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WO-A1-96/30353
WO-A2-01/42216
WO-A2-01/90070
WO-A2-02/22611
WO-A2-02/42278
WO-A2-96/30395
WO-A2-2008/008264
US-A- 3 341 331
US-A- 3 640 716
US-A- 4 269 936
US-A- 5 976 858
US-B2- 7 256 198

Fortsættes ...

MAHESH UTTAMCHANDANI ET AL: "Activity-based fingerprinting and inhibitor discovery of cysteine proteases in a microarray", CHEMICAL COMMUNICATIONS, no. 15, 1 January 2007 (2007-01-01), page 1518, XP55023694, ISSN: 1359-7345, DOI: 10.1039/b702826a

SU LING NG ET AL: ""Click" synthesis of small-molecule inhibitors targeting caspases", ORGANIC & BIOMOLECULAR CHEMISTRY, vol. 6, no. 5, 1 January 2008 (2008-01-01) , page 844, XP55023693, ISSN: 1477-0520, DOI: 10.1039/b718304f

MERRIFIELD, R.B.: 'Solid phase peptide synthesis. I. The synthesis of a tetrapeptide' J. AM. CHEM. SOC. vol. 85, no. 14, 1963, pages 2149 - 2154, XP000994867

WINSSINGER N ET AL: "Profiling protien function with small molecule microarrays", PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES, NATIONAL ACADEMY OF SCIENCES, US, vol. 99, no. 17, 20 August 2002 (2002-08-20), pages 11139-11144, XP003012517, ISSN: 0027-8424, DOI: 10.1073/PNAS.172286899

TALANIAN R V ET AL: "Caspases as targets for anti-inflammatory and anti-apoptotic drug discovery", JOURNAL OF MEDICINAL CHEMISTRY, AMERICAN CHEMICAL SOCIETY, US, vol. 43, no. 18, 7 September 2000 (2000-09-07), pages 3351-3371, XP002223863, ISSN: 0022-2623, DOI: 10.1021/JM000060F

DESCRIPTION

FIELD OF INVENTION

[0001] The present invention relates to chemical compounds and their pharmaceutical uses. More particularly, the invention relates to selective inhibitors of caspases and their uses for the prevention and/or treatment of various diseases and conditions in subjects.

BACKGROUND OF INVENTION

[0002] Caspases comprise a family of cysteine protease enzymes with a well-known role as key mediators in apoptosis signaling pathways and cell disassembly. Interleukin converting enzyme (ICE), also known as Caspase-1, was the first identified caspase. In humans, 11 other known caspases have been further identified. Caspases have been classified in two general groups according to their effects: proapoptotic (caspase-2, 3, 6, 7, 8, 9, 10) and proinflammatory (caspase-1, 4, 5, 11, 12) caspases. The proapoptotic caspases have been divided in initiators (caspase-2, 8, 9, 10) also known as group II, and executioners (caspase-3, 6, 7) of the apoptotic process or group III. The Interleukin converting enzyme (ICE) also known as Caspase-1 has a proinflammatory role only.

[0003] There is growing evidence demonstrating the role of caspases in very diverse pathologies. For instance it is known that proapoptotic caspases are involved in the pathogenesis of many cardiovascular disorders. Some proapoptotic caspases such as caspase -8 also possess non-apoptotic function that may contribute to tumor progression. Caspase-1 plays an important role in response to pathogenic infection as well as in inflammatory and autoimmune disorders. In addition, caspase-1 activity is increased in retinas of diabetic patients and it constitutes a critical regulator of cardiomyocyte programmed cell death in the mammalian heart. Caspases also plays a role in neurodegenerative diseases and trauma. For instance, it has been shown that the caspase-3 cascade is highly activated due to the traumatic spinal cord injury. Finally, the activation of caspase-1 and caspase-3 in Amyotrophic Lateral Sclerosis (ALS) patients and the activation of caspase-7, -8, and -9 in a mouse model at end stage of ALS have been reported. Increased levels of apoptosis and caspase activity (especially caspase-3) are reported to be frequently observed at sites of cellular damage in both acute (e.g. Sepsis, myocardial infarction(MI), Ischemic Stroke, Spinal cord injury (SCI), traumatic Brain Injury (TBI)) and neurodegenerative disease (e.g. Alzheimer's, Parkinson's and Huntington's diseases, and multiple sclerosis (MS)).

[0004] Since caspases are involved in a number of diseases, several compounds and methods have been developed to inactivate them. For example, the broad irreversible caspase inhibitor benzyloxycarbonyl-Val-Ala-Asp-fluoromethylketone (z-VAD-fmk) was protective and efficiently blocked death receptor-mediated liver injury in animal models (Rodriguez et al. (1996), J Exp Med. 1996 Nov 1; 184(5):2067-72). Myocardial infarction and the resulting death of myocytes was ameliorated by z-VAD-fmk and related peptide inhibitors in animal models (Yaoita et al., 91998) Circulation 97: 276-281). There have been also a lot of efforts for identifying inhibitors of peptidase. For instance, Hanzlik and Thompson (J. Med. Chem. (1984), 27(6), 711-712) describe vinyllogous amino acid esters for inactivating thiol proteases. Thompson et al. (J. Med. Chem. (1986), 29(1), 104-111) describe carboxyl-modified amino acids and peptides as protease inhibitors. Liu and Hanzlik have prepared a series of peptidyl Michael acceptors with different electron withdrawing groups with different recognition and binding groups as inactivators against papain, a member of the cysteine proteinase family. Similarly, US patent No. 5,976,858 and 6,287,840 to Palmer et al. describes irreversible cysteine protease inhibitors containing vinyl groups conjugated to electron withdrawing groups. However, these and other compounds are not effective against caspases, because caspases are among the most specific endopeptidases.

[0005] Given their role in several diseases and conditions, there is a need for compounds capable of selectively targeting either a specific caspase or a group of caspases. There is also a need for effective pharmaceutical compositions and method of treatment for caspase-related diseases.

[0006] The present invention addresses these needs for novel therapies, new treatment methods, compounds, and pharmaceutical compositions.

[0007] Additional features of the invention will be apparent from review of the disclosure, figures and description of the invention below.

BRIEF SUMMARY OF THE INVENTION

[0008] The present invention relates to compounds according to any of Formula IA, II, IIA, III, or IIIA as defined herein, compositions thereof and methods for the prevention and/or treatment of caspase-related diseases in subjects. Particular aspects of the invention relates to use of compounds according to any of Formula IA, II, IIA, III, or IIIA as defined herein.

[0009] Also described herein are compounds according to any of Formula I, as defined herein, compositions thereof and methods for

the prevention and/or treatment of caspase-related diseases in subjects. Particular aspects of the invention relates to use of compounds according to Formula I, as defined herein.

[0010] One aspect of the invention concerns a method for preventing and/or treating a caspase-related disease in a subject in need thereof, comprising administering to said subject an effective amount of a compound represented by any of Formula IA, II, IIA, III, or IIIA as defined herein.

[0011] Also described herein is a method for preventing and/or treating a caspase-related disease in a subject in need thereof, comprising administering to said subject an effective amount of a compound represented by Formula I, as defined herein.

[0012] One aspect of the invention concerns the use of a compound represented by any of Formula IA, II, IIA, III, or IIIA as defined herein for preventing and/or treating of caspase-related diseases in a subject in need thereof.

[0013] Also described herein is the use of a compound represented by Formula I, as defined herein for preventing and/or treating of caspase-related diseases in a subject in need thereof.

[0014] Another related aspect of the invention concerns the use of a compound represented by any of Formula IA, II, IIA, III, or IIIA as defined herein for the manufacture of a medication for preventing and/or treating of caspase-related diseases in a subject in need thereof.

[0015] Also described herein is the use of a compound represented by Formula I, as defined herein for the manufacture of a medication for preventing and/or treating of caspase-related diseases in a subject in need thereof.

[0016] One aspect of the invention concerns a method of treating excessive apoptosis affected by caspase activity in a cell or a tissue, the method comprising: contacting the cell or tissue with an effective amount of one or more compounds represented by any of Formula IA, II, IIA, III, or IIIA as defined herein, so as to treat the excessive apoptosis.

[0017] Also described herein is a method of treating excessive apoptosis affected by caspase activity in a cell or a tissue, the method comprising: contacting the cell or tissue with an effective amount of one or more compounds represented by Formula I, as defined herein, so as to treat the excessive apoptosis.

[0018] One particular aspect of the invention concerns the use of a compound represented by any of Formula IA, II, IIA, III, or IIIA as defined herein for use in apoptosis mediated diseases.

[0019] Also described herein is the use of a compound represented by Formula I, as defined herein for use in apoptosis mediated diseases.

[0020] Caspase-related disease as defined herein are selected from the group consisting of apoptosis mediated diseases, IL-1 mediated diseases, inflammatory diseases, autoimmune diseases, autoinflammatory diseases, proliferative diseases, infectious diseases, degenerative diseases, retinal disorders, inflammatory peritonitis, osteoarthritis, pancreatitis, asthma, respiratory distress syndrome, rheumatoid arthritis, systemic lupus erythematosus, scleroderma, Grave's disease, autoimmune gastritis, diabetes, autoimmune hemolytic anemia, autoimmune neutropenia, thrombocytopenia, hepatitis, inflammatory bowel disease, crohn's disease, psoriasis, dermatitis, Graft vs host disease, organ transplant rejection, osteoporosis, leukemias and related disorders, multiple myeloma-related diseases, metastatic melanomas, Kaposi's sarcoma, sepsis, septic shock, Alzheimer's disease, Parkinson's disease, Huntington's disease, cerebral ischemia, epilepsy, myocardial ischemia, acute and chronic heart disease, myocardial infarction, congestive heart failure, atherosclerosis, spinal muscular atrophy, amyotrophic lateral sclerosis, multiple sclerosis, HIV-related encephalitis, aging, neurological damage due to stroke, ulcerative colitis, traumatic brain injury, spinal cord injury, hepatitis-B, hepatitis-C, hepatitis-G, liver-related diseases, renal disease, and HIV infection.

[0021] Specific examples of compounds according to the invention are represented in **Table 1**.

[0022] The invention also provides methods and strategies of targeting caspases. In one embodiment the approach consists of designing a suicide substrate leading to a permanent inhibition of the caspase. Preferably, the approach consists of designing a substrate that is recognizable enough for caspases, especially one or more specific caspase(s), to fit into it, to be potentially cleaved at a specific position in a way that makes the caspase enzyme irreversibly linked to the substrate thereby leading to a permanent inhibition of the caspase. In some embodiments, the suicide substrates of this invention are vinyl electron withdrawing group (EWG).

[0023] Further aspects of the invention will be apparent to a person skilled in the art from the following description, and claims and generalizations therein.

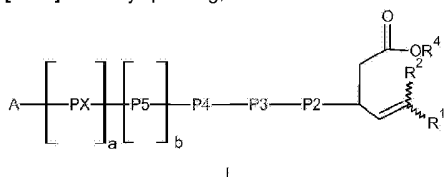
DETAILED DESCRIPTION OF THE INVENTION

A) General overview of the invention

[0024] The present inventors have discovered compounds that have beneficial pharmaceutical properties and that these compounds may be effective for use in caspase-mediated diseases such as sepsis, myocardial infarction, ischemic stroke, spinal cord injury (SCI), traumatic brain injury (TBI) and neurodegenerative disease (e.g. multiple sclerosis (MS) and Alzheimer's, Parkinson's, and Huntington's diseases).

B) Compounds of the invention

[0025] Broadly speaking, as described herein there is a compound represented by Formula I



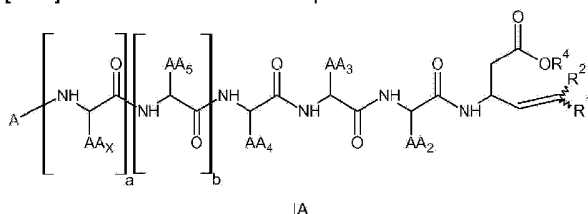
wherein A, PX, P5, P4, P3, P2, R¹, R², a and b are as defined hereinabove and hereinbelow;

or a prodrug, or a pharmaceutically acceptable salt to allow the drug to penetrate the cell membrane or the compound is labeled with a detectable label or an affinity tag thereof.

[0026] The line "-" when located between P2, P3, P4, P5 and PX represents a peptide bond or a peptidomimetic bond; The PX, P5, P4, P3, P2 amino acid residues are normally linked via a peptide bond, that is, a peptidic carbamoyl group, i.e. --CONH--. However, peptidomimetic bonds are also contemplated, such as CH₂-NH, CO-CH₂, azapeptide and retro-inverso bonds.

[0027] The R¹ and R² that are bonded to the vinyl group can be either in the cis configuration or the trans configuration, as represented by the wavy lines. In one example, R¹ is configured to be trans such that the electron withdrawing capability of the R¹ group is stabilized.

[0028] The invention concerns compounds of Formula IA :

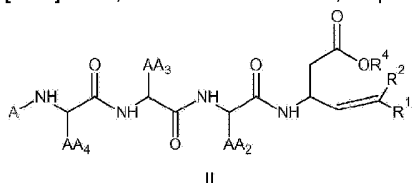


wherein A, AA_x, AA₅, AA₄, AA₃, AA₂, R¹, R², a and b are as defined hereinabove and hereinbelow;

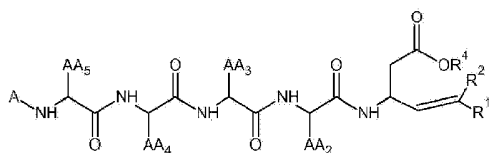
or a pharmaceutically acceptable salt to allow the drug to penetrate the cell membrane, or the compound is labeled with a detectable label or an affinity tag thereof.

[0029] Also described herein are prodrugs of the compounds of the invention.

[0030] Thus, when a and b are both 0, the present invention includes compounds of Formula II :

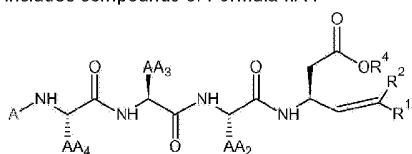


wherein A, AA₄, AA₃, AA₂, R¹, and R² are as defined hereinabove and hereinbelow. Furthermore, when a is 0 and b is 1, the present invention includes compounds of Formula III



III

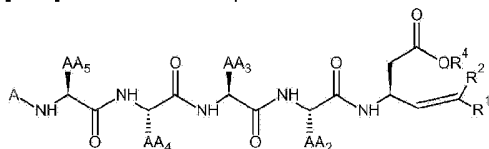
wherein A, AA₅, AA₄, AA₃, AA₂, R¹, and R² are as defined hereinabove and hereinbelow. One subset of compounds of Formula II includes compounds of Formula IIA :



IIA

wherein A, AA₄, AA₃, AA₂, R¹, and R² are as defined hereinabove and hereinbelow.

[0031] One subset of compounds of Formula III includes compounds of Formula IIIA :



IIIA

wherein A, AA₅, AA₄, AA₃, AA₂, R¹, and R² are as defined hereinabove and hereinbelow.

a and b:

[0032] In one subset of compounds of the invention, a is 0 or 1; and b is 0 or 1 provided that when b is 0, a is 0.

[0033] In one example, a and b are both 0.

[0034] In another example, a is 0 and b is 1.

1. A:

In one subset, A is

1. 1) H,
2. 2) C₁-C₆ alkyl,
3. 3) aryl,
4. 4) heteroaryl,
5. 5) heterocyclyl,
6. 6) PhCH₂OC(O)-;
7. 7) R³-C(O)O-, or
8. 8) R³-S(O)₂-;

wherein R³ is

1. 1) C₁-C₆ alkyl,
2. 2) aryl,
3. 3) heteroaryl, or
4. 4) heterocyclyl;.

[0035] In one example, A is H.

[0036] In one example, A is R³-OC(O)-.

[0037] In one example, A is $\text{PhCH}_2\text{OC(O)-}$.

R¹:

In one subset, R¹ is an electron withdrawing group (EWG) selected from

1. 1) aryl,
2. 2) heteroaryl,
3. 3) heterocyclyl,
4. 4) C₂-C₆ alkene-R²⁰,
5. 5) SO₂R⁵,
6. 6) SO₃R⁵,
7. 7) SO₂R⁵,
8. 8) SONHR⁵,
9. 9) SO₂NHR⁵,
10. 10) CN,
11. 11) CO₂R⁵,
12. 12) COR⁵,
13. 13) PO₃R⁵,
14. 14) PO(OR⁵)₂, or
15. 15) PO(OR⁵),

wherein the aryl, the heteroaryl, or the heterocyclyl are optionally substituted with one or more R³⁰.

R²:

In one subset, R² is

1. 1) R¹; or
2. 2) H,
3. 3) halogen,
4. 4) haloalkyl,
5. 5) C₁-C₆ alkyl,
6. 6) C₂-C₆ alkene,
7. 7) C₃-C₇ cycloalkyl,
8. 8) OR⁹;
9. 9) OCOR⁶,
10. 10) OCO₂R⁶,
11. 11) NR⁷R⁸,
12. 12) NH₂SO₂R⁶,
13. 13) NHCOR⁶,
14. 14) aryl,
15. 15) heteroaryl, or
16. 16) heterocyclyl;

wherein R¹, R⁶, R⁷, R⁸ and R⁹ are as defined hereinabove and hereinbelow.

[0038] In one example, R² is H.

[0039] In another example, R² is halogen.

[0040] In yet another example, R² is Cl.

R⁴:

In one subset, R^4 is

- 1) H, or
- 2) methyl, ethyl, propyl, or tert-butyl.

In one example, R^4 is H.

R^5 :

In one subset, R^5 is

- 1) H,
- 2) C_1 - C_6 alkyl,
- 3) C_2 - C_6 alkene,
- 4) C_3 - C_7 cycloalkyl,
- 5) aryl,
- 6) heteroaryl,
- 7) heterocyclyl, or
- 8) any optionally protected (D) or (L) amino acid residue.

R^6 :

In one subset, R^6 is

- 1) any (D) or (L) amino acid residue,
- 2) C_1 - C_6 alkyl,
- 3) C_3 - C_7 cycloalkyl,
- 4) aryl,
- 5) heteroaryl, or
- 6) heterocyclyl,

in which the alkyl or the cycloalkyl are optionally substituted with one or more R^{10} substituents; and in which the aryl, heteroaryl or heterocyclyl are optionally substituted with one or more R^{20} substituents.

R^7 and R^8 :

In one subset, R^7 and R^8 are independently selected from:

- 1) H,
- 2) C_1 - C_6 alkyl,
- 3) C_3 - C_7 cycloalkyl,
- 4) haloalkyl,
- 5) aryl,
- 6) heteroaryl, or
- 7) heterocyclyl,

wherein the alkyl and the cycloalkyl are optionally substituted with one or more R^{10} substituents, and the aryl, the heteroaryl and the heterocyclyl are optionally substituted with one or more R^{20} substituents.

R^9 :

In one subset, R^9 is

- 1) H,
- 2) C_1 - C_6 alkyl,
- 3) C_3 - C_7 cycloalkyl,
- 4) aryl,
- 5) heteroaryl, or
- 6) heterocyclyl,

in which the alkyl or the cycloalkyl are optionally substituted with one or more R^{10} substituents; and in which the aryl, heteroaryl or

heterocyclyl are optionally substituted with one or more R^{20} substituents.

R^{10} :

In one subset, R^{10} is independently selected from:

1. 1) halogen,
2. 2) C_1 - C_6 alkyl,
3. 3) C_3 - C_7 cycloalkyl,
4. 4) haloalkyl,
5. 5) aryl,
6. 6) heteroaryl,
7. 7) heterocyclyl,
8. 8) OR^9 ,
9. 9) $S(O)_mR^9$,
10. 10) NR^7R^8 ,
11. 11) COR^9 ,
12. 12) $C(O)OR^9$,
13. 13) $OC(O)R^9$,
14. 14) $SC(O)R^9$,
15. 15) $CONR^7R^8$, or
16. 16) $S(O)_2NR^7R^8$;

wherein R^7 , R^8 , R^9 are as defined hereinabove and hereinbelow;

m is an integer of 0, 1, or 2.

R^{20} :

In one subset, R^{20} is independently selected from:

1. 1) halogen,
2. 2) NO_2 ,
3. 3) CN,
4. 4) C_1 - C_6 alkyl,
5. 5) haloalkyl,
6. 6) C_3 - C_7 cycloalkyl,
7. 7) OR^7 ,
8. 8) NR^7R^8 ,
9. 9) SR^7 ,
10. 10) aryl,
11. 11) heteroaryl,
12. 12) heterocyclyl,
13. 13) SO_2R^5 ,
14. 14) SO_3R^5 ,
15. 15) SOR^5 ,
16. 16) $SONHR^5$,
17. 17) SO_2NHR^5 ,
18. 18) PO_3R^5 ,
19. 19) $PO(OR^5)_2$,
20. 20) $PO(OR^5)$,
21. 21) COR^7 ,
22. 22) CO_2R^7 ,
23. 23) $S(O)_mR^7$,
24. 24) $CONR^7R^8$, or
25. 25) $S(O)_2NR^7R^8$,

wherein the alkyl and the cycloalkyl are optionally substituted with one or more R^6 substituents; and wherein the aryl, the heteroaryl, or the heterocyclyl are optionally substituted with one or more R^{30} .
wherein R^7 , R^8 and m are as defined hereinabove and hereinbelow.

R^{30} :

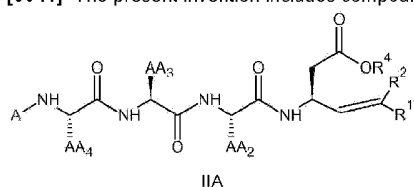
In one subset, R^{30} is

1. 1) NO_2 ,
2. 2) C_2-C_6 alkene- R^{20} ,
3. 3) SO_2R^5 ,
4. 4) SOR^5 ,
5. 5) $SONHR^5$,
6. 6) SO_2NHR^5 ,
7. 7) CN ,
8. 8) CO_2R^5 ,
9. 9) COR^5 ,
10. 10) PO_3R^5 ,
11. 11) $PO(OR^5)_2$, or
12. 12) $PO(OR^5)$;

wherein R^5 and R^{20} are as defined hereinabove and hereinbelow.

Caspase 3 inhibitors

[0041] The present invention includes compounds of Formula IIA:



wherein

AA_2 is the amino acid side chain of Val, Leu, Pro, Met, Ala, Thr, His.

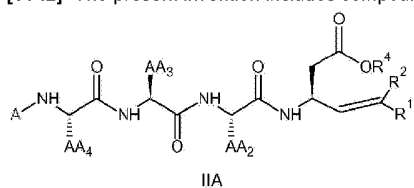
AA_3 is the amino acid side chain of Trp, Tyr, Ala, Asp, Glu, Gln, Phe, Ser, Thr, Val, Tyr, Gly, Leu; or AA_3 is phenylglycine, indanylglycine, or Ala-(2'-quinolyl);

AA_4 is the amino acid side chain of Asp;

and wherein A, R^1 , R^2 and R^4 are as defined hereinabove and hereinbelow.

Caspase 8/Caspase 9 inhibitors

[0042] The present invention includes compounds of Formula IIA:



wherein

AA₂ is the amino acid side chain of Thr, His, Val, Trp, Ile, or Ala

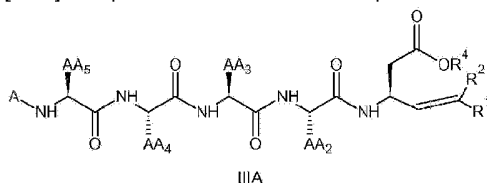
AA₃ is the amino acid side chain of Glu or AA₃ is Ala-(2'-quinolyl);

AA₄ is the amino acid side chain of Ile, Leu, Glu, Asp, Ala, Pro or Val;

and wherein A, R¹, R² and R⁴ are as defined hereinabove and hereinbelow.

Caspase 2 inhibitors

[0043] The present invention includes compounds of Formula IIIA



wherein

AA₂ is the amino acid side chain of Ala, Ser, Lys or Val;

AA₃ is the amino acid side chain of Val, Glu, Thr, or Gln;

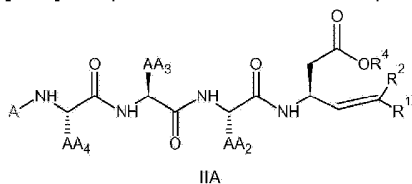
AA₄ is the amino acid side chain of Asp, or Leu;

AA₅ is the amino acid side chain of Val or Leu;

and wherein A, R¹, R² and R⁴ are as defined hereinabove and hereinbelow.

Caspase 1 inhibitors

[0044] The present invention includes compounds of Formula IIA (caspase 1 inhibitors)



wherein

AA₂ is the amino acid side chain of Val, Ala, Thr, or His;

AA₃ is the amino acid side chain of Glu, Gln, Asp, Ala, Gly, Thr, Val, Trp; or AA₃ is phenylglycine or indanylglycine;

AA₄ is the amino acid side chain of Tyr, Trp, Phe, or Asp;

and wherein A, R¹, R² and R⁴ are as defined hereinabove and hereinbelow.

[0045] Compounds and intermediate compounds synthesized according to the present invention include those in Table 1 :

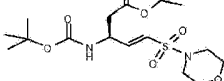
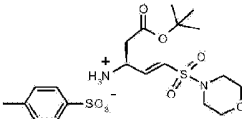
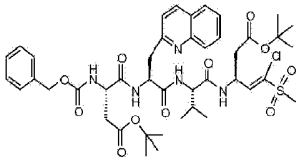
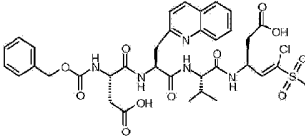
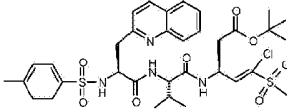
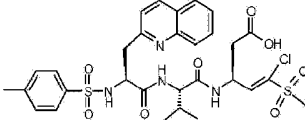
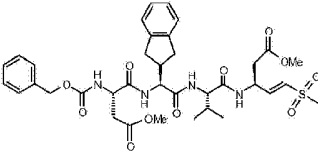
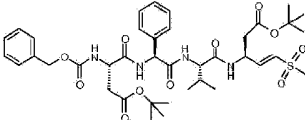
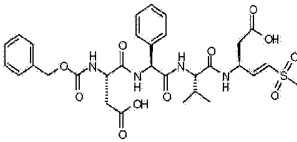
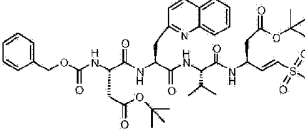
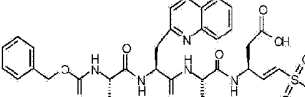
TABLE 1

CPD N°	CPD NAME	STRUCTURE
1	Fmoc-Ala(2'-quinolyl)-Val-OAllyl	
2	Ala(2'quinolyl)-Val-OAllyl	
3	Cbz-Asp(O-tBu)-Ala(2'-quinolyl)-Val-OAllyl	
4	Cbz-Asp(O-tBu)-Ala(2'-quinolyl)-ValOH	
5	Ts-Ala(2'-quinolyl)-Val-OH	
6	Ts-Ala(2'-quinolyl)-OH	
7	Ts-Ala(2'-quinolyl)-Val-OAllyl	
8	Z-Asp(OtBu)-Tyr(OtBu)-Val-Asp(OtBu)methyl vinyl sulfone	
9	Fmoc-Indanylglycine-Val-OAllyl	
10	Indanylglycine-Val-OAllyl	
11		

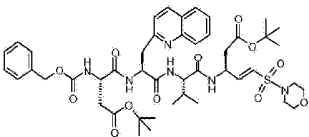
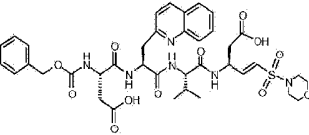
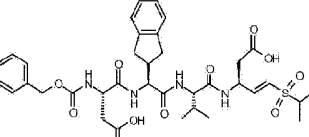
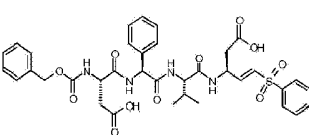
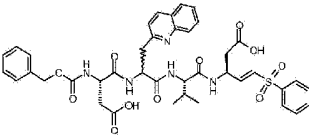
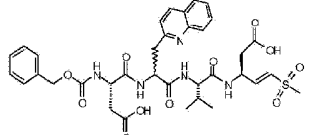
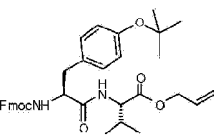
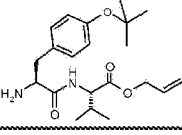
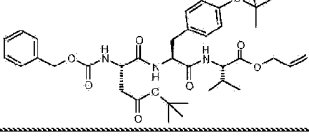
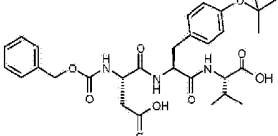
CPD N°	CPD NAME	STRUCTURE
	Cbz-Asp(O-tBu)-Indanylglycine-Val-OAllyl	
12	Cbz-Asp(O-tBu)-Indanylglycine-Val-OH	
13	Fmoc-Phg-Val-OAllyl	
14	Phg-Val-OAllyl	
15	Z-Asp(β-tert-Butyl)- Phg-Val-OAllyl.	
16	Z-Asp(β-tert-Butyl)-Phg-Val-OH.	
17	Fmoc-Glu(O-tBu)-Val-OAllyl	
18	Glu(O-tBu)-Val-OAllyl	
19	Cbz-Asp(O-tBu)-Glu(O-tBu)-Val-OAllyl	
20	Cbz-Asp(O-tBu)-Glu(O-tBu)-Val-OH	
21	Diethyl chloro(methylsulfone) methylphosphonate	
22		

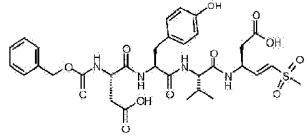
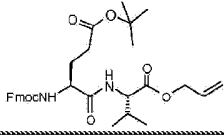
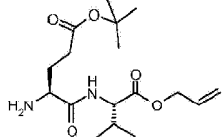
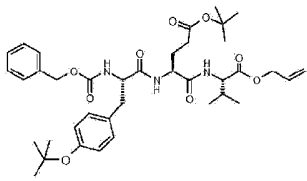
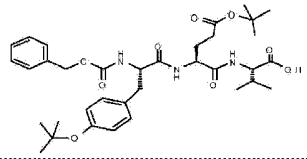
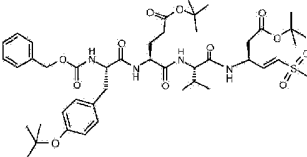
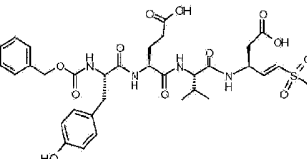
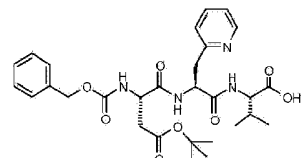
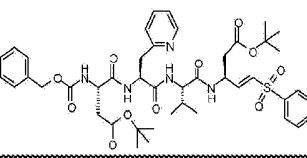
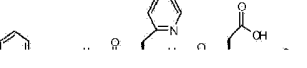
CPD N°	CPD NAME	STRUCTURE
	Boc-Asp (β-tert-butyl) achlorovinyl methylsulfone	
23	Asp(β-tert-butyl)achlorovinyl methylsulfone tosyl salt	
24	Diethyl chloro(phenylsulfone)methylphosphonate	
25	Asp(β-tert-butyl)achlorovinyl phenylsulfone	
26	Asp(β-tert-butyl)achlorovinyl phenylsulfone tosyl salt	
27	Boc-Aspartimol(β-Methyl)	
28	Boc-Asp(β-Methyl)-H	
29	Boc-Asp(β-Methyl)methyl vinyl sulfone	
30	Boc-Asp(β-Methyl)methyl vinyl sulfone tosyl salt	
31	Diethyl (methylsulfone)methylphosphonate	
32	Boc-Asp(β-tert-butyl)methyl vinyl sulfone	
33		

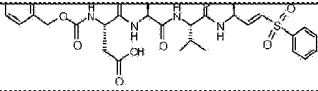
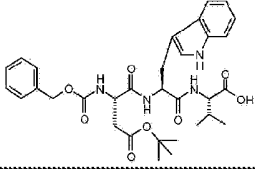
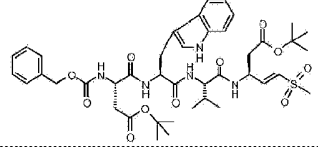
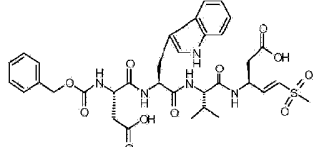
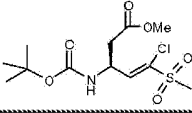
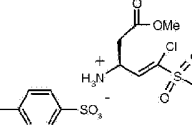
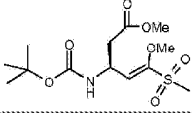
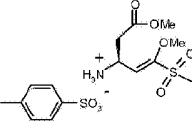
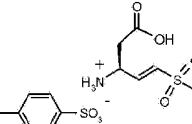
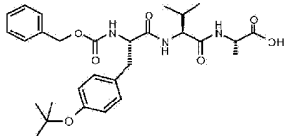
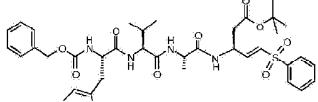
CPD N°	CPD NAME	STRUCTURE
	Asp(β -tert-butyl)methyl vinyl sulfone tosyl salt	
34	Diethyl phenylsulfonylmethylphosphonate	
35	Boc-Asp(β -tert-Butyl)-H	
36	Boc-Asp-vinyl phenyl sulfone	
37	AspVinyl phenyl sulfone tosyl salt	
38	Diethyl (phenoxysulfone)methylphosphonate	
39	Asp(β -tert-butyl) phenoxy vinyl sulfone	
40	Asp(β -tert-butyl) phenoxy vinyl sulfone tosyl salt	
41	Diethyl (isopropylsulfone)methylphosphonate	
42	BocAsp(β -tert-butyl) isopropyl vinyl sulfone	
43	Asp(β -tert-butyl)isopropyl vinyl sulfone tosyl salt	
44	Diethyl (morpholinesulfone)methylphosphonate	
45		

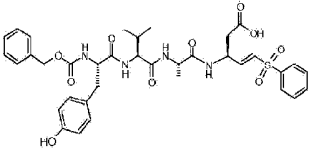
CPD N°	CPD NAME	STRUCTURE
	Boc-Asp(β -tert-butyl)morpholine vinyl sulfone	
46	Asp(β -tert-butyl)morpholine vinyl sulfone tosyl salt	
47	Z-Asp(β -tert-Butyl)-Ala(2'-quinolyl)-Val-Asp(β -tert-Butyl) α chlorovinyl methylsulfone	
48	Z-Asp-Ala(2'-quinolyl)-Val-Asp- α chlorovinyl methylsulfone	
49	Ts-Ala(2'quinolyl)-Val-Asp(β -tert-Butyl)- α chlorovinyl methylsulfone	
50	Ts-Ala(2'quinolyl)-Val-Asp- α chlorovinyl methylsulfone	
51	Z-Asp(β -methyl)-Indanylglycine-Val-Asp(β -methyl)methyl vinyl sulfone	
52	Z-Asp(β -tert-Butyl)-Phg-Val-Asp(β -tert-Butyl)methyl vinyl sulfone	
53	Z-Asp-Phg-Val- Aspmethyl vinyl sulfone	
54	Z-Asp(β -tert-Butyl)-Al(2'-quinolyl)-Val-Asp(β -tert-Butyl)methyl vinyl sulfone	
55		

CPD N°	CPD NAME	STRUCTURE
	Z-Asp-Ala(2'-quinolyl)-Val-Aspmethyl vinyl sulfone	
56	Z-Asp(β-tert-Butyl)-Indanylglycine-Val-Asp(β-tert-Butyl)methyl vinyl sulfone	
57	Z-Asp-Indanylglycine-Val-Aspmethyl vinyl sulfone	
58	Z-Asp(β-tert-Butyl)-Glu(β-tert-Butyl)-Val-Asp(β-tert-Butyl)methyl vinyl sulfone	
59	Z-Asp-Glu-Val-Aspmethyl vinyl sulfone	
60	Z-Val-Asp(β-tert-Butyl)methyl vinyl sulfone	
61	Z-Val-Aspmethyl vinyl sulfone	
62	Z-Asp(β-tert-Butyl)-Ala(2'-quinolyl)-Val-Asp(β-tert-Butyl)phenyl vinyl sulfone	
63	Z-Asp-Ala(2'-quinolyl)-Val-Aspphenyl vinyl sulfone	
64	Z-Asp(β-tert-Butyl)-Ala(2'-quinolyl)-Val-Asp(β-tert-Butyl)phenoxy vinyl sulfone	
65	Z-Asp-Ala(2'-quinolyl)-Val-Aspphenoxy vinyl sulfone	

CPD N°	CPD NAME	STRUCTURE
66	Z-Asp(β-tert-Butyl)-Ala(2'-quinolyl)-Val-Asp(β-tert-Butyl)morpholine vinyl sulfone	
67	Z-Asp-Ala(2'-quinolyl)-Val-Aspmorpholine vinyl sulfone	
68	Z-Asp-Indanylglycine-Val-Aspisopropyl vinyl sulfone	
69	Z-Asp-Phg-Val-Asp-phenyl vinylsulfone	
70	Z-Asp-(D,L Ala(2'-quinolyl))-Val-Aspphenyl vinylsulfone	
71	Z-Asp-(D,L Ala(2'-quinolyl))-Val-Aspmethyl vinylsulfone	
72	Fmoc-Tyr(O-tBu)-Val-Oallyl	
73	Tyr(O-tBu)-Val-Oallyl	
74	Z-Asp(O-tBu)-Tyr(O-tBu)-Val-Oallyl	
75	Z-Asp-Tyr(O-tBu)-Val-OH	

CPD N°	CPD NAME	STRUCTURE
76	Z-Asp-Tyr-Val-Aspmethyl vinyl sulfone	
77	Fmoc-Glu(O-tBu)-Val-Oallyl	
78	Glu(O-tBu)-Val-Oallyl	
79	Z-Tyr(O-tBu)-Glu(O-tBu)-Val-Oallyl	
80	Z-Tyr(O-tBu)-Glu(O-tBu)-Val-OH	
81	Z-Tyr(O-tBu)-Glu(O-tBu)-Val-Asp(OtBu)methyl vinyl sulfone	
82	Z-Tyr-Glu-Val-Aspmethyl vinyl sulfone	
83	Z-Asp(O-tBu)-Ala-(2'pyridine)-Val-OH	
84	Z-Asp(O-tBu)-Ala-(2'pyridine)-Val-Asp(O-tBu)phenyl vinyl sulfone	
85		

CPD N°	CPD NAME	STRUCTURE
	Z-Asp-Ala(2'-pyridyl)-Val-Aspphenyl vinylsulfone	
86	Z-Asp(O-tBu)-Trp-Val-OH	
87	Z-Asp(O-tBu)-Trp-Val-Asp(OtBu)methyl vinyl sulfone	
88	Z-Asp-Trp-Val-Aspmethyl vinyl sulfone	
89	Boc-Asp(β-methyl)αchlorovinyl methylsulfone	
90	Asp(β-methyl)αchlorovinyl methylsulfone tosyl salt	
91	Boc-Asp(β-methyl)αmethoxyvinyl methylsulfone	
92	Asp(β-methyl)αmethoxyvinyl methylsulfone tosyl salt	
93	Aspmethyl vinyl sulfone tosyl salt	
94	Z-Tyr(OtBu)-Val-Ala-OH	
95		

CPD N°	CPD NAME	STRUCTURE
	Z-Tyr(OtBu)-Val-Ala-Asp(OtBu) phenyl vinyl sulfone	
96	Z-Tyr-Val-Ala-Asp phenyl vinyl sulfone	

Definitions

[0046] Unless otherwise specified, the following definitions apply:

The singular forms "a", "an" and "the" include corresponding plural references unless the context clearly dictates otherwise.

[0047] As used herein, the term "comprising" is intended to mean that the list of elements following the word "comprising" are required or mandatory but that other elements are optional and may or may not be present.

[0048] As used herein, the term "consisting of" is intended to mean including and limited to whatever follows the phrase "consisting of". Thus the phrase "consisting of" indicates that the listed elements are required or mandatory and that no other elements may be present.

[0049] As used herein, the term "alkyl" is intended to include both branched and straight chain saturated aliphatic hydrocarbon groups having the specified number of carbon atoms, for example, C₁-C₆ as in C₁-C₆ - alkyl is defined as including groups having 1,2,3,4,5 or 6 carbons in a linear or branched arrangement. Examples of C₁-C₆-alkyl and C₁-C₄ alkyl as defined above include methyl, ethyl, n-propyl, *i*-propyl, *n*-butyl, *t*-butyl, *i*-butyl, pentyl and hexyl.

[0050] As used herein, the term, "alkenyl" is intended to mean unsaturated straight or branched chain hydrocarbon groups having the specified number of carbon atoms therein, and in which at least two of the carbon atoms are bonded to each other by a double bond, and having either E or Z regiochemistry and combinations thereof. For example, C₂-C₆ as in C₂-C₆ alkenyl is defined as including groups having 1, 2, 3, 4, 5, or 6 carbons in a linear or branched arrangement, at least two of the carbon atoms being bonded together by a double bond. Examples of C₂-C₆ alkenyl include ethenyl (vinyl), 1-propenyl, 2-propenyl, 1- butenyl.

[0051] As used herein, the term "cycloalkyl" is intended to mean a monocyclic saturated aliphatic hydrocarbon group having the specified number of carbon atoms therein, for example, C₃-C₇ as in C₃-C₇ cycloalkyl is defined as including groups having 3,4,5,6, or 7 carbons in a monocyclic arrangement. Examples of C₃-C₇ cycloalkyl as defined above include cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl and cycloheptyl.

[0052] As used herein, the term "halo" or "halogen" is intended to mean fluorine, chlorine, bromine and iodine.

[0053] As used herein, the term "haloalkyl" is intended to mean an alkyl as defined above, in which each hydrogen atom may be successively replaced by a halogen atom. Examples of haloalkyls include CH₂F, CHF₂ and CF₃.

[0054] As used herein, the term "aryl", either alone or in combination with another radical, means a carbocyclic aromatic monocyclic group containing 6 carbon atoms which may be further fused to a second 5- or 6-membered carbocyclic group which may be aromatic, saturated or unsaturated. Aryl includes phenyl, indanyl, 1-naphthyl, 2-naphthyl and tetrahydronaphthyl. The aryls may be connected to another group either at a suitable position on the cycloalkyl ring or the aromatic ring.

[0055] As used herein, the term "heteroaryl" is intended to mean a monocyclic or bicyclic ring system of up to ten atoms, wherein at least one ring is aromatic, and contains from 1 to 4 hetero atoms selected from the group consisting of O, N, and S. The heteroaryl substituent may be attached either via a ring carbon atom or one of the heteroatoms. Examples of heteroaryl groups include thienyl, benzimidazolyl, benzo[b]thienyl, furyl, benzofuranyl, pyranlyl, isobenzofuranyl, chromenyl, xanthenyl, 2H-pyrrolyl, pyrrolyl, imidazolyl, pyrazolyl, pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, indolizynyl, isoindolyl, 3H-indolyl, indolyl, indazolyl, purinyl, 4H-quinolizynyl, isoquinolyl, quinolyl, phthalazinyl, naphthyridinyl, quinoxalinyl, quinazolinyl, cinnolynyl, pteridinyl, isothiazolyl, isochromanyl, chromanyl, isoxazolyl, furazanyl, indolynyl, isoindolynyl, thiazolo[4,5-b]-pyridine, and fluoroscein derivatives.

[0056] As used herein, the term "heterocycle", "heterocyclic" or "heterocyclyl" is intended to mean a 5, 6, or 7 membered non-aromatic ring system containing from 1 to 4 heteroatoms selected from the group consisting of O, N and S. Examples of heterocycles include pyrrolidinyl, tetrahydrofuranyl, piperidyl, pyrrolinyl, piperazinyl, imidazolidinyl, morpholinyl, imidazolynyl, pyrazolidinyl, and pyrazolinyl.

[0057] As used herein, the term "electron withdrawing group (EWG)" is intended to mean a functional group that allows nucleophilic attack by the thiol-group of a caspase at the alkene bond of the inhibitor as a result of the electron withdrawing properties of the EWG. The EWG is conjugated with the alkene bond, such that the electron withdrawing properties of the EWG allow nucleophilic attack by a caspase at the alkene bond, i.e. the alkene bond and the EWG are electronically conjugated. Thus, the covalent bond between the alkene bond and the EWG is a direct one, without intervening moieties that would prevent the electron withdrawing properties of the EWG from being exerted on the alkene bond.

[0058] As used herein, the term "detectable label" is intended to mean a group that may be linked to a compound of the present invention to produce a probe or to a caspase, such that when the probe is associated with the caspase, the label allows either direct or indirect recognition of the probe so that it may be detected, measured and quantified.

[0059] As used herein, the term "affinity tag" is intended to mean a ligand or group, which is linked to either a compound of the present invention or to a caspase to allow another compound to be extracted from a solution to which the ligand or group is attached.

[0060] As used herein, the term "probe" is intended to mean a compound of Formula I, IA, II, IIA, III, or IIIA, which is labeled with either a detectable label or an affinity tag, and which is capable of binding, either covalently or non-covalently, to a caspase. When, for example, the probe is non-covalently bound, it may be displaced by a test compound. When, for example, the probe is bound covalently, it may be used to form cross-linked adducts, which may be quantified and inhibited by a test compound.

[0061] As used herein, the term "optionally substituted with one or more substituents" or its equivalent term "optionally substituted with at least one substituent" is intended to mean that the subsequently described event of circumstances may or may not occur, and that the description includes instances where the event or circumstance occurs and instances in which it does not. The definition is intended to mean from zero to five substituents.

[0062] If the substituents themselves are incompatible with the synthetic methods of the present invention, the substituent may be protected with a suitable protecting group (PG) that is stable to the reaction conditions used in these methods. The protecting group may be removed at a suitable point in the reaction sequence of the method to provide a desired intermediate or target compound. Suitable protecting groups and the methods for protecting and de-protecting different substituents using such suitable protecting groups are well known to those skilled in the art; examples of which may be found in T. Greene and P. Wuts, *Protecting Groups in Chemical Synthesis* (3rd ed.), John Wiley & Sons, NY (1999). Examples of protecting groups used throughout include Fmoc, Bn, Boc, CBz and COC₃. In some instances, a substituent may be specifically selected to be reactive under the reaction conditions used in the methods of this invention. Under these circumstances, the reaction conditions convert the selected substituent into another substituent that is either useful in an intermediate compound in the methods of this invention or is a desired substituent in a target compound.

[0063] Three and single letter abbreviations for α -amino acids used throughout are as follows:

Amino acid	Abbreviation	Abbreviation
α -Amino butyric acid	Abu	-
Alanine	Ala	A
Arginine	Arg	R
Aspartic acid	Asp	D
Asparagine	Asn	N
Cysteine	Cys	C
Glutamic acid	Glu	E
Glutamine	Gln	Q
Glycine	Gly	G
Isoleucine	Ile	I
Histidine	His	H
Leucine	Leu	L
Lysine	Lys	K
Methionine	Met	M
Phenylalanine	Phe	F
Proline	Pro	P
Serine	Ser	S

Amino acid	Abbreviation	Abbreviation
Threonine	Thr	T
Tryptophan	Trp	W
Tyrosine	Tyr	Y
Valine	Val	V

[0064] A series of non natural amino acids which may be used in place of natural amino acids is 3-amino 2-hydroxy pyridine; (2furyl)alanine; 1amino 1cyclohexane carboxylic acid; (2thienyl)alanine; 2-Aminobenzoic acid (2-Abz); 2PyridylAlanine; 1Amino 1Cyclopentanecarboxylic acid; 2-Aminobutyric acid (2Abu); 3Amino3phenylpropionic acid; Aminocyclopentane carboxylic acid (ACPC); 4-Aminomethylbenzoic acid (Amb); Aminoisobutiric acid (Aib); p-Benzoyl-1-phenylalanine (Bpa); AllylGlycine; 4-Aminomethyl cyclohexane carboxylic acid (Amc); Cyclohexyl-alanine (Cha); deltaValine; deltaLeucine; Cyanobbutylalanine (Cba); Indanylglycine (Igl); 3-(2-naphthyl)alanine (1-Nal); Biphenylalanine (Bip); Hydroxyproline (Hyp); Isonipecotic acid (Inp); Norvaline (Nva); 4-Iodophenylalanine (Phe(PI)); 4-nitroPhenylalanine; 4MethylPhenylalanine; 4MethylPhealanine; Homophenylalanine (hPhe); 4-aminophenylalanine (Phe4NH(Boc)); phenyl glycine; Pipecolic acid (Pip); propargylglycine; Thioproline (Thz); ButylGlycine (Tle); 3-NitroTyrosine.

[0065] As used herein, the term "residue" when referring to α -amino acids is intended to mean a radical derived from the corresponding α -amino acid by eliminating the hydroxyl of the carboxy group and one hydrogen of the α -amino group. For example, the terms Gln, Ala, Gly, Ile, Arg, Asp, Phe, Ser, Leu, Cys, Asn, and Tyr represent the residues of L-glutamine, L-alanine, glycine, L-isoleucine, L-arginine, L-aspartic acid, L-phenylalanine, L-serine, L-leucine, L-cysteine, L-asparagine, and L-tyrosine, respectively.

[0066] As used herein the term "amino acid side chain" is intended to mean the part of an amino acid's chemistry that differentiates it from other amino acids. Amino acid structure includes a carboxyl group, an amine group plus the individual side chain. Each amino acid has a unique side chain. This is applied to unnatural amino acids as well. This side chain may exist in protected form or not.

[0067] As used herein, the term "prodrug" is intended to mean a compound that may be converted under physiological conditions or by solvolysis to a biologically active compound of the present invention. Thus, the term "prodrug" refers to a precursor of a compound of the invention that is pharmaceutically acceptable. A prodrug may be inactive or display limited activity when administered to a subject in need thereof, but is converted in vivo to an active compound of the present invention. Typically, prodrugs are transformed in vivo to yield the compound of the invention, for example, by hydrolysis in blood or other organs by enzymatic processing. The prodrug compound often offers advantages of solubility, tissue compatibility or delayed release in the subject (see, Bundgard, H., Design of Prodrugs (1985), pp. 7-9, 21-24 (Elsevier, Amsterdam). The definition of prodrug includes any covalently bonded carriers which release the active compound of the invention in vivo when such prodrug is administered to a subject. Prodrugs of a compound of the present invention may be prepared by modifying functional groups present in the compound of the invention in such a way that the modifications are cleaved, either in routine manipulation or in vivo, to a parent compound of the invention.

[0068] As used herein, the term "pharmaceutically acceptable salt" is intended to mean both acid and base addition salts.

[0069] As used herein, the term "pharmaceutically acceptable acid addition salt" is intended to mean those salts which retain the biological effectiveness and properties of the free bases, which are not biologically or otherwise undesirable, and which are formed with inorganic acids such as hydrochloric acid, hydrobromic acid, sulfuric acid, nitric acid, phosphoric acid, and organic acids such as acetic acid, trifluoroacetic acid, propionic acid, glycolic acid, pyruvic acid, oxalic acid, maleic acid, malonic acid, succinic acid, fumaric acid, tartaric acid, citric acid, benzoic acid, cinnamic acid, mandelic acid, methanesulfonic acid, ethanesulfonic acid, p-toluenesulfonic acid, salicylic acid.

[0070] As used herein, the term "pharmaceutically acceptable base addition salt" is intended to mean those salts which retain the biological effectiveness and properties of the free acids, which are not biologically or otherwise undesirable. These salts are prepared from addition of an inorganic base or an organic base to the free acid. Salts derived from inorganic bases include the sodium, potassium, lithium, ammonium, calcium, magnesium, iron, zinc, copper, manganese, aluminum salts. Salts derived from organic bases include salts of primary, secondary, and tertiary amines, substituted amines including naturally occurring substituted amines, cyclic amines and basic ion exchange resins, such as isopropylamine, trimethylamine, diethylamine, triethylamine, tripropylamine, ethanolamine, 2-dimethylaminoethanol, 2-diethylaminoethanol, dicyclohexylamine, lysine, arginine, histidine, caffeine, procaine, hydrabamine, choline, betaine, ethylenediamine, glucosamine, methylglucamine, theobromine, purines, piperazine, piperidine, N-ethylpiperidine, polyamine resins.

[0071] The compounds of the present invention, or their pharmaceutically acceptable salts may contain one or more asymmetric centers, chiral axes and chiral planes and may thus give rise to enantiomers, diastereomers, and other stereoisomeric forms and may be defined in terms of absolute stereochemistry, such as (R)- or (S)- or, as (D)- or (L)- for amino acids. The present invention is intended to include all such possible isomers, as well as, their racemic and optically pure forms. Optically active (+) and (-), (R)- and (S)-, or (D)- and (L)-isomers may be prepared using chiral synthons or chiral reagents, or resolved using conventional techniques,

such as reverse phase HPLC. The racemic mixtures may be prepared and thereafter separated into individual optical isomers or these optical isomers may be prepared by chiral synthesis. The enantiomers may be resolved by methods known to those skilled in the art, for example by formation of diastereoisomeric salts which may then be separated by crystallization, gas-liquid or liquid chromatography, selective reaction of one enantiomer with an enantiomer specific reagent. It will also be appreciated by those skilled in the art that where the desired enantiomer is converted into another chemical entity by a separation technique, an additional step is then required to form the desired enantiomeric form. Alternatively specific enantiomers may be synthesized by asymmetric synthesis using optically active reagents, substrates, catalysts, or solvents or by converting one enantiomer to another by asymmetric transformation.

[0072] Certain compounds of the present invention may exist as a mix of epimers. Epimers means diastereoisomers that have the opposite configuration at only one of two or more stereogenic centres present in the respective compound.

[0073] Certain compounds of the present invention may exist in Zwitterionic form and the present invention includes Zwitterionic forms of these compounds and mixtures thereof.

[0074] In addition, the compounds of the invention also may exist in hydrated and anhydrous forms. Hydrates of the compound of any of the formulas described herein are included as compounds of the invention. In a further embodiment, the compound according to any of the formulas described herein is a monohydrate. In one embodiment, the compound of the invention comprises about 10% or less, about 9 % or less, about 8% or less, about 7% or less, about 6% or less, about 5% or less, about 4% or less, about 3% or less, about 2% or less, about 1% or less, about 0.5% or less, about 0.1% or less by weight of water. In another embodiment, the compounds of the invention comprise, about 0.1% or more, about 0.5% or more, about 1% or more, about 2% or more, about 3% or more, about 4% or more, about 5% or more, or about 6% or more by weight of water.

C) Methods of preparation

[0075] General methods for the synthesis of the compounds of the present invention are shown below.

[0076] Those skilled in the art will readily appreciate that a number of methods are available for the preparation of the compounds of the present invention.

[0077] The starting materials and reagents used in preparing these compounds are either available from commercial suppliers such as Sigma-Aldrich Chemicals, Anaspec or chemipex.

[0078] The starting materials and the intermediates of the reaction may be isolated and purified if desired using conventional techniques such chromatography (Biotage flash chromatography), filtration, distillation, etc. Such materials can be characterized using conventional analytical methods such as NMR and LCMS.

[0079] The coupling step is carried out in the presence of a suitable coupling agent such as, , diisopropyl carbodiimide (DPC), anhydride mixte (isobutylchloroformate), O-benzotriazole-1-yloxytrispyrrolidinophosphonium hexafluorophosphate (PyBOP), O-benzotriazol-1-yl-N,N,N,N tetramethyluroniumhexafluoro-phosphate (HBTU), O-(7-azabenzotriazol-1-yl)-1,1,3,3-tetramethyluronium hexafluoro-phosphate (HATU), 1-hydroxybenzotriazole (HOBT) in the presence of 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (EDC). A base such as N,N-diisopropylethylamine, triethyl amine, or N-methylmorpholine. The reaction is carried out at 20 °C except for anhydride mix formation (isobutyl chloroformate, -5/-13 °C) to avoid any racemisation of amino acids.

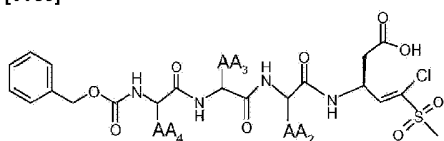
[0080] Removal of the amino protecting group is carried out but by piperidine in dichloromethane (Fmoc protecting group). The removal of β -tert-butyl carboxylic acid is carried out by TFA. Selective removal of N-Boc from B-tert butyl aspartic acid is carried out by PTSA or TFA at 0°C.

[0081] Vinyl sulfone function is elucidated as an example only in the following examples. It will be recognized by a skilled person in the art, that other vinyl electron-withdrawing group (EWG) may be used in this invention. Benzyloxycarbonyl (Z) is elucidated as an example of linkers that may facilitate the penetration of the drug into cells. It is recognized that other X-R3 may be used as well. The allyl group was introduced directly by the use of commercially available AA₂-OAllyl protected form or synthesized from the the corresponding amino acid AA₂-OH.

[0082] This is a convergent synthesis, which consists of synthesizing two different fragments (suicide inhibitor linker and the peptide or peptidomimetic fragment) prior to the 'coupling step'.

A compound of Formula 1a can be prepared by the procedure described bellow

[0083]

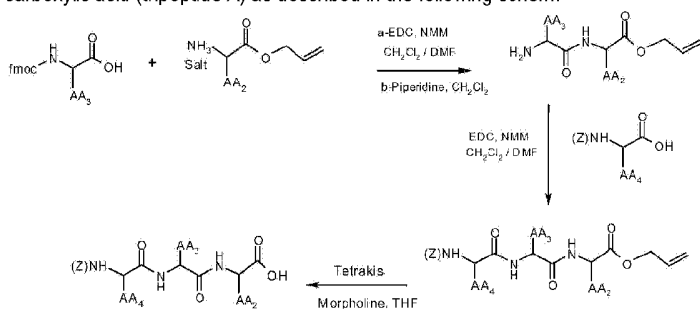


Formula 1a

Left arm synthesis:

[0084] The allyl group was introduced directly by the use of commercially available AA₂-Oallyl protected form or synthesized from the protected amino acid fmoc-AA₂-OH and allyl alcohol (EDC, DMAPcat, NMM, CH₂Cl₂/DMF (5/1)) followed with fmoc deprotection (piperidine, CH₂Cl₂).

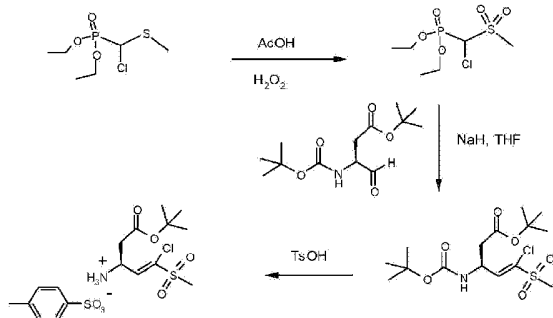
[0085] The AA₂-Oallyl is coupled to Fmoc-AA₃-OH with the coupling reagent 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (EDC) and subsequent fmoc deprotection with piperidine gives AA₃-AA₂-Oallyl (intermediate A) ready for coupling with (Z) AA₄-OH gave (Z) AA₄-AA₃-AA₂-Oallyl (intermediate B) which upon removal of allyl group (tetrakis) liberate the C-terminal carboxylic acid (tripeptide A) as described in the following scheme.

**Right arm synthesis**

[0086] The suicide substrate proposed in the following scheme is Asp α-chlorovinyl methylsulfone.

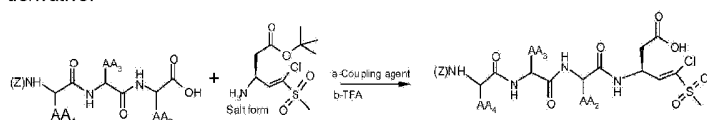
[0087] The common intermediate Boc-Asp(B-tert-butyl)-H is synthesized from Boc-Asp(B-tert-butyl)-N-hydroxysuccinimide ester as reported by (Mancuso A et al., 1981, William R. Ewing et al., 1999 and Won Bum Jang. 2004).

[0088] Treatment of the aldehyde with sodium anion of Diethyl chloro(methylsulfone)methylphosphonate results in the corresponding Boc-Asp (β-tert-butyl) α-chlorovinyl-methylsulfone in the manner of Wadsworth and Emmons.

**Coupling step synthesis**

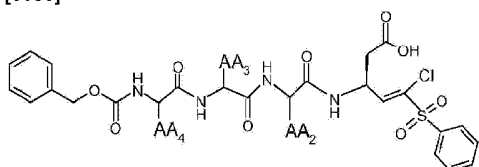
[0089] The coupling step between Asp α chlorovinyl methylsulfone salt and the peptide in the presence of coupling reagents such (isobutyl chloroformate, NMM) or (HOBT, NMM, EDC; Dragovich P., S., 1999) results in Asp α-chlorovinyl methylsulfone peptide

derivative.



A compound of Formula 1b can be prepared by the procedure described below

[0090]

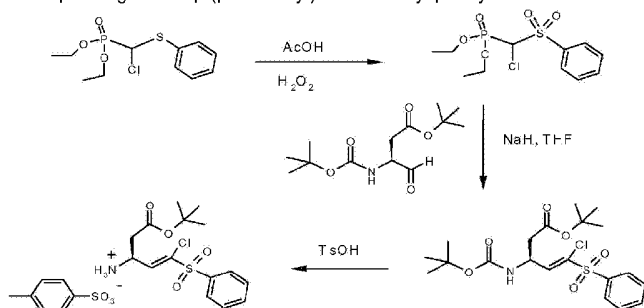


Formula 1b

Right arm synthesis

[0091] The suicide substrate proposed in the following scheme is Asp α -chlorovinyl phenylsulfone.

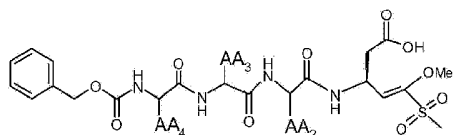
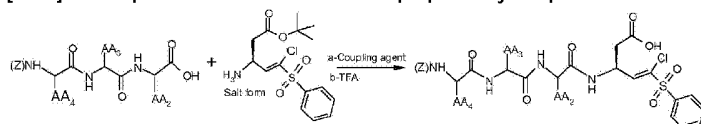
[0092] Treatment of the aldehyde with sodium anion of Diethyl chloro(methylsulfone)phenyl phosphonate results in the corresponding Boc-Asp (β -tert-butyl) α -chlorovinyl phenylsulfone in the manner of Wadsworth and Emmons.



Coupling step synthesis

[0093] The coupling step between Asp α -chlorovinyl phenylsulfone salt and the peptide in the presence of coupling reagents such (isobutyl chloroformate, NMM) or (HOBT, NMM, EDC) results in Asp α -chlorovinyl phenylsulfone peptide derivative.

[0094] A compound of Formula 1c can be prepared by the procedure described below



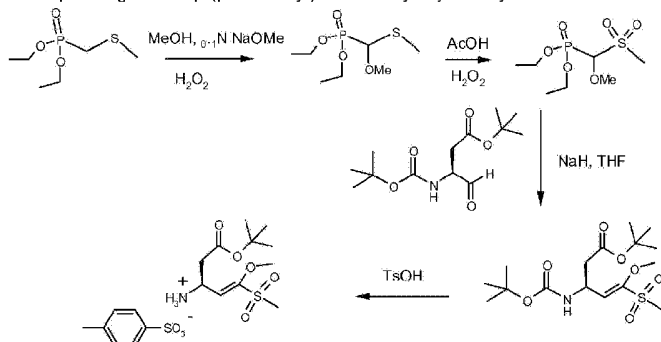
Formula 1c

Right arm synthesis

[0095] The suicide substrate proposed in the following schem is Asp α -methoxyvinyl methylsulfone.

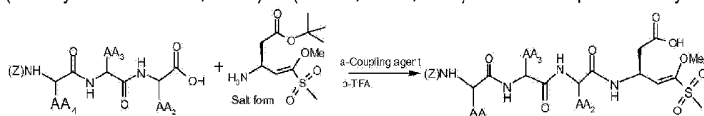
[0096] Oxidation of diethyl methylthiomethylphosphonate include anodic oxidation in NaOMe/MeOH (saturated with CO₂) to give the O, S-acetal of diethoxyphosphinylformaldehyde, which upon oxidation may give the corresponding sulfone as Diethyl methoxy(methylsulfone)phenyl phosphonate (Shankaran, K et al. 2001).

[0097] Treatment of the aldehyde with sodium anion of Diethyl methoxy (methylsulfone)phenyl phosphonate should result in the corresponding Boc-Asp (β -tert-butyl) α -methoxyvinyl methylsulfone in the manner of Wadsworth and Emmons.



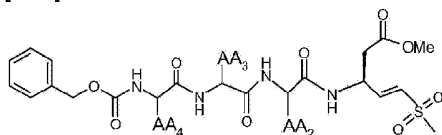
Coupling step synthesis

[0098] The coupling step between Asp α -methoxyvinyl methylsulfone salt and the peptide in the presence of coupling reagents such (isobutyl chloroformate, NMM) or (HOBT, NMM, EDC) results in Asp α -methoxyvinyl methylsulfone peptide derivative.



A compound of Formula 1d can be prepared by the procedure described below

[0099]



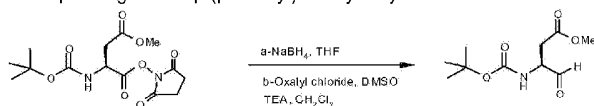
Formula 1d

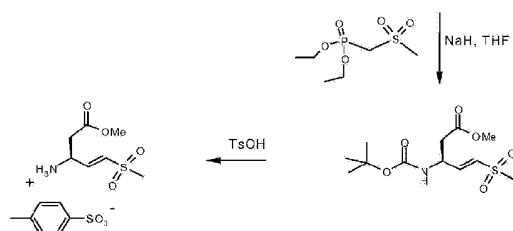
Right arm synthesis

[0100] The suicide substrate proposed in the following schem is Asp(B-Methyl) methyl vinylsulfone.

[0101] Boc-Asp(B-Methyl)-N-hydroxysuccinimide ester was reduced to the corresponding alcohol (NaBH₄, THF), a subsequent oxidation (oxalyl chloride, DMSO) gave the aldehyde: Boc-Asp(B-Methyl)-H.

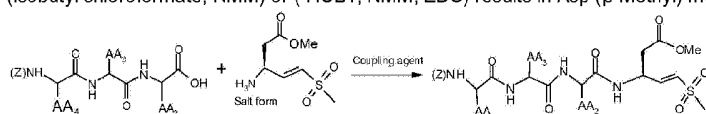
[0102] Treatment of Boc-Asp(B-Methyl)-H with sodium anion of Diethyl (methylsulfone) methyl phosphonate results in the corresponding Boc-Asp (β -Methyl) methyl vinylsulfone in the manner of Wadsworth and Emmons.



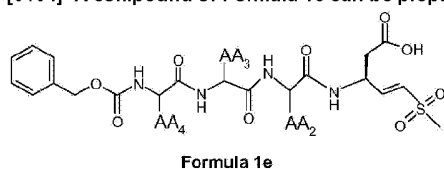


Coupling step synthesis

[0103] The coupling step between Asp (β-Methyl) methyl vinylsulfone salt and the peptide in the presence of coupling reagents such (isobutyl chloroformate, NMM) or (HOBT, NMM, EDC) results in Asp (β-Methyl) methyl vinylsulfone peptide derivative.



[0104] A compound of Formula 1e can be prepared by the procedure described below

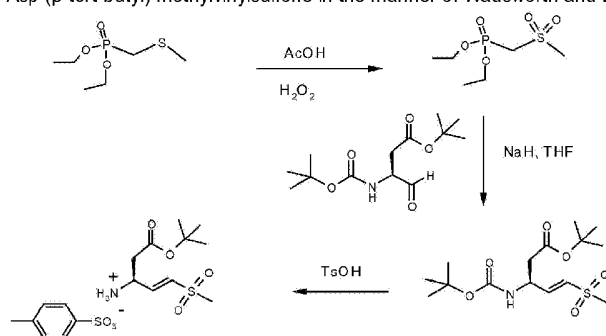


Right arm synthesis

[0105] The suicide substrate proposed in the following schem is Asp methyl vinylsulfone.

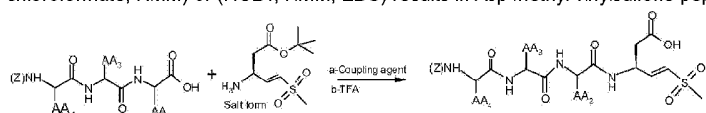
[0106] The common intermediate Boc-Asp(B-tert-butyl)-H is synthesized from Boc-Asp(B-tert-butyl)-N-hydroxysuccinimide ester as reported by (Mancuso A et al., 1981, William R. Ewing et al., 1999 and Won Bum Jang. 2004).

[0107] Treatment of the aldehyde with sodium anion of Diethyl (methylsulfonyl) methyl phosphonate results in the corresponding Boc-Asp (B-tert-butyl) methylvinylsulfone in the manner of Wadsworth and Emmons, 1961 and Palmer et al. 1995.

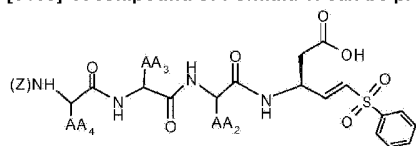


Coupling step synthesis

[0108] The coupling step between Asp methyl vinylsulfone salt and the peptide in the presence of coupling reagents such (isobutyl chloroformate, NMM) or (HOBT, NMM, EDC) results in Asp methyl vinylsulfone peptide derivative.



[0109] A compound of Formula 1f can be prepared by the procedure described below



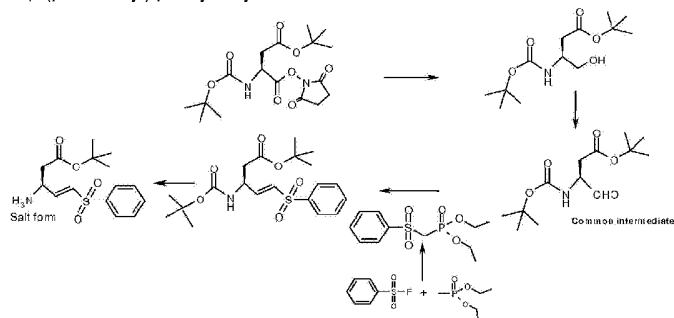
Formula 1f

Right arm synthesis

[0110] The suicide substrate proposed in the following schem is Asp phenyl vinylsulfone.

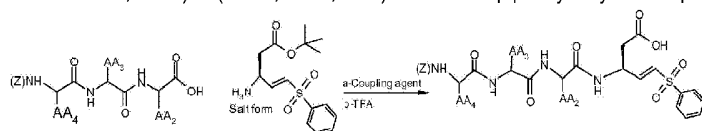
[0111] Diethyl phenylsulfonylmethylphosphonate was obtained in one step from benzenesulfonyl fluoride and triethyl phosphorane in the presence of lithium hexamethyldisilazide (Won Bum Jang et al., 1998).

[0112] Treatment of the aldehyde with sodium anion of Diethyl (phenyl sulfone) methylphosphonate results in the corresponding Boc-Asp (β-tert-butyl) phenyl vinylsulfone in the manner of Wadsworth and Emmons, 1961 and Palmer et al. 1995.

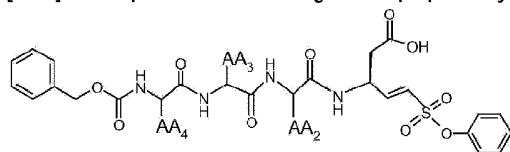


Coupling step synthesis

[0113] The coupling step between Asp phenyl vinylsulfone salt and the peptide in the presence of coupling reagents such (isobutyl chloroformate, NMM) or (HOBT, NMM, EDC) results in Asp phenyl vinylsulfone peptide derivative.



[0114] A compound of Formula 1g can be prepared by the procedure described below



Formula 1g

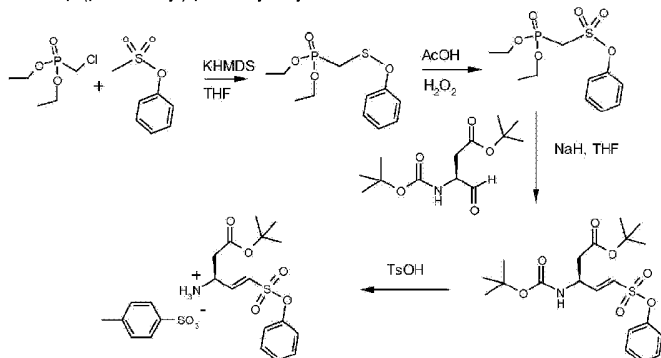
Right arm synthesis

[0115] The suicide substrate proposed in the following schem is Asp phenoxy vinylsulfone.

[0116] Diethyl (phenoxysulfone) methylphosphonate was obtained from methanesulfonyl phenoxy and diethyl chlorophosphonate in

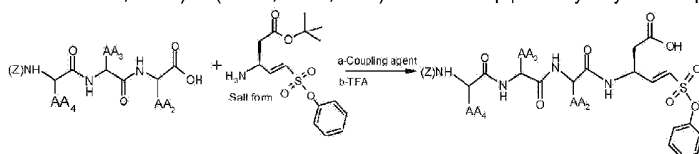
the presence of potassium bis(trimethylsilyl)amide. A subsequent oxidation (AcOH, H₂O₂) gave the corresponding sulfone.

[0117] Treatment of the aldehyde with sodium anion of Diethyl (phenoxy sulfone) methylphosphonate results in the corresponding Boc-Asp (β-tert-butyl) phenoxy vinylsulfone in the manner of Wadsworth and Emmons.

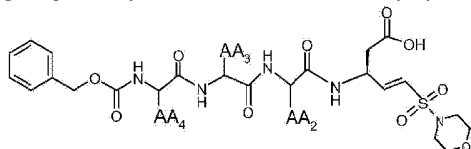


Coupling step synthesis

[0118] The coupling step between Asp phenoxy vinylsulfone salt and the peptide in the presence of coupling reagents such (isobutyl chloroformate, NMM) or (HOBT, NMM, EDC) results in Asp phenoxy vinylsulfone peptide derivative.



[0119] A compound of Formula 1h can be prepared by the procedure described bellow



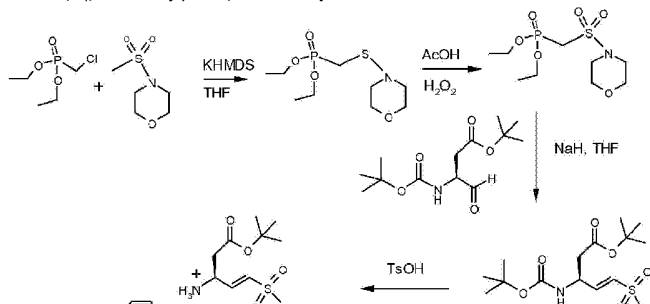
Formula 1h

Right arm synthesis

[0120] The suicide substrate proposed in the following schem is Asp morpholine vinylsulfone.

[0121] Diethyl (morpholinesulfone) methylphosphonate was prepared from methane sulfonyl morpholine and chloromethylphosphonate in the presence of potassium bis(trimethylsilyl) amide.

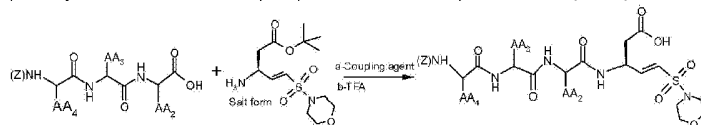
[0122] Treatment of the aldehyde with sodium anion of Diethyl (morpholinesulfone) methylphosphonate results in the corresponding Boc-Asp (β-tert-butyl) morpholino vinylsulfone in the manner of Wadsworth and Emmons.



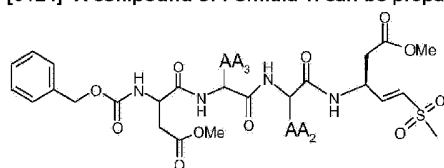


Coupling step synthesis

[0123] The coupling step between Asp morpholine vinylsulfone salt and the peptide in the presence of coupling reagents such (isobutyl chloroformate, NMM) or (HOBT, NMM, EDC) results in Asp morpholine vinylsulfone peptide derivative



[0124] A compound of Formula 1i can be prepared by the procedure bellow



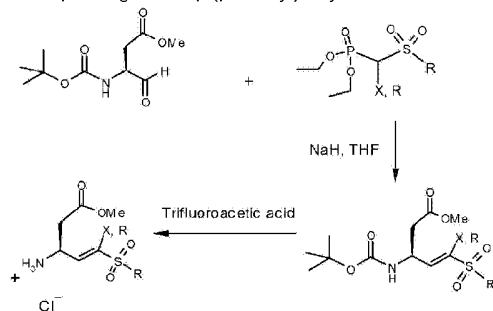
Formula 1i

Right arm synthesis

[0125] The suicide substrate proposed in the following schem is Asp(B-Methyl)- vinylsulfone derivatives.

[0126] Boc-Asp(B-Methyl)-N-hydroxysuccinimide ester was reduced to the corresponding alcohol (NaBH₄, THF), a subsequent oxidation (oxalyl chloride , DMSO) gave the aldehyde Boc-Asp(B-Methyl)-H.

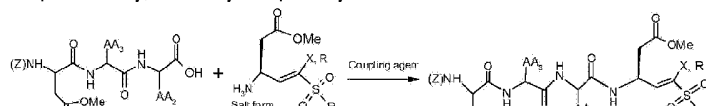
[0127] Treatment of Boc-Asp(B-Methyl)-H with sodium anion of Diethyl (methylsulfone) phosphonate derivatives results in the corresponding Boc-Asp (β-Methyl) vinylsulfone derivatives in the manner of Wadsworth and Emmons.



Coupling step synthesis

[0128] The coupling step between Asp (β-Methyl) vinylsulfone derivatives salt and the caspase-3 designed peptide in the presence of coupling reagents such (isobutyl chloroformate, NMM) or (HOBT, NMM, EDC) results in Z-Asp(β-Methyl)-AA₃-AA₂-Asp (β-Methyl) vinylsulfone peptide derivatives.

[0129] This approach will allow the synthesis of a variety of pro-drug vinylsulfone derivatives that could be easily obtained by choosing the appropriate combination (X, R) and applying the appropriate method mentioned above. Such combination may enhance cell permeability, selectivity and potency.





[0130] All acid, salt and other ionic and non-ionic forms of the compounds described are included as compounds of the invention. For example, if a compound is shown as an acid herein, the salt forms of the compound are also included. Likewise, if a compound is shown as a salt and the acid forms are also included.

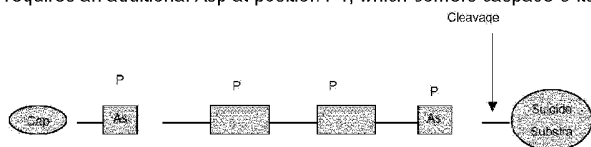
[0131] In certain embodiments, the compounds of the present invention are represented by generalized Formula IA, II, IIA, III, or IIIA, or a pharmaceutically acceptable salt thereof:

Also described herein are compounds represented by generalized Formula I, or a pharmaceutically acceptable salt and/or prodrug thereof.

D) Development of specific Caspase inhibitors

[0132] An additional aspect of the invention relates to a method for designing caspase inhibitors. Following the same approach as outlined hereinafter, those skilled in the art will appreciate that it is conceivable to further improve the potency of the compounds of Formula II, e.g. Z-Asp-Phg-Val-AspVSMethyl (53), against caspase-3 and inhibit selectively additional caspases such as caspase-2, caspase-8, caspase-9 and caspase-1.

[0133] As is known, all caspases cleave substrates to the right of the aspartic acid amino acid in position P1. However, caspase-3 requires an additional Asp at position P4, which confers caspase-3 its specificity:



[0134] As shown in **Table 2** hereinafter, suicide substrates such as AspVSMethyl (Compound 93) and Asp(OtBu)VSMethyl (Compound 33) are devoid of any activity against caspase-3. The substrate z-Asp(OtBu)-Phg-val-OH (Compound 16) is also devoid of any activity against caspase-3. However the fusion product of peptide and suicide substrate, z-Asp-Phg-Val-AspVSMethyl (Compound 53), proved to be a very potent inhibitor of caspase-3 with an IC₅₀ 30-90nM. Inhibition is selective since even though caspase-7 belongs to the same group than caspase-3, the IC₅₀ value for this caspase was about 12 fold higher. Replacement of Phg at position P3 with Ala (2'-quinolyl) enhanced the selectivity further (about 56 fold difference, see Compound 55).

[0135] As observed with z-Val-AspVSMethyl (Compound 61), the deletion of both Aspartic acid and Ala(2'-quinolyl) at P4 and P3 positions abolished completely the activity against caspase-3. The same result was observed after the deletion of Aspartic acid only in the example of z-Ala (2'-quinolyl)-Val-Asp alpha chlorovinyl methyl sulfone (Compound 50) compared with z-Asp-Ala(2'-quinolyl)-Val-Asp alpha chlorovinyl methyl sulfone (Compound 48).

Changes at position P3 and P2

[0136] The amino acids at both position P3 and P2 can serve to selectively target caspase-3 as observed but also to selectively target other caspases. The following examples highlight this possibility: (1) The replacement of Ala(2'-quinolyl) (Compound 55) with indalylglycine at P3 position lead to z-Asp-indalylglycine-Val-AspVSMethyl (Compound 57). This substitution enhanced the inhibitory effect against the group III of caspases (caspase-3 (30-90nM) and 7 (0.18-0.30 uM) and, with about 29 fold less efficiency, against group 1 (caspase-1 (0.6-1.2 uM)). (2) The presence of Trp at P3 position (z-Asp-Trp-Val-AspVSMethyl; Compound 88) retained the selectivity against caspase-3 (30-90 nM) and produced an additional activity against caspase-1 (0.6-1.2 uM). This molecule possesses therefore the capability to inhibit selectively two caspases belonging to two different groups, namely the proinflammatory and the proapoptotic group. (3) The presence of Glutamic acid at P-3 position (z-Asp-Glu-Val-AspVSMethyl; Compound 59) retained the selectivity against caspase-3 (IC₅₀ 20nM) and produced an additional activity against caspase-7 (IC₅₀ 42 nM) and caspase-9 (509 nM). This molecule therefore possesses the capability to inhibit selectively two caspases belonging to two different groups, namely initiator and executioner caspases.

Changes in position P4

[0137] Changes in position P4 can also affect the selectivity of a given inhibitor. The amino acid that has been showed to fit well into the corresponding caspase-1 pockets at position P4 is Tyrosine. Therefore, z-Tyr-Val-Ala-AspVS phenyl (Compound 96) was tested against different caspases, and it proved to be selective against caspase-1 (IC₅₀ 1.2-1.5 μM). The amino acid that has been showed to fit well into the corresponding caspase-1 pockets at position P3 is glutamic acid. Therefore, z-Tyr-Glu-Ala-AspVS methyl (Compound 82) was tested against different caspases and the inhibition of caspase-1 was enhanced to 0.5 μM.

[0138] These specific examples demonstrate that it is possible to make selective caspase-3 inhibitors based on sequences recognized by group III caspases. Following the same approach as outlined hereinafter, it is conceivable to inhibit selectively additional caspases and to further improve the potency against selected caspases.

Design for selective caspase-3 inhibitors

[0139] In one embodiment, the method comprises synthesizing compounds having the following general **Formula D1**:



wherein

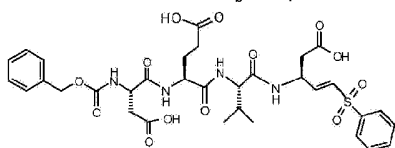
P2 is selected from the following amino acids: V, L, P, M, A, T, and H;

P3 is selected from the following amino acids: Phg, E, Indanylglycine, W, Y, A, D, Ala-(2'-quinolyl), Q, F, S, T, V, Y, G, L;

D_a is (D) or (L) aspartic acid.

D_b is the side chain of (D) or (L) aspartic acid.

[0140] The following compound (DEVD-vinyl phenyl sulfone) is an example a compounds having a sequence (i.e. **D_a** is Cbz-aspartic acid; **P3** is Glu; **P2** is Val; **D_b** is Asp; and the suicide substrate is vinyl phenyl sulfone) designed to selectively inhibit caspase-3:



Design for selective Caspase-8/Caspase-9 inhibitors

[0141] In one embodiment, the method comprises synthesizing compounds having the following general **Formula D2**:



where

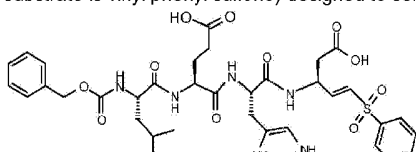
P2 is selected from the following amino acids: T, H, V, W, I, and A;

P3 is selected from the following amino acids: E, Ala(2'-quinolyl);

P4 is selected from the following amino acids: I, L, E, D, A, P, and V;

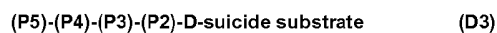
D is the side chain of (D) or (L) aspartic acid and the suicide substrate is vinyl phenyl sulfone.

[0142] The following is an example of a compound having a sequence (i.e. **P4** is Cbz-L; **P3** is Glu; **P2** is His; **D** is Asp and the suicide substrate is vinyl phenyl sulfone) designed to selectively inhibit caspase-8:



Design for selective caspase-2 inhibitors

[0143] In one embodiment, the method comprises synthesizing compounds having the following general **Formula D3**:



where

P2 is selected from the following amino acids: A, S, K and V;

P3 is selected from the following amino acids: V, E, T and Q;

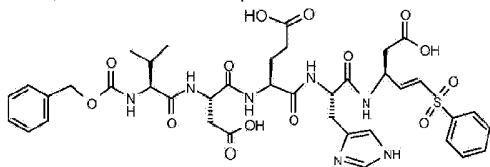
P4 is selected from the following amino acids: D and L;

P5 is selected from the following amino acids: V and L;

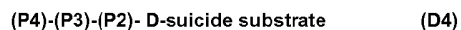
D is the side chain of (D) or (L) aspartic acid; and

the suicide substrate is selected from the group consisting of vinyl phenyl sulfone.

[0144] The following is an example of a compound (VDEHD- vinylphenyl sulfone) having a sequence (i.e. P5 is Cbz Val; P4 is Asp; P3 is Glu; P2 is His and D is Asp. The suicide substrate is vinyl phenyl sulfone)., which is designed to selectively inhibit caspase-2:

**Design for selective Caspase-1 inhibitors**

[0145] In one embodiment, the method comprises synthesizing compounds having the following general **Formula D4**:



where

P2 is selected from the following amino acids: V, A, T, and H.

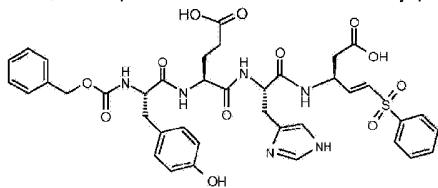
P3 is selected from the following amino acids: E, Q, D, A, G, T, V, Ala(2'-quinolyl), indanylglycine, and W;

P4 is selected from the following amino acids: Y, W, F, and D;

D is the side chain of (D) or (L) aspartic acid; and

the suicide substrate is selected from the group consisting of vinyl phenyl sulfone.

[0146] The following is an example of a compound (YEHD- vinylphenyl sulfone) having a sequence (i.e. P4 is Cbz Tyr; P3 is Glu; P2 is His; D is Asp and the suicide substrate is vinyl phenyl sulfone, which is designed to selectively inhibit caspase-1:



D) Pharmaceutical applications

[0147] As indicated hereinbefore and exemplified hereinafter, the compounds of the invention have beneficial pharmaceutical properties and these compounds may have pharmaceutical applications in the prevention and/or treatment of various diseases and conditions in a subject. Medical and pharmaceutical applications contemplated by the inventors include caspase-mediated diseases. In addition, the compounds of the present invention may have useful benefits on cells *in vitro* such as promoting cell survival or the health of the cells.

[0148] The term "subject" includes living organisms in which blood disorders, renal failure, inflammatory-related diseases associated with high blood pressure, and/or oxidative stress-related disorders, can occur, or which are susceptible to such conditions. The term "subject" includes animals (e.g., mammals (e.g., cats, dogs, horses, pigs, cows, goats, sheep, rodents (e.g., mice or rats), rabbits, squirrels, bears, primates (e.g., chimpanzees, monkeys, gorillas, and humans))), as well as avians (e.g. chickens, ducks, Peking ducks, geese), and transgenic species thereof. Preferably, the subject is a mammal. More preferably, the subject is a human. Even more preferably, the subject is a human patient in need of treatment.

[0149] The term "caspase-mediated disease" includes all diseases, disorder and/or conditions in which any one or more of caspase-1, -2, -3, -4, -5, -6, -7, -8, -9, -10, -11, -12, plays a significant role. In some embodiments, the caspase-mediated disease mainly involves executioner caspases (caspase-3, 6, 7). In another embodiment, the caspase-mediated disease mainly involves initiators (caspase-2, 8, 9, 10). In some embodiments, a compound of the invention shows a high specificity towards one particular caspase. In another embodiment, a compound of the invention is able to inhibit two groups of caspases. Yet, in another embodiment, a compound of the invention even is able to inhibit two specific caspases belonging to two different groups of caspases.

[0150] Examples of caspase-mediated disease according to the invention include apoptosis mediated diseases, IL-1 mediated diseases, inflammatory diseases, autoimmune diseases, autoinflammatory diseases, proliferative diseases, infectious diseases, degenerative diseases, retinal disorders, inflammatory peritonitis, osteoarthritis, pancreatitis, asthma, respiratory distress syndrome, rheumatoid arthritis, systemic lupus erythematosus, scleroderma, Grave's disease, autoimmune gastritis, diabetes, autoimmune hemolytic anemia, autoimmune neutropenia, thrombocytopenia, hepatitis, inflammatory bowel disease, crohn's disease, psoriasis, dermatitis, Graft vs host disease, organ transplant rejection, osteoporosis, leukemias and related disorders, multiple myeloma-related diseases, metastatic melanomas, Kaposi's sarcoma, sepsis, septic shock, Alzheimer's disease, Parkinson's disease, Huntington's disease, cerebral ischemia, epilepsy, myocardial ischemia, acute and chronic heart disease, myocardial infarction, congestive heart failure, atherosclerosis, spinal muscular atrophy, amyotrophic lateral sclerosis, multiple sclerosis, HIV-related encephalitis, aging, neurological damage due to stroke, ulcerative colitis, traumatic brain injury, spinal cord injury, hepatitis-B, hepatitis-C, hepatitis-G, liver-related diseases, renal disease, and HIV infection.

[0151] As used herein, "preventing" or "prevention" is intended to refer to at least the reduction of likelihood of the risk of (or susceptibility to) acquiring a disease or disorder (i.e., causing at least one of the clinical symptoms of the disease not to develop in a patient that may be exposed to or predisposed to the disease but does not yet experience or display symptoms of the disease). Biological and physiological parameters for identifying such patients are provided herein and are also well known by physicians.

[0152] The terms "treatment" or "treating" of a subject includes the application or administration of a compound of the invention to a subject (or application or administration of a compound of the invention to a cell or tissue from a subject) with the purpose of stabilizing, curing, healing, alleviating, relieving, altering, remedying, less worsening, ameliorating, improving, or affecting the disease or condition, the symptom of the disease or condition, or the risk of (or susceptibility to) the disease or condition. The term "treating" refers to any indicia of success in the treatment or amelioration of an injury, pathology or condition, including any objective or subjective parameter such as abatement; remission; lessening of the rate of worsening; lessening severity of the disease; stabilization, diminishing of symptoms or making the injury, pathology or condition more tolerable to the subject; slowing in the rate of degeneration or decline; making the final point of degeneration less debilitating; or improving a subject's physical or mental well-being. In some embodiments, the term "treating" can include increasing a subject's life expectancy and/or delay before additional treatments are required.

[0153] Addressing caspase-mediated diseases is among the medical and pharmaceutical applications contemplated by present invention. Therefore, in one of its aspects the present invention relates to compounds and compositions for use in the prevention and/or treatment of a caspase-mediated disease in a subject, preferably a human patient in need thereof.

[0154] Also described herein are methods for the prevention and/or treatment of a caspase-mediated disease in a subject, preferably a human patient in need thereof.

[0155] Another aspect of the invention relates to the use of the compounds described herein for inhibiting a caspase or a caspase-like protein in a cell, comprising contacting the caspase or caspase-like protein with an effective amount of a caspase inhibitor according to the invention.

[0156] In some embodiments, the subject may be suffering from a viral infection. Therefore, the invention also relates to caspase inhibitors according to the invention (or pharmaceutical compositions comprising the same) for use in the prophylaxis or therapy of a viral infection. This may be helpful for inhibiting a cellular caspase thereby inhibiting virus multiplication.

[0157] Also described herein is a method for the prophylaxis or therapy of a viral infection, comprising administering to a subject in need thereof an effective dose of a caspase inhibitor according to the invention (or a pharmaceutical composition comprising the same). This may be helpful for inhibiting a cellular caspase thereby inhibiting virus multiplication.

[0158] Also of particular interest is the caspase inhibitors according to the invention (or pharmaceutical compositions comprising the same) for use in the treatment of excessive apoptosis affected by caspase activity in a cell or a tissue.

[0159] Also described herein is a method for use in the treatment of excessive apoptosis affected by caspase activity in a cell or a tissue, comprising contacting a cell or tissue with an effective amount of one or more caspase inhibitors according to the invention (or a pharmaceutical composition comprising the same).

[0160] Also of particular interest is the caspase inhibitors according to the invention (or pharmaceutical compositions comprising the same) for use in stimulating stem cell proliferation by preventing some of the stem cells from entering a partial or complete apoptosis cycle.

[0161] It is conceivable that the compounds of the invention be also effective in inhibiting additional families of proteases, including, serine peptidases, cysteine peptidases, aspartic peptidases, metallo-peptidases, and other peptidases of unknown catalytic type. For a more elaborate listing of proteases that may be inhibited by the compounds defined herein, see ZBIGNIEW GRZONKA. Cysteine protease. Industrial Enzymes, 181-195, Chapter 11, 2007 Springer.

[0162] In order to evaluate, assess, and/or confirm the efficacy of the compounds and/or compositions of the invention, serial measurements can be determined. Quantitative assessment of caspase functions and parameters of caspase dysfunction are well known in the art. Examples of assays for the determination of caspases activity are provided in the Exemplification section.

[0163] The compounds according to the invention can be further analyzed, tested or validated for their ability to cross the Blood Brain Barrier BBB as so desired. Many *in-vitro*, *in-vivo* and *in-silico* methods may be employed during drug development to mimic the BBB (Lohmann et al. (2002) Predicting blood-brain barrier permeability of drugs: evaluation of different in vitro assays. J Drug Target 10:263-276; Nicolazzo et al. (2006) Methods to assess drug permeability across the blood-brain barrier. J Pharm Pharmacol 58:281-293). *In-vitro* models include primary endothelial cell culture and immortalized cell lines such as Caco-2, BMEC, MDCK. These cells are useful as a screening method and can appropriately rank compounds in order of BBB permeability. *In vivo* models such as the internal carotid artery single injection or perfusion, intravenous bolus injection, brain efflux index and intracerebral microdialysis provide more accurate information regarding brain uptake, and these can be complemented with novel imaging techniques (such as magnetic resonance imaging and positron emission tomography), although such methods are not suited to high-throughput permeability assessment.

[0164] In certain instances described herein, at least some of the prodrugs administered generates the corresponding pharmaceutical compound only after absorption by the gastrointestinal tract and/or only once it has reached the brain, i.e. after it has passed the blood brain barrier (BBB).

E) Pharmaceutical compositions and formulations

[0165] A related aspect of the invention concerns pharmaceutical compositions comprising one or more of the compounds of the invention described herein. As indicated hereinbefore, the compounds of the invention may be useful in preventing and/or treating caspase-mediated disease, and more particularly sepsis, myocardial infarction, ischemic stroke, spinal cord injury (SCI), traumatic brain injury (TBI) and neurodegenerative disease (e.g. multiple sclerosis (MS) and Alzheimer's, Parkinson's, and Huntington's diseases).

[0166] As used herein, the term "*therapeutically effective amount*" means the amount of compound that, when administered to a subject for treating or preventing a particular disorder, disease or condition, is sufficient to effect such treatment or prevention of that disorder, disease or condition. Dosages and therapeutically effective amounts may vary for example, depending upon a variety of factors including the activity of the specific agent employed, the age, body weight, general health, gender, and diet of the subject, the time of administration, the route of administration, the rate of excretion, and any drug combination, if applicable, the effect which the practitioner desires the compound to have upon the subject and the properties of the compounds (e.g. bioavailability, stability, potency, toxicity, etc), and the particular disorder(s) the subject is suffering from. In addition, the therapeutically effective amount may depend on the subject's blood parameters (e.g. lipid profile, insulin levels, glycemia), the severity of the disease state, organ function, or underlying disease or complications. Such appropriate doses may be determined using any available assays including the assays described herein. When one or more of the compounds of the invention is to be administered to humans, a physician may for

example, prescribe a relatively low dose at first, subsequently increasing the dose until an appropriate response is obtained.

[0167] As used herein, the term "*pharmaceutical composition*" refers to the presence of at least one compound of the invention according to any one of Formula IA, II, IIA, III, or IIIA as defined herein and at least one pharmaceutically acceptable vehicle. Examples of representative compounds of the invention include the compounds in **Table 1** and pharmaceutically acceptable salts thereof.

[0168] "*Pharmaceutically acceptable vehicle*" refers to a diluent, adjuvant, excipient, or carrier with which a compound is administered. The term "pharmaceutically acceptable" refers to drugs, medicaments, inert ingredients etc., which are suitable for use in contact with the tissues of humans and lower animals without undue toxicity, incompatibility, instability, irritation, allergic response, commensurate with a reasonable benefit/risk ratio. It preferably refers to a compound or composition that is approved or approvable by a regulatory agency of the Federal or state government or listed in the U.S. Pharmacopoeia or other generally recognized pharmacopoeia for use in animals and more particularly in humans. The pharmaceutically acceptable vehicle can be a solvent or dispersion medium containing, for example, water, ethanol, polyol (for example, glycerol, propylene glycol, and liquid polyethylene glycol), suitable mixtures thereof, and vegetable oils. Additional examples of pharmaceutically acceptable vehicles include: Water for Injection USP; aqueous vehicles such as, Sodium Chloride Injection, Ringer's Injection, Dextrose Injection, and Sodium Chloride Injection; and Lactated Ringer's Injection; water-miscible vehicles such as, ethyl alcohol, polyethylene glycol, and polypropylene glycol; and non-aqueous vehicles such as, corn oil, cottonseed oil, peanut oil, sesame oil, ethyl oleate, isopropyl myristate, and benzyl benzoate. Prevention of the action of microorganisms can be achieved by various antibacterial and antifungal agents, for example, parabens, chlorobutanol, phenol, ascorbic acid, thimerosal. In many cases, isotonic agents are included, for example, sugars, sodium chloride, or polyalcohols such as mannitol and sorbitol, in the composition. Prolonged absorption of injectable compositions can be brought about by including in the composition an agent which delays absorption, for example, aluminum monostearate or gelatin. Nanoparticles, liposomes, and antibodies conjugated to nanoparticles or combinations thereof, are also contemplated as pharmaceutically acceptable vehicles.

[0169] In some embodiments, the compositions of the invention comprise an effective amount of a compound of the Formula IA, II, IIA, III, or IIIA as described hereinbefore, preferably compound Z-Asp-Indanylglycine-Val-Aspmethyl vinyl sulfone (57); Z-Asp-Glu-Val-Aspmethyl vinyl sulfone (59) Z-Asp-Ala(2'-quinolyl)-Val-Aspmethyl vinyl sulfone (55); Z-Asp-Phg-Val-Aspmethyl vinyl sulfone (53); Z-Asp-Ala(2'-quinolyl)-Val-Asp- α -chlorovinyl methylsulfone (48); Z-Asp(β -methyl)-Indanylglycine-Val-Asp(β -methyl)methyl vinyl sulfone (51) Z-Asp-Tyr-Val-Aspmethyl vinyl sulfone (76) Z-Asp-Trp-Val-Aspmethyl vinyl sulfone (88) or a pharmaceutically acceptable salt thereof.

[0170] In some embodiments the invention pertains to pharmaceutical compositions for preventing and/or treating diseases or other medical conditions in which at least one caspase is significantly involved that include one or more compounds of Formula IA, II, IIA, III, or IIIA as defined herein.

[0171] In some embodiments the invention pertains to pharmaceutical compositions for preventing and/or treating diseases or other medical conditions in which at least one caspase is significantly involved, the composition comprising one or more compounds of Formula IA, II, IIA, III, or IIIA as defined herein.

[0172] The compounds of the invention may be formulated prior to administration into pharmaceutical compositions using available techniques and procedures. For instance, the pharmaceutical compositions may be formulated into suitable administration (orally, parenterally, (intravascular (IV), intraarterial (IA), intramuscular (IM), depo-IM, subcutaneous (SC), and depo SC), sublingually, intranasally (inhalation), intrathecally, topically, or rectally.

[0173] Preferably, the compound(s) of the invention can be orally administered. The formulations may conveniently be presented in unit dosage form and may be prepared by any methods well known in the art of pharmacy. Methods of preparing these formulations or compositions include the step of bringing into association a compound of the present invention with a pharmaceutically acceptable vehicle (e.g. an inert diluent or an assimilable edible carrier) and, optionally, one or more accessory ingredients. In general, the formulations are prepared by uniformly and intimately bringing into association a compound of the present invention with liquid carriers, or finely divided solid carriers, or both, and then, if necessary, shaping the product. The amount of the therapeutic agent in such therapeutically useful compositions is such that a suitable dosage will be obtained.

[0174] Formulations of the invention suitable for oral administration may be in the form of capsules (e.g. hard or soft shell gelatin capsule), cachets, pills, tablets, lozenges, powders, granules, pellets, dragees, e.g., coated (e.g., enteric coated) or uncoated, or as a solution or a suspension in an aqueous or non-aqueous liquid, or as an oil-in-water or water-in-oil liquid emulsion, or as an elixir or syrup, or as pastilles or as mouth washes, each containing a predetermined amount of a compound of the present invention as an active ingredient. A compound of the present invention may also be administered as a bolus, electuary or paste, or incorporated directly into the subject's diet. Moreover, in certain embodiments these pellets can be formulated to (a) provide for instant or rapid drug release (i.e., have no coating on them); (b) be coated, e.g., to provide for sustained drug release over time; or (c) be coated with an enteric coating for better gastrointestinal tolerability. Coating may be achieved by conventional methods, typically with pH or time-dependent coatings, such that the compound(s) of the invention is released in the vicinity of the desired location, or at various times to

extend the desired action. Such dosage forms typically include one or more of cellulose acetate phthalate, polyvinylacetate phthalate, hydroxypropyl methyl cellulose phthalate, ethyl cellulose, waxes, and shellac.

[0175] In solid dosage forms for oral administration a compound of the present invention may be mixed with one or more pharmaceutically acceptable carriers, such as sodium citrate or dicalcium phosphate, or any of the following: fillers or extenders, such as starches, lactose, sucrose, glucose, mannitol, or silicic acid; binders, such as, for example, carboxymethylcellulose, alginates, gelatin, polyvinyl pyrrolidone, sucrose or acacia; humectants, such as glycerol; disintegrating agents, such as agar-agar, calcium carbonate, potato or tapioca starch, alginic acid, certain silicates, and sodium carbonate; solution retarding agents, such as paraffin; absorption accelerators, such as quaternary ammonium compounds; wetting agents, such as, for example, cetyl alcohol and glycerol monostearate; absorbents, such as kaolin and bentonite clay; lubricants, such as talc, calcium stearate, magnesium stearate, solid polyethylene glycols, sodium lauryl sulfate, and mixtures thereof; and coloring agents. In the case of capsules, tablets and pills, the pharmaceutical compositions may also comprise buffering agents. Solid compositions of a similar type may also be employed as fillers in soft and hard-filled gelatin capsules using such excipients as lactose or milk sugars, as well as high molecular weight polyethylene glycols.

[0176] Peroral compositions typically include liquid solutions, emulsions, suspensions. The pharmaceutically acceptable vehicles suitable for preparation of such compositions are well known in the art. Typical components of carriers for syrups, elixirs, emulsions and suspensions include ethanol, glycerol, propylene glycol, polyethylene glycol, liquid sucrose, sorbitol and water. For a suspension, typical suspending agents include methyl cellulose, sodium carboxymethyl cellulose, tragacanth, and sodium alginate; typical wetting agents include lecithin and polysorbate 80; and typical preservatives include methyl paraben and sodium benzoate. Peroral liquid compositions may also contain one or more components such as sweeteners, flavoring agents and colorants disclosed above.

[0177] Pharmaceutical compositions suitable for injectable use may include sterile aqueous solutions (where water soluble) or dispersions and sterile powders for the extemporaneous preparation of sterile injectable solutions or dispersions. In all cases, the composition must be sterile and must be fluid to the extent that easy syringability exists. It must be stable under the conditions of manufacture and storage and must be preserved against the contaminating action of microorganisms such as bacteria and fungi. Sterile injectable solutions can be prepared by incorporating the therapeutic agent in the required amount in an appropriate solvent with one or a combination of ingredients enumerated above, as required, followed by filtered sterilization. Generally, dispersions are prepared by incorporating the therapeutic agent into a sterile vehicle which contains a basic dispersion medium and the required other ingredients from those enumerated above. In the case of sterile powders for the preparation of sterile injectable solutions, the methods of preparation are vacuum drying and freeze-drying which yields a powder of the active ingredient (*i.e.*, the therapeutic agent) plus any additional desired ingredient from a previously sterile-filtered solution thereof.

[0178] Pharmaceutical formulations are also provided which are suitable for administration as an aerosol, by inhalation. These formulations comprise a solution or suspension of the desired compound of any Formula herein or a plurality of solid particles of such compound(s). The desired formulation may be placed in a small chamber and nebulized. Nebulization may be accomplished by compressed air or by ultrasonic energy to form a plurality of liquid droplets or solid particles comprising the agents or salts. The liquid droplets or solid particles should have a particle size in the range of about 0.5 to about 5 microns. The solid particles can be obtained by processing the solid agent of any Formula described herein, or a salt thereof, in any appropriate manner known in the art, such as by micronization. The size of the solid particles or droplets will be, for example, from about 1 to about 2 microns. In this respect, commercial nebulizers are available to achieve this purpose. A pharmaceutical formulation suitable for administration as an aerosol may be in the form of a liquid, the formulation will comprise a water-soluble agent of any Formula described herein, or a salt thereof, in a carrier which comprises water. A surfactant may be present which lowers the surface tension of the formulation sufficiently to result in the formation of droplets within the desired size range when subjected to nebulization.

[0179] The compositions of this invention can also be administered topically to a subject, *e.g.*, by the direct laying on or spreading of the composition on the epidermal or epithelial tissue of the subject, or transdermally *via* a "patch". Such compositions include, for example, lotions, creams, solutions, gels and solids. These topical compositions may comprise an effective amount, usually at least about 0.1%, or even from about 1% to about 5%, of a compound of the invention. Suitable carriers for topical administration typically remain in place on the skin as a continuous film, and resist being removed by perspiration or immersion in water. Generally, the carrier is organic in nature and capable of having dispersed or dissolved therein the therapeutic agent. The carrier may include pharmaceutically acceptable emollients, emulsifiers, thickening agents and solvents.

[0180] Other compositions useful for attaining systemic delivery of the subject agents include sublingual, buccal and nasal dosage forms. Such compositions typically comprise one or more of soluble filler substances such as sucrose, sorbitol and mannitol; and binders such as acacia, microcrystalline cellulose, carboxymethyl cellulose and hydroxypropyl methyl cellulose. Glidants, lubricants, sweeteners, colorants, antioxidants and flavoring agents disclosed above may also be included.

[0181] The compound(s) of the invention may also be administered parenterally, intraperitoneally, intravenously, intraspinally, intrathecally or intracerebrally. For such compositions, the compound(s) of the invention can be prepared in glycerol, liquid polyethylene glycols, and mixtures thereof and in oils. Under ordinary conditions of storage and use, these preparations may contain a preservative to prevent the growth of microorganisms.

[0182] The use of the compounds according to the invention may also include co-administration of the at least one compound according to the invention, or a pharmaceutically acceptable salt thereof together with the administration of another therapeutically effective agent. Therefore, an additional aspect of the invention relates to methods of concomitant therapeutic treatment of a subject, comprising administering to a subject in need thereof an effective amount of a first agent and a second agent, wherein the first agent is as defined in Formula IA, II, IIA, III, or IIIA and the second agent is for the prevention or treatment of any one of disorder or disease indicated hereinbefore. As used herein, the term "concomitant" or "concomitantly" as in the phrases "concomitant therapeutic treatment" or "concomitantly with" includes administering a first agent in the presence of a second agent. A concomitant therapeutic treatment method includes methods in which the first, second, third or additional agents are co-administered. A concomitant therapeutic treatment method also includes methods in which the first or additional agents are administered in the presence of a second or additional agents, wherein the second or additional agents, for example, may have been previously administered. A concomitant therapeutic treatment method may be executed step-wise by different actors. For example, one actor may administer to a subject a first agent and as a second actor may administer to the subject a second agent and the administering steps may be executed at the same time, or nearly the same time, or at distant times, so long as the first agent (and/or additional agents) are after administration in the presence of the second agent (and/or additional agents). The actor and the subject may be the same entity (e.g. a human).

[0183] Accordingly, the invention also relates to the use of the compounds according to the invention for preventing, reducing or eliminating a symptom or complication of any one of the above mentioned disease or condition. The use comprises administering, to a subject in need thereof, a first pharmaceutical composition comprising at least one compound of the invention and a second pharmaceutical composition comprising one or more additional active ingredients, wherein all active ingredients are administered in an amount sufficient to inhibit, reduce, or eliminate one or more symptoms or complications of the disease or condition to be treated. In one aspect, the administration of the first and second pharmaceutical composition is temporally spaced apart by at least about two minutes. Preferably the first agent is a compound of Formula IA, II, IIA, III, or IIIA as defined herein, or a pharmaceutically acceptable salt thereof. The second agent may be selected from the following list of compounds: Z-Asp-Phg-Val-Aspmethyl vinyl sulfone (53)

F) Screening assays

[0184] The compounds of the present invention may also be used in screening methods. For instance, these compounds may be used in methods for tracking activity of caspases *in vitro* and/or *in vivo*. The compounds of the present invention may also be helpful for identifying other compounds that bind to a caspase active site. In some embodiments, the compounds of the invention are labeled (e.g. radioactively labeled). Radiolabeled compounds may also be useful in diagnostic assays.

[0185] There are a number of ways in which to determine the binding of a compound of the present invention to the caspase. In one embodiment the caspase is bound to a support, and a labeled compound of the invention is added to the assay. Alternatively, the compound of the invention may be bound to the support and the caspase is added.

[0186] The compounds of the invention may also be used as competitors to screen for additional drug candidates or test compounds. As used herein, the terms "drug candidate" or "test compounds" are used interchangeably and describe any molecule, for example, protein, oligopeptide, small organic molecule, polysaccharide, polynucleotide to be tested for bioactivity.

[0187] Typically, the signals that are detected in the assay (e.g. *in vitro*, *in vivo* and/or diagnostic) may include radioactivity. Detectable labels useful in performing screening assays in this invention include radioisotopes such as tritium and ^{125}I .

[0188] Also described herein are assays where the signals that are detected include fluorescence, resonance energy transfer, time resolved fluorescence, fluorescence polarization, plasma resonance, or chemiluminescence depending on the nature of the label. Detectable labels useful in performing such assays include a fluorescent label such as Fluorescein, Oregon green, dansyl, rhodamine, tetramethyl rhodamine, texas red, Eu^{3+} ; a chemiluminescent label such as luciferase; colorimetric labels; and enzymatic markers. Also described herein are screening assays utilizing affinity tags including biotin and polyhistidine.

F) Kits

[0189] Also described herein are kits comprising the compound(s) of the invention, optionally including a container (e.g. packaging, a box, a vial, etc). The kit may be commercially used according to the methods described herein and may include instructions for use in a method of the invention. Additional kit components may include acids, bases, buffering agents, inorganic salts, solvents, antioxidants, preservatives, or metal chelators. The additional kit components are present as pure compositions, or as aqueous or organic solutions that incorporate one or more additional kit components. Any or all of the kit components optionally further comprise buffers.

[0190] The compound(s) of the invention may or may not be administered to a patient at the same time or by the same route of

administration. Therefore, the methods of the invention encompass kits which, when used by the medical practitioner, can simplify the administration of appropriate amounts of two or more active ingredients to a patient.

[0191] A typical kit comprises a unit dosage form of a at least one compound according to the invention, e.g., a compound of Formula I, IA, II, IIA, III, or IIIA as defined herein or a pharmaceutically acceptable salt thereof, and a unit dosage form of at least one additional active ingredient. Examples of additional active ingredients that may be used in conjunction with the compounds according to the invention, include any of the compounds that could be used in combination with the compound(s) of the invention as indicated herein before.

[0192] Kits can further comprise devices that are used to administer the active ingredients. Examples of such devices include syringes, drip bags, patches, inhalers, enemas, and dispensers for the administration of suppository formulations.

[0193] Kits can further comprise pharmaceutically acceptable vehicles that can be used to administer one or more active ingredients. For example, if an active ingredient is provided in a solid form that must be reconstituted for parenteral administration, the kit can comprise a sealed container of a suitable vehicle in which the active ingredient can be dissolved to form a particulate-free sterile solution that is suitable for parenteral administration. Examples of pharmaceutically acceptable vehicles are provided hereinbefore.

[0194] Headings are included herein for reference and to aid in locating certain sections. These headings are not intended to limit the scope of the concepts described therein under, and these concepts may have applicability in other sections throughout the entire specification.

[0195] Unless otherwise indicated, all numbers expressing quantities of ingredients, reaction conditions, concentrations, properties, and so forth used in the specification and claims are to be understood as being modified in all instances by the term "about." At the very least, each numerical parameter should at least be construed in light of the number of reported significant digits and by applying ordinary rounding techniques. Accordingly, unless indicated to the contrary, the numerical parameters set forth in the present specification and attached claims are approximations that may vary depending upon the properties sought to be obtained. Notwithstanding that the numerical ranges and parameters setting forth the broad scope of the embodiments are approximations, the numerical values set forth in the specific examples are reported as precisely as possible. Any numerical value, however, inherently contain certain errors resulting from variations in experiments, testing measurements, statistical analyses and such.

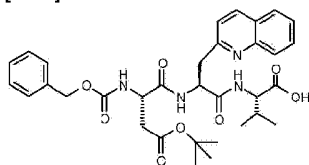
[0196] Those skilled in the art will recognize, or be able to ascertain using no more than routine experimentation, numerous equivalents to the specific procedures, embodiments, claims, and examples described herein. The invention is further illustrated by the following examples.

EXAMPLES

[0197] The Examples set forth herein below provide exemplary methods for the preparation of certain representative compounds encompassed by general Formula IA, II, IIA, III, or IIIA. Some Examples provide exemplary uses of certain representative compounds of the invention. Also provided are exemplary methods for assaying the compounds of the invention for *in vitro* and *in vivo* efficacy.

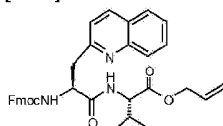
Example I: Synthesis of compound 4 (Cbz-Asp(O-tBu)-Ala(2'-quinolyl)-ValOH)

[0198]



a) Fmoc-Ala(2'-quinolyl)-Val-OAllyl

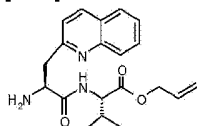
[0199]



[0200] Fmoc-Ala (2'-quinolyl)-OH (0.152 g, 0.347 mmol) was solvated in DMF (1 mL) and CH₂Cl₂ (0.9 mL). L-Val allylester toluene-4-sulfonate (0.115 g, 1.01 eq) in 0.3 mL of CH₂Cl₂ was added, followed with 4-methylmorpholine (0.04 mL, 1.05 eq) and EDC (0.0681 g, 1.02eq). The mixture was stirred for 3 hours, and then it was extracted using CH₂Cl₂/ brine. The organic layer was dried over MgSO₄, filtered off and concentrated to dryness.

b) Ala(2'-quinolyl)-Val-OAllyl

[0201]

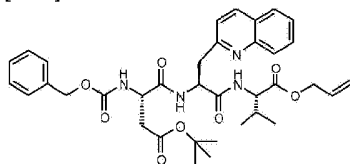


[0202] The Fmoc-Ala(2'-quinolyl)-Val(O-Allyl) (0.425 g, 0.737 mmol) was solvated in CH₂Cl₂ (40 mL) followed by dropwise addition of piperidine (0.6 mL, 8.24 eq). After 40 min the mixture was evaporated to dryness under vacuum. The product was then purified on silica using a gradient of MeOH/ CH₂Cl₂ (0 to 10 %) to get 0.188 g of desired compound.

NMR ¹H (CDCl₃, 400 MHz) δ: 8.18 (s, 1H, NH); 8.08 (d, 1H, J= 8.4 Hz); 8.03 (d, 1H, J= 8.47 Hz); 7.79 (d, 1H, J= 7.91 Hz); 7.71-7.68 (m, 1H); 7.52-7.49 (m, 1H); 7.35 (d, 1H, J= 8.42 Hz); 5.92-5.85 (m, 2H); 5.34-5.22 (m, 2H); 4.63-4.58 (m, 2H); 4.55-4.52 (m, 1H); 3.95-3.93 (m, 1H); 3.52 (dd, 1H, J=3.95 Hz); 3.25-3.20 (m, 1H); 2.20-2.13 (m, 1H); 2.01 (m, 2H, NH₂); 0.85 (m, 6H).

c) Cbz-Asp(O-tBu)-Ala(2'-quinolyl)-Val-OAllyl

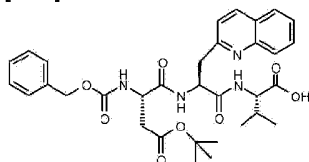
[0203]



[0204] Ala(2'-quinolyl)-Val(O-Allyl) (0.054 g, 0.1519 mmol) was solvated in CH₂Cl₂ (0.7 mL) followed by addition of L-Asp (O-tBu)-OH (0.053 g, 1.08 eq) and 4- methylmorpholine (0.018 mL, 1.08 eq) and finally DPC (0.025 mL, 1.06 eq). The mixture was stirred for 1 hour. It was then subjected to a liquid extraction dichloromethane / brine. The organic layer was dried over MgSO₄, concentrated to vacuum and purified on silica using a gradient of MeOH/ CH₂Cl₂ (0 to 10 %) to get 0.084 g of the desired compound.

d) Cbz-Asp(O-tBu)-Ala(2'-quinolyl)-ValOH

[0205]



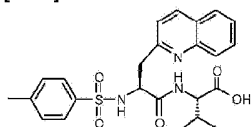
[0206] Cbz-Asp (OtBu)-Ala(2'-quinolyl)-Val (OAllyl) (0.082 g, 0.1241 mmol) was dissolved in THF (3.5 mL) and rotary evaporated to dryness, the sample was then redissolved in THF (3.5 mL) and vacuum aspirated (3* 1 min of aspiration), followed by replacement of the atmosphere with Argon. Morpholine (0.04 mL, 3.7 eq) was added, followed by Pd(PPh₃)₄ (0.0171 g, 0.12 eq). The sample flask was then covered with a tin foil and kept under stirring for 3 days under Argon. The compound was evaporated to dryness and the obtained residue was subjected to purification on C₁₈ using a gradient MeOH/solution of H₂O at pH= 3.5 (0 to 100 %) to get 0.065 g of the desired compound.

NMR ¹H (CD₃OD, 400 MHz) δ: 8.18 (d, 1H, J= 8.4 Hz); 8.02 (d, 1H, J= 8.49 Hz); 7.84 (d, 1H, J= 8.08 Hz); 7.63 (t, 1H, J= 7.51 Hz);

7.52 (t, 1H, $J = 7.39$ Hz); 7.41 (d, 1H, $J = 8.3$ Hz); 7.40-7.30 (m, 5H); 5.03-4.93 (m, 3H); 4.50-4.48 (m, 1H); 4.26 (d, 1H, $J = 4.9$ Hz); 3.48-3.34 (m, 2H); 2.73-2.68 (m, 1 H); 2.54-2.49 (m, 1 H); 2.15-2.11 (m, 1 H); 1.36 (m, 9H); 0.87-0.84 (m, 6H).

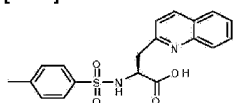
Example 2: Synthesis of compound 5 (Ts-Ala(2'-quinolyl)-Val-OH)

[0207]



a) Ts-Ala(2'-quinolyl)-OH

[0208]

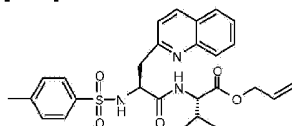


[0209] To L-Ala(2'-quinolyl)-OH (0.060 g, 0.277 mmol) was added H₂O (0.4 mL) and THF (0.15 mL). The mixture was stirred for 2 min before adding TEA (0.074 ml, 1.93 eq). The mixture was allowed to reach 0 °C, before adding Tosyl chloride (0.052 g, 1 eq) in THF (0.4 mL) in a dropwise manner. The mixture was then allowed to warm to room temperature and stirred for 16 hours. The mixture was diluted with EtOAc (8 mL) and H₂O (2 mL), then it was acidified with HCl 1N (dropwise addition) to reach pH 3/4. The mixture was extracted with EtOAc, dried over MgSO₄ and evaporated to dryness, to get upon addition of MeOH (0.5 mL) 30 mg of precipitate; an other portion could be obtained from the solution.

NMR ¹H (DMSO, 400 MHz) δ : 8.30-8.23 (m, 1 H, NH); 8.14 (d, 1 H, $J = 8.37$ Hz); 7.89 (d, 1H, $J = 8.0$ Hz); 7.78 (d, 1H, $J = 8.39$ Hz); 7.70 (t, 1H, $J = 7.2$ Hz); 7.55 (t, 1H, $J = 7.33$ Hz); 7.32-7.27 (m, 3H); 6.88 (d, 2H, $J = 8.05$ Hz); 4.28 (s, 1H); 3.4-3.03 (m, 2H); 2.13 (s, 3H).

b) Ts-Ala(2'-quinolyl)-Val-OAllyl

[0210]



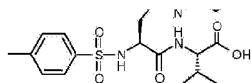
[0211] Ts- Ala(2'-quinolyl)-OH (0.027g, 0.076 mmol) was solvated in CH₂Cl₂ followed by addition of L- Val-OAllyl ester toluene-4-sulfonate (0.025 g, 1.02 eq), 4-methylmorpholine (0.017 mL, 1.91 eq), DMAP (0.0012 g, 0.13 eq) and finally EDC (0.015g, 1.04 eq). The progress of the reaction was followed by TLC. The mixture was extracted with CH₂Cl₂/ brine. The organic layer was dried over MgSO₄, filtered off and concentrated to dryness. The obtained residue was purified on silica using a gradient EtOAc/Hexane (5 to 60%) to get 10 mg of the desired compound.

NMR ¹H (CDCl₃, 400 MHz) δ : 7.99 (m, 2H); 7.89 (d, 1H, $J = 8.93$ Hz); 7.76 (d, 1H, $J = 8.06$ Hz); 7.72 (t, 1H, $J = 8.16$ Hz); 7.64 (m, 2H); 7.53 (t, 1H, $J = 7.61$ Hz); 7.45 (d, 1H, $J = 6.86$ Hz); 7.16 (d, 1H, $J = 8.34$ Hz); 7.08 (m, 2H); 5.86-5.79 (m, 1H); 5.27 (dd, 1H, $J = 17.25$ Hz, $J = 1.36$ Hz); 5.20 (dd, 1H, $J = 10.4$ Hz, $J = 1.0$ Hz); 4.58-4.5 (m, 2H); 4.38 (dd, 1H, $J = 8.78$ Hz, $J = 5.02$ Hz); 4.22 (q, 1H, $J = 5.58$ Hz); 3.41 (dd, 1H, $J = 15.50$ Hz, $J = 5.23$ Hz); 3.14 (dd, 1H, $J = 15.53$ Hz, $J = 5.64$ Hz); 2.33 (s, 3H); 2.16-2.00 (m, 1H); 0.72 (d, 3H, $J = 6.8$ Hz); 0.65 (d, 3H, $J = 6.85$ Hz).

c) Ts-Ala(2'-quinolyl)-Val-OH

[0212]



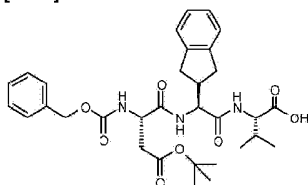


[0213] The Ts-Ala(2'-quinoyl)-OAllyl (0.010 g, 0.02 mmol) was dissolved in THF (3.5 mL) and rotary evaporated to dryness, the sample was then redissolved in THF (3.5 mL) and vacuum aspirated (3* 1 min of aspiration), followed by replacement of the atmosphere with Argon. Pd(PPh₃)₄ (0.0036 g, 0.16 eq) was added under Argon, followed by Morpholine (0.007mL, 4.1 eq). The sample flask was then covered with a tin foil and kept under stirring for 3 days under Argon. The compound was evaporated to dryness and the obtained residue was subjected to purification on C₁₈ using a gradient: MeOH/H₂O (15 to 100 %) to get 0.006 g of the desired compound.

NMR ^1H (CD_3OD , 400 MHz) δ : 8.05 (d, 1H, J = 8.44 Hz); 7.88-7.84 (m, 2H); 7.76-7.72 (m, 1H); 7.58-7.55 (m, 1H); 7.33-7.29 (m, 3H); 6.78-6.76 (m, 2H); 4.35 (dd, 1H, J = 10.43 Hz, J = 3.38 Hz); 4.19 (d, 1H, J = 4.6 Hz); 3.44-3.03 (m, 2H); 2.20-2.12 (m, 1H); 2.12 (s, 3H); 0.93-0.89 (m, 6H).

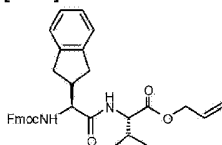
Example 3: Sythesis of compound 12 (Cbz-Asp(O-tBu)-Indanylglycine-Val-OH)

[0214]



a) Fmoc-Indanylglycine-Val-OAllyl

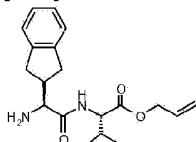
[0215]



[0216] Fmoc Indanylglycine (0.54 g, 1.315 mmol) was solvated in CH₂Cl₂ (5 mL) and DMF (1.8 mL) followed with the addition of L-Val allylester (0.437 g, 1 eq), 4-methylmorpholine (0.15 ml, 1.04 eq) and 4 min latter DMAP (14.5 mg, 0.09 eq) then EDC (0.265 g, 1.05 eq). The mixture was stirred for 1 hour 45 min. Then it was extracted using EtOAc/ brine. The organic layer was dried over MgSO₄, filtered off and concentrated to dryness.

b) Indanylglycine-Val-OAllyl

[0217]



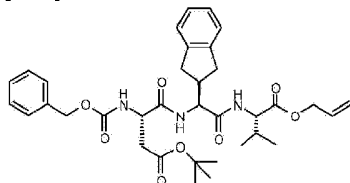
[0218] Fmoc- Indanylglycine-Val-OAllyl (0.758 g, 1.31 mmol) was solvated in CH₂Cl₂ (5mL) and DMF (1.8 mL) followed by dropwise addition of piperidine (1.05 mL, 8.1 eq) over 30 seconds. After 45 min, the mixture was subjected to extraction CH₂Cl₂/ brine (30/15 mL) and saturated NH₄Cl (5 mL). The organic layer was dried over MgSO₄, concentrates and purified by silica using a gradient of MeOH/ CH₂Cl₂ (0 to 5 %) to get 0.45 g of the desired compound.

NMR ^1H (CDCl_3 , 400 MHz) δ : 7.80 (d, 1H, NH); 7.20-7.17 (m, 2H); 7.15-7.11 (m, 2H); 5.96-5.88 (m, 1H); 5.37-5.25 (m, 2H); 4.68-4.58

(m, 3H); 3.53 (d, 1H, $J = 5.05$ Hz); 3.12-2.8 (m, 5H); 2.27-2.20 (m, 1H); 0.98 (d, 3H, $J = 6.85$ Hz); 0.94 (d, 3H, $J = 6.88$ Hz).

c) Cbz-Asp(O-*t*Bu)-Indanylglycine-Val-OAllyl

[0219]

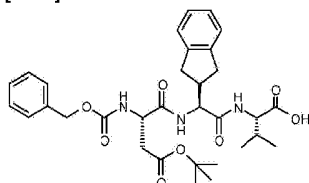


[0220] Indanylglycine-Val-OAllyl (0.46 g, 1.3 mmol) and Z-L-Asp(O-*t*Bu)-OH (0.425 g, 1.3 mmol) were solvated in CH_2Cl_2 (4.5 mL) followed by addition of 4-methylmorpholine (1.45 mL, 1.01 eq), DMAP (14 mg, 0.09 eq) then EDC (0.251 g, 1.01 eq). The mixture was stirred for 1 hour 40 min. Then it was extracted using CH_2Cl_2 /brine. The organic layer was dried over MgSO_4 , concentrates and purified by silica using a gradient of EtOAc/Hexane (10 to 80 %) to get 0.578 g of the desired compound.

NMR ^1H (CD_3OD , 400 MHz) δ : 7.35-7.25 (m, 5H); 7.15-7.07 (m, 4H); 6.0-5.92 (m, 1H); 5.36 (dd, 1H, $J = 17.17$ Hz, $J = 1.429$ Hz); 5.24 (dd, 1H, $J = 10.45$ Hz, $J = 1.251$ Hz); 5.09-5.036 (m, 2H); 4.77-4.61 (m, 2H); 4.54-4.50 (m, 2H); 4.33 (d, 1H, $J = 6.04$ Hz); 2.97-2.52 (m, 7H); 2.20-2.011 (m, 1H); 1.41 (s, 9H); 0.97 (d, 3H, $J = 1.78$ Hz); 0.96 (d, 3H, $J = 1.78$ Hz).

d) Cbz-Asp(O-*t*Bu)Indanylglycine-Val-OH

[0221]

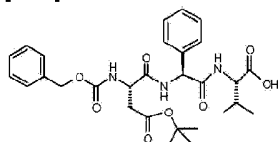


[0222] Z-Asp (O-*t*Bu)-Indanylglycine-Val-(OAllyl) (0.102 g, 0.1611 mmol) was dissolved in THF (6 mL) and rotary evaporated to dryness, the sample was then redissolved in THF (6 mL) and vacuum aspirated (3* 1 min of aspiration), followed by replacement of the atmosphere with Argon. $\text{Pd}(\text{PPh}_3)_4$ (0.017 g, 0.151 0.094 eq) was added in one shot, the flask was then evacuated with Argon. Morpholine (0.06 mL, 4.28 eq) was added and the flask covered with a tin foil. The mixture was kept under stirring for 2.5 days under Argon. The sample was evaporated to dryness and the obtained residue was subjected to purification on C_{18} using a gradient MeOH/ solution of H_2O at pH= 3.5 (10 to 100 %) to get 0.046 g of the desired compound.

NMR ^1H (CD_3OD , 400 MHz) δ : 7.4-7.23 (m, 5H); 7.18-7.03 (m, 4H); 5.20-5.0 (m, 2H); 4.58-4.50 (m, 2H); 4.3 (d, 1H, $J = 5.5$ Hz); 3.0-2.7 (m, 6H); 2.6-2.50 (m, 1H); 1.4 (s, 9H); 0.97 (d, 3H, $J = 2.06$ Hz); 0.96 (d, 3H, $J = 2.18$ Hz).

Example 4: Synthesis of compound 16 (Z- Asp(β -tert-butyl)-Phg-Val -OH)

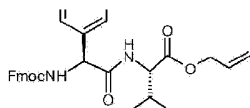
[0223]



a) Fmoc-Phg-Val-OAllyl

[0224]

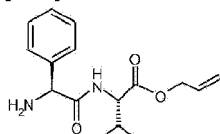




[0225] Fmoc-PhgOH (1g, 2.678 mmol) was dissolved in a mix of anhydrous dichloromethane and DMF (9 ml / 0.5 ml). Val(OAllyl) tosyl salt form (0.804 g, 2.44 mmol) in 1 ml of dichloromethane was added followed by diisopropylcarbodiimide (0.414 ml, 2.44 mmol) and N-methyl morpholine (0.270 ml, 2.44 mmol). The mixture was stirred for 2.20 hrs at RT, then filtered off on a path of celite (1 cm) and washed with dichloromethane. The filtrate was concentrated and the obtained residue was used as a crude material for the next step.

b) Phg-Val-OAllyl

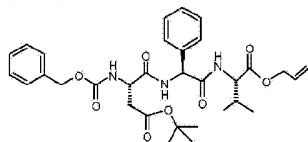
[0226]



[0227] 1.45 g of the previous crude material (Fmoc-Phg-Val -OAllyl) was dissolved in a solution of 20% piperidine in dichloromethane (8.5 ml) and stirred at room temperature for 45 minutes. The mixture was then concentrated under vacuum, diluted with dichloromethane and filtered off through a path of celite (1 cm). The solvent was evaporated to dryness and then purified on silica gel, eluting first with ethyl acetate/hexane (20 %) followed with a gradient of dichloromethane/methanol (0 to 10%) to get 0.72 g of the N-unprotected peptide Phg-Val -OAllyl.

c) Z-Asp(β-tert-Butyl)- Phg-Val-OAllyl.

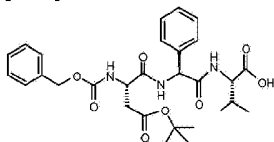
[0228]



[0229] The Phg-Val-OAllyl (0.89 g, 2.74 mmol) was dissolved in anhydrous dichloromethane (7 ml). Then Z-Asp(β-tert-butyl)-OH (0.89g, 2.74 mmol) in dichloromethane (2 ml) was added followed by diisopropyl carbodiimide (0.424 ml, 2.74 mmol). The mixture was stirred at RT for 2.45 hrs and then diluted with dichloromethane. The organic phase was washed twice with brine and dried over anhydrous magnesium sulphate. The solvent was evaporated and the obtained residue was purified on silica gel (gradient: ethyl acetate / hexane) to afford 0.4 g of Z-Asp(OtBu)-Phg-Val-OAllyl.

d) Z-Asp(β-tert-Butyl)-Phg-Val-OH.

[0230]



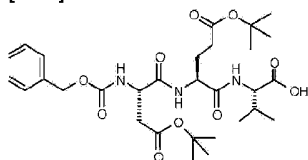
[0231] Z-Asp(β-tert-butyl)-Phg-Val-OAllyl (0.064 g; 0.107 mmol) was dissolved under argon in dry THF (3 ml, inhibitor free). The solvent was degassed three times under argon before adding morpholine (28 ul, 3 eq), followed by Tetrakis (13 mg). The mixture was stirred for 3.5 days at room temperature. The mixture was then concentrated under vacuum (12 mbar) and purified on silica gel (gradient: methanol/dichloromethane: 1 to 14 %) to afford 46 mg of the desired Z-Asp(β-tert-Butyl)-Phg-Val-OH.

NMR ^1H (CD_3OD , 400 MHz) δ 7.43 (m, 2 H); 7.34-7.26 (m, 8H); 5.57 (s, 1H); 5.10 (s, 2 H); 4.60 (dd, J = 8.28; 5.57 Hz, 1 H); 4.24 (s, 1

H); 2.81 (m, 1 H); 2.58 (m, 1 H); 1.41 (s, 9H); 0.96 (t, J= 6Hz, 6 H).
LCMS negative (M-H) = 554.2

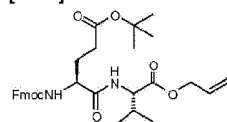
Example 5 : Synthesis of compound 20 Cbz-Asp(O-tBu)-Glu(O-tBu)-Val-OH

[0232]



a) Fmoc-Glu(O-tBu)-Val-OAllyl

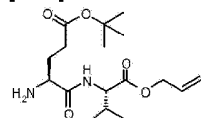
[0233]



[0234] Fmoc- Glu (O-tBu)-OH (0.549g, 1.292 mmol) was solvated in CH_2Cl_2 (5 mL) followed by addition of L- Val- allyl ester toluene-4- sulfonate (0.426 g, 1 eq), 4-methylmorpholine (0.145 mL, 1.02 eq), DMAP (11.53 mg, 0.13 eq) and finally EDC (0.252g, 1.02 eq). The EDC vial was washed with CH_2Cl_2 (0.5 mL*2) and added to the reaction mixture. After 1 hour 10 min of stirring, the reaction mixture was extracted with CH_2Cl_2 (30 mL)/ brine (5 mL). The organic layer was dried over MgSO_4 , filtered off and concentrated. The obtained residue was purified on silica using a gradient Hex/ EtOAc (0 to 40 %) to get 0.396 g of the desired compound.

b) Glu(O-tBu)-Val-OAllyl

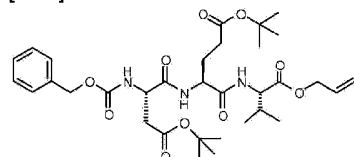
[0235]



[0236] Fmoc- Glu(O-tBu)-Val(OAllyl) (0.394 g, 0.698 mmol) was solvated in CH_2Cl_2 (4 mL) followed by addition of piperidine (0.550 mL, 7.98 eq). After 40 min, the mixture was evaporated to dryness and co-evaporated with CH_2Cl_2 (20 mL*2) followed with high vacuum for 10 min to remove the excess of piperidine. The sample was purified on silica with a gradient of MeOH / CH_2Cl_2 (0 to 7 %) to get 0.191 g of the desired compound.

c) Cbz-Asp(O-tBu)-Glu(O-tBu)-Val-OAllyl

[0237]

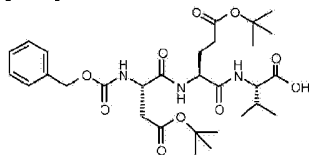


[0238] Glu-Val (OAllyl) (0.189 g, 0.55 mmol) was solvated in CH_2Cl_2 (2.5 ml) followed addition of Z-Asp (OtBu)-OH (0.187 g, 1.05 eq), DMAP (5.98 mg, 0.088 eq) and 4-methylmorpholine (0.065 ml, 1.07 eq), then EDC (0.109 g, 1.03 eq). The vial was rinsed with 0.5 ml of dichloromethane. The mixture was stirred for 2 hour at room temperature. Then it was extracted using CH_2Cl_2 / brine. The organic

layer was dried over MgSO_4 , filtered off and concentrated. The obtained residue was purified on silica using a gradient EtOAc/Hex (10 to 100 %) to get 0.300 g of the desired compound.

d) Cbz-Asp(O-*t*Bu)-Glu(O-*t*Bu)-Val-OH

[0239]

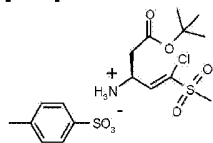


[0240] Cbz Asp (O-*t*Bu)- Glu (O*t*Bu)- Val- (OAllyl) (0.298 g, 0.46 mmol) was dissolved in THF (10 mL) and rotary evaporated to dryness, the sample was then redissolved in THF (10 mL) and vacuum aspirated (3* 1 min of aspiration), followed by replacement of the atmosphere with Argon. $\text{Pd}(\text{PPh}_3)_4$ (0.052 g, 0.1 eq) was added in one shot, the flask was then evacuated with Argon. Morpholine (0.14 mL, 3.49 eq) was added and the flask covered with a tin foil. The mixture was kept under stirring for 2.5 days under Argon. The sample was evaporated to dryness and the obtained residue was subjected to purification on C_{18} using a gradient MeOH/ solution of H_2O at pH= 3.5 (10 to 100 %) to get 0.150 g of the desired compound.

NMR ^1H (CD_3OD , 400 MHz) δ : 7.4-7.25 (m, 5H); 5.20-5.05 (m, 2H); 4.6-4.05 (m, 1H); 4.47-4.38 (m, 1H); 4.25-4.18 (m, 1H); 2.88-2.75 (m, 1H); 2.68-2.55 (m, 1H); 2.42-2.25 (m, 2H); 2.22-2.08 (m, 2H); 1.96-1.80 (m, 1H); 1.45 (2s, 18H); 0.93 (t, 6H).

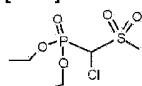
Example 6 : Synthesis of compound 23 (Asp (β -*tert*-butyl) Chlorovinyl-methyl vinyl sulfone tosyl salt)

[0241]



a) Diethyl chloro(methylsulfone) methylphosphonate

[0242]

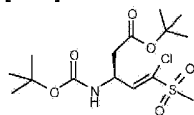


[0243] Diethyl (methylthio)methylphosphonate (1.45 g, 6.25 mmol) was solvated in acetic acid (5 mL, 14 eq) follow by addition of hydrogen peroxide (1.98 mL, 2.8 eq). The sample was then placed in a oil bath preheated to 70 °C, vigour evolution of gaz shortly followed. After 30 min the reaction was allowed to reach room temperature. Then NaHCO_3 was added in a small portion until the pH become neutral. The sample was then vacuum aspirated followed by extraction with Ether (30 mL). The organic layer was then washed with 20% of citric acid (5 mL) and then brine (2*5 mL). The organic layer was dried over MgSO_4 and purified on silica using a gradient EtOAc/Hexane (20 to 100 %) to get 0.552 g of the desired compound.

NMR ^1H (CDCl_3 , 400 MHz) δ : 4.21-4.13 (m, 4H); 2.88-2.75 (m, 1H); 2.68 (d, 2H, $J = 12.83$ Hz); 2.29 (s, 3H); 1.34 (t, 6H).

b) Boc-Asp (β -*tert*-butyl) achlorovinyl methylsulfone

[0244]

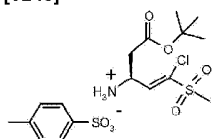


[0245] Diethyl chloro(methylsulfonyl)methylphosphonate (0.327 g, 1.24 mmol) was solvated in THF (5mL) and the solution was allowed to reach -78 °C. NaH 60% (0.0519 g, 1.04 eq), which had been washed with Ether anhydrous (3*0.9 mL), was then added in suspension in THF (1.5 mL). The vial containing NaH in suspension was washed with THF (0.4 mL*2) and it was added to the solution. The mixture was stirred for 25 min, then Boc Asp (O-*t*Bu)-H (0.326 g, 1eq) solvated in THF (3 mL) was added dropwise to the solution over 1 min. The vial was rinsed with THF (0.3 mL*2) and added to the reaction mixture. After 45 min of stirring, the solution was quenched with a solution of saturated ammonium chloride (5 mL) and extracted with EtOAc (30 mL). The organic layer was washed with brine, dried over MgSO₄, filtered and concentrated. The residue was purified on silica using a gradient of EtOAc/Hex (0 to 40 %) to get first the *cis* isomer and then 0.183 g of the *trans* isomer compound.

NMR ¹H (EDCl₃, 400 MHz) δ: 7.1 (d, 1 H, *J*= 8.06 Hz); 5.5 (m, 1 H); 4.8 (m, 1 H); 3.05 (s, 3H); 2.62 (m, 2H); 1.45 (2s, 18H).

c) Asp(β-*tert*-butyl)achlorovinyl methylsulfonyl tosyl salt

[0246]

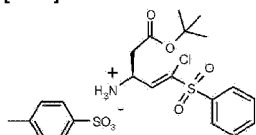


[0247] Boc-Asp (β-*tert*-butyl) Chlorovinyl-methyl vinyl sulfone (0.182 g, 0.4937 mmol) was solvated in CH₂Cl₂ (0.7 mL) followed by the addition of Et₂O (0.7 mL). *p*-Toluene sulfonic Acid monohydrate (0.101 g, 1.07 eq) was added in one shot. After 15 hours of stirring at room temperature, it was then diluted with Ether (8 mL) and filtered off. The white solid was then dried over vacuum, 0.12 g of the desired compound was obtained.

NMR ¹H (DMSO, 400 MHz) δ: 8.2 (bs, 3H, NH₃); 7.48 (d, 2H, *J*=8.04 Hz); 7.12 (d, 2H, *J*=7.89 Hz); 7.02 (d, 1H, *J*=9.4 Hz); 4.4-4.3 (m, 1H); 3.2 (s, 3H); 2.85 (dd, 1H, *J*= 16.39 Hz, *J*= 5.55 Hz); 2.75 (dd, 1H, *J*= 16.47 Hz, *J*= 7.63 Hz); 2.28 (s, 3H); 1.42 (s, 9H).

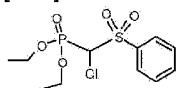
Example 7: Synthesis of compound 26 (Asp(β-*tert*-butyl)achlorovinyl phenylsulfone tosyl salt)

[0248]



a) Diethyl chloro(phenylthio)methylphosphonate

[0249]

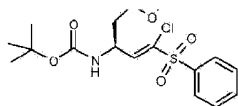


[0250] Diethyl chloro(phenylthio)methylphosphonate (0.919 g, 3.12 mmol) was solvated in acetic acid (1.8 mL), followed by addition of hydrogen peroxide (0.78 mL, 2.8 eq). The sample was then placed in a oil bath preheated to 70 °C. An additional portion of hydrogen peroxide (0.21 mL) was added after 5 min. After 30 min the reaction was allowed to reach room temperature. The sample was extracted with AcOEt /5% NaHCO₃ (30/5 mL). The organic layer was dried over MgSO₄, filtered and concentrated. The residue was purified on silica using a gradient EtOAc/Hexane (10 to 80 %) to get 0.627 g of the desired compound.

b) Asp(β-*tert*-butyl)achlorovinyl phenylsulfone

[0251]



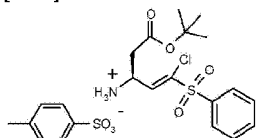


[0252] Diethyl chloro(phenylsulfone)methylphosphonate (0.458 g, 1.03 eq) was solvated in THF (3mL) and the solution was allowed to reach -78 °C. NaH 60% (0.016 g, 1.11 eq), which had been washed with Ether anhydrous (3*0.9 mL), was then added in suspension in THF (1.5 mL). The vial containing NaH in suspension was washed with THF (0.4 mL*2) and added to the solution. The mixture was stirred for 25 min, then BocAsp(OtBu)-H (0.372 g, 1.36 mmol) solvated in THF (0.5 mL) was added dropwise to the solution over 1 min. The vial was rinsed with THF (2* 0.5 ml) and added to the reaction mixture. After 45 min of stirring, the solution was quenched with a solution of ammonium chloride saturated (5 mL) and extracted with EtOAc (30 mL). The organic layer was washed with brine, dried over MgSO₄, filtered and concentrated. The residue was purified on silica using a gradient of Hex/ EtOAc (0 to 30 %), repurified using a gradient of MeOH/CH₂Cl₂(0 to 3 %) to elute first the cis product and then the trans product. Finally, a small amount of the desired compound was obtained.

NMR ¹H (CDCl₃, 400 MHz) δ: 8.13 (m, 2H); 7.67 (t, 1H, J=7.42 Hz); 7.57 (t, 2H); 6.55 (d, 1H, J=9.19 Hz); 5.75 (bs, 1H); 5.64 (bs, 1H); 2.86-2.77 (m, 2H); 1.47 (s, 9H); 1.45 (s, 9H).

c) Asp(β-tert-butyl)achlorovinyl phenylsulfone tosyl salt

[0253]

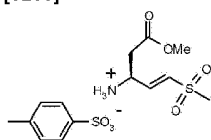


[0254] Asp(β-tert-butyl) achlorovinyl phenylsulfone (0.018 g, 0.04 mmol) was solvated in CH₂Cl₂ (0.1 mL) followed by the addition of Et₂O (0.1 mL). p-Toluene sulfonic Acid monohydrate (0.05 g, 0.1 eq) was added in one shot. After 15 hours of stirring at room temperature, it was then evaporated to dryness without heating to get 18 mg of the desired compound.

NMR ¹H (DMSO, 400 MHz) δ: 8.3 (s, 3H, NH₃); 8.08-8.02 (m, 2H); 7.8 (t, 1H, J= 7.46 Hz); 7.76 (t, 2H, J= 8.08 Hz); 7.48 (d, 2H, J= 8.06 Hz); 7.1 (d, 2H, J= 7.81 Hz); 6.73 (d, 1H, J= 10.09 Hz); 5.3 (m, 1H); 2.87 (dd, 1H, J= 16.67 Hz, J= 6.13 Hz); 2.78 (dd, 1H, J= 16.77 Hz, J= 7.01 Hz); 2.28 (s, 3H); 1.43 (s, 9H).

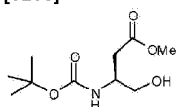
Example 8 : Synthesis of compound 30 (Asp(β-Methyl) methyl vinyl sulfone tosyl salt)

[0255]



a) Boc-Aspartimol(β-Methyl)

[0256]



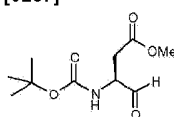
1. α-Boc-L-Asp (β-Methyl)-OH (2.5 g, 0.0101 mol) was solvated in ethyl acetate (12.5 ml) and chilled at 0°C. N-hydroxysuccinimide (1.163 g, 1eq) was added, followed with a dropwise addition of DCC (10.1 ml, 1 eq, 1M in CH₂Cl₂). The mixture was allowed to reach room temperature overnight (20 h), it was then diluted with ethyl acetate and filtered off on celite and washed with ethyl acetate (80 ml total volume). The organic layer was washed with 5% NaHCO₃ (2* 15 ml), brine (2 * 25 ml), dried over MgSO₄ and concentrated to dryness to get 3.58 g of the desired compound.

2. b-Boc-L-Asp (β -Methyl)-N-hydroxysuccinimide (1.81 g, 0.00526 mol) was dissolved in 24 mL of anhydrous THF under Argon. The mixture was chilled to 0 °C, NaBH_4 (0.5g, 2.51 eq) was added portion wise over a period of 25 min. The mixture was allowed to reach room temperature, and stirred for an extra 4 hours. A solution of ice water/ brine (1/1, 15 mL) was added dropwise at 0 °C followed by caution addition of citric acid (0.5 M, 40 mL). The biphasic mixture was stirred and the product was extracted with EtOAc (4*40 mL). The combined organic layer were washed with 5% NaHCO_3 (15 mL) and brine (15 mL), dried over MgSO_4 , filtered off and concentrated under vacuum. The crude material was then purified on silica with a gradient of CH_2Cl_2 / MeOH (0 to 5 %) to get 0.7 g of the desired compound.

NMR ^1H (CDCl_3 , 400 MHz) δ : 5.20 (bs, 1H); 4.02-3.95 (m, 1H); 3.88-3.64 (m, 2H); 3.70 (s, 3H); 2.63 (d, 2H, J = 5.86 Hz); 2.0 (bs, 1 H, OH); 1.44 (s, 9H).

b) Boc-Asp(β -Methyl)-H

[0257]

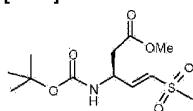


[0258] Oxalyl chloride 2M in CH_2Cl_2 (1.242 mL, 1.7 eq) dissolved in CH_2Cl_2 (2.4 mL) was cooled to -65 °C. A solution of DMSO (0.4mL, 3.9 eq) in CH_2Cl_2 (0.92 mL) was added dropwise over 20 min at -65 °C. Boc Asp (OCH_3)- CH_2OH in CH_2Cl_2 (2.6 mL) was added dropwise over a period of 20 min and the reaction was stirred for an extra 15 min at -65 °C. TEA (1.33 mL, 6.54 eq) in CH_2Cl_2 (1.42 mL) was added dropwise over 20 min. The reaction was left for an extra 55 min at -65 °C /-70 °C then quenched at this temperature with ether/ 0.5 N KHSO_4 (30/5 mL). The organic layer was washed 3 times with 0.5N KHSO_4 (3*5 mL) then brine, dried over MgSO_4 , filtered off and concentrate, to get 0.29 g of the desired compound.

NMR ^1H (CDCl_3 , 400 MHz) δ : 9.65 (s, 1H); 5.61 (d, 1H, J = 7.04 Hz); 4.38-4.35 (q, 1H, J = 4.3 Hz); 3.70 (s, 3H); 3.01 (dd, 1H, J = 17.41 Hz, J = 4.49 Hz); 2.83 (dd, 1H, J = 17.41 Hz, J = 4.69 Hz); 1.46 (s, 9H).

c) Boc-Asp(β -Methyl)methyl vinyl sulfone

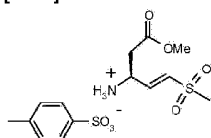
[0259]



[0260] NaH 95% (0.038g, 1.2 eq), which had been washed with Ether anhydrous (3*0.9 mL), was suspended in THF (1mL). The NaH solution was added dropwise at 0 °C to Diethyl (methylsulfone) methylphosphonate (0.280 g, 1.08eq) which was dissolved in a solution of THF (10 mL). The mixture was stirred for 20 min, then Boc Asp (OMethyl)-H (0.290 g, 1.255 mmol) solvated in THF (2.5 mL) was added dropwise to the solution over 1 min. After 15 min at 0 °C, the reaction was allowed to reach room temperature. After 1 hour of stirring, the solution was quenched with a solution of saturated ammonium chloride (10 mL) and extracted with EtOAc (30 mL). The organic layer was washed with brine (20 mL), dried over MgSO_4 , filtered and concentrate. The residue was purified on silica using a gradient of EtOAc /Hex (10 to 80 %) to elute first the cis isomer then the trans isomer (0.174 g).

d) Boc-Asp(β -Methyl)methyl vinyl sulfone tosyl salt

[0261]

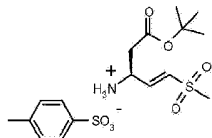


[0262] Boc-Asp(β -Methyl) methyl vinyl sulfone (0.172 g, 0.56 mmol) was solvated in CH_2Cl_2 (0.44 mL) followed by the addition of Et_2O (0.44 mL). *p*-Toluene sulfonic Acid monohydrate (0.19 g, 1.04 eq) was added in one shot. After 15 hours of stirring at room temperature, it was then diluted with Ether (2 mL) and filtered off. The white solid was then dried over vacuum, 0.140 g of the desired compound was obtained.

NMR ^1H (DMSO, 400 MHz) δ : 8.20 (bs, 3H, NH_3); 7.47-7.45 (m, 2H); 7.11 (dd, 2H, $J = 8.41$ Hz, $J = 0.58$ Hz); 7.03 (dd, 1H, $J = 15.45$ Hz, $J = 1.17$ Hz); 6.73 (dd, 1H, $J = 15.45$ Hz, $J = 6.26$ Hz); 4.34 (q, 1H, $J = 6.45$ Hz); 3.66 (s, 3H); 3.05 (s, 3H); 2.92-2.82 (m, 2H); 2.28 (s, 3H).

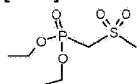
Example 9 :Synthesis of compound 33 (Asp(β -tert-butyl)methyl vinyl sulfone tosyl salt)

[0263]



a) Diethyl (methylsulfone)methylphosphonate

[0264]

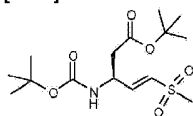


[0265] Diethyl (methylthio)methylphosphonate (4.0180 g, 20.3 mmol) was solvated in acetic acid (14 mL, 12 eq) followed by dropwise addition (10 min) of hydrogen peroxide (5.9 mL, 2.56 eq). The sample was then heated to 70 °C, vigorous evolution of gas shortly followed. After 25 min the reaction was allowed to reach room temperature. Then NaHCO_3 was added in a small portion until the pH became neutral. The sample was then vacuum aspirated followed by extraction with Ether. The organic layer was then washed with 20% of citric acid (5 mL) and then brine (2*10 mL). The organic layer was dried over MgSO_4 and purified on silica using a gradient EtOAc/Hexane (10 to 100 %) to get 3.237 g of the desired compound.

NMR ^1H (DMSO, 400 MHz) δ : 4.18 (d, 2H, $J = 16.62$ Hz); 4.10-4.03 (m, 4H); 3.10 (s, 3H); 1.23 (t, 6H, $J = 7.04$ Hz).

b) Boc-Asp(β -tert-butyl)methyl vinyl sulfone

[0266]

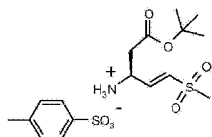


[0267] NaH 60% (0.087g, 1.11 eq), which had been washed with Ether anhydrous (3*0.9 mL), was suspended in THF (1mL). The NaH solution was added dropwise to Diethyl (methylsulfone)methylphosphonate (0.485 g, 1.08 eq) which was dissolved in a 0 °C solution of THF (20 mL). The mixture was stirred for 20 min, then Boc Asp (OtBu)-H (0.537 g, 1eq) solvated in THF (2 mL) was added dropwise to the solution over 1 min. After 10 min at 0 °C, the reaction was allowed to reach room temperature. After 1 hour of stirring, the solution was quenched with a solution of ammonium chloride saturated (45 mL) and extracted with EtOAc (100 mL). The organic layer was washed with brine (20 mL), dried over MgSO_4 , filtered and concentrate. The residue was purified on silica using a gradient of Hex/ EtOAc (10 to 80 %) to elute 0.05 g of the cis compound and 0.480 g of the trans compound.

NMR ^1H (CDCl_3 , 400 MHz) δ : 6.86 (dd, 1H, $J = 15.08$ Hz, $J = 4.60$ Hz); 6.51 (dd, 1H, $J = 15.14$ Hz, $J = 1.38$ Hz); 5.41 (bs, 1H); 4.7 (bs, 1H); 2.91 (s, 3H); 2.64-2.52 (qd, 2H); 1.43 (s, 18H).

c) Asp(β -tert-butyl)methyl vinyl sulfone tosyl salt

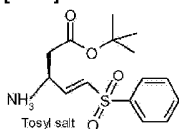
[0268]



[0269] Boc-Asp(β -tert-butyl) methyl vinyl sulfone (0.158 g, 0.4534 mmol) was solvated in CH_2Cl_2 (0.7 mL) followed by the addition of Et_2O (0.7 mL). p-Toluene sulfonic Acid monohydrate (0.0878 g, 1.02 eq) was added in one shot. The use of excess of PTSA hydrate cleaves both boc and tert-butyl groups unlike what was reported by Palmer. After 15 hours of stirring at room temperature, it was then diluted with Ether (5 mL) and filtered off. The white solid was then dried over vacuum, 0.121 g of the desired compound was obtained. NMR ^1H (DMSO, 400 MHz) δ : 8.18 (bs, 3H, NH_3); 7.47 (d, 2H, $J = 8.14$ Hz); 7.10 (d, 2H, $J = 7.85$ Hz); 7.02 (d, 1H, $J = 15.42$ Hz); 6.70 (dd, 1H, $J = 15.40$ Hz, $J = 6.61$ Hz); 4.27 (q, 1H, $J = 6.3$ Hz); 3.04 (s, 3H); 2.81-2.70 (qd, 2H); 2.28 (s, 3H); 1.42 (s, 9H).

Example 10 : Synthesis of compound 37 (Asp-vinyl phenyl sulfone salt form)

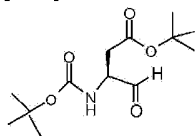
[0270]



[0271] The commercially available N-tBoc-L-Asp(β -tert-Butyl)-O-succinimide was reduced to the corresponding alcohol in the presence of sodium borohydride in THF, as described in the literature (Ramond J. Begeron et al., 1999)

a) Boc-Asp(β -tert-Butyl)-H

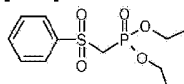
[0272]



[0273] The alcohol is then oxidized to the corresponding Boc-L-Asp(β -tert-Butyl)-H in the presence of oxalyl chloride, DMSO and TEA in dichloromethane at -70°C as described in the literature (William R. Ewing et al., 1999; Won Bum Jang. 2004 and Mancuso A et al., 1981)

b) Diethyl phenylsulfonylmethylphosphonate

[0274]



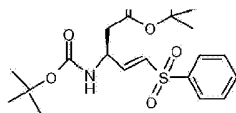
[0275] The precursor of phenyl vinyl sulfone was obtained in one step from benzenesulfonyl fluoride and triethyl phosphorane in the presence of lithium hexamethyldisilazide at -78°C to get diethyl phenylsulfonylmethylphosphonate as described in the literature (Won Bum Jang et al., 1998).

Boc-Asp(β -tert-Butyl)-vinyl phenyl sulfone (Gang Wang et al., 2003; Marion G. Gotz et al., 2004; Palmer, James T et al., 1995).

c) Boc-Asp-vinyl phenyl sulfone

[0276]



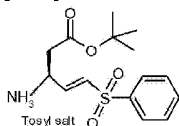


[0277] Sodium hydride (40 mg (60%), 1.09 eq) was added to a solution of diethyl phenylsulfonylmethylphosphonate (0.279g, 1.09 eq) in dry THF (5.6 ml) at 0 °C. The mixture was stirred for 20 minutes before adding, drop-wise, a solution of Boc-L-Asp(O-t-Bu)-H (0.24 g, 0.876 mmol) in 1.6 ml of THF. The mixture was stirred for 1.15 h at RT, then poured into a mix of ethyl acetate and ammonium chloride saturated solution (45/15 ml). The organic layer was dried over magnesium sulfate and the solvent was evaporated to dryness. The crude material was purified on silica gel (gradient: ethyl acetate/hexane) to make the desired compound with a high chemical yield.

NMR ^1H (CD_3OD , 400 MHz) δ 7.87 (d, J = 7.62 Hz, 2 H); 7.61 (t, J = 7.35 Hz, 1H); 7.53 (t, J = 7.68 Hz, 2H); 6.92 (dd, J = 15.08 and 4.28 Hz, 1H); 6.46 (d, J = 15.12 Hz, 1H); 5.34 (m, 1 H); 4.68 (m, 1 H); 2.63-2.52 (m, 2 H); 1.40 (s, 18H).

d) Asp Vinyl phenyl sulfone tosyl salt

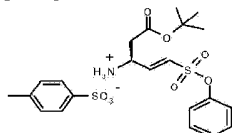
[0278]



[0279] Boc-L-Asp(β -tert-Butyl)-Vinyl phenyl sulfone (0.1 g, 0.243 mmol) was dissolved in a mix of dichloromethane and ether (0.7 / 0.7 ml), then PTSA hydrate (1eq) was added. The use of excess of PTSA hydrate cleaves both boc and tert-butyl groups unlike what was reported by Palmer. The mixture was stirred at room temperature overnight. Then, it was diluted with ether (8 ml). The white precipitate was filtered off and dried to yield to the desired compound as a white powder.

Example 11: Synthesis of compound 40 (Asp(β -tert-butyl) phenoxy vinylsulfone tosyl salt)

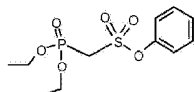
[0280]



[0281] To a solution of phenol (1.62 g, 17.2 mmol) in Et_2O (50 mL) at -10 °C was added TEA (3.6 mL, 1.5 eq). After 15 min a solution of methanesulfonylchloride (1.6 mL, 1.2 eq) in Et_2O (4 mL) was added dropwise over 50 min. Then the solution was allowed to warm at room temperature and an additional portion of Et_2O (5mL) was added. The reaction was quenched by addition of 1 N HCl (12 mL, 4 C), the organic layer was then washed with saturated NaHCO_3 , brine and dried over MgSO_4 , filtered off and concentrate to give an oil which was recrystallised for $\text{CH}_2\text{Cl}_2/\text{Hex}$ (1/1). The resulting solid was filtered and the residual solvent removed.

a) Diethyl (phenoxysulfone)methylphosphonate

[0282]



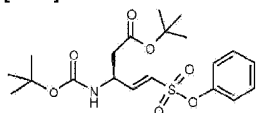
[0283] To a solution of methanesulfonyl phenoxy (1.938 g, 11.25 mmol) in THF (8 mL) at -78 °C, was added dropwise a solution of potassium bis(trimethylsilyl)amide (2.36 g, 1 eq) in 11 mL of THF over a period of 40 min. Then the reaction was stirred for an extra 5 min. Diethyl chlorophosphonate (0.95 mL, 0.59 eq) was added dropwise over 7 min. After 1 hour, the reaction was quenched by dropwise addition of a solution of Acetic Acid (0.645 mL, 0.59 eq) over 5 min. The solution was allowed to warm to room temperature

and the solvent was removed in vacuum. The product was extract with CH_2Cl_2 (30 mL) and H_2O (10 mL), dried over MgSO_4 , filtered off and concentrate. The residue was purified on silica using a gradient EtOAc/Hexane (12 to 100 %) to get 1.046 g of the desired compound.

NMR ^1H (CDCl_3 , 400 MHz) δ : 7.44-7.41 (m, 2H); 7.36-7.32 (m, 3H); 4.31-4.25 (m, 4H); 3.81 (d, 2H, J = 17.14 Hz); 1.38 (t, 6H, J = 7.11 Hz).

b) Asp(β -tert-butyl) phenoxy vinyl sulfone

[0284]

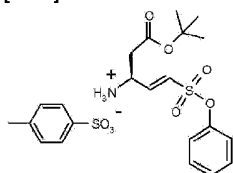


[0285] Diethyl (phenoxy sulfone)methylphosphonate (0.36 g, 1.07 mmol) was solvated in THF (10mL) and the solution was allowed to reach -0 °C. NaH 60% (0.0539 g, 1.22 eq), which had been washed with Ether anhydrous (3*0.9 mL), was then added in suspension in THF (1 mL). The vial containing NaH in suspension was washed with THF (1 mL) and it was added to the solution. The mixture was stirred for 20 min, then Boc Asp (O-*t*Bu)-H (0.30 g, 1eq) solvated in THF (2 mL) was added dropwise to the solution over 1 min. The vial was rinsed with THF (1 mL) and added to the reaction mixture. After 2 hours of stirring at room temperature, the solution was quenched with a solution of ammonium chloride saturated (5 mL) and extracted with EtOAc (30 mL). The organic layer was washed with brine, dried over MgSO_4 , filtered off and concentrated. The residue was purified on silica using a gradient of EtOAc/Hexane (0 to 30%, 30 to 80 %) to get 0.359 g of the desired compound.

NMR ^1H (CDCl_3 , 400 MHz) δ : 7.38 (t, 2H, J = 7.96 Hz); 7.29 (t, 1H, J = 7.46 Hz); 7.23-7.21 (m, 2H); 6.79 (dd, 1H, J = 15.14 Hz, J = 4.67 Hz); 6.49 (dd, 1H, J = 15.18 Hz, J = 1.31 Hz); 5.37 (bs, 1H); 4.65 (bs, 1H); 2.60 (dd, 1H, J = 16.04 Hz, J = 5.46 Hz); 2.50 (dd, 1H, J = 16.04 Hz, J = 5.63 Hz); 1.46 (s, 9H); 1.43 (s, 9H).

c) Asp(β -tert-butyl) phenoxy vinyl sulfone tosyl salt

[0286]

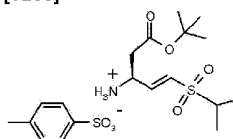


[0287] Asp(β -tert-butyl) phenoxy vinyl sulfone (0.187 g, 0.444 mmol) was solvated in CH_2Cl_2 (0.7mL) followed by the addition of Et_2O (0.7 mL). *p*-Toluene sulfonic Acid monohydrate (0.084 g, 1.01 eq) was added in one shot. After 15 hours of stirring at room temperature, it was then diluted with Ether (8 mL) and filtered off. The while solid was then dried over vacuum to get the desired compound.

NMR ^1H (DMSO , 400 MHz) δ : 8.25 (bs, 3H, NH_3); 7.49-7.46 (m, 4H); 7.39 (t, 1H, J = 7.4 Hz); 7.31-7.29 (m, 2H); 7.22-7.17 (m, 1H); 7.10 (d, 2H, J = 7.88 Hz); 6.82 (dd, 1H, J = 15.41 Hz, J = 6.15 Hz); 4.35 (m, 1 H); 2.82-2.73 (m, 2H); 2.28 (s, 3H); 1.41 (s, 9H).

Example 12: Synthesis of compound 43 (Asp(β -tert-butyl) isopropyl vinyl sulfone tosyl salt)

[0288]



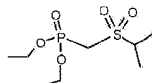
[0289] Chloromethyl isopropylsulfide (12.54 g, 100.6 mmol) was heated to 110 °C follow by dropwise addition of triethylphosphonyl (21 mL, 1.10 eq). After stirring for 8 hours, the reaction was allowed to reach to room temperature. The sample was purified by

distillation (110 °C / 6 mmbar) to get 4.7 g of Diethyl (isopropylthio)methylphosphonate.

NMR ^1H (CDCl_3 , 400 MHz) δ : 4.21-4.13 (m, 4H); 3.19-3.14 (m, 1H); 2.76 (d, 2H, J = 14.35 Hz); 1.34-1.27 (m, 12H).

Example 13: Synthesis of compound 41 (Diethyl (isopropylsulfone) methylphosphonate)

[0290]

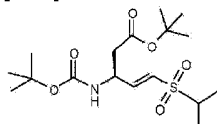


[0291] Diethyl (isopropylthio)methylphosphonate (4.746 g, 0.02 mol) was solvated in acetic acid (14.5 mL, 12.06 eq) followed by dropwise addition over 5min of hydrogen peroxide (6 mL, 2.52 eq). The sample was then heated to 70 °C, vigor evolution of gas shortly followed. After 30 min the reaction was allowed to reach room temperature. Then NaHCO_3 was added in a small portion until the pH become neutral. The sample was then vacuum aspirated followed by extraction with Ether (30 mL). The organic layer was then washed with 20% of citric acid (5 mL) and then brine (2*10 mL). The organic layer was combined and extract with CH_2Cl_2 / water, dried over MgSO_4 and purified on silica using a gradient EtOAc/Hexane (20 to 100 %) to get 4.689 g of the desired compound.

NMR ^1H (CDCl_3 , 400 MHz) δ : 4.23-4.50 (m, 4H); 3.67-3.63 (m, 1H); 3.53 (d, 2H, J = 16.82 Hz); 1.37-1.30 (m, 12H).

Example 14 : Synthesis of compound 42 (BocAsp(β -tert-butyl) isopropyl vinyl sulfone)

[0292]

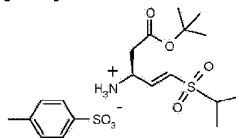


[0293] Diethyl (isopropylsulfone)methylphosphonate (0.036 g, 0.1427 mmol) was solvated in THF (0.2mL) and the solution was allowed to reach 0 °C. NaH 60% (5.38 mg, 1.05 eq), which had been washed with Ether anhydrous (3*0.9 mL), was then added in suspension in THF (1.2 mL). The mixture was stirred for 25 min, then Boc Asp (O-*t*Bu)-H (0.0351 g, 1eq) solvated in THF (0.6 mL) was added dropwise to the solution over 1 min. After 45 min of stirring, the solution was quenched with a solution of ammonium chloride saturated (2 mL) and extracted with EtOAc (10 mL). The organic layer was washed with brine, dried over MgSO_4 , filtered and concentrated. The residue was purified on silica using a gradient of EtOAc/Hexane (5 to to 60 %) to get 29.3 mg of the desired compound.

NMR ^1H (CDCl_3 , 400 MHz) δ : 6.82 (dd, 1H, J = 15.24 Hz, J = 4.78 Hz); 6.38 (dd, 1H, J = 15.17 Hz, J = 1.41 Hz); 5.43 (m, 1H); 4.67 (s, 1H); 3.07-3.0 (m, 1H); 2.64 (dd, 1H, J = 16.07 Hz, J = 5.5 Hz); 2.57 (dd, 1H, J = 16.12 Hz, J = 5.52 Hz); 1.48-1.32 (m, 24H).

Example 15 : Synthesis of compound 43 (Asp(β -tert-butyl)isopropyl vinyl sulfone tosyl salt)

[0294]

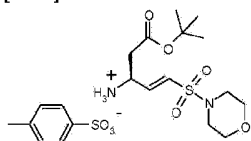


[0295] Boc-Asp(β -tert-butyl) isopropyl vinyl sulfone (0.028 g, 0.0741 mmol) was solvated in CH_2Cl_2 (0.2mL) followed by the addition of Et_2O (0.2 mL). *p*-Toluene sulfonic Acid monohydrate (0.0154 g, 1.09 eq) was added in one shot. After 15 hours of stirring at room temperature, it was then diluted with Ether (8 mL) and filtered off. The while solid was then dried over vacuum, 11 mg of the desired compound was obtained.

NMR ^1H (DMSO , 400 MHz) δ : 8.21 (bs, 3H, NH_3); 7.46 (d, 2H, J = 7.81 Hz); 7.10 (d, 2H, J = 7.85 Hz); 6.92-6.85 (m, 1H); 6.71 (dd, 1H, J = 15.42 Hz, J = 6.34 Hz); 4.30 (m, 1H); 3.23-3.13 (m, 1 H); 2.85-2.72 (m, 2H); 2.28 (s, 3H); 1.42 (s, 9H); 1.21 (m, 6H).

Example 16 : Synthesis of compound 46 (Asp(β -tert-butyl) morpholine vinyl sulfone tosyl salt)

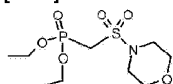
[0296]



[0297] To a solution of morpholine (1.4 mL, 16.07 mmol) in CH_2Cl_2 (30 mL) at -10°C was added TEA (3.4 mL, 1.5 eq). After 15 min a solution of methanesulfonylchloride (1.5 mL, 1.2 eq) in CH_2Cl_2 (4 mL) was added dropwise over 40 min. Then the solution was allowed to warm at room temperature and an additional portion of Et_2O (5mL) was added. The reaction was quenched by addition of 1 N HCl (12 mL, 4 C), the organic layer was then washed with saturated NaHCO_3 , brine and dried over MgSO_4 , filtered off and concentrate to give an oil which was purified on silica using a gradient of MeOH/ CH_2Cl_2 (0 to 5 %) to get 0.831 g of Diethyl (morpholinethio)methylphosphonate.

Example 16: Synthesis of compound 44 (Diethyl (morpholinesulfone)methylphosphonate)

[0298]

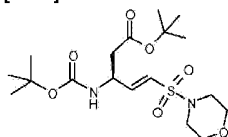


[0299] To a solution of methane sulfonyl morpholine (0.719 g, 4.356 mmol) in THF (4 mL) at -78°C , was added dropwise a solution of potassium bis(trimethylsilyl)amide (0.748 g, 0.86 eq) in 11 mL of THF over a period of 40 min. Then the reaction was stirred for an extra 5 min. Diethyl chloromethylphosphonate (0.37 mL, 0.57 eq) was added dropwise over 7 min. After 1 hour, the reaction was quenched by dropwise addition of a solution of Acetic Acid (0.226 mL, 0.77 eq) over 5 min. The solution was allowed to warm to room temperature and the solvent was removed in vacuum. The product was extract with CH_2Cl_2 (30 mL) and H_2O (10 mL), dried over MgSO_4 , filtered off and concentrate. The residue was purified on silica using a gradient MeOH/ CH_2Cl_2 (0 to 4 %) to get 0.247g of the desired compound.

NMR ^1H (CDCl_3 , 400 MHz) δ : 4.26-4.20 (m, 4H); 3.76 (t, 4H, $J = 4.6$ Hz); 3.51 (d, 2H, $J = 17.33$ Hz); 3.34 (t, 4H, $J = 4.74$ Hz); 1.37 (t, 6H, $J = 7.11$ Hz).

Example 17 : Synthesis of compound 45 (Boc-Asp(β -tert-butyl)morpholine vinyl sulfone)

[0300]

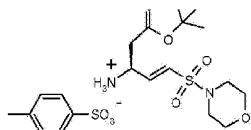


[0301] Diethyl (morpholinesulfone)methylphosphonate (0.103 g, 1.07 mmol) was solvated in THF (1 mL) and the solution was allowed to reach -10°C . NaH 60% (0.013 g, 1.04 eq), which had been washed with Ether anhydrous (3*0.9 mL), was then added in suspension in THF (1 mL). The vial containing NaH in suspension was washed with THF (0.5 mL) and it was added to the solution. The mixture was stirred for 20 min, then Boc Asp (OtBu)-H (0.208 g, 1eq) solvated in THF (1.5 mL) was added dropwise to the solution over 1 min, the vial was washed with DMF (0.5 ml) and added to the solution. After 3 hours of stirring at room temperature, the solution was quenched with a solution of ammonium chloride saturated (5 mL) and extracted with EtOAc (30 mL). The organic layer was washed with brine, dried over MgSO_4 , filtered off and concentrated. The residue was purified on silica using a gradient of first Hex/ EtOAc (5 to 80 %) then CH_2Cl_2 / MeOH (0 to 10 %) to get 0.075 g of the desired compound.

Example 18 : Synthesis of compound 46 (Asp(β -tert-butyl)morpholine vinyl sulfone tosyl salt)

[0302]

o 1 2

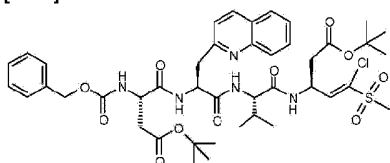


[0303] Boc-Asp(β -tert-butyl) morpholine vinyl sulfone (0.075 g, 0.178 mmol) was solvated in CH_2Cl_2 (0.2 mL) followed by the addition of Et_2O (0.2 mL). *p*-Toluene sulfonic Acid monohydrate (0.034 g, 1.0 eq) was added in one shot. After 15 hours of stirring at room temperature, it was then diluted with Ether (0.5 mL) and filtered off. The white solid was then dried over vacuum, 0.045 g of the desired compound was obtained.

NMR ^1H (DMSO, 400 MHz) δ : 8.20 (bs, 3H, NH_3); 7.47 (d, 2H, $J = 7.9$ Hz); 7.10 (d, 2H, $J = 7.89$ Hz); 6.88 (d, 1H, $J = 15.33$ Hz); 6.60 (dd, 1H, $J = 15.00$ Hz, $J = 6.36$ Hz); 4.28 (s, 1H); 3.65 (t, 4H, $J = 4.43$ Hz); 3.02 (m, 4H); 2.84-2.76 (m, 2H); 2.28 (s, 3H); 1.42 (s, 9H).

Example 18 : Synthesis of compound 47 (Z-Asp(β -tert-Butyl)-Ala(2'-quinolyl)-Val-Asp(β -tert-Butyl)-achlorovinyl methylsulfone)

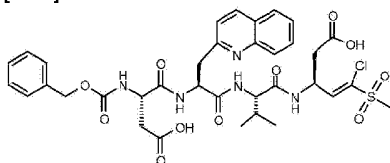
[0304]



[0305] The Z-Asp(β -tert-Butyl)-Ala(2'-quinolyl)-Val-OH (18 mg, 0.029 mmol) is dissolved in a mix of dichloromethane and DMF (0.39 ml / 0.13 ml). The mixture was allowed to reach -15/-20 °C before adding N-methyl morpholine (8 μl) followed 3 min latter with isobutyl chloroformate (7 μl). The mixture was stirred at -15 °C for 10 minutes. Next, Asp (β -tert-butyl) achloromethyl-methylsulfone tosyl salt (13 mg, 1eq) was added in one shot, followed by N-methyl morpholine (8 μl). The mixture was stirred 35 minutes at -15/-20 °C and then diluted with 5 ml of dichloromethane and quenched with 1.5 ml of a saturated solution of sodium bicarbonate. The organic layer was dried over magnesium sulphate, the solvent was evaporated to dryness. The obtained residue was purified on silica gel (gradient: ethyl acetate / hexane: 5 to 100%) to afford 12 mg of the desired compound.

Example 19: Synthesis of compound 48 (Z-Asp-Ala(2'-quinolyl)-Val-Asp-achlorovinyl methylsulfone)

[0306]



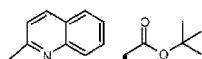
[0307] Z-Asp(β -tert-Butyl)-Ala(2'-quinolyl)-Val-Asp(β -tert-Butyl)-achlorovinyl methylsulfone (11.7 mg) was dissolved in dichloromethane (0.24 ml), followed by addition of trifluoroacetic acid (0.35 ml). The mixture was stirred at room temperature overnight (15 h). It was then quenched with diethyl ether (5 ml), the solvent was removed under vacuum. The obtained residue was diluted again with ether (5 ml) and the process was repeated twice. The precipitate was washed with 2* 1 ml of ether, dried to give 9 mg of the desired compound.

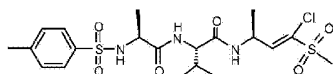
NMR ^1H (CD_3OD , 400 MHz) δ : 8.6-7.25 (m, 11H); 7.07 (d, 1H, $J = 8.61$ Hz); 5.03-4.06 (m, 4H); 4.45-4.42 (m, 1H); 4.09-4.05 (m, 1H); 3.65-3.49 (m, 2H); 3.07 (s, 3H); 2.84-2.7 (m, 2H); 2.71-2.66 (m, 2H); 2.00 (m, 1H); 0.85 (t, 6H, $J = 8.2$ Hz).

LCMS (M-H $^+$) = 772.4.

Example 20 : Synthesis of compound 49 (Ts-Ala(2'quinolyl)-Val-Asp(β -tert-Butyl)-achlorovinyl methylsulfone)

[0308]

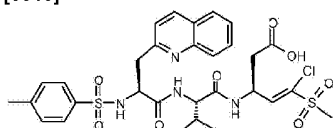




[0309] The Ts-Ala (2'-quinolyl)-OH (5.6 mg, 0.0123 mmol) was dissolved in a mix of CH_2Cl_2 / DMF (0.15/ 0.13 mL). The mixture was allowed to reach - 15 °C / - 20 °C (ice MeOH bath) before adding NMM (0.004 mL) followed 3 min later with isobutylchloroformate (0.004 mL). The mixture was stirred for 10 min at -15 °C before adding Asp (β-tert-butyl) Chlorovinyl-methyl vinyl sulfone tosyl salt (6.3 mg, 1 eq) in one shot, followed with 4-methylmorpholine (0.004 mL). The mixture was stirred at - 15 °C / -20 °C for 30 min, it was then diluted with CH_2Cl_2 (5 mL), then water (2 mL) and the mixture was allowed to reach room temperature, extracted. The organic layer was dried over MgSO_4 , concentrated and purified by silica using Hex/ EtOAc (15 to 100%) to get 6 mg of the desired compound.

Example 21 : Synthesis of compound 50 (Ts-Ala(2'-quinolyl)-Val-Asp- αchlorovinyl methylsulfone)

[0310]

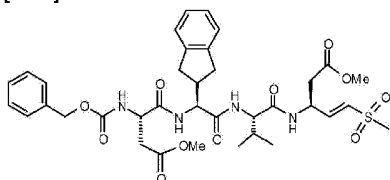


[0311] Ts-Ala(2'-quinolyl)-Val-Asp(β-tert-Butyl)- αchlorovinyl methylsulfone (6 mg) was dissolved in dichloromethane (0.15 ml), followed by addition of trifluoroacetic acid (0.2 ml). The mixture was stirred at room temperature overnight (15 h). It was then quenched with diethyl ether (5 ml) and the solvent was removed under vacuum. The obtained residue was diluted again with ether and the process was repeated twice. The precipitate was washed with 2* 1 ml of ether, dried to give 4 mg of the desired compound.

NMR ^1H (CD_3OD , 400 MHz) δ : 8.37 (m, 1 H); 8.01-7.99 (m, 1 H); 7.96-7.94 (m, 1 H); 7.90-7.94 (m, 1H); 7.90-7.88 (m, 1H); 7.71-7.63 (m, 1H); 7.46 (m, 1H); 7.37 (d, 2H, J = 8.22 Hz); 7.08 (d, 1H, J = 8.46 Hz); 6.88 (d, 2H, J = 8.0 Hz); 5.04-5.01 (m, 1 H); 4.37-4.34 (m, 1 H); 4.08-4.07 (m, 1 H); 3.44-3.4 (m, 1 H); 3.2 (m, 1 H); 3.08 (s, 3H); 2.83-2.78 (m, 1 H); 2.73-2.684 (m, 1H); 2.17 (s, 3H); 2.07-2.03 (m, 1H); 0.91-0.89 (m, 6H).

Example 22: Synthesis of compound 51 (Z-Asp(β-methyl)-Indanylglycine-Val-Asp(β-methyl)methyl vinyl sulfone)

[0312]



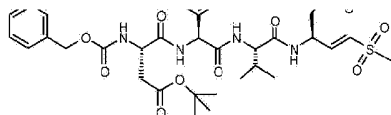
[0313] The Z-Asp(β-methyl)-Indanylglycine-Val-OH (16.7 mg, 0.0301 mmol) is dissolved in a mix of THF and DMF (0.5 ml / 0.1 ml). The mixture was allowed to reach -15/-20 °C before adding N-methyl morpholine (9 μl) followed 3 min latter with isobutyl chloroformate (8 μl). The mixture was stirred at -15 °C for 10 minutes. Next, Asp (β-methyl) methyl vinyl sulfone tosyl salt (12 mg, 1 eq) was added in one shot, the vial was washed with THF (0.1 ml) and added to the solution, followed by the addition of N-methyl morpholine (9 μl). The mixture was stirred 35 minutes at -15 /-20 °C and then diluted with 7 ml of dichloromethane and quenched with 1.5 ml of a saturated solution of sodium bicarbonate. After that, the organic layer was dried over magnesium sulphate. The solvent was evaporated to dryness. The obtained residue was purified on silica gel (ethyl acetate / hexane: 40% then dichloromethane/methanol: 5 to 15%) to afford 14 mg of Z-Asp(β-methyl)-Indanylglycine-Val- Asp(β-methyl)-methyl vinyl sulfone.

NMR ^1H (DMSO, 400 MHz) δ : 8.31 (d, NH, J = 7.82 Hz); 8.11 (d, NH, J = 8.8 Hz); 7.93 (d, NH, J = 8.21 Hz); 7.69 (d, NH, J = 8.02 Hz); 7.37-7.08 (m, 9H); 6.74 (dd, 1H, J = 15.45 Hz, J = 4.30 Hz); 6.67 (d, 1H, J = 15.59 Hz); 5.05 (s, 2H); 4.99 (m, 1H); 4.44 (m, 2H); 4.11 (m, 1 H); 3.57 (s, 6H); 2.98 (s, 3H); 2.95-2.5 (m, 9H); 2.00-1.94 (m, 1H); 0.84 (t, 6H, J = 6.06 Hz).

Example 23 : Synthesis of compound 52 (Z-Asp(β-tert-Butyl)-Phg-Val-Asp(β-tert-Butyl)methyl vinyl sulfone)

[0314]

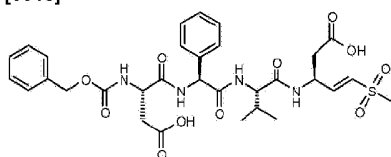




[0315] The Z-Asp(β-tert-Butyl)-Phg-Val-OH (17 mg, 0.0306 mmol) is dissolved in a mix of dichloromethane and DMF (0.39 ml / 0.13 ml). The mixture was allowed to reach -15/-20 °C before adding N-methyl morpholine (8 μl) followed 3 min latter with isobutyl chloroformate (8 μl). The mixture was stirred at -15 °C for 10 minutes. Next, Asp (β-term-butyl) methyl vinyl sulfone tosyl salt (12 mg, 1 eq) was added in one shot, followed by N-methyl morpholine (8 μl). The mixture was stirred 35 minutes at -15 /-20 °C and then diluted with 5 ml of dichloromethane and quenched with 1.5 ml of a saturated solution of sodium bicarbonate. The organic layer was dried over magnesium sulphate. The solvent was evaporated to dryness. The obtained residue was purified on silica gel (gradient: ethyl acetate / hexane: 5 to 100%) to afford 8 mg of Z-Asp(β-tert-Butyl)-Phg-Val- Asp(β-tert-butyl)-methyl vinyl sulfone.

Example 24 : Synthesis of compound 53 (Z-Asp-Phg-Val- Asp-methyl vinyl sulfone)

[0316]



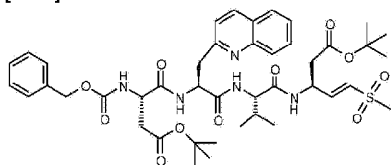
[0317] Z-Asp(β-tert-Butyl)-Phg-Val-Asp(β-tert-butyl)-methyl vinyl sulfone (6.2 mg) was dissolved in dichloromethane (0.16 ml), followed by addition of trifluoroacetic acid (0.22 ml). The mixture was stirred at room temperature overnight (15 h). It was then quenched with diethyl ether (5 ml), then the solvent was removed under vacuum. The obtained residue was diluted again with ether (5 ml) and the process was repeated twice. The crude material was diluted once more with ether, the filtrate was removed and the precipitate was washed with 2* 1 ml of ether, dried to give 5 mg of the desired compound.

NMR ¹H (CD₃OD, 400 MHz) δ: 7.46-7.29 (m, 10H); 6.83 (dd, 1H, J= 15.22 Hz, J= 4.60 Hz); 6.68 (dd, 1H, J= 15.229 Hz, J= 1.36 Hz); 5.38 (s, 1H); 5.10 (q, 2H, J= 12.73 Hz); 4.94-4.90 (m, 1H); 4.56 (t, 1 H, J= 6.75 Hz); 4.10 (d, 1 H, J= 7.07 Hz); 2.95 (s, 3H); 2.98-2.88 (m, 1H); 2.76-2.60 (m, 3H); 2.21-2.01 (m, 1H); 0.98 (t, 6H, J= 6.3 Hz).

LCMS (M-H⁺)= 673.7

Example 26 : Synthesis of compound 54 (Z-Asp(β-tert-Butyl)-Al(2'-quinolyl)-Val-Asp(β-tert- Butyl)methyl vinyl sulfone)

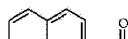
[0318]

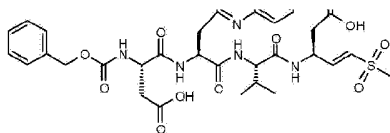


[0319] The Z-Asp(β-tert-Butyl)- Al(2'-quinolyl)-Val-OH (19 mg, 0.0306 mmol) is dissolved in a mix of dichloromethane and DMF (0.4 ml / 0.14 ml). The mixture was allowed to reach - 15/-20 °C before adding N-methyl morpholine (9 μl) followed 3 min latter with isobutyl chloroformate (9 μl). The mixture was stirred at -15 °C for 10 minutes. Next, Asp (β-tert-butyl) methyl vinyl sulfone tosyl salt (13.1 mg, 1 eq) was added in one shot, followed by N-methyl morpholine (9 μl). The mixture was stirred 35 minutes at -15 /-20 °C and then diluted with 5 ml of dichloromethane and quenched with 1.5 ml of a saturated solution of sodium bicarbonate. After that, the organic layer was washed with sodium bicarbonate and dried over magnesium sulphate. The solvent was evaporated to dryness. The obtained residue was purified on silica gel (gradient: ethyl acetate / hexane: 5 to 100%) to afford 13.1 mg of Z-Asp(β-tert-Butyl)- Al(2'-quinolyl)-Val- Asp(β-tert-butyl)-methyl vinyl sulfone.

Example 27 : Synthesis of compound 55 (Z-Asp-Ala(2'-quinolyl)-Val- Aspmethyl vinyl sulfone)

[0320]





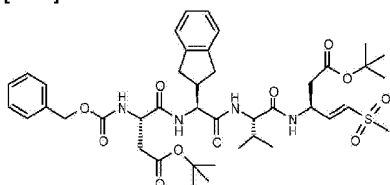
[0321] Z-Asp(β-tert-Butyl)-Phg-Val-Asp(β-tert-butyl)-methyl vinyl sulfone (13.1 mg) was dissolved in dichloromethane (0.32 ml), followed by addition of trifluoroacetic acid (0.44 ml). The mixture was stirred at room temperature overnight (15 h). It was then quenched with diethyl ether (5 ml), then the solvent was removed under vacuum. The obtained residue was diluted again with ether and the process was repeated twice. The precipitate was washed with 2* 1 ml of ether, dried to give 12 mg of the desired compound.

NMR ¹H (CD₃OD, 400 MHz) δ: 8.68 (m, 1H); 8.17 (d, 1H, J= 8.9 Hz); 8.1 (d, 1H, J= 8.2 Hz); 7.93 (t, 1H, J= 7.85 Hz); 7.78 (m, 2H); 7.34-7.29 (m, 5H); 6.88 (dd, 1H, J= 15.20 and 4.75 Hz); 6.7 (d, 1H, J= 15.20 Hz); 5.03-4.83 (m, 4H); 4.44 (t, 1H, J= 6.45 Hz); 4.1 (d, 1H, J= 7.15 Hz); 3.70 (m, 1H); 3.52 (m, 1H); 2.98 (s, 3H); 2.90-2.68 (m, 4H); 0.92 (d, 3H, J= 6.9Hz), 0.89 (d, 3H, J= 6.7Hz).

LCMS (M-H⁺)= 738.3

Example 28 :Synthesis of compound 56 (Z-Asp(β-tert-Butyl)-Indanylglycine-Val-Asp(β-tert-Butyl)methyl vinyl sulfone)

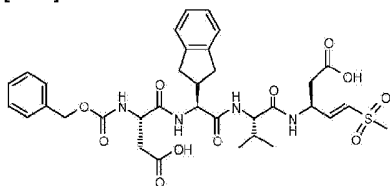
[0322]



[0323] The Z-Asp(β-tert-Butyl)-Indanylglycine -Val-OH (21 mg, 0.0353 mmol) is dissolved in a mix of dichloromethane and DMF (0.44 ml / 0.15 ml). The mixture was allowed to reach - 15/-20 °C before adding N-methyl morpholine (10 μl) followed by isobutyl chloroformate (9 μl). The mixture was stirred at -15 °C for 10 minutes. Next, Asp (β-tert-butyl) methyl vinyl sulfone tosyl salt (15 mg, 1 eq) was added in one shot, followed by N-methyl morpholine (10 μl). The mixture was stirred 35 minutes at -15 /-20 °C and then diluted with 5 ml of dichloromethane and quenched with 1.5 ml of a saturated solution of sodium bicarbonate. After that, the organic layer was washed with sodium bicarbonate and dried over magnesium sulphate. The solvent was evaporated to dryness. The obtained residue was purified on silica gel (ethyl acetate / hexane: 40% then dichloromethane/methanol: 5 to 15%) to afford 10 mg of Z-Asp(β-tert-Butyl)- indanylglycine-Val-Asp(β-tert-butyl)-methyl vinyl sulfone.

Example 30 : Synthesis of compound 57 (Z-Asp-Indanylglycine-Val- Aspmethyl vinyl sulfone)

[0324]

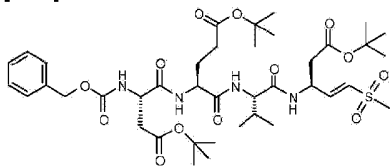


[0325] Z-Asp(β-tert-Butyl)-Indanylglycine-Val-Asp(β-tert-butyl)-methyl vinyl sulfone (11 mg) was dissolved in dichloromethane (0.26 ml), followed by addition of trifluoroacetic acid (0.36 ml). The mixture was stirred at room temperature overnight (15 h). It was then quenched with diethyl ether (5 ml) and the solvent was removed under vacuum. The obtained residue was diluted again with ether and the process was repeated twice. The precipitate was washed with 2* 1 ml of ether, dried to give 9 mg of the desired compound.

NMR ¹H (DMSO, 400 MHz) δ: 12.35 (bs, 2H, 2*CO₂H); 8.21 (d, 1H, J= 7.78 Hz); 8.06 (m, 1H); 7.89 (d, 1H, J= 7.34 Hz); 7.61 (d, 1H, J= 7.10 Hz); 7.34-7.31 (m, 5H); 7.17-6.95 (m, 4H); 6.75 (dd, 1H, J= 15.33 Hz, J= 4.8 Hz); 6.64 (d, 1H, J= 15.58 Hz); 4.99 (s, 2H); 4.81 (m, 1H); 4.46-4.40 (m, 2H); 4.12 (t, 1H, J= 6.28 Hz); 2.97 (s, 3H); 2.82-2.44 (m, 9H); 0.85 (t, 6H, J= 5.95 Hz).

Example 32 : Synthesis of compound 58 (Z-Asp(β-tert-Butyl)-Glu(β-tert-Butyl)-Val-Asp(β-tert-Butyl)methyl vinyl sulfone)

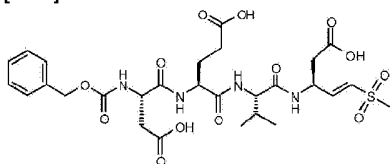
[0326]



[0327] The Z-Asp(β -tert-Butyl)-Glu(β -tert-Butyl)-Val-OH (18 mg, 0.029 mmol) is dissolved in a mix of dichloromethane and DMF (0.39 ml / 0.13 ml). The mixture was allowed to reach -15/-20 °C before adding N-methyl morpholine (8 μ l) followed 3 min latter with isobutyl chloroformate (7 μ l). The mixture was stirred at -15 °C for 10 minutes. Next, Asp (β -tert-butyl) methyl vinyl sulfone tosyl salt (13mg) was added in one shot, followed by N-methyl morpholine (8 μ l). The mixture was stirred 35 minutes at -15 /-20 °C and then diluted with 5 ml of dichloromethane and quenched with 1.5 ml of a saturated solution of sodium bicarbonate. After that, the organic layer was dried over magnesium sulphate. The solvent was evaporated to dryness. The obtained residue was purified on silica gel (ethyl acetate / hexane: 40% then dichloromethane/methanol: 5 to 15%) to afford 24 mg of the desired compound.

Example 33 : Synthesis of compound 59 (Z-Asp-Glu-Val-Aspmethyl vinyl sulfone)

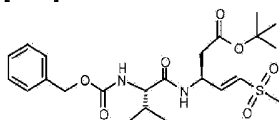
[0328]



[0329] Z-Asp(β -tert-Butyl)-Glu(β -tert-Butyl)-Val-Asp(β -tert-butyl)-methyl vinyl sulfone (22.7 mg) was dissolved in dichloromethane (0.55 ml), followed by addition of trifluoroacetic acid (0.7 ml). The mixture was stirred at room temperature overnight (15 h). It was then quenched with diethyl ether (7 ml) and the solvent was removed under vacuum. The obtained residue was diluted again with ether (5 ml) and the process was repeated twice. The precipitate was washed with 2* 1 ml of ether, dried to give 19 mg of the desired compound.

Example 34: Synthesis of compound 60 Z-Val-Asp(β -tert-Butyl)methyl vinyl sulfone

[0330]

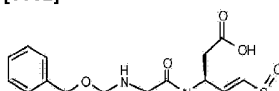


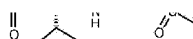
[0331] The Z-Val-OH (15 mg, 0.059 mmol) is dissolved in a mix of dichloromethane and DMF (0.5 ml / 0.2 ml). The mixture was allowed to reach -15/-20 °C before adding N-methyl morpholine (10 μ l) followed by isobutyl chloroformate (9 μ l). The mixture was stirred at -15 °C for 10 minutes. Next, Asp (β -tert-butyl) methyl vinyl sulfone tosyl salt (25mg, 1eq) was added in one shot, followed by N-methyl morpholine (10 μ l). The mixture was stirred 35 minutes at -15 /-20 °C and then diluted with 5 ml of dichloromethane and quenched with 1.5 ml of a saturated solution of sodium bicarbonate. After that, the organic layer was dried over magnesium sulphate. The solvent was evaporated to dryness. The obtained residue was purified on silica gel (gradient: methanol/ dichloromethane: 0 to 15%) to afford 24 mg of Z-Val- Asp-methyl vinyl sulfone.

NMR ^1H (CD_3OD , 400 MHz) δ : 7.38-7.29 (m, 5H); 6.82 (m, 1H); 6.69 (m, 1H); 5.10 (m, 2H); 4.96 (m, 1 H); 3.91 (m, 1 H); 2.94 (s, 3H); 2.8-2.6 (m, 2H); 2.1-2.0 (m, 1 H); 1.43 (s, 9H); 0.95 (t, 6H, J = 6.7 Hz).

Example 35 : Synthesis of compound 61 (Z-Val-Asp-methyl vinyl sulfone)

[0332]



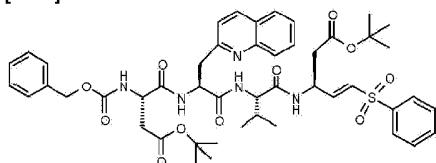


[0333] Z-Val-Asp(β -tert-butyl)-methyl vinyl sulfone (25 mg) was dissolved in dichloromethane (0.6 ml), followed by addition of trifluoroacetic acid (0.45 ml). The mixture was stirred at room temperature overnight (15 h). It was then quenched with diethyl ether (7 ml), the solvent was removed under vacuum. The obtained residue was diluted again with ether and the process was repeated twice. The precipitate was washed with 2* 1 ml of ether, dried to give 22 mg of the desired compound.

NMR ^1H (CD_3OD , 400 MHz) δ : 7.38-7.27 (m, 5H); 6.86 (m, 1H); 6.68 (m, 1H); 5.11 (m, 2H); 5.03 (m, 1H); 3.92 (m, 1H); 2.94 (s, 3H); 2.88-2.59 (m, 2H); 2.09-2.01 (m, 1H); 0.95 (t, 6H, J = 6.49 Hz).

Example 36 : Synthesis of compound 62 (Z-Asp(β -tert-Butyl)-Ala(2'-quinolyl)-Val-Asp(β -tert-Butyl)phenyl vinyl sulfone)

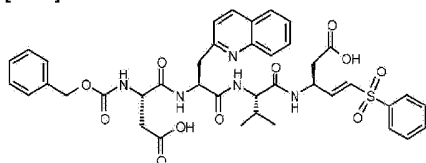
[0334]



[0335] The Z-Asp(β -tert-Butyl)-Ala(2'-quinolyl)-Val-OH (19 mg, 0.0306 mmol) is dissolved in a mix of dichloromethane and DMF (0.39 ml / 0.13 ml). The mixture was allowed to reach -15/-20 °C before adding N-methyl morpholine (8 μl) followed by isobutyl chloroformate (7 μl). The mixture was stirred at -15 °C for 10 minutes. Next, Asp (β -tert-butyl) phenyl vinyl sulfone tosyl salt (11 mg) was added in one shot, followed by N-methyl morpholine (8 μl). The mixture was stirred 35 minutes at -15/-20 °C and then diluted with 5 ml of dichloromethane and quenched with 1.5 ml of a saturated solution of sodium bicarbonate. After that, the organic layer was dried over magnesium sulphate. The solvent was evaporated to dryness. The obtained residue was purified on silica gel (gradient: ethyl acetate / hexane: 5 to 100%) to afford 13.7 mg of the desired compound.

Example 37: Synthesis of compound 63 (Z-Asp-Ala(2'-quinolyl)-Val-Aspphenyl vinyl sulfone)

[0336]

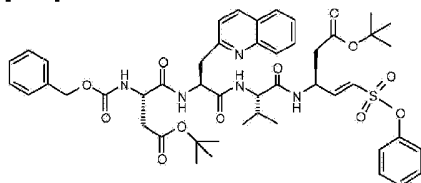


[0337] Z-Asp(β -tert-Butyl)-Ala(2'-quinolyl)-Val-Asp(β -tert-Butyl)-phenyl vinyl sulfone (13.7 mg) was dissolved in dichloromethane (0.3 ml), followed by addition of trifluoroacetic acid (0.4 ml). The mixture was stirred at room temperature overnight (15 h). It was then quenched with diethyl ether (5 ml), the solvent was removed under vacuum. The obtained residue was diluted again with ether and the process was repeated twice. The precipitate was washed with 2* 1 ml of ether, dried to give 10 mg of the desired compound.

NMR ^1H (CD_3OD , 400 MHz) δ : 8.58 (m, 1H); 8.11 (d, 1H, J = 8.75 Hz); 8.04 (d, 1H, J = 7.99 Hz); 7.85 (d, 3H, J = 7.25 Hz); 7.72-7.6 (m, 3H); 7.56 (t, 2H, J = 7.83 Hz); 7.31-7.28 (m, 5H); 6.93 (dd, 1H, J = 15.24 Hz, J = 5.15 Hz); 6.65 (d, 1H, J = 14.87 Hz); 4.97-4.80 (m, 4H); 4.40 (m, 1H); 4.21-4.02 (m, 1H); 3.63-3.50 (m, 1H); 3.47-3.44 (m, 1H); 2.82-2.6 (m, 4H); 2.15-1.98 (m, 1H); 0.81 (t, 6H, J = 6.38 Hz).
LCMS ($\text{M}-\text{H}^+$) = 800.5

Example 38 : Synthesis of compound 64 (Z-Asp(β -tert-Butyl)-Ala(2'-quinolyl)-Val-Asp(β -tert-Butyl)phenoxy vinyl sulfone)

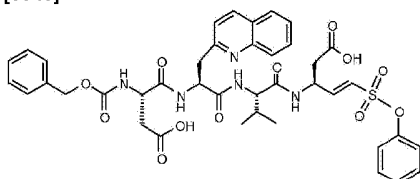
[0338]



[0339] The Z-Asp(β -tert-Butyl)-Ala(2'-quinolyl)-Val-OH (10 mg, 0.016 mmol) is dissolved in a mix of dichloromethane and DMF (0.24 ml / 0.080 ml). The mixture was allowed to reach -15/-20 °C before adding N-methyl morpholine (5 μ l) followed 3 min latter with isobutyl chloroformate (5 μ l). The mixture was stirred at -15 °C for 10 minutes. Next, Asp (β -tert-butyl) phenoxy vinyl sulfone tosyl salt (11.3 mg, 0.023 mmol) was added in one shot, the vial was washed with DMF (0.04 ml), followed by the addition of N-methyl morpholine (5 μ l). The mixture was stirred 35 minutes at -15/-20 °C and then diluted with 5 ml of dichloromethane and quenched with 1.5 ml of a saturated solution of sodium bicarbonate. After that, the organic layer was dried over magnesium sulphate. The solvent was evaporated to dryness. The obtained residue was purified on silica gel (gradient: ethyl acetate / hexane: 12 to 100%) to afford 4.7 mg of the desired compound.

Example 39 : Synthesis of compound 65 (Z-Asp-Ala(2'-quinolyl)-Val-Asp phenoxy vinyl sulfone)

[0340]

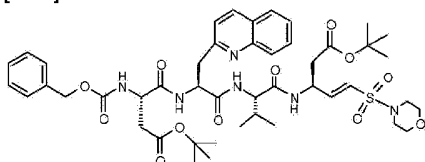


[0341] Z-Asp(β -tert-Butyl)-Ala(2'-quinolyl)-Val-Asp(β -tert-Butyl)-phenoxy vinyl sulfone (4.7 mg) was dissolved in dichloromethane (0.15 ml), followed by addition of trifluoroacetic acid (0.2 ml). The mixture was stirred at room temperature overnight (15 h). It was then quenched with diethyl ether (5 ml) and the solvent was removed under vacuum. The obtained residue was diluted again with ether (5 ml) and the process was repeated twice. The precipitate was washed with 2* 1 ml of ether, dried to give 4 mg of the desired compound.

NMR ^1H (CD_3OD , 400 MHz) δ : 8.25 (d, 1H, J = 7.9 Hz); 8.05 (d, 1H, J = 8.66 Hz); 7.88 (d, 1H, J = 8.25 Hz); 7.63 (d, 1H, J = 6.82 Hz); 7.53 (t, 1H, J = 7.46 Hz); 7.45-7.33 (m, 3H); 7.33-7.20 (m, 8H); 6.75 (dd, 1H, J = 15.33 Hz, J = 4.33 Hz); 6.65 (d, 1H, J = 15.55 Hz); 5.1-4.8 (m, 4H); 4.49 (t, 1H, J = 6.17 Hz); 4.06 (d, 1H, J = 6.45 Hz); 3.6-3.4 (m, 2H); 2.9 (dd, 1H, J = 17.02 Hz, J = 5.39 Hz); 2.73 (dd, 1H, J = 16.57 Hz, J = 6.53 Hz); 2.59 (d, 2H, J = 6.97 Hz); 2.08 (m, 1 H); 0.75 (m, 6H).

Example 40 : Synthesis of compound 66 (Z-Asp(β -tert-Butyl)-Ala(2'-quinolyl)-Val-Asp(β -tert-Butyl)morpholine vinyl sulfone)

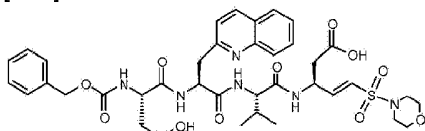
[0342]



[0343] Z-Asp(β -tert-Butyl)-Ala(2'-quinolyl)-Val-OH (13 mg, 0.0209 mmol) is dissolved in a mix of dichloromethane and DMF (0.28/0.094 ml). The mixture was allowed to reach -15/-20 °C before adding N-methyl morpholine (6 μ l) followed by isobutyl chloroformate (5 μ l). The mixture was stirred at -15 °C for 10 minutes. Next, Asp (β -tert-butyl) morpholine vinyl sulfone tosyl salt (10.2 mg, 1 eq) was added in one shot, followed by N-methyl morpholine (6 μ l). The mixture was stirred 35 minutes at -15/-20 °C and then diluted with 5 ml of dichloromethane and quenched with 1.5 ml of a saturated solution of sodium bicarbonate. After that, the organic layer was washed with sodium bicarbonate and dried over magnesium sulphate. The solvent was evaporated to dryness. The obtained residue was purified on silica gel (gradient: ethyl acetate / hexane: 12 to 100%) to afford 10 mg of the desired compound.

Example 41 : Synthesis of compound 67 (Z-Asp-Ala(2'-quinolyl)-Val-Aspmorpholine vinyl sulfone)

[0344]





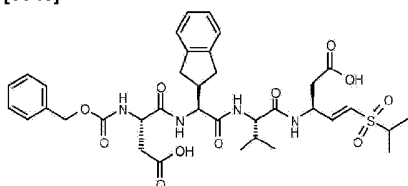
[0345] Z-Asp(β-tert-Butyl)-Ala(2'-quinolyl)-Val-Asp(β-tert-Butyl)-morpholine vinyl sulfone

(4.7 mg) was dissolved in dichloromethane (0.15 ml), followed by addition of trifluoroacetic acid (0.2 ml). The mixture was stirred at room temperature overnight (15 h). It was then quenched with diethyl ether (5 ml) and the solvent was removed under vacuum. The obtained residue was diluted again with ether and the process was repeated twice. The precipitate was washed with 2* 1 ml of ether, dried to give 4 mg of the desired compound.

NMR ¹H (CD₃OD, 400 MHz) δ: 8.39 (s, 1H); 8.08 (d, 1H, J= 8.56 Hz); 7.94 (d, 1H, J= 8.29 Hz); 7.74 (m, 1H); 7.66-7.55 (m, 2H); 7.31-7.29 (m, 5H); 6.69 (dd, 1H, J= 14.98 Hz, J= 4.59 Hz); 6.41 (d, 1H, J= 14.86 Hz); 5.0-4.7 (m, 4H); 4.45 (m, 1H); 4.05 (d, 1H, J= 6.65 Hz); 3.70-3.64 (m, 4H); 3.55-3.44 (m, 2H); 3.16-3.08 (m, 4H); 2.89-2.70 (m, 4H); 2.17-2.07 (m, 1H); 0.85-0.81 (m, 6H).
LCMS (M-H⁺) = 809.6

Example 42 : Synthesis of compound 68 (Z-Asp-Indanylglycine-Val-Aspisopropyl vinyl sulfone)

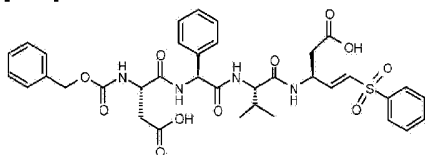
[0346]



[0347] This compound was synthesized from Asp(β-tert-butyl)isopropyl vinyl sulfone tosyl salt (43) and Cbz-Asp(O-*t*Bu)Indanylglycine-Val-OH (12) via anhydride mixte method as for Z-Asp-Indanylglycine-Val- Aspmethyl vinyl sulfone (57).

Example 43 : Synthesis of compound 69 (Z-Asp-Phg-Val- Asp-phenyl vinylsulfone)

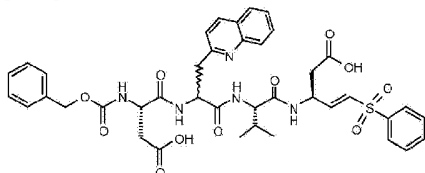
[0348]



[0349] This compound was synthesized from Z-Asp(β-tert-Butyl)-Phg-Val-OH (16) and Asp(O*t*Bu)-Vinyl phenyl sulfone tosyl salt (37) via anhydride mixte method as for Z-Asp-Ala(2'-quinolyl)-Val-Aspphenyl vinyl sulfone (63).

Example 44: Synthesis of compound 70 (Z-Asp-(D, L Ala(2'-quinolyl))-Val-Aspphenyl vinylsulfone)

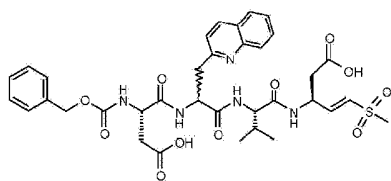
[0350]



[0351] This compound was synthesized as the corresponding Z-Asp-Ala(2'-quinolyl)-Val-Aspphenyl vinyl sulfone (63).

Example 45: Synthesis of compound 71 (Z-Asp-(D, L Ala(2'-quinolyl))-Val-Aspmethyl vinylsulfone)

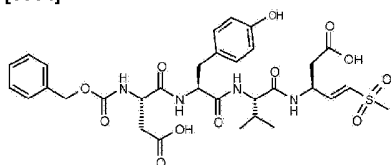
[0352]



[0353] This compound was synthesized as the corresponding Z-Asp-Ala(2'-quinolyl)-Val-Aspmethyl vinyl sulfone (55).

Example 46 : Synthesis of compound 76 (Z-Asp-Tyr-Val-Aspmethyl vinyl sulfone)

[0354]

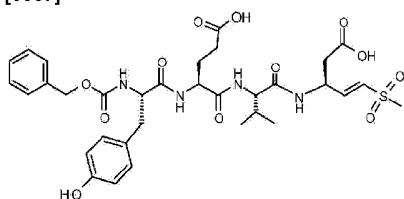


[0355] This compound was synthesized from Z-Asp-Tyr(OtBu)-Val-OH (75) and Asp(OtBu)-Vinyl methy sulfone tosyl salt (33) via anhydride mixte method as for Z-Asp-Phg-Val-Aspmethyl vinyl sulfone (53).

[0356] Z-Asp-Tyr(OtBu)-Val-OH was synthesized as Cbz-Asp(O-tBu)-Glu(O-tBu)-Val-OH (20).

Example 47 : Synthesis of compound 82 (Z-Tyr-Glu-Val-Aspmethyl vinyl sulfone)

[0357]

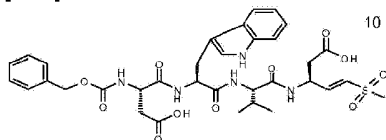


[0358] This compound was synthesized from Z-Tyr(OtBu)-Glu(OtBu)-Val-OH (80) and Asp(OtBu)-Vinyl methy sulfone tosyl salt (33) via anhydride mixte method as for Z-Asp-Phg-Val-Aspmethyl vinyl sulfone (53).

Z-Tyr(OtBu)-Glu(OtBu)-Val-OH (80) was synthesized as Cbz-Asp(O-tBu)-Glu(O-tBu)-Val-OH (20).

Example 48 : Synthesis of compound 83 (Z-Asp-Trp-Val-Aspmethyl vinyl sulfone)

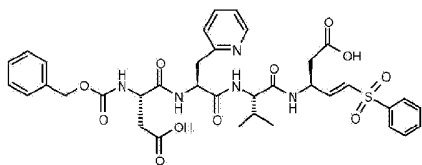
[0359]



[0360] This compound was synthesized from Z-Asp(OtBu)-Trp-Val-OH (86) and Asp(OtBu)-Vinyl methy sulfone tosyl salt (33) via anhydride mixte method as for Z-Asp-Phg-Val-Aspmethyl vinyl sulfone (53).

Example 49: Synthesis of compound 85 (Z-Asp-Ala(2'-pyridyl)-Val-Aspmethyl vinylsulfone)

[0361]



[0362] This compound was synthesized from Z-Asp(OtBu)-Ala(2'-pyridyl)-Val-OH (83) and Asp(OtBu)-Vinyl phenyl sulfone tosyl salt (37) via anhydride mixture method as for Z-Asp-Ala(2'-quinolyl)-Val-Asp(phenyl vinyl sulfone) (63).

Example 50: Selectivity of exemplary compounds for caspase-3 relative to caspase-1, caspase-5, caspase-7, and caspase-9

[0363] Selectivity of compound 55, compound 63, compound 48, compound 57, compound 88 toward caspase-1 (pro-inflammatory group), caspase-5 (group I), caspase-9 (group II) and caspase-3 and caspase-7 (group III) was evaluated by using fluorometric methods using the Caspase-1, -3, -5, -7, -9 Inhibitor Drug Screening Kit™ (Catalog #: K151-100, K153-100, K155-100, K157-100, K159-100 respectively, BioVision™). Briefly, using instructions of the manufacturer, a wide range of different concentrations of the compound: 3333, 1000, 333, 100, 33, 10, 3, and 1nM (final concentration) was added directly to the reaction mixtures containing the substrate and the enzyme in a final volume of 10 µl. After a 30-minute incubation at 37°C, the liberation of AFC was measured as an endpoint assay using the Flexstation3™ (Molecular Devices) with an excitation wavelength of 400 nm and an emission wavelength of 505 nm. The level of inhibition of caspase-1, -3, -5, -7, -9 activity was determined by comparison of the relative fluorescence intensity in samples with or without the compound. Results are summarized in **Table 2** herein after.

[0364] As shown in **Table 2**, Compound 55 showed an inhibitory effect on both caspase-3 and -7 activity. However, based on the IC₅₀ values calculated, it was about 200 fold more selective of Caspase-3 over Caspase-7 (see Table 2). No significant inhibitory activity of the compound was observed for caspase-1, -5, and -9 at the tested dose-range. Inhibition of Caspase-1, -5, and -9 was achieved (25%, 21%, and 57% respectively) but at extremely high concentration (about 10,000nM).

[0365] Similar to Compound 55, Compound 63 also showed high selectivity in caspase-3 activity inhibition relative to caspase-1, -5, -7, and -9. Compound 63 also showed an inhibitory effect on caspase-3 and -7, with about a 50 fold selectivity of caspase-3 over caspase-7. No significant inhibitory activity was observed for caspase-1, -5, and -9 at the tested dose-range of Compound 63. The data indicates that both Compound 55 and Compound 63 are highly potent and selective compounds for caspase-3 activity inhibition.

[0366] Compound 48 was able to inhibit specific groups of caspases. The compound was able to inhibit caspase-3, -7, and -9 with IC₅₀ values of about 8-30 nM, 0.4-0.9 µM and 1-1.8 µM respectively. Thus Compound 48 shows inhibition of Group III caspases (caspase-3 and -7) with high potency, and inhibition of Group II caspase (caspase-9) with a lower yet significant efficacy.

[0367] From the inhibition profile of Compound 57 on caspase-1, -3, -5, -7, and -9 activity, it can be appreciated that Compound 57 is a potent inhibitor of Group III caspases (caspase-3 and -7). Compound 57 inhibited caspase-3 and -7 activities with IC₅₀ values of about 30-90 nM and about 180-300 nM respectively. Therefore, data indicates that Compound 48 and Compound 57 are not specific inhibitors for caspase-3 activity, but are in fact inhibitors targeted at specific groups of caspases.

[0368] Compound 88 has a dual inhibition effect on both caspase-1 and caspase-3 activity. It is a potent inhibitor of caspase-3 activity, with an IC₅₀ value of about 30-90 nM. It also inhibits caspase-1 activity with an IC₅₀ value of about 0.6-1.2 µM.

Example 51: *In vitro* inhibitory activity of Caspase-3 inhibitors

[0369] To test the efficacy of caspase-3 inhibitors at the cellular level, the ability of selected compounds to inhibit the proteolytic cleavage of PARP (poly ADP-ribose polymerase) was evaluated in live Hela cells.

[0370] Briefly, in this assay Hela cells are seeded in 96 well plates and incubated for 4 hours with staurosporine, a well characterized inducer of apoptosis, alone or together with different concentrations of compound (50, 25, 10 and 3µM). After formaldehyde-based fixation, the cells are stained with a fluorescein-labeled anti-cleaved PARP antibody (Cell signaling, Cat#: 9547) and counterstained with Hoechst33342 (Invitrogen, Cat#: H3570) to mark all nuclei. Fluorescence images are taken on a Cellomics™ microscope system (Thermo Scientific, Pittsburgh, USA) with the Hoechst stain in the blue channel and the cleaved PARP antibody stain in the green channel. The percentage of cleaved PARP positive cells is determined by calculating the ratio between nuclei with a cleaved PARP antibody staining above a certain threshold and all (Hoechst positive) nuclei. The efficacy of caspase-3 inhibition is determined by calculating the ratio between cleaved PARP positive cells after staurosporine incubation together with compounds and staurosporine incubation without compounds. Results are summarized in **Table 2**.

[0371] As shown in **Table 2**, results of this assay show that compounds that inhibited Caspase-3 activity in the enzymatic assay with an IC₅₀ below 100nM were generally also effective in inhibiting PARP cleavage *in vitro*, although a major factor contributing to a compound's activity / effectivity in this assay is the compound's cell membrane permeability coefficient (the higher the permeability coefficient, the greater the amount of the compound in the media reaching inside the cell). Certain modifications in the molecular composition of the compounds improved the inhibitory activity further; for example Compound **51**, Compound **53**, Compound **57**, and Compound **76** reduced the percentage of PARP positive cells after staurosporine treatment by more than 50%, with Compound **51** and Compound **57** being the most effective compounds reaching values of about 65% inhibition, and Compound **53** also being highly effective relative to **DEVD-fmk** (positive control compound with low caspase-3 selectivity but highly cell membrane permeable with general caspase inhibitory activity), and Compounds **48**, **55**, and **59** also showing effectivity especially relative to **DEVD-fmk**. These results indicate that inhibitory compounds identified in a primary enzymatic assay screen retain their activity in a cellular environment and that molecular modifications allow to further improve their activity as caspase-3 inhibitors.

Table 2: Activity assays for caspase inhibitors. Values given are approximations of the average values obtained from the assays.

No	Compound Name	Caspase enzymatic activity, IC ₅₀ (uM)					% inhibition of PARP cleavage at 50uM compound	% inhibition of PARP cleavage at 50uM DEVDfMK (positive control)	% of inhibition with compound relative to DEVDfMK	IC ₅₀ (uM) Glo3 /7
		Caspase 1	Caspase 3	Caspase 5	Caspase 7	Caspase 9				
48	Z-Asp-Ala(2'-quinolyl)-Val-Asp alpha chlorovinyl methylsulfone	>3.33	0.01	>3.33	0.4	1.2	45	70	65	0.1
50	Tosyl-Ala(2'-quinolyl)-Val-Asp alpha chlorovinyl methylsulfone	>3.33	>3.33							
36	Boc-Asp(O-tBu)VSphenyl		>2							
37	Asp(O-tBu)VSphenyl salt		>2							
16	Z-Asp(O-tBu)-Phg-Val-OH		>2							
51	Z-Asp(O-Methyl)-Indanylglycine-Val-Asp(O-Methyl)VS methyl	>3.0	1	>3.0	>3.0	>3.0	65	80	80	
53	Z-Asp-Phg-Val-AspVS methyl	2	0.06	>10	0.8	>10	55	70	80	0.1
93	AspVS methyl salt		>10							
33	Asp(O-tBu)VS methyl salt		>10							
71	Z-Asp-d,l Ala(2'-quinolyl)-Val-AspVS methyl	>10	0.1	>10	8.9	>10				
55	Z-Asp-Ala(2'-quinolyl)-Val-AspVS methyl	>10	0.01	>10	1.4	>3.33	40	65	60	0.1
65	Z-Asp-Ala(2'-quinolyl)-Val-AspVS phenoxy	1.2	0.01	>3.33	0.4	0.3	30	75	40	0.2
70	Z-Asp-d,l Ala(2'-quinolyl)-Val-AspVS phenyl		0.3	>10	>10	>10				
63	Z-Asp-Ala(2'-quinolyl)-Val-AspVS phenyl	>3.33	0.04	>3.33	1.8	>3.33	10	70	15	0.9
66	Z-Asp-Ala(2'-quinolyl)-Val-AspVS morpholine		0.2				25			
85	Z-Asp-Ala(2'-pyridyl)-Val-AspVS phenyl	>3.33	0.03	>3.33	1.0	>3.33	40			

No	Compound	Caspase enzymatic activity, IC50 (uM)					% inhibition of PARP cleavage at 50uM compound	% inhibition of PARP cleavage at 50uM DEVDFMK (positive control)	% of inhibition with compound relative to DEVDFMK	IC50 (uM) Glo3 /7
	Name	Caspase 1	Caspase 3	Caspase 5	Caspase 7	Caspase 9				
69	Z-Asp-Phg-Val-AspVSphenyl		0.1							
76	Z-Asp-Tyr-Val-AspVSmethyl	>3.33	0.03	>3.33	0.7	>3.33	55	90	60	0.04
57	Z-Asp-Indanylglycine-Val-AspVSmethyl	0.9	0.03	>3.33	0.2	>3.33	65	90	70	0.1
88	Z-Asp-Trp-Val-AspVSmethyl	0.7	0.04	>3.33	1.1	>3.33	45	90	50	
59	Z-Asp-Glu-Val-AspVSmethyl	1.4	0.02	>3.33	0.04	0.5	45	65	70	
61	Z-Val-AspVSmethyl	>3.33	>3.33	>3.33	>3.33	>3.33	5	70	5	
68	Z-Asp-Indanylglycine-Val-AspVSisopropyl	>3.33	0.2	>3.33	1.5	>3.33	30	60	50	
96	Z-Tyr-Val-Ala-AspVSphenyl	1.3	>10	>3.33	>10	>10				
82	Z-Tyr-Glu-Val-AspVSmethyl	0.5	0.3	>3	>5	1.6				
	DEVD-fmk (positive control compound)	0.4	0.6	>3.33	1	0.5	85 (20 uM IC50)			

REFERENCES CITED IN THE DESCRIPTION

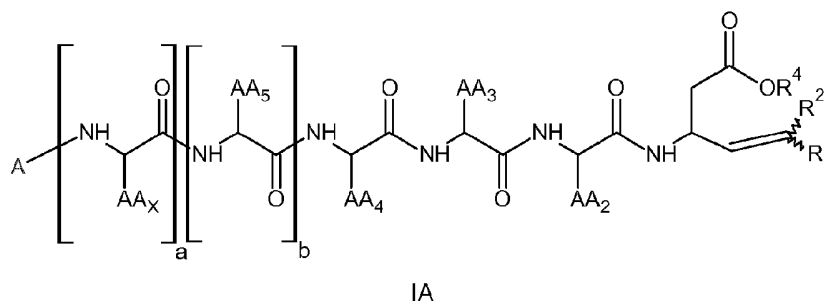
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Patent documents cited in the description

- [US5976850A \[0004\]](#)
- [US6267540B \[0004\]](#)

Non-patent literature cited in the description

- RODRIGUEZ et al. J Exp Med., 1996, vol. 184, 52067-72 [0004]
- YAOITA et al. Circulation, vol. 97, 276-281 [0004]
- HANZLIK THOMPSON J. Med. Chem., 1984, vol. 27, 6711-712 [0004]
- THOMPSON et al. J. Med. Chem., 1986, vol. 29, 1104-111 [0004]
- T. GREENE P. WUTS Protecting Groups in Chemical Synthesis John Wiley & Sons 1999 0000 [0002]
- BUNDGARD, H. Design of Prodrugs Elsevier 1985 00007-921-24 [0007]
- DRAGOVICH P., S. HOB T, NMM, EDC, 1999, [0009]
- Cysteine protease ZBIGNIEW GRZONKA Industrial Enzymes Springer 2007 0000 181-195 [0101]
- LOHMANN et al. Predicting blood-brain barrier permeability of drugs: evaluation of different in vitro assays J Drug Target, 2002, vol. 10, 263-276 [0103]
- NICOLAZZO et al. Methods to assess drug permeability across the blood-brain barrier J Pharm Pharmacol, 2006, vol. 58, 281-293 [0103]

Patentkrav**1. Forbindelse med formel IA**

hvor

5 a er 0 eller 1;

b er 0 eller 1 forudsat at når b er 0, a er 0;

A er

1) H,

2) C₁-C₆ alkyl,

10 3) aryl,

4) heteroaryl,

5) heterocyclyl,

6) R³-OC(O)-,

7) PhCH₂OC(O)-, eller

15 8) R³-S(O)₂-;

AA₂ er (R)- eller (S)-aminosyresidekæden af Val, Leu, Pro, Met, Ala, Thr, His, Ser eller Lys;

AA₃ er (R)- eller (S)-aminosyresidekæden af Trp, Tyr, Ala, Asp, Gln, Glu, Phe, Ser, Thr, Val, Gly, Leu, His, eller Ile, eller AA₃ er phenylglycin, indanylglycin, eller Ala-(2'-quinolyl);

5 AA₄ er (R)- eller (S)-aminosyresidekæden af Asp, Ile, Leu, Glu, Ala, Val, Tyr, Trp, Phe, eller Pro;

AA₅, når til stede, er (R)- eller (S)-aminosyresidekæden af Val eller Leu;

AA_x, når til stede, er (R)- eller (S)-aminosyresidekæden af hvilken som helst D- eller L-aminosyrerest;

bølgelinjen betegner enten cis- eller trans-stilling af R¹ og R²;

10 R¹ er

1) aryl,

2) heteroaryl,

3) heterocyclyl,

4) C₂-C₆ alken-R²⁰,

15 5) SO₂R⁵,

6) SO₃R⁵,

7) SO⁵,

8) SONHR⁵,

9) SO₂NHR⁵,

20 10) CN,

11) CO²R⁵,

12) COR⁵,

13) PO_3R^5 ,

14) $\text{PO}(\text{OR}^5)_2$, eller

15) $\text{PO}(\text{OR}^5)$,

5 hvor aryl, heteroaryl, eller heterocyclyl eventuelt er substitueret med én eller flere R^{30} ;

R^2 er

1) R^1 ; eller

2) H,

3) halogen,

10 4) haloalkyl,

5) C_1 - C_6 alkyl,

6) C_2 - C_6 alken,

7) C_3 - C_7 cycloalkyl,

8) OR^9 ;

15 9) OCOR^6 ,

10) OCO_2R^6 ,

11) NR^7R^8 ,

12) NHSO_2R^6 ,

13) NHCOR^6 ,

20 14) aryl,

15) heteroaryl, eller

16) heterocyclyl;

R³ er

1) C₁-C₆ alkyl,

2) aryl,

5 3) heteroaryl, eller

4) heterocyclyl;

R⁴ er

1) H, eller

2) C₁-C₆ alkyl;

10 R⁵ er

1) H,

2) C₁-C₆ alkyl,

3) C₂-C₆ alken,

4) C₃-C₇ cycloalkyl,

15 5) aryl,

6) heteroaryl,

7) heterocyclyl, eller

8) en hvilken som helst beskyttet (D)- eller (L)-aminosyrerest;

R⁶ er

20 1) en hvilken som helst (D)- eller (L)-aminosyrerest,

5

2) C₁-C₆ alkyl,3) C₃-C₇ cycloalkyl,

4) aryl,

5) heteroaryl, eller

5 6) heterocyclyl,

hvor alkyl eller cycloalkyl eventuelt er substitueret med én eller flere R¹⁰ substituent; og hvor aryl, heteroaryl eller heterocyclyl eventuelt er substitueret med én eller flere R²⁰ substituent;

R⁷ og R⁸ er uafhængigt valgt fra:

10 1) H,

2) C₁-C₆ alkyl,3) C₃-C₇ cycloalkyl,

4) haloalkyl,

5) aryl,

15 6) heteroaryl, eller

7) heterocyclyl,

hvor alkyl og cycloalkyl eventuelt er substitueret med én eller flere R¹⁰ substituent, og aryl, heteroaryl og heterocyclyl eventuelt er substitueret med én eller flere R²⁰ substituent;

20 R⁹ er

1) H,

2) C₁-C₆ alkyl,

3) C₃-C₇ cycloalkyl,

4) aryl,

5) heteroaryl, eller

6) heterocyclyl,

5 hvor alkyl eller cycloalkyl eventuelt er substitueret med én eller flere R¹⁰ substituent; og hvor aryl, heteroaryl eller heterocyclyl eventuelt er substitueret med én eller flere R²⁰ substituent;

R¹⁰ er uafhængigt valgt fra:

1) halogen,

10 2) C₁-C₆ alkyl,

3) C₃-C₇ cycloalkyl,

4) haloalkyl,

5) aryl,

6) heteroaryl,

15 7) heterocyclyl,

8) OR⁹,

9) S(O)_mR⁹,

10) NR⁷R⁸,

11) COR⁹,

20 12) C(O)OR⁹,

13) OC(O)R⁹,

14) SC(O)R^9 ,

15) CONR^7R^8 , eller

16) $\text{S(O)}_2\text{NR}^7\text{R}^8$;

R^{20} er uafhængigt valgt fra:

5 1) halogen,

2) NO_2 ,

3) CN ,

4) $\text{C}_1\text{-C}_6$ alkyl,

5) haloalkyl,

10 6) $\text{C}_3\text{-C}_7$ cycloalkyl,

7) OR^7 ,

8) NR^7R^8 ,

9) SR^7 ,

10) aryl,

15 11) heteroaryl,

12) heterocyclyl,

13) SO_2R^5 ,

14) SO_3R^5 ,

15) SOR^5 ,

20 16) SONHR^5 ,

- 17) SO_2NHR^5 ,
- 18) PO_3R^5 ,
- 19) $\text{PO}(\text{OR}^5)_2$,
- 20) $\text{PO}(\text{OR}^5)$,
- 5 21) COR^7 ,
- 22) CO_2R^7 ,
- 23) $\text{S}(\text{O})_m\text{R}^7$,
- 24) CONR^7R^8 , eller
- 25) $\text{S}(\text{O})_2\text{NR}^7\text{R}^8$,
- 10 hvor alkyl og cycloalkyl eventuelt er substitueret med én eller flere R^6 substituent; og hvor aryl, heteroaryl, eller heterocyclyl eventuelt er substitueret med én eller flere R^{30} ;
- R^{30} er
- 1) NO_2 ,
- 15 2) $\text{C}_2\text{-C}_6$ alken- R^{20} ,
- 3) SO_2R^5 ,
- 4) SOR^5 ,
- 5) SONHR^5 ,
- 6) SO_2NHR^5 ,
- 20 7) CN ,
- 8) CO_2R^5 ,

9) COR⁵,

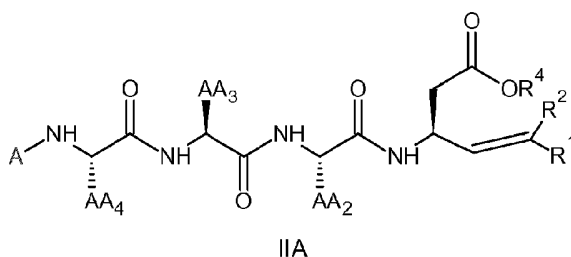
10) PO₃R⁵,

11) PO(OR⁵)₂, eller

12) PO(OR⁵);

5 eller et farmaceutisk acceptabelt salt deraf, eller hvor forbindelsen er mærket med en detekterbar markør deraf, hvor den detekterbare markør er en radioisotop.

2. Forbindelse med formel IIA



10

hvor:

A er

1) H,

2) C₁-C₆ alkyl,

15

3) aryl,

4) heteroaryl,

5) heterocyclyl,

6) R³-OC(O)-,

7) PhCH₂OC(O)-, eller

20

8) R³-S(O)₂-;

AA₂ er aminosyresidekæden af Val, Leu, Pro, Met, Ala, Thr, His, Ser, eller Lys,

AA₃ er aminosyresidekæden af Trp, Tyr, Ala, Asp, Gln, Phe, Ser, Thr, Val, Gly, Leu, eller

5 AA₃ er phenylglycin, indanylglycin, eller Ala-(2'-quinolyl),

AA₄ er aminosyresidekæden af Asp; eller

AA₂ er aminosyresidekæden af Thr, His, Val, Trp, Ile, eller Ala,

AA₃ er aminosyresidekæden af Glu eller AA₃ er Ala-(2'-quinolyl),

AA₄ er aminosyresidekæden af Ile, Leu, Glu, Asp, Ala, Pro eller Val; eller

10 AA₂ er aminosyresidekæden af Val, Ala, Thr, eller His,

AA₃ er aminosyresidekæden af Glu, Gln, Asp, Ala, Gly, Thr, Val, Trp, eller

AA₃ er phenylglycin eller indanylglycin,

AA₄ er aminosyresidekæden af Tyr, Trp, Phe, eller Asp;

Og hvor

15 R¹ er

1) aryl,

2) heteroaryl,

3) heterocyclyl,

4) C₂-C₆ alken-R²⁰,

20 5) SO₂R⁵,

6) SO₃R⁵,

7) SOR⁵,

8) SONHR^5 ,

9) SO_2NHR^5 ,

10) CN ,

11) CO^2R^5 ,

5 12) COR^5 ,

13) PO_3R^5 ,

14) $\text{PO}(\text{OR}^5)_2$, eller

15) $\text{PO}(\text{OR}^5)$,

hvor aryl, heteroaryl, eller heterocyclyl eventuelt er substitueret med én eller flere

10 R^{30} ;

R^2 er

1) R^1 ; eller

2) H ,

3) halogen,

15 4) haloalkyl,

5) C_1 - C_6 alkyl,

6) C_2 - C_6 alken,

7) C_3 - C_7 cycloalkyl,

8) OR^9 ;

20 9) OCOR^6 ,

10) OCO_2R^6 ,

- 11) NR^7R^8 ,
- 12) NHSO_2R^6 ,
- 13) NHCOR^6 ,
- 14) aryl,
- 5 15) heteroaryl, eller
- 16) heterocyclyl;
- R^3 er
- 1) $\text{C}_1\text{-C}_6$ alkyl,
- 2) aryl,
- 10 3) heteroaryl, eller
- 4) heterocyclyl;
- R^4 er
- 1) H, eller
- 2) $\text{C}_1\text{-C}_6$ alkyl;
- 15 R^5 er
- 1) H,
- 2) $\text{C}_1\text{-C}_6$ alkyl,
- 3) $\text{C}_2\text{-C}_6$ alken,
- 4) $\text{C}_3\text{-C}_7$ cycloalkyl,
- 20 5) aryl,

6) heteroaryl,

7) heterocyclyl, eller

8) en hvilken som helst beskyttet (D)- eller (L)-aminosyrerest;

R⁶ er

5 1) en hvilken som helst (D)- eller (L)-aminosyrerest,

2) C₁-C₆ alkyl,

3) C₃-C₇ cycloalkyl,

4) aryl,

5) heteroaryl, eller

10 6) heterocyclyl,

hvor alkyl eller cycloalkyl eventuelt er substitueret med én eller flere R¹⁰ substituent; og hvor aryl, heteroaryl eller heterocyclyl eventuelt er substitueret med én eller flere R²⁰ substituent;

R⁷ og R⁸ er uafhængigt valgt fra:

15 1) H,

2) C₁-C₆ alkyl,

3) C₃-C₇ cycloalkyl,

4) haloalkyl,

5) aryl,

20 6) heteroaryl, eller

7) heterocyclyl,

hvor alkyl og cycloalkyl eventuelt er substitueret med én eller flere R^{10} substituent, og aryl, heteroaryl og heterocyclyl eventuelt er substitueret med én eller flere R^{20} substituent;

R^9 er

- 5 1) H,
- 2) C_1 - C_6 alkyl,
- 3) C_3 - C_7 cycloalkyl,
- 4) aryl,
- 5) heteroaryl, eller
- 10 6) heterocyclyl,

hvor alkyl eller cycloalkyl eventuelt er substitueret med én eller flere R^{10} substituent; og hvor aryl, heteroaryl eller heterocyclyl eventuelt er substitueret med én eller flere R^{20} substituent;

R^{10} er uafhængigt valgt fra:

- 15 1) halogen,
- 2) C_1 - C_6 alkyl,
- 3) C_3 - C_7 cycloalkyl,
- 4) haloalkyl,
- 5) aryl,
- 20 6) heteroaryl,
- 7) heterocyclyl,
- 8) OR^9 ,

15

9) $S(O)_mR^9$,10) NR^7R^8 ,11) COR^9 ,12) $C(O)OR^9$,5 13) $OC(O)R^9$,14) $SC(O)R^9$,15) $CONR^7R^8$, eller16) $S(O)_2NR^7R^8$; R^{20} er uafhængigt valgt fra:

10 1) halogen,

2) NO_2 ,3) CN ,4) C_1 - C_6 alkyl,

5) haloalkyl,

15 6) C_3 - C_7 cycloalkyl,7) OR^7 ,8) NR^7R^8 ,9) SR^7 ,

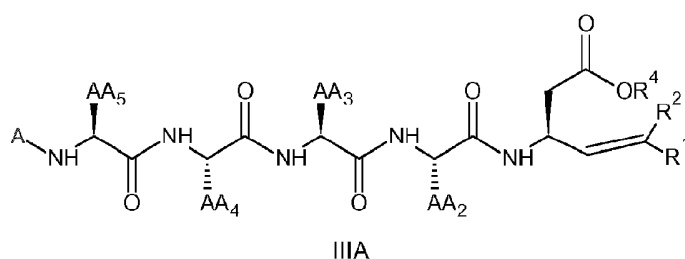
10) aryl,

20 11) heteroaryl,

- 12) heterocyclyl,
- 13) SO_2R^5 ,
- 14) SO_3R^5 ,
- 15) SOR^5 ,
- 5 16) SONHR^5 ,
- 17) SO_2NHR^5 ,
- 18) PO_3R^5 ,
- 19) $\text{PO}(\text{OR}^5)_2$,
- 20) $\text{PO}(\text{OR}^5)$,
- 10 21) COR^7 ,
- 22) CO_2R^7 ,
- 23) $\text{S}(\text{O})_m\text{R}^7$,
- 24) CONR^7R^8 , eller
- 25) $\text{S}(\text{O})_2\text{NR}^7\text{R}^8$,
- 15 hvor alkyl og cycloalkyl eventuelt er substitueret med én eller flere R^6 substituent; og hvor aryl, heteroaryl, eller heterocyclyl eventuelt er substitueret med én eller flere R^{30} ;
- R^{30} er
- 1) NO_2 ,
- 20 2) $\text{C}_2\text{-C}_6$ alken- R^{20} ,
- 3) SO_2R^5 ,

- 4) SOR^5 ,
- 5) SONHR^5 ,
- 6) SO_2NHR^5 ,
- 7) CN ,
- 5 8) CO_2R^5 ,
- 9) COR^5 ,
- 10) PO_3R^5 ,
- 11) $\text{PO}(\text{OR}^5)_2$, eller
- 12) $\text{PO}(\text{OR}^5)$;
- 10 eller et farmaceutisk acceptabelt salt deraf, eller hvor forbindelsen er mærket med en detekterbar markør deraf, hvor den detekterbare markør er en radioisotop.

3. Forbindelsen ifølge krav 1, hvor forbindelsen er en forbindelse med formel IIIA



15 hvor

AA_2 er aminosyresidekæden af Ala, Ser, Lys eller Val;

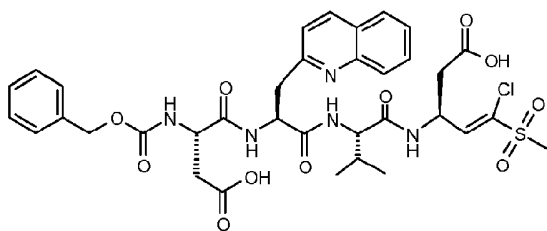
AA_3 er aminosyresidekæden af Val, Glu, Thr, eller Gln;

AA_4 er aminosyresidekæden af Asp, eller Leu;

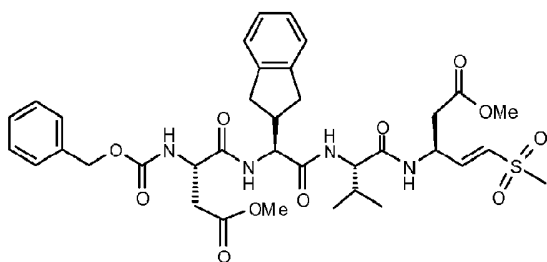
AA_5 er aminosyresidekæden af Val eller Leu;

20 og hvor A, R^1 , R^2 og R^4 er som defineret i krav 1.

4. En forbindelse, ifølge krav 1, valgt fra gruppen bestående af:

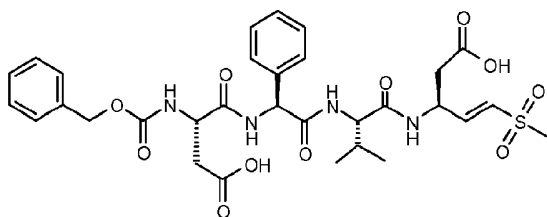


Forbindelse #48;

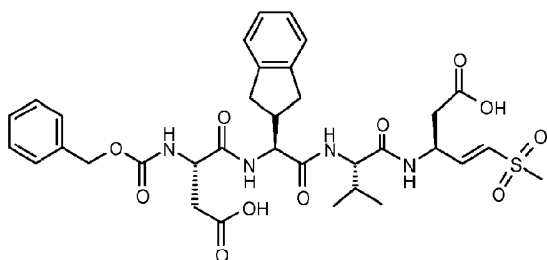


5

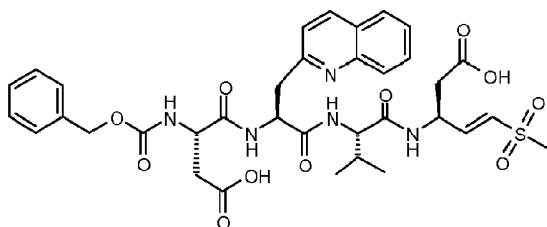
Forbindelse #51;



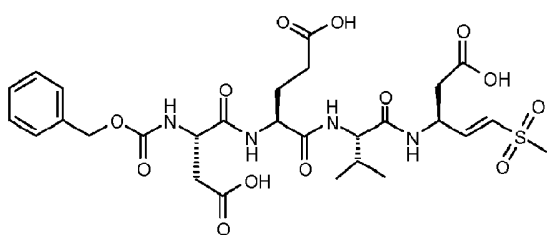
Forbindelse #53;



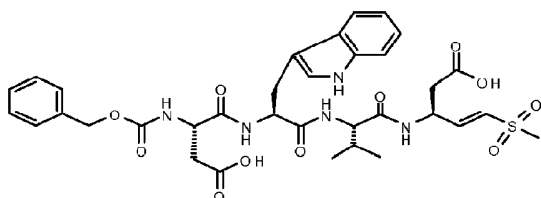
10 Forbindelse #57;



Forbindelse #55;



Forbindelse #59; og



Forbindelse #88.

5

5. Farmaceutisk sammensætning til forebyggelse og/eller behandling af en caspase-relateret sygdom i et individ med behov derfor, omfattende en effektiv mængde af en forbindelse ifølge et hvilket som helst af kravene 1-4.

10 **6.** Forbindelse ifølge et hvilket som helst af kravene 1-4 for forebyggelse og/eller behandling af en caspase-relateret sygdom i et individ med behov derfor.

7. Forbindelsen til anvendelse ifølge krav 6, hvor den caspase-relaterede sygdom er valgt fra gruppen bestående af apoptosis-medierede sygdomme, IL-1-

15 medierede sygdomme, inflammatoriske sygdomme, autoimmune sygdomme, autoinflammatoriske sygdomme, proliferative sygdomme, infektionssygdomme, degenerative sygdomme, nethindeligelser, inflammatorisk peritonitis, osteoarthritis, pancreatitis, astma, respiratorisk distress syndrom, reumatoid arthritis, systemisk lupus erythematosus, scleroderma, Grave's sygdom,

20 autoimmune gastritis, diabetes, autoimmun hæmolytisk anæmi, autoimmun neutropeni, thrombocytopeni, hepatitis, inflammatorisk tarmsygdom, Crohn's sygdom, psoriasis, dermatitis, Graft vs host sygdom, eller gentransplantatafstødning, osteoporose, leukæmier og relaterede lidelser, multiple myelom-relaterede sygdomme, metastatiske melanomer, Kaposi's

25 sarkom, sepsis, septisk chok, Alzheimer's sygdom, Parkinson's sygdom, Huntington's sygdom, cerebral iskæmi, epilepsi, myokardial iskæmi, akut og kronisk hjertesygdom, hjerteinfarkt, kongestiv hjertefejl, atherosclerose, spinal muskelatrofi, amyotrof lateral sclerose, multipel sclerose, HIV-relateret

encephalitis, ældning, neurologisk skade på grund af slagtilfælde, ulcerativ colitis, traumatisk hjerteskaade, rygmarvsskaade, hepatitis-B, hepatitis-C, hepatitis-G, lever-relaterede sygdomme, nyresygdom, og HIV-infektion.

- 5 **8.** Anvendelse af en effektiv mængde af én eller flere forbindelser ifølge et hvilket som helst af kravene 1-4 til behandling af overdreven apoptosis forårsaget af caspaseaktivitet i en *ex vivo*-celle eller et *ex vivo*-væv.

- 9.** Forbindelse ifølge et hvilket som helst af kravene 1-4 til anvendelse i
10 behandling af overdreven apoptosis forårsaget af caspaseaktivitet i en *in vivo*-celle eller et *in vivo*-væv.

- 10.** Forbindelsen ifølge et hvilket som helst af kravene 1-4, hvor det farmaceutisk acceptable salt er et der tillader forbindelsen af penetrere celler.