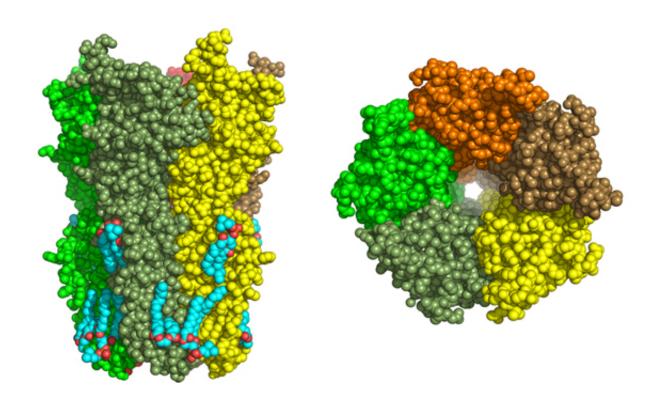


Alzheimer's: Nicotinic receptors as a new therapeutic target

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Representation of space in the nicotinic receptor. Credit: Institut Pasteur

Several scientific studies have indicated that nicotine may be beneficial for memory function. Scientists from the Institut Pasteur and the CNRS set out to shed further light on the properties attributed to nicotine - which is known to have an adverse effect on health - by determining the precise structure of the nicotinic receptors in the hippocampus region of the brain. Using mouse models for Alzheimer's disease, they identified



the $\beta 2$ subunit of the nicotinic receptor as a target that, if blocked, prevents the memory loss associated with Alzheimer's. These results were published in *Neurobiology of Aging* on August 12, 2016.

Alzheimer's disease is characterized by two types of lesion: amyloid plaques and neurofibrillary degeneration. Amyloid <u>beta peptide</u>, naturally present in the brain, builds up over the years as a result of genetic and environmental factors until it forms amyloid plaques. This build-up is toxic for nerve cells and leads to a loss of neuronal structure and to what is known as "neurofibrillary" degeneration, which in turn results in the death of nerve cells.

By the time <u>amyloid plaques</u> appear in the brain, patients' <u>memory loss</u> is already considerable and irreversible. The aim of current research into this disease is therefore to detect amyloid beta peptide at an earlier stage, when it is still soluble, before the plaques appear. In this study, scientists from the Integrative Neurobiology of Cholinergic Systems Unit (Institut Pasteur / CNRS), directed by Uwe Maskos, investigated the toxic effects of a build-up of amyloid beta peptide in the hippocampus and the role of the nicotinic receptor in this same region of the brain.

Acetylcholine receptors, also known as <u>nicotinic receptors</u>, are found in the cell membrane and are sensitive to neurotransmitters. They act like pores for communication between the cell's internal environment and the outside world (see diagram). These receptors are involved in several functions of the nervous system, especially in controlling voluntary movements, memory, attention, sleep, pain and anxiety. Nicotine is an agonist for these receptors, meaning that it can act on these targets instead of acetylcholine.

Nine genes code for nicotinic receptor subunits in the hippocampus, and four of these subunits are part of the receptor involved in this process (β 2 twice, α 2, α 4 and α 5). The scientists are working to determine the



exact composition of this pentamer so that they can assess its suitability as a pharmaceutical target on which therapeutic molecules can be tested.

They decided to focus specifically on the role of the $\beta 2$ subunit of the nicotinic receptor. To do this, they created a mouse model in which the $\beta 2$ subunit was inactivated by blocking the gene coding for it. Memory tests performed on this model showed that the individuals were protected from the <u>toxic effects</u> of amyloid beta peptide and did not develop the cognitive deficit associated with Alzheimer's disease.

The scientists were therefore able to prove that the $\beta 2$ subunit of the nicotinic receptor is the direct target of soluble <u>amyloid beta peptide</u>.

"Characterizing this new therapeutic target will enable us to test molecules that are capable of blocking the $\beta 2$ subunit. The aim is to find a therapeutic molecule that resembles nicotine but does not have the same harmful effects (dependence, premature cell aging, increased heart rate, effects on the gastroenteric system, etc.)," explained Uwe Maskos, the main author of the study.

This research is protected by patent. It is supported by the Institut Pasteur and the CNRS and has also received funding from the BrainTrain European project, the French Foundation for Medical Research, the Greater Paris region and the Gilbert Lagrue foundation.

More information: Sylvia Lombardo et al, A role for β2* nicotinic receptors in a model of local amyloid pathology induced in dentate gyrus, *Neurobiology of Aging* (2016). DOI: 10.1016/j.neurobiologing.2016.06.005

Provided by Institut Pasteur



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