

Title (en)

A141S AND G399S MUTATION IN THE OMI/HTRA2 PROTEIN IN PARKINSON'S DISEASE

Title (de)

A141S- UND G399S-MUTATION IM OMI/HTRA2-PROTEIN BEI MORBUS PARKINSON

Title (fr)

MUTATION EN A141S ET G399S DANS LA PROTEINE OMI/HTRA2 DANS LA MALADIE DE PARKINSON

Publication

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Application

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Priority

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Abstract (en)

[origin: WO2005071107A1] The invention relates to a method for the diagnosis of Parkinson's disease in a human being, to nucleic acid molecules used in said method, to the uses thereof in order to detect a nucleic acid molecule coding for human Omi/HtrA2 protein and/or for amplifying the human Omi/HtrA2 gene; to the use of a nucleic acid molecule which codes Omi/HtrA2 protein or sections thereof which is genetically modified in relation to the wild type, and/or one such protein and/or sections thereof for the diagnosis of Parkinson's disease and/or predisposition thereof; a nucleic acid molecule which codes for a human Omi/HtrA2 protein which is genetically modified in relation to the wild type on the 141 and/or 399 amino acid position, in addition to corresponding sections thereof; a host, preferably a transgenic non-human mammal, wherein one such nucleic acid molecule is introduced; a (poly)peptide which is coded by one such nucleic acid molecule; a method which is used to detect substances binding Omi/HtrA2 protein which is genetically modified in relation to the wild type; a substance discovered by means of said method in addition to, preferably, a pharmaceutical composition comprising one such substance. The invention further relates to a kit comprising at least one of the above-mentioned nucleic-acid molecules.

IPC 8 full level

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CPC (source: EP US)

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Citation (search report)

See references of WO 2005071107A1

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