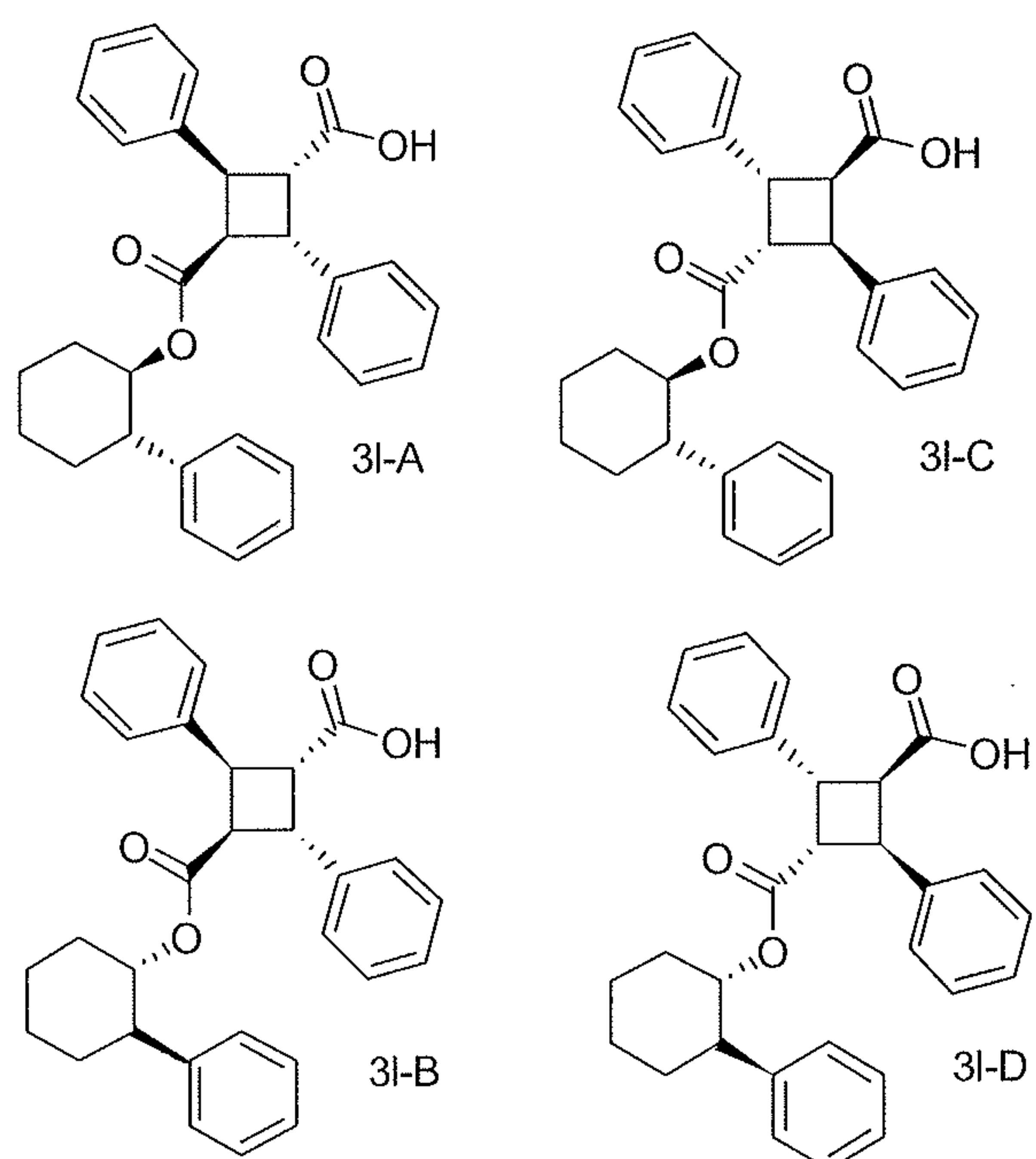


Compound **4a** was synthesized from commercially available 4-hydroxy-3-methoxycinnamic acid with modification in the mono-esterification step, where water-soluble carbodiimide, EDC.HCl, was used as the condensation reagent (**Scheme 4**).



Scheme 4. Synthesis of **4a**.

Compound **31** (racemic) was found to have a binding affinity to FABP5 better than **3A** by a factor of 4-5. However, **31** has four possible stereoisomers due to the chiral center in its ester moiety as well as α -truxillic acid moiety. Therefore, isolation of these four stereoisomers of **31** was attempted by using two pure enantiopures of 2-phenyl-1-cyclohexanol and fractional recrystallization. This efforts led to the successful isolations of two pure stereoisomers **31-A** and **31-B**, bearing (1*R*,2*S*)-2-phenylcyclohex-1-yl and (1*S*,2*R*)-2-phenylcyclohex-1-yl groups, respectively. Also, a diastereomer mixture of **31-A** and **31-C**, as well as **31-B** and **31-D**, was obtained and all those samples were subjected to the fluorescence displacement assays. Interestingly, all these four samples showed the same binding affinity to FABP5 within the standard deviation. The procedures for the synthesis of **31** stereoisomers is illustrated in Scheme 5 and the structures of each diastereomer are recited in Scheme 6.

Scheme 5. Synthesis of **31**.5 Scheme 6. Structures of each diastereomer **31-A** to **31-D**.

Example 2. Determination of K_i

Determination of K_i was derived from DAUDA or NBD-stearate displacement assays (see Table 1).

5 **Table 1.** In vitro affinities (K_i) of FABP inhibitors

(**1**: unsubstituted and substituted α -truxillic acid; **3**: α -truxillic acid mono ester; **4**: substituted α -truxillic acid mono ester)

Compound	FABP3 K_i	FABP5 K_i	FABP7 K_i
3	$2.70 \pm 0.42 \mu\text{M}$	$0.81 \pm 0.09 \mu\text{M}$	$0.45 \pm 0.07 \mu\text{M}$
3-A	$3.26 \pm 0.70 \mu\text{M}$	$0.78 \pm 0.14 \mu\text{M}$	$0.89 \pm 0.24 \mu\text{M}$
3-B	$2.82 \pm 0.10 \mu\text{M}$	$0.80 \pm 0.14 \mu\text{M}$	$0.66 \pm 0.16 \mu\text{M}$
3a	$>10 \mu\text{M}$	$3.81 \pm 0.53 \mu\text{M}$	$0.53 \pm 0.12 \mu\text{M}$
3b	$>10 \mu\text{M}$	$2.15 \pm 0.10 \mu\text{M}$	$1.14 \pm 0.06 \mu\text{M}$
3c	$>10 \mu\text{M}$	$2.42 \pm 0.18 \mu\text{M}$	$1.65 \pm 0.21 \mu\text{M}$
3d	$>10 \mu\text{M}$	$1.58 \pm 0.16 \mu\text{M}$	$1.25 \pm 0.03 \mu\text{M}$
3e	$1.18 \pm 0.10 \mu\text{M}$	$1.34 \pm 0.21 \mu\text{M}$	$0.94 \pm 0.34 \mu\text{M}$
3f	$>10 \mu\text{M}$	$0.89 \pm 0.15 \mu\text{M}$	$0.78 \pm 0.12 \mu\text{M}$
3g	$0.70 \pm 0.42 \mu\text{M}$	$0.77 \pm 0.08 \mu\text{M}$	$0.35 \pm 0.12 \mu\text{M}$
3h	$9.75 \pm 0.79 \mu\text{M}$	$0.85 \pm 0.22 \mu\text{M}$	$0.74 \pm 0.17 \mu\text{M}$
3i	$3.93 \pm 0.16 \mu\text{M}$	$2.52 \pm 0.36 \mu\text{M}$	$2.27 \pm 0.03 \mu\text{M}$
3j	$3.52 \pm 0.53 \mu\text{M}$	$1.59 \pm 0.43 \mu\text{M}$	$0.54 \pm 0.18 \mu\text{M}$
3k	$2.98 \pm 0.85 \mu\text{M}$	$0.80 \pm 0.11 \mu\text{M}$	$0.54 \pm 0.02 \mu\text{M}$
3l	$1.08 \pm 0.37 \mu\text{M}$	$0.21 \pm 0.02 \mu\text{M}$	$0.40 \pm 0.03 \mu\text{M}$
3l-A	---	$0.20 \pm 0.03 \mu\text{M}$	---
3l-B	---	$0.18 \pm 0.03 \mu\text{M}$	---
3l-C	---	$0.21 \pm 0.02 \mu\text{M}$	---
3l-D	---	$0.18 \pm 0.01 \mu\text{M}$	---
3m	$>10 \mu\text{M}$	$1.57 \pm 0.15 \mu\text{M}$	$2.41 \pm 0.09 \mu\text{M}$
3o	$2.82 \pm 0.18 \mu\text{M}$	$0.97 \pm 0.18 \mu\text{M}$	$1.12 \pm 0.45 \mu\text{M}$
3p	$4.94 \pm 0.31 \mu\text{M}$	$3.92 \pm 0.75 \mu\text{M}$	$1.03 \pm 0.22 \mu\text{M}$
3q	$>10 \mu\text{M}$	$2.56 \pm 0.16 \mu\text{M}$	$2.70 \pm 0.62 \mu\text{M}$
3r	$>10 \mu\text{M}$	$>10 \mu\text{M}$	$1.59 \pm 0.24 \mu\text{M}$
3s	$>10 \mu\text{M}$	$2.17 \pm 0.32 \mu\text{M}$	$0.50 \pm 0.11 \mu\text{M}$
3t	$>10 \mu\text{M}$	$>10 \mu\text{M}$	$1.06 \pm 0.07 \mu\text{M}$

4a	1.06 ± 0.19 μ M	>10 μ M	2.12 ± 0.19 μ M
4b	0.69 ± 0.17 μ M	0.55 ± 0.05 μ M	0.67 ± 0.04 μ M
4d	2.30 ± 0.47	>10 μ M	1.06 ± 0.34
4e	0.40 ± 0.08 μ M	0.68 ± 0.06 μ M	0.40 ± 0.03 μ M
4f	>10 μ M	1.70 ± 0.33 μ M	>10 μ M
4g	>10 μ M	1.23 ± 0.18 μ M	6.32 ± 0.96 μ M
4h	>10 μ M	2.76 ± 0.16 μ M	>10 μ M
4j	>10 μ M	1.72 ± 0.12 μ M	>10 μ M
4k	>10 μ M	0.89 ± 0.05 μ M	3.54 ± 0.77 μ M

*All values derived from DAUDA, ANS, or NBD-stearate fluorescence binding assays. K_i values represent an average $\pm$ S.E. of at least three independent experiments.

5 The compounds of Table 1 display unexpected selectivity toward FABP3, FABP 5 or FABP 7. Compounds contained herein are selective for one of FABP3, FABP5 or FABP7, or are selective for both FABP5 and FABP7, but not for FABP3.

10 **Example 3. AEA uptake in cells**

It was previously shown that FABPs are intracellular carriers that shuttle endocannabinoids and related N-acylethanolamines to intracellular sites, such as FAAH for hydrolysis (Kaczocha, M. et al. 2209; Kaczocha, M. et al. 2012). Pharmacological or genetic inhibition
 15 of FABPs reduces AEA catabolism in cells, confirming an essential role for these proteins in endocannabinoid inactivation. Therefore, it is examined as to whether the present FABP inhibitors reduce FABP-mediated AEA uptake in cells.

20 Compounds shown in Table 1 inhibit cellular AEA accumulation and fail to reduce AEA uptake in cells bearing a knockdown of FABP5, the main FABP expressed in HeLa cells. Additionally, the compounds fail to inhibit FAAH. Accordingly, compounds shown in Table 1 selectively inhibit FABP.

Example 4. Antinociceptive and anti-inflammatory effects in mice.

Similar to cannabinoid receptor agonists, inhibitors of endocannabinoid inactivation produce anti-inflammatory and antinociceptive effects (Cravatt, B.F. et al. 2001; Lichtman, A.H. 1990). Importantly, FAAH inhibitors lack the untoward psychotropic effects of cannabinoid receptor agonists (Cravatt, B.F. et al. 2004), highlighting the therapeutic advantage of pharmacologically targeting endocannabinoid inactivation. Because inhibition of AEA transport to FAAH reduces AEA inactivation, we hypothesized that FABP inhibitors may likewise possess antinociceptive and anti-inflammatory properties. The present compounds are examined using two nociceptive models: the formalin test and carrageenan-induced thermal hyperalgesia.

In the formalin test, injection of formalin results in the induction of two temporally distinct phases of pain with the first phase (0-5 min) representing nociceptor activation and the second phase (15-45 min) representing inflammatory pain and central sensitization. Compounds 4-listed in Table 1 reduce nocifensive behavior during the first phase of the formalin test.

The present compounds alleviate inflammatory pain induced by intraplantar injection of λ -carrageenan. Compounds listed in Table 1 (20 mg/kg, i.p. is used) reduce carrageenan-induced thermal hyperalgesia and paw edema.

Example 5. Additional *In vivo* mouse studies and Pharmacokinetic Data

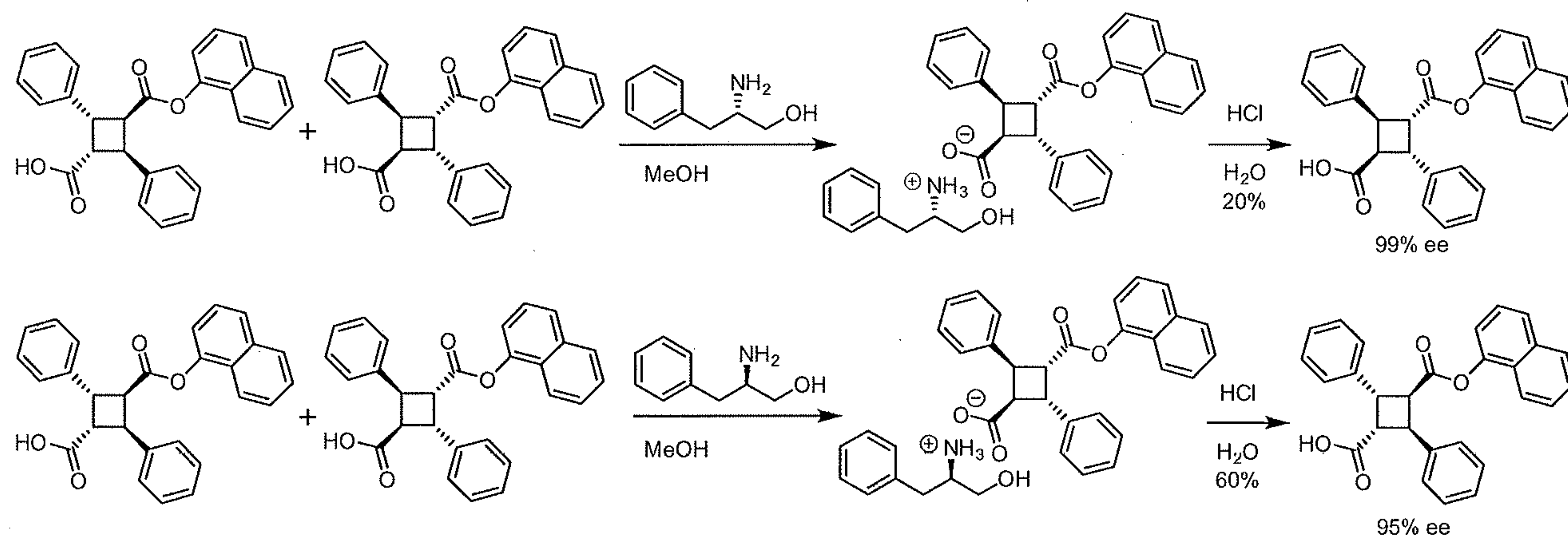
Compound listed in Table 1 are administered by a single injection and plasma and brain concentrations were analyzed over 24 hours. Compounds shown in Table 1 also reduce the first and second phases of formalin-induced nociception in mice. Compounds listed in Table 1 reduce acetic acid-induced writhing in mice. Compounds listed in Table 1 also elevate brain levels of the endocannabinoid anandamide (AEA).

Example 6. Optical Resolution of α -Truxillic Acid Monoesters

As shown below (Scheme 7), starting from a racemic mixture, high enantiomeric excess of each of the two enantiomers of compound **17** can be achieved by chiral resolution via recrystallization.

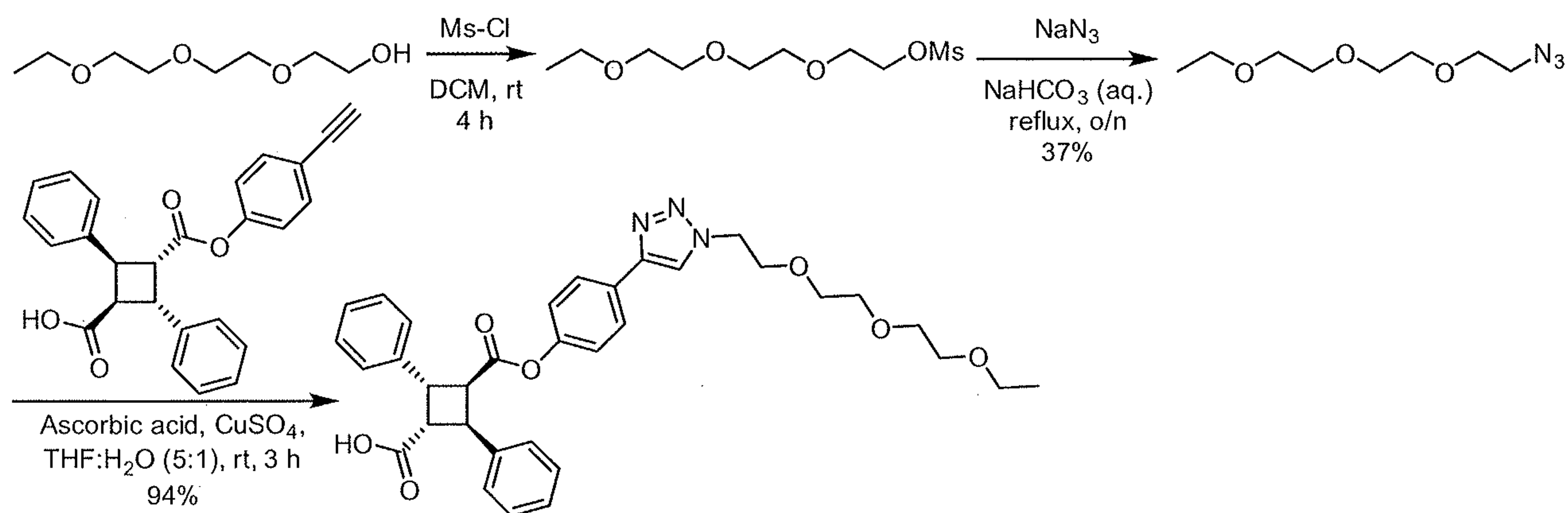
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Scheme 7.

**Example 7. Synthesis of a PEG analogue**

10 As shown below (Scheme 8), PEG analogues may be prepared via click reaction. The example below shows a PEG analogue which was prepared from a PEG-azide and an alkyne-containing α -Truxillic Acid Monoester.

Scheme 8.



15

Discussion

The design of truxillic acid based compounds targeting early stage (neurogenic pain response) and late stage (inflammatory pain response) has coincidentally been studied recently based on the structure of the natural product (-)-incarvillateine. This natural product was first isolated from the plant species *Incarvillea sinensis*, which has been known in traditional Chinese medicine to treat rheumatism and pain. Interestingly, the isolated (-)-incarvillateine was found to possess antinociceptive properties on the same level as morphine.

Recently research (Chi, Y. et al. 2005) was carried out to identify potential leads for commercialization purposes focusing on di-esters and di-carboxylic acid derivatives. Despite the lack of target identification, α and β -truxillic acid di-ester and di-carboxylic acid derivatives were designed, synthesized and tested against early and late stage pain in formalin induced mouse models to study the effects of each analogue in-vivo. The results of the SAR study clearly showed that α -truxillic acid alone provided the best antinociceptive agent in both early and late stage pain. Although the para-hydroxyl functionalization on the phenyl rings slightly improved late stage pain relief it drastically reduced early stage pain relief.

As described herein, α -2,4-diphenyl-cyclobutane-1,3-dicarboxylic acid mono-esters are reversible inhibitors of FABPs. These compounds bind to FABPs and block the shuttling of endocannabinoids within the cell and thereby increase the endogenous levels of the endocannabinoid anandamide by circumventing degradation by FAAH. Increased levels of anandamide result in the activation of the CB1 pathway leading to antinociceptive pain relief and reduction of inflammation which is shown in a formalin induced mouse model (*in-vivo* results).

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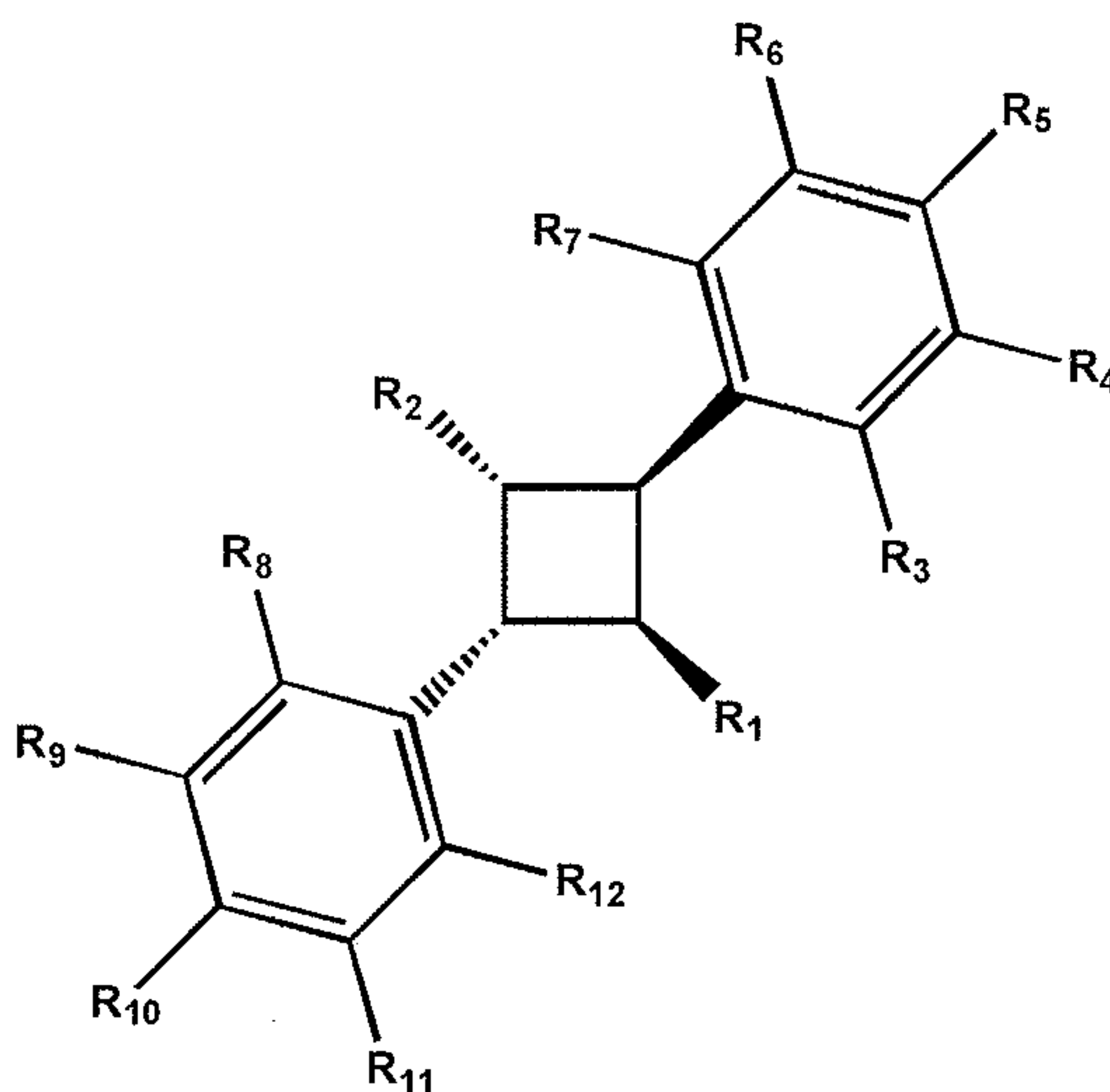
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What is claimed is:

1. A compound of having the structure:



wherein

one of R_1 or R_2 is $-C(=O)OH$ and the other of R_1 or R_2 is $-C(=O)OR_{13}$ or $-C(=O)O\text{-alkyl-}R_{14}$,

wherein

R_{13} is cycloalkyl, aryl or heteroaryl, and

R_{14} is CF_3 , cycloalkyl, cycloheteroalkyl, aryl or heteroaryl; and $R_3, R_4, R_5, R_6, R_7, R_8, R_9, R_{10}, R_{11}$ and R_{12} are each independently, H $-OR_{15}$ or halogen

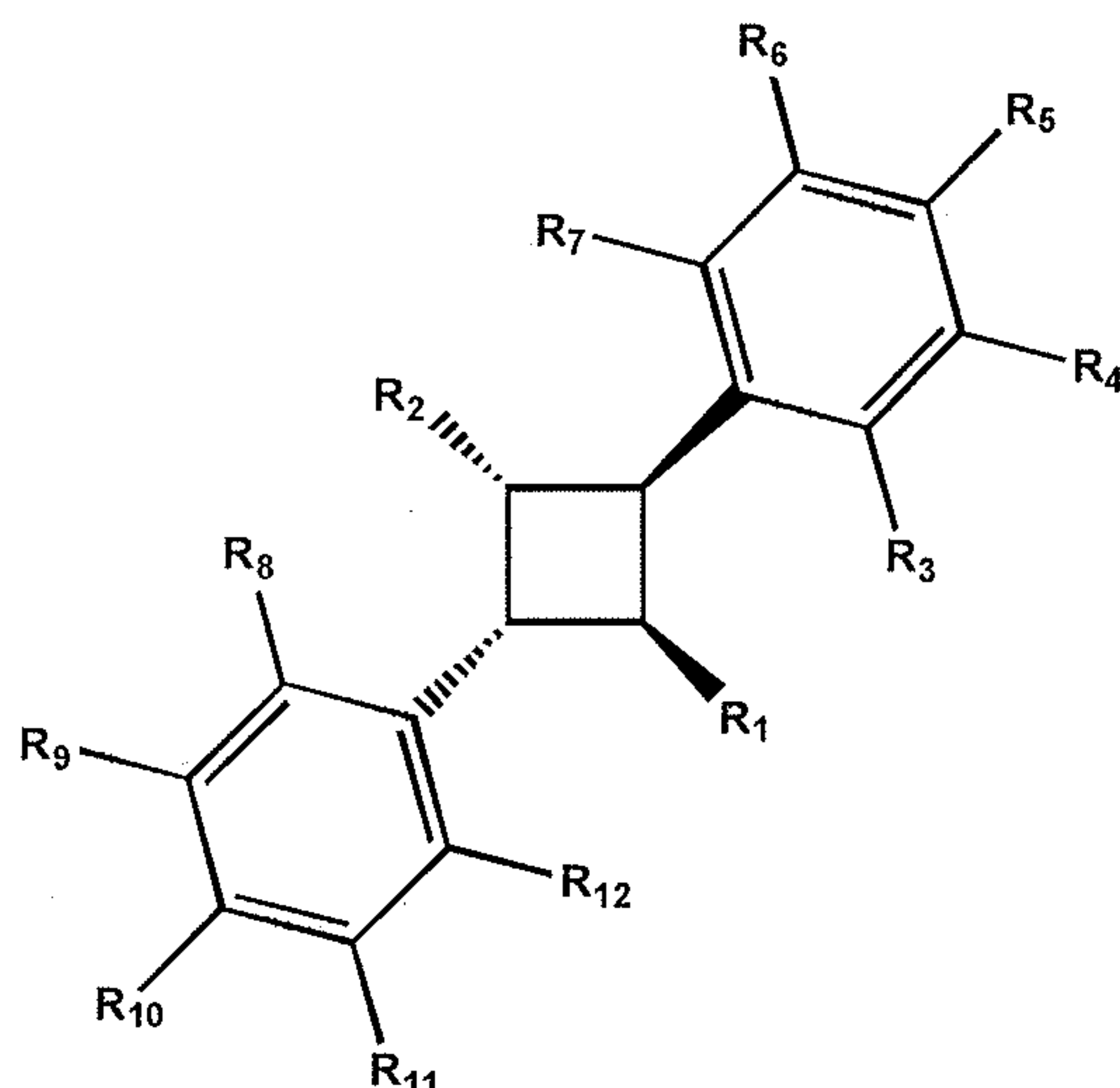
wherein R_{15} is H, C_{1-10} alkyl, C_{2-10} alkenyl, C_{2-10} alkynyl, aryl, or heteroaryl,

wherein when one of R_1 or R_2 is $-C(=O)OH$ and $R_3, R_4, R_5, R_6, R_7, R_8, R_9, R_{10}, R_{11}$ and R_{12} are each H, then the other of R_1 or R_2 is other than $-C(=O)OR_{13}$ where R_{13} is 1-naphthalene or 2-naphthalene or $-C(=O)O\text{-alkyl-}R_{14}$ where the alkyl is a branched C_2 alkyl and the R_{14} is phenyl,

or an enantiomer or racemate thereof;

or a pharmaceutically acceptable salt thereof.

2. The compound of claim 1 having the structure:



wherein

one of R_1 or R_2 is $-C(=O)OH$ and the other of R_1 or R_2 is $-C(=O)OR_{13}$ or $-C(=O)O\text{-alkyl-}R_{14}$,

wherein

R_{13} is cycloalkyl, aryl or heteroaryl, and

R_{14} is CF_3 , cycloalkyl, aryl or heteroaryl; and

R_3 , R_4 , R_5 , R_6 , R_7 , R_8 , R_9 , R_{10} , R_{11} and R_{12} are each independently, H $-OR_{15}$ or halogen

wherein R_{15} is H, C_{1-10} alkyl, C_{2-10} alkenyl, C_{2-10} alkynyl, aryl, or heteroaryl,

wherein when one of R_1 or R_2 is $-C(=O)OH$ and R_3 , R_4 , R_5 , R_6 , R_7 , R_8 , R_9 , R_{10} , R_{11} and R_{12} are each H, then the other of R_1 or R_2 is other than $-C(=O)OR_{13}$ where R_{13} is 1-naphthalene or 2-naphthalene, or $-C(=O)O\text{-alkyl-}R_{14}$ where the alkyl is a branched C_2 alkyl and the R_{14} is phenyl,

or an enantiomer or racemate thereof;

or a pharmaceutically acceptable salt thereof.

3. The compound of claim 1 or 2, wherein when one of R_1 or R_2 is $-C(=O)OH$ and R_3 , R_4 , R_5 , R_6 , R_7 , R_8 , R_9 , R_{10} , R_{11} and R_{12} are each H, then the other of R_1 or R_2 is other than $-C(=O)OR_{13}$ where R_{13} is tolyl, 1-naphthalene or

2-naphthalene, or $-C(=O)O\text{-alkyl-R}_{14}$ where the alkyl is a branched C_2 alkyl and the R_{14} is phenyl.

4. The compound of claim 1 or 2, wherein when one of R_1 or R_2 is $-C(=O)OH$ and the other of R_1 or R_2 is $-C(=O)OR_{13}$ where R_{13} is 1-naphthalene or 2-naphthalene, then one of $R_3, R_4, R_5, R_6, R_7, R_8, R_9, R_{10}, R_{11}$ and R_{12} is other than $-H$.

5. The compound of claim 1 or 2, wherein when one of R_1 or R_2 is $-C(=O)OH$ and the other of R_1 or R_2 is $-C(=O)OR_{13}$ where R_{13} is 1-naphthalene or 2-naphthalene, then two of $R_3, R_4, R_5, R_6, R_7, R_8, R_9, R_{10}, R_{11}$ and R_{12} is other than $-H$.

6. The compound of claim 1 or 2, wherein when one of R_1 or R_2 is $-C(=O)OH$ and the other of R_1 or R_2 is $-C(=O)OR_{13}$ where R_{13} is 1-naphthalene or 2-naphthalene, then four of $R_3, R_4, R_5, R_6, R_7, R_8, R_9, R_{10}, R_{11}$ and R_{12} is other than $-H$.

7. The compound of any one of claims 1-5,
 wherein
 one of R_1 or R_2 is $-C(=O)R_{13}$,
 wherein R_{13} is cycloalkyl, aryl or heteroaryl; and
 the other of R_1 or R_2 is $-C(=O)OH$.

8. The compound of any one of claims 1-5,
 wherein
 one of R_1 or R_2 is $-C(=O)O\text{-alkyl-R}_{14}$,
 wherein R_{14} is CF_3 , cycloalkyl, aryl or heteroaryl; and
 the other of R_1 or R_2 is $-C(=O)OH$.

9. The compound of claim 8,
 wherein
 one of R_1 or R_2 is $-C(=O)O\text{-(C}_{1-6}\text{ alkyl)-R}_{14}$,
 wherein R_{14} is CF_3 , cycloalkyl, aryl or heteroaryl; and
 the other of R_1 or R_2 is $-C(=O)OH$.

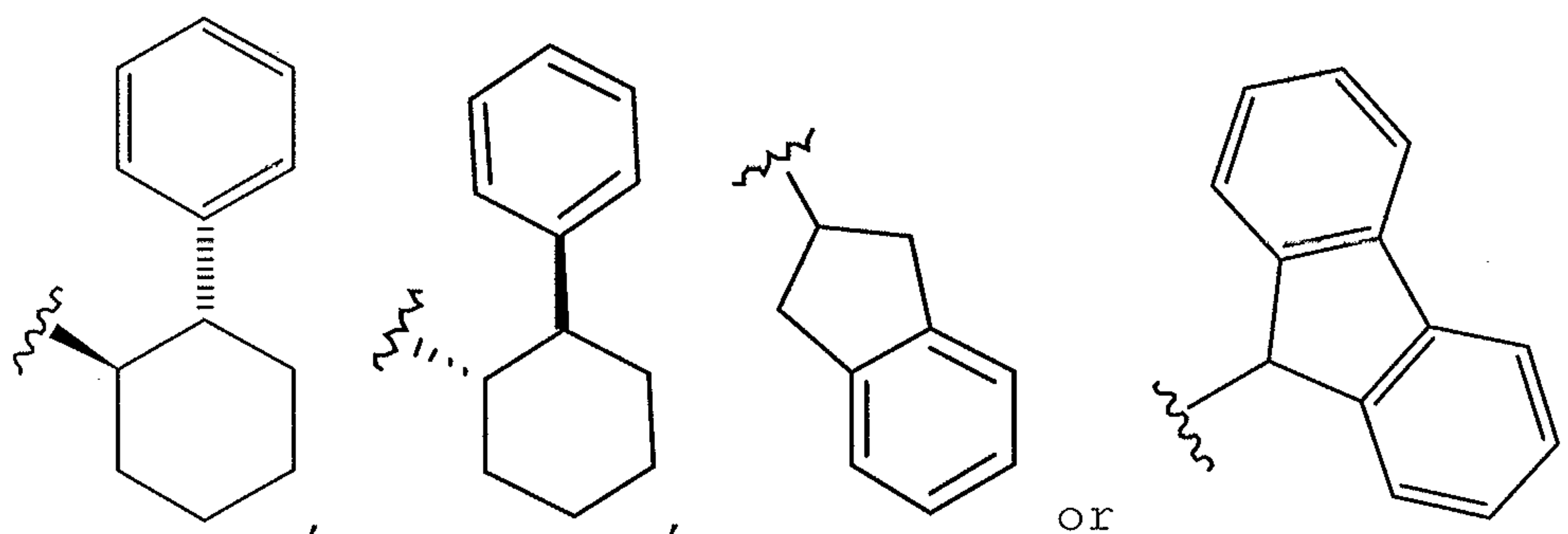
10. The compound of claim 9,
wherein
one of R₁ or R₂ is -C(=O)O-CH₃-R₁₄,
wherein R₁₄ is CF₃, cycloalkyl, aryl or heteroaryl; and
the other of R₁ or R₂ is -C(=O)OH.
11. The compound of any one of claims 7-10, wherein the aryl is a substituted aryl.
12. The compound of claim 10, wherein the aryl is substituted with a halogen, -OH, heteroaryl, C₂-C₆ alkynyl or -O(alkyl).
13. The compound of claim 11, wherein the aryl is substituted with a amide, aryl or hydroxyaryl.
14. The compound of claim 11, wherein the aryl is substituted with a F, Cl, Br, -OH, triazolyl, C₂ alkynyl or -OCH₃.
15. The compound of claim 11, wherein the aryl is substituted with a F, Cl, Br, -OH, I, -NHC(O)CH₃, triazolyl, C₂ alkynyl, phenyl, o-hydroxyphenyl or -OCH₃.
16. The compound of any one of claims 7-10, wherein the heteroaryl is a substituted heteroaryl.
17. The compound of claim 16, wherein the heteroaryl is substituted with a halogen, -OH, heteroaryl, C₂-C₆ alkynyl or -O(alkyl).
18. The compound of claim 16, wherein the heteroaryl is substituted with an amide, aryl or hydroxyaryl.
19. The compound of claim 16, wherein the heteroaryl is substituted with an F, Cl, Br, -OH, triazolyl, C₂ alkynyl or -OCH₃.

20. The compound of claim 16, wherein the heteroaryl is substituted with a F, Cl, Br, -OH, I, -NHC(O)CH₃, triazolyl, C₂ alkynyl, phenyl, o-hydroxyphenyl or -OCH₃.

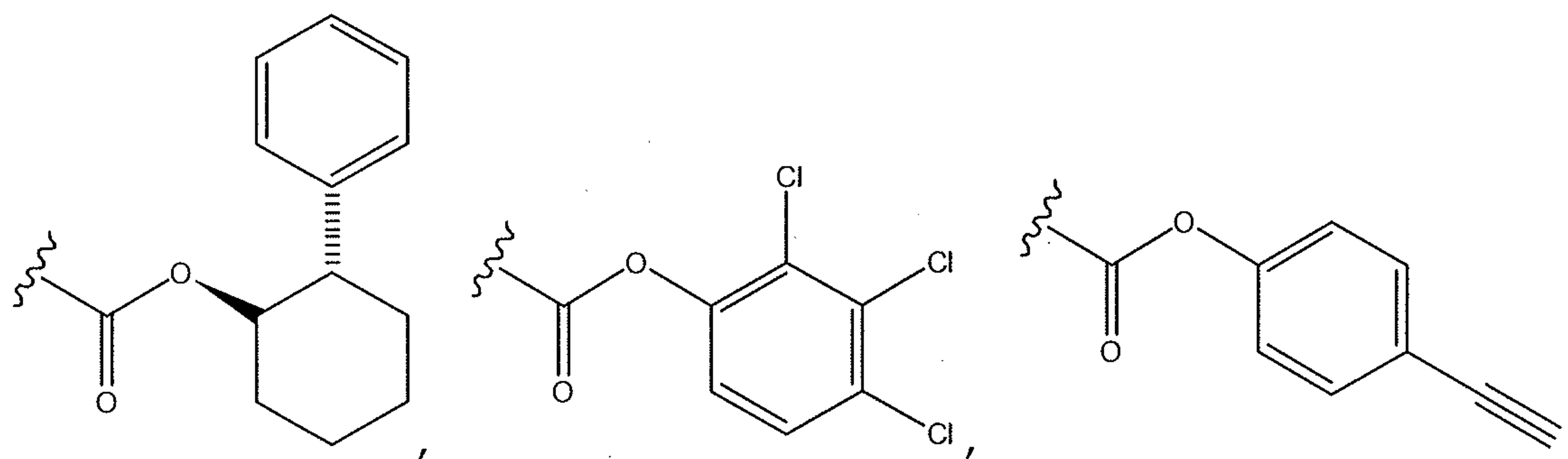
21. The compound of any one of claims 7-10, wherein the cycloalkyl is a substituted cycloalkyl.

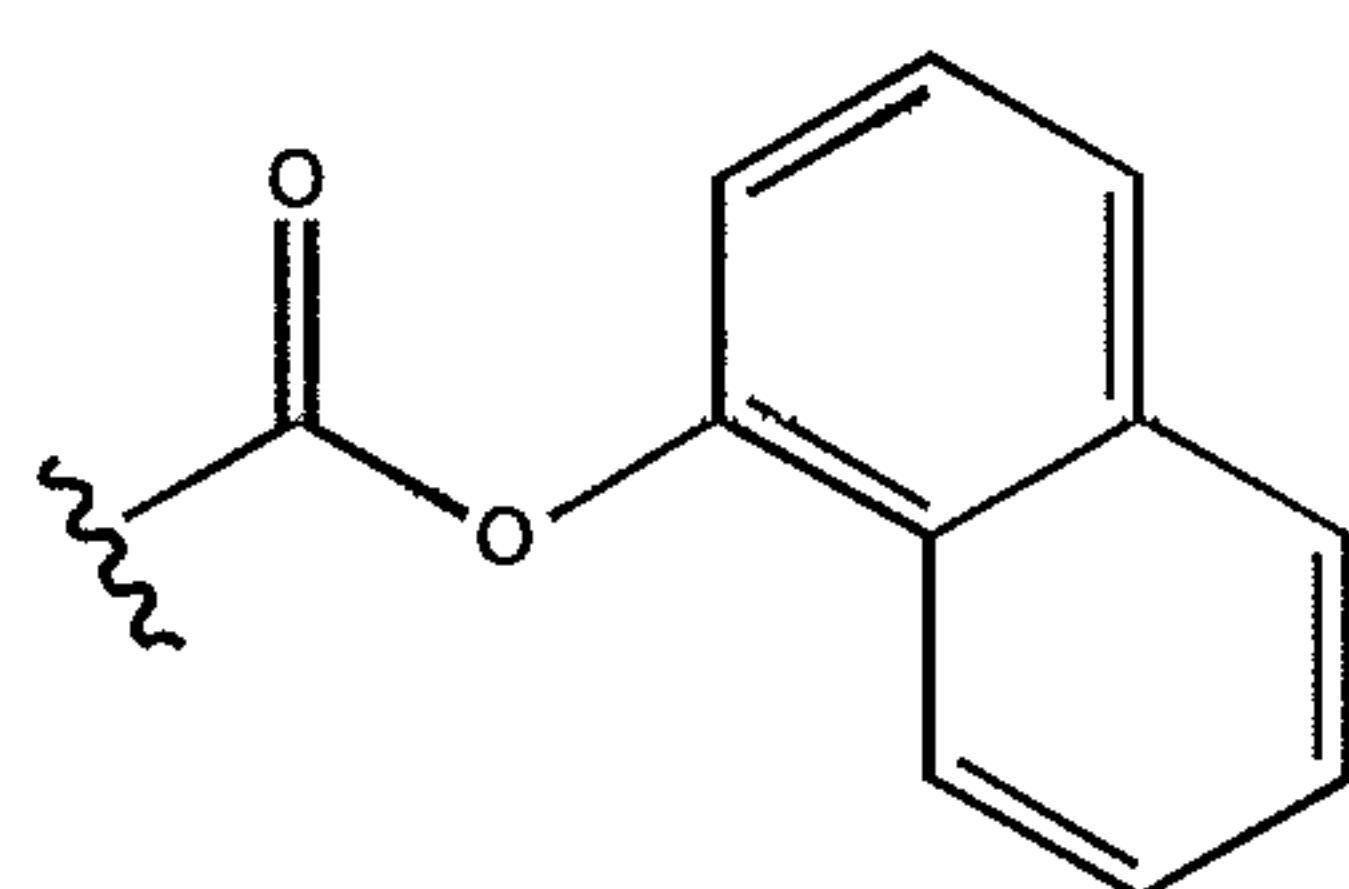
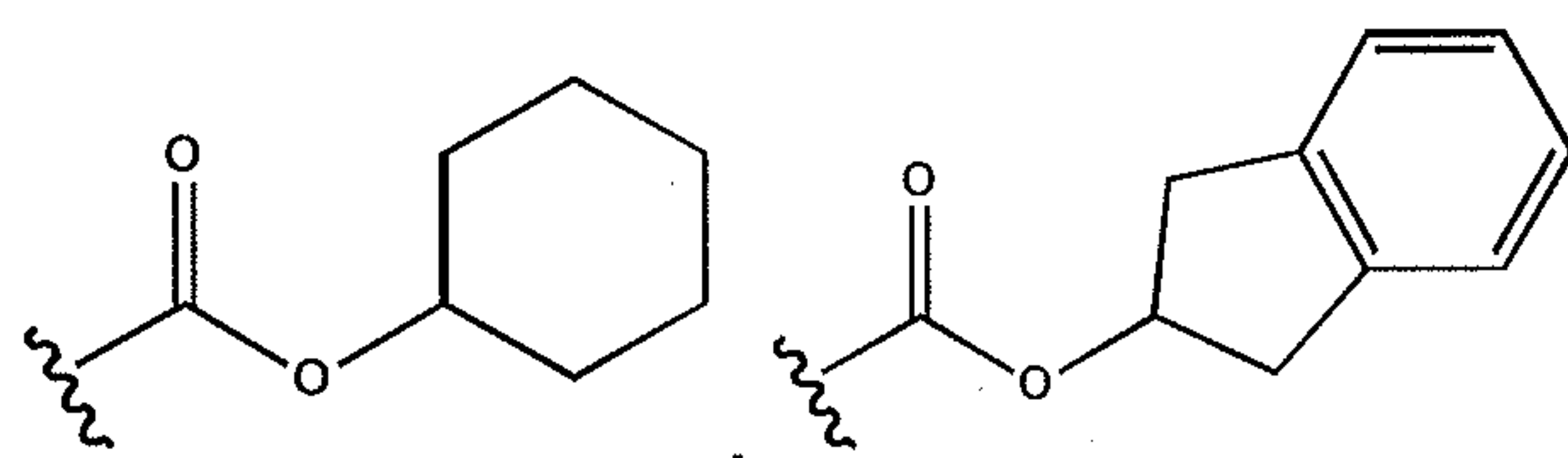
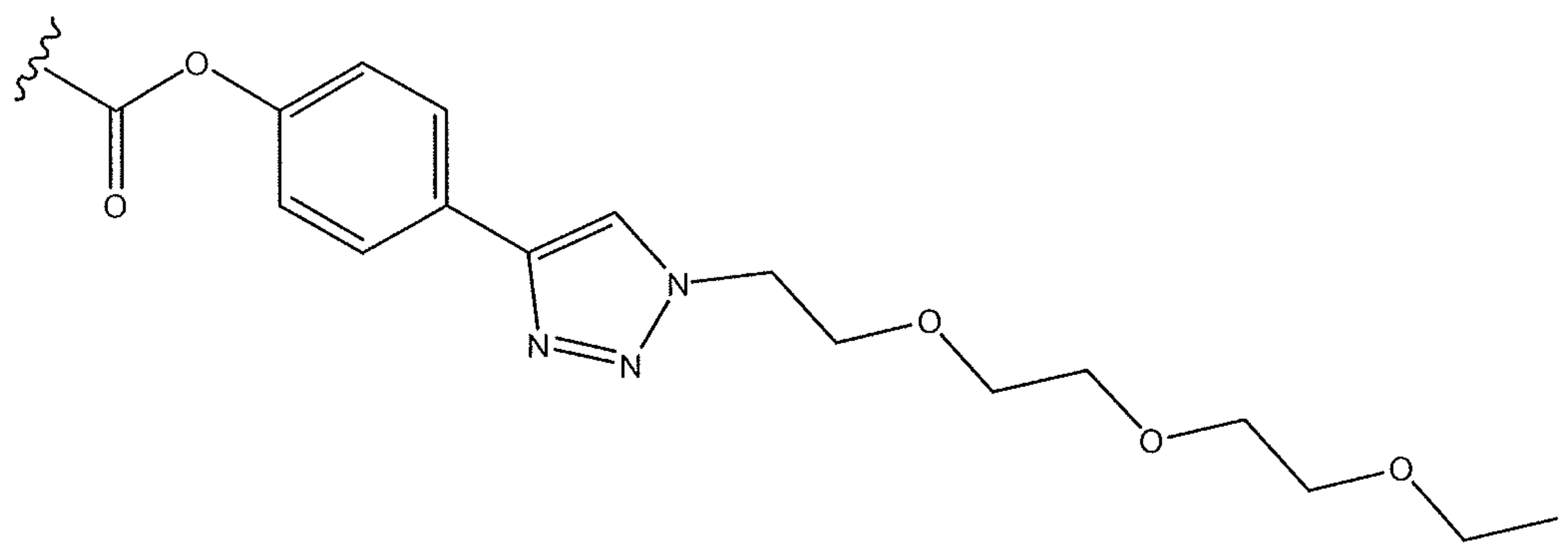
22. The compound of claim 21, wherein the cycloalkyl is substituted with a phenyl or a fused benzo group.

23. The compound of claim 22, wherein the cycloalkyl is:

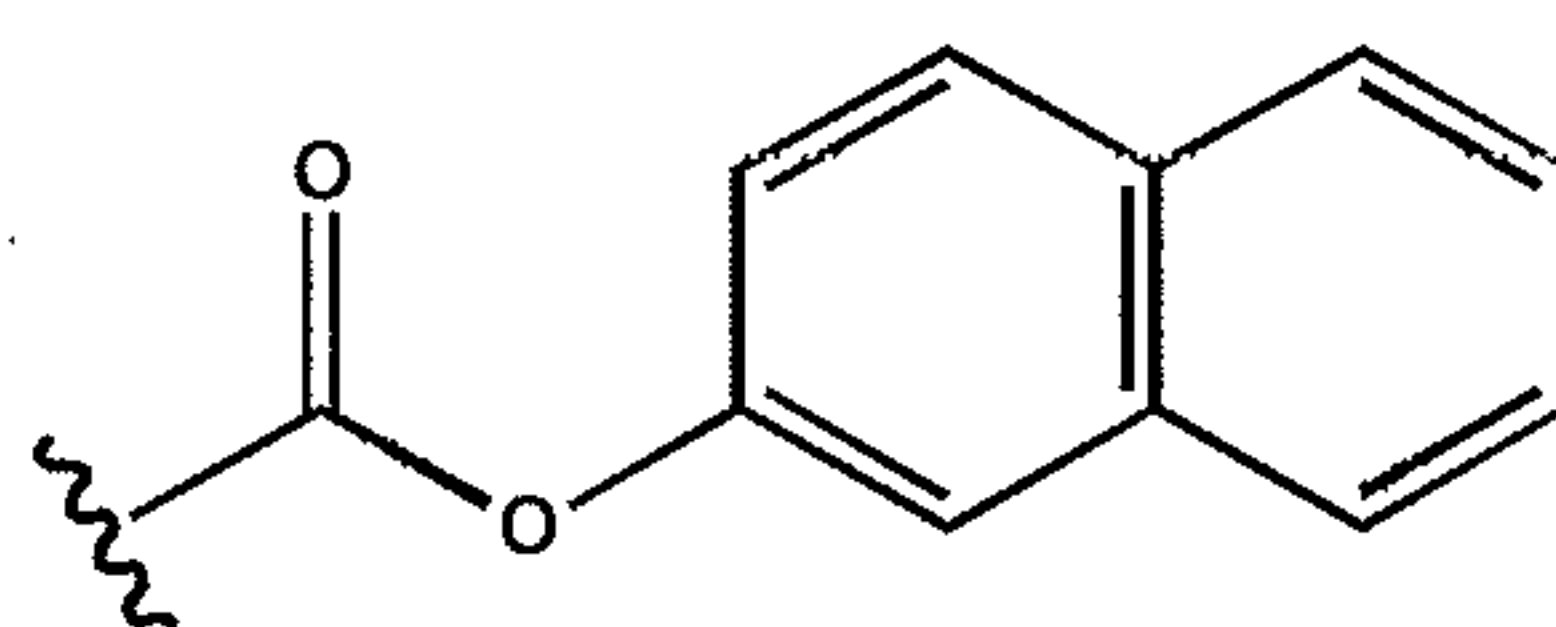


24. The compound of any one of claims 1-6,
wherein
one of R₁ or R₂ is





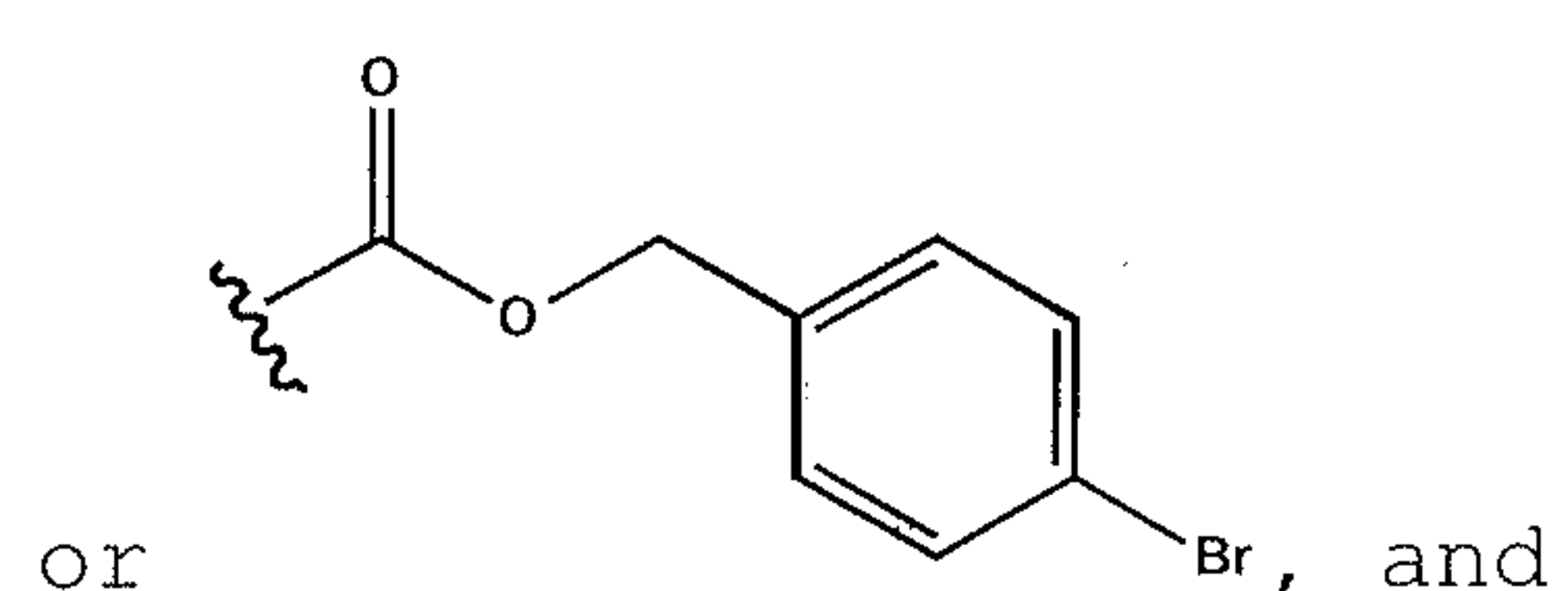
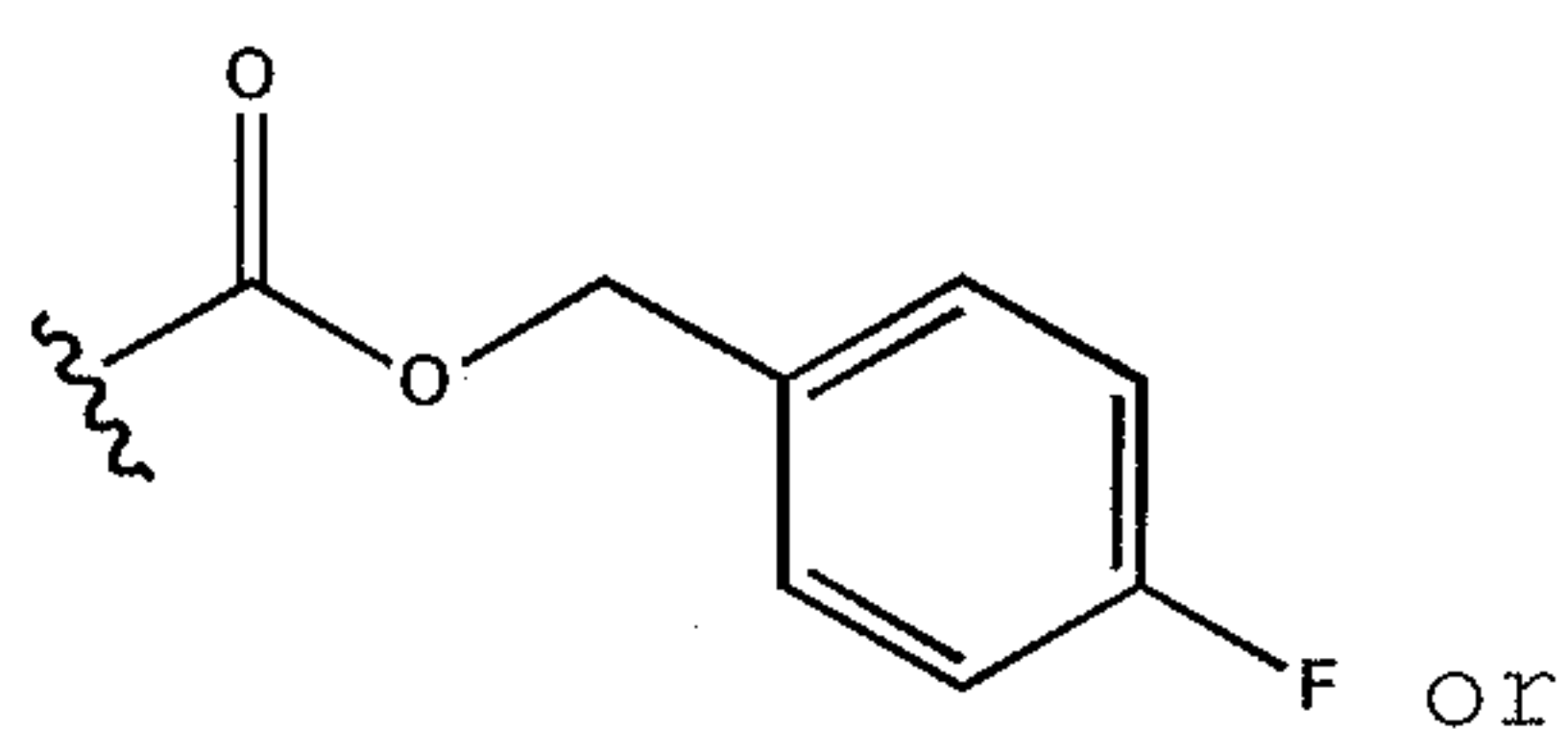
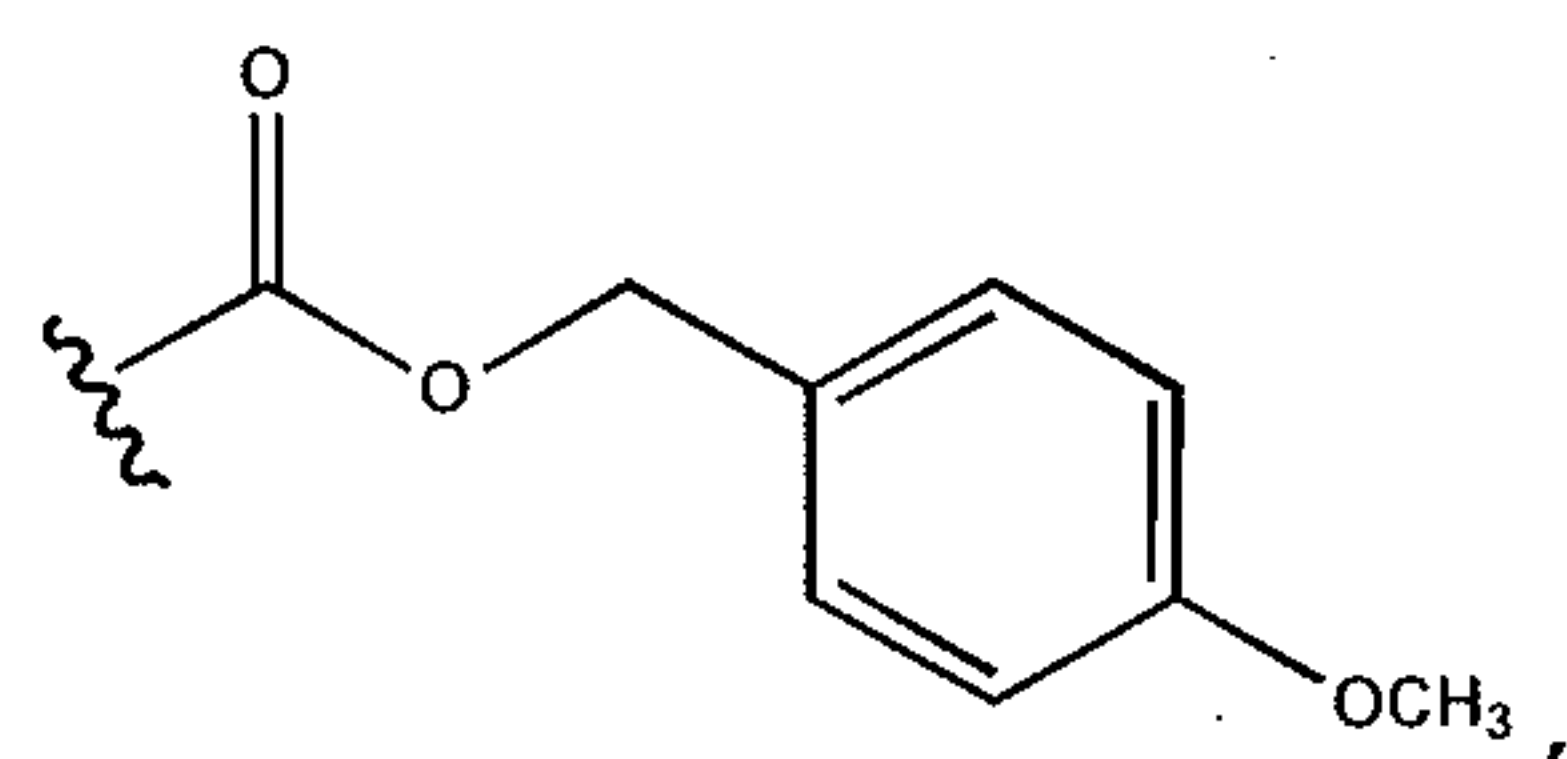
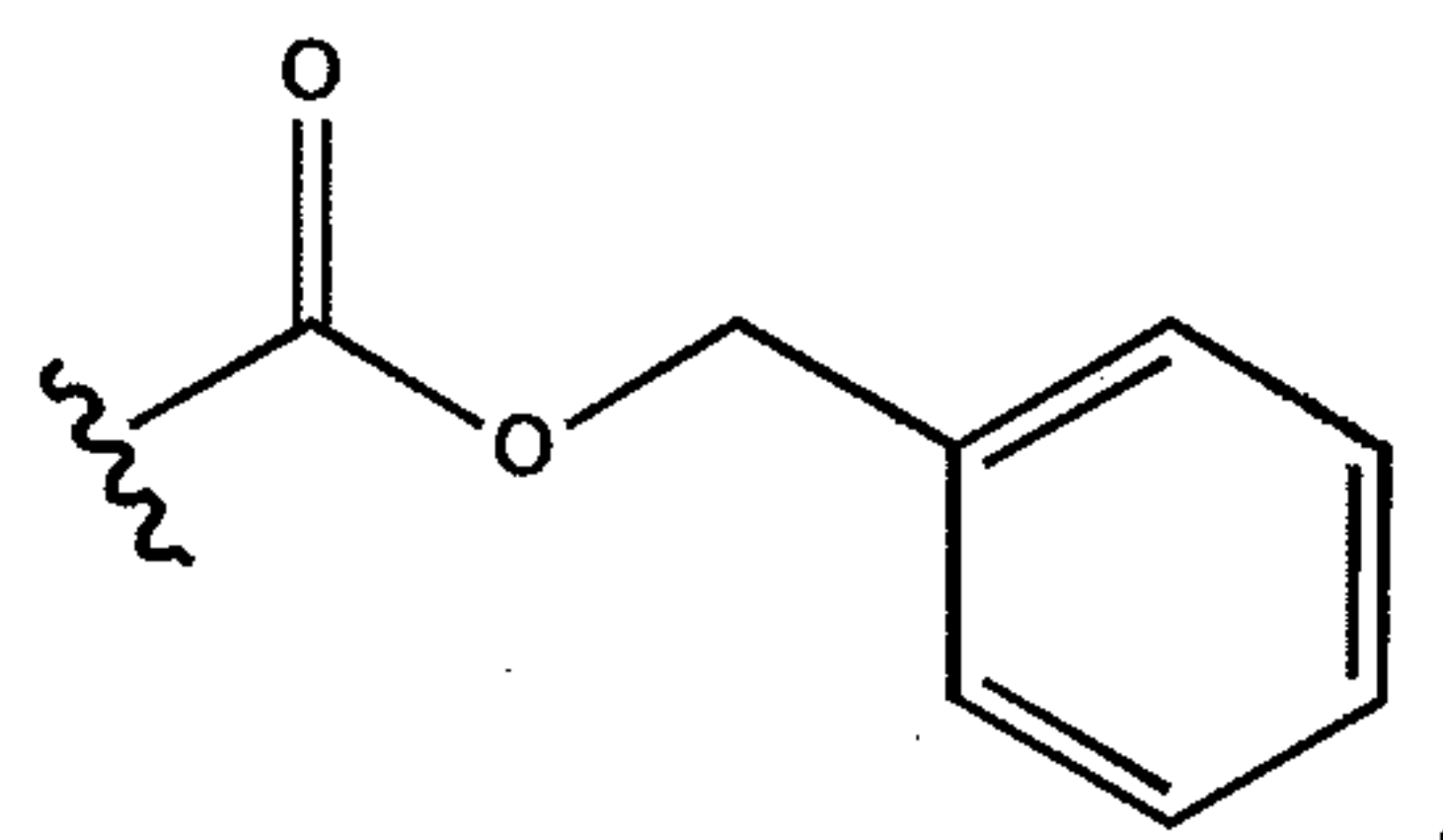
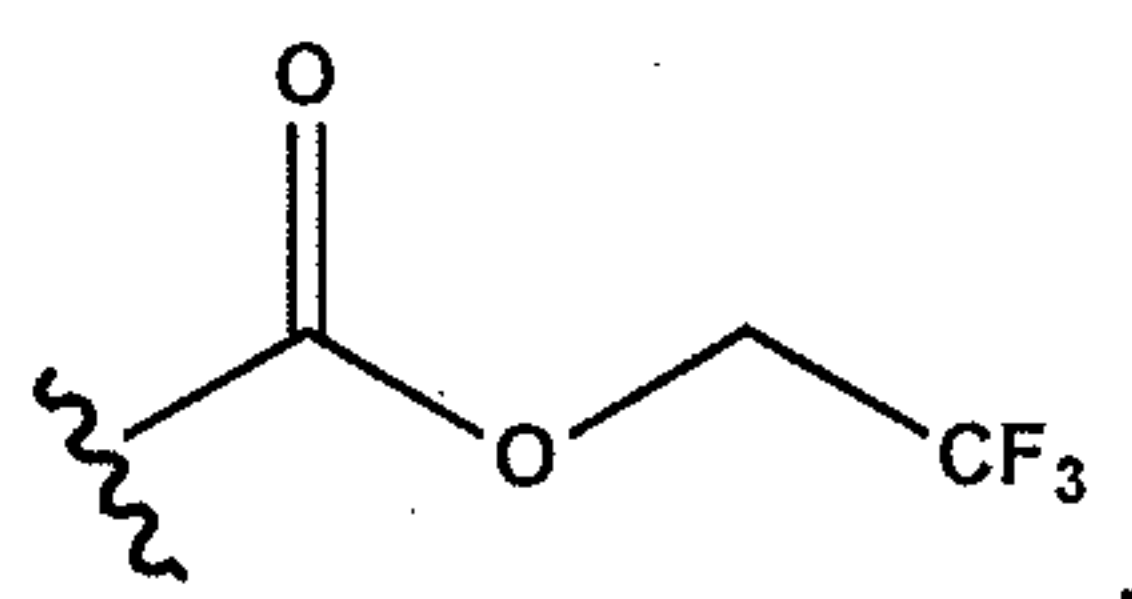
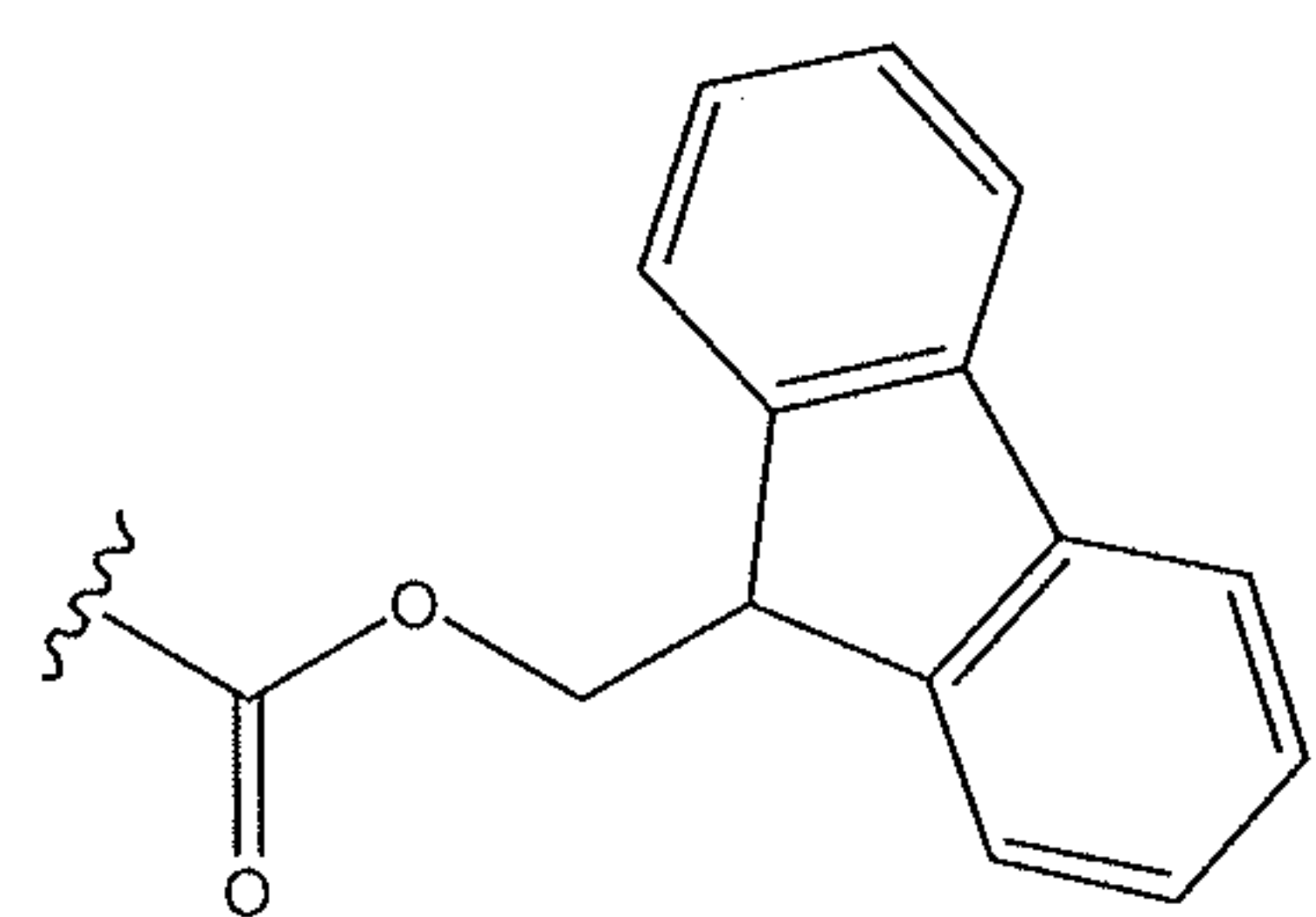
or



, and

the other of R_1 or R_2 is $-C(=O)OH$.

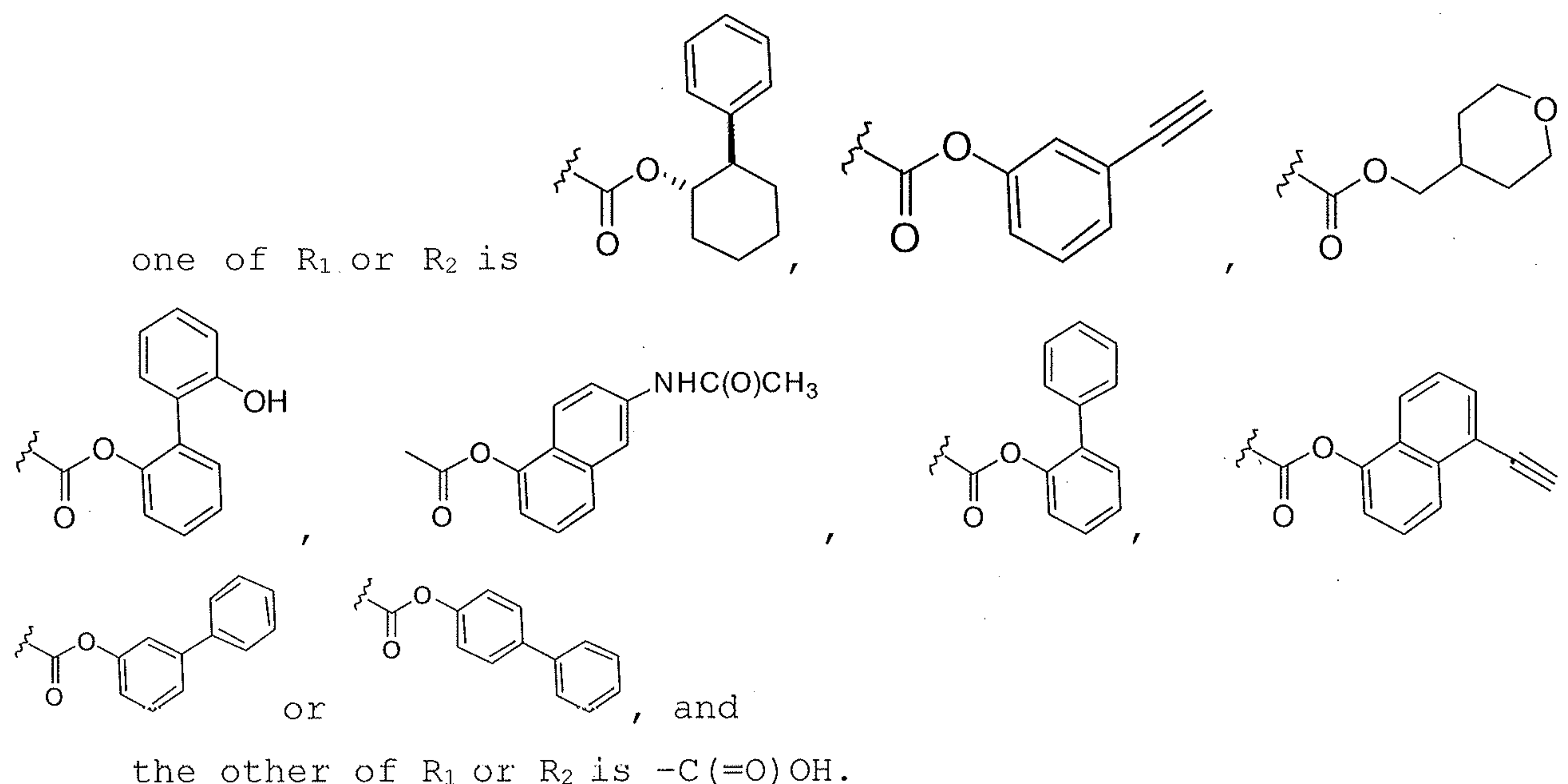
25. The compound of any one of claims 1-6,
 wherein
 one of R_1 or R_2 is



, and

the other of R_1 or R_2 is $-C(=O)OH$.

26. The compound of any one of claims 1-6,
wherein



27. The compound of any one of claims 1-26, wherein R₃, R₄, R₅, R₆, R₇, R₈, R₉, R₁₀, R₁₁ and R₁₂ are each independently, -H, halogen or -OR₁₅,
wherein R₁₅ is -H or C₁₋₁₀ alkyl.

28. The compound of any one of claims 1-26, wherein R₃, R₄, R₅, R₆, R₇, R₈, R₉, R₁₀, R₁₁ and R₁₂ are each independently, -H or -NO₂.

29. The compound of any one of claims 1-26, wherein one of R₃, R₄, R₅, R₆, R₇, R₈, R₉, R₁₀, R₁₁ and R₁₂ is other than -H.

30. The compound of any one of claims 1-26, wherein two of R₃, R₄, R₅, R₆, R₇, R₈, R₉, R₁₀, R₁₁ and R₁₂ are other than -H.

31. The compound of any one of claims 1-26, wherein four of R₃, R₄, R₅, R₆, R₇, R₈, R₉, R₁₀, R₁₁ and R₁₂ are other than -H.

32. The compound of any one of claims 1-26, wherein R₃, R₄, R₆, R₇, R₈, R₉, R₁₁ and R₁₂ are each -H; and R₅ and R₁₀ are each halogen or -OR₁₅,

wherein R₁₅ is -H, C₁₋₁₀ alkyl, C₂₋₁₀ alkenyl, C₂₋₁₀ alkynyl, aryl, or heteroaryl.

33. The compound of any one of claims 1-26, wherein R₄, R₅, R₆, R₉, R₁₀ and R₁₁ are each -H; R₃ and R₁₂ are each -H, halogen or -OR₁₅; and R₇ and R₈ are each halogen or -OR₁₅,

wherein R₁₅ is -H, C₁₋₁₀ alkyl, C₂₋₁₀ alkenyl, C₂₋₁₀ alkynyl, aryl, or heteroaryl.

34. The compound of any one of claims 1-26, wherein R₃, R₄, R₇, R₈, R₉ and R₁₂ are each -H; and R₅, R₆, R₁₀ and R₁₁ are each halogen or -OR₁₅,

wherein R₁₅ is -H, C₁₋₁₀ alkyl, C₂₋₁₀ alkenyl, C₂₋₁₀ alkynyl, aryl, or heteroaryl.

35. The compound of any one of claims 1-26, wherein R₃, R₄, R₅, R₆, R₈, R₉, R₁₀ and R_{1a} are each -H; and R₇ and R₁₂ are each NO₂.

36. The compound of claim 1,

wherein

one of R₁ or R₂ is -C(=O)OH and the other of R₁ or R₂ is -C(=O)OR₁₃ or -C(=O)O-alkyl-R₁₄,

wherein

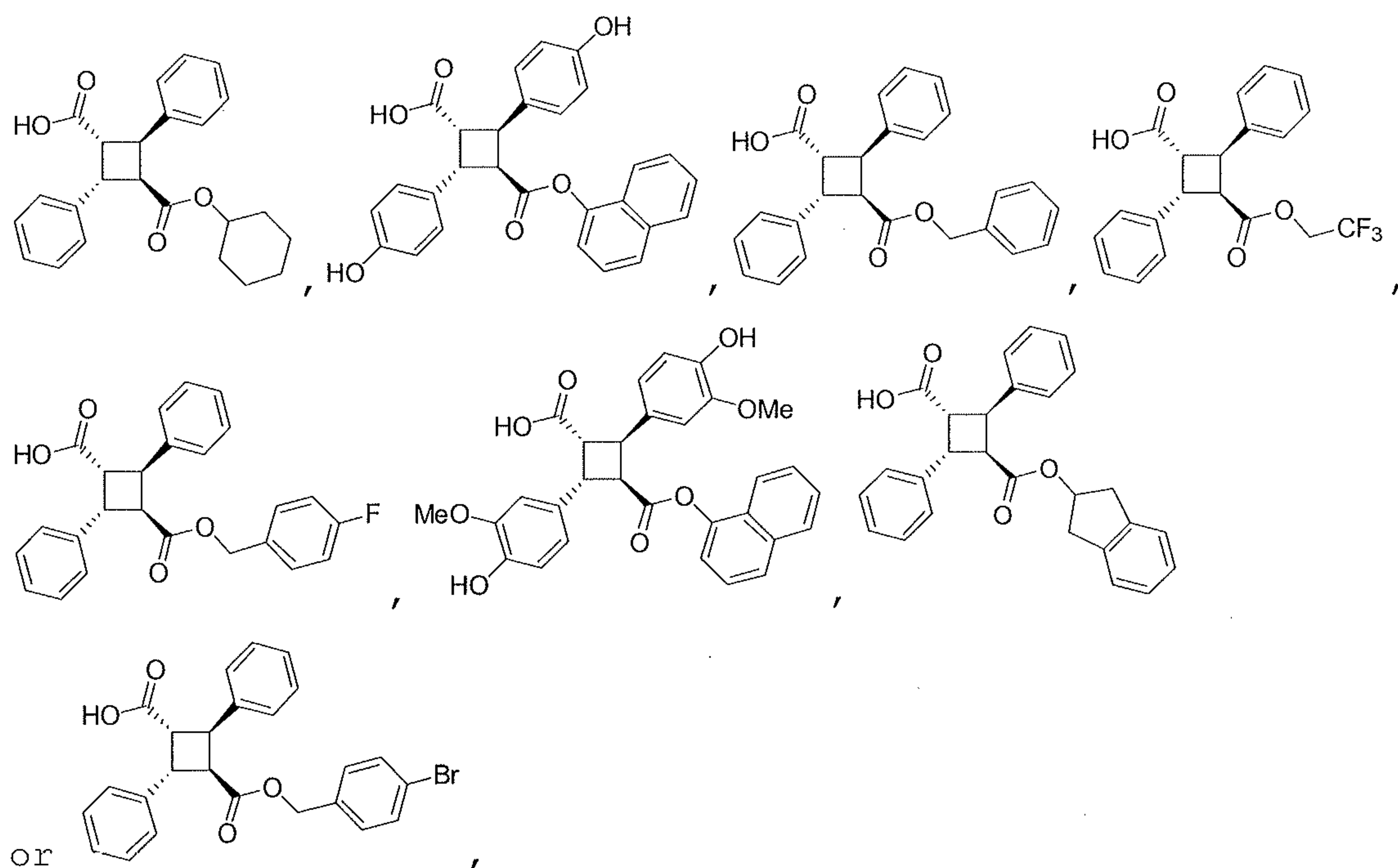
R₁₃ is cycloalkyl or aryl, and

R₁₄ is CF₃, cycloheteroalkyl or aryl; and

R₃, R₄, R₅, R₆, R₇, R₈, R₉, R₁₀, R₁₁ and R₁₂ are each independently, H or -OR₁₅,

wherein R₁₅ is H or C₁₋₁₀ alkyl.

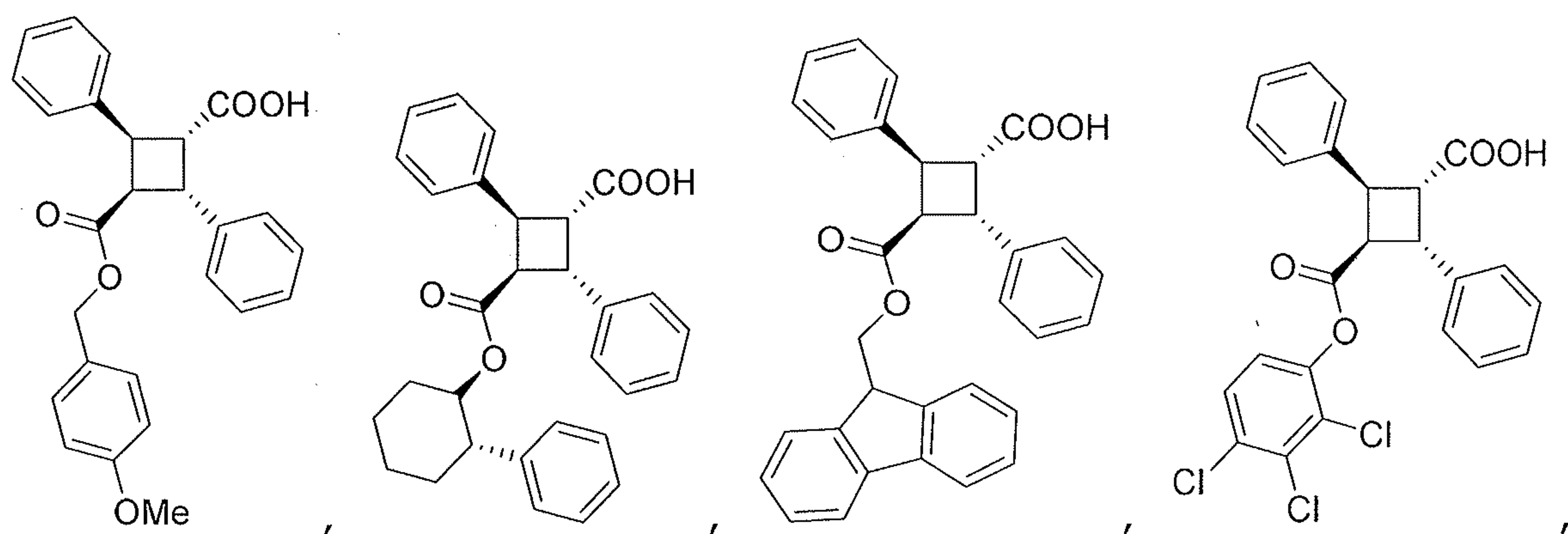
37. The compound of claim 1 having the structure:

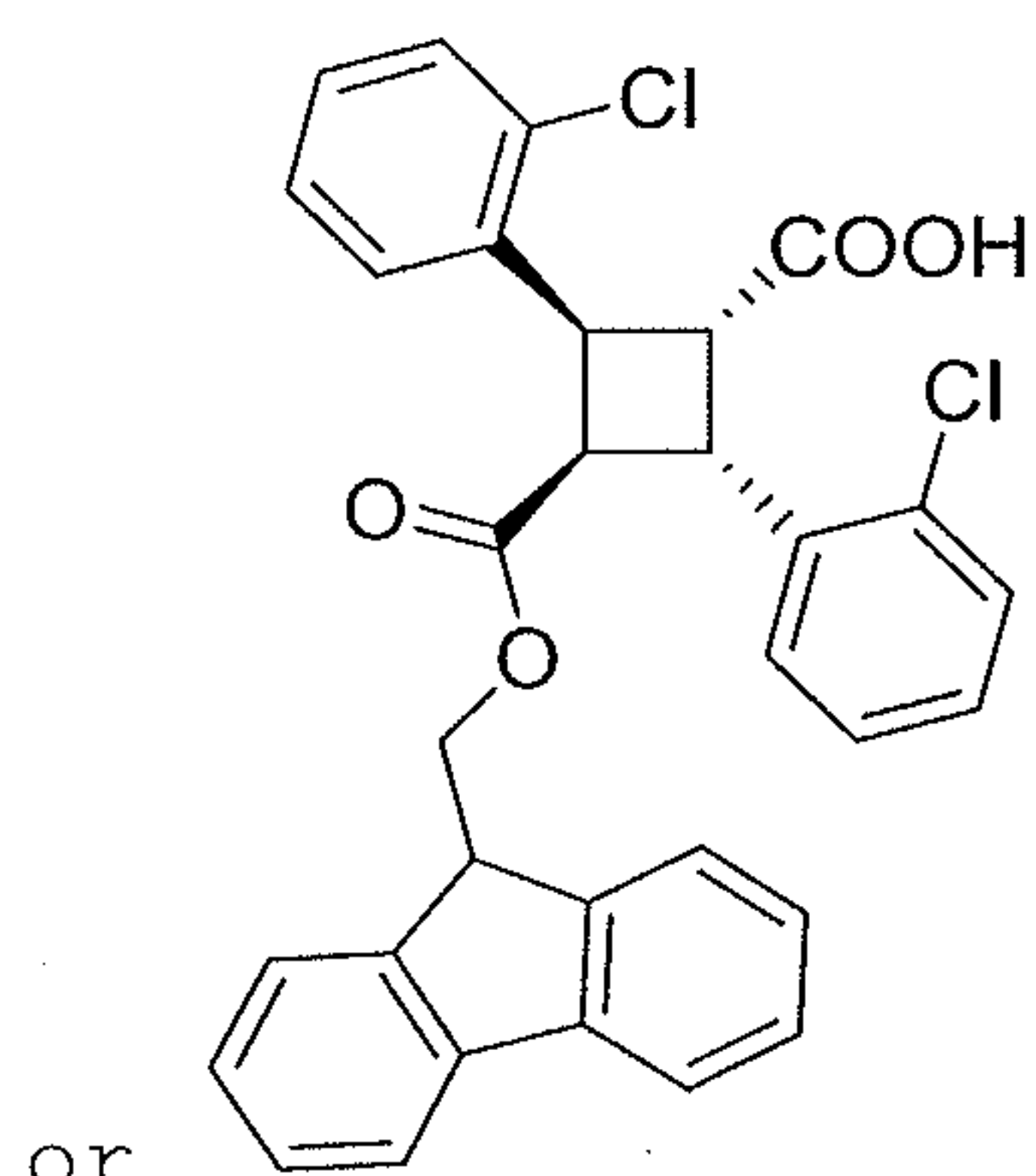
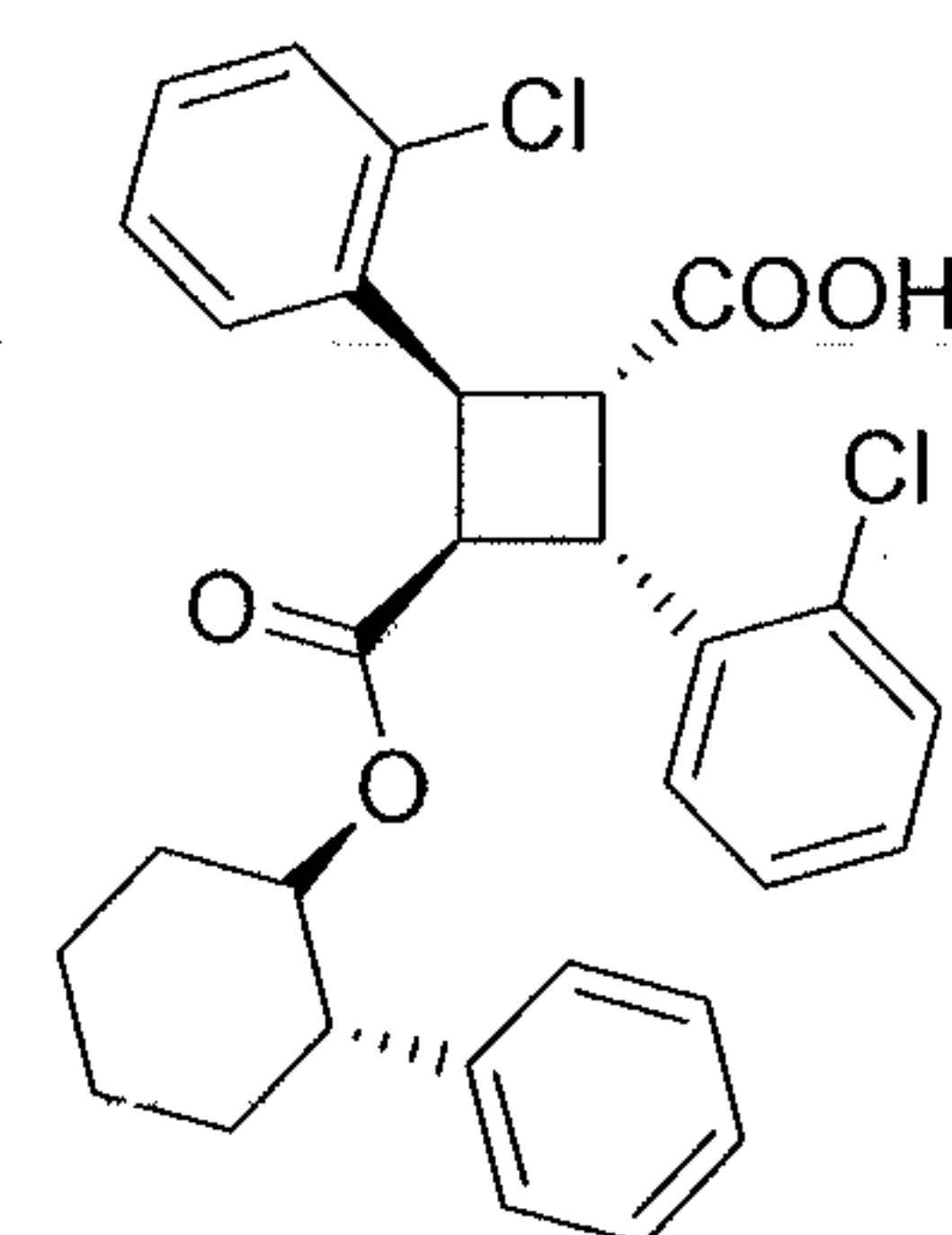
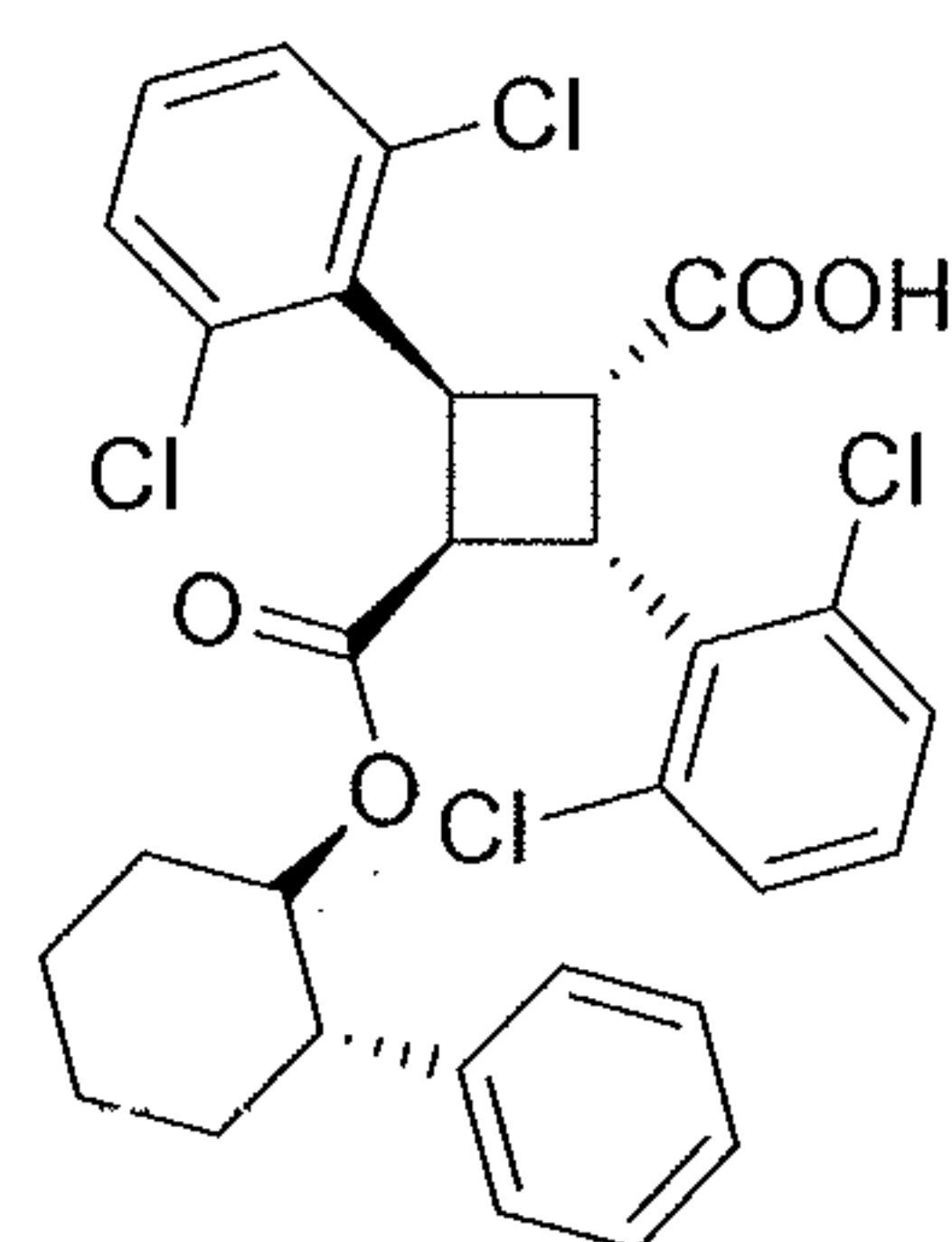
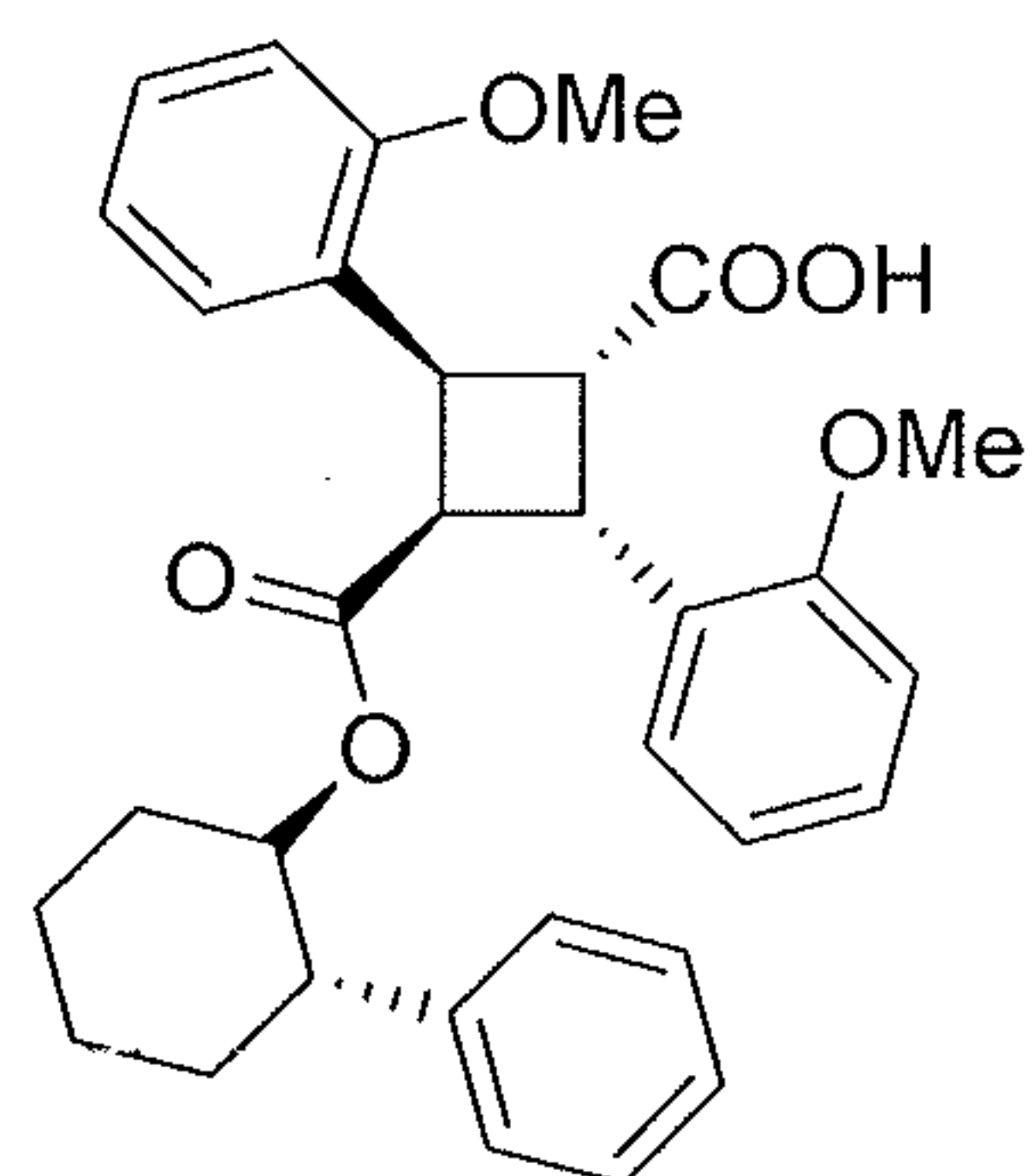
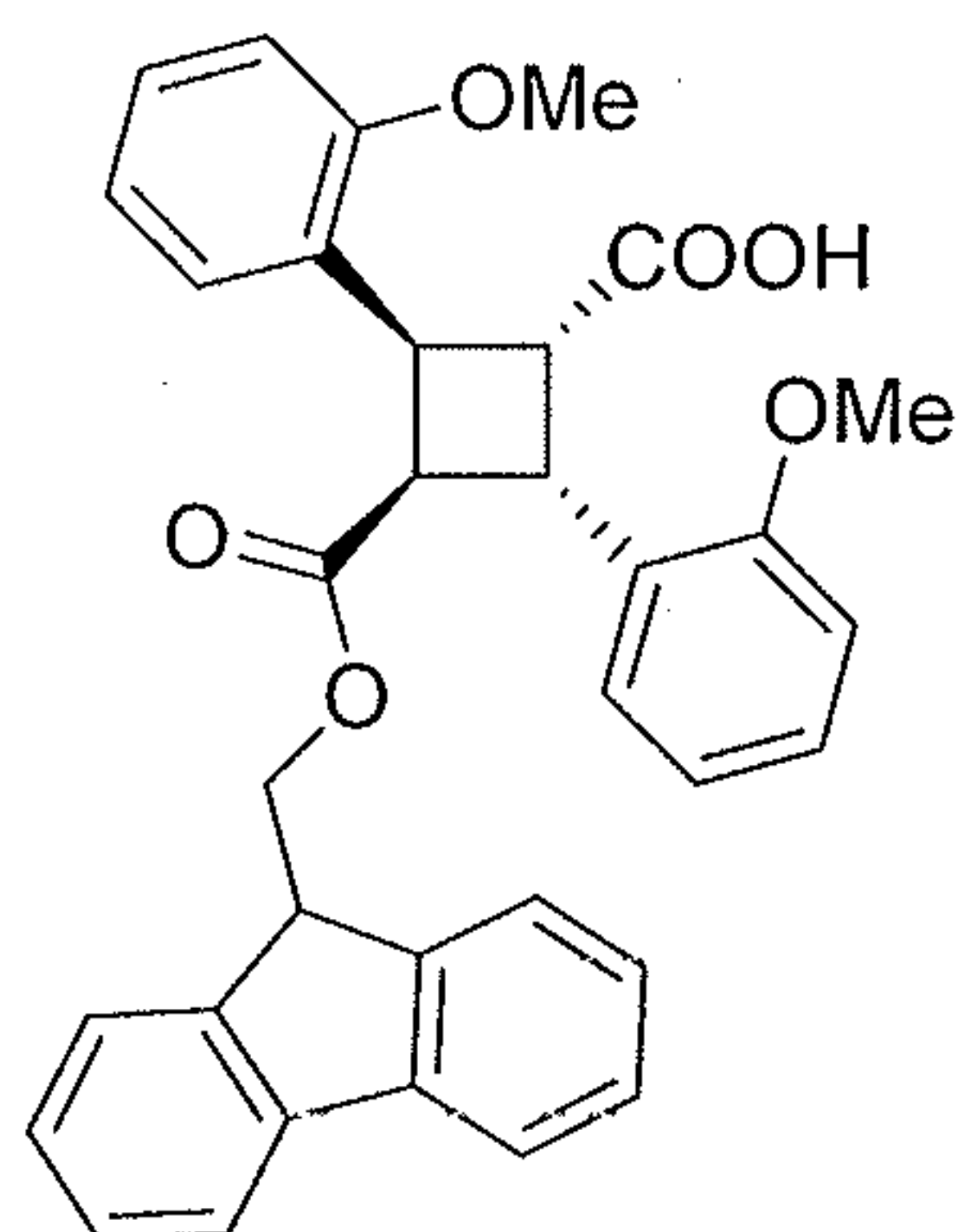
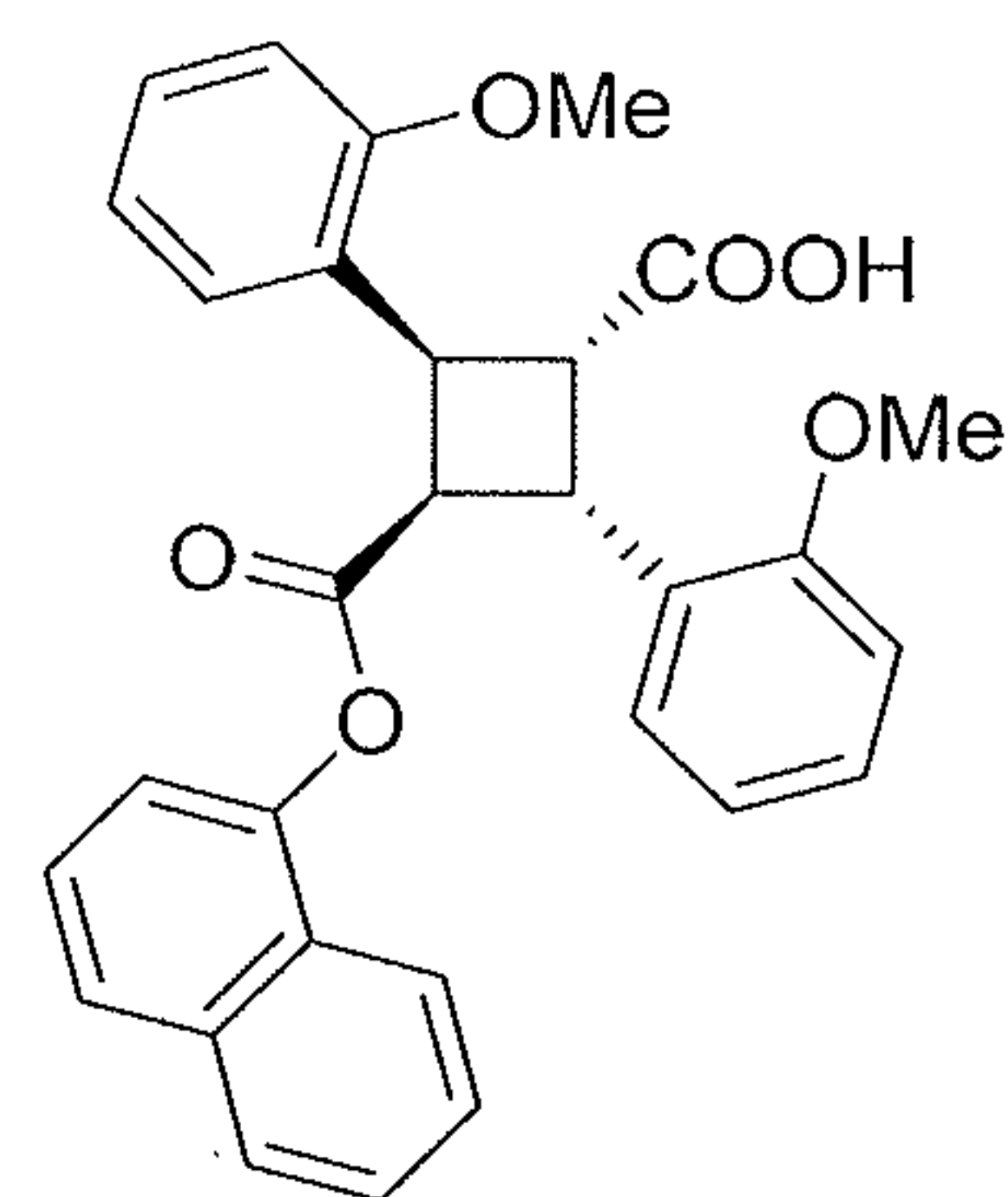
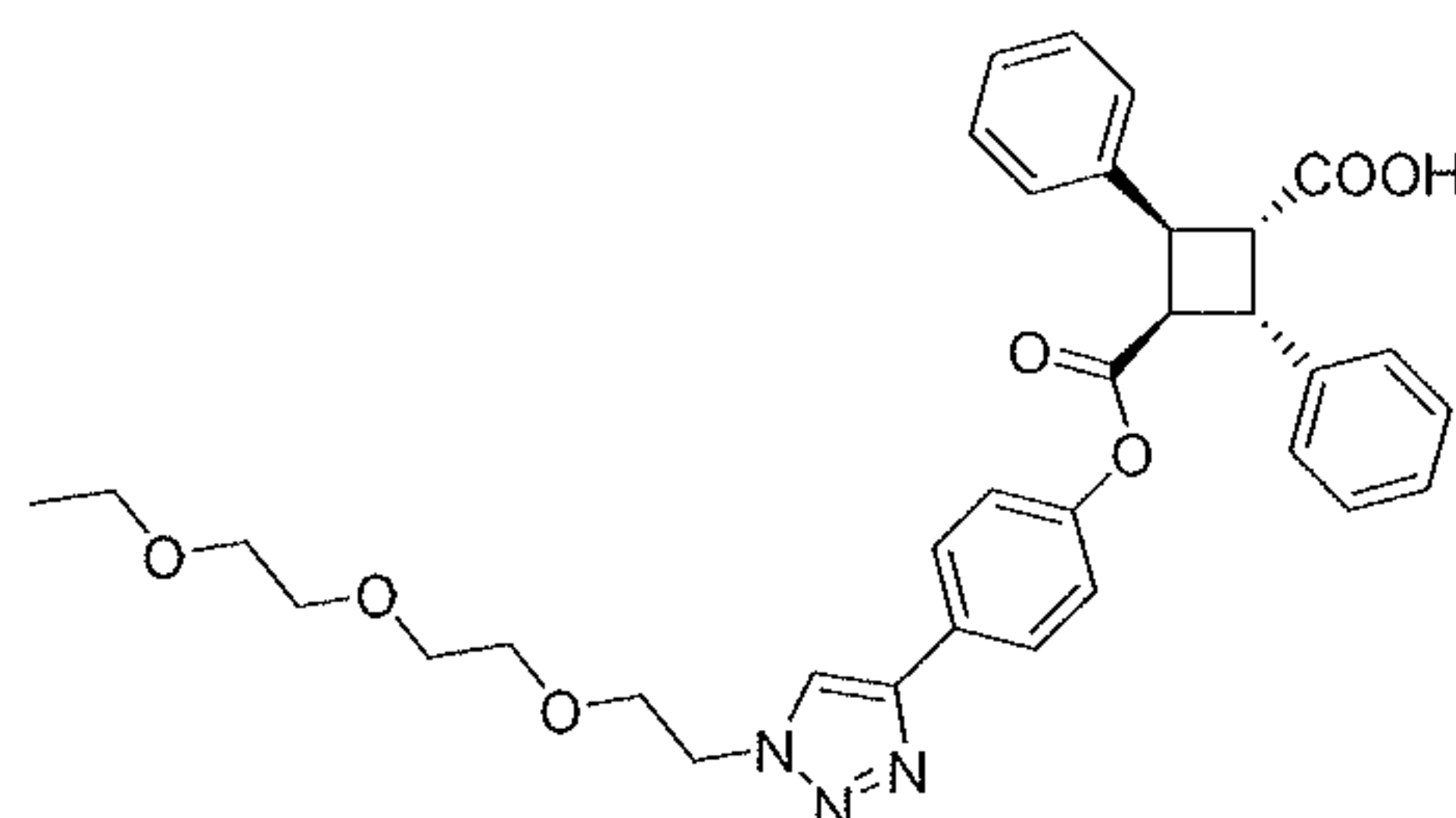
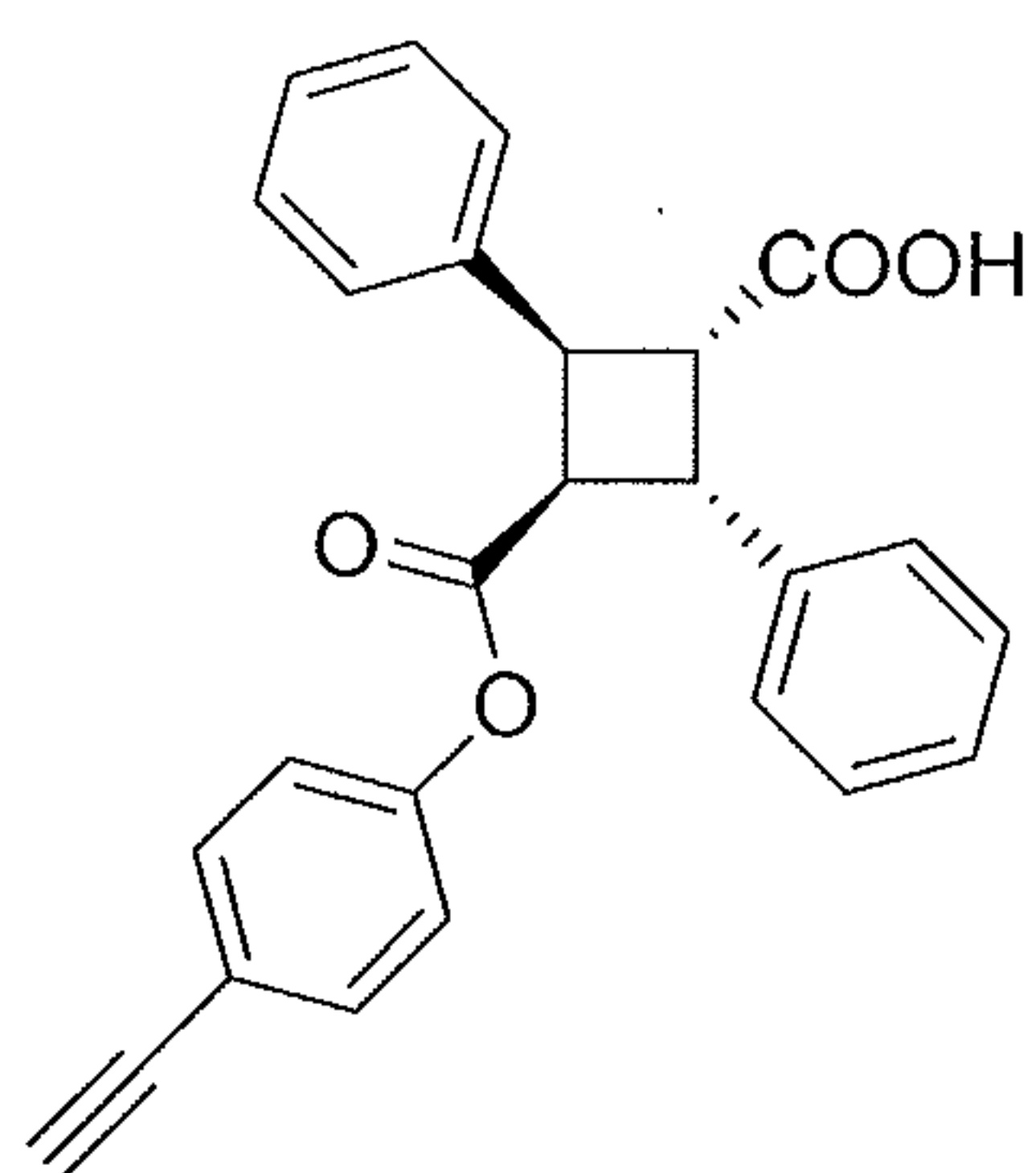


or an enantiomer or racemate thereof;

or a pharmaceutically acceptable salt thereof.

38. The compound of claim 1 having the structure:

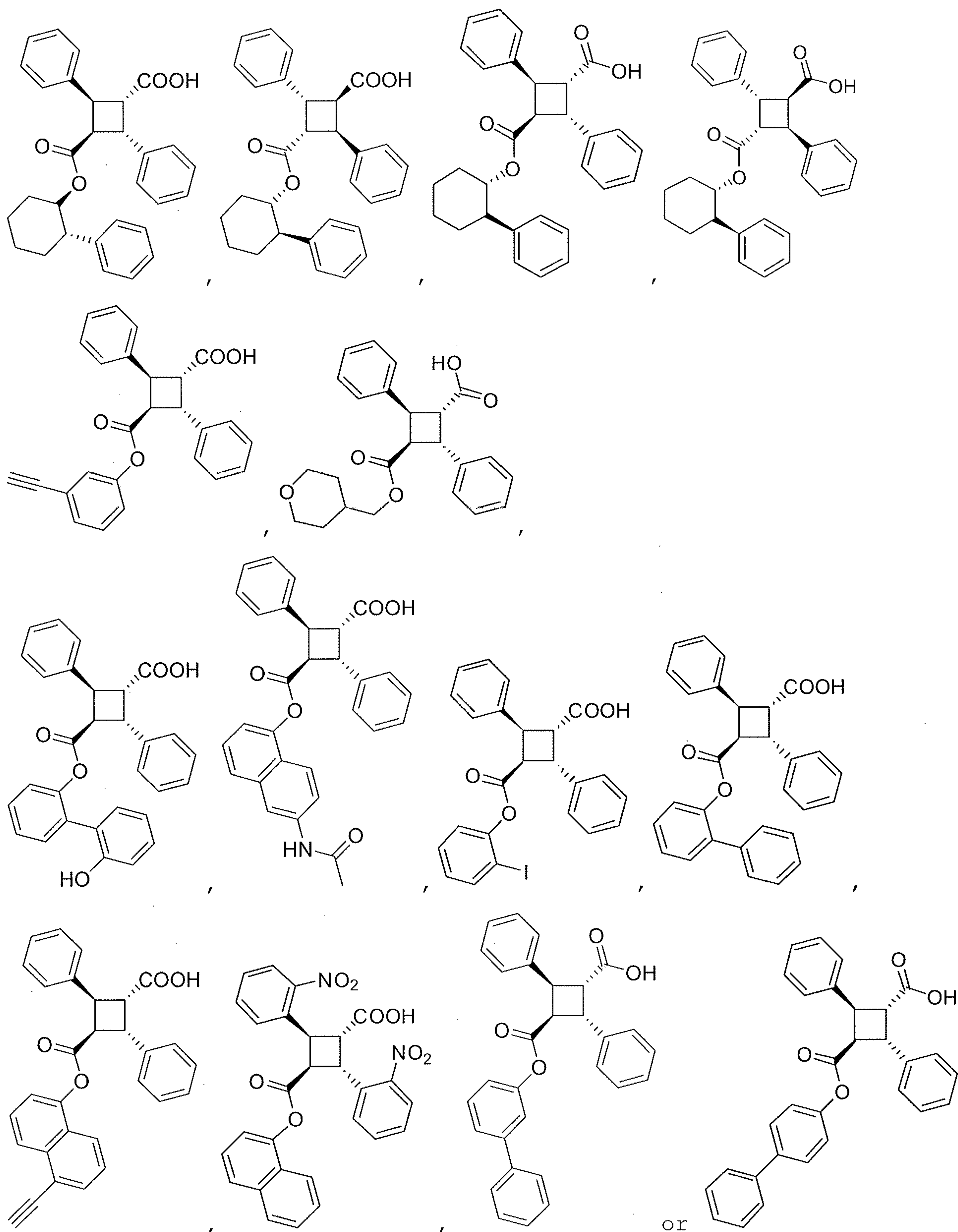




or enantiomer or racemate thereof;

or a pharmaceutically acceptable salt thereof.

39. The compound of claim 1 having the structure:



or enantiomer or racemate thereof;

or a pharmaceutically acceptable salt thereof.

40. The compound of any one of claims 1-39, wherein the compound is the (S,S) enantiomer.

41. The compound of any one of claims 1-39, wherein the compound is the (R,R) enantiomer.

42. A pharmaceutical composition comprising the compound of any one of claims 1-41 and a pharmaceutically acceptable carrier.

43. A method of inhibiting the activity of a Fatty Acid Binding Protein (FABP) comprising contacting the FABP with the compound of any one of claims 1-41.

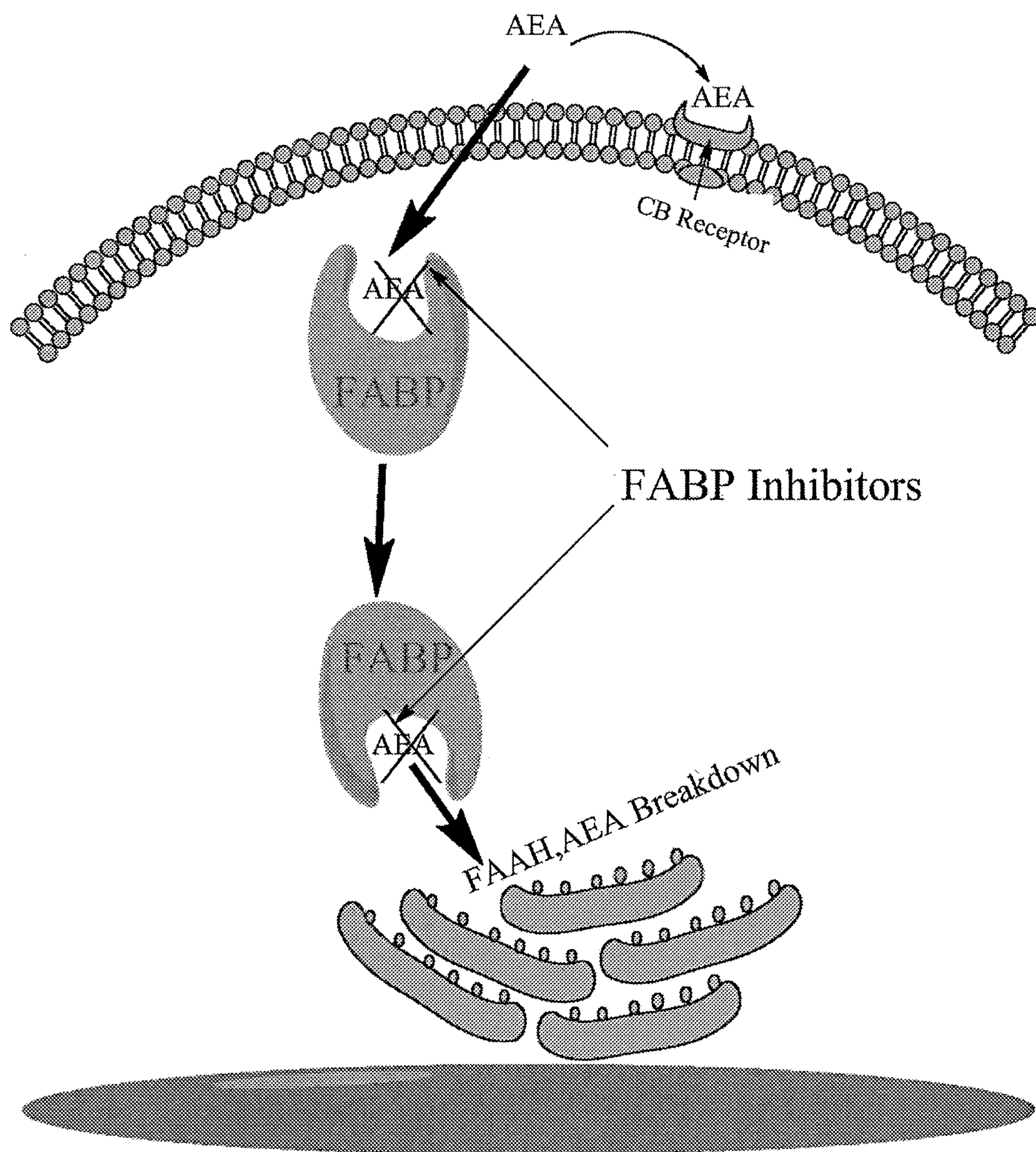
44. The method of claim 43, wherein the compound inhibits binding of an FABP ligand to the FABP.

45. The method of claim 44, wherein the FABP ligand is an endocannabinoid ligand.

46. The method of claim 44, wherein the FABP ligand is anandamide (AEA) or 2-arachidonoylglycerol (2-AG).

47. The composition of claim 42, wherein the composition comprises a mixture of enantiomers enriched in (S,S) enantiomer.

48. The composition of claim 42, wherein the composition comprises a mixture of enantiomers enriched in (R,R) enantiomer.



AEA (Anandamide) FAAH (fatty acid amide hydrolase)
FABP (fatty acid binding protein), CB (cannabinoid receptor)

Fig. 1

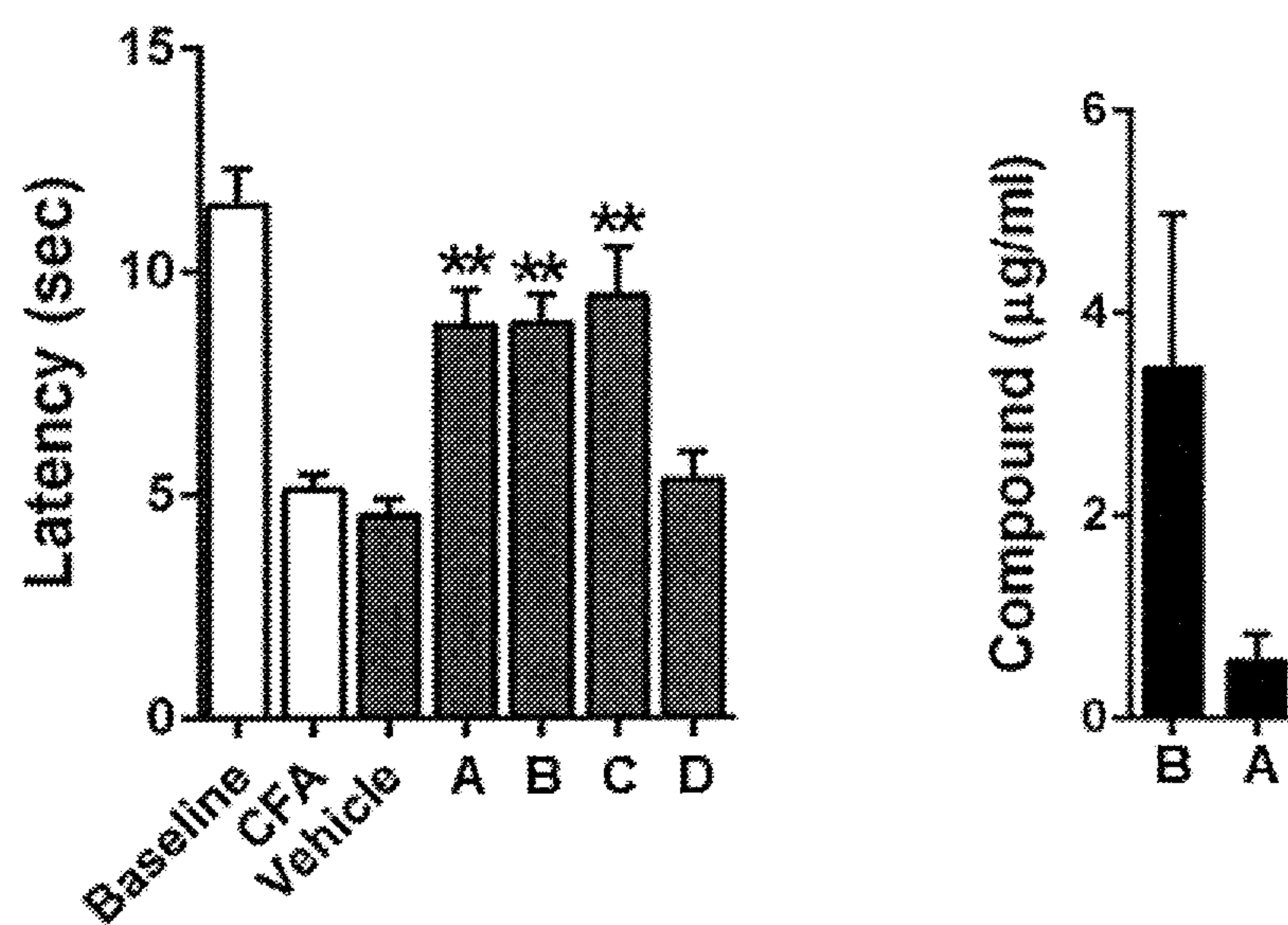


Fig. 2

