

Title (en)

INTERACTION OF ALPHA-CONOTOXIN PEPTIDES WITH NEURONAL NICOTINIC ACETYLCHOLINE RECEPTORS

Title (de)

WECHSELWIRKUNG VON ALPHA-CONOTOXINPEPTIDEN MIT NEURONALEN NIKOTINISCHEN ACETYLCHOLIN REZEPTOREN

Title (fr)

INTERACTION DE PEPTIDES D'ALPHA-CONOTOXINE AVEC DES RECEPTEURS NEURAUUX D'ACETYLCHOLINE NICOTINIQUE

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Application

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Abstract (en)

[origin: WO9921878A1] This invention relates to derivatives of the conopeptide MII, an alpha -4/7 conotoxin peptide, in which amino acid residues are substituted as described herein while maintaining the basic activity of MII. The present invention also relates to the discovery of the 3-dimensional structure of MII, and the relationship of its structure to its specificity to the alpha 3 beta 2 subtype of the neuronal nicotinic acetylcholine receptor (nAChR). The present invention also relates to computer based programs for the expression of the three-dimensional structure of MII and peptide analogs, peptide mimetics or non-peptide mimetics thereof. The structural characteristics may be correlated with biological activity to enable the design of alpha -4/7 conotoxin peptide analogs and peptide mimetics which demonstrate the same specificity to neuronal nAChR. Such analogs and peptide mimetics are useful as cardiovascular agents and for treating or detecting small-cell lung carcinoma (SCLC).

[origin: WO9921878A1] This invention relates to derivatives of the conopeptide MII, an alpha -4/7 conotoxin peptide, in which amino acid residues are substituted as described herein while maintaining the basic activity of MII. The present invention also relates to the discovery of the 3-dimensional structure of MII, and the relationship of its structure to its specificity to the alpha 3 beta 2 subtype of the neuronal nicotinic acetylcholine receptor (nAChR). The present invention also relates to computer based programs for the expression of the three-dimensional structure of MII and peptide analogs, peptide mimetics or non-peptide mimetics thereof. The structural characteristics may be correlated with biological activity to enable the design of alpha -4/7 conotoxin peptide analogs and peptide mimetics which demonstrate the same specificity to neuronal nAChR. Such analogs and peptide mimetics are useful as cardiovascular agents and for treating or detecting small-cell lung carcinoma (SCLC).

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