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(54) Title: COMPOUNDS WHICH INHIBIT NEURONAL EXOCYTOSIS (II)

(57) Abstract: Compounds of general formula (I): R₁-W_n-X_m-AA₁-AA₂-AA₃-AA₄-AA₅-AA₆-AA₇-Y_p-Z_q-R₂, their stereoisomers, mixtures thereof and/or their cosmetically or pharmaceutically acceptable salts, preparation processes, cosmetic or pharmaceutical compositions which contain them and their use in medicine, particularly in the treatment and/or prevention of pain, inflammation, itching, neurological, compulsive and/or neuropsychiatric diseases and/or disorders and in processes of treatment and/or care of the skin, hair and/or mucous membranes mediated by neuronal exocytosis.

COMPOUNDS WHICH INHIBIT NEURONAL EXOCYTOSIS (II)**FIELD OF THE INVENTION**

This invention refers to compounds capable of inhibiting neuronal exocytosis and 5 cosmetic or pharmaceutical compositions which contain these compounds useful in the treatment of those conditions, disorders and/or diseases which require the inhibition of neuronal exocytosis, such as muscle spasticity, pain, inflammation, perspiration, facial asymmetry and/or facial wrinkles, preferably expression wrinkles.

10 BACKGROUND OF THE INVENTION

Botulinum toxins (also known as botulinum neurotoxins) are neurotoxins produced by the gram-positive bacteria *Clostridium botulinum*. They act by causing the paralysis of the muscles through the inhibition of the release of acetylcholine in the presynaptic axon terminal of the neuromuscular junction (synaptic transmission), thus preventing 15 nerve transmission and muscle contraction. The paralyzing effects of the muscles of the botulinum toxin have been used both for therapeutic purposes as well as for cosmetic effects. The controlled administration of the botulinum toxin has been used for the treatment of a wide range of conditions, disorders and diseases, such as disorders and diseases of the urinary bladder (EP 2273976 A2), premature ejaculation (US 20 2011/052636 A1), priapism (US 6776991 B2), ulcers and gastroesophageal reflux (US 7238357 B2), disorders and diseases associated with hyper- and hypothyroidism (US 6740321 B2), primary hyperparathyroid disorders and diseases (US 6974793 B2), perspiration and hyperhidrosis (US 6974578 B2 and US 6683049 B2), inflammatory 25 eye disorders and diseases (US 7465458 B2 and US 7220422 B2), strabismus (US 6841156), otic disorders and diseases (US 6265379 B2 and US 6358926 B2), excess cerumen secretion (US 2010/028385), neuropsychiatric disorders and diseases such as Alzheimer's, anxiety, schizophrenia, mania, depression (US 7811587 B2), different compulsive disorders and diseases such as obsessions, compulsive skin picking, Tourette's syndrome, trichotillomania (US 7393537 B2), cerebral paralysis (US 30 6939852 B2), gonadotropin-related disorders and diseases (WO 02/074327), different cancers (US 6139845 B2, US7838007 B2), neoplasms (US 7709440 B2), different types of pain including headaches, migraines, fibromyalgia, arthritis or neuropathic pain

among others (US 2010/266638, US 7811586 B2, US 7704524 B2, US 7704511 B2, US 7468189 B2, US 7255866 B2, US 7091176 B2, US 6887476 B2, US 6869610 B2, US 6838434 B2, US 6641820 B2, US 6623742 B2, US 6565870 B1, US 6500436 B1, US 6458365 B1, US 6423319 B1, US 6113915 A and US 5714468 A), neurogenic inflammation (US 6063768 B2), different disorders and diseases of the autonomic nervous system such as otitis and sinusoidal disorders (US 5766605 A), disorders and diseases of the smooth muscle (US 5437291 A), nerve impingements (US 2003/0224019), epilepsy (US 7357934 B2), dystonia (US 6872397 B2), trembling (US 6861058 B2), Parkinson's disease (US 6620415 B2), dizziness (US 7270287 B2), osteoporosis (WO 2011/038015), different disorders and diseases of the skin such as calluses, warts, ulcers and lesions on the skin (US 8048423 B2, US 2011/206731), psoriasis and dermatitis (US 5670484 A), vascular hyperreactivity and rosacea (WO 2010/114828), acne (WO 03/011333), hair growth and maintenance (US 6299893 B1), facial wrinkles (US 7255865 B2), ptosis of the eyebrows and forehead (US 2011/280978) or drooping mouth corners (US 6358917 B1) among others.

However, the toxicity inherent in botulinum toxin causes its administration, in a wide range of doses, to result in undesired secondary effects, such as immunogenic responses, cephalgias, nausea, paralysis or muscle weakness, respiratory failure, and in more extreme cases even the death of the subject treated [FDA News, February 20 8, 2008, "FDA Notifies Public of Adverse Reactions Linked to Botox Use"; Coté, T.R. et al. "Botulinum toxin type A injections: Adverse events reported to the US Food and Drug Administration in therapeutic and cosmetic cases" *J. Amer. Acad. Derm.* 2005, 53 (3), 407-415]. These severe secondary effects, together with the high cost of the treatment, seriously limits the application of botulinum toxin with therapeutic or cosmetic purposes, being relegated to chronic applications and/or diseases for which there is no suitable treatment. There is, therefore, a pressing need to develop molecules which imitate the paralyzing effects of botulinum toxins but which are equipped with much simpler and more stable molecular structures that do not induce immune reactions, and whose cost of production is affordable. Molecules of a peptide nature comply with these properties.

At a molecular level, botulinum toxins are proteases which degrade neuronal proteins that are involved in the exocytosis mechanism activated by the calcium ion [Schiavo G. et al. "Bases Moleculares del tétanos y del botulismo" *Investigación y Ciencia* 1996, 234, 46-55; Montecucco C. and Schiavo G. "Mechanism of action of tetanus and

botulinum neurotoxins” Mol. Microbiol. 1994, 13, 1-8; Schiavo G. et al. “Tetanus and botulinum neurotoxins are zinc proteases specific for components of the neuroexocytosis apparatus” Ann. NY Acad. Sci. 1994, 710, 65-75]. For example, botulinum toxin A, the most commonly used in clinics to treat the symptomatology of 5 spasmodic diseases and in cosmetics due to its applications in the elimination of facial wrinkles and facial asymmetry, breaks down the neuronal protein SNAP-25. This protein SNAP-25 plays a key role in neurosecretion since it is involved in the formation of a protein complex (known by the name of SNARE or fusion complex) which manages and controls the release of acetylcholine accumulated in vesicles. The 10 nucleus of this fusion complex is comprised of the proteins SNAP-25 and syntaxin, located in the presynaptic plasma membrane, and the synaptobrevin protein of the VAMP family of proteins, located in the vesicular plasma membrane [Calakos N. and Scheller R.H. “Synaptic vesicle biogenesis, docking and fusion: a molecular description” Physiol. Rev. 1996, 76, 1-29; Sutton R.B. et al. “Crystal structure of a 15 SNARE complex involved in synaptic exocytosis at 2.4Å resolution” Nature 1998, 395, 347-353]. The principal function of the fusion complex is to bring the neurotransmitter (acetylcholine) loaded vesicle closer to and place it in contact with the presynaptic plasma membrane [Calakos N. and Scheller R.H. “Synaptic vesicle biogenesis, docking and fusion: a molecular description” Physiol. Rev. 1996, 76, 1-29; Sutton R.B. 20 et al. “Crystal structure of a SNARE complex involved in synaptic exocytosis at 2.4Å resolution” Nature 1998, 395, 347-353]. In this way, in response to an increase in the concentration of calcium, the fusion of both plasma membranes will be favored, thus producing the release of the neurotransmitter. Therefore, this vesicle docking and fusion protein SNARE complex constitutes a key target for controlling neurosecretion. 25 The truncation of any of the proteins which form the fusion complex prevents their assembling and, therefore, inhibits vesicle release and inhibits neuronal exocytosis.

It is known in the prior art that certain peptides derived from the protein sequences which form the SNARE complex are capable of inhibiting neuronal exocytosis, such as peptides derived from the amino and carboxy-terminal domains of the protein SNAP-25 30 [Apland J.P. et al. “Peptides that mimic the carboxy-terminal domain of SNAP-25 block acetylcholine release at an aplysia synapse” J. Appl. Toxicol. 1999, 19, Suppl.1: S23-S26; Mehta P.P. et al. “SNAP-25 and synaptotagmin involvement in the final Ca^{2+} -dependent triggering of neurotransmitter exocytosis” Proc. Natl. Acad. Sci. USA 1996, 93, 10471-10476; Ferrer-Montiel A.V. et al. “The 26-mer peptide released from 35 cleavage by botulinum neurotoxin E inhibits vesicle docking” FEBS Lett. 1998, 435, 84-

88; Gutierrez L.M. et al. "A peptide that mimics the carboxy-terminal domain of SNAP-25 blocks Ca^{2+} -dependent exocytosis in chromaffin cells" *FEBS Lett.* 1995, 372, 39-43; Gutierrez L.M. et al. "A peptide that mimics the C-terminal sequence of SNAP-25 inhibits secretory vesicle docking in chromaffin cells" *J. Biol. Chem.* 1997, 272, 2634-5 2639; Blanes-Mira C et al. "Small peptides patterned after the N-terminus domain of SNAP-25 inhibit SNARE complex assembly and regulated exocytosis" *J. Neurochem.* 2004, 88, 124-135], the peptides derived from the sequence of syntaxin amino acids [Martin F. et al. "Inhibition of insulin release by synthetic peptides show that the H3 region at the C-terminal domain of syntaxin-1 is crucial for Ca^{2+} -but not for guanosine 10 5'-[gammathio]thriphosphate-induced secretion" *Biochem. J.* 1996, 320, 201-205], of the synaptobrevin [Cornille F. "Inhibition of neurotransmitter release by synthetic prolinerich peptides shows that the N-terminal domain of vesicle-associated membrane protein/synaptobrevin is critical for neuro-exocytosis" *J. Biol. Chem.* 1995, 270, 16826-16830], of the synaptotagmin [Mehta P.P. et al. "SNAP-25 and synaptotagmin 15 involvement in the final Ca^{2+} -dependent triggering of neurotransmitter exocytosis" *Proc. Natl. Acad. Sci. USA* 1996, 93, 10471-10476] and of the protein snapin [Illardi J.M. et al. "Snapin: A SNARE associated protein implicated in synaptic transmission" *Nat. Neurosci.* 1999, 2, 119-124]. Similarly, synthetic peptides obtained by rational design or by searching synthetic libraries which are capable of inhibiting neuronal exocytosis by 20 interfering in the formation of the SNARE complex have also been described [Blanes-Mira C. et al. "Identification of SNARE complex modulators that inhibit exocytosis form an α -helixconstrained combinatorial library" *Biochem J.* 2003, 375, 159-166].

The industrial application of this type of compounds has been limited. The document EP 2318033 A2 describes the use of peptides derived from SNAP-25 for the treatment 25 of pain and inflammation, and the document EP 1856139 A2 describes the use of peptides derived from SNAP-25 chemically modified to increase their bioavailability for the treatment of different diseases for which the treatment with botulinum toxin has shown effectiveness, among them the treatment of hyperhidrosis. Similarly, the cosmetic industry has made significant efforts to develop compounds which imitate the 30 action of botulinum toxins with use in the treatment and prevention of the formation of expression wrinkles [Blanes-Mira C. et al. "A synthetic hexapeptide (Argireline[®]) with anti-wrinkle activity" *Int. J. Cosmetic Sci.* 2002, 24, 303-310]. In particular, peptides derived from the amino terminal fragment of the protein SNAP-25 which have anti-wrinkle effects are described in the documents EP 1180524 A1 and EP 2123673 A1, 35 international application WO 97/34620 also describes peptides derived from the

sequence of amino acids of the protein SNAP-25, in particular from its carboxy-terminal region, or from the synaptobrevin or the syntaxin capable of inhibiting neuronal exocytosis, and international application WO 2011/048443 describes peptides derived from the subunit c of the membrane component of V-ATPase capable of inhibiting neuronal exocytosis through its bonding to synaptobrevin and its potential application as anti-wrinkle treatment.

Thus, this invention provides an alternative to the existing needs and comprises the discovery of peptide sequences not derived from the protein SNAP-25 which are capable of inhibiting neuronal exocytosis.

10

DESCRIPTION OF THE INVENTION

This invention provides an alternative to the above-mentioned problem. Surprisingly, the authors of this invention have found that neuronal exocytosis can be inhibited by certain compounds not derived from the protein SNAP-25 and which are an alternative to the existing compounds in the prior art. These compounds are useful for the treatment and/or care of conditions, disorders and/or diseases which improve or are prevented by the inhibition of neuronal exocytosis.

Definitions

20 In order to facilitate the comprehension of this invention, the meanings of some terms and expressions as they are used in the context of the invention are included.

In the context of this invention "skin" is understood as the layers which comprise it, from the uppermost layer or stratum corneum to the lowermost layer or hypodermis, both inclusive. These layers are composed of different types of cells such as 25 keratinocytes, fibroblasts, melanocytes, mastocytes, neurones and/or adipocytes, among others. The term "skin" also comprises the scalp.

The term "treatment", as used in the context of this specification when it is not accompanied by the qualifications "cosmetic, non-therapeutic", means the administration of a compound according to the invention to alleviate or eliminate a 30 disease or disorder or reduce or eliminate one or more symptoms associated with this

disease or disorder. The term “treatment” also covers the ability to alleviate or eliminate the physiological consequences of the disease or disorder.

When the term “treatment” is accompanied by the qualifications “cosmetic, non-therapeutic” they refer to the application of the compound to the skin, hair and/or mucous membranes in particular with the aim of improving the cosmetic qualities of the skin, hair and/or mucous membranes such as and not restricted to, their level of hydration, elasticity, firmness, shine, tone or texture, among others. The term “care” in this invention refers to the maintenance of the qualities of the skin, hair and/or mucous membranes. These qualities are subject to improvement and maintained through a cosmetic treatment and/or care of the skin, hair and/or mucous membranes both in healthy subjects as well as those which present diseases and/or disorders of the skin and/or mucous membranes, such as and not restricted to, ulcers and lesions on the skin, psoriasis, dermatitis, acne or rosacea, among others.

The term “prevention”, as used in this invention, refers to the ability of a compound of the invention to prevent, delay or hinder the appearance or development of a disease or disorder before its appearance.

In the context of this invention, the term “aging” refers to the changes experienced by the skin with age (chronoaging) or through exposure to the sun (photoaging) or to extreme environmental climatic conditions of cold or wind, chemical contaminants or pollutants, and includes all the external visible and/or perceptible changes through touch, such as and not restricted to, the development of discontinuities on the skin such as wrinkles, fine lines, expression lines, stretch marks, furrows, irregularities or roughness, increase in the size of pores, loss of hydration, loss of elasticity, loss of firmness, loss of smoothness, loss of the capacity to recover from deformation, loss of resilience, sagging of the skin such as sagging cheeks, the appearance of bags under the eyes or the appearance of a double chin, among others, changes to the color of the skin such as marks, reddening, bags or the appearance of hyperpigmented areas such as age spots or freckles among others, anomalous differentiation, hyperkeratinization, elastosis, keratosis, hair loss, orange peel skin, loss of collagen structure and other histological changes of the stratum corneum, of the dermis, epidermis, vascular system (for example the appearance of spider veins or telangiectasias) or of those tissues close to the skin, among others. The term “photoaging” groups together the set of processes due to the prolonged exposure of the skin to ultraviolet radiation which result in the premature aging of the skin, and it presents the same physical characteristics as

aging, such as and not restricted to, flaccidity, sagging, changes to the color or irregularities in the pigmentation, abnormal and/or excessive keratinization. The sum of several environmental factors such as exposure to tobacco smoke, exposure to pollution, and climatic conditions such as cold and/or wind also contributes to the aging 5 of the skin.

In this description the abbreviations used for amino acids follow the recommendations of the 1983 IUPAC-IUB Commission of Biochemical Nomenclature specified in *Eur. J. Biochem.*, (1984), 138, 937.

Thus, for example, Phe represents $\text{NH}_2\text{-CH}(\text{CH}_2\text{-C}_5\text{H}_6)\text{-COOH}$, Phe- represents 10 $\text{NH}_2\text{-CH}(\text{CH}_2\text{-C}_5\text{H}_6)\text{-CO-}$, -Phe represents $-\text{NH}\text{-CH}(\text{CH}_2\text{-C}_5\text{H}_6)\text{-COOH}$ and -Phe- represents $-\text{NH}\text{-CH}(\text{CH}_2\text{-C}_5\text{H}_6)\text{-CO-}$. Therefore, the hyphen, which represents the peptide bond, eliminates the OH in the 1-carboxyl group of the amino acid (represented here in the conventional non-ionized form) when situated to the right of the symbol, and eliminates the H of the 2-amino group of the amino acid when 15 situated to the left of the symbol; both modifications can be applied to the same symbol (see Table 1).

Table 1. Structures of the amino acid residues and their nomenclature in one and three-letter code

Name	Residue	Symbol	Residue
Asparaginyl -Asn- N		Glutaminyl -Gln- Q	
Histidyl -His- H		Arginyl -Arg- R	

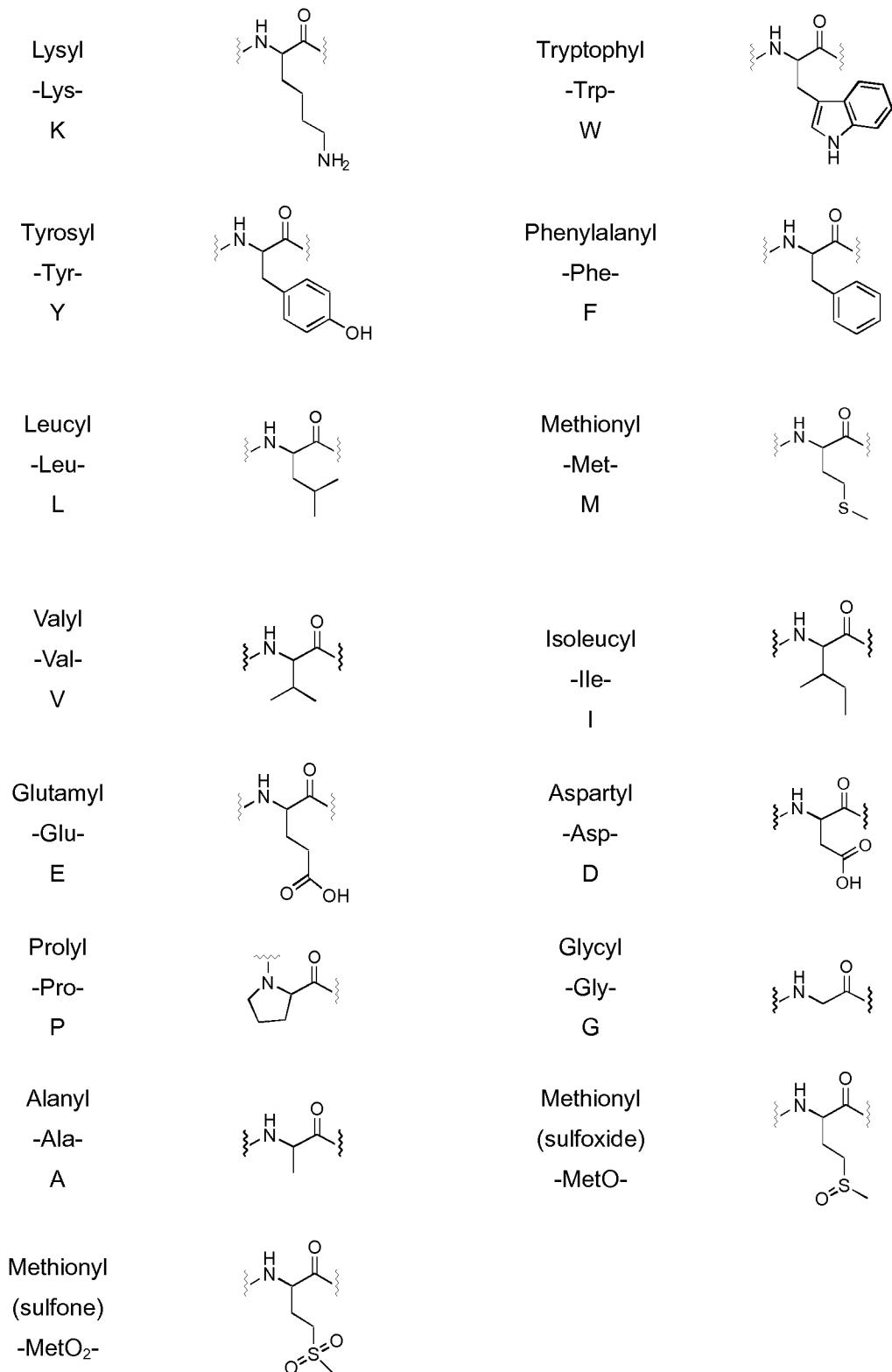


Table 1

The abbreviation “-MetO-“ is used in this invention to designate the amino acid residue methionyl(sulfoxide). The amino acid residue methionyl(sulfoxide) can be incorporated

into the compounds of the invention using the commercial amino acid methionine(sulfoxide) or can be obtained *in situ* in the compounds of the invention by oxidation of the methionyl residue.

The abbreviation “-MetO₂” is used in this invention to designate the amino acid residue 5 methionyl(sulfone). The amino acid residue methionyl(sulfone) can be incorporated into the compounds of the invention using the commercial amino acid methionine(sulfone) or can be obtained *in situ* in the compounds of the invention by oxidation of the methionyl residue or the methionyl(sulfoxide) residue.

The abbreviation “Ac-” is used in this description to designate the acetyl group 10 (CH₃-CO-), the abbreviation “Palm-” is used to designate the palmitoyl group (CH₃-(CH₂)₁₄-CO-) and the abbreviation “Myr-” is used to designate the myristoyl group (CH₃-(CH₂)₁₂-CO-).

The term “non-cyclic aliphatic group” is used in this invention to cover alkyl, alkenyl and alkynyl groups, linear or branched.

15 The term “alkyl group” refers to a linear or branched saturated group which has between 1 and 24, preferably between 1 and 16, more preferably between 1 and 14, even more preferably between 1 and 12, yet more preferably 1, 2, 3, 4, 5 or 6 carbon atoms and is bound to the rest of the molecule by a simple bond, including, for example and not restricted to, methyl, ethyl, isopropyl, isobutyl, *tert*-butyl, heptyl, octyl, decyl, 20 dodecyl, lauryl, hexadecyl, octadecyl, amyl, 2-ethylhexyl, 2-methylbutyl, 5-methylhexyl and similar.

The term “alkenyl group” refers to a group, linear or branched, which has between 2 and 24, preferably between 2 and 16, more preferably between 2 and 14, even more preferably between 2 and 12, yet more preferably 2, 3, 4, 5 or 6 carbon atoms, with one 25 or more double carbon-carbon bonds, preferably with 1, 2 or 3 double carbon-carbon bonds, conjugated or unconjugated, which is bound to the rest of the molecule by a simple bond, including, for example and not restricted to, the vinyl group (-CH₂=CH₂), allyl (-CH₂-CH=CH₂), oleyl, linoleyl and similar.

The term “alkynyl group” refers to a group, linear or branched, which has between 2 30 and 24, preferably between 2 and 16, more preferably between 2 and 14, even more preferably between 2 and 12, yet more preferably 2, 3, 4, 5 or 6 carbon atoms, with one or more triple carbon-carbon bonds, preferably 1, 2 or 3 triple carbon-carbon bonds,

conjugated or unconjugated, which is bound to the rest of the molecule by a simple bond, including, for example and not restricted to, the ethynyl group, 1-propynyl, 2-propynyl, 1-butynyl, 2-butynyl, 3-butynyl, pentynyl, such as 1-pentynyl, and similar. The alkynyl groups can also contain one or more double carbon-carbon bonds,

5 including for example and not restricted to, the group but-1-en-3-ynyl, pent-4-en-1-ynyl and similar.

The term “alycyclic group” is used in this invention to cover, for example and not restricted to, cycloalkyl or cycloalkenyl or cycloalkynyl groups.

The term “cycloalkyl” refers to a saturated mono- or polycyclic aliphatic group which

10 has between 3 and 24, preferably between 3 and 16, more preferably between 3 and 14, even more preferably between 3 and 12, yet more preferably 3, 4, 5 or 6 carbon atoms and is bound to the rest of the molecule by a simple bond, including, for example and not restricted to, cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, methyl cyclohexyl, dimethyl cyclohexyl, octahydroindene, decahydronaphthalene,

15 dodecahydrophenalene and similar.

The term “cycloalkenyl” refers to a non-aromatic mono- or polycyclic aliphatic group which has between 5 and 24, preferably between 5 and 16, more preferably between 5 and 14, even more preferably between 5 and 12, yet more preferably 5 or 6 carbon atoms, with one or more double carbon-carbon bonds, preferably 1, 2 or 3 double

20 carbon-carbon bonds, conjugated or unconjugated, bound to the rest of the molecule by a simple bond, including, for example and not restricted to, the cyclopent-1-en-1-yl group and similar.

The term “cycloalkynyl” refers to a non-aromatic mono- or polycyclic aliphatic group which has between 8 and 24, preferably between 8 and 16, more preferably between 8 and 14, even more preferably between 8 and 12, yet more preferably 8 or 9 carbon atoms, with one or more triple carbon-carbon bonds, preferably 1, 2 or 3 triple carbon-carbon bonds, conjugated or unconjugated, bound to the rest of the molecule by a simple bond, including, for example and not restricted to, the cyclooct-2-en-1-yl group and similar. The cycloalkynyl groups can also contain one or more double carbon-

30 carbon bonds, including for example and not restricted to, the cyclooct-4-en-2-ynyl group and similar.

The term “aryl group” refers to an aromatic group which has between 6 and 30, preferably between 6 and 18, more preferably between 6 and 10, even more preferably

between 6 or 10 carbon atoms, which comprises 1, 2, 3 or 4 aromatic rings, bound by a carbon-carbon bond or condensed, including, for example and not restricted to, phenyl, naphthyl, diphenyl, indenyl, phenanthryl or anthranyl, among others; or an aralkyl group.

- 5 The term “aralkyl group” refers to an alkyl group substituted by an aromatic group, with between 7 and 24 carbon atoms and including, for example and not restricted to, -(CH₂)₁₋₆-phenyl, -(CH₂)₁₋₆-(1-naphthyl), -(CH₂)₁₋₆-(2-naphthyl), -(CH₂)₁₋₆-CH(phenyl)₂ and similar.

The term “heterocyclyl group” refers to a hydrocarbonated ring of 3-10 members, in 10 which one or more of the atoms in the ring, preferably 1, 2 or 3 of the atoms in the ring, is a different element to carbon, such as nitrogen, oxygen or sulfur and can be saturated or unsaturated. For the purposes of this invention, the heterocycle can be a monocyclic, bicyclic or tricyclic system, which can include systems of condensed rings; and the nitrogen, carbon or sulfur atoms can optionally be oxidized in the radical 15 heterocycle; the nitrogen atom can optionally be quaternized; and the radical heterocyclyl can be partially or completely saturated or aromatic. The greatest preference is for the term heterocyclyl to refer to a ring of 5 or 6 members. Examples of saturated heterocyclic groups are dioxane, piperidine, piperazine, pyrrolidine, morpholine and thiomorpholine. Examples of aromatic heterocyclic groups, also known 20 as heteroaromatic groups are pyridine, pyrrol, furane, thiophene, benzofuran, imidazolin, quinolein, quinolina, pyridazin and naphthyridine.

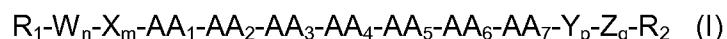
The term “heteroarylalkyl group” refers to an alkyl group substituted by a substituted or 25 unsubstituted aromatic heterocyclyl group, the alkyl group having from 1 to 6 carbon atoms and the aromatic heterocyclyl group between 2 and 24 carbon atoms and from 1 to 3 atoms other than carbon and including, for example and not restricted to, -(CH₂)₁₋₆-imidazolyl, -(CH₂)₁₋₆-triazolyl, -(CH₂)₁₋₆-thienyl, -(CH₂)₁₋₆-furyl, -(CH₂)₁₋₆-pyrrolidinyl and similar.

As is understood in this technical field, there can be a certain level of substitution of the 30 aforementioned groups. Therefore, there can be substitution in any of the groups of this invention where specifically stated. The references in this document to substituted groups in the groups of this invention indicate that the specified radical can be substituted in one or more positions available by one or more substituents, preferably in 1, 2 or 3 positions, more preferably in 1 or 2 positions, yet more preferably in 1

position. These substituents include, alkyl C₁-C₄; hydroxyl; alcoxyl C₁-C₄; amino; aminoalkyl C₁-C₄; carbonyloxy C₁-C₄; oxycarbonyl C₁-C₄; halogen such as fluoride, chlorine, bromine and iodine; cyano; nitro; azide; alkylsulfonyl C₁-C₄; thiol; alkylthio C₁-C₄; aryloxy such as phenoxy; -NR_b(C=NR_b)NR_bR_c; wherein R_b and R_c are 5 independently selected from the group formed by H, alkyl C₁-C₄, alkenyl C₂-C₄, alkynyl C₂-C₄, cycloalkyl C₃-C₁₀, aryl C₆-C₁₈, aralkyl C₇-C₁₇, heterocyclyl of 3-10 members or protective group of the amino group.

Compounds in the invention

10 A first aspect of the invention refers to a compound of general formula (I):



its stereoisomers, mixtures thereof and/or its cosmetically or pharmaceutically acceptable salts, characterized in that:

15 AA₁ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₂ is selected from the group formed by -His-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₃ is selected from the group formed by -Leu-, -Ile-, -Phe- and -Tyr-;

AA₄ is selected from the group formed by -Lys-, -His-, -Arg-, -Gln-, -Leu-, -Ile-, -Met-, -MetO- and -MetO₂-;

20 AA₅ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₆ is selected from the group formed by -Phe-, -Trp-, -Ile- and -Val-;

AA₇ is selected from the group formed by -Met-, -MetO- and -MetO₂-;

25 W, X, Y, Z are amino acids and are independently selected from amongst themselves;

n, m, p and q are independently selected from amongst themselves and have a value of 0 or 1;

n+m+p+q is smaller or equal to 2;

with the condition that it is not Trp-Lys-Lys-His-Leu-Leu-Lys-Ile-Met (SEQ ID NO:111);

R₁ is selected from the group formed by H, a polymer derived from polyethylene glycol, a non-cyclic substituted or unsubstituted aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl and R₅-CO-, wherein R₅ is selected from the group formed by H, a non-cyclic substituted or unsubstituted aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl, substituted or unsubstituted heterocyclyl and substituted or unsubstituted heteroarylalkyl;

R₂ is selected from the group formed by -NR₃R₄, -OR₃ and -SR₃, wherein R₃ and R₄ are independently selected from the group formed by H, a polymer derived from polyethylene glycol, a non-cyclic substituted or unsubstituted aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, and substituted or unsubstituted aralkyl; and

R₁ and R₂ are not α -amino acids.

Groups R₁ and R₂ are bound to the amino-terminal (N-terminal) and carboxy-terminal (C-terminal) ends of the peptide sequences respectively.

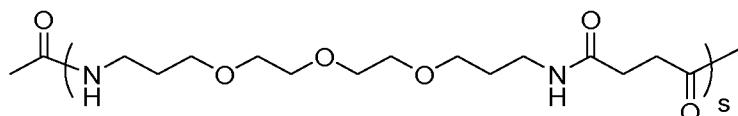
In accordance with a preferred embodiment of this invention R₁ is selected from the group formed by H, a polymer derived from polyethylene glycol and R₅-CO-, wherein R₅ is selected from the group formed by substituted or unsubstituted alkyl radical C₁-C₂₄, substituted or unsubstituted alkenyl C₂-C₂₄, substituted or unsubstituted alkynyl C₂-C₂₄, substituted or unsubstituted cycloalkyl C₃-C₂₄, substituted or unsubstituted cycloalkenyl C₅-C₂₄, substituted or unsubstituted cycloalkynyl C₈-C₂₄, substituted or unsubstituted aryl C₆-C₃₀, substituted or unsubstituted aralkyl C₇-C₂₄, substituted or unsubstituted heterocyclyl ring of 3-10 members, and substituted or unsubstituted heteroarylalkyl of 2 to 24 carbon atoms and 1 to 3 atoms other than carbon and an alkyl chain of 1 to 6 carbon atoms and R₅-CO- is not an α -amino acid. More preferably, R₁ is selected from the group formed by H, a polymer derived from polyethylene glycol with a molecular weight comprised between 200 and 35000 Daltons, acetyl, *tert*-butanoyl, prenyl, hexanoyl, 2-methylhexanoyl, cyclohexanecarboxyl, octanoyl,

decanoyl, lauroyl, myristoyl, palmitoyl, stearoyl, oleoyl and linoleoyl. Even more preferably, R₁ is H, acetyl, lauroyl, myristoyl or palmitoyl. In an even more preferred embodiment, R₁ is acetyl or palmitoyl.

In accordance with another preferred embodiment, R₂ is selected from the group formed by -NR₃R₄, -OR₃, -SR₃, wherein R₃ and R₄ are independently selected from the group formed by H, a polymer derived from polyethylene glycol, substituted or unsubstituted alkyl C₁-C₂₄, substituted or unsubstituted alkenyl C₂-C₂₄, substituted or unsubstituted alkynyl C₂-C₂₄, substituted or unsubstituted cycloalkyl C₃-C₂₄, substituted or unsubstituted cycloalkenyl C₅-C₂₄, substituted or unsubstituted cycloalkynyl C₈-C₂₄, substituted or unsubstituted aryl C₆-C₃₀, substituted or unsubstituted aralkyl C₇-C₂₄, substituted or unsubstituted heterocyclyl ring of 3-10 members, and substituted or unsubstituted heteroarylalkyl of 2 to 24 carbon atoms and 1 to 3 atoms other than carbon wherein the alkyl chain is of 1 to 6 carbon atoms and -NR₃R₄ is not an α-amino acid. Optionally, R₃ and R₄ can be bound by a saturated or unsaturated carbon-carbon bond, forming a cycle with the nitrogen atom. More preferably R₂ is -NR₃R₄ or -OR₃. More preferably, R₃ and R₄ are independently selected from the group formed by H, a polymer derived from polyethylene glycol with a molecular weight comprised between 200 and 35000 Daltons, methyl, ethyl, hexyl, dodecyl, or hexadecyl. Even more preferably R₃ and R₄ are independently selected from the group formed by H, methyl, ethyl, hexyl, dodecyl, or hexadecyl. Even more preferably R₃ is H and R₄ is selected from the group formed by H, methyl, ethyl, hexyl, dodecyl, and hexadecyl. In accordance with an even more preferred embodiment, R₂ is selected from -OH and -NH₂.

In accordance with another embodiment of this invention R₁ is selected from the group formed by H, acetyl, lauroyl, myristoyl or palmitoyl, preferably R₁ is selected from the group formed by H, acetyl and palmitoyl and R₂ is selected from the group formed by -OH and -NH₂.

In accordance with another particular embodiment the most preferred structures of the polymer derived from polyethylene glycol are the group (-CH₂-CH₂-O)_r-H in which r is a number comprised between 4 and 795 and the group



where s is a number comprised between 1 and 125.

In accordance with another embodiment of this invention n, m, p and q are 0.

- In accordance with another embodiment of this invention AA₁ is selected from the group formed by -Arg-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-, AA₂ is selected from the group formed by -His-, -Gln-, -Asn-, -Glu- and -Asp-, AA₃ is selected from the group formed by -Leu- and -Phe-, AA₄ is selected from the group formed by -His-, -Lys-, -Gln- and -Leu-, AA₅ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-, AA₆ is selected from the group formed by -Trp- and -Val- and AA₇ is selected from the group formed by -Met-, -MetO- and -MetO₂. More preferably, AA₄ is selected from the group formed by -Lys- and -Leu-, and AA₆ is -Trp-. Even more preferably, n, m, p and q are 0. Additionally, the inventors of the present invention have found that when AA₂ is selected from the group formed by -Asn- and -Glu-, the obtained compound is more chemically stable compared with other aminoacids in AA₂.
- 15 In accordance with another embodiment of this invention R₁ is selected from the group formed by H, acetyl, lauroyl, myristoyl and palmitoyl, AA₁ is selected from the group formed by -L-Arg-, -L-Lys-, -L-Gln-, -L-Asn-, -L-Glu- and -L-Asp-, AA₂ is selected from the group formed by -L-His-, -L-Gln-, -L-Asn-, -L-Glu- and -L-Asp-, AA₃ is selected from the group formed by -L-Leu- and -L-Phe-, AA₄ is selected from the group formed by -L-His-, -L-Lys-, -L-Gln- and -L-Leu-, AA₅ is selected from the group formed by -L-Arg-, -L-His-, -L-Lys-, -L-Gln-, -L-Asn-, -L-Glu- and -L-Asp-, AA₆ is selected from the group formed by -L-Trp- and -L-Val-, AA₇ is selected from the group formed by -L-Met-, -L-MetO- and -L-MetO₂, and R₂ is selected from the group formed by -NR₃R₄ and -OR₃ where R₃ and R₄ are independently selected from H, methyl, ethyl, 20 hexyl, dodecyl and hexadecyl. More preferably, AA₄ is selected from the group formed by -L-Lys- and -L-Leu-, and AA₆ is -L-Trp-. More preferably, R₁ is acetyl or palmitoyl and R₂ is -NH₂. Even more preferably, n, m, p and q are 0.
- 25 In accordance with another embodiment of this invention R₁ is selected from the group formed by H, acetyl, lauroyl, myristoyl and palmitoyl, AA₁ is selected from the group formed by -L-Asn-, -L-Glu- and -L-Asp-, AA₂ is selected from the group formed by -L-His- and -L-Asp-, AA₃ is selected from the group formed by -L-Leu- and -L-Phe-, AA₄ is selected from the group formed by -L-Lys-, -L-Gln- and -L-Leu-, AA₅ is selected from the group formed by -L-Arg-, -L-Gln-, -L-Asn- and -L-Asp-, AA₆ is selected from

the group formed by -L-Trp-, AA₇ is selected from the group formed by -L-Met-, -L-MetO- and -L-MetO₂-, and R₂ is selected from the group formed by -NR₃R₄ and -OR₃ where R₃ and R₄ are independently selected from H, methyl, ethyl, hexyl, dodecyl and hexadecyl. More preferably, R₁ is acetyl or palmitoyl and R₂ is -NH₂.

- 5 Even more preferably, n, m, p and q are 0.

In accordance with another embodiment of this invention R₁ is selected from the group formed by H, acetyl, lauroyl, myristoyl and palmitoyl, AA₁ is -L-Glu-, AA₂ is selected from the group formed by -L-Asn-, -L-Glu-, -L-Gln- and -L-Asp-, AA₃ is -L-Leu-, AA₄ is -L-Lys-, AA₅ is -L-Arg-, AA₆ is -L-Trp-, AA₇ is selected from the group formed

- 10 by -L-Met-, -L-MetO- and -L-MetO₂-, and R₂ is selected from the group formed by -NR₃R₄ and -OR₃ where R₃ and R₄ are independently selected from H, methyl, ethyl, hexyl, dodecyl and hexadecyl. More preferably, R₁ is acetyl or palmitoyl and R₂ is -NH₂. Even more preferably, n, m, p and q are 0.

In accordance with another embodiment of this invention R₁ is selected from the group

- 15 formed by H, acetyl, lauroyl, myristoyl or palmitoyl, AA₁ is -L-Glu-, AA₂ is -L-His-, AA₃ is -L-Leu-, AA₄ is -L-Lys-, AA₅ is -L-Gln-, AA₆ is -L-Trp-, AA₇ is -L-Met-, -L-MetO- or -L-MetO₂- and R₂ is -NR₃R₄ or -OR₃ where R₃ and R₄ are independently selected from H, methyl, ethyl, hexyl, dodecyl and hexadecyl, preferably R₂ is -OH or -NH₂. More preferably, R₁ is acetyl or palmitoyl and R₂ is -NH₂. Even more preferably, n, m, p and q are 0.

In accordance with another embodiment of this invention R₁ is selected from the group formed by H, acetyl, lauroyl, myristoyl or palmitoyl, AA₁ is -L-Glu-, AA₂ is -L-Asp-, AA₃ is -L-Leu-, AA₄ is -L-Lys-, AA₅ is -L-Arg-, AA₆ is -L-Trp-, AA₇ is -L-Met-, -L-MetO- or -L-MetO₂- and R₂ is -NR₃R₄ or -OR₃ where R₃ and R₄ are

- 25 independently selected from H, methyl, ethyl, hexyl, dodecyl and hexadecyl, preferably R₂ is -OH or -NH₂. More preferably, R₁ is acetyl or palmitoyl and R₂ is -NH₂. Even more preferably, n, m, p and q are 0.

In accordance with another embodiment of this invention R₁ is selected from the group

formed by H, acetyl, lauroyl, myristoyl or palmitoyl, AA₁ is -L-Asn-, AA₂ is -L-His-, AA₃ is -L-Phe-, AA₄ is -L-Leu-, AA₅ is -L-Asn-, AA₆ is -L-Trp-, AA₇ is -L-Met-, -L-MetO- or -L-MetO₂- and R₂ is -NR₃R₄ or -OR₃ where R₃ and R₄ are

- 30 independently selected from H, methyl, ethyl, hexyl, dodecyl and hexadecyl, preferably R₂ is -OH or -NH₂. More preferably, R₁ is acetyl or palmitoyl and R₂ is -NH₂.

R_2 is $-OH$ or $-NH_2$. More preferably, R_1 is acetyl or palmitoyl and R_2 is $-NH_2$. Even more preferably, n , m , p and q are 0.

In accordance with another embodiment of this invention R_1 is selected from the group formed by H, acetyl, lauroyl, myristoyl or palmitoyl, AA_1 is $-L\text{-Asp-}$, AA_2 is $-L\text{-His-}$, AA_3 is $-L\text{-Phe-}$, AA_4 is $-L\text{-Leu-}$, AA_5 is $-L\text{-Asp-}$, AA_6 is $-L\text{-Trp-}$, AA_7 is $-L\text{-Met-}$, $-L\text{-MetO-}$ or $-L\text{-MetO}_2$ and R_2 is $-NR_3R_4$ or $-OR_3$ where R_3 and R_4 are independently selected from H, methyl, ethyl, hexyl, dodecyl and hexadecyl, preferably R_2 is $-OH$ or $-NH_2$. More preferably, R_1 is acetyl or palmitoyl and R_2 is $-NH_2$. Even more preferably, n , m , p and q are 0.

10 In particular, the compounds in the invention which inhibit neuronal exocytosis, represented according to the formula (I) are selected from the group of peptide sequences outlined in Table 2, in which their sequence identifier is detailed:

SEQUENCE	IDENTIFIER
EHLKQWM	SEQ ID NO:1
QDFLHWM	SEQ ID NO:2
QHLHNVM	SEQ ID NO:3
KDIKNWM	SEQ ID NO:4
DHLKQFM	SEQ ID NO:5
EHMKQYM	SEQ ID NO:6
KDLKEWM	SEQ ID NO:7
EHLKKWM	SEQ ID NO:8
DHLKEWM	SEQ ID NO:9
KDLRRWM	SEQ ID NO:10
DDLKRYM	SEQ ID NO:11
QHLKEWM	SEQ ID NO:12
RHFLQWM	SEQ ID NO:13
EDLKRWM	SEQ ID NO:14
NHFLNWM	SEQ ID NO:15
RHLKDWM	SEQ ID NO:16
DHFNQRM	SEQ ID NO:17

EDFQQHM	SEQ ID NO:18
EHFLEFM	SEQ ID NO:19
KHLQQIM	SEQ ID NO:20
NHFLKWM	SEQ ID NO:21
DHLKKWM	SEQ ID NO:22
NDLKDWM	SEQ ID NO:23
DHFQQRM	SEQ ID NO:24
DHFLDWM	SEQ ID NO:25
DHFLQWM	SEQ ID NO:26
QHLKNWM	SEQ ID NO:27
RDLIRKM	SEQ ID NO:28
NHFNNKM	SEQ ID NO:29
EHFLQWM	SEQ ID NO:30
NHLKNWM	SEQ ID NO:31
NHLKHWM	SEQ ID NO:32
QHFLRWM	SEQ ID NO:33
DHFLNWM	SEQ ID NO:34
EHLRQWM	SEQ ID NO:35
EHIKQWM	SEQ ID NO:36
EHMRQVM	SEQ ID NO:37
EHIMHWM	SEQ ID NO:38
DHMNRVM	SEQ ID NO:39
DDMKRWM	SEQ ID NO:40
NHYLNWM	SEQ ID NO:41
QHFINRM	SEQ ID NO:42
NHFVNYM	SEQ ID NO:43
QHFLNFM	SEQ ID NO:44
EHYQQRM	SEQ ID NO:45
RHLVQMM	SEQ ID NO:46
HHLIQMM	SEQ ID NO:47

HDIVRHM	SEQ ID NO:48
KHMMQIM	SEQ ID NO:49
Glu-His-Leu-Lys-Gln-Trp-MetO	SEQ ID NO:50
Gln-Asp-Phe-Leu-His-Trp-MetO	SEQ ID NO:51
Gln-His-Leu-His-Asn-Val-MetO	SEQ ID NO:52
Glu-Asp-Leu-Lys-Arg-Trp-MetO	SEQ ID NO:53
Asn-His-Phe-Leu-Asn-Trp-MetO	SEQ ID NO:54
Asp-His-Phe-Gln-Gln-Arg-MetO	SEQ ID NO:55
Asp-His-Phe-Leu-Asp-Trp-MetO	SEQ ID NO:56
Asp-His-Phe-Leu-Asn-Trp-MetO	SEQ ID NO:57
Glu-His-Leu-Lys-Gln-Trp-MetO ₂	SEQ ID NO:58
Glu-Asp-Leu-Lys-Arg-Trp-MetO ₂	SEQ ID NO:59
Asp-His-Phe-Gln-Gln-Arg-MetO ₂	SEQ ID NO:60
Asp-His-Phe-Leu-Asp-Trp-MetO ₂	SEQ ID NO:61
EEHLKQWM	SEQ ID NO:62
EHLKQWMR	SEQ ID NO:63
RQDFLHWM	SEQ ID NO:64
QDFLHWMH	SEQ ID NO:65
DQHLHNVM	SEQ ID NO:66
QHLHNVMK	SEQ ID NO:67
EQHLKEWM	SEQ ID NO:68
QHLKEWMR	SEQ ID NO:69
DEDLKRW	SEQ ID NO:70
EDLKRW	SEQ ID NO:71
ENHFLNWM	SEQ ID NO:72
NHFLNWMH	SEQ ID NO:73
DDHLKKWM	SEQ ID NO:74
DHLKKWMR	SEQ ID NO:75
DDHFQQRM	SEQ ID NO:76

DHFQQRMR	SEQ ID NO:77
EDHFLDWM	SEQ ID NO:78
DHFLDWMK	SEQ ID NO:79
ENHLKNWM	SEQ ID NO:80
NHLKNWMH	SEQ ID NO:81
DDHFLNWM	SEQ ID NO:82
DHFLNWMR	SEQ ID NO:83
Glu-His-Leu-Lys-Gln-Trp-MetO-Arg	SEQ ID NO:84
Asp-Glu-Asp-Leu-Lys-Arg-Trp-MetO	SEQ ID NO:85
Asn-His-Phe-Leu-Asn-Trp-MetO-Arg	SEQ ID NO:86
Asp-His-Phe-Gln-Gln-Arg-MetO-Arg	SEQ ID NO:87
Asp-His-Phe-Leu-Asp-Trp-MetO-Lys	SEQ ID NO:88
Glu-His-Leu-Lys-Gln-Trp-MetO ₂ -Arg	SEQ ID NO:89
Asp-Glu-Asp-Leu-Lys-Arg-Trp-MetO ₂	SEQ ID NO:90
Asp-His-Phe-Gln-Gln-Arg-MetO ₂ -Arg	SEQ ID NO:91
Asp-His-Phe-Leu-Asp-Trp-MetO ₂ -Lys	SEQ ID NO:92
EEHLKQWMR	SEQ ID NO:93
EHLKQWMRR	SEQ ID NO:94
DEQHLHNVM	SEQ ID NO:95
QHLHNVMRR	SEQ ID NO:96
EEDLKRWMM	SEQ ID NO:97
DEEDLKRWMM	SEQ ID NO:98
QRNHFLNWM	SEQ ID NO:99
NHFLNWMMR	SEQ ID NO:100
EDHFQQRML	SEQ ID NO:101
DDHFQQRMR	SEQ ID NO:102
DHFLDWMRR	SEQ ID NO:103
DHDHFLDWM	SEQ ID NO:104
EHFLQWMRM	SEQ ID NO:105

DEHFLQWMV	SEQ ID NO:106
Glu-His-Leu-Lys-Gln-Trp-MetO-Arg-Lys	SEQ ID NO:107
Gln-Arg-Asn-His-Phe-Leu-Asn-Trp-MetO	SEQ ID NO:108
Glu-His-Leu-Lys-Gln-Trp-MetO ₂ -Arg-Lys	SEQ ID NO:109
Gln-Arg-Asn-His-Phe-Leu-Asn-Trp-MetO ₂	SEQ ID NO:110
EDVKRWM	SEQ ID NO:112
EQLKRWM	SEQ ID NO:113
DDVKKFM	SEQ ID NO:114
DNLKRFM	SEQ ID NO:115
EELKRWM	SEQ ID NO:116
ENLKRWM	SEQ ID NO:117
EDVRRWM	SEQ ID NO:118
EHFLEWM	SEQ ID NO:119
DEIHKWM	SEQ ID NO:120
ENLRRWM	SEQ ID NO:121
DNLHKYM	SEQ ID NO:122
EQIKHFM	SEQ ID NO:123
DNMRRFM	SEQ ID NO:124
EEMKRWM	SEQ ID NO:125
DQMKHYM	SEQ ID NO:126
Glu-Glu-Leu-Lys-Arg-Trp-MetO	SEQ ID NO:127
Asp-Asp-Val-His-Arg-Trp-MetO	SEQ ID NO:128
Glu-Asn-Leu-Lys-Arg-Trp-MetO	SEQ ID NO:129
Asp-Glu-Val-Arg-His-Tyr-MetO	SEQ ID NO:130
Glu-Glu-Leu-Lys-Arg-Trp-MetO ₂	SEQ ID NO:131
Glu-Asn-Leu-Lys-Arg-Trp-MetO ₂	SEQ ID NO:132
Asp-Gln-Ile-Arg-Lys-Phe-MetO ₂	SEQ ID NO:133
EEQLKRWM	SEQ ID NO:134

EQLKRWMR	SEQ ID NO:135
REELKRW	SEQ ID NO:136
EELKRW	SEQ ID NO:137
DENLKRWM	SEQ ID NO:138
ENLKRWMK	SEQ ID NO:139
EDNLKRFM	SEQ ID NO:140
DNLKRFMR	SEQ ID NO:141
EEQIKHFM	SEQ ID NO:142
EQIKHFMH	SEQ ID NO:143
Glu-Glu-Leu-Lys-Arg-Trp-MetO-Arg	SEQ ID NO:144
Glu-Asn-Leu-Lys-Arg-Trp-MetO-Arg	SEQ ID NO:145
Glu-Glu-Leu-Lys-Arg-Trp-MetO ₂ -Arg	SEQ ID NO:146
Glu-Asn-Leu-Lys-Arg-Trp-MetO ₂ -Arg	SEQ ID NO:147
EEQLKRW	SEQ ID NO:148
EQLKRW	SEQ ID NO:149
DEELKRW	SEQ ID NO:150
EELKRW	SEQ ID NO:151
QRENLKRWM	SEQ ID NO:152
ENLKRWM	SEQ ID NO:153
QREQIKHFM	SEQ ID NO:154
EQIKHFMMR	SEQ ID NO:155
Glu-Glu-Leu-Lys-Arg-Trp-MetO-Arg-Lys	SEQ ID NO:156
Gln-Arg-Glu-Asn-Leu-Lys-Arg-Trp-MetO	SEQ ID NO:157
Glu-Asn-Leu-Lys-Arg-Trp-MetO ₂ -Arg-Lys	SEQ ID NO:158
Gln-Arg-Glu-Glu-Leu-Lys-Arg-Trp-MetO ₂	SEQ ID NO:159

Table 2

their stereoisomers, mixtures thereof and/or their cosmetically or pharmaceutically acceptable salts.

- The compounds of this invention can exist as stereoisomers or mixtures of stereoisomers; for example, the amino acids which comprise them can have the configuration L-, D-, or be racemic independently of each other. Therefore, it is possible to obtain isomeric mixtures as well as racemic mixtures or diastereomeric mixtures, or pure diastereomers or enantiomers, depending on the number of asymmetric carbons and on which isomers or isomeric mixtures are present. The preferred structures of the compounds of the invention are pure isomers, i.e., enantiomers or diastereomers.
- 10 For example, when it is stated that AA₁ can be -Lys-, it is understood that AA₁ is selected from -L-Lys-, -D-Lys- or mixtures of both, racemic or non-racemic. The preparation procedures described in this document enable the person skilled in the art to obtain each of the stereoisomers of the compound of the invention by choosing the amino acid with the right configuration.
- 15 In the context of this invention, the term "amino acids" includes the amino acids encoded by the genetic code as well as non-encoded amino acids, whether they are natural or not. Examples of non-encoded amino acids are, without restriction, citrulline, ornithine, sarcosine, desmosine, norvaline, 4-aminobutyric acid, 2-aminobutyric acid, 2-aminoisobutyric acid, 6-aminohexanoic acid, 1-naphthylalanine, 2-naphthylalanine,
- 20 2-aminobenzoic acid, 4-aminobenzoic acid, 4-chlorophenylalanine, 2,3-diaminopropionic acid, 2,4 diaminobutyric acid, cycloserine, carnitine, cystine, penicillamine, pyroglutamic acid, thienylalanine, hydroxyproline, allo-isoleucine, allo-threonine, isonipecotic acid, isoserine, phenylglycine, statin, β -alanine, norleucine, N-methyl amino acids, α -amino acids and β -amino acids, among others, as well as
- 25 their derivatives. A list of non-natural amino acids can be found in the article "*Unusual amino acids in peptide synthesis*" by D.C. Roberts and F. Vellaccio, in *The Peptides*, Vol. 5 (1983), Chapter VI, Gross E. and Meienhofer J., Eds., Academic Press, New York, USA or in the commercial catalogs of the companies specialized in the field.
- In the context of this invention, when n, m, p or q are not 0 it is clearly understood that
- 30 the nature of W, X, Y and/or Z does not hinder the activity of the compounds of the invention, but that it contributes to the inhibition of neuronal exocytosis or has no effect on it.

The cosmetically and pharmaceutically acceptable salts of the peptides provided by this invention are also found within the field of this invention. The term "cosmetically or pharmaceutically acceptable salts" means a salt recognized for its use in animals and more specifically in human beings, and includes salts used to form base addition salts, 5 either they are inorganic, such as and not restricted to, lithium, sodium, potassium, calcium, magnesium, manganese, copper, zinc or aluminum among others, either they are organic, such as and not restricted to, ethylamine, diethylamine, ethylenediamine, ethanolamine, diethanolamine, arginine, lysine, histidine or piperazine among others, or acid addition salts, either they are organic, such as and not restricted to, acetate, 10 citrate, lactate, malonate, maleate, tartrate, fumarate, benzoate, aspartate, glutamate, succinate, oleate, trifluoroacetate, oxalate, pamoate or gluconate among others, or inorganic, such as and not restricted to, chloride, sulfate, borate or carbonate, among others. The nature of the salt is not critical, provided that it is cosmetically or pharmaceutically acceptable. The cosmetically or pharmaceutically acceptable salts of 15 the peptides of the invention can be obtained by the conventional methods, well known in the prior art [*Berge S.M. et al., "Pharmaceutical Salts", (1977), J. Pharm. Sci., 66, 119*].

Preparation procedures of the compounds of the invention

- 20 Synthesis of the compounds of the invention, their stereoisomers, mixtures thereof and/or their cosmetically or pharmaceutically acceptable salts can be carried out according to conventional methods, known in the prior art, such as using solid phase peptide synthesis methods [*Stewart J.M. y Young J.D., "Solid Phase Peptide Synthesis, 2nd edition", (1984), Pierce Chemical Company, Rockford, Illinois; Bodanzsky M. y Bodanzsky A., "The practice of Peptide Synthesis", (1994), Springer Verlag, Berlin; LloydWilliams P. et al., "Chemical Approaches to the Synthesis of Peptides and Proteins", (1997), CRC, Boca Raton, FL, USA*], synthesis in solution, enzymatic synthesis [*Kullmann W. "Proteases as catalysts for enzymic syntheses of opioid peptides", (1980), J.Biol.Chem., 255(17), 82348238*] or any combination thereof.
- 25 30 Compounds can also be obtained by fermentation of a strain of bacteria, modified or unmodified, by genetic engineering with the objective of producing the desired sequences, or by controlled hydrolysis of proteins with animal, fungal, or preferably plant origins, which free peptide fragments which contain, at least, the desired sequence.

For example, a method of obtaining the compounds (I) of the invention, their stereoisomers and mixtures thereof comprises the stages of:

- coupling of an amino acid, with the *N*-terminal end protected and the *C*-terminal end free, with an amino acid with the *N*-terminal end free and the *C*-terminal end protected or bound to a solid support ;
- 5 —elimination of the group protecting the *N*-terminal end;
- repetition of the coupling sequence and elimination of the group protecting the *N*-terminal end until the desired peptide sequence is obtained;
- elimination of the group protecting the *C*-terminal end or cleavage of the solid support.
- 10

Preferably, the *C*-terminal end is bound to a solid support and the procedure is carried out in solid phase and, therefore, comprises the coupling of an amino acid with the protected *N*-terminal end and the free *C*-terminal end with an amino acid with the *N*-terminal end free and the *C*-terminal end bound to a polymeric support; elimination of the group protecting the *N*-terminal end; and repetition of this sequence as many times as is necessary to thus obtain the compound of the desired length, finally followed by the cleavage of the synthesized compound from the original polymeric support.

The functional groups of the side chains of the amino acids are maintained conveniently protected with temporary or permanent protective groups throughout synthesis, and can be unprotected simultaneously or orthogonally to the process of cleavage of the peptide from the polymeric support.

Alternatively, solid phase synthesis can be carried out using a convergent strategy coupling a peptide with the polymeric support or with a peptide or amino acid previously bound to the polymeric support. Convergent synthesis strategies are widely known by persons skilled in the art and are described in *Lloyd-Williams P. et al.*, "Convergent Solid-Phase Peptide Synthesis", (1993), *Tetrahedron*, 49(48), 11065-11133.

The procedure can comprise the additional stages of deprotection of the *N*-terminal and *C*-terminal ends and/or cleavage of the peptide from the polymeric support in an indiscriminate order, using standard procedures and conditions known in the prior art, after which the functional groups of these ends can be modified. The optional

modification of the *N*-terminal and *C*-terminal ends can be carried out with the peptide of formula (I) anchored to the polymeric support or once the peptide has been separated from the polymeric support.

5 Optionally, R_1 can be introduced by the reaction of the *N*-terminal end of the compound of the invention with a R_1 -X compound, wherein R_1 has the aforementioned meaning and X is a leaving group, such as and not restricted to, the tosyl group, the mesyl group and halogen groups among others; through a nucleophilic substitution reaction, in the presence of an adequate base and solvent, wherein the fragments that have the functional groups not involved in the *N*-C bond formation are suitably 10 protected with temporary or permanent protective groups.

15 Optionally and/or additionally, the R_2 radicals can be introduced by the reaction of a compound HR_2 wherein R_2 is $-OR_3$, $-NR_3R_4$ or $-SR_3$, with a complementary fragment which corresponds to the compound of formula (I) in which R_2 is $-OH$ in the presence of an adequate solvent and a base such as, *N,N*-diisopropylethylamine (DIEA) or triethylamine or an additive such as 1-hydroxybenzotriazole (HOBt) or 1-hydroxyazabenzotriazole (HOAt) and a dehydrating agent, such as a carbodiimide, a 20 uronium salt, a phosphonium salt or amidinium salt, among others, or by prior formation of an acyl halide with, for example, thionyl chloride, and thereby obtaining a peptide according to the invention of general formula (I), wherein the fragments that have the functional groups not involved in the *N*-C bond formation are suitably protected with temporary or permanent protective groups, or alternatively other R_2 radicals may be introduced by simultaneous incorporation to the cleavage process of the peptide from the polymeric support.

25 A person skilled in the art would easily understand that the deprotection/cleavage steps of the *C*-terminal and *N*-terminal ends and their subsequent derivatization can be performed in a different order, according to the processes known in the prior art.

30 The term "protective group" relates to a group which blocks an organic functional group and which can be removed in controlled conditions. The protective groups, their relative reactivities and the conditions in which they remain inert are known to the person skilled in the art.

Examples of representative protective groups for the amino group are amides, such as amide acetate, amide benzoate, amide pivalate; carbamates such as benzyloxycarbonyl (Cbz or Z), 2-chlorobenzyl (ClZ), para-nitrobenzyloxycarbonyl

(pNZ), *tert*-butyloxycarbonyl (Boc), 2,2,2-trichloroethyloxycarbonyl (Troc), 2-(trimethylsilyl)ethyloxycarbonyl (Teoc), 9-fluorenylmethyloxycarbonyl (Fmoc) or allyloxycarbonyl (Alloc), Trityl (Trt), methoxytrityl (Mtt), 2,4-dinitrophenyl (Dnp), *N*-[1-(4,4-dimethyl-2,6-dioxocyclohex-1-ylidene)ethyl (Dde), 1-(4,4-dimethyl-2,6-dioxocyclohexylidene)-3-methylbutyl (ivDde), 1-(1-adamantyl)-1-methylethoxycarbonyl (Adpoc), among others, preferably Boc or Fmoc.

Examples of representative protective groups for the carboxyl group are esters, such as the *tert*-butyl ester (tBu), allyl ester (All), triphenylmethyl ester (Trt ester), cyclohexyl ester (cHx), benzyl ester (Bzl), *ortho*-nitrobenzyl ester, *para*-nitrobenzyl ester, *para*-methoxybenzyl ester, trimethylsilylethyl ester, 2-phenylisopropyl ester, fluorenylmethyl ester (Fm), 4-(*N*-[1-(4,4-dimethyl-2,6-dioxocyclohexylidene)-3-methylbutyl]amino) benzyl ester (Dmab), among others; preferred protective groups of the invention are the All, tBu, cHx, Bzl and Trt esters .

The side chains of the trifunctional amino acids can be protected during the synthetic process with temporary or permanent protective groups orthogonal to the protective groups of the *N*-terminal and *C*-terminal ends.

The hydroxyl group of the tyrosine side chain can be protected with the 2-bromobenzylloxycarbonyl group (2-BrZ), tBu, All, Bzl or 2,6-dichlorobenzyl (2,6-diClZ) among others. The histidine side chain can be protected by a protective group selected from the group formed by Tos, Dnp, methyl (Me), Boc, benzyloxymethyl (Bom), Bzl, Fmoc, Mts, Trt and Mtt. The amide group of the glutamine and asparagine side chain can be protected by the Trt group or xanthyl group (Xan) or can be used unprotected. For the protection of the carboxyl group of the aspartic acid and glutamic acid side chain esters can be used such as tBu ester, All ester, triphenylmethyl ester (Trt ester), cHx ester, Bzl ester, *ortho*-nitrobenzyl ester, *para*-nitrobenzyl ester, *para*-methoxybenzyl ester, trimethylsilylethyl ester, 2-phenylisopropyl ester, Fm ester or Dmab ester, among others. The arginine side chain can be protected by a protective group selected from the group formed by Tos, 4-methoxy-2,3,6-trimethylbenzenesulfonyl (Mtr), Alloc, nitro, 2,2,4,6,7-pentamethyldihydrobenzofuran-5-sulfonyl (Pbf) and 2,2,5,7,8-pentamethylchroman-6-sulfonyl (Pmc). The indole group of the tryptophan side chain can be protected by the formyl group (For), Boc, Mts or can be used unprotected. For the protection of the amino groups of the lysine side chains amides can be used, such as amide acetate, amide benzoate, amide pivalate; carbamates such as Cbz or Z, ClZ, pNZ, Boc, Troc, Teoc, Fmoc or Alloc, Trt, Mtt, Dnp, Dde, ivDde,

Adpac, among others. The methionine side chain can be protected in sulfoxide form, in sulfone form or used without protection. The methionyl(sulfoxide) and methionyl(sulfone) side chains are not protected.

In a preferred embodiment, the protective group strategy used is the strategy wherein
5 the amino groups are protected by Boc, the carboxyl groups are protected by Bzl, cHx or All, the tyrosine side chain is protected with 2-BrZ or Bzl, the histidine side chain is protected by the Tos or Bom group, the aspartic acid and glutamic acid side chains are protected with Bzl, cHx or All, glutamine and asparagine are used without protection in their side chain, methionine is used without protection in its side chain, the arginine
10 side chain is protected by Tos, the tryptophan side chain is protected by For or Mts and the lysine side chain is protected by ClZ, Fmoc or Alloc.

In another preferred embodiment, the protective group strategy used is the strategy wherein the amino groups are protected by Fmoc, the carboxyl groups are protected by the tBu, All or Trt esters, the tyrosine side chain is protected by tBu, the histidine side
15 chain is protected by the Trt or Mtt group, the aspartic acid and glutamic acid side chains are protected with tBu or All, glutamine and asparagine are protected by the Trt group in their side chain, methionine is used without protection in its side chain, the arginine side chain is protected by Pmc or Pbf, the tryptophan side chain is protected by Boc or is used unprotected, and the lysine side chain is protected by Boc, Trt or
20 Alloc.

Examples of these and other additional protective groups, their introduction and removal, can be found in the literature [*Atherton B. and Sheppard R.C., "Solid Phase Peptide Synthesis: A practical approach"*, (1989), *IRL Oxford University Press*]. The term "protective groups" also includes the polymeric supports used in solid phase
25 synthesis.

When synthesis takes place totally or partially in solid phase, the possible solid supports used in the procedure of the invention involve polystyrene supports, polyethylene glycol grafted to polystyrene and similar, such as and not restricted to, *p*-methylbenzhydrylamine resins (MBHA) [*Matsueda G.R. et al., "A 30 p-methylbenzhydrylamine resin for improved solid phase synthesis of peptide amides"*, (1981), *Peptides*, 2, 4550], 2-chlorotrityl resins [*Barlos K. et al., "Darstellung geschützter PeptidFragmente unter Einsatz substituierter TriphenylmethylHarze"*, (1989), *Tetrahedron Lett.*, 30, 39433946; *Barlos K. et al., "Veresterung von partiell*

geschützten PeptidFragmenten mit Harzen. Einsatz von 2-Chlorotritylchlorid zur Synthese von Leu1Gastrin I", (1989), *Tetrahedron Lett.*, 30, 3947-3951], TentaGel® resins (Rapp Polymere GmbH), ChemMatrix® resins (Matrix Innovation, Inc) and similar, which may or may not include a labile linker, such as

5 5-(4-aminomethyl-3,5-dimethoxyphenoxy) valeric acid (PAL) [Albericio F. et al., "Preparation and application of the 5-(4-(9-fluorenylmethyloxycarbonyl) aminomethyl-3,5-dimethoxy-phenoxy)valeric acid (PAL) handle for the solid phase synthesis of C-terminal peptide amides under mild conditions", (1990), *J. Org. Chem.*, 55, 3730-3743], 2-[4-aminomethyl-(2,4-dimethoxyphenyl)] phenoxyacetic acid (AM)

10 [Rink H., "Solid phase synthesis of protected peptide fragments using a trialkoxy-diphenyl-methylester resin", (1987), *Tetrahedron Lett.*, 28, 3787-3790], Wang [Wang S.S., "p-Alkoxybenzyl Alcohol Resin and p-Alkoxybenzyloxycarbonylhydrazide Resin for Solid Phase Synthesis of Protected Peptide Fragments", (1973), *J. Am. Chem. Soc.*, 95, 1328-1333] and similar, which enable simultaneous deprotection and cleavage of

15 the peptide from the polymeric support.

Cosmetic or pharmaceutical compositions of the invention

The compounds of the invention can be administered to inhibit neuronal exocytosis by any means which causes contact between the compounds and the site of action in a

20 mammal's body, preferably that of a human being, and in the form of a composition which contains them.

To this regard, another aspect of the invention is a cosmetic or pharmaceutical composition which comprises at least one compound of general formula (I), its stereoisomers, mixtures thereof, and/or its cosmetically or pharmaceutically acceptable

25 salts together with at least one cosmetically or pharmaceutically acceptable adjuvant. These compositions can be prepared by conventional means known to persons skilled in the art [“Harry's Cosmeticology”, Seventh edition, (1982), Wilkinson J.B., Moore R.J., ed. Longman House, Essex, GB].

The compounds of this invention have variable solubility in water, according to the

30 nature of their amino acid sequence or any possible modifications in the N-terminal and/or C-terminal ends. Therefore, the compounds of this invention can be incorporated into the compositions by aqueous solution, and those which are not soluble in water can be solubilized in cosmetically or pharmaceutically acceptable

conventional solvents such as and not restricted to, ethanol, propanol, isopropanol, propylene glycol, glycerin, butylene glycol or polyethylene glycol or any combination thereof.

The 5 cosmetically or pharmaceutically effective amount of the compounds of the invention which should be administered, as well as their dosage, will depend on numerous factors, including age, state of the patient, the nature or severity of the condition, disorder or disease to be treated and/or cared for, the route and frequency of administration and the particular nature of the compounds to be used.

“Cosmetically and pharmaceutically effective amount” is understood to mean a non-toxic but sufficient amount of the compound or compounds of the invention to provide the desired effect. The compounds of the invention are used in the cosmetic or pharmaceutical composition of this invention at cosmetically or pharmaceutically effective concentrations to achieve the desired effect; in a preferred form with regards to the total weight of the composition, between 0.00000001% (in weight) and 20% (in weight); preferably between 0.000001% (in weight) and 15% (in weight), more preferably between 0.0001% (in weight) and 10% (in weight) and even more preferably between 0.0001% (in weight) and 5% (in weight).

The 20 compounds of general formula (I), their stereoisomers, mixtures thereof and/or their cosmetic or pharmaceutically acceptable salts, can also be incorporated into cosmetic or pharmaceutical delivery systems and/or sustained release systems.

The term “delivery systems” relates to a diluent, adjuvant, excipient or carrier with which the compound of the invention is administered. These cosmetic or pharmaceutical carriers can be liquids, such as water, oils or surfactants, including those of petroleum, animal, plant or synthetic origin, such as and not restricted to, 25 peanut oil, soybean oil, mineral oil, sesame oil, castor oil, polysorbates, sorbitan esters, ether sulfates, sulfates, betaines, glycosides, maltosides, fatty alcohols, nonoxynols, poloxamers, polyoxyethylenes, polyethylene glycols, dextrose, glycerol, digitonin and similar. A person skilled in the art knows the diluents, adjuvants or excipients which can be used in the different delivery systems in which the compound of the invention can 30 be administered.

The term “sustained release” is used in a conventional sense relating to a delivery system of a compound which provides the gradual release of this compound during a

period of time and preferably, although not necessarily, with relatively constant compound release levels over a long period of time.

- Examples of delivery or sustained release systems include, without restriction, liposomes, mixed liposomes, oleosomes, niosomes, ethosomes, milliparticles, 5 microparticles, nanoparticles and solid lipid nanoparticles, nanostructured lipid carriers, sponges, cyclodextrins, vesicles, micelles, mixed micelles of surfactants, surfactant-phospholipid mixed micelles, millispheres, microspheres and nanospheres, lipospheres, millicapsules, microcapsules and nanocapsules, as well as in microemulsions and nanoemulsions, which can be added to achieve a greater 10 penetration of the active principle and/or improve its pharmacokinetic and pharmacodynamic properties. Preferred delivery or sustained release systems are liposomes, surfactant-phospholipid mixed micelles, microemulsions, more preferably water-in-oil microemulsions with an internal structure of reverse micelle and nanocapsules containing microemulsions.
- 15 The sustained release systems can be prepared by methods known in the prior art, and the compositions which contain them can be administered, for example, by topical or transdermal administration, including adhesive patches, non-adhesive patches, occlusive patches and microelectric patches, or by systemic administration, for example and not restricted to, oral or parenteral route, including nasal, rectal or 20 subcutaneous implantation or injection, or direct implantation or injection into a specific body part, and preferably should release a relatively constant quantity of the peptides of the invention. The amount of compound contained in the sustained release system will depend, for example, on where the composition is to be administered, the kinetics and duration of the release of the compound of the invention, as well as the nature of 25 the condition, disorder and/or disease to be treated and/or cared for.

The compounds of this invention can also be adsorbed on solid organic polymers or solid mineral supports such as and not restricted to, talc, bentonite, silica, starch or maltodextrin among others.

- 30 The compositions which contain the compounds of general formula (I), their stereoisomers, mixtures thereof and/or their cosmetically or pharmaceutically acceptable salts can also be incorporated into fabrics, non-woven fabrics and medical devices which are in direct contact with the skin, thus releasing the compounds of the invention whether by biodegradation of the binding system to the fabric, non-woven

fabric or medical device, or by friction between them and the body, due to bodily moisture, the skin's pH or body temperature. Furthermore, the compounds of the invention can be incorporated into the fabrics and non-woven fabrics used to make garments that are in direct contact with the body. Preferably, the fabrics, non-woven
5 fabrics and medical devices containing the compounds of the invention are used for the treatment and/or care of conditions, disorders and/or diseases which improve or are prevented by the inhibition of neuronal exocytosis.

Examples of fabrics, non-woven fabrics, garments, medical devices and means for immobilizing the compounds to them, among which are the delivery systems and/or the
10 sustained release systems described above, can be found in the literature and are known in the prior art [Schaab C.K. (1986) *HAPPI* May 1986; Nelson G., "Application of microencapsulation in textiles", (2002), *Int. J. Pharm.*, 242(1-2), 55-62; "Biofunctional Textiles and the Skin" (2006) *Curr. Probl. Dermatol.* v.33, *Hippler U.C. and Elsner P.*, eds. S. Karger AG, Basel, Switzerland; Malcolm R.K. et al., "Controlled release of a
15 model antibacterial drug from a novel self-lubricating silicone biomaterial", (2004), *J. Cont. Release*, 97(2), 313-320]. The preferred fabrics, non-woven fabrics, garments and medical devices are bandages, gauzes, t-shirts, socks, tights, underwear, girdles, gloves, diapers, sanitary napkins, dressings, bedspreads, wipes, adhesive patches, non-adhesive patches, occlusive patches, microelectric patches and/or face masks.
20 The cosmetic or pharmaceutical compositions which contain the compounds of the invention, their stereoisomers, mixtures thereof and/or their cosmetically or pharmaceutically acceptable salts, can be used in different types of compositions of topical or transdermal application which optionally include cosmetically or pharmaceutically acceptable excipients necessary for formulating the desired
25 administration form. A person skilled in the art knows the different excipients which can be used in the cosmetic or pharmaceutical compositions which contain the compounds of the invention.

The compositions of topical or transdermal application can be produced in any solid, liquid or semi-solid formulation, such as and not restricted to, creams, multiple
30 emulsions such as and not restricted to, oil and/or silicone in water emulsions, water-in-oil and/or silicone emulsions, water/oil/water or water/silicone/water type emulsions, and oil/water/oil or silicone/water/silicone type emulsions, anhydrous compositions, aqueous dispersions, oils, milks, balsams, foams, lotions, gels, cream gels, hydroalcoholic solutions, hydroglycolic solutions, hydrogels, liniments, sera, soaps,

shampoos, conditioners, serums, polysaccharide films, ointments, mousses, pomades, powders, bars, pencils and sprays or aerosols (sprays), including leave-on and rinse-off formulations. These topical or transdermal application formulations can be incorporated using techniques known by the person skilled in the art into different types 5 of solid accessories such as and not restricted to, bandages, gauzes, t-shirts, socks, tights, underwear, girdles, gloves, diapers, sanitary napkins, dressings, bedspreads, wipes, adhesive patches, non-adhesive patches, occlusive patches, microelectric patches or face masks, or they can be incorporated into different make-up products such as make-up foundation, such as fluid foundations and compact foundations, 10 make-up removal lotions, make-up removal milks, under-eye concealers, eye shadows, lipsticks, lip protectors, lip gloss and powders, among others.

The cosmetic or pharmaceutical compositions of the invention may include agents which increase the percutaneous absorption of the compounds of this invention, such as and not restricted to, dimethyl sulfoxide, dimethylacetamide, dimethylformamide, 15 surfactants, azone (1-dodecylazacycloheptane-2-one), alcohol, urea, ethoxydiglycol, acetone, propylene glycol or polyethylene glycol, among others. Furthermore, the cosmetic or pharmaceutical compositions of this invention can be applied to local areas to be treated by means of iontophoresis, sonophoresis, electroporation, microelectric patches, mechanical pressure, osmotic pressure gradient, occlusive cure, 20 microinjections or needle-free injections by means of pressure, such as injections by oxygen pressure, or any combination thereof, to achieve a greater penetration of the compound of the invention. The application area will be determined by the nature of the condition, disorder and/or disease to be treated and/or cared for.

Furthermore, the cosmetic compositions containing the compounds of general formula 25 (I), their stereoisomers, mixtures thereof and/or their cosmetically or pharmaceutically acceptable salts can be used in different types of formulations for oral administration, preferably in the form of oral cosmetics or drugs, such as and not restricted to, capsules, including gelatin capsules, soft capsules, hard capsules, tablets, including sugar coated tablets, pills, powders, granules, chewing gum, solutions, suspensions, 30 emulsions, syrups, elixirs, polysaccharide films, jellies or gelatins, and any other form known by the person skilled in the art. In a particular embodiment, the compounds of the invention can be incorporated into any form of functional food or fortified food, such as and not restricted to, dietary bars or compact or non-compact powders. These powders can be dissolved in water, soda, dairy products, soy derivatives or can be

incorporated into dietary bars. The compounds of this invention can be formulated with common excipients and adjuvants for oral compositions or food supplements, such as and not restricted to, fat components, aqueous components, humectants, preservatives, texturizing agents, flavors, aromas, antioxidants and colorants common 5 in the food industry.

- Cosmetic or pharmaceutical compositions containing the compounds of general formula (I), their stereoisomers, mixtures thereof and/or their cosmetically or pharmaceutically acceptable salts can also be administered, as well as by topical or transdermal route, by any other appropriate route, such as oral or parenteral route, for 10 which they will include the pharmaceutically acceptable excipients necessary for the formulation of the desired administration form. In the context of this invention, the term “parenteral” includes nasal, auricular, ophthalmic, rectal, urethral, vaginal, subcutaneous, intradermal, intravascular injections such as intravenous, intramuscular, intraocular, intravitreous, intracorneal, intraspinal, intramedullary, intracranial, 15 intracervical, intracerebral, intrameningeal, intraarticular, intrahepatic, intrathoracic, intratracheal, intrathecal and intraperitoneal, and any other similar injection or infusion technique. A person skilled in the art knows the different means by which the cosmetic or pharmaceutical compositions which contain the compounds of the invention can be administered.
- 20 Among the cosmetically or pharmaceutically acceptable excipients and/or adjuvants contained in the cosmetic or pharmaceutical compositions described in this invention are additional ingredients commonly used in cosmetic or pharmaceutical compositions, such as and not restricted to, other agents which inhibit neuronal exocytosis, other anticholinergic agents, other agents which inhibit muscular contraction, other anti-aging 25 agents, other anti-wrinkle agents, other antiperspirant agents, other anti-inflammatory agents and/or analgesics, other anti-itching agents, calming agents, anesthetic agents, inhibitors of acetylcholine-receptor aggregation, agents that inhibit acetylcholinesterase, skin relaxant agents, melanin synthesis stimulating or inhibiting agents, whitening or depigmenting agents, propigmenting agents, self-tanning agents, 30 NO-synthase inhibiting agents, 5 α -reductase inhibiting agents, lysyl- and/or prolyl hydroxylase inhibiting agents, antioxidants, free radical scavengers and/or agents against atmospheric pollution, reactive carbonyl species scavengers, anti-glycation agents, antihistamine agents, antiviral agents, antiparasitic agents, emulsifiers, emollients, organic solvents, liquid propellants, skin conditioners, humectants,

substances that retain moisture, alpha hydroxy acids, beta hydroxy acids, moisturizers, epidermal hydrolytic enzymes, vitamins, amino acids, proteins, pigments or colorants, dyes, biopolymers, gelling polymers, thickeners, surfactants, softening agents, emulsifiers, binding agents, preservatives, agents able to reduce or treat the bags under the eyes, exfoliating agents, keratolytic agents, flaking agents, antimicrobial agents, antifungal agents, fungistatic agents, bactericidal agents, bacteriostatic agents, agents stimulating the synthesis of dermal or epidermal macromolecules and/or capable of inhibiting or preventing their degradation, collagen synthesis-stimulating agents, elastin synthesis-stimulation agents, decorin synthesis-stimulation agents, laminin synthesis-stimulation agents, defensin synthesis-stimulating agents, chaperone synthesis-stimulating agents, cAMP synthesis-stimulating agents, AQP-3 modulating agents, aquaporin synthesis modulating agents, proteins from the aquaporin family, hyaluronic acid synthesis-stimulating agents, glycosaminoglycan synthesis-stimulating agents, fibronectin synthesis-stimulating agents, sirtuin synthesis-stimulating agents, sirtuin activating agents, heat shock proteins, heat shock protein synthesis-stimulating agents, agents stimulating the synthesis of lipids and components of the stratum corneum, ceramides, fatty acids, agents that inhibit collagen degradation, matrix metalloproteinase inhibitory agents, agents that inhibit elastin degradation, agents that inhibit serine proteases such as kallikreins, leukocyte elastase or cathepsin G, agents stimulating fibroblast proliferation, agents stimulating keratinocyte proliferation, agents stimulating adipocyte proliferation, agents stimulating melanocyte proliferation, agents stimulating keratinocyte differentiation, agents stimulating or delaying adipocyte differentiation, antihyperkeratosis agents, comedolytic agents, anti-psoriasis agents, DNA repair agents, DNA protecting agents, stabilizers, agents for the treatment and/or care of sensitive skin, firming agents, anti-stretch mark agents, binding agents, agents regulating sebum production, lipolytic agents or agents stimulating lipolysis, adipogenic agents, agents modulating PGC-1 α expression, agents modulating the activity of PPAR γ , agents which increase or reduce the triglyceride content of adipocytes, anti-cellulite agents, agents which inhibit the activity of PAR-2, agents stimulating healing, coadjuvant healing agents, agents stimulating reepithelialization, coadjuvant reepithelialization agents, cytokine growth factors, agents acting on capillary circulation and/or microcirculation, agents stimulating angiogenesis, agents that inhibit vascular permeability, venotonic agents, agents acting on cell metabolism, agents to improve dermal-epidermal junction, agents inducing hair growth, hair growth inhibiting or retardant agents, hair loss retardant agents, preservatives, perfumes, cosmetic and/or

absorbent and/or body odor masking deodorants, chelating agents, plant extracts, essential oils, marine extracts, agents obtained from a biotechnological process, mineral salts, cell extracts, sunscreens and organic or mineral photoprotective agents active against ultraviolet A and/or B rays and/or infrared A rays, or mixtures thereof,

5 provided that they are physically and chemically compatible with the rest of components in the composition and particularly with the compounds of the invention. Likewise, the nature of these additional ingredients should not unacceptably alter the benefits of the compounds of this invention. The nature of these additional ingredients can be synthetic or natural, such as plant extracts, or come from a biotechnological

10 procedure, or from a combination of a synthetic procedure and a biotechnological procedure. Additional examples can be found in *CTFA International Cosmetic Ingredient Dictionary & Handbook, 12th Edition* (2008). In the context of this invention, biotechnological procedure is understood to be any procedure to produce the active ingredient, or part of it, in an organism, or in part of it.

15 An additional aspect of this invention refers to a cosmetic or pharmaceutical composition which comprises a cosmetically or pharmaceutically effective quantity of at least one compound general formula (I), its stereoisomers, mixtures thereof and/or its cosmetically or pharmaceutically acceptable salts, as well as a cosmetically or pharmaceutically effective quantity of at least one extract, synthetic compound or

20 product of biotechnological origin which is an anti-wrinkle agent and/or anti-aging agent such as and not restricted to the extracts or hydrolyzed extracts of *Vitis vinifera*, *Rosa canina*, *Curcuma longa*, *Theobroma cacao*, *Ginkgo biloba*, *Leontopodium alpinum* or *Dunaliella salina* among others, Matrixyl® [INCI: Palmitoyl Pentapeptide-4], Matrixyl® 3000® [INCI: Palmitoyl Tetrapeptide-7, Palmitoyl Oligopeptide], Matrixyl® Synthe'6™

25 [INCI: Glycerin, Water, Hydroxypropyl Cyclodextrin, Palmitoyl Tripeptide-38], Essenskin™ [INCI: calcium hydroxymethionine], Renovage [INCI: teprenone], Resistem™ [INCI: Globularia Cordifolia Ferment] or Dermaxyl® [INCI: Palmitoyl Oligopeptide] marketed by Sederma/Croda, Vialox® [INCI: Pentapeptide-3], Syn® Ake® [INCI: Dipeptide Diaminobutyroyl Benzylamide Diacetate], Syn®-Coll [INCI: Palmitoyl

30 Tripeptide-5], Phytaluronate [INCI: Locust Bean (*Ceratonia siliqua*) Gum] or Preregen® [INCI: Glycine soja (Soybean) Protein, Oxido Reductases] marketed by Pentapharm/DSM, Myoxinol™ [INCI: Hydrolyzed *Hibiscus esculentus* Extract], Syniorage™ [INCI: Acetyl Tetrapeptide-11], Dermican™ [INCI: Acetyl Tetrapeptide-9] or DN AGE™ LS [INCI: Cassia alata leaf Extract] marketed by Laboratoires

35 Sérobiologiques/Cognis/BASF, Algisum C® [INCI: Methylsilanol Mannuronate] or

Hydroxyprolisilane CN® [INCI: Methylsilanol Hydroxyproline Aspartate] marketed by Exsymol, Argireline® [INCI: Acetyl Hexapeptide-8], SNAP-7 [INCI: Acetyl Heptapeptide-4], SNAP-8 [INCI: Acetyl Octapeptide-3], Leuphasyl® [INCI: Pentapeptide-18], Inyline™ [INCI: Acetyl Hexapeptide-30], Aldenine® [INCI: Hydrolized Wheat Protein, Hydrolized 5 Soy Protein, Tripeptide-1], Preventhelia™ [INCI: Diaminopropionyl Tripeptide-33], Decorinyl® [INCI: Tripeptide-10 Citrulline], Decorinol® [INCI: Tripeptide-9 Citrulline], Trylagen® [INCI: Pseudoalteromonas Ferment Extract, Hydrolyzed Wheat Protein, Hydrolyzed Soy Protein, Tripeptide-10 Citrulline, Tripeptide-1], Eyeseryl® [INCI: Acetyl Tetrapeptide-5], Peptide AC29 [INCI: Acetyl Tripeptide-30 Citrulline], Relistase™ [INCI: 10 Acetylarginyltriptophyl Diphenylglycine], Thermostressine® [INCI: Acetyl Tetrapeptide-22], Lipochroman™ [INCI: Dimethylmethoxy Chromanol], Chromabright™ [INCI: Dimethylmethoxy Chromanyl Palmitate], Antarcticine® [INCI: Pseudoalteromonas Ferment Extract], dGlyage™ [INCI: Lysine HCl, Lecithin, Tripeptide-9 Citrulline], Vilastene™ [INCI: Lysine HCl, Lecithin, Tripeptide-10 Citrulline], Hyadisine™ [INCI: 15 Pseudoalteromonas Ferment Extract], Hyanify™ [proposed INCI: Saccharide Isomerate], Diffuporine™ [INCI: Acetyl Hexapeptide-37], Silusyne™ [INCI: Soybean (Glycine Soja) Oil, Sorbitan Sesquioleate, Isohexadecane, Sodium Hyaluronate, Lauryldimonium Hydroxypropyl Hydrolized Soy Protein, Acetyl Hexapeptide-39] or Adifyline™ [INCI: Acetyl Hexapeptide-38] marketed by Lipotec, Kollaren® [INCI: 20 Tripeptide-1, Dextran] marketed by Institut Europeen de Biologie Cellulaire, Collaxyl® IS [INCI: Hexapeptide-9], Laminixyl IS™ [INCI: Heptapeptide], Orsirtine™ GL [INCI: Oryza sativa (Rice) Extract], D'Orientine™ IS [INCI: Phoenix dactylifera (Date) Seed Extract], Phytoquintescine™ [INCI: Einkorn (Triticum monococcum) Extract] or Quintescine™ IS [INCI: Dipeptide-4] marketed by Vincience/ISP/Ashland, BONT-L- 25 Peptide [INCI: Palmitoyl Hexapeptide-19] marketed by Infinitec Activos, Deepaline™ PVB [INCI: Palmitoyl hydrolyzed Wheat Protein] or Sepilift® DPHP [INCI: Dipalmitoyl Hydroxyproline] marketed by Seppic, Gatuline® Expression [INCI: *Acmella oleracea* Extract], Gatuline® In-Tense [INCI: *Spilanthes acmella* Flower Extract] or Gatuline® Age Defense 2 [INCI: *Juglans regia* (Walnut) Seed Extract] marketed by Gattefosse, 30 Thalassine™ [INCI: Algae Extract] marketed by Biotechmarine, ChroNOLine™ [INCI: Caprooyl Tetrapeptide-3] or Thymulen-4 [INCI: Acetyl Tetrapeptide-2] marketed by Atrium/Unipex Innovations, EquiStat [INCI: *Pyrus malus* Fruit Extract, *Glycine soja* Seed Extract] or Juvenesce [INCI: Ethoxydiglycol and Caprylic Triglycerid, Retinol, Ursolic Acid, Phytonadione, Ilomastat] marketed by Coletica/Engelhard/BASF, Ameliox 35 [INCI: Carnosine, Tocopherol, *Silybum Marianum* Fruit Extract] or PhytoCellTec Malus

- Domestica [INCI: *Malus domestica* Fruit Cell Culture] marketed by Mibelle Biochemistry, Bioxilift [INCI: *Pimpinella anisum* Extract] or SMS Anti-Wrinkle® [INCI: *Annona squamosa* Seed Extract] marketed by Silab, antagonists of the Ca^{2+} channel such as and not restricted to, alverine, manganese or magnesium salts, certain 5 secondary or tertiary amines, retinol and its derivatives, idebenone and its derivatives, Coenzyme Q10 and its derivatives, boswellic acid and its derivatives, GHK and its derivatives and/or salts, carnosine and its derivatives, DNA repair enzymes such as and not restricted to, photolyase or T4 endonuclease V, or chloride channel agonists among others, and/or mixtures thereof.
- 10 An additional aspect of this invention relates to a cosmetic or pharmaceutical composition which comprises a cosmetically or pharmaceutically effective amount of at least one compound of general formula (I), its stereoisomers, mixtures thereof and/or its cosmetically or pharmaceutically acceptable salts, and, in addition, a cosmetically or pharmaceutically effective amount of at least one natural extract or essential oil which
- 15 is an anti-itching agent, for example and not restricted to, extracts of *Abelmoschus esculentus*, *Actaea alba*, *Aglaia odorata*, *Alkanna tinctoria*, *Althaea officinalis*, *Altingia excelsa*, *Andropogon virginicus*, *Aralia nudicaulis*, *Aralia racemosa*, *Argemone mexicana*, *Barleria prionitis*, *Camelia sinensis*, *Caesalpinia digyna*, *Campsis grandiflora*, *Carissa congesta*, *Carthamus oxyacantha*, *Cassia tora*, *Chrysanthemum 20 indicum*, *Cimicifuga racemosa*, *Cinnamomum camphora*, *Clematis vitalba*, *Cuscuta reflexa*, *Diospyros peregrina*, *Enicostema axillare*, *Hammamelis virginiana*, *Jatropha multifida*, *Lavandula officinalis*, *Lavandula latifolia*, *Liquidambar orientalis*, *Lithospermum officinale*, *Madhuca longifolia*, *Martynia annua*, *Medicago sativa*, *Michelia champaca*, *Mikania glomerata*, *Mimosa pudica*, *Oryza sativa*, *Phaseolus 25 vulgaris*, *Phyllanthus urinaria*, *Phyllanthus virgatus*, *Pistacia vera*, *Polygonum hydropiper*, *Quercus ilex*, *Rauvolfia caffra*, *Ricinus communis*, *Rubus idaeus*, *Sagittaria sagittifolia*, *Sandoricum koetjape*, *Sapindus mukorossi*, *Schleichera oleosa*, *Sesbania grandiflora*, *Spondias dulcis*, *Tilia sp.*, *Toona ciliata*, *Tragia involucrata*, *Trichosanthes quinquangularis*, *Vaccaria pyramidata*, *Ventilago madraspatana*, *Veratrum album* or 30 *Xanthium strumarium* among others or as well as at least one synthetic compound or product of biotechnological origin which is an anti-itching agent, for example and not restricted to, mepyramine (pyrilamine), antazoline, diphenhydramine, carbinoxamine, doxylamine, clemastine, dimenhydrinate, pheniramine, chlorphenamine (chlorpheniramine), dexchlorpheniramine, bronpheniramine, triprolidine, cyclizine, 35 chlorcyclizine, hydroxyzine, meclizine, cetirizine, levocetirizine, promethazine,

thenaldine, alimemazine (trimeprazine), cyproheptadine, azatidine, ketotifen, acrivastine, astemizole, cetirizine, loratadine, desloratadine, mizolastine, terfenadine, fexofenadine, azelastine, levocabastine, olopatadine, corticosteroids such as cortisone, hydrocortisone, dexamethasone, prednisone; Neutrazen™ [INCI: Water, Butylene Glycol, Dextran, Palmitoyl Tripeptide-8] marketed by Atrium /Unipex Innovations, Meliprene® [INCI: Dextran, Acetyl Heptapeptide-1] marketed by Institut Européen de Biologie Cellulaire/Unipex Innovations, Delisens™ [proposed INCI: Acetyl Hexapeptide-46] marketed by Lipotec, Skinasensyl™ [INCI: Acetyl Tetrapeptide-15] marketed by Laboratoires Sérobiologiques/Cognis/BASF, SymSitive® 1609 [INCI: 4-t-Butylcyclohexanol] marketed by Symrise, Symbiocell™ [INCI: Extract from *Cestrum Latifolium*] marketed by BASF, Gatuline® Derma-Sensitive [INCI: Octyldodecyl Myristate, *Capparis Spinosa* Fruit Extract] marketed by Gattefossé or MAXnolia [INCI: Magnolia Officinalis Bark Extract, *Vitis Vinifera/Vitis Vinifera* (Grape) Seed Extract, Tocopherol] marketed by Mibelle Biochemistry among others, or mixtures thereof.

15 An additional aspect of this invention relates to a cosmetic or pharmaceutical composition containing a cosmetically or pharmaceutically effective quantity of at least one compound of general formula (I), its stereoisomers, mixtures thereof and/or its cosmetically or pharmaceutically acceptable salts, and also a cosmetically or pharmaceutically effective amount of at least one anti-inflammatory agent and/or analgesic selected, for example and not restricted to, from the group formed by the extract of madecassoside, extract of echinacin, amaranth seed oil, sandalwood oil, peach tree leaf extract, extract of *Aloe vera*, *Arnica montana*, *Artemisia vulgaris*, *Asarum maximum*, *Calendula officinalis*, *Capsicum*, *Centipeda cunninghamii*, *Chamomilla recutita*, *Crinum asiaticum*, *Hamamelis virginiana*, *Harpagophytum procumbens*, *Hypericum perforatum*, *Lilium candidum*, *Malva sylvestris*, *Melaleuca alternifolia*, *Origanum majorana*, *Origanum vulgare*, *Prunus laurocerasus*, *Rosmarinus officinalis*, *Salix alba*, *Silybum marianum*, *Tanacetum parthenium*, *Thymus vulgaris*, *Uncaria guianensis* or *Vaccinium myrtillus*, mometasone furoate, prednisolone, non-steroidal anti-inflammatory drugs including cyclooxygenase or lipoxygenase inhibitors, 30 benzylamine, acetylsalicylic acid, rosmarinic acid, ursolic acid, glycyrrhizinate derivatives, α -bisabolol, azulene and analogs, sericoside, ruscogenin, escin, escoline, rutin and analogs, hydrocortisone, clobetasol, dexamethasone, halobetasol, diflorasone, fluocinonide, amcinonide, triamcinolone, fluticasone, fluocinolone, flurandrenolide, prednicarbate, prednisone, paracetamol, amoxiprin, benorilate, choline 35 salicylate, faislamine, methyl salicylate, magnesium salicylate, salsalate, diclofenac,

aceclofenac, acemetacin, bromfenac, etodolac, indomethacin, oxamethacin, proglumetacin, sulindac, tolmetin, ibuprofen, dexibuprofen, carprofen, fenbufen, fenoprofen, flurbiprofen, ketoprofen, dexketoprofen, ketorolac, loxoprofen, naproxen, miroprofen, oxaprozin, pranoprofen, tiaprofenic acid, suprofen, mefenamic acid, 5 meclofenamate, meclofenamic acid, flufenamic acid, tolfenamic acid, nabumetone, phenylbutazone, azapropazone, clofezone, kebuzone, metamizole, mofebutazone, oxyphenbutazone, phenazone, sulfinpyrazone, piroxicam, lornoxicam, meloxicam, tenoxicam, celecoxib, etoricoxib, lumiracoxib, parecoxib, rofecoxib, valdecoxib, nimesulide, naproxcinod, fluproquazone or licofelone, omega-3 and omega-6 fatty 10 acids, morphine, codeine, oxycodone, hydrocodone, diamorphine, pethidine, tramadol, buprenorphine, benzocaine, lidocaine, chlorprocaine, tetracaine, procaine, amitriptyline, carbamazepine, gabapentine, pregabalin, bisabolol, Neutrazen™ [INCI: Water, Butylene Glycol, Dextran, Palmitoyl Tripeptide-8] marketed by Atrium/Unipex Innovations, Delisens™ [proposed INCI: Acetyl Hexapeptide-46] marketed by Lipotec, 15 Meliprene® [INCI: Dextran, Acetyl Heptapeptide-1] marketed by Institut Européen de Biologie Cellulaire/Unipex Innovations, Skinasensyl™ [INCI: Acetyl Tetrapeptide-15] or Anasensyl™ [INCI: Mannitol, Ammonium Glycyrrhizate, Caffeine, Hippocastanum (Horse Chestnut) Extract] marketed by Laboratoires Serobiologiques/Cognis/BASF, Calmosensine™ [INCI: Acetyl Dipeptide-1] marketed by Sederma/Croda, coenzyme 20 Q10 or alkylglycerin ethers, among others, or mixtures thereof.

Another additional aspect of this invention relates to a cosmetic or pharmaceutical composition containing a cosmetically or pharmaceutically effective quantity of at least one compound of general formula (I), its stereoisomers, mixtures thereof and/or its 25 cosmetically or pharmaceutically acceptable salts, and also a cosmetically or pharmaceutically effective amount of at least one agent which inhibits neuronal exocytosis, anticholinergic agent, inhibitor of acetylcholine-receptor aggregation and/or a muscle contraction inhibitor selected, for example and not restricted to, from the group formed by extracts of *Atropa belladonna*, *Hyoscyamus niger*, *Mandragora officinarum*, *Chondodendron tomentosum*, plants from the *Brugmansias* genus, or from 30 the *Daturas* genus, *Clostridium botulinum* toxin, peptides derived from the protein SNAP-25, peptides derived from the protein synaptotagmin, peptides derived from the protein syntaxin, peptides derived from the protein synaptobrevin, peptides derived from the protein snapin, baclofen, carbidopa, levodopa, bromocriptine, chlorphenesin, chlorzoxazone, donepezil, mephenoxalone, reserpine, tetrabenazine, dantrolene, 35 thiocolchicoside, tizanidine, chlonidine, procyclidine, glycopyrrolate, atropine,

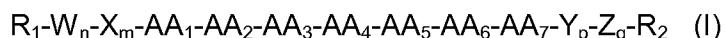
hyoscyamine, benz tropine, scopolamine, prometazine, diphenhydramine, dimenhydrinate, dicyclomine, cyclobenzaprine, orphenadrine, flavoxate, cyclopentolate, ipratropium, oxybutynin, pirenzepine, tiotropium, trihexyphenidyl, tolterodine, tropicamide, solifenacin, darifenacin, mebeverine, trimethaphan, atracurium, 5 cisatracurium, doxacurium, fazadinium, metocurine, mivacurium, pancuronium, pipecuronium, rapacuronium, tubocurarine, dimethyl tubocurarine, rocuronium, vecuronium, suxamethonium, 18-methoxycoronaridine, carisoprodol, febarbamate, meprobamate, metocarbamol, phenprobamate, tibamate, anticonvulsant agents such as levetiracetam, stiripentol, phenobarbital, methylphenobarbital, pentobarbital, 10 metarbital, barbexaclone, primidone, carbamazepine, oxcarbazepine, benzodiazepines, for example and not restricted to, clonazepam, cloxazolam, clorazepate, diazepam, flutoprazepam, lorazepam, midazolam, nitrazepam, nimetazepam, fenazepam, temazepam, tetrazepam, clobazam, Argireline[®] [INCI: Acetyl Hexapeptide-8], SNAP-7 [INCI: Acetyl Heptapeptide-4], SNAP-8 [INCI: Acetyl 15 Octapeptide-3], Leuphasyl[®] [INCI: Pentapeptide-18] or Inyline[™] [INCI: Acetyl Hexapeptide-30] marketed by Lipotec, BONT-L-Peptide [INCI: Palmitoyl Hexapeptide-19] marketed by Infinitec Activos, and Vialox[®] [INCI: Pentapeptide-3] or Syn[®] Ake[®] [INCI: Dipeptide Diaminobutyroyl Benzylamide Diacetate] marketed by Pentapharm/DSM among others, or mixtures thereof.

20 Another additional aspect of this invention relates to a cosmetic or pharmaceutical composition containing a cosmetically or pharmaceutically effective quantity of at least one compound of general formula (I), its stereoisomers, mixtures thereof and/or its cosmetically or pharmaceutically acceptable salts, and also a cosmetically or pharmaceutically effective amount of at least one cosmetic and/or absorbent and/or 25 body odor masking deodorant and/or antiperspirant agent, perfume and/or perfumed oil selected, for example and not restricted to, from the group formed by the complex zinc salt of ricinoleic acid, derived from abiotic acid, salvia essence, chamomile essence, carnation essence, lemon balm essence, mint essence, cinnamon leaf essence, lime blossom essence, juniper berry essence, vetiver essence, frankincense essence, 30 galbanum essence, labdanum essence, lavender essence, peppermint essence, benzoin, bergamot, dihydromyrcenol, lilial, lyral, citronellol, lemon essence, mandarin essence, orange essence, lavender essence, muscatel, geranium bourbon essence, aniseed, cilantro, cumin, juniper, extracts of fleur-de-lis, lily, roses, jasmine, neroli; benzyl acetate, *p*-*tert*-butylcyclohexyl acetate, linalyl acetate, phenylethyl acetate, 35 ethylmethylphenyl glycinate, linalyl benzoate, benzyl formate, allylcyclohexyl

propionate, stiraryl propionate, benzyl salicylate, benzylethylether, linear alkanes with from 8 to 18 carbon atoms, citral, ricinoleic acid, citronellal, citronellyl oxyacetaldehyde, cyclamen aldehyde, hydroxycitronellal, bourgeonal, ionones, methyl cedryl ketone, anethole, eugenol, isoeugenol, geraniol, linalool, terpineol, phenylethylalcohol, 5 α -hexylcinnamaldehyde, geraniol, benzylacetone, cyclamenaldehyde, ambroxan, indole, hedione, sandelice, cyclovertal, β -damascone, allyl amyl glycolate, dihydromyrcenol, phenoxyethyl isobutyrate, cyclohexyl salicylate, phenylacetic acid, geranyl acetate, romilate, irotyl, floramate, aluminum salts such as alum, aluminum chloride, aluminum chlorohydrate, aluminum dichlorohydrate, aluminum 10 sesquichlorohydrate, aluminum hydroxy allantoinate, aluminum chlorotartrate, aluminum and zirconium trichlorohydrate, aluminum and zirconium tetrachlorohydrate, aluminum and zirconium pentachlorohydrate and/or mixtures thereof, Leuphasyl® [INCI: Pentapeptide-18], SNAP-7 [INCI: Acetyl Heptapeptide-4], SNAP-8 [INCI: Acetyl Octapeptide-3], Argireline® [INCI: Acetyl Hexapeptide-8] or Inyline™ [INCI: Acetyl 15 Hexapeptide-30] marketed by Lipotec, Vialox® [INCI: Pentapeptide 3] or Syn® Ake® [INCI: Dipeptide Diaminobutyroyl Benzylamide Diacetate] marketed by Pentapharm/DSM and BONT-L-Peptide [INCI: Palmitoyl Hexapeptide-19] marketed by Infinitec Activos among others, or mixtures thereof.

20 Applications

In another aspect, this invention relates to a compound of general formula (I):



its stereoisomers, mixtures thereof and/or its cosmetically or pharmaceutically acceptable salts, characterized in that:

25 AA₁ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₂ is selected from the group formed by -His-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₃ is selected from the group formed by -Leu-, -Ile-, -Phe- and -Tyr-;

AA₄ is selected from the group formed by -Lys-, -His-, -Arg-, -Gln-, -Leu-, -Ile-, -Met-, -MetO- and -MetO₂-, 30

AA₅ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₆ is selected from the group formed by -Phe-, -Trp-, -Ile- and -Val-;

AA₇ is selected from the group formed by -Met-, -MetO- and -MetO₂-;

5 W, X, Y, Z are amino acids and are independently selected from amongst themselves;

n, m, p and q are independently selected from amongst themselves and have a value of 0 or 1;

n+m+p+q is smaller or equal to 2;

10 with the condition that it is not SEQ ID NO:111;

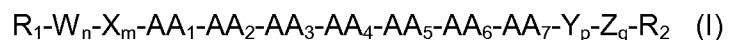
R₁ is selected from the group formed by H, a polymer derived from polyethylene glycol, substituted or unsubstituted non-cyclic aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl and R₅-CO-, wherein R₅ is selected from the group formed by H, substituted or unsubstituted non-cyclic aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl, substituted or unsubstituted heterocyclyl and substituted or unsubstituted heteroarylalkyl;

20 R₂ is selected from the group formed by -NR₃R₄, -OR₃ and -SR₃, wherein R₃ and R₄ are independently selected from the group formed by H, a polymer derived from polyethylene glycol, substituted or unsubstituted non-cyclic aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl; and

25 R₁ and R₂ are not α -amino acids,

for its use in medicine.

In another aspect, the invention relates to a compound of general formula (I):



its stereoisomers, mixtures thereof and/or its cosmetically or pharmaceutically acceptable salts, characterized in that:

AA₁ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

5 AA₂ is selected from the group formed by -His-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₃ is selected from the group formed by -Leu-, -Ile-, -Phe- and -Tyr-;

AA₄ is selected from the group formed by -Lys-, -His-, -Arg-, -Gln-, -Leu-, -Ile-, -Met-, -MetO- and -MetO₂-;

10 AA₅ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₆ is selected from the group formed by -Phe-, -Trp-, -Ile- and -Val-;

AA₇ is selected from the group formed by -Met-, -MetO- and -MetO₂-;

W, X, Y, Z are amino acids and are independently selected from amongst themselves;

15 n, m, p and q are independently selected from amongst themselves and have a value of 0 or 1;

n+m+p+q is smaller or equal to 2;

20 R₁ is selected from the group formed by H, a polymer derived from polyethylene glycol, substituted or unsubstituted non-cyclic aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl and R₅-CO-, wherein R₅ is selected from the group formed by H, substituted or unsubstituted non-cyclic aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl, substituted or unsubstituted heterocyclyl and substituted or 25 unsubstituted heteroarylalkyl;

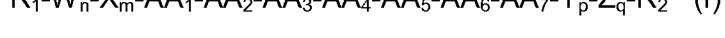
R₂ is selected from the group formed by -NR₃R₄, -OR₃ and -SR₃, wherein R₃ and

R₄ are independently selected from the group formed by H, a polymer derived from polyethylene glycol, substituted or unsubstituted non-cyclic aliphatic group,

substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl; and

R_1 and R_2 are not α -amino acids,

- 5 for its use in the treatment and/or prevention of pain, inflammation, itching, hyperhidrosis and/or of neurological, compulsive and/or neuropsychiatric diseases and/or disorders which are improved by the inhibition of neuronal exocytosis, selected from the group formed by muscular spasticity, dystonia, focal dystonia, blepharospasm, torsion dystonia, cervical dystonia or torticollis, laryngeal dystonia or spasmodic
- 10 dysphonia, oromandibular dystonia, extremity dystonia, writer's cramp, musician's cramp, foot dystonia, bruxism, facial scoliosis, hemifacial spasm, tics, strabismus, segmentary dystonia, Meige's syndrome, multifocal dystonia, hemidystonia, dopamine-responsive dystonia, Segawa's dystonia, trembling, Parkinson's disease, nerve impingements, Alzheimer's disease and Tourette's syndrome.
- 15 In another aspect, the invention relates to a compound of general formula (I):



its stereoisomers, mixtures thereof and/or its cosmetically or pharmaceutically acceptable salts, characterized in that:

AA₁ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₂ is selected from the group formed by -His-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₃ is selected from the group formed by -Leu-, -Ile-, -Phe- and -Tyr-;

AA₄ is selected from the group formed by -Lys-, -His-, -Arg-, -Gln-, -Leu-, -Ile-, -Met-, -MetO- and -MetO₂-;

25 AA₅ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₆ is selected from the group formed by -Phe-, -Trp-, -Ile- and -Val-;

AA₇ is selected from the group formed by -Met-, -MetO- and -MetO₂-;

W, X, Y, Z are amino acids and are independently selected from amongst themselves;

n, m, p and q are independently selected from amongst themselves and have a value of 0 or 1;

5 n+m+p+q is smaller or equal to 2;

10 R_1 is selected from the group formed by H, a polymer derived from polyethylene glycol, substituted or unsubstituted non-cyclic aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl and R_5 -CO-, wherein R_5 is selected from the group formed by H, substituted or unsubstituted non-cyclic aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl, substituted or unsubstituted heterocyclyl and substituted or unsubstituted heteroarylalkyl;

15 R_2 is selected from the group formed by $-NR_3R_4$, $-OR_3$ and $-SR_3$, wherein R_3 and R_4 are independently selected from the group formed by H, a polymer derived from polyethylene glycol, substituted or unsubstituted non-cyclic aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl; and

20 R_1 and R_2 are not α -amino acids,

R₁ and R₂ are not α -amino acids,

for its use in the treatment of the skin, hair and/or mucous membranes.

In another aspect, the invention relates to the use of a compound of general formula (I):



25 its stereoisomers, mixtures thereof and/or its cosmetically or pharmaceutically acceptable salts, characterized in that:

AA₁ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₂ is selected from the group formed by -His-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₃ is selected from the group formed by -Leu-, -Ile-, -Phe- and -Tyr-;

AA₄ is selected from the group formed by -Lys-, -His-, -Arg-, -Gln-, -Leu-, -Ile-, -Met-, -MetO- and -MetO₂-;

AA₅ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₆ is selected from the group formed by -Phe-, -Trp-, -Ile- and -Val-;

AA₇ is selected from the group formed by -Met-, -MetO- and -MetO₂-;

W, X, Y, Z are amino acids and are independently selected from amongst themselves;

10 n, m, p and q are independently selected from amongst themselves and have a value of 0 or 1;

n+m+p+q is smaller or equal to 2;

R₁ is selected from the group formed by H, a polymer derived from polyethylene glycol, substituted or unsubstituted non-cyclic aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl and R₅-CO-, wherein R₅ is selected from the group formed by H, substituted or unsubstituted non-cyclic aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl, substituted or unsubstituted heterocyclyl and substituted or unsubstituted heteroarylalkyl;

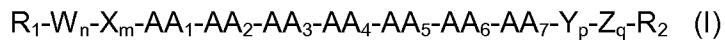
R₂ is selected from the group formed by -NR₃R₄, -OR₃ and -SR₃, wherein R₃ and R₄ are independently selected from the group formed by H, a polymer derived from polyethylene glycol, substituted or unsubstituted non-cyclic aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl; and

R₁ and R₂ are not α -amino acids,

for the cosmetic, non-therapeutic treatment and/or care of the skin, hair and/or mucous membranes, in particular for the prevention of aging and/or photoaging of the skin, hair

and/or mucous membranes, the treatment and/or prevention of wrinkles and/or expression wrinkles, treatment and/or prevention of perspiration, treatment and/or care of disorders of the skin selected from the group formed by calluses, warts, treatment stimulating hair growth and/or prevention of hair loss.

5 In another aspect the invention relates to a compound of general formula (I):



its stereoisomers, mixtures thereof and/or its cosmetically or pharmaceutically acceptable salts, characterized in that :

AA₁ is selected from the group formed
10 by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₂ is selected from the group formed by -His-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₃ is selected from the group formed by -Leu-, -Ile-, -Phe- and -Tyr-;

AA₄ is selected from the group formed
by -Lys-, -His-, -Arg-, -Gln-, -Leu-, -Ile-, -Met-, -MetO- and -MetO₂;
15

AA₅ is selected from the group formed
by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₆ is selected from the group formed by -Phe-, -Trp-, -Ile- and -Val-;

AA₇ is selected from the group formed by -Met-, -MetO- and -MetO₂;

W, X, Y, Z are amino acids and are independently selected from amongst
20 themselves;

n, m, p and q are independently selected from amongst themselves and have a
value of 0 or 1;

n+m+p+q is smaller or equal to 2;

R₁ is selected from the group formed by H, a polymer derived from polyethylene
25 glycol, substituted or unsubstituted non-cyclic aliphatic group, substituted or
unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or
unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, substituted or
unsubstituted aralkyl and R₅-CO-, wherein R₅ is selected from the group formed
by H, substituted or unsubstituted non-cyclic aliphatic group, substituted or

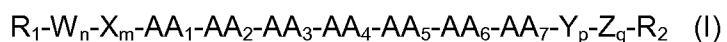
unsubstituted alicyclyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl, substituted or unsubstituted heterocyclyl and substituted or unsubstituted heteroarylalkyl;

R₂ is selected from the group formed by -NR₃R₄, -OR₃ and -SR₃, wherein R₃ and R₄ are independently selected from the group formed by H, a polymer derived from polyethylene glycol, substituted or unsubstituted non-cyclic aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl; and

R₁ and R₂ are not α -amino acids,

for its use in the inhibition of neuronal exocytosis.

Alternatively, in another aspect, the invention relates to a method of treatment and/or prevention of pain, inflammation, itching, hyperhidrosis and/or of neurological, compulsive and/or neuropsychiatric diseases and/or disorders which are improved by the inhibition of neuronal exocytosis, selected from the group formed by muscular spasticity, dystonia, focal dystonia, blepharospasm, torsion dystonia, cervical dystonia or torticollis, laryngeal dystonia or spasmodic dysphonia, oromandibular dystonia, extremity dystonia, writer's cramp, musician's cramp, foot dystonia, bruxism, facial scoliosis, hemifacial spasm, tics, strabismus, segmentary dystonia, Meige's syndrome, multifocal dystonia, hemidystonia, dopamine-responsive dystonia, Segawa's dystonia, trembling, Parkinson's disease, nerve impingements, Alzheimer's disease and Tourette's syndrome, which comprises the administration of a cosmetically or pharmaceutically effective quantity of at least one compound of general formula (I):



its stereoisomers, mixtures thereof and/or its cosmetically or pharmaceutically acceptable salts, characterized in that:

AA₁ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₂ is selected from the group formed by -His-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₃ is selected from the group formed by -Leu-, -Ile-, -Phe- and -Tyr-;

AA₄ is selected from the group formed by -Lys-, -His-, -Arg-, -Gln-, -Leu-, -Ile-, -Met-, -MetO- and -MetO₂;

AA₅ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

5 AA₆ is selected from the group formed by -Phe-, -Trp-, -Ile- and -Val-;

AA₇ is selected from the group formed by -Met-, -MetO- and -MetO₂;

W, X, Y, Z are amino acids and are independently selected from amongst themselves;

10 n, m, p and q are independently selected from amongst themselves and have a value of 0 or 1;

n+m+p+q is smaller or equal to 2;

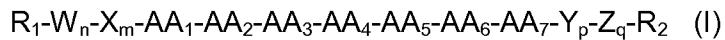
15 R₁ is selected from the group formed by H, a polymer derived from polyethylene glycol, substituted or unsubstituted non-cyclic aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl and R₅-CO-, wherein R₅ is selected from the group formed by H, substituted or unsubstituted non-cyclic aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl, substituted or unsubstituted heterocyclyl and substituted or unsubstituted heteroarylalkyl;

20 R₂ is selected from the group formed by -NR₃R₄, -OR₃ and -SR₃, wherein R₃ and R₄ are independently selected from the group formed by H, a polymer derived from polyethylene glycol, substituted or unsubstituted non-cyclic aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl; and

R₁ and R₂ are not α -amino acids.

In another aspect, the invention relates to a method of treatment and/or care of the skin, hair and/or mucous membranes which comprises the administration of a

cosmetically or pharmaceutically effective quantity of at least one compound of general formula (I):



its stereoisomers, mixtures thereof and/or its cosmetically or pharmaceutically acceptable salts, characterized in that :

AA₁ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₂ is selected from the group formed by -His-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₃ is selected from the group formed by -Leu-, -Ile-, -Phe- and -Tyr-;

AA₄ is selected from the group formed by -Lys-, -His-, -Arg-, -Gln-, -Leu-, -Ile-, -Met-, -MetO- and -MetO₂-;

AA₅ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₆ is selected from the group formed by -Phe-, -Trp-, -Ile- and -Val-;

AA₇ is selected from the group formed by -Met-, -MetO- and -MetO₂-;

W, X, Y, Z are amino acids and are independently selected from amongst themselves;

n, m, p and q are independently selected from amongst themselves and have a value of 0 or 1;

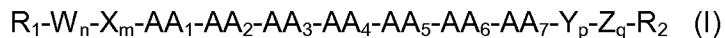
n+m+p+q is smaller or equal to 2;

R₁ is selected from the group formed by H, a polymer derived from polyethylene glycol, substituted or unsubstituted non-cyclic aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl and R₅-CO-, wherein R₅ is selected from the group formed by H, substituted or unsubstituted non-cyclic aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl, substituted or unsubstituted heterocyclyl and substituted or unsubstituted heteroarylalkyl;

R₂ is selected from the group formed by -NR₃R₄, -OR₃ and -SR₃, wherein R₃ and R₄ are independently selected from the group formed by H, a polymer derived from polyethylene glycol, substituted or unsubstituted non-cyclic aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl; and

R₁ and R₂ are not α -amino acids.

In another aspect the invention relates to a method of inhibition of neuronal exocytosis which comprises the administration of a cosmetically or pharmaceutically effective quantity of at least a compound of general formula (I):



its stereoisomers, mixtures thereof and/or its cosmetically or pharmaceutically acceptable salts, characterized in that :

AA₁ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₂ is selected from the group formed by -His-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₃ is selected from the group formed by -Leu-, -Ile-, -Phe- and -Tyr-;

AA₄ is selected from the group formed by -Lys-, -His-, -Arg-, -Gln-, -Leu-, -Ile-, -Met-, -MetO- and -MetO₂-;

AA₅ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₆ is selected from the group formed by -Phe-, -Trp-, -Ile- and -Val-;

AA₇ is selected from the group formed by -Met-, -MetO- and -MetO₂-;

W, X, Y, Z are amino acids and are independently selected from amongst themselves;

n, m, p and q are independently selected from amongst themselves and have a value of 0 or 1;

n+m+p+q is smaller or equal to 2 ;

- R₁ is selected from the group formed by H, a polymer derived from polyethylene glycol, substituted or unsubstituted non-cyclic aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl and R₅-CO-, wherein R₅ is selected from the group formed by H, substituted or unsubstituted non-cyclic aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl, substituted or unsubstituted heterocyclyl and substituted or unsubstituted heteroarylalkyl;
- R₂ is selected from the group formed by -NR₃R₄, -OR₃ and -SR₃, wherein R₃ and R₄ are independently selected from the group formed by H, a polymer derived from polyethylene glycol, substituted or unsubstituted non-cyclic aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl; and

R₁ and R₂ are not α -amino acids.

In accordance with a preferred embodiment, R₁ is selected from the group formed by H, a polymer derived from polyethylene glycol and R₅-CO-, is selected from the group formed by substituted or unsubstituted alkyl radical C₁-C₂₄, substituted or unsubstituted alkenyl C₂-C₂₄, substituted or unsubstituted alkynyl C₂-C₂₄, substituted or unsubstituted cycloalkyl C₃-C₂₄, substituted or unsubstituted cycloalkenyl C₅-C₂₄, substituted or unsubstituted cycloalkynyl C₈-C₂₄, substituted or unsubstituted aryl C₆-C₃₀, substituted or unsubstituted aralkyl C₇-C₂₄, substituted or unsubstituted heterocyclyl ring of 3-10 members, and substituted or unsubstituted heteroarylalkyl of 2 to 24 carbon atoms and 1 to 3 atoms other than carbon and an alkyl chain of 1 to 6 carbon atoms and R₅-CO is not an α -amino acid. More preferably, R₁ is selected from the group formed by H, a polymer derived from polyethylene glycol of a molecular weight comprised between 200 and 35000 Daltons, acetyl, *tert*-butanoyl, prenyl, hexanoyl, 2-methylhexanoyl, cyclohexanecarboxyl, octanoyl, decanoyl, lauroyl, myristoyl, palmitoyl, stearoyl, oleoyl and linoleoyl. Even more preferably, R₁ is H, acetyl, lauroyl, myristoyl or palmitoyl. In an even more preferable embodiment, R₁ is acetyl or palmitoyl.

In accordance with another preferred embodiment, R₂ is selected from the group formed by -NR₃R₄, -OR₃, -SR₃, wherein R₃ and R₄ are independently selected from the

group formed by H, a polymer derived from polyethylene glycol, substituted or unsubstituted alkyl C₁-C₂₄, substituted or unsubstituted alkenyl C₂-C₂₄, substituted or unsubstituted alkynyl C₂-C₂₄, substituted or unsubstituted cycloalkyl C₃-C₂₄, substituted or unsubstituted cycloalkenyl C₅-C₂₄, substituted or unsubstituted cycloalkynyl C₈-C₂₄,
5 substituted or unsubstituted aryl C₆-C₃₀, substituted or unsubstituted aralkyl C₇-C₂₄, substituted or unsubstituted heterocyclyl ring of 3-10 members, and substituted or unsubstituted heteroarylalkyl of 2 to 24 carbon atoms and 1 to 3 atoms other than carbon wherein the alkyl chain is of 1 to 6 carbon atoms and -NR₃R₄ is not an α-amino acid. Optionally, R₃ and R₄ can be bound by a saturated or unsaturated carbon-carbon
10 bond, forming a cycle with the nitrogen atom. More preferably R₂ is -NR₃R₄ or -OR₃. More preferably, R₃ and R₄ are selected from the group formed by H, a polymer derived from polyethylene glycol of a molecular weight comprised between 200 and 35000 Daltons, methyl, ethyl, hexyl, dodecyl and hexadecyl. Even more preferably R₃ is H and R₄ is selected from the group formed by H, methyl, ethyl, hexyl, dodecyl and
15 hexadecyl. In accordance with an even more preferred embodiment, R₂ is selected from -OH and -NH₂.

In accordance with another preferred embodiment AA₁ is selected from the group formed by -Arg-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-, AA₂ is selected from the group formed by -His-, -Gln-, -Asn-, -Glu- and -Asp-, AA₃ is selected from the group formed
20 by -Leu-and -Phe-, AA₄ is selected from the group formed by -His-, -Lys-, -Gln- and -Leu-, AA₅ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-, AA₆ is selected from the group formed by -Trp- and -Val- and AA₇ is selected from the group formed by -Met-, -MetO- and -MetO₂- . More preferably, AA₄ is selected from the group formed
25 by -Lys- and -Leu-, and AA₆ is -Trp-. Even more preferably, n, m, p and q are 0.

In accordance with another embodiment of this invention R₁ is selected from the group formed by H, acetyl, lauroyl, myristoyl and palmitoyl, AA₁ is selected from the group formed by -L-Arg-, -L-Lys-, -L-Gln-, -L-Asn-, -L-Glu- and -L-Asp-, AA₂ is selected from the group formed by -L-His-, -L-Gln-, -L-Asn-, -L-Glu- and -L-Asp-, AA₃ is selected from
30 the group formed by -L-Leu- and -L-Phe-, AA₄ is selected from the group formed by -L-His-, -L-Lys-, -L-Gln- and -L-Leu-, AA₅ is selected from the group formed by -L-Arg-, -L-His-, -L-Lys-, -L-Gln-, -L-Asn-, -L-Glu- and -L-Asp-, AA₆ is selected from the group formed by -L-Trp- and -L-Val-, AA₇ is selected from the group formed by -L-Met-, -L-MetO- and -L-MetO₂-, and R₂ is selected from the group formed

by -NR₃R₄ and -OR₃ where R₃ and R₄ are independently selected from H, methyl, ethyl, hexyl, dodecyl and hexadecyl. More preferably, AA₄ is selected from the group formed by -L-Lys- and -L-Leu-, and AA₆ is -L-Trp-. More preferably, R₁ is acetyl or palmitoyl and R₂ is -NH₂. Even more preferably, n, m, p and q are 0.

- 5 In accordance with another even more preferred embodiment R₁ is selected from the group formed by H, acetyl, lauroyl, myristoyl and palmitoyl, AA₁ is selected from the group formed by -L-Asn-, -L-Glu- and -L-Asp-, AA₂ is selected from the group formed by -L-His- and -L-Asp-, AA₃ is selected from the group formed by -L-Leu- and -L-Phe-, AA₄ is selected from the group formed by -L-Lys-, -L-Gln- and -L-Leu-, AA₅ is selected from the group formed by -L-Arg-, -L-Gln-, -L-Asn- and -L-Asp-, AA₆ is selected from the group formed by -L-Trp-, AA₇ is selected from the group formed by -L-Met-, -L-MetO- and -L-MetO₂- and R₂ is selected from the group formed by -NR₃R₄ and -OR₃ wherein R₃ and R₄ are independently selected from H, methyl, ethyl, hexyl, dodecyl and hexadecyl.
- 10 15 In accordance with another embodiment of this invention R₁ is selected from the group formed by H, acetyl, lauroyl, myristoyl and palmitoyl, AA₁ is -L-Glu-, AA₂ is selected from the group formed by -L-Asn-, -L-Glu-, -L-Gln- and -L-Asp-, AA₃ is -L-Leu-, AA₄ is -L-Lys-, AA₅ is -L-Arg-, AA₆ is -L-Trp-, AA₇ is selected from the group formed by -L-Met-, -L-MetO- and -L-MetO₂-, and R₂ is selected from the group formed by -NR₃R₄ and -OR₃ where R₃ and R₄ are independently selected from H, methyl, ethyl, hexyl, dodecyl and hexadecyl. More preferably, R₁ is acetyl or palmitoyl and R₂ is -NH₂. Even more preferably, n, m, p and q are 0.

In a preferred embodiment, the itching is selected from itching associated with conditions, diseases and/or disorders, for example and not restricted to, dermatitis, atopic dermatitis, contact dermatitis, diaper dermatitis, dermatitis herpetiformis, photodermatoses, photosensitivity, pregnancy related dermatitis, menopause related dermatitis, eczema, sensitive skin, psoriasis, chickenpox, herpes, herpes zoster, Netherton's syndrome, peeling skin syndrome, lichen planus, acne, dandruff, seborrhea, seborrheic dermatitis, alopecia, athlete's foot, candidiasis, hemorrhoids, vaginal itching, perianal itching, anogenital itching, sunburn, hives, pruritic otitis, itchy eyes, senile pruritus, aquagenic pruritus, prurigo nodularis, prurigo planus, pityriasis rosea, xerosis and dry skin, or itching associated with dialysis, infection from human immunodeficiency virus, malignant neoplasms, Hodgkin's disease, leukemia, myeloma, lymphoma, solid tumors, adenocarcinoma, lung cancer, hepatic diseases, jaundice,

cholestasis, liver failure, cirrhosis, polycythemia, hypereosinophilic syndrome, essential thrombocythemia, myelodysplastic syndrome, iron-deficiency anemia, systemic lupus erythematosus, endocrine diseases, thyroid diseases, hyperthyroidism, hypothyroidism, parathyroid diseases, diabetes mellitus, kidney diseases, kidney failure, uremia, parasitic diseases, scabies, lice, intestinal worms, allergic reactions, allergies to medicines, food allergies, allergies to chemical products, exposure to poisonous plants, exposure to insect bites, chemotherapy, stress and anxiety, among others.

In another particular embodiment, the pain is selected, for example and not restricted to, from the group formed by acute pain, chronic pain, nociceptive pain, neuropathic pain, inflammatory pain, visceral pain, abdominal pain, digestive system pain, respiratory system pain, urogenital system pain, endocrine system pain, heart pain, pancreatic pain, liver pain, pain due to gallstones, cholestasis, intestinal pain, stomach pain, pain due to duodenal ulcers, pain due to esophagitis, pain due to 10 gastroesophageal reflux, spleen pain, blood vessel pain, thalamic pain syndrome, irritable bowel syndrome, pain associated with Crohn's disease, pain associated with ulcerative colitis, diverticulitis, gastrointestinal mucositis, headache, tension-type headache, sinusitis-associated headache, migraine, eye pain, dry eye syndrome, post-operative pain, post-operative pain due to surgical incisions, post-operative pain due to 15 bone grafting, post-operative pain due to bone substitution, post-operative pain due to infections, post-operative pain due to limb amputations, pain due to bone fractures, pain due to cancer, pain due to bone cancer, pain associated with benign bone tumors, pain associated with osteoid osteomas, pain associated with osteoblastomas, pain due to cancer treatment, pain due to chemotherapy, pain due to emesis, pain due to emesis 20 as a consequence of chemotherapy treatment, musculoskeletal pain, spastic muscle pain, fibromyalgia, complex regional pain syndrome, psychogenic pain, neuralgic pain, pain due to demyelinating diseases, neck pain associated with cervical dystonia, back pain, lumbago, sciatica, neurogenic inflammation, neuritis, causalgia, touch sensitivity, sensitivity to cold, sensitivity to heat, cutaneous irritation, post-hair removal cutaneous 25 irritation, post-shaving cutaneous irritation, psoriasis, sensitive skin, dermatitis, atopic dermatitis, contact dermatitis, diaper dermatitis, seborrheic dermatitis, eczema, lichen planus, burns, sunburn, arthritis, rheumatoid arthritis, osteoarthritis, psoriatic arthritis, uveitis, pain due to nerve lesions, neuralgia, postherpetic neuralgia, neuropathies, peripheral neuropathies, phantom pains, allodynia, hyperalgesia, cold hyperalgesia, 30 pain due to carpal tunnel syndrome, burning pain, Grierson-Gopalan syndrome (better 35

known as burning feet syndrome), burning mouth syndrome, paresthesias, Fabry's disease, facial pain, trigeminal neuralgia, neuropathic pain due to diabetes, neuropathic pain due to AIDS, orofacial pain, dental pain, pain due to tooth extraction, pain due to wisdom tooth extraction, tooth sensitivity to cold, tooth sensitivity to heat, oral 5 mucositis, temporomandibular joint pain, joint pain caused by gout, pain associated with tattooing or tattoo removal processes, pain due to bunions, testicular pain, myofascial pain, urinary bladder pain, urinary tract pain, cystitis, pain due to kidney stones, renal colic, vulvar pain, vaginal pain, post-birth pain, menstrual pain, scrotal pain, perineal pain, pelvic pain or hypersensitivity, cutaneous pain or irritation after 10 surgery, after intense pulsed light treatment (IPL), after treatment with monochromatic pulsed light (laser), after a treatment with chemical flaking agents or after overexposure to external aggressive agents and pain due to chronic alcoholism, among others.

In another particular embodiment, inflammation is selected, for example and not restricted to, from the group formed by neurogenic inflammation, joint inflammation, 15 tendon inflammation, muscular inflammation, sepsis, vascular inflammation, respiratory inflammation, chronic obstructive pulmonary disease, rhinitis, allergic rhinitis, asthma, otitis, intestinal inflammation, Crohn's disease, pancreatitis, hepatitis, conditions related to chronic inflammation, to acute inflammation, nephritis, systemic lupus erythematosus, arthritis, rheumatoid arthritis, adult and juvenile rheumatoid arthritis, 20 Still's disease, psoriatic arthritis, osteoarthritis, arthritis caused by gout, rheumatoid spondylitis, glomerulonephritis, neuritis, inflammation of the nerve tissue, multiple sclerosis, immunological system disorders, Sjögren's syndrome, atherosclerosis, myocarditis, pericarditis, vasculitis, inflammatory skin conditions, psoriasis, sensitive skin, dermatitis, atopic dermatitis, contact dermatitis, diaper dermatitis, seborrheic 25 dermatitis, eczema, rosacea, acne, hyperproliferative skin disease, burns, sunburn, cutaneous inflammation after surgery, after intense pulsed light treatment (IPL), after treatment with monochromatic pulsed light (laser), after a treatment with chemical flaking agents or after overexposure to external aggressive agents, inflammation of the vaginal mucous membranes, vulvodynia, vaginitis, inflammation of the oral mucous 30 membranes, gingivitis, periodontitis, inflammatory eye diseases, uveitis, ocular and vernal conjunctivitis, sarcoidosis, peptic ulcers, urticaria, bullous pemphigoid, scleroderma, fibrosis, angioedema, anaphylaxis, alopecia, hepatic cirrhosis, restenosis, polymyalgia rheumatica, seronegative spondyloarthropathies, including ankylosing 35 spondylitis and Reiter's disease, dermatomyositis, inclusion body myositis, polymyositis and lymphangioleiomyomatosis, among others.

In another particular embodiment, the neurological, compulsive and neuropsychiatric diseases and/or disorders which are improved by inhibition of neuronal exocytosis are selected, for example and not restricted to, from the group formed by muscular spasticity, dystonia, and more specifically focal dystonia such as blepharospasm, 5 torsion dystonia, cervical dystonia or torticollis, laryngeal dystonia or spasmodic dysphonia, oromandibular dystonia, extremity dystonia such as writer's cramp, musician's cramp or foot dystonia, bruxism, facial scoliosis, hemifacial spasm, tics and/or strabismus; segmentary dystonia, Meige's syndrome, multifocal dystonia, hemidystonia, dopamine-responsive dystonia, Segawa's dystonia, trembling, 10 Parkinson's disease, nerve impingements, Alzheimer's disease and Tourette's syndrome, among others.

In another particular embodiment, the cosmetic, non-therapeutic treatment and/or care of the skin, is a treatment and/or prevention of aging and/or photoaging, treatment and/or prevention of wrinkles and/or expression wrinkles, treatment and/or prevention 15 of perspiration, treatment and/or care of skin disorders selected, for example and not restricted to, from the group formed by calluses and warts, among others. In an even more particular embodiment in its treatment and/or care of the facial skin.

In another particular embodiment, the treatment and/or care of the hair is a hair-growth stimulating and/or hair loss prevention treatment.

20 In another particular embodiment, the treatment and/or prevention of perspiration, and the treatment and/or prevention of hyperhidrosis, are treatments and/or preventions of perspiration or axillary, facial, genital, palmar or plantar hyperhidrosis.

In another aspect, the compounds of the invention can be administered by any means that causes contact between the compounds and the site of action in a mammal's body, 25 preferably that of a human being, and in the form of a composition which contains them. The administration of the compounds of this invention is carried out topically, transdermally, orally or parenterally. In a more particular aspect topical or transdermal application is carried out by iontophoresis, sonophoresis, electroporation, mechanical pressure, osmotic pressure gradient, occlusive cure, microinjections, by needle-free 30 injections by means of pressure, by microelectric patches, face masks or any combination thereof.

The frequency of the application can vary greatly, depending on the needs of each subject, with a recommendation of an application from once a month to ten times a day,

preferably from once a week to four times a day, more preferably from three times a week to twice a day, even more preferably once a day.

EXAMPLES OF ADMINISTRATION

- 5 The following specific examples provided here illustrate the nature of this invention. These examples are included for illustrative purposes only and should not be construed as limitations on the invention claimed herein.

General Methodology

Abbreviations

- 10 The abbreviations used for amino acids follow the 1983 IUPAC-IUB Joint Commission on Biochemical Nomenclature recommendations outlined in *Eur. J. Biochem.*, (1984), 138, 9-37.

®, resin; 2,6-diClZ, 2,6-dichlorobenzyl; 2-BrZ, 2-bromobenzylloxycarbonyl; 2-ClTrt-®, 2-chlorotriyl resin; Ac, acetyl; Adpoc, 1-(1-adamantyl)-1-methylethoxy-carbonyl: Ala, alanine; All, allyl; Alloc, allyloxycarbonyl; AM, 2-[4-aminomethyl-(2,4-dimethoxyphenyl)] phenoxyacetic acid; Arg, arginine; Asn, asparagine; Asp, aspartic acid; Boc, *tert*-butyloxycarbonyl; Bom, benzyloxymethyl; Brz, bromobenzylloxycarbonyl; BSA, bovine serum albumin; Bzl, benzyl; Cbz, benzyloxycarbonyl; cHx, cyclohexyl; ClZ, 2-chlorobenzyl; Cys, cysteine; C-terminal, carboxy-terminal; DCM, dichloromethane, 20 methylene chloride; Dde, *N*-[1-(4,4-dimethyl-2,6-dioxocyclohex-1-ylidene)ethyl; DIEA, *N,N*'-diisopropylethylamine; DIPCDI, *N,N*'-diisopropylcarbodiimide; Dmab, 4-(*N*-[1-(4,4-dimethyl-2,6-dioxocyclohexylidene)-3-methylbutyl]amino)benzyl; DMF, *N,N*-dimethylformamide; Dnp, 2,4-dinitrophenyl; EDTA, ethylenediaminetetraacetic acid; equiv, equivalent; ESI-MS, electrospray ionization mass spectrometry; Fm, fluorenylmethyl; Fmoc, 9-fluorenylmethyloxycarbonyl; For, formyl; Gln, glutamine; Glu, glutamic acid; Gly, glycine; GST, glutathione S-transferase; His, histidine; HOAt, 1-hydroxyazabenzotriazole; HOBt, 1-hydroxybenzotriazole; HPLC, high performance liquid chromatography; Ile, isoleucine; INCI, International Nomenclature of Cosmetic Ingredients; IPL, Intense Pulsed Light; ivDde, 30 1-(4,4-dimethyl-2,6-dioxocyclohexylidene)-3-methyl-butyl; LDMA-25, lauryldimonium hydrolyzed protein; Leu, leucine; Lys, lysine; MBHA, *p*-methylbenzhydrylamine; Me, methyl; MeCN, acetonitrile; MeOH, methanol; Met, methionine; MetO,

methionine(sulfoxide); MetO₂, methionine(sulfone); MLV, multilamellar vesicles; Mtr, 4-methoxy-2,3,6-trimethylbenzenesulfonyl; Mts, mesitylenesulfonyl; Mtt, methoxytrityl or methyltrityl; Myr, myristoyl; MW, molecular weight; N-terminal, amino terminal; PAL, 5-(4-aminomethyl-3,5-dimethoxyphenoxy)valeric acid; Palm, palmitoyl; Pbf, 5 2,2,4,6,7-pentamethyldihydrobenzofuran-5-sulfonyl; PEG, polyethylene glycol; Phe, phenylalanine; Pmc, 2,2,5,7,8-pentamethylchroman-6-sulfonyl; pNZ, p-nitrobenzyloxycarbonyl; Pro, proline; q.s., quantity sufficient; q.s.p., quantity sufficient for; SEM, standard error of the mean; Ser, serine; SNAP-25, synaptosomal-associated protein 25; SNARE, Soluble NSF Attachment Protein Receptor; tBu, *tert*-butyl; Teoc, 10 2-(trimethylsilyl)ethyloxycarbonyl; TFA, trifluoroacetic acid; THF, tetrahydrofuran; Thr, threonine; TIS, triisopropylsilane; Tos, tosyl or *p*-toluenesulfonyl; TPA, 12-O-tetradecanoylphorbol-13-acetate; Troc, 2,2,2-trichloroethoxycarbonyl; Trp, tryptophan; Trt, triphenylmethyl or trityl; Tyr, tyrosine; IUPAC, International Union of Pure and Applied Chemistry; IUB, International Union of Biochemistry; ULV, unilamellar 15 vesicles; UVA, ultraviolet radiation A; UVB, ultraviolet radiation B; Val, valine; VAMP, vesicle-associated membrane proteins; Xan, xanthyl; Z, benzyloxycarbonyl.

Chemical Synthesis

All synthetic processes were carried out in polypropylene syringes fitted with porous 20 polyethylene discs. All the reagents and solvents were synthesis quality and were used without any additional treatment. The solvents and soluble reagents were removed by suction. The Fmoc group was removed with piperidine-DMF (2:8, v/v) (1 x 1 min, 1 x 5 min; 5 mL/g resin) [LloydWilliams P. et al., "Chemical Approaches to the Synthesis of Peptides and Proteins", (1997), CRC, Boca Raton (FL, USA)]. Washes between stages 25 of deprotection, coupling, and, again, deprotection, were carried out with DMF (3 x 1 min) each time using 10 mL solvent/g resin. The coupling reactions were performed with 3 mL solvent/g resin. The control of the couplings was performed by carrying out the ninhydrin test [Kaiser E. et al., "Color test for detection of free terminal amino groups in the solid-phase synthesis of peptides", (1970), *Anal. Biochem.*, 34(2), 595- 30 598] or chloranil test [Christensen T., "A Qualitative Test for Monitoring Coupling Completeness in Solid Phase Peptide Synthesis Using Chloranil", (1979), *Acta Chem. Scand.*, 33B, 763-766]. All synthetic reactions and washes were carried out at 25 °C.

The HPLC chromatographic analysis was carried out with a Shimadzu equipment (Kyoto, Japan) using a reversed-phase column thermostatized at 30°C (250 x 4.0 mm,

Kromasil C₈, 5 µm, Akzo Nobel, Sweden). The elution was carried out using a gradient of acetonitrile (+0.07% TFA) in water (+0.1% TFA) at a flow rate of 1 mL/min and detection was carried out at 220 nm. The electrospray ionization mass spectrometry was carried out in a WATERS Alliance ZQ 2000 detector using a mixture of MeCN:H₂O 5 4:1 (+0.1% TFA) as the mobile phase and a flow rate of 0.2 mL/min.

EXAMPLE 1

10 *Obtaining Fmoc-W_n-X_m-AA₁-AA₂-AA₃-AA₄-AA₅-AA₆-AA₇-Y_p-Z_q-AM-MBHA-®, wherein AA₁ is -L-Glu- or -L-Asp-; AA₂ is -L-Asn-, -L-Glu- or -L-Asp-; AA₃ is -L-Leu-, -L-Val- or -L-Met-; AA₄ is -L-Lys- or -L-His-; AA₅ is -L-Lys-, -L-His- or -L-Arg-; AA₆ is -L-Tyr- or -L-Trp-; AA₇ is -L-Met- or -L-MetO-; and n, m, p and q are 0.*

15 5 mmol (1 equiv) of the Fmoc-AM-MBHA resin with a functionalization of 0.73 mmol/g were treated with piperidine-DMF according to the described general protocol in order to remove the Fmoc group. 2.5 equiv of Fmoc-L-Met-OH or Fmoc-L-MetO-OH were incorporated onto the deprotected resin in the presence of 2.5 equiv of DIPCDI and 2.5 equiv of HOBt, using DMF as a solvent, for 1 hour.

The resins were then washed as described in the general methods and the deprotection treatment of the Fmoc group was repeated to couple the next amino acid. The Fmoc N-terminal group was deprotected as described in the general methods and 20 2.5 equiv of Fmoc-L-Tyr(tBu)-OH or Fmoc-L-Trp(Boc)-OH; 2.5 equiv of Fmoc-L-Lys(Boc)-OH-, Fmoc-L-His(Trt)-OH or Fmoc-L-Arg(Pbf)-OH; 2.5 equiv of Fmoc-L-Lys(Boc)-OH- or Fmoc-L-His(Trt)-OH; 2.5 equiv of Fmoc-L-Leu-OH, Fmoc-L-Val-OH or Fmoc-L-Met-OH; 2.5 equiv of Fmoc-L-Asn(Trt)-OH, Fmoc-L-Glu(OtBu)-OH or Fmoc-L-Asp(OtBu)-OH and finally 2.5 equiv of 25 Fmoc-L-Glu(OtBu)-OH or Fmoc-L-Asp(OtBu)-OH were sequentially coupled onto the peptidyl resins for 1 hour, each coupling in the presence of 2.5 equiv of HOBt and 2.5 equiv of DIPCDI and using DMF as a solvent.

After the synthesis, the peptidyl resins were washed with DCM (5 x 3 min) and dried under nitrogen stream.

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EXAMPLE 2

Prophetic synthesis of Fmoc-W_n-X_m-AA₁-AA₂-AA₃-AA₄-AA₅-AA₆-AA₇-Y_p-Z_q-O-2-CITrt-®, wherein AA₁ is -L-Glu- or -L-Asp-; AA₂ is -L-Gln-, -L-Asn-, -L-Glu- or -L-Asp-; AA₃

is -L-Leu-, -L-Ile-, -L-Val-, -L-Met-, -L-MetO- or -L-MetO₂-, AA₄ is -L-Lys-, -L-His- or -L-Arg-; AA₅ is -L-Lys-, -L-His- or -L-Arg-; AA₆ is -L-Phe-, -L-Tyr- or -L-Trp-; AA₇ is -L-Met-, -L-MetO- or -L-MetO₂-, and n, m, p and q are 0.

- 5 8.8 mmol (1 equiv) of Fmoc-L-Met-OH, Fmoc-L-MetO-OH or Fmoc-L-MetO₂-OH dissolved in 55 mL of DCM to which 0.86 equiv of DIEA are added, are coupled to the dry 2-chlorotriptyl resin (5.5 g; 8.8 mmol). They are stirred for 5 min, after which 1.66 equiv of DIEA are added. The mixture is allowed to react for 40 min. The remaining chloride groups are blocked by treatment with 4.4 mL of MeOH.
- 10 The N-terminal Fmoc group is deprotected as described in the general methods and 2.5 equiv of Fmoc-L-Phe-OH, Fmoc-L-Tyr(tBu)-OH or Fmoc-L-Trp(Boc)-OH are coupled onto the peptidyl resins in presence of 2.5 equiv of DIPCDI and 2.5 equiv of HOBr, using DMF as a solvent, for 1 hour. The resins are then washed as described in the general methods and the deprotection treatment of the Fmoc group is repeated to 15 couple the next amino acid. Following the protocols described, 2.5 equiv of Fmoc-L-Lys(Boc)-OH, Fmoc-L-His(Trt)-OH or Fmoc-L-Arg(Pbf)-OH; 2.5 equiv of Fmoc-L-Lys(Boc)-OH, Fmoc-L-His(Trt)-OH or Fmoc-L-Arg(Pbf)-OH; 2.5 equiv of Fmoc-L-Leu-OH, Fmoc-L-Ile-OH, Fmoc-L-Val-OH, Fmoc-L-Met-OH, Fmoc-L-MetO-OH or Fmoc-L-MetO₂-OH; 2.5 equiv of Fmoc-L-Gln(Trt)-OH, Fmoc-L-Asn(Trt)-OH, 20 Fmoc-L-Glu(OtBu)-OH or Fmoc-L-Asp(OtBu)-OH and finally 2.5 equiv of Fmoc-L-Glu(OtBu)-OH or Fmoc-L-Asp(OtBu)-OH are sequentially coupled in the presence of 2.5 equiv of HOBr and 2.5 equiv of DIPCDI in each coupling.

After the synthesis, the peptidyl resins are washed with DCM (5 x 3 min) and dried under nitrogen stream.

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EXAMPLE 3

Obtaining Fmoc-W_n-X_m-AA₁-AA₂-AA₃-AA₄-AA₅-AA₆-AA₇-Y_p-Z_q-AM-MBHA-®, wherein AA₁ is -L-Arg-, -L-Lys-, -L-Gln-, -L-Asn-, -L-Glu- or -L-Asp-; AA₂ is -L-His- or -L-Asp-; AA₃ is -L-Leu-, -L-Met- or -L-Phe-; AA₄ is -L-Lys-, -L-His-, -L-Gln- or -L-Leu-; AA₅ is -L-Arg-, -L-His-, -L-Lys-, -L-Gln-, -L-Asn-, -L-Glu- or -L-Asp-; AA₆ is -L-Tyr-, -L-Trp-, -L-Val- or -L-Arg-; AA₇ is -L-Met- or -L-MetO-; and n, m, p and q are 0.

5 mmol (1 equiv) of the Fmoc-AM-MBHA resin with a functionalization of 0.73 mmol/g were treated with piperidine-DMF according to the described general protocol in order to remove the Fmoc group. 2.5 equiv of Fmoc-L-Met-OH or Fmoc-L-MetO-OH were incorporated onto the deprotected resin in the presence of 2.5 equiv of DIPCDI and 2.5 equiv of HOBr, using DMF as a solvent, for 1 hour.

The resins were then washed as described in the general methods and the deprotection treatment of the Fmoc group was repeated to couple the next amino acid. The Fmoc N-terminal group was deprotected as described in the general methods and 2.5 equiv of Fmoc-L-Tyr(tBu)-OH, Fmoc-L-Trp(Boc)-OH, Fmoc-L-Val-OH or 10 Fmoc-L-Arg(Pbf)-OH; 2.5 equiv de Fmoc-L-Arg(Pbf)-OH, Fmoc-L-His(Trt)-OH, Fmoc-L-Lys(Boc)-OH, Fmoc-L-Gln(Trt)-OH, Fmoc-L-Asn(Trt)-OH, Fmoc-L-Glu(OtBu)-OH or Fmoc-L-Asp(OtBu)-OH; 2.5 equiv of Fmoc-L-Lys(Boc)-OH-, Fmoc-L-His(Trt)-OH, Fmoc-L-Gln(Trt)-OH or Fmoc-L-Leu-OH,; 2.5 equiv of 15 Fmoc-L-Leu-OH, Fmoc-L-Met-OH or Fmoc-L-Phe-OH; 2.5 equiv of Fmoc-L-His(Trt)-OH or Fmoc-L-Asp(OtBu)-OH and finally 2.5 equiv of Fmoc-L-Arg(Pbf)-OH, Fmoc-L-Lys(Boc)-OH, Fmoc-L-Gln(Trt)-OH, Fmoc-L-Asn(Trt)-OH, Fmoc-L-Glu(OtBu)-OH or Fmoc-L-Asp(OtBu)-OH were sequentially coupled onto the peptidyl resins for 1 hour, each coupling in the presence of 2.5 equiv of HOBr and 2.5 equiv of DIPCDI and using DMF as a solvent.

20 After the synthesis, the peptidyl resins were washed with DCM (5 x 3 min) and dried under nitrogen stream.

EXAMPLE 4

Prophetic synthesis of Fmoc-W_n-X_m-AA₁-AA₂-AA₃-AA₄-AA₅-AA₆-AA₇-Y_p-Z_q-O-2-CITrt-®,
25 *wherein AA₁ is -L-Arg-, -L-His-, -L-Lys-, -L-Gln-, -L-Asn-, -L-Glu- or -L-Asp-; AA₂ is -L-His-, -L-Lys- or -L-Asp-; AA₃ is -L-Leu-, -L-Ile-, -L-Met-, -L-MetO-, -L-MetO₂-, -L-Phe- or -L-Tyr-; AA₄ is -L-Lys-, -L-His-, -L-Arg-, -L-Gln-, -L-Asn-, -L-Leu-, -L-Ile-, -L-Met-, -L-MetO-, -L-MetO₂- or -L-Val-; AA₅ is -L-Arg-, -L-His-, -L-Lys-, -L-Gln-, -L-Asn-, -L-Glu- or -L-Asp-; AA₆ is -L-Phe-, -L-Tyr-, -L-Trp-, -L-Met-, -L-MetO-, -L-MetO₂-, -L-Ile-, -L-Val-, -L-His-, -L-Lys- or -L-Arg-; AA₇ is -L-Met-, -L-MetO- or -L-MetO₂-; and n, m, p and q are 0.*

8.8 mmol (1 equiv) of Fmoc-L-Met-OH, Fmoc-L-MetO-OH or Fmoc-L-MetO₂-OH dissolved in 55 mL of DCM to which 0.86 equiv of DIEA are added, are coupled to the

dry 2-chlorotriyl resin (5.5 g; 8.8 mmol). They are stirred for 5 min, after which 1.66 equiv of DIEA are added. The mixture is allowed to react for 40 min. The remaining chloride groups are blocked by treatment with 4.4 mL of MeOH.

The *N*-terminal Fmoc group is deprotected as described in the general methods and 5 2.5 equiv of Fmoc-L-Phe-OH, Fmoc-L-Tyr(tBu)-OH, Fmoc-L-Trp(Boc)-OH, Fmoc-L-Met-OH, Fmoc-L-MetO-OH, Fmoc-L-MetO₂-OH, Fmoc-L-Ile-OH, Fmoc-L-Val-OH, Fmoc-L-His(Trt)-OH, Fmoc-L-Lys(Boc)-OH or Fmoc-L-Arg(Pbf)-OH are coupled onto the peptidyl resins in presence of 2.5 equiv of DIPCDI and 2.5 equiv 10 of HOBr, using DMF as a solvent, for 1 hour. The resins are then washed as described in the general methods and the deprotection treatment of the Fmoc group is repeated 15 to couple the next amino acid. Following the protocols described, 2.5 equiv of Fmoc-L-Arg(Pbf)-OH, Fmoc-L-His(Trt)-OH, Fmoc-L-Lys(Boc)-OH, Fmoc-L-Gln(Trt)-OH, Fmoc-L-Asn(Trt)-OH, Fmoc-L-Glu(OtBu)-OH or Fmoc-L-Asp(OtBu)-OH; 2.5 equiv of 20 Fmoc-L-Lys(Boc)-OH, Fmoc-L-His(Trt)-OH, Fmoc-L-Arg(Pbf)-OH, Fmoc-L-Gln(Trt)-OH, Fmoc-L-Asn(Trt)-OH, Fmoc-L-Leu-OH, Fmoc-L-Ile-OH, Fmoc-L-Met-OH, Fmoc-L-MetO-OH, Fmoc-L-MetO₂-OH or Fmoc-L-Val-OH; 2.5 equiv of 25 Fmoc-L-Leu-OH, Fmoc-L-Ile-OH, Fmoc-L-Met-OH, Fmoc-L-MetO-OH, Fmoc-L-MetO₂-OH, Fmoc-L-Phe-OH or Fmoc-L-Tyr(tBu)-OH; 2.5 equiv of Fmoc-L-His(Trt)-OH, Fmoc-L-Lys(Boc)-OH or Fmoc-L-Asp(OtBu)-OH and finally 2.5 equiv of Fmoc-L-Arg(Pbf)-OH, Fmoc-L-His(Trt)-OH, Fmoc-L-Lys(Boc)-OH, Fmoc-L-Gln(Trt)-OH, Fmoc-L-Asn(Trt)-OH, Fmoc-L-Glu(OtBu)-OH or Fmoc-L-Asp(OtBu)-OH are sequentially coupled in the presence of 2.5 equiv of HOBr and 2.5 equiv of DIPCDI in each coupling.

After the synthesis, the peptidyl resins are washed with DCM (5 x 3 min) and dried 25 under nitrogen stream.

EXAMPLE 5

*General process for cleavage of Fmoc *N*-terminal protective group.*

The *N*-terminal Fmoc group of the peptidyl resins obtained in examples 1 and 3 was 30 deprotected as described in the general methods (20% piperidine in DMF, 1 x 5 min + 1 x 20 min). The peptidyl resins were washed with DMF (5 x 1 min), DCM (4 x 1 min), diethyl ether (4 x 1 min) and dried under vacuum.

Prophetically, using the same protocol the *N*-terminal Fmoc group of the peptidyl resins obtained in examples 2 and 4 are deprotected, washed and dried.

EXAMPLE 6

- 5 *Process for introducing the R₁ palmitoyl group onto the peptidyl resins obtained in Example 5 for SEQ ID NO:25.*

10 equiv of palmitic acid pre-dissolved in DMF (1 mL) were added onto 1 mmol (1 equiv) of the peptidyl resins obtained in Example 5 for SEQ ID NO:25, in the presence of 10 equiv of HOBr and 10 equiv of DIPCDI. They were allowed to react for 15 hours, 10 after which the resins were washed with THF (5 x 1 min), DCM (5 x 1 min), DMF (5 x 1 min), MeOH (5 x 1 min), DMF (5 x 1 min) THF (5 x 1 min), DMF (5 x 1 min), DCM (4 x 1 min), ether (3 x 1 min), and were dried under vacuum.

Prophetically, following the same protocol the R₁ palmitoyl group is introduced onto the other peptidyl resins obtained in Examples 1 to 4 after deprotecting the *N*-terminal 15 following the protocol described in Example 5.

EXAMPLE 7

- Process for introducing the R₁ acetyl group onto the peptidyl resins obtained in Example 5.*

20 1 mmol (1 equiv) of the peptidyl resins obtained in Example 5 was treated with 25 equiv of acetic anhydride in the presence of 25 equiv of DIEA, using 5 mL of DMF as a solvent. They were allowed to react for 30 min, after which the peptide resins were washed with DMF (5 x 1 min), DCM (4 x 1 min), diethyl ether (4 x 1 min) and were dried under vacuum.

25 Prophetically, following the same protocol the R₁ acetyl group is introduced onto the peptidyl resins prophetically obtained in Examples 5.

EXAMPLE 8

- Cleavage process of the peptidyl resins obtained in Examples 6 and 7 from the*

30 *polymeric support.*

200 mg of the dried peptidyl resins obtained in Examples 6 and 7 were treated with 5 mL of TFA:TIS:H₂O (90:5:5) for 2 hours at room temperature under stirring. The filtrates were collected onto 50 mL cold diethyl ether, filtered through polypropylene syringes fitted with porous polyethylene discs, and washed 5 times with 50 mL diethyl ether. The final precipitates were dried under vacuum.

HPLC analysis of the obtained peptides in gradients of MeCN (+0.07% TFA) in H₂O (+0.1% TFA) showed a purity exceeding 80% in all cases. The identity of the peptides obtained was confirmed by ESI-MS.

Prophetically, following the same protocol the cleavage from the polymeric support is performed for the other peptidyl resins prophetically obtained in Examples 5 to 7.

EXAMPLE 9

Prophetic cleavage process from the polymeric support and functionalization with R₂ substituted amine: Obtaining Ac-W_n-X_m-AA₁-AA₂-AA₃-AA₄-AA₅-AA₆-AA₇-Y_p-Z_q-NH-(CH₂)₁₅-CH₃, wherein AA₁ is -L-Glu- or -Asp-; AA₂ is -L-Gln-, -L-Asn-, -L-Glu- or -L-Asp-; AA₃ is -L-Leu-, -L-Ile-, -L-Val-, -L-Met-, -L-MetO-, -L-MetO₂-; AA₄ is -L-Lys-, -L-His- or -L-Arg-; AA₅ is -L-Lys-, -L-His- or -L-Arg-; AA₆ is -L-Phe-, -L-Tyr- or -L-Trp-; AA₇ is -L-Met-, -L-MetO- or -L-MetO₂-; and n, m, p and q are 0.

The compounds Ac-W_n-X_m-AA₁-AA₂-AA₃-AA₄-AA₅-AA₆-AA₇-Y_p-Z_q-OH with fully protected side chains are obtained by treating 150 mg of the peptidyl resins Ac-W_n-X_m-AA₁-AA₂-AA₃-AA₄-AA₅-AA₆-AA₇-Y_p-Z_q-O-2-CITrt-® from Example 7, previously desiccated under vacuum in the presence of KOH, with 3 mL of a 3% solution of TFA in DCM for 5 min. The filtrates are collected onto 50 mL of cold diethyl ether and the treatment is repeated three times. The ether solutions are evaporated to dryness at reduced pressure and room temperature, the precipitates are redissolved in 50% of MeCN in H₂O and they are lyophilized. 10 mg of the obtained crude peptides are weighed in a flask and 3 equiv of hexadecylamine and 25 mL of anhydrous DMF are added. 2 equiv of DIPCDI are added and allowed to react under magnetic stirring at 47 °C. The reactions are monitored by HPLC until disappearance of the initial products. The solvents are evaporated to dryness and co-evaporated twice with DCM. The obtained residues [Ac-W_n-X_m-AA₁-AA₂-AA₃-AA₄-AA₅-AA₆-AA₇-Y_p-Z_q-NH-(CH₂)₁₅-CH₃ with fully protected side chains] are redissolved in 25 mL of a mixture of TFA-DCM-

anisole (49:49:2) and allowed to react for 30 min at room temperature. 250 mL of cold diethyl ether are added, the solvents are evaporated under reduced pressure and two additional co-evaporations with ether are carried out. The residues are dissolved in a mixture of 50% MeCN in H₂O and lyophilized.

- 5 Purity of the obtained peptides is determined by HPLC analysis in gradients of MeCN (+0.07% TFA) in H₂O (+0.1% TFA). The identity of the peptides obtained is confirmed by ESI-MS.

EXAMPLE 10

- 10 *Prophetic cleavage process from the polymeric support and functionalization with R₂ substituted amine: Obtaining Ac-W_n-X_m-AA₁-AA₂-AA₃-AA₄-AA₅-AA₆-AA₇-Y_p-Z_q-NH-(CH₂)₁₅-CH₃, wherein AA₁ is -L-Arg-, -L-His-, -L-Lys-, -L-Gln-, -L-Asn-, -L-Glu- or -Asp-; AA₂ is -L-His-, -L-Lys- or -L-Asp-; AA₃ is -L-Leu-, -L-Ile-, -L-Met-, -L-MetO-, -L-MetO₂-, -L-Phe- or -L-Tyr-; AA₄ is -L-Lys-, -L-His-, -L-Arg-, -L-Gln-, -L-Asn-, -L-Leu-, -L-Ile-, -L-Met-, -L-MetO-, -L-MetO₂- or -L-Val-; AA₅ is -L-Arg-, -L-His-, -L-Lys-, -L-Gln-, -L-Asn-, -L-Glu- or -L-Asp-; AA₆ is -L-Phe-, -L-Tyr-, -L-Trp-, -L-Met-, -L-MetO-, -L-MetO₂-, -L-Ile-, -L-Val-, -L-His-, -L-Lys- or -L-Arg-; AA₇ is -L-Met-, -L-MetO- or -L-MetO₂-; and n, m, p and q are 0.*

- 20 The compounds Ac-W_n-X_m-AA₁-AA₂-AA₃-AA₄-AA₅-AA₆-AA₇-Y_p-Z_q-OH with fully protected side chains are obtained by treating 150 mg of the prophetic peptidyl resins Ac-W_n-X_m-AA₁-AA₂-AA₃-AA₄-AA₅-AA₆-AA₇-Y_p-Z_q-O-2-CITr-® from Example 7, previously desiccated under vacuum in the presence of KOH, with 3 mL of a 3% solution of TFA in DCM for 5 min. The filtrates are collected onto 50 mL of cold diethyl ether and the treatment is repeated three times. The ether solutions are evaporated to dryness at reduced pressure and room temperature, the precipitates are redissolved in 50% of MeCN in H₂O and they are lyophilized. 10 mg of the obtained crude peptides are weighed in a flask and 3 equiv of hexadecylamine and 25 mL of anhydrous DMF are added. 2 equiv of DIPCDI are added and allowed to react under magnetic stirring at 47° C. The reactions are monitored by HPLC until disappearance of the initial products.
- 25 The solvents are evaporated to dryness and co-evaporated twice with DCM. The obtained residues [Ac-W_n-X_m-AA₁-AA₂-AA₃-AA₄-AA₅-AA₆-AA₇-Y_p-Z_q-NH-(CH₂)₁₅-CH₃ with fully protected side chains] are redissolved in 25 mL of a mixture of TFA-DCM-anisole (49:49:2) and allowed to react for 30 min at room temperature. 250 mL of cold diethyl ether are added, the solvents are evaporated under reduced pressure and two

additional co-evaporations with ether are carried out. The residues are dissolved in a mixture of 50% MeCN in H₂O and lyophilized.

Purity of the obtained peptides is determined by HPLC analysis in gradients of MeCN (+0.07% TFA) in H₂O (+0.1% TFA). The identity of the peptides obtained is confirmed
5 by ESI-MS.

EXAMPLE 11

Following the protocols described in examples 1 to 10, routinely varying the nature of the reagents and the peptide sequences, the following compounds included in the
10 scope of this invention were also obtained.

Identifier	Average MW	Experimental MW
Ac-SEQ ID NO:1-NH ₂	1012.20	1012.13 \pm 0.11
Ac-SEQ ID NO:2-NH ₂	1017.18	1017.88 \pm 0.92
Ac-SEQ ID NO:3-NH ₂	919.08	918.47 \pm 0.83
Ac-SEQ ID NO:8-NH ₂	1012.24	1012.52 \pm 0.29
Ac-SEQ ID NO:9-NH ₂	999.16	999.19 \pm 0.05
Ac-SEQ ID NO:12-NH ₂	1012.20	1012.23 \pm 0.05
Ac-SEQ ID NO:13-NH ₂	1058.28	1057.81 \pm 0.72
Ac-SEQ ID NO:14-NH ₂	1018.21	1018.08 \pm 0.50
Ac-SEQ ID NO:15-NH ₂	1002.16	1001.10 \pm 1.88
Ac-SEQ ID NO:16-NH ₂	1026.23	1026.40 \pm 0.18
Ac-SEQ ID NO:21-NH ₂	1016.24	1016.10 \pm 0.20
Ac-SEQ ID NO:22-NH ₂	998.22	998.29 \pm 0.15
Ac-SEQ ID NO:23-NH ₂	962.09	961.63 \pm 0.68
Ac-SEQ ID NO:24-NH ₂	1002.12	1002.13 \pm 0.04
Ac-SEQ ID NO:25-NH ₂	1004.13	1003.44 \pm 0.78
Palm-SEQ ID NO:25-NH ₂	1200.51	1200.46 \pm 0.21
Ac-SEQ ID NO:26-NH ₂	1017.18	1017.07 \pm 0.15
Ac-SEQ ID NO:27-NH ₂	997.19	997.51 \pm 0.55
Ac-SEQ ID NO:30-NH ₂	1031.20	1031.22 \pm 0.19

Ac-SEQ ID NO:31-NH ₂	983.16	984.16 ± 1.01
Ac-SEQ ID NO:32-NH ₂	1006.20	1006.23 ± 0.14
Ac-SEQ ID NO:33-NH ₂	1058.28	1057.71 ± 0.72
Ac-SEQ ID NO:34-NH ₂	1003.15	1001.52 ± 2.22
Ac-SEQ ID NO:50-NH ₂	1028.20	1028.14 ± 0.10
Ac-SEQ ID NO:53-NH ₂	1034.20	1034.14 ± 0.09
Ac-SEQ ID NO:55-NH ₂	1018.12	1018.24 ± 0.49
Ac-SEQ ID NO:56-NH ₂	1020.13	1020.59 ± 1.72
Ac-SEQ ID NO:116-NH ₂	1032.23	1032.18 ± 0.12
Ac-SEQ ID NO:117-NH ₂	1017.22	1017.18 ± 0.07

Table 3

EXAMPLE 12

5 *Study of the inhibition of the SNARE complex formation with detection of the complex by ELISA*

With the aim of determining the capacity of inhibition of the SNARE complex formation by the compounds of the invention, the competitive inhibition of the compounds compared to SNAP-25 was studied with regards to the formation of this complex. The proportion of SNARE complex formed was determined by the ELISA technique, using 10 one of the proteins from the complex bound to GST. In a 96-well plate VAMP was immobilized (using a 0.037 µM solution) and subsequently the free spaces were blocked with BSA (3%). Parallel to this process, SNAP-25 bound to GST (0.0185 µM), syntaxin (0.037 µM) and a compound of the invention (2.5 mM and 0.5 mM compounds) were incubated for 1 hour. After incubation, the samples were transferred 15 to the plate with immobilized VAMP and were incubated for 1 hour to allow the formation of the SNARE complex. Afterward, the plate was washed and the complex was detected by a primary antibody anti-GST (Antibody anti-GST epitope TAG, Fisher Cat. no:PA1-982A). The reading was carried out at a wavelength of 490 nm in a TECAN GENios spectrophotometric reader.

20 Table 4 shows the results of the competitive inhibition of the formation of the SNARE complex by the compounds of the invention versus SNAP-25. The percentage of

inhibition of the formation of the complex is inversely proportional to the quantity of SNARE complex spectrophotometrically detected.

Compound	% inhibition formation SNARE complex	
	Concentration of compound 2.5 mM	0.5 mM
Ac-SEQ ID NO:1-NH ₂	63	33
Ac-SEQ ID NO:2-NH ₂	64	26
Ac-SEQ ID NO:3-NH ₂	64	31
Ac-SEQ ID NO:8-NH ₂	44	11
Ac-SEQ ID NO:12-NH ₂	71	20
Ac-SEQ ID NO:13-NH ₂	20	Not studied
Ac-SEQ ID NO:14-NH ₂	67	43
Ac-SEQ ID NO:15-NH ₂	42	48
Ac-SEQ ID NO:22-NH ₂	72	18
Ac-SEQ ID NO:23-NH ₂	60	10
Ac-SEQ ID NO:24-NH ₂	55	41
Ac-SEQ ID NO:25-NH ₂	71	40
Ac-SEQ ID NO:27-NH ₂	14	Not studied
Ac-SEQ ID NO:30-NH ₂	71	24
Ac-SEQ ID NO:31-NH ₂	59	26
Ac-SEQ ID NO:32-NH ₂	19	Not studied
Ac-SEQ ID NO:34-NH ₂	85	27
Ac-SEQ ID NO:56-NH ₂	Not studied	9
Ac-SEQ ID NO:116-NH ₂	69	
Ac-SEQ ID NO:117-NH ₂	65	

Table 4

5 EXAMPLE 13

Study of the inhibition of the SNARE complex formation with detection of the complex by electrophoresis

VAMP (6 μ M), syntaxin (6 μ M) and the compound of the invention (5 mM or 1 mM compounds) were incubated for 3 hours. Subsequently, SNAP-25 (0.6 μ M) was added

and the mixture was incubated for an additional 15 hours to allow the formation of the SNARE complex. After incubation, the loading buffer (Laemli Simple Buffer) was added and the mixture was analyzed by 10% acrylamide SDS-PAGE in gel. The amount of complex was determined by MediaCybernetics image analysis software Image-Pro Plus.

Table 5 shows the results of the inhibition of the formation of the SNARE complex. The percentage of inhibition of the formation of the complex is inversely proportional to the quantity of SNARE complex detected.

Compound	% inhibition formation SNARE complex	
	Concentration of compound 5 mM	1 mM
Ac-SEQ ID NO:1-NH ₂	100	59
Ac-SEQ ID NO:2-NH ₂	80	37
Ac-SEQ ID NO:14-NH ₂	100	83
Ac-SEQ ID NO:15-NH ₂	100	50
Ac-SEQ ID NO:24-NH ₂	100	48
Ac-SEQ ID NO:25-NH ₂	100	83
Ac-SEQ ID NO:34-NH ₂	59	14

10

Table 5

EXAMPLE 14

Quantification of the release of noradrenaline induced by TPA/ionomycin in a neuroblastoma cell line by ELISA

The induction of the release of noradrenaline with TPA (12-O-tetradecanoylphorbol-13-acetate)/ionomycin enables direct measurement of neuronal exocytosis. For the study of the inhibitory effect of the compounds of the invention on the release of noradrenaline, cells of a human neuroblastoma cell line were pre-incubated (1×10^6 cells/well) for 60 minutes with the compound of the invention (1 μ M, 10 μ M, 100 μ M, 500 μ M or 1 mM compounds), and afterward the release of noradrenaline was induced. The release of the noradrenaline neurotransmitter was induced by an 8 minute pre-treatment with a solution of 12-O-tetradecanoylphorbol-13-acetate (TPA) 100 nM, which mobilized the intracellular vesicles which contained the neurotransmitter; followed by a 5 minute incubation with TPA/ionomycin (100 nM/10 μ M), which induced

the release of the neurotransmitter contained in these vesicles. The quantity of neurotransmitter released into the growth medium was quantified by ELISA (Noradrenaline ELISA kit, IBL International ref. RE59261), in an assay mediated by specific antibodies against noradrenaline and completed by an enzymatic reaction 5 based on the reaction of alkaline phosphatase, which resulted in a quantifiable color indication. For this, absorbance at 405 nm was measured in Thermo Scientific Multiskan Ascent equipment.

The blocking of the SNARE complex by the compounds of the invention lead to an inhibition of neuronal exocytosis and therefore, a decrease in the levels of released 10 noradrenaline (Table 6).

TREATMENT	DOSIS	% RELEASED NORADRENALINE	
		Average	SEM
TPA/ION		100.00	3.11
Ac-SEQ ID NO:1-NH ₂	1 µM	96.14	5.61
	10 µM	93.07	3.40
	100 µM	79.59	2.04
	500 µM	49.08	1.49
	1000 µM	42.48	0.89
Ac-SEQ ID NO:24-NH ₂	1 µM	90.63	2.38
	10 µM	97.06	1.76
	100 µM	88.95	2.15
	500 µM	78.11	2.92
	1000 µM	67.90	2.79
Ac-SEQ ID NO:25-NH ₂	1 µM	99.59	1.81
	10 µM	94.48	1.76
	100 µM	74.79	3.66
	500 µM	48.07	3.79
	1000 µM	41.05	1.84
Ac-SEQ ID NO:14-NH ₂	10 µM	84.96	7.24
	100 µM	85.97	8.38
	1000 µM	45.78	2.08
Ac-SEQ ID NO:116-NH ₂	1 µM	87.59	4.72
	10 µM	83.30	4.58
	100 µM	62.24	3.53
	500 µM	52.82	2.26
	1000 µM	43.10	1.22
Ac-SEQ ID NO:117-NH ₂	1 µM	87.99	5.20
	10 µM	83.11	4.82
	100 µM	70.76	2.77

	500 μ M	58.37	2.82
	1000 μ M	54.64	3.53

Table 6

EXAMPLE 15

Preparation of a cosmetic facial composition containing Ac-SEQ ID NO:1-NH₂.

- 5 In a suitable vessel the components of phase A were dissolved. Next, carbomer was slowly added (INCI: CARBOMER) (phase A1), under stirring, until it was completely dissolved. The mixture was heated to 70-75 °C.

In another vessel the components of phase B were mixed together and the mixture was heated to 70-75 °C.

- 10 Next the mixture of phase B was added to the aqueous solution of Phases A+A1, stirred with a turbine to form an emulsion.

The compound Ac-SEQ ID NO:1-NH₂, ALDENINE® C (INCI: WATER (AQUA), HYDROLIZED SOY PROTEIN, HYDROLYZED WHEAT PROTEIN, XANTHAN GUM, TRIPEPTIDE-1), aqueous solution of LEUPHASYL® (INCI: WATER (AQUA),

- 15 GLYCERIN, PENTAPEPTIDE-18, CAPRYLYL GLYCOL), SILICONE DC 200 (INCI: DIMETHICONE) and SILICONE DC 245 (INCI: CYCLOPENTASILOXANE) (phase C). were added stepwise under stirring to the previous emulsion at 40 °C until homogenization. Next, perfume (INCI: FRAGRANCE (PARFUM)) (phase D), was slowly added under stirring until homogenized.

- 20 Finally, the pH of the mixture was adjusted to 5.5-7.0 with aqueous solution of 20% sodium hydroxide .

Cosmetic facial composition		
Phase	INGREDIENT	% in weight
A	WATER (AQUA)	q.s.p. 100
A	DISODIUM EDTA	0.30
A	PHENONIP® (INCI: PHENOXYETHANOL, METHYLPARABEN, ETHYLPARABEN, BUTYLPARABEN, PROPYLPARABEN, ISOBUTYLPARABEN): PHENOXYETHANOL,	
		0.610

	METHYLPARABEN	0.130
	ETHYLPARABEN	0.033
	BUTYLPARABEN	0.033
	PROPYLPARABEN	0.017
	ISOBUTYLPARABEN	0.017
A	IMIDAZOLIDINYL UREA	0.20
A	GLYCERIN	3.00
A1	CARBOMER	0.60
B	MINERAL OIL (PARAFFINUM LIQUIDUM)	7.00
B	BHT	0.05
B	CETYL ALCOHOL	0.80
B	BEESWAX (CERA ALBA)	0.50
B	STEARYL ALCOHOL	1.50
B	LIPOMULSE® 165 (INCI: GLYCERYL STEARATE, PEG-100 STEARATE):	
	GLYCERYL STEARATE	1.25
	PEG-100 STEARATE	1.25
B	DIMETHYLMETHOXY CHROMANOL	0.01
B	BENZOPHENONE-3	0.20
B	ETHYLHEXYL METHOXYCINNAMATE	0.90
B	CETEARYL ALCOHOL	1.00
C	Ac-SEQ ID NO:1-NH ₂	0.10
C	ALDENINE® C (INCI: WATER (AQUA), HYDROLIZED SOY PROTEIN, HYDROLYZED WHEAT PROTEIN, XANTHAN GUM, TRIPEPTIDE-1):	
	WATER (AQUA)	1.195
	HYDROLIZED SOY PROTEIN	0.500
	HYDROLYZED WHEAT PROTEIN	0.300
	XANTHAN GUM	0.003
	TRIPEPTIDE-1	0.002
C	LEUPHASYL® SOLUTION (INCI: WATER (AQUA), GLYCERIN, PENTAPEPTIDE-18, CAPRYLYL GLYCOL):	
	WATER (AQUA)	4.473
	GLYCERIN	0.500
	PENTAPETIDE-18	0.003
	CAPRYLYL GLYCOL	0.024
C	DIMETHICONE	0.10
C	CYCLOPENTASILOXANE	1.00

D	FRAGRANCE (PARFUM)	0.05
E	20% SODIUM HYDROXIDE	q.s.p. pH 5.5 - 7

Table 7

EXAMPLE 16

Prophetic preparation of a cosmetic facial composition containing Ac-SEQ ID NO:24-NH₂

- 5 In a suitable vessel the components from phase A are dissolved and the mixture is heated to 70-75 °C.

Next the mixture of phase B is added to the aqueous solution of Phase A under turbine stirring to form an emulsion. The emulsion is heated to 40 °C and the compound Ac-SEQ ID NO:24-NH₂, aqueous solution of ARGIRELINE® (INCI: WATER (AQUA),

- 10 ACETYL HEXAPEPTIDE-8) and ANTARTICINE® (INCI: WATER (AQUA), PSEUDOALTEROMONAS FERMENT EXTRACT) (phase C) is added under stirring. Finally, the perfume (INCI: FRAGRANCE (PARFUM)) (phase D) is added.

Cosmetic facial composition		
Phase	INGREDIENT	% in weight
A	WATER (AQUA)	q.s.p. 100
A	GLYCERIN	3.00
A	MAGNESIUM SULFATE	1.00
A	PHENONIP® (INCI: PHENOXYETHANOL, METHYLPARABEN, ETHYLPARABEN, BUTYLPARABEN, PROPYLPARABEN, ISOBUTYLPARABEN):	
	PHENOXYETANOL	0.363
	METHYLPARABEN	0.077
	ETHYLPARABEN	0.020
	BUTYLPARABEN	0.020
	PROPYLPARABEN	0.010
	ISOBUTYLPARABEN	0.010
A	DISODIUM EDTA	0.15
A	IMIDAZOLIDINYL UREA	0.1
B	MINERAL OIL (PARAFFINUM LIQUIDUM)	10.00
B	ETHYLHEXYL COCOATE	10.00
B	CETYL PEG/PPG-10/1 DIMETHICONE	2.00
B	BEESWAX (CERA ALBA)	1.50

B	HYDROGENATED CASTOR OIL	1.00
B	TOCOPHERYL ACETATE	0.10
C	Ac-SEQ ID NO:24-NH ₂	0.10
C	ARGIRELINE [®] SOLUTION (INCI: WATER (AQUA), ACETYL HEXAPEPTIDE-8):	
	WATER (AQUA)	9.995
	ACETYL HEXAPEPTIDE-8	0.005
	ANTARTICINE [®] (INCI: WATER (AQUA),	
C	PSEUDOALTEROMONAS FERMENT EXTRACT):	
	WATER (AQUA)	3.750
	PSEUDOALTEROMONAS FERMENT EXTRACT	1.250
D	FRAGRANCE (PARFUM)	0.10

Table 8

EXAMPLE 17

Prophetic preparation of a face cream containing Ac-SEQ ID NO:25-NH₂.

- 5 In a suitable vessel pentylene glycol [INCI: PENTYLENE GLYCOL], benzyl alcohol [INCI: BENZYL ALCOHOL], INYLINETM [INCI: ACETYL HEXAPEPTIDE-30] and the compound Ac-SEQ ID NO:25-NH₂ (phase A) are dissolved in water under constant, light stirring. Once homogenized, carbomer [INCI: CARBOMER] (phase A1) is added and the mixture is stirred until completely dissolved. Next, potassium cetyl phosphate 10 [INCI: POTASSIUM CETYL PHOSPHATE] (phase A2) is added until it is dispersed and the whole mixture is heated to 70-75 °C.

In a separate vessel the components of phase B are mixed together, heated to 70-75 °C and, once homogenized, phase B is added little by little to phase A under constant stirring.

- 15 With the mixture at about 50 °C, phase C is slowly added maintaining stirring. Phase D is then added until homogenization.

Once the mixture is homogenized, the pH is adjusted to 6.0-6.5 with phase E. Finally, the perfume is added (phase F).

Face cream composition		
Phase	INGREDIENT	% in weight

A	WATER (AQUA)	q.s.p. 100
A	PENTYLENE GLYCOL	5.0
A	BENZYL ALCOHOL	0.40
A	Ac-SEQ ID NO:25-NH ₂	0.10
A	ACETYL HEXAPEPTIDE-30	0.05
A1	CARBOMER	0.50
A2	POTASSIUM CETYL PHOSPHATE	0.50
B	SOYBEAN (GLYCINE SOJA) OIL	5.00
B	PHYTOCREAM 2000 [®] (INCI: GLYCERYL STEARATE, CETEARYL ALCOHOL, POTASSIUM PALMITOYL HYDROLYZED WHEAT PROTEIN):	
	GLYCERYL STEARATE	2.050
	CETEARYL ALCOHOL	2.050
	POTASSIUM PALMITOYL HYDROLYZED WHEAT PROTEIN	0.900
B	ETHYLHEXYL COCOATE	2.00
B	PHENOXYETHANOL	0.90
C	DIMETHICONE	1.00
D	SEPIGEL™ 305 (INCI: POLYACRYLAMIDE, WATER (AQUA), C13-14 ISOPARAFFIN, LAURETH-7)	
	POLYACRYLAMIDE	0.400
	WATER (AQUA)	0.340
	C13-14 ISOPARAFFIN	0.200
	LAURETH-7	0.060
E	20% SODIUM HYDROXIDE	q.s.p. pH 6.0 – 6.5
F	FRAGRANCE (PARFUM)	0.20

Table 9

EXAMPLE 18

Preparation of an antiperspirant serum containing Ac-SEQ ID NO:14-NH₂.

- 5 In a suitable vessel Phase B was stirred until it dissolved. In another vessel, the ingredients from phase C were heated until completely dissolved, and phase C was added, under stirring, to phase B.

In a separate vessel the compound Ac-SEQ ID NO:14-NH₂ and PHENONIP[®] [INCI: PHENOXYETHANOL, METHYLPARABEN, ETHYLPARABEN, BUTYLPARABEN,

- 10 PROPYLPARABEN, ISOBUTYLPARABEN] (phase A) were dissolved in water, under stirring.

Finally, phase A was added to the mixture of phases B+C, under stirring until homogenized.

Antiperspirant serum composition		
Phase	INGREDIENT	% in weight
A	WATER (AQUA)	0.50
A	Ac-SEQ ID NO:14-NH ₂	0.50
	PHENONIP® (INCI: PHENOXYETHANOL,	
	METHYLPARABEN, ETHYLPARABEN, BUTYLPARABEN,	
A	PROPYLPARABEN, ISOBUTYLPARABEN):	
	PHENOXYETHANOL	0.36
	METHYLPARABEN	0.08
	ETHYLPARABEN	0.02
	BUTYLPARABEN	0.02
	PROPYLPARABEN	0.01
	ISOBUTYLPARABEN	0.01
B	SILICONA 9040 (INCI: CYCLOPENTASILOXANE,	
	DIMETHICONE CROSSPOLYMER):	
	CYCLOPENTASILOXANE	43.265
	DIMETHICONE CROSSPOLYMER	7.635
	CYCLOPENTASILOXANE	31.36
B	FRAGRANCE (PARFUM)	0.13
C	ETHYLHEXYL COCOATE	12.43
C	C24-28 ALKYL DIMETHICONE	2.25
C	LECITHIN	1.38
C	BHT	0.05

Table 10

5 EXAMPLE 19

Preparation of a cosmetic facial composition containing Ac-SEQ ID NO:116-NH₂.

In a suitable vessel the components of phase A were dissolved. Next, carbomer was slowly added (INCI: CARBOMER) (phase A1), under stirring, until it was completely dissolved. The mixture was heated to 70-75 °C.

- 10 In another vessel the components of phase B were mixed together and the mixture was heated to 70-75 °C.

Next the mixture of phase B was added to the aqueous solution of Phases A+A1, stirred with a turbine to form an emulsion.

The compound Ac-SEQ ID NO:116-NH₂, ALDENINE® C (INCI: WATER (AQUA), HYDROLIZED SOY PROTEIN, HYDROLYZED WHEAT PROTEIN, XANTHAN GUM, 5 TRIPEPTIDE-1), aqueous solution of LEUPHASYL® (INCI: WATER (AQUA), GLYCERIN, PENTAPEPTIDE-18, CAPRYLYL GLYCOL), SILICONE DC 200 (INCI: DIMETHICONE) and SILICONE DC 245 (INCI: CYCLOPENTASILOXANE) (phase C). were added stepwise under stirring to the previous emulsion at 40 °C until homogenization. Next, perfume (INCI: FRAGRANCE (PARFUM)) (phase D), was 10 slowly added under stirring until homogenized.

Finally, the pH of the mixture was adjusted to 5.5-7.0 with aqueous solution of 20% sodium hydroxide .

Cosmetic facial composition		
Phase	INGREDIENT	% in weight
A	WATER (AQUA)	q.s.p. 100
A	DISODIUM EDTA	0.30
A	PHENONIP® (INCI: PHENOXYETHANOL, METHYLPARABEN, ETHYLPARABEN, BUTYLPARABEN, PROPYLPARABEN, ISOBUTYLPARABEN):	
	PHENOXYETHANOL,	0.610
	METHYLPARABEN	0.130
	ETHYLPARABEN	0.033
	BUTYLPARABEN	0.033
	PROPYLPARABEN	0.017
	ISOBUTYLPARABEN	0.017
A	IMIDAZOLIDINYL UREA	0.20
A	GLYCERIN	3.00
A1	CARBOMER	0.60
B	MINERAL OIL (PARAFFINUM LIQUIDUM)	7.00
B	BHT	0.05
B	CETYL ALCOHOL	0.80
B	BEESWAX (CERA ALBA)	0.50
B	STEARYL ALCOHOL	1.50
B	LIPOMULSE® 165 (INCI: GLYCERYL STEARATE, PEG-100	

STEARATE):		
	GLYCERYL STEARATE	1.25
	PEG-100 STEARATE	1.25
B	DIMETHYLMETHOXY CHROMANOL	0.01
B	BENZOPHENONE-3	0.20
B	ETHYLHEXYL METHOXYCINNAMATE	0.90
B	CETEARYL ALCOHOL	1.00
C	Ac-SEQ ID NO:116-NH ₂	0.10
C	ALDENINE [®] C (INCI: WATER (AQUA), HYDROLIZED SOY PROTEIN, HYDROLYZED WHEAT PROTEIN, XANTHAN GUM, TRIPEPTIDE-1):	
	WATER (AQUA)	1.195
	HYDROLIZED SOY PROTEIN	0.500
	HYDROLYZED WHEAT PROTEIN	0.300
	XANTHAN GUM	0.003
	TRIPEPTIDE-1	0.002
C	LEUPHASYL [®] SOLUTION (INCI: WATER (AQUA), GLYCERIN, PENTAPEPTIDE-18, CAPRYLYL GLYCOL):	
	WATER (AQUA)	4.473
	GLYCERIN	0.500
	PENTAPETIDE-18	0.003
	CAPRYLYL GLYCOL	0.024
C	DIMETHICONE	0.10
C	CYCLOPENTASILOXANE	1.00
D	FRAGRANCE (PARFUM)	0.05
E	20% SODIUM HYDROXIDE	q.s.p. pH 5.5 - 7

Table 11

EXAMPLE 20

Preparation of a cosmetic facial composition containing Ac-SEQ ID NO:14-NH₂

In a suitable vessel the components from phase A were dissolved and the mixture was
5 heated to 70-75 °C.

Next the mixture of phase B was added to the aqueous solution of Phase A under turbine stirring to form an emulsion. The emulsion was heated to 40 °C and the compound Ac-SEQ ID NO:14-NH₂, aqueous solution of ARGIRELINE[®] (INCI: WATER (AQUA), ACETYL HEXAPEPTIDE-8) and ANTARTICINE[®] (INCI: WATER (AQUA),

PSEUDOALTEROMONAS FERMENT EXTRACT) (phase C) were added under stirring. Finally, the perfume (INCI: FRAGRANCE (PARFUM)) (phase D) was added.

Cosmetic facial composition		
Phase	INGREDIENT	% in weight
A	WATER (AQUA)	q.s.p. 100
A	GLYCERIN	3.00
A	MAGNESIUM SULFATE	1.00
A	PHENONIP [®] (INCI: PHENOXYETHANOL, METHYLPARABEN, ETHYLPARABEN, BUTYLPARABEN, PROPYLPARABEN, ISOBUTYLPARABEN): PHENOXYETHANOL	0.363
	METHYLPARABEN	0.077
	ETHYLPARABEN	0.020
	BUTYLPARABEN	0.020
	PROPYLPARABEN	0.010
	ISOBUTYLPARABEN	0.010
A	DISODIUM EDTA	0.15
A	IMIDAZOLIDINYL UREA	0.1
B	MINERAL OIL (PARAFFINUM LIQUIDUM)	10.00
B	ETHYLHEXYL COCOATE	10.00
B	CETYL PEG/PPG-10/1 DIMETHICONE	2.00
B	BEESWAX (CERA ALBA)	1.50
B	HYDROGENATED CASTOR OIL	1.00
B	TOCOPHERYL ACETATE	0.10
C	Ac-SEQ ID NO:14-NH ₂	0.10
C	ARGIRELINE [®] SOLUTION (INCI: WATER (AQUA), ACETYL HEXAPEPTIDE-8): WATER (AQUA)	9.995
	ACETYL HEXAPEPTIDE-8	0.005
	ANTARTICINE [®] (INCI: WATER (AQUA),	
C	PSEUDOALTEROMONAS FERMENT EXTRACT): WATER (AQUA)	3.750
	PSEUDOALTEROMONAS FERMENT EXTRACT	1.250
D	FRAGRANCE (PARFUM)	0.10

Table 12

Preparation of a face cream containing Ac-SEQ ID NO:117-NH₂.

In a suitable vessel pentylene glycol [INCI: PENTYLENE GLYCOL], benzyl alcohol [INCI: BENZYL ALCOHOL], INYLINE™ [INCI: ACETYL HEXAPEPTIDE-30] and the compound Ac-SEQ ID NO:117-NH₂ (phase A) were dissolved in water under constant, 5 light stirring. Once homogenized, carbomer [INCI: CARBOMER] (phase A1) was added and the mixture was stirred until completely dissolved. Next, potassium cetyl phosphate [INCI: POTASSIUM CETYL PHOSPHATE] (phase A2) was added until it was dispersed and the whole mixture was heated to 70-75 °C.

In a separate vessel the components of phase B were mixed together, heated to 70-75 10 °C and, once homogenized, phase B was added little by little to phase A under constant stirring.

With the mixture at about 50 °C, phase C was slowly added maintaining stirring. Phase D was then added until homogenization.

Once the mixture was homogenized, the pH was adjusted to 6.0-6.5 with phase E. 15 Finally, the perfume was added (phase F).

Face cream composition		
Phase	INGREDIENT	% in weight
A	WATER (AQUA)	q.s.p. 100
A	PENTYLENE GLYCOL	5.0
A	BENZYL ALCOHOL	0.40
A	Ac-SEQ ID NO:117-NH ₂	0.10
A	ACETYL HEXAPEPTIDE-30	0.05
A1	CARBOMER	0.50
A2	POTASSIUM CETYL PHOSPHATE	0.50
B	SOYBEAN (GLYCINE SOJA) OIL	5.00
B	PHYTOCREAM 2000 [®] (INCI: GLYCERYL STEARATE, CETEARYL ALCOHOL, POTASSIUM PALMITOYL HYDROLYZED WHEAT PROTEIN):	
	GLYCERYL STEARATE	2.050
	CETEARYL ALCOHOL	2.050
	POTASSIUM PALMITOYL HYDROLYZED WHEAT PROTEIN	0.900
B	ETHYLHEXYL COCOATE	2.00
B	PHENOXYETHANOL	0.90
C	DIMETHICONE	1.00

D	SEPIGEL™ 305 (INCI: POLYACRYLAMIDE, WATER (AQUA), C13-14 ISOPARAFFIN, LAURETH-7)	
	POLYACRYLAMIDE	0.400
	WATER (AQUA)	0.340
	C13-14 ISOPARAFFIN	0.200
	LAURETH-7	0.060
E	20% SODIUM HYDROXIDE	q.s.p. pH 6.0 – 6.5
F	FRAGRANCE (PARFUM)	0.20

Table 13

EXAMPLE 22

Prophetic preparation of a face cream containing Palm-SEQ ID NO:1-NH₂

- 5 In a suitable vessel pentylene glycol (PENTYLENE GLYCOL), benzyl alcohol (INCI: BENZYL ALCOHOL), glycerol (INCI: GLYCEROL), urea (INCI: UREA) and the compound Palm-SEQ ID NO:1-NH₂ (phase A) are dissolved in water. Next, carbomer (INCI: CARBOMER) (phase A1) is slowly added, under stirring, until it is completely dissolved. The mixture is heated to 70-75 °C.
- 10 The mixture of phase B is added to the aqueous solution of Phases A+A1 under turbine stirring to form an emulsion.

The perfume (INCI: FRAGRANCE (PARFUM)) (phase C) is added to the previous emulsion at 40 °C.

- 15 Finally, the pH of the mixture is adjusted to 6.0-6.5 with aqueous solution of 20% sodium hydroxide when required.

Face cream composition		
Phase	INGREDIENT	% in weight
A	WATER (AQUA)	q.s.p. 100
A	PENTYLENE GLYCOL	5.0
A	BENZYL ALCOHOL	0.40
A	GLYCERIN	1.00
A	UREA	1.00
A	Palm-SEQ ID NO:1-NH ₂	0.10
A1	CARBOMER	0.60

B	LIPOMULSE [®] 165 (INCI: GLYCERYL STEARATE, PEG-100 STEARATE):	
	GLYCERYL STEARATE	1.500
	PEG-100 STEARATE	1.500
B	CETEARETH-25	2.00
B	SHEA BUTTER (BUTYROSPERMUM PARKII)	2.00
B	CAPRYLIC/CAPRIC TRIGLYCERIDE	1.00
B	ETHYLHEXYL COCOATE	7.00
B	PHENOXYETHANOL	0.90
B	DIMETHICONE	1.00
C	FRAGRANCE (PARFUM)	0.20
D	20% SODIUM HYDROXIDE	q.s.p. pH 6.0 – 6.5

Table 14

EXAMPLE 23*Preparation of liposomes containing Ac-SEQ ID NO:116-NH₂*

Dipalmitoylphosphatidylcholine (DPPC) was weighed and dissolved in chloroform. The solvent was evaporated to dryness until a fine layer of phospholipid was obtained, and this layer was hydrated by treatment with an aqueous solution which contained the peptide Ac-SEQ ID NO:116-NH₂ at the desired concentration (containing PHENONIP[®]) at 55 °C, obtaining the MLV liposomes. The ULV liposomes were obtained by submerging the MLV liposomes in an ultrasound bath at 55 °C for 8 cycles of 2 minutes at intervals of 5 minutes.

INGREDIENT	% IN WEIGHT
WATER (AQUA)	q.s.p. 100
DIPALMITOYLPHOSPHATIDYLCHOLINE	4.00
15 Ac-SEQ ID NO:116-NH ₂	0.20
PHENONIP [®] (INCI: PHENOXYETHANOL, METHYLPARABEN, ETHYLPARABEN, BUTYLPARABEN, PROPYLPARABEN, ISOBUTYLPARABEN):	
ISOBUTYLPARABEN	0.01
PHENOXYETHANOL	0.36
20 METHYLPARABEN	0.08
ETHYLPARABEN	0.02
BUTYLPARABEN	0.02
PROPYLPARABEN	0.01

Table 15

EXAMPLE 24

Preparation of coacervation capsules containing the compound Palm-SEQ ID NO:25-NH₂

- 5 a) *Preparation of an emulsion of the compound Palm-SEQ ID NO:25-NH₂*
 In a suitable vessel the peptide was dissolved in water (phase A) heating the mixture to 70 °C. In a separate vessel soybean oil [INCI: SOYBEAN (GLYCINE SOJA) OIL], Abil EM 90 [INCI: CETYL PEG/PPG-10/1 DIMETHICONE] and Span 65 [INCI: SORBITAN TRISTEARATE] (phase B) were mixed together, heating the mixture to 80 °C until the
 10 Span 65 dissolved. Once melted, phase A was added to phase B slowly under intense stirring with a turbine. Once the components were mixed together, the mixture was stirred until it reached room temperature.

Phase	INGREDIENT	% in weight
A	PURIFIED WATER	56.00
B	SOYBEAN (GLYCINE SOJA) OIL	33.00
B	CETYL PEG/PPG-10/1 DIMETHICONE	5.00
B	SORBITAN TRISTEARATE	4.00
A	Palm-SEQ ID NO:25-NH ₂	2.00

Table 16

- 15 b) *Preparation of a microfluidized emulsion of the compound Palm-SEQ ID NO:25-NH₂*

The components of phase A were mixed together in water: Zemea Propanediol [INCI: PROPANEDIOL], phenoxyethanol [INCI: PHENOXYETHANOL], Structure XL [INCI: HYDROXYPROPYL STARCH PHOSPHATE], Amigel [INCI: SCLEROTIUM GUM] and
 20 powdered hyaluronic acid [INCI: SODIUM HYALURONATE], the mixture was heated to 70 °C under stirring. In another vessel, the emulsion from section a), and the components of phase B: Massocare HD [INCI: ISOHEXADECANE], Arlacel 83 [INCI: SORBITAN SESQUIOLEATE], LIPOCHROMAN™ [INCI: DIMETHYLMETHOXY CHROMANOL] were mixed together, and the mixture was heated to 80 °C under stirring.
 25

Once this temperature was reached, phase B was added to phase A very slowly under intense stirring with a turbine.

The sample was passed, without cooling, through a microfluidizer for three cycles at an entrance pressure of 80 bar and an exit pressure of 15000 psi, maintaining the operating temperature between 65 and 75 °C. Once microfluidized, the emulsion was stirred with a rotor until room temperature was reached.

5

Phase	INGREDIENT	% in weight
A	WATER (AQUA)	70.36
B	Emulsion section a)	10.95
B	ISOHEXADECANE	5.48
A	PROPANEDIOL	5.48
B	SORBITAN SESQUIOLEATE	4.38
A	PHENOXYETHANOL	2.85
A	HYDROXYPROPYL STARCH PHOSPHATE	0.33
A	SCLEROTIUM GUM	0.11
B	DIMETHYLMETHOXY CHROMANOL	0.05
A	SODIUM HYALURONATE	0.01

Table 17

c) *Obtaining coacervate capsules containing the compound Palm-SEQ ID NO:25-NH₂*

- 10 In a vessel the emulsion from section b) (phase A) was weighed. In another vessel Sensomer CI 50 [INCI: WATER (AQUA); STARCH HYDROXYPROPYLTRIMONIUM CHLORIDE; UREA; SODIUM LACTATE; SODIUM CHLORIDE; SODIUM BENZOATE] was dissolved in water (phase B). Phase B was added to phase A under intense stirring.
- 15 Amigel [INCI: SCLEROTIUM GUM] (phase C) was added to the previous mixture very slowly and under intense stirring. The mixture was stirred for 2 hours to obtain good hydration of the gum.

20 Next, Structure XL [INCI: HYDROXYPROPYL STARCH PHOSPHATE] (phase D) was added, maintaining the stirring for another hour to obtain the complete hydration of the biopolymers added.

Finally, Sepigel 305 [INCI: POLYACRYLAMIDE; WATER (AQUA); C13-14 ISOPARAFFIN; LAURETH-7] (phase E) was added, maintaining the stirring until a homogenous suspension was obtained. The average size of the capsules in

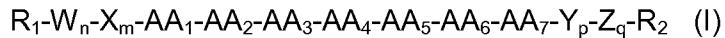
suspension obtained determined by Dynamic Laser Light Scattering was approximately 300 nm.

Phase	INGREDIENT	% in weight
A	Emulsion section b)	91.30
B	WATER (AQUA)	6.00
D	HYDROXYPROPYL STARCH PHOSPHATE	1.50
C	SCLEROTIUM GUM	0.75
E	SEPIGEL™ 305 (INCI: POLYACRYLAMIDE, WATER (AQUA), C13-14 ISOPARAFFIN, LAURETH-7): POLYACRYLAMIDE	0.10
	WATER (AQUA)	0.085
	C13-14 ISOPARAFFIN	0.05
	LAURETH-7	0.015
B	SENSOMER CI 50 [INCI: WATER (AQUA); STARCH HYDROXYPROPYLTRIMONIUM CHLORIDE; UREA; SODIUM LACTATE; SODIUM CHLORIDE; SODIUM BENZOATE]: WATER (AQUA)	0.137
	STARCH HYDROXYPROPYLTRIMONIUM CHLORIDE	0.048
	UREA	0.006
	SODIUM LACTATE	0.004
	SODIUM CHLORIDE	0.004
	SODIUM BENZOATE	0.001

Table 18

CLAIMS

1. A compound of general formula (I):



its stereoisomers, mixtures thereof and/or its cosmetically or pharmaceutically acceptable salts, characterized in that:

AA₁ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₂ is selected from the group formed by -His-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₃ is selected from the group formed by -Leu-, -Ile-, -Phe- and -Tyr-;

AA₄ is selected from the group formed by -Lys-, -His-, -Arg-, -Gln-, -Leu-, -Ile-, -Met-, -MetO- and -MetO₂-;

AA₅ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₆ is selected from the group formed by -Phe-, -Trp-, -Ile- and -Val-;

AA₇ is selected from the group formed by -Met-, -MetO- and -MetO₂-;

W, X, Y, Z are amino acids and are independently selected from amongst themselves;

n, m, p and q are independently selected from amongst themselves and have a value of 0 or 1;

20 n+m+p+q is smaller or equal to 2 ;

with the condition that it is not SEQ ID NO:111;

25 R₁ is selected from the group formed by H, a polymer derived from polyethylene glycol, a non-cyclic substituted or unsubstituted aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl and R₅-CO-, wherein R₅ is selected from the group formed by H, a non-cyclic substituted or unsubstituted aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted aryl, substituted or

unsubstituted aralkyl, substituted or unsubstituted heterocyclyl and substituted or unsubstituted heteroarylalkyl;

R₂ is selected from the group formed by -NR₃R₄, -OR₃ and -SR₃, wherein R₃ and R₄ are independently selected from the group formed by H, a polymer derived from polyethylene glycol, a non-cyclic substituted or unsubstituted aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, and substituted or unsubstituted aralkyl; and

R₁ and R₂ are not α -amino acids.

- 10 2. Compound according to claim 1, wherein AA₁ is selected from the group formed by -Arg-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-, AA₂ is selected from the group formed by -His-, -Gln-, -Asn-, -Glu- and -Asp-, AA₃ is selected from the group formed by -Leu- and -Phe-, AA₄ is selected from the group formed by -His-, -Lys-, -Gln- and -Leu-, AA₅ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-, AA₆ is selected from the group formed by -Trp- and -Val- and AA₇ is selected from the group formed by -Met-, -MetO- and -MetO₂.
- 15 3. Compound according to claim 2, wherein AA₄ is selected from the group formed by -Lys- and -Leu-, and AA₆ is -Trp-.
- 20 4. Compound according to claim 3, wherein n, m, p and q are 0.
5. Compound according to claim 2, wherein R₁ is selected from the group formed by H, acetyl, lauroyl, myristoyl and palmitoyl, AA₁ is selected from the group formed by -L-Asn-, -L-Glu- and -L-Asp-, AA₂ is selected from the group formed by -L-His- and -L-Asp-, AA₃ is selected from the group formed by -L-Leu- and -L-Phe-, AA₄ is selected from the group formed by -L-Lys-, -L-Gln- and -L-Leu-, AA₅ is selected from the group formed by -L-Arg-, -L-Gln-, -L-Asn- and -L-Asp-, AA₆ is selected from the group formed by -L-Trp-, AA₇ is selected from the group formed by -L-Met-, -L-MetO- and -L-MetO₂, and R₂ is selected from the group formed by -NR₃R₄ and -OR₃ wherein R₃ and R₄ are independently selected from H, methyl, ethyl, hexyl, dodecyl and hexadecyl.

6. Compound according to claim 2, wherein R₁ is selected from the group formed by H, acetyl, lauroyl, myristoyl and palmitoyl, AA₁ is -L-Glu-, AA₂ is selected from the group formed by -L-Asn-, -L-Glu-, -L-Gln- and -L-Asp-, AA₃ is -L-Leu-, AA₄ is -L-Lys-, AA₅ is -L-Arg-, AA₆ is -L-Trp-, AA₇ is selected from the group formed by -L-Met-, -L-MetO- and -L-MetO₂-, and R₂ is selected from the group formed by -NR₃R₄ and -OR₃ where R₃ and R₄ are independently selected from H, methyl, ethyl, hexyl, dodecyl and hexadecyl.
5
7. Compound according to claim 5, wherein R₁ is selected from the group formed by H, acetyl, lauroyl, myristoyl or palmitoyl, AA₁ is -L-Glu-, AA₂ is -L-His-, AA₃ is -L-Leu-, AA₄ is -L-Lys-, AA₅ is -L-Gln-, AA₆ is -L-Trp-, AA₇ is -L-Met-, -L-MetO- or -L-MetO₂- and R₂ is -NR₃R₄ or -OR₃ wherein R₃ and R₄ are independently selected from H, methyl, ethyl, hexyl, dodecyl and hexadecyl.
10
8. Compound according to claim 5, wherein R₁ is selected from the group formed by H, acetyl, lauroyl, myristoyl or palmitoyl, AA₁ is -L-Glu-, AA₂ is -L-Asp-, AA₃ is -L-Leu-, AA₄ is -L-Lys-, AA₅ is -L-Arg-, AA₆ is -L-Trp-, AA₇ is -L-Met-, -L-MetO- or -L-MetO₂- and R₂ is -NR₃R₄ or -OR₃ wherein R₃ and R₄ are independently selected from H, methyl, ethyl, hexyl, dodecyl and hexadecyl.
15
9. Compound according to claim 5, wherein R₁ is selected from the group formed by H, acetyl, lauroyl, myristoyl or palmitoyl, AA₁ is -L-Asn-, AA₂ is -L-His-, AA₃ is -L-Phe-, AA₄ is -L-Leu-, AA₅ is -L-Asn-, AA₆ is -L-Trp-, AA₇ is -L-Met-, -L-MetO- or -L-MetO₂- and R₂ is -NR₃R₄ or -OR₃ wherein R₃ and R₄ are independently selected from H, methyl, ethyl, hexyl, dodecyl and hexadecyl.
20
10. Compound according to claim 5, wherein R₁ is selected from the group formed by H, acetyl, lauroyl, myristoyl or palmitoyl, AA₁ is -L-Asp-, AA₂ is -L-His-, AA₃ is -L-Phe-, AA₄ is -L-Leu-, AA₅ is -L-Asp-, AA₆ is -L-Trp-, AA₇ is -L-Met-, -L-MetO- or -L-MetO₂- and R₂ is -NR₃R₄ or OR₃ wherein R₃ and R₄ are independently selected from H, methyl, ethyl, hexyl, dodecyl and hexadecyl.
25
11. Cosmetic or pharmaceutical composition which comprises at least one compound of general formula (I), its stereoisomers, mixtures thereof and/or its cosmetically or pharmaceutically acceptable salts, according to any of claims 1 to 10, together with at least one cosmetically or pharmaceutically acceptable excipient or adjuvant.
30

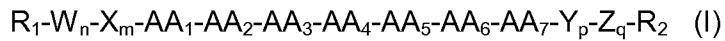
12. Composition according to claim 11, wherein this compound of general formula (I), its stereoisomers, mixtures thereof and/or its cosmetically or pharmaceutically acceptable salts, is incorporated into a cosmetically or pharmaceutically acceptable delivery system or sustained release system selected from the group formed by 5 liposomes, mixed liposomes, oleosomes, niosomes, ethosomes, milliparticles, microparticles, nanoparticles, solid lipid nanoparticles, nanostructured lipid carriers, sponges, cyclodextrins, vesicles, micelles, mixed micelles of surfactants, surfactant-phospholipid mixed micelles, millispheres, microspheres, nanospheres, lipospheres, millicapsules, microcapsules, nanocapsules, microemulsions and 10 nanoemulsions or is adsorbed on a solid organic polymer or solid mineral support selected from the group formed by talc, bentonite, silica, starch and maltodextrin.
13. Composition according to any of claims 11 or 12, wherein this composition is presented in a formulation selected from the group formed by creams, multiple emulsions, anhydrous compositions, aqueous dispersions, oils, milks, balsams, 15 foams, lotions, gels, cream gels, hydroalcoholic solutions, hydroglycolic solutions, hydrogels, liniments, sera, soaps, shampoos, conditioners, serums, polysaccharide films, ointments, mousses, pomades, powders, bars, pencils, sprays, aerosols, capsules, gelatin capsules, soft capsules, hard capsules, tablets, sugar coated tablets, pills, powders, granules, chewing gum, solutions, suspensions, emulsions, 20 syrups, elixirs, jellies and gelatins.
14. Composition according to any of claims 11 to 13, wherein this composition also comprises at least one cosmetically or pharmaceutically acceptable adjuvant selected from the group formed by agents which inhibit neuronal exocytosis, 25 anticholinergic agents, agents which inhibit muscular contraction, anti-aging agents, anti-wrinkle agents, antiperspirant agents, anti-inflammatory agents and/or analgesics, anti-itching agents, calming agents, anesthetic agents, inhibitors of acetylcholine-receptor aggregation, agents that inhibit acetylcholinesterase, skin relaxant agents, melanin synthesis stimulating or inhibiting agents, whitening or depigmenting agents, propigmenting agents, self-tanning agents, NO-synthase 30 inhibiting agents, 5 α -reductase inhibiting agents, lysyl- and/or prolyl hydroxylase inhibiting agents, antioxidants, free radical scavengers and/or agents against atmospheric pollution, reactive carbonyl species scavengers, anti-glycation agents, antihistamine agents, antiviral agents, antiparasitic agents, emulsifiers, emollients, organic solvents, liquid propellants, skin conditioners, humectants, substances that

retain moisture, alpha hydroxy acids, beta hydroxy acids, moisturizers, epidermal hydrolytic enzymes, vitamins, amino acids, proteins, pigments, colorants, dyes, biopolymers, gelling polymers, thickeners, surfactants, softening agents, emulsifiers, binding agents, preservatives, agents able to reduce or treat bags under the eyes, exfoliating agents, keratolytic agents, flaking agents, antimicrobial agents, antifungal agents, fungistatic agents, bactericidal agents, bacteriostatic agents, agents stimulating the synthesis of dermal or epidermal macromolecules and/or capable of inhibiting or preventing their degradation, collagen synthesis-stimulating agents, elastin synthesis-stimulation agents, decorin synthesis-stimulation agents, laminin synthesis-stimulation agents, defensin synthesis-stimulating agents, chaperone synthesis-stimulating agents, cAMP synthesis-stimulating agents, AQP-3 modulating agents, aquaporin synthesis modulating agents, proteins from the aquaporin family, hyaluronic acid synthesis-stimulating agents, glycosaminoglycan synthesis-stimulating agents, fibronectin synthesis-stimulating agents, sirtuin synthesis-stimulating agents, sirtuin activating agents, heat shock proteins, heat shock protein synthesis-stimulating agents, agents stimulating the synthesis of lipids and components of the stratum corneum, ceramides, fatty acids, agents that inhibit collagen degradation, matrix metalloproteinase inhibitory agents, agents that inhibit elastin degradation, agents that inhibit serine proteases, agents stimulating fibroblast proliferation, agents stimulating keratinocyte proliferation, agents stimulating adipocyte proliferation, agents stimulating melanocyte proliferation, agents stimulating keratinocyte differentiation, agents stimulating or delaying adipocyte differentiation, antihyperkeratosis agents, comedolytic agents, anti-psoriasis agents, DNA repair agents, DNA protecting agents, stabilizers, agents for the treatment and/or care of sensitive skin, firming agents, anti-stretch mark agents, binding agents, agents regulating sebum production, lipolytic agents or agents stimulating lipolysis, adipogenic agents, agents modulating PGC-1 α expression, agents modulating PPAR γ , agents which increase or reduce the triglyceride content of adipocytes, anti-cellulite agents, agents which inhibit the activity of PAR-2, agents stimulating healing, coadjuvant healing agents, agents stimulating reepithelialization, coadjuvant reepithelialization agents, cytokine growth factors, agents acting on capillary circulation and/or microcirculation, agents stimulating angiogenesis, agents that inhibit vascular permeability, venotonic agents, agents acting on cell metabolism, agents to improve dermal-epidermal junction, agents inducing hair

growth, hair growth inhibiting or retardant agents, hair loss retardant agents, preservatives, perfumes, cosmetic and/or absorbent and/or body odor masking deodorants, chelating agents, plant extracts, essential oils, marine extracts, agents obtained from a biotechnological process, mineral salts, cell extracts, sunscreens and organic or mineral photoprotective agents active against ultraviolet A and/or B rays and/or infrared A rays, or mixtures thereof.

5 15. Compound of general formula (I), its stereoisomers, mixtures thereof and/or its cosmetically or pharmaceutically acceptable salts, as defined in any of claims 1 to 10, for its use in medicine.

10 16. A compound of general formula (I),



its stereoisomers, mixtures thereof and/or its cosmetically or pharmaceutically acceptable salts, characterized in that:

15 AA₁ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₂ is selected from the group formed by -His-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₃ is selected from the group formed by -Leu-, -Ile-, -Phe- and -Tyr-;

AA₄ is selected from the group formed by -Lys-, -His-, -Arg-, -Gln-, -Leu-, -Ile-, -Met-, -MetO- and -MetO₂-;

20 AA₅ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₆ is selected from the group formed by -Phe-, -Trp-, -Ile- and -Val-;

AA₇ is selected from the group formed by -Met-, -MetO- and -MetO₂-;

25 W, X, Y, Z are amino acids and are independently selected from among themselves;

n, m, p and q are independently selected from amongst themselves and have a value of 0 or 1;

n+m+p+q is smaller or equal to 2;

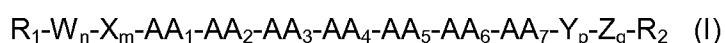
R₁ is selected from the group formed by H, a polymer derived from polyethylene glycol, a non-cyclic substituted or unsubstituted aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl and R₅-CO-, wherein R₅ is selected from the group formed by H, a non-cyclic substituted or unsubstituted aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl, substituted or unsubstituted heterocyclyl and substituted or unsubstituted heteroarylalkyl;

R₂ is selected from the group formed by -NR₃R₄, -OR₃ and -SR₃, wherein R₃ and R₄ are independently selected from the group formed by H, a polymer derived from polyethylene glycol, a non-cyclic substituted or unsubstituted aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, and substituted or unsubstituted aralkyl; and

R₁ and R₂ are not α -amino acids,

for its use in the treatment and/or prevention of pain, inflammation, itching, hyperhidrosis and/or of neurological, compulsive and/or neuropsychiatric diseases and/or disorders which are improved by the inhibition of neuronal exocytosis selected from the group formed by muscular spasticity, dystonia, focal dystonia, blepharospasm, torsion dystonia, cervical dystonia or torticollis, laryngeal dystonia or spasmodic dysphonia, oromandibular dystonia, extremity dystonia, writer's cramp, musician's cramp, foot dystonia, bruxism, facial scoliosis, hemifacial spasm, tics, strabismus, segmentary dystonia, Meige's syndrome, multifocal dystonia, hemidystonia, dopamine-responsive dystonia, Segawa's dystonia, trembling, Parkinson's disease, nervous impingements, Alzheimer's disease and Tourette's syndrome.

17. Use of a compound of general formula (I),



its stereoisomers, mixtures thereof and/or its cosmetically or pharmaceutically acceptable salts, characterized in that:

AA₁ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₂ is selected from the group formed by -His-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₃ is selected from the group formed by -Leu-, -Ile-, -Phe- and -Tyr-;

5 AA₄ is selected from the group formed by -Lys-, -His-, -Arg-, -Gln-, -Leu-, -Ile-, -Met-, -MetO- and -MetO₂-;

AA₅ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₆ is selected from the group formed by -Phe-, -Trp-, -Ile- and -Val-;

10 AA₇ is selected from the group formed by -Met-, -MetO- and -MetO₂-;

W, X, Y, Z are amino acids and are independently selected from among themselves;

n, m, p and q are independently selected from amongst themselves and have a value of 0 or 1;

15 n+m+p+q is smaller or equal to 2;

20 R₁ is selected from the group formed by H, a polymer derived from polyethylene glycol, a non-cyclic substituted or unsubstituted aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl and R₅-CO-, wherein R₅ is selected from the group formed by H, a non-cyclic substituted or unsubstituted aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl, substituted or unsubstituted heterocyclyl and substituted or unsubstituted heteroarylalkyl;

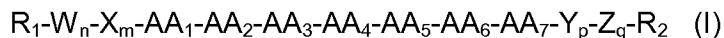
25 R₂ is selected from the group formed by -NR₃R₄, -OR₃ and -SR₃, wherein R₃ and R₄ are independently selected from the group formed by H, a polymer derived from polyethylene glycol, a non-cyclic substituted or unsubstituted aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, and substituted or unsubstituted aralkyl; and

R_1 and R_2 are not α -amino acids,

for the cosmetic, non-therapeutic treatment and/or care of the skin, hair and/or mucous membranes.

18. Use according to claim 17, wherein the cosmetic, non-therapeutic treatment and/or care of the skin, hair and/or mucous membranes is a treatment and/or prevention of aging and/or photoaging, treatment and/or prevention of wrinkles and/or expression wrinkles, treatment and/or prevention of perspiration, treatment and/or care of disorders of the skin selected from the group formed by calluses, warts, treatment stimulating hair growth and/or prevention of hair loss.

- 10 19. Compound of general formula (I),



its stereoisomers, mixtures thereof and/or its cosmetically or pharmaceutically acceptable salts, characterized in that:

15 AA₁ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₂ is selected from the group formed by -His-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₃ is selected from the group formed by -Leu-, -Ile-, -Phe- and -Tyr-;

AA₄ is selected from the group formed by -Lys-, -His-, -Arg-, -Gln-, -Leu-, -Ile-, -Met-, -MetO- and -MetO₂-;

20 AA₅ is selected from the group formed by -Arg-, -His-, -Lys-, -Gln-, -Asn-, -Glu- and -Asp-;

AA₆ is selected from the group formed by -Phe-, -Trp-, -Ile- and -Val-;

AA₇ is selected from the group formed by -Met-, -MetO- and -MetO₂-;

25 W, X, Y, Z are amino acids and are independently selected from among themselves;

n, m, p and q are independently selected from amongst themselves and have a value of 0 or 1;

n+m+p+q is smaller or equal to 2;

- R₁ is selected from the group formed by H, a polymer derived from polyethylene glycol, a non-cyclic substituted or unsubstituted aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl and R₅-CO-, wherein R₅ is selected from the group formed by H, a non-cyclic substituted or unsubstituted aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted aryl, substituted or unsubstituted aralkyl, substituted or unsubstituted heterocyclyl and substituted or unsubstituted heteroarylalkyl;
- R₂ is selected from the group formed by -NR₃R₄, -OR₃ and -SR₃, wherein R₃ and R₄ are independently selected from the group formed by H, a polymer derived from polyethylene glycol, a non-cyclic substituted or unsubstituted aliphatic group, substituted or unsubstituted alicyclyl, substituted or unsubstituted heterocyclyl, substituted or unsubstituted heteroarylalkyl, substituted or unsubstituted aryl, and substituted or unsubstituted aralkyl; and
- R₁ and R₂ are not α -amino acids,

for its use in the inhibition of neuronal exocytosis.

INTERNATIONAL SEARCH REPORT

International application No
PCT/EP2013/057656

A. CLASSIFICATION OF SUBJECT MATTER
INV. A61K8/64 C07K7/06
ADD.

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
A61K C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPO-Internal, CHEM ABS Data, BIOSIS, EMBASE, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	KR 101 046 426 B1 (ECODERM INC [KR]) 4 July 2011 (2011-07-04) the whole document -----	1-17
X	WO 2006/050611 A1 (UNIV BRITISH COLUMBIA [CA]; HANCOCK ROBERT E W [CA]; HILPERT KAI [CA]) 18 May 2006 (2006-05-18) sequence 1234 -----	14,17
Y	EP 2 123 673 A1 (LIPOTEC SA [ES]) 25 November 2009 (2009-11-25) cited in the application examples 1-7 -----	1-17
Y	WO 2011/009626 A1 (LIPOTEC SA [ES]; GARCIA SANZ NURIA [ES]; VAN DEN NEST WIM [ES]; CARREN) 27 January 2011 (2011-01-27) claim 1; examples 1,2,7,8,16; table 2 ----- -/-	1-17

Further documents are listed in the continuation of Box C.

See patent family annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier application or patent but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
6 June 2013	01/07/2013
Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016	Authorized officer Pinheiro Vieira, E

INTERNATIONAL SEARCH REPORT

International application No
PCT/EP2013/057656

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	BLANES-MIRA C ET AL: "A SYNTHETIC HEXAPEPTIDE (ARGIRELINE) WITH ANTIWRINKLE ACTIVITY", INTERNATIONAL JOURNAL OF COSMETIC SCIENCE, KLUWER ACADEMIC PUBLISHERS, DORDRECHT, NL, vol. 24, no. 5, 1 January 2002 (2002-01-01), pages 303-310, XP009045291, ISSN: 0142-5463, DOI: 10.1046/J.1467-2494.2002.00153.X page 306; figures 1-4 -----	1-17
Y	BLANES-MIRA C ET AL: "Identification of SNARE complex modulators that inhibit exocytosis from an alpha-helix-constrained combinatorial library", BIOCHEMICAL JOURNAL, THE BIOCHEMICAL SOCIETY, LONDON, GB, vol. 375, no. 1, 1 October 2003 (2003-10-01), pages 159-166, XP002402317, ISSN: 0264-6021, DOI: 10.1042/BJ20030509 page 163; figures 2-6 -----	1-17
1		

INTERNATIONAL SEARCH REPORT

International application No.

PCT/EP2013/057656

Box No. I Nucleotide and/or amino acid sequence(s) (Continuation of item 1.c of the first sheet)

1. With regard to any nucleotide and/or amino acid sequence disclosed in the international application and necessary to the claimed invention, the international search was carried out on the basis of:
 - a. (means)
 on paper
 in electronic form
 - b. (time)
 in the international application as filed
 together with the international application in electronic form
 subsequently to this Authority for the purpose of search
2. In addition, in the case that more than one version or copy of a sequence listing and/or table relating thereto has been filed or furnished, the required statements that the information in the subsequent or additional copies is identical to that in the application as filed or does not go beyond the application as filed, as appropriate, were furnished.
3. Additional comments:

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No PCT/EP2013/057656
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Patent document cited in search report	Publication date	Patent family member(s)		Publication date
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