Abstract:

Title: NEW ANTIBODIES SPECIFIC OF THE β-AMYLOID PEPTIDES AND THEIR USES AS DIAGNOSTIC AGENTS OR DRUGS

Abstract: The present invention relates to a monoclonal antibody which specifically binds to the N-terminal region of Aβx peptide, x being comprised from 11 to 42, and recognises neither Aβx1-40nor Aβx1-42 and which presents a high affinity with respect to Aβx peptide, such as determined by an immunological complex formation between the monoclonal antibody and the peptide Aβx.
NEW ANTIBODIES SPECIFIC OF THE β-AMYLOID PEPTIDES AND THEIR USES AS DIAGNOSTIC AGENTS OR DRUGS

The present invention relates to new antibodies specific of the β-amyloid peptides and their uses as diagnostic agents or drugs.

Amyloidosis refers to a pathological condition in a mammal characterized by the presence of amyloid fibers. Amyloid is a generic term referring to a group of diverse but specific protein deposits. All amyloid deposits have common morphologic properties, stain with specific dyes (e.g. Congo red), and have a characteristic red-green birefringent appearance in polarized light after staining. Different amyloids are also characterized by the type of protein present in the deposit. For example, neurodegenerative diseases such as scrapie, bovine spongiform encephalitis, Creutzfeldt-Jakob disease and the like are characterized by the appearance and accumulation of a protease-resistant form of prion protein (referred to as AScr or PrP-27) in the central nervous system. Similarly, Alzheimer's disease, another neurodegenerative disorder, is characterized by neuritic plaques and neurofibrillary tangles. In this case, the plaque and blood vessel amyloid is formed by the deposition of fibrillar β-amyloid protein.

Alzheimer's disease (AD) is the most common type of senile dementia and is believed to be responsible for 40-60% of all cases of dementia. The incidence of AD increases with age, affecting 1 out of 10 persons older than age 65 and nearly 1 out of 2 persons older than age 85. Overall, the natural history of the disease can be characterized as an irreversibly progressive brain disorder that ultimately results in devastating memory loss, profound behavioural and personality changes, and severely damaged cognitive abilities. These impairments are related to the underlying death of brain cells and the breakdown of communication between them. In view of the large expenses for health care systems that must provide institutional and ancillary care for the AD patients, the impact of AD on society and on national economies is enormous.


(i) at the intracellular level, the neuronal cytoskeleton in AD patients is progressively disrupted and replaced by neurofibrillary tangles (NFTs) composed of paired
helical filaments (PHF);
(ii) at the extracellular level, amyloid plaques are formed by deposits of fibrillary β-amyloid (Aβ).

Aβ is a major component of the senile plaques. Aβ is a small peptide found mainly in two sizes, consisting of 40 (Aβ_{40}) and 42 (Aβ_{42}) amino acids respectively, and in minor amounts in other sizes. Aβ is known to be metabolised from the proteolytic cleavage of APP (Saido, 2000). Degradation of amyloid-β peptide: a key to Alzheimer pathogenesis, prevention and therapy. Neurosci. News 3: 52-62), a large transmembrane protein with known, although not completely clear, neurotrophic functions (Seo et al, 2001). Effects of nicotine on APP secretion and Abeta- or CT(105)-induced toxicity. Biol. Psychiatry 49: 240-247). APP can be cleaved via two main routes, a major non-amyloidogenic route and a minor second, amyloidogenic route that yields Aβ as ultimate product.

The main pathway for catabolism of APP is through cleavage by α-secretase at a single site in APP near the center of the β-amyloid peptide region (Esch et al., 1990). Cleavage of amyloid beta peptide during constitutive processing of its precursor. Science 248: 1122-1124; Sisodia, (1992), Beta-amyloid precursor protein cleavage by a membrane-bound protease. Proc. Natl. Acad. Sci. USA 89: 6075-6079). The products yielded by this route are a large N-terminal region of APP (APPs α) and a membrane associated C-terminal fragment (C83), which is subsequently hydrolysed by γ-secretase to yield the nearly unknown small p3 peptide. This is the non-amyloidogenic route because the cleavage site is located approximately in the middle of the Aβ sequence, with no possibility of Aβ formation. The second APP processing pathway is the N- and C-terminal cleavage of APP by β- and γ-secretase (Fig. 1). The resulting molecules of these two proteolytic steps are the central fragments of APP, Aβ_{10} and Aβ_{42}, Aβ_{10} being the more abundant of the whole Aβ formed (Conde, (2002), β-amyloid peptide as a target for treatment of Alzheimer's disease. Expert Opin. Ther. Patents 12: 503-512). β-secretase cleaves at the amino terminus of the β-amyloid peptide and occurs first, followed by γ-secretase, which releases the carboxy terminus of the peptide. This statement is based upon the observation that C-terminal fragments produced by β-secretase cleavage are readily apparent in cells, whereas APP fragments corresponding to a single C-terminal γ cleavage are not (Haass et al., (1992), Amyloid beta-peptide is produced


Immunotherapy for Alzheimer with antibodies directed to the β-amyloid peptide is a

However since β-amyloid is a normal constituent of normal tissue and biological fluids severe side effects have halted the first clinical trials (Orgogozo et al., (2003), Subacute meningoencephalitis in a subset of patients with AD after Abeta 42 immunization. Neurology 61: 46-54).

It has been shown by Sergeant et al. (Sergeant et al., (2003), Truncated beta-amyloid peptide species in pre-clinical Alzheimer's disease as new targets for the vaccination approach. Journal of Neurochemistry 85: 1581-1591) that 60% of all Aβ species of early amyloid deposits are amino -truncated Aβ species.

The international application WO 2004/029630 discloses a monoclonal antibody which specifically recognises Aβn-x peptides and does not recognise Aβi-x peptide (x being 40 or 42).

The peptides used for immunization are the first 5 to 7 human amino-acids of the β secretase_1 cleavage site (the β secretase cleaves the APP protein at Glu 11). Nevertheless, the Aβn-x peptides are not the Aβ peptides observed at the very early stages of amyloid deposition (Sergeant et al., Truncated beta-amyloid peptide species in pre-clinical Alzheimer's disease as new targets for the vaccination approach. Journal of Neurochemistry 85, 1581-1591 (2003). Moreover, Aβn-x is not a pathological species as it is produced from cleavage of β secretase and N-truncated forms of Aβ42 are much more abundant than the full-size Aβ42 and Aβn-x species.

International application WO 2004/013172 relates to polyclonal antibodies directed toward truncated beta-amyloid peptide species Aβm-n, m being comprised from 1 to 10 and n being comprised from m + 3 to m + 15. The peptides used for immunization are Aβn-i2, Aβi-13, Aβn-i8, Aβn-i6. Nevertheless antibodies of this application are polyclonal, having a moderate affinity.

Murayama K. S. et al. (Murayama K. S. et al., (2007), A novel monoclonal antibody specific for the amino -truncated β-amyloid Aβ54-42 produced from caspase-cleaved amyloid precursor protein, 161: 244-249) disclose a monoclonal antibody obtained with peptide
A βs-12 immunization, and that recognises specifically A βs_4o and not A βi_4o.

Two other antibodies are described in this paper:
- mouse monoclonal antibody 4G8 specific for A β17-24;
- rabbit polyclonal antibody Ab-I specific for A β15-30.

Nevertheless these two antibodies are not specific and recognize A βs_4o and A βi_4o.

One of the aims of the present invention is to provide an antibody which specifically binds to N-terminal region of A β8-x peptide and does not recognise A βi-x (x being 40 or 42) and is able to specifically recognise the early stages peptides of β-amyloid deposit.

A further aim of the present invention is to provide synthetic peptides useful to produce an immune response against N-truncated peptides of A β and consequently useful for the prevention or the treatment of Alzheimer’s disease.

The present invention also relates to a process of preparation in order to obtain an antibody which specifically binds to N-terminal region of A β8-x peptide.

The present invention further relates to a method for determining amyloid burden in mammals.

A further aim of the present invention is to provide a method for determining, in a mammal, the susceptibility to a disease associated with A β formation and/or aggregation such as Alzheimer’s disease, for determining, in a mammal, the risk of developing a disease associated with β-amyloid formation and/or aggregation such as Alzheimer’s disease, for screening of the clearance of β-amyloid deposition in mammal, or for predicting the level of β-amyloid burden in a mammal.

The present invention also relates to therapeutic or vaccine compositions comprising an antibody specific to N-terminal region of A β8-x peptide or comprising synthetic peptides with a free N-terminal-end mimicking the free N-terminal-end of N-truncated A β peptides, useful for the preparation of a drug or a vaccine intended for the prevention or the treatment of Alzheimer disease.

The present invention further relates the use of an antibody for the preparation of a drug or a vaccine intended for the prevention or the treatment of Alzheimer disease.

Therefore, the present invention relates to an antibody which specifically binds to the N-terminal region of A β8-x peptide, x being comprised from 11 to 42, and recognises neither A βs_4o nor A βi_42.
The term "antibody" is used to denote polyclonals or monoclonals specific to \( \text{A}\beta_x \) and also include fragments or molecules which mimick the monoclonals specific to \( \text{A}\beta_x \), and in particular epitope binding fragment. Fragments or molecules may be derived from monoclonals by recombinant DNA techniques or by enzymatic or chemical methods and may exhibit similar binding characteristics compared to the monoclonal for an antigen fragment.

By "polyclonal antibody" is meant an antibody derived from different B-cell lines.

By "monoclonal antibody" is meant an antibody coming from only one type of cell, the hybridoma cell.

By "hybridoma" cell is meant a cell fusion which will continually produce antibodies, i.e. tumor cells that can replicate endlessly which are fused with mammalian cells.

The antibodies of the present invention include both the full length antibodies discussed above, as well as epitope-binding fragments thereof. As used herein, "antibody fragments" include any portion of an antibody that retains the ability to bind to the epitope recognized by the full length antibody, generally termed "epitope-binding fragments." Examples of antibody fragments include, but are not limited to, Fab, Fab' and F(ab')2, Fd, single-chain Fvs (scFv), single-chain antibodies, disulfide-linked Fvs (dsFv) and fragments comprising either a VL or VH region. Epitope-binding fragments, including single-chain antibodies, may comprise the variable region(s) alone or in combination with the entirety or a portion of the following: hinge region, CH1, CH2, and CH3 domains.

Such fragments may contain one or both Fab fragments or the F(ab')2 fragment. Further, the fragments may be or may combine members of any one of the following immunoglobulin classes: IgG, IgM, IgA, IgD, or IgE, and the subclasses thereof.

Fab and F(ab')2 fragments may be produced by proteolytic cleavage, using enzymes such as papain (Fab fragments) or pepsin (F(ab')2 fragments).

The "single-chain FVs" ("scFvs") fragments are epitope-binding fragments that contain at least one fragment of an antibody heavy chain variable region (\( V_H \)) linked to at least one fragment of an antibody light chain variable region (\( V_L \)). The linker may be a short, flexible peptide selected to assure that the proper three-dimensional folding of the \( V_L \) and \( V_H \) regions occurs once they are linked so as to maintain the target molecule binding-specificity of the whole antibody from which the single-chain antibody fragment is derived. The
carboxyl terminus of the $V_L$ or $V_H$ sequence may be covalently linked by a linker to the amino acid terminus of a complementary $V_L$ or $V_H$ sequence.

Single-chain antibody fragments of the present invention contain amino acid sequences having at least one of the variable or complementarity determining regions (CDRs) of the whole antibodies described in this specification, but lack some or all of the constant domains of those antibodies. These constant domains are not necessary for antigen binding, but constitute a major portion of the structure of whole antibodies. Single-chain antibody fragments may therefore overcome some of the problems associated with the use of antibodies containing a part or all of a constant domain. For example, single-chain antibody fragments tend to be free of undesired interactions between biological molecules and the heavy-chain constant region, or other unwanted biological activity. Additionally, single-chain antibody fragments are considerably smaller than whole antibodies and may therefore have greater capillary permeability than whole antibodies, allowing single-chain antibody fragments to localize and bind to target antigen-binding sites more efficiently. Also, antibody fragments can be produced on a relatively large scale in prokaryotic cells, thus facilitating their production. Furthermore, the relatively small size of single-chain antibody fragments makes them less likely to provoke an immune response in a recipient than whole antibodies.

Single-chain antibody fragments may be generated by molecular cloning, antibody phage display library or similar techniques well known to the skilled artisan. These proteins may be produced, for example, in eukaryotic cells or prokaryotic cells, including bacteria. The epitope-binding fragments of the present invention can also be generated using various phage display methods known in the art. In phage display methods, functional antibody domains are displayed on the surface of phage particles which carry the polynucleotide sequences encoding them. In particular, such phage can be utilized to display epitope-binding domains expressed from a repertoire or combinatorial antibody library (e.g., human or murine). Phage expressing an epitope-binding domain that binds the antigen of interest can be selected or identified with antigen, e.g., using labelled antigen bound or captured to a solid surface or bead. Phages used in these methods are typically filamentous phage including fd and M13 binding domains expressed from phage with Fab, Fv or disulfide-stabilized Fv antibody domains recombinantly fused to either the phage gene III or gene VIII protein.

Examples of phage display methods that can be used to make the epitope-binding

After phage selection, the regions of the phage encoding the fragments can be isolated and used to generate the epitope-binding fragments through expression in a chosen host, including mammalian cells, insect cells, plant cells, yeast, and bacteria, using recombinant DNA technology, e.g., as described in detail below. For example, techniques to recombinantly produce Fab, Fab' and F(ab')2 fragments can also be employed using methods known in the art such as those disclosed in WO 92/22324; Mullinax et al, 1992, *BioTechniques*, **12**(6): 864-869; Sawai et al, 1995, *AJRI*, **34**: 26-34; and Better et al, 1988, *Science*, **240**:1041-1043; said references incorporated by reference in their entireties. Examples of techniques which can be used to produce single-chain Fvs and antibodies include those described in U.S. Pat. Nos. 4,946,778 and 5,258,498; Huston et al, 1991, *Methods in Enzymology*, **203**: 46-88; Shu et al, 1993, *PNAS*, **90**: 7995-7999; Skerra et al, 1988, *Science*, **240**:1038-1040.

Also included within the scope of the invention are functional equivalents of the antibodies specifically disclosed in the present application. The term "functional equivalents" includes antibodies with homologous sequences, chimeric antibodies, artificial antibodies and modified antibodies, for example, wherein each functional equivalent is defined by its ability to specifically bind to the N-terminal region of Aβ4-8 peptide as defined above. The skilled artisan will understand that there is an overlap in the group of molecules termed "antibody fragments" and the group termed "functional equivalents." Methods of producing functional equivalents are known to the person skilled in the art and are disclosed, for example, in WO 93/21319, EP 239,400; WO 89/09622; EP 338,745; and EP 332,424, which are incorporated in their respective entireties by reference.

Artificial antibodies include scFv fragments, diabodies, triabodies, tetrabodies and
mru (see reviews by Winter, G. and Milstein, C., 1991, Nature, 349: 293-299; Hudson, P.J., 1999, Current Opinion in Immunology, 11: 548-557), each of which has antigen-binding ability. In the single chain Fv fragment (scFv), the V_{H} and VL domains of an antibody are linked by a flexible peptide. Typically, this linker peptide is about 15 amino acid residues long. If the linker is much smaller, for example 5 amino acids, diabodies are formed, which are bivalent scFv dimers. If the linker is reduced to less than three amino acid residues, trimeric and tetrameric structures are formed that are called triabodies and tetrabodies. The smallest binding unit of an antibody is a CDR, typically the CDR2 of the heavy chain which has sufficient specific recognition and binding that it can be used separately. Such a fragment is called a molecular recognition unit or mru. Several such mrus can be linked together with short linker peptides, therefore forming an artificial binding protein with higher avidity than a single mru.

The functional equivalents of the present application also include modified antibodies, e.g., antibodies modified by the covalent attachment of any type of molecule to the antibody. For example, modified antibodies include antibodies that have been modified, e.g., by glycosylation, acetylation, pegylation, phosphorylation, amidation, derivatization by known protecting/blocking groups, proteolytic cleavage, linkage to a cellular ligand or other protein, etc. The covalent attachment does not prevent the antibody from generating an anti-idiotypic response. These modifications may be carried out by known techniques, including, but not limited to, specific chemical cleavage, acetylation, formylation, metabolic synthesis of tunicamycin, etc. Additionally, the modified antibodies may contain one or more non-classical amino acids.

Functional equivalents may be produced by interchanging different CDRs on different chains within different frameworks. Thus, for example, different classes of antibody are possible for a given set of CDRs by substitution of different heavy chains, whereby, for example, IgG1-4, IgM, IgA1-2, IgD, IgE antibody types and isotypes may be produced. Similarly, artificial antibodies within the scope of the invention may be produced by embedding a given set of CDRs within an entirely synthetic framework.

Functional equivalents may be readily produced by mutation, deletion and/or insertion within the variable and/or constant region sequences that flank a particular set of CDRs, using a wide variety of methods known in the art.
The antibody specific for said N-terminal region of $\text{A}_{8-\times} \text{B}_{\beta}$ peptide can be detected by an immunoassay. As used herein, an "immunoassay" is an assay that utilizes an antibody to specifically bind to the antigen (i.e. the N-terminal region of $\text{A}_{8-\times} \text{B}_{\beta}$ peptide). The immunoassay is thus characterized by detection of specific binding of proteins to antibodies.

The expressions "specifically binds to", "specific recognition", "specifically recognizing", "specifically reacting with" or "specifically forming an immunological reaction with" refer to a binding reaction by the antibody to the N-terminal region of $\text{A}_{8-\times} \text{B}_{\beta}$ peptide, which is determinative of the presence of the N-terminal region of $\text{A}_{8-\times} \text{B}_{\beta}$ peptide in the sample tested, in the presence of a heterogeneous population of other proteins and/or other biologies. The specificity can be determined by a Luminex assay. Using this assay, antibodies of the invention present a high specificity on $\text{A}_{8-\times} \text{B}_{\beta}$ peptide, i.e. the Mean Fluorescence Intensity (MFI) obtained with an antibody is much more higher on a $\text{A}_{8-\times} \text{B}_{\beta}$ peptide than on a non-specific peptide like $\text{A}_{6-\times} \text{B}_{\gamma}$ peptide, for example MFI = 1822 with TeiAl1 on $\text{A}_{8-\times} \text{B}_{\beta}$ peptide and only 24 on $\text{A}_{8-\times} \text{B}_{\gamma}$ peptide (see example 3 and table 3).


Thus, under the designated immunoassay conditions, the specified antibody preferentially binds to a N-terminal region of $\text{A}_{8-\times} \text{B}_{\beta}$ peptide of the invention while binding to other proteins or protein isoforms does not occur in significant amounts.

In particular, the specified antibody does not bind to $\text{A}_{8-\times} \text{B}_{\beta}$ peptide and therefore would not present the severe side effects observed with antibodies against $\text{A}_{8-\times} \text{B}_{\beta}$ peptide when used for therapeutic purposes (see example 5).

Such a response can be an active response induced by administration of immunogen or a passive response induced by administration of antibody or primed T-cells. A cellular immune response is elicited by the presentation of polypeptide epitopes in association with
Class I or Class II MHC molecules to activated antigen-specific CD4 T helper cells and/or CD8+ cytotoxic T-cells. The response may also involve activation of monocytes, macrophages, NK cells, basophils, dendritic cells, astrocytes, microglia cells, eosinophils or components of innate immunity.

An "immunogenic agent" or "immunogen" is capable of inducing an immunological response directed against itself upon administration to a recipient mammal, optionally in conjunction with an adjuvant.

In a preferred embodiment, said antibody presents a high specificity for the free N-terminal end of $A\beta_{8-10}$ peptide.

The expression "free N-terminal end" refers to an unblocked N-terminal end i.e. an amino-acid having a NH$_2$ terminal end.

Antibodies of the invention can be polyclonal having a high specificity or monoclonal having a high specificity.

In another preferred embodiment, said antibody presents a high affinity with respect to $A\beta_{8-10}$ peptide.

The term "affinity" refers to the strength of the binding of the antibody to the N-terminal region of $A\beta_{8-10}$ peptide, i.e., how tightly the antibody binds to the N-terminal region of $A\beta_{8-10}$ peptide.

Antibodies of the invention can be polyclonal having a high affinity or monoclonal having a high affinity.

The affinity of the monoclonal antibody of the invention to the N-terminal region of $A\beta_{8-10}$ peptide is determined by the bridging assay test (see example 3). OD values below 1 represent low affinity and above 1 show high affinity of the monoclonals to its target.

In another advantageous embodiment, antibodies of the invention can be polyclonal with a high specificity and a high affinity, or monoclonal with a high specificity and a high affinity.

In a more preferred embodiment, said antibody specifically target parenchymal amyloid deposits of $A\beta_{8-10}$ peptide in the brain and does not interact with vascular amyloid deposits.

The induction of an immune response is "active" when an immunogen is administered to induce antibodies or T-cells reactive against the immunogen. The induction
of an immune response is "passive" when an antibody is administered that itself binds to the N-terminal truncated Aβ8–x peptide in the mammal.

One of the side-effects of the passive immunization is the frequency of microhemorrhages. Such increase in the number of microhemorrhages may be explained by the fixation of injected antibodies to the aggregated Aβ peptides within vessel walls (see exemple 5).

Therefore, antibodies of the invention specifically targeting parenchymal amyloid deposits and not vascular amyloid deposits would not present the severe side effects observed with antibodies against Aβ42 peptide (see example 5).

In a preferred embodiment, the present invention relates to an antibody wherein x is comprised from 15 to 42, in particular a monoclonal antibody.

In a preferred embodiment, the present invention relates to a monoclonal antibody which specifically binds to the N-terminal region of Aβ8–x peptide, wherein the variable region comprises one of the following pairs of amino acid sequences, corresponding respectively to the light and heavy chain:

Areas in grey corresponds to the Complementarity Determining Regions of the light chain (CDR-Lx) or the heavy chain (CDR-Hx)

- Antibody TeiA 1.6 (secreted by hybridoma IGH521)

**Light chain variable region:**

<table>
<thead>
<tr>
<th>CDR-L1</th>
<th>CDR-L2</th>
<th>CDR-L3</th>
</tr>
</thead>
<tbody>
<tr>
<td>SSLTVTAGEKVTMSCKSSQLLAGRYQKNYLTYWQPKPGQPPKLIYWAST</td>
<td>RDSGVDRFTGSGSTDFTLTISSVQAESLAVYQCNDTYPLTFAG</td>
<td></td>
</tr>
</tbody>
</table>

(SEQ ID NO : 1)

**Heavy chain variable region:**

<table>
<thead>
<tr>
<th>CDR-H1</th>
<th>CDR-H2</th>
<th>CDR-H3</th>
</tr>
</thead>
<tbody>
<tr>
<td>GGLVQPGGSLRLSCAISGFTSFYMWERQPGKRLWIAASRNKANDYT</td>
<td>EYSASVKGVRIVSRDTSQSLYQMNLRAEDTAYTYTAMDYWGQ</td>
<td></td>
</tr>
</tbody>
</table>
GTSVTSS (SEQ ID NO: 2)

• Antibody TeiA 1.7 (secreted by hybridoma IGH522)

Light chain variable region:

CDR-L1

SSLTVTAGEKVTMNCKSSQKLLNSGHQYHHTLTFQQKPGQPPKLLIYWAST

CDR-L2

CDR-L3

RISGVPDFRFIGSGSTDTFTLTINSVQAEDLAVYYCQNYRYTILTFGAG

(SEQ ID NO:3)

Heavy chain variable region:

CDR-H1

GGLVQPGGSLRLSCATSGFTFSDFYMIWVRQPPWRLEWIAASRØKAKØTIT

CDR-H2

EYSASYKGRFIVSRDTQSIFYLQMNLRSDETAIYYCATYFSYAMDYWGLT

CDR-H3

SVTVSS (SEQ ID NO: 4)

• Antibody TeiA 1.8 (secreted by hybridoma IGH523)

Light chain variable region:

CDR-L1

SSLAVTAGERVTSCKSSLTLNNSGQNTNYLTWYQQKPGQPPKLLIYWAST

CDR-L2

CDR-L3

ISGVPDFRFTGSGTDFTLTISSVQAEDLAVYYCQKDYSYPLTFGAG

(SEQ ID NO:5)

Heavy chain variable region:

CDR-H1

GGLVQPGGSLRLSCATAGFTFDQYMSWVRQPPGKALEWLATIRNKAKGFT

CDR-H2

CDR-H3

HYSASYKGRFTISRDNQSILYLMSTLRAGDSATYYCAVYGNYAAMPYWG
QGTSVNVS\textit{(SEQ ID NO : 6)}

- Antibody TeiA 2b.6 (secreted by hybridoma IGH524)

Light chain variable region:

\begin{align*}
\text{CDR-L1} & \quad \text{CDR-L2} \\
\text{SSLTVTAGEKVTMSCKSSQSLFMGRQTM} & \quad \text{LTFWQPQRPPGQAPKLLIYWASTR} \\
\text{CDR-L3} & \\
\text{GSGVPDRFTGS} & \quad \text{GILYLSAQMSALRAEDTAIYYCAIYRR} \\
\text{TEYSASVKGR} & \quad \text{YAMDYWGQ} \\
\end{align*}

Heavy chain variable region:

\begin{align*}
\text{CDR-H1} & \quad \text{CDR-H2} \\
\text{GGLVQPGGS} & \quad \text{LRLSCATSGFTFTDFYMEVRQPPG} \\
\text{CDR-H3} & \quad \text{KRLEWIAASR} \\
\text{TEYSASVKGR} & \quad \text{FIVSRDTSQGILYLQ} \\
\text{GTS} & \quad \text{MSALRAEDTAIYYCAIY} \\
\text{TVSS (SEQ ID NO : 7)} & \quad \text{YAMDYWGQ} \\
\text{GTS} & \quad \text{TVSS (SEQ ID NO : 8)} \\
\end{align*}

- Antibody TeiA 1.1 (secreted by hybridoma IGH525)

Light chain variable region:

\begin{align*}
\text{CDR-L1} & \quad \text{CDR-L2} \\
\text{SSLTVTAGEKVTMSC} & \quad \text{SSQSLFNSGTQTM} \\
\text{CDR-L3} & \quad \text{MYiLTWYQQKPQPPKL} \\
\text{GSGVPDRFTGS} & \quad \text{LTVSSVQAEDLA} \\
\text{TEYSASVKGR} & \quad \text{VYQCNDTY} \\
\text{GTS} & \quad \text{TVSS (SEQ ID NO:9)} \\
\text{GTS} & \quad \text{TVSS (SEQ ID NO : 9)} \\
\end{align*}

Heavy chain variable region:

\begin{align*}
\text{CDR-H1} & \quad \text{CDR-H2} \\
\text{GGLVQPGGS} & \quad \text{LRLSCATSGF} \\
\text{CDR-H3} & \quad \text{FTFDFTLVSSVQAED} \\
\end{align*}
As used herein, an "antibody" refers to a protein consisting of one or more polypeptides substantially encoded by immunoglobulin genes or fragments of immunoglobulin genes.

In a preferred embodiment, the CDR of the light and heavy chain of the variable region of the antibody defined above comprises one of the following amino acid sequences:

- Antibody TeiA 1.6 (IGH521 sequence)
  CDR of the light chain variable region:
  CDR-L1:
  KSSQSSLAGRYQKNYLT (SEQ ID NO : 11)
  CDR-L2:
  WASTRESG (SEQ ID NO : 12)
  CDR-L3:
  QNDYTYFILT (SEQ ID NO : 13)

  CDR of the heavy chain variable region:
  CDR-H1:
  GPTFSDFYMI (SEQ ID NO : 14)
  CDR-H2:
  ASRNKANDYTEYSASVKG (SEQ ID NO : 15)
  CDR-H3
  YHDYAMDY (SEQ ID NO : 16)

- Antibody TeiA 1.7 (IGH522 sequence)
  CDR of the light chain variable region:
  CDR-L1:
  KSSQSSLNLNSGNQVNYLT (SEQ ID NO : 17)
  CDR-L2:
  WASTRESG (SEQ ID NO : 18)
CDR-L3:
QNDYRYPLT (SEQ ID NO : 19)

CDR of the heavy chain variable region:
CDR-H1:
GFTFSDFYM1 (SEQ ID NO : 14)
CDR-H2:
ASRDKAKDYTTEYSASVKO (SEQ ID NO : 20)
CDR-H3:
YFSYAMDY (SEQ ID NO : 21)

- Antibody TeiA 1.8 (IGH523 sequence)

CDR of the light chain variable region:
CDR-L1:
KSSLTLLNSGSQTNYLT (SEQ ID NO : 22)
CDR-L2:
WASTRESG (SEQ ID NO : 18)
CDR-L3:
QNDYSYPLT (SEQ ID NO : 23)

CDR of the heavy chain variable region:
CDR-H1:
OFTFTDQYMS (SEQ ID NO : 24)
CDR-H2:
TMNKAKGFTTIYSASVKG (SEQ ID NO : 25)
CDR-H3:
YGNYAMDY (SEQ ID NO : 26)

- Antibody TeiA 2b.6 (IGH524 sequence)
KSSQSLFNSGitQTNYLT (SEQ ID NO : 27)
CDR-L2:
WASTRGS (SEQ ID NO : 28)
CDR-L3:
QKDYTYTiLT (SEQ ID NO : 13)

CDR of the heavy chain variable region:
CDR-H1:
GFTFTDFYME (SEQ ID NO : 29)
CDR-H2:
ASRNKANOYTTEYSASVKG (SEQ ID NO : 30)
CDR-H3:
YRYYAMDY (SEQ ID NO : 31)

• Antibody TeiA 1.1 (IGH525 sequence)

CDR of the light chain variable region:
CDR-L1:
TSSQSLFNSTQTMYLT (SEQ ID NO : 32)
CDR-L2:
WASTRESG (SEQ ID NO : 18)
CDR-L3:
QNDYTYFLT (SEQ ID NO : 13)

CDR of the heavy chain variable region:
CDR-H1:
GFTFTDFYME (SEQ ID NO : 33)
CDR-H2:
ASRNKNYDYKTEYSASVKG (SEQ ID NO : 34)
CDR-H3:
YRHYAMDY (SEQ ID NO : 35)

The CDRs of the present invention include not only those completely identical but also variants so long as the specificity to Aβ_{x} peptide is maintained. That is, the CDR amino
acid sequences in which one or more amino acid residues are modified may also be used as the CDR sequence. The modified amino acid residues in the amino acid sequences of the CDR variant are preferably 30% or less, more preferably 20% or less, most preferably 10% or less, within the entire CDR.

Therefore, any antibody, fragment, molecule or ligand comprising at least one of the indicated CDR's or homologous sequences can be used.

The CDRs are of primary importance for epitope recognition and antibody binding. However, changes may be made to the residues that comprise the CDRs without interfering with the ability of the antibody to recognize and bind its cognate epitope. For example, changes that do not affect epitope recognition, yet increase the binding affinity of the antibody for the epitope may be made.

Several studies have surveyed the effects of introducing one or more amino acid changes at various positions in the sequence of an antibody, based on the knowledge of the primary antibody sequence, on its properties such as binding and level of expression (Yang, W. P. et al., 1995, J. Mol. Biol., 254: 392-403; Rader, C. et al., 1998, Proc. Natl. Acad. Sci. USA, 95: 8910-8915; Vaughan, T. J. et al., 1998, Nature Biotechnology, 16: 535-539).


By a similar directed strategy of changing one or more amino acid residues of the antibody, the antibody sequences described in this invention can be used to develop
antibodies which specifically bind to the N-terminal region of Aβ₅₋ₓ peptide as defined above with improved functions, including improved affinity to the N-terminal region of Aβ₅₋ₓ peptide.

Preferred amino acid substitutions are those which: (1) reduce susceptibility to proteolysis, (2) reduce susceptibility to oxidation, (3) alter binding affinity for forming protein complexes, and (4) confer or modify other physico-chemical or functional properties of such analogs. Analogs can include various muteins of a sequence other than the naturally-occurring peptide sequence. For example, single or multiple amino acid substitutions (preferably conservative amino acid substitutions) may be made in the naturally-occurring sequence (preferably in the portion of the polypeptide outside the domain(s) forming intermolecular contacts. A conservative amino acid substitution should not substantially change the structural characteristics of the parent sequence (e.g., a replacement amino acid should not tend to break a helix that occurs in the parent sequence, or disrupt other types of secondary structure that characterizes the parent sequence). Examples of art-recognized polypeptide secondary and tertiary structures are described in Proteins, Structures and Molecular Principles (Creighton, Ed., W. H. Freeman and Company, New York (1984)); Introduction to Protein Structure (C. Branden and J. Tooze, eds., Garland Publishing, New York, N. Y. (1991)); and Thornton et al., 1991, Nature, 354: 105, which are each incorporated herein by reference.

Improved antibodies also include those antibodies having improved characteristics that are prepared by the standard techniques of animal immunization, hybridoma formation and selection for antibodies with specific characteristics.

It is also possible to use cell lines specifically engineered for production of improved antibodies. In particular, these lines have altered regulation of the glycosylation pathway, resulting in antibodies which are poorly fucosylated or even totally defucosylated. Such cell lines and methods for engineering them are disclosed in e.g. Shinkawa et al. (2003, J. Biol. Chem. 278(5): 3466-3473), Ferrara et al. (2006, J. Biol Chem. 281(8): 5032-5036; 2006, Biotechnol Bioeng. 93(5): 851-61), EP 1331266, EP 1498490, EP 1498491, EP 1676910, EP 1792987, and WO 99/54342.

In another preferred embodiment, the present invention relates to an antibody which specifically binds to the N-terminal region of Aβ₅₋ₓ peptide, said antibody being labelled with
a compound chosen from the group comprising: a radionuclide, a fluor, an enzyme label, an enzyme substrate, an enzyme co-factor, enzyme inhibitor and a hapten.

The particular label or detectable group used in the assay is generally not a critical aspect of the invention, so long as it does not significantly interfere with the specific binding of the antibody used in the assay. The detectable group can be any material having a detectable physical or chemical property. Such detectable labels have been well developed in the field of immunoassays and, in general, almost any label useful in such methods can be applied to the method of the present invention.

Thus, a label is any composition detectable by spectroscopic, photochemical, biochemical, immunochemical, electrical, optical, radiological or chemical means. Useful labels in the present invention include but are not limited to magnetic beads (e.g. Dynabeads™), fluorescent dyes (e.g. fluorescein isothiocyanate, texas red, rhodamine), radiolables (e.g. \(^3\)H, \(^125\)I, \(^35\)S, \(^14\)C, or \(^32\)P), enzymes (e.g. horseradish peroxidase, alkaline phosphatase and others commonly used in an ELISA), and colorimetric labels such as colloidal gold, colored glass or plastic (e.g. polystyrene, polypropylene, latex, etc.) beads.

The label may be coupled directly or indirectly to the desired component of the assay according to methods well known in the art. As indicated above, a wide variety of labels may be used, with the choice of label depending on the sensitivity required, the ease of conjugation with the compound, stability requirements, the available instrumentation and disposal provisions. Non-radioactive labels are often attached by indirect means.

Generally, a ligand molecule (e.g. biotin) is covalently bound to the antibody. The ligand then binds to an anti-ligand (e.g. streptavidin) molecule, which is either inherently detectable or covalently bound to a signal system, such as a detectable enzyme, a fluorescent compound, or a chemiluminescent compound. A number of ligands and anti-ligands can be used. Where a ligand has a natural anti-ligand, for example, biotin, thyroxine, and Cortisol, it can be used in conjunction with the labeled, naturally occurring anti-ligands. Alternatively, a haptenic or antigenic compound can be used in combination with an antibody.

The antibodies can also be conjugated directly to signal-generating compounds, for example, by conjugation with an enzyme or fluorophore. Enzymes of interest as labels will primarily be hydrolases, particularly phosphatases, esterases and glycosidases, or oxidoreductases, particularly peroxidases.

Fluorescent compounds include fluorescein and its derivatives, rhodamine and its
derivatives, dansyl, umbelliferone, etc. Chemiluminescent compounds include luciferin, and 2,3-dihydrophtalazinediones, for example, luminol. A review of other labeling or signal producing systems is available in US patent No. 4,391,904.

Means for detecting labels are well known in the art. Thus, for example, where the label is a radioactive label, means for detection include a scintillation counter or photographic film as in autoradiography. Where the label is a fluorescent label, it may be detected by exciting the fluorophore with the appropriate wavelength of light and detecting the resulting fluorescence. The fluorescence may be detected visually, by means of a photographic film, by the use of electronic detectors such as charge coupled devices (CCDs) or photomultipliers and the like.

Similarly, enzyme labels may be detected by providing the appropriate substrates for the enzyme and detecting the resulting reaction product. Finally simple colorimetric labels may be detected simply by observing the color associated with the label.

In a preferred embodiment, the monoclonal antibody is a humanised antibody.

By "humanised antibody" is meant a genetically engineered antibody in which the minimum mouse part from a murine antibody is transplanted onto a human antibody; generally humanized antibodies are 5-10% mouse and 90-95% human.

Humanized antibodies have the advantage to counter the HAMA (human Antibodies directed against mouse antibodies) and HACA (human antibodies directed against chimeric antibodies) responses seen with murine and chimeric antibodies and exhibit minimal or no response of the human immune system against them.

According to another aspect, the present invention concerns hybridomas producing monoclonal antibodies as above defined, i.e. which specifically binds to N-terminal region of Aβ1-40 peptide and does not recognise neither Aβ1-40 nor Aβ1-42, and in particular the variable region of which comprises one of the pairs of the amino acid sequences defined above and which presents a high specificity.

In a preferred embodiment, the above defined hybridoma have been deposited on August 23, 2007, at:

BCCM / LMBP Plasmid Collection
Department of Molecular Biology
Ghent University
'Fiers-Schell-Van Montagu’ building
Technologiepark 927
under the following Accession No:

TeiA 1.6 or 2.6F4C2 (IGH521) -> LMBP 6594CB
TeiA 1.7 or 2.8A3F8 (IGH522) -> LMBP 6595CB
TeiA 1.8 or 1.3B12H3 (IGH523) -> LMBP 6596CB
TeiA 2b.6 or 2.13E5E4 (IGH524) --> LMBP 6597CB
TeiA 1.1 or 3.46B I0E7 (IGH 525) -> LMBP 6598CB

According to another aspect, the present invention refers to a peptide preparation to generate an immune response giving rise to an antibody production which is efficient to reduce the amyloid deposits and to isolate a monoclonal antibody.

By "peptide preparation" is meant a short synthetic peptide with a free N-terminal-end which mimics the free N-terminal-end of N-truncated Aβ peptides.

The peptide used is the following: Aβ 8-x mimicking peptide: SGYGVHHGC-KLH

Whereas KLH is keyhole limpet hemocyanin which is coupled to the cysteine by disulfide bridge. The sequence corresponding to Aβ is underlined and followed by spacer amino-acid that is a glycine. Aβ 8-x is similar to IGP-21 19 (PG 127) Table 2

The peptide preparation was mixed in phosphate saline buffer and added with Freund adjuvant for intraperitoneal injections (Figure 2). After 24 weeks, the immune response was analyzed by TAPIR (Figure 3) and the effect on the amyloid load was determined by western blotting (Figure 4).

According to another aspect, the present invention relates to a process of preparation of the above defined antibody which specifically binds to the N-terminal region of Aβ8-x peptide and does not recognise Aβ42, x being comprised from 11 to 42, in particular 15-42, which present a high specificity, and comprising an immunisation step of an appropriate animal with a Aβ8-x peptide and a T-helper epitope, in particular with a Aβ8-x peptide fused with a T-helper epitope, or a Aβ8-x branched peptide, in particular a Aβ8-15 peptide.

The expression "Aβ8-x peptide fused with a T-helper epitope" refers to the linking of the Aβ8-x peptide with a T-helper epitope according to Livingston et al., (2002) containing a terminal cysteine for coupling to KLH.

The expression "Aβ8-x branched peptide" refers to a Aβ8-x peptides linked with a peptide spacer containing a terminal cysteine for coupling to KLH.
It was not obvious for the person skilled in the art to prepare the above defined antibodies because following a conventional process, i.e. immunization with five peptides (Aβ1-8, Aβ5-13, Aβ6-14, APs-15, and Aβp9-17), no specific antibody secreting hybridomas could be isolated and therefore it was necessary to immunize with a Aβ8-x peptide and a T-helper epitope, in particular a Aβp8-x peptide fused with a T-helper epitope, or to immunize with a Aβp8-x branched peptide.

In a preferred embodiment, the present invention relates to the process of preparation of an antibody above defined, wherein said antibody binds specifically to the N-terminal region of Aβp8-15 peptide, does not recognise APi_42 and which presents a high affinity with respect to Aβp8-15 peptide, such as determined on Western Blot.

A "Western blot" is a method to detect a specific protein in a given sample of tissue homogenate or extract.

According to another aspect, the present invention relates to an antibody which binds specifically to the N-terminal region of Aβp8-x peptide, such as obtained by a process defined above.

According to another aspect, the present invention relates to a method for determining in vitro amyloid burden in a mammal, comprising the following steps:

(i) quantifying the level of N-terminal truncated Aβp8-x in a body fluid of said mammal, using the antibody as defined above,

(ii) comparing the level of antibody of said mammal to those obtained with a control mammal, and

(iii) deducing from step (ii) if said mammal is suffering from a neurological disease provided the N-terminal truncated Aβp8-x level is modified with respect to the level measured in the control mammal, in particular is higher than the level measured in the control mammal.

The mammal examined in the present invention may be a non-human mammal, such as (but not limited to) a cow, a pig, a sheep, a goat, a horse, a monkey, a rabbit, a hare, a dog, a cat, a mouse, a rat, an elk, a deer, or a tiger. In a preferred embodiment, the mammal is a primate.

In a preferred embodiment, the mammal of the above defined method is a human, more preferably the mammal is a human adult.
In another preferred embodiment, the present invention relates to the above defined method wherein specificity and sensitivity of said antibody toward Aβ_{42} is higher than 60%, preferably comprised from about 60 to about 100%, more preferably comprised above 80%.

The term "sensitivity" refers to the degree of detection of Aβ_{42} peptide that the method can detect. (See Neurobiology of aging, Vol 19, N°.2, pO9-16, 1998: Consensus report of the working group on: "Molecular and biochemical markers of AD"). This working group sets standards for diagnostic kit in AD and mentions that sensitivity and specificity should be > 80%.

In another preferred embodiment, said body fluid of the above defined method is cerebrospinal fluid (CSF) or blood.

The term "cerebrospinal fluid" or "CSF" is intended to include whole cerebrospinal fluid or derivatives of fractions thereof well known to those skilled in the art. Thus, a cerebrospinal fluid sample can include various fractionated forms of cerebrospinal fluid or can include various diluents added to facilitate storage or processing in a particular assay. Such diluents are well known to those skilled in the art and include various buffers, preservatives and the like.

According to another aspect, the present invention relates to a method for determining, in a mammal, the susceptibility to a disease associated with β-amyloid formation and/or aggregation such as Alzheimer's disease, for determining, in a mammal, the risk of developing a disease associated with β-amyloid formation and/or aggregation such as Alzheimer's disease, for screening of the clearance of β-amyloid deposition in a mammal, or for predicting the level of β-amyloid burden in a mammal, said method comprising the following steps:

(i) determining, in said mammal, the amount of peptide Aβ_{42} with an antibody defined above,

(ii) comparing the amount determined in step (i) with the amount of antibody specific of said N-terminal region of Aβ_{42} peptide in a control mammal, and

(iii) concluding from the comparison in step (ii), whether the mammal is susceptible to a disease associated with β-amyloid formation and/or aggregation such as Alzheimer's disease, whether the mammal is at risk of developing a disease associated with β-amyloid formation and/or aggregation such as Alzheimer's disease, whether the β-amyloid deposition in a mammal is cleared, or what the level of β-
amyloid is in said mammal.

An increase in the level of N-terminal truncated Aβ₈₋ₓ in the brain of the tested mammal, for example, could be an indication of the mammal being susceptible to or at risk of developing a disease associated with β-amyloid formation and/or aggregation. It could also indicate that the Aβ deposition in the mammal is not yet cleared.

Increased levels of N-terminal truncated Aβ₈₋ₓ in certain body fluids after vaccination or therapy, are an indication of the level of Aβ burden (DeMattos et al, 2002). N-terminal APP soluble fragment will mainly be found in certain body fluids. The presence of these N-terminal APP soluble fragments indicates an aberrant cleavage of APP, resulting in the formation of N-terminal truncated Aβ variants and, consequently, in an increased susceptibility to or risk of developing a disease associated with β-amyloid formation and/or aggregation by the mammal.

In a preferred embodiment, the amount of antibody specific of the N-terminal region of Aβ₈₋ₓ peptide using the above defined method is determined on a tissue sample obtained from said mammal.

By "tissue" is meant brain tissue.

According to another aspect, the present invention relates to a kit comprising at least one buffer, and at least one detection compound, at least one N-truncated Aβ₈₋ₓ specific antibody as defined above.

In a preferred embodiment, the kit defined above, further comprises a preferably labelled second antibody which binds to the above defined antibody.

For example, the antibodies can be bound directly to a solid substrate where they are immobilized. These immobilized antibodies then capture the N-terminal truncated Aβ₈₋ₓ peptide of the invention present in the sample, which are subsequently detected with a second antibody.

In another aspect, the present invention relates to a therapeutic composition comprising as active ingredient an above defined antibody, or synthetic peptides with a free N-terminal-end mimicking the free N-terminal-end of N-truncated Aβ peptides, in association with a pharmaceutically acceptable vehicle.

The amount of antibody to be administered or delivered to an individual should be sufficient to cause a significant reduction in βamyloid levels in the brain of the individual. The appropriate amount will depend upon various parameters (e.g. the particular antibody
used, the weight of the individual and the levels of endogenous β amyloid) and is to be determined on the case by case basis.
The dosage and frequency of administration can also vary depending on whether the treatment is prophylactic or therapeutic.

In a preferred embodiment, the above defined therapeutic composition is suitable for the administration to an individual of a dose of an antibody from 1 mg/kg/day to 200 mg.

Patients amenable to treatment include individuals at risk of disease but not showing symptoms, as well as patients presently showing symptoms. In the case of Alzheimer's disease, virtually anyone is at risk of suffering from Alzheimer's disease if he or she lives long enough. Therefore, the present antibodies can be administered prophylactically to the general population without any assessment of the risk of the subject patient. The present antibodies are especially useful for individuals who do have a known genetic risk of Alzheimer's disease. Such individuals include those having relatives who have experienced this disease and those whose risk is determined by analysis of genetic or biochemical markers.

Administration of the antibodies according to the present invention to an individual can be made via intravenous administration.

Another way of delivering to the brain is via direct infusion of the antibodies according to the present invention into the brain of the individual.

According to another aspect, the invention relates to a vaccine composition comprising as active ingredient an above defined antibody, fragments or derivatives thereof, or synthetic peptides with a free N-terminal-end mimicking the free N-terminal-end of N-truncated Aβ peptides, in association with a pharmaceutically acceptable vehicle.

In a preferred embodiment, the above defined vaccine composition is suitable for the administration to an individual of a dose of an antibody from 1 mg/kg/day to 200 mg/kg/day.

The vaccine or therapeutic compositions of the present invention induce an immune response against the specific N-terminal truncated Aβ_{8-16} peptide of the invention.

According to another aspect, the present invention relates to the use of at least one of the above defined antibodies, for the preparation of a drug or a vaccine intended for the prevention or the treatment of Alzheimer disease.

As used herein, the term "preventing a disease" means inhibiting or reversing the onset of the disease, inhibiting or reversing the initial signs of the disease (i.e. formation
and/or aggregation of Aβ variants), inhibiting the appearance of clinical symptoms of the disease.

As used herein, the term "treating a disease" includes substantially inhibiting the disease, substantially slowing or reversing the progression of the disease, substantially ameliorating clinical symptoms of the disease or substantially preventing the appearance of clinical symptoms of the disease.

According to another aspect, the present invention relates to the use of at least one of the above defined antibodies, for the preparation of a drug or a vaccine intended for the clearance of β amyloid burden.

The term "clearance of β amyloid burden" means that the β amyloid burden is eliminated from the brain tissue. Clearance of amyloid deposits in the brain of AD patients using vaccination against Aβ peptide is a novel approach that opens treatment perspectives (Schenk et al., 2001, Immunotherapy with beta-amyloid for Alzheimer's disease: a new frontier. DNA Cell Biol. 20: 679-681).

According to another aspect, the present invention relates to a method of clearance of β-amyloid burden in a mammal comprising the administration of an above defined composition to the said mammal.

According to another aspect, the present invention relates to the use of peptide composition defined above for induction of an immune response in a mammal being affected by or susceptible to develop an Alzheimer disease.

DESCRIPTION OF THE FIGURES

Figure 1 represents the partial amino acid sequence of APP770, displaying the amino acid sequence of Aβ with the cc-, β-, and γ-secretase cleavage sites indicated.

Figure 2A and 2B represent the schedule of intraperitoneal injections of the peptide preparation (2A) and the antibody titer measured for each bleeding (2B).

x-axis: serum dilution
y-axis: optical density

Figure 3A to 3D represent the use of immunized mice sera to detect or not amyloid deposits in brain tissue of double transgenic APPxPS1 mice:
3A and 3B: Not responding mouse serum (magnification: x25 and x100 respectively).

3C and 3D: Trunc8 immunized mouse serum (magnification: x25 and x100 respectively).

Figure 4A and 4B represent the Aβ load measured by western-blotting in immunized and control mice (4A) and the histogram represents the efficiency of immunization expressed the percentage of Aβ-42 load to the control condition (PBS).

Figure 5 the schematic overview of primer location for the light chain and the heavy chain.

Figure 6 represents a 2D gel analysis of a formic acid extract of a human Alzheimer brain and a mixture of "full-size" synthetic Aβ peptides (Aβ2-42, Aβ3-42, Aβ4-42, Aβ5-42, Aβ6-42, Aβ7-42, Aβ8-42, Aβ9-42) immunoblots obtained with 7Gl 2 equivalent to 21F12 (Aβ1-42) as described by Sergeant et al. (2003), TeiA 1.1, TeiA 1.8, and TeiA 2b.6).

Figure 7 represents immuno-capture antibody of 4D7A3 (a 42-C-terminal specific antibody) and TeiA 2b.6, TeiA1.8, TeiA1.7 and TeiA1.6.

Figures 8A and 8B represent the specificity of the monoclonal antibody TeiA1.6 for parenchymal amyloid. 8A: labelling of both parenchymal (arrows) and vascular amyloid deposits (arrowheads) with a classical Aβ antibody 6E10. 8B: labelling of the only parenchymal (arrows) amyloid deposits with a 8-truncated Aβ (TeiAl.6) and not the perivascular deposits (arrowheads) in an adjacent brain section.

Figures 9A to 9J represent results from the intracranial injection (right hippocampus) of 4G8 antibody (commercial monoclonal antibody) to mice n°47, 7 month old. 9A, 9D and 9G: position of brain sections with regard to the injection point.

9B, 9E and 9H: immunohistochemistry images of the corresponding brain sections showing amyloid peptide deposits detected with the "revealing" antibody, 6E10. 9C, 9F and 9I: amyloid peptide load calculated from images 9B, 9E and 9H, respectively, in
different brain subregions in each hemisphere (Hipp: Hippocampus, CxI: cortical region 1(dorsal), Cx2: cortical region 2 (lateral), Cx3: cortical region 3(latero-ventral), Th: thalamic). Ratio: stained area / total area of that region.
L: left, R: right (injected).

9J: amyloid peptide load calculated in subiculum in each hemisphere by immunohistochemistry in the brain section H only.

Figures 10A to 10J represent the intracranial injection (right hippocampus) of TeiAl.6 antibody to mice nº 17, 7 month old.

10B, 10E and 10H: immunohistochemistry images of the corresponding brain sections showing amyloid peptide deposits detected with the "revealing" antibody, 4G8.

10C, 10F and 10I: amyloid peptide load calculated from images 10B, 10E and 10H, respectively, in different brain subregions in each hemisphere (Hipp: Hippocampus, CxI: cortical region 1(dorsal), Cx2: cortical region 2 (lateral), Cx3: cortical region 3(latero-ventral), Th: thalamic). Ratio: stained area / total area of that region.

L: left, R: right

10J: amyloid peptide load calculated in subiculum in each hemisphere by immunohistochemistry in the brain section B and E only.

Figure 11 represents the ratio of amyloid load between the injected (TeiAl.6 antibody) and non injected (control) in the different brain subregions (Hipp: Hippocampus, CxI: cortical region 1(dorsal), Cx2: cortical region 2 (lateral), Cx3: cortical region 3(latero-ventral), Th: thalamic) after intracranial injection (right hippocampus) of TeiAl.6 monoclonal antibody. This is the average of ratios for 4 animals with three brain sections quantified for each (data represents means +/- SEM).

Figures 12A to 12J represent the intracranial injection (right hippocampus) of TeiAl.8 antibody to mice nº 58, 7 month old.

12B, 12E and 12H: immunohistochemistry images of the corresponding brain sections showing amyloid peptide deposits detected with the "revealing" antibody, 4G8.
12C, 12F and 12I: Amyloid peptide load calculated from images 12B, 12E and 12H, respectively, in different brain subregions in each hemisphere (Hipp: Hippocampus, Cx1: cortical region 1(dorsal), Cx2: cortical region 2 (lateral), Cx3: cortical region 3(lateral-ventral), Th: thalamic). Ratio: stained area / total area of that region.

L: left, R: right

12J: Amyloid peptide load calculated in subiculum in each hemisphere by immuunohistochemistry in the brain section H only.

Figure 13 represents the ratio of amyloid load between the injected (TeiAl.8 antibody) and non injected (control) in the different brain subregions (Hipp: Hippocampus, Cx1: cortical region 1(dorsal), Cx2: cortical region 2 (lateral), Cx3: cortical region 3(lateral-ventral), Th: thalamic) after intracranial injection (right hippocampus) of TeiAl.6 monoclonal antibody. This is the average of ratios for 4 animals with three brain sections quantified for each (data represents means +/- SEM).

EXAMPLES

Example 1: Immunization of double transgenic mice with N-trunc 8 peptide preparation and consequences on the brain amyloid load

Double APP Swedish London x Presenilin 1 transgenic mice (Blanchard et al., 2003 Exp Neurology 184 :247; woo 1209777) were injected every three weeks with 50 μg of N-Trunc 8 peptides (Figure 2A). The whole duration of immunization was of 21 weeks. As negative and positive controls, series of mice were injected with phosphate buffer saline or aggregated Aβ1-42 peptide, respectively. The antibody titer was determined by direct ELISA against the Trunc 8 peptides (Figure 2B).

Sera from fifth bleeding of immunized mice was used to perform tissue amyloid plaque immunoreactivity (TAPIR) (Christoph Hock, Roger M. Nitsch, Clinical Observations with AN-1792 Using TAPIR Analyses Neurodegenerative Diseases 2005;2:273-276) (Figure 3). The serum from a non-responding mouse was used as a negative control. Amyloid deposits were detected with the sera obtained from mice immunized with Trunc-8 peptides.
Consequences of immunization on the Aβ load was examined using formic acid extracts of Aβ peptides and detection by Western-blotting as previously described (Casas et al., 2004) (Figure 4A). The amount total Aβ-42 was measured and compared to control condition (PBS) and expressed as the percentage to the control condition (100%). The histogram represents the quantifications for each experimental condition (Figure 4B).

Example 2: Characterization of monoclonal antibodies variable regions from hybridoma's IGH524, IGH525, IGH521, IGH522, IGH523

The results of DNA sequence analysis was evaluated by translation of the appropriate open reading frame to amino acid sequence and alignment with consensus antibody heavy and light chain framework regions.

Data analysis

Raw sequencing data (DNA chromatograms) are generated with Sequencing Analysis Software v5.2 (Applied Biosystems) and the KB basecaller v1.2 (Applied Biosystem) and interpreted and edited using Sequencher 4.1.2. In general, double-stranded sequencing results were assembled and the consensus sequence was linked to the Innogenetics Lotus Notes Custom Sequencing Service Request (CSSR) database and stored with the assigned CSSR project number.

Results

RNA isolation, RT-PCR, cloning and deposit.

Table 1 shows for each hybridoma/MAb the origin and source of the cells used for RNA extraction, and shows for each heavy or light antibody chain the corresponding primer combination which successfully resulted in a specific clonable PCR fragment.

Sequence analysis

For each variable region, DNA sequence analysis and subsequent alignment revealed a possible consensus for each hybridoma/MAb. Complementary-determining regions (CDR) were identical for all clones specifying one variable region.

An overview and alignment of the obtained final consensus sequences is given in appendix 1. Theoretically predicted CDR loops are indicated (based on consensus sequence rules).

• The complementarity-determining regions (CDR) as marked in the consensus sequences were assigned based on a set of public available rules from the Kabat...
definition (Reczko et al., 1995) or a public available analysis tool for modelling
(Honegger et al., 2001). The CDRs are marked for explorative/informal use only.

IGH524. TeiA 2b.6

The results obtained for the heavy and light chain of MAb TeiA 2b.6 (2.13E5E4) isolated from hybridoma IGH524, were clear with only minor ambiguities and/or differences located mainly in framework regions. The complete variable regions have been determined and the N-terminal end (including the largest part of CDR1) of both mature antibody chains were confirmed by N-terminal amino acid sequencing of the purified antibody.

IGH521 (TeiA 1.6UGH522 (TeiA 1.7), IGH523 (TeiA 1.8), IGH525 (TeiA 1.1)

The results for all heavy and light chains of MAb TeiA 1.6 (2.6F4C2, IGH521), TeiA 1.7 (2.8A3F8, IGH522), TeiA 1.8 (1.3B12H3, IGH523) and TeiA 1.1 (3.46B10E7, IGH525) were also clear. Eight sequences of cloned PCR products were aligned and in least three identical sequences lead to the consensus sequence. The complete variable regions have been determined by alignment with the sequence obtained from hybridoma IGH524.

Table 1 PCR primers

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<td>CCGTTTCAGCTCCAGGCTTGGTCCC</td>
<td>Orlandi et al</td>
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Kabat et al. (Sequences of proteins of immunological interest. National Institutes of Health Publication No. 91-3242, 5th ed., 1991, United States Department of Health and Human
Services, Bethesda, MD)

Coloma et al. (Novel vectors for the expression of antibody molecules regions generated by polymerase chain reaction. J. Immunol Methods, 1992; 152(1):89-104)

Orlandi et al. (Cloning immunoglobulin variable domains for expression by the polymerase chain reaction. Proc Natl Acad Sci U S A. 1989 May;86(10):3833-7)

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<tr>
<th>IGH</th>
<th>Name</th>
<th>Ab Chain</th>
<th>Primer pair</th>
<th>ICCG</th>
</tr>
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<tbody>
<tr>
<td>524</td>
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<td>Light</td>
<td>1009565/18696, 18700/1010500</td>
<td>6152</td>
</tr>
<tr>
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<td>TeiA 1.6 (2.6F4C2)</td>
<td>Light</td>
<td>18698/18696, 18700/19735</td>
<td>6233</td>
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<tr>
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<tr>
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<td>Light</td>
<td>18698/18694, 18700/1010500</td>
<td>6235</td>
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<tr>
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<td>6268</td>
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</tbody>
</table>

**appendix 1:**

**IGH524 sequence**

**Light chain variable region:**

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<tr>
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</thead>
<tbody>
<tr>
<td>SSLTVTAGEKVTMSCKSSQSLFNSORQTMYLTWFQQRPGQ APKLLIYWASTEGSGVP</td>
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</table>

<table>
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<tr>
<th>CDR-L3</th>
</tr>
</thead>
<tbody>
<tr>
<td>DRFTGSOSGTEFTLTISSVQAEDLAVYYCQNDYTPTTFAG (SEQ ID NO: 7)</td>
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</tbody>
</table>

**Heavy chain variable region:**

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<thead>
<tr>
<th>CDR-H1</th>
<th>CDR-H2</th>
</tr>
</thead>
<tbody>
<tr>
<td>GGLVQPGGSLRLSCATSGFTFTDFYMEWVRQPAGKRLWIAASRNKANGYTTEYSA</td>
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</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>CDR-H3</th>
</tr>
</thead>
<tbody>
<tr>
<td>SVKGRFIVSRDTSQGILYLMQMSALRAEDTAIYYCAIYRYYAMDYWQGTSVTVSS (SEQ ID NO: 8)</td>
</tr>
</tbody>
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IGH521 sequence
Light chain variable region:

CDR-L1

CDR-L2

SSLTVTAGEKVTMSCKSSQSSLAAORYQEMYLTWYQQKPGQPPKLLIYWASXRøSGV

CDR-L3

PDRFTGSGSTDFLTISSVQAEDLAVYYCQNDYTYPPLTFAG (SEQ ID NO : 1)

Heavy chain variable region:

CDR-H1

CDR-H2

GGLVQPGGSRLSCAISGFTSFDFYMIVWRQPPGKRLEWIAASNKANDYTEYSAS

CDR-H3

VKGRFIIVSRDTSQSILYLQMNAEDTAYYCATYHDYAMDYWGQGTSVTSS

(SEQ ID NO : 2)

IGH525 sequence
Light chain variable region:

CDR-L1

CDR-L2

SSLTVTAGEKVTMSCf SSQSLFNSTQTMYiLTWYQQKPGQPPKLLIYWASmESOVP

CDR-L3

DRFTGSGSTDFLTISSVQAEDLAVYYCQMDYTYPPLTFAG(SEQ ID NO:9)

Heavy chain variable region:

CDR-H1

CDR-H2

GGLVQPGGSRLSCATSFGFTSFDFIEWVRQPPGKRLEWITASRNKNYDYKTEYSAS

CDR-H3

VKGRFIIVSRDTSQSILYLQMNAEDTAYYCAIYRBYAMDYWGQGTSVTSS

(SEQ ID NO : 10 )

IGH522 sequence
Light chain variable region:
CDR-L1
SSLTVTAGEKVTMNCKSSQNLLLNSGQNYLTWFQQKPQPKLLIYWASTRESGV

CDR-L2

CDR-L3
PDRFIGSGGTDFTLTINSVQAEDLAVYYCQNDYRYPLTFGAG (SEQ ID NO:3)

5

Heavy chain variable region:

CDR-H1
GGLVQPGGSLRLSCATAGFTFYFSDYMEWVRQPPRLEWIAASiIDKAiCDYTTIYSA

CDR-H2

CDR-H3
SmORFIVSRDTSQSYFLQMNALRSEDTAIYYCATYFSYAMDYWGLGTSVTVSS (SEQ ID NO: 4)

IGH523 sequence

Light chain variable region:

CDR-L1
SSLAVTAGERVTMSCKSSLNLNSGQTYLTWYQQKPQPKLLIYWASTRESGV

CDR-L2

CDR-L3
DRFTGSGGTDFTLTISSVQAEDLAVYYCQNDYSYPLTFGAG (SEQ ID NO:5)

10

Heavy chain variable region:

CDR-H1
GGLVQPGGSLRLSCATAGFTFYFSDYMEWVRQPPRLEWIAASiIDKAiCDYTTIYSA

CDR-H2

CDR-H3
SVKGRFTISRDNSQILQMSTLRAGDSATYCAVYGNAMYGDYWQGTSVNVS (SEQ ID NO: 6)

Example 3: N-truncated 8-specific Aβ antibodies and (limited) characterization during cloning

Fifteen Balb-C mice were injected with a mixture of 5 short synthetic Aβ peptides (50 µg per mouse KLH-coupled peptides). One mouse died for an unknown reason. The peptides
correspond to the first eight N-terminal residues of respectively $\text{A} \beta_{i,s}$, $\text{A} \beta_{5,13}$, $\text{A} \beta_{6,14}$, $\text{A} \beta_{s,i5}$, and $\text{A} \beta_{g,i7}$ (see Table 2). The peptides also contained a C-terminal residue for coupling to KLH. After 5 injections titration of the sera was done in a 'coatings assay' of a mixture of peptides. Peptides were coated as a streptavidine-biotinylated peptide complex (peptides (IGP-2258, see Table 2) or as a BSA (bovine Serum Albumin)-peptide complex (PG-Nr see Table 2) and an anti-mouse antibody coupled to HRP (Jackson goat anti-mouse HRP, Cat No 115-035-071) was used for detection. Although titers were low (not shown) a first mouse was sacrificed and a fusion was performed. No specific antibody secreting hybridoma's has been isolated.

Therefore sets of mice were boosted with 'modified peptides'. Three mice were injected with the original peptide mixture, two mice were further injected with IGP-21 19 KLH-coupled peptide (see Table 2).

Peptides corresponding to $\text{A} \beta_{s,i5}$ were the more immunogenic out of the mixture of five, three additional peptides were thus synthesized. One corresponds to $\text{A} \beta_{i,s}$, fused to a T-helper epitope (PGPGP (Livingston et al, 2002); IGP-2406 (Table 2) and a C-terminal cysteine residue for coupling to KLH. The other peptide also contained another T-helper epitope (DGDGD (McMillan et al., 1983); IGP-2258 (Table 2). Finally a branched peptide containing a C-terminal cysteine for coupling was also synthesized (IGP-2407 (Table X).

Each time two mice were immunized with the newly synthesized peptides. The $\text{A} \beta_{s,i5}$ peptide was also coupled to E1 particles (WO 2004/013172) and used for boosting in the last two mice. Titers were again monitored with a 'coatings-assay' (results not shown). Titers to $\text{A} \beta_{s,i5}$ were indeed improved in the mice boosted with the T-helper peptides and branched peptide and it was decided to use all three surviving mice for fusion. One of the mice boosted with the branched peptide died.

Table 2: Sequence of the peptides used and their Innogenetics reference number.

<table>
<thead>
<tr>
<th>Name</th>
<th>Innogenetics ref number</th>
<th>Sequence</th>
</tr>
</thead>
<tbody>
<tr>
<td>$\text{A} \beta_{i,8}$</td>
<td>IGP-2062</td>
<td>DiAEFRHDS$ _{8}$GC</td>
</tr>
<tr>
<td>$\text{A} \beta_{s,i2}$</td>
<td>IGP-2121</td>
<td>R$_s$HDSGYEVi$ _{2}$GC</td>
</tr>
</tbody>
</table>
Spleen of both mice were prepared and fused to SP2/0 cells. After plating, 66 plates (±3000 clones) were screened. During subcloning a limited number of clones, 24, were characterized using the biotinylated peptides IGP-2258 and IGP-2259 in a bridging assay and a Luminex assay. In the bridging assay, BSA coupled peptide IGP-128, PG127 (see Table 3) was used to capture one binding site of the antibody and a biotinylated peptide was used to detect the captured antibody in a so-called bridging assay. This assay gives an indication on the affinity of the antibody: a high-affinity antibody will give a higher signal than a lower affinity antibody. Indeed 'two classes' of antibodies have been identified.

To determine the specificity of the antibody a peptide was used which is shifted two amino-acids N-terminal as compared to the AB_{8-15}, the AB_{6-13} peptide. The biotinylated versions of these peptides are used in order to capture it efficiently to avidin Luminex beads. After washing, the antibodies were revealed by an anti-mouse phyco-erythrine antibody. The results presented in Table 3 are raw data expressed as Mean Fluorescence Intensity (MFI). A value below 10 means below background, so for all antibodies tested with 'low affinity' (bridging assay OD<1), no reaction on the non-specific peptide (IGP-2259) was observed.

For the 'high-affinity antibody' a small signal was measurable on the non-specific peptide, but with a small difference between the antibodies. From the 'high-affinity' class of antibodies, three antibodies were chosen for subcloning, one IgG2b subtype and two IgG1, while from the 'low affinity' antibodies two IgG1 antibodies were selected, resulting in five antibodies for full characterization.

<table>
<thead>
<tr>
<th>Peptide</th>
<th>IGPs</th>
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<tbody>
<tr>
<td>AB_{6-13}</td>
<td>IGP-2120</td>
</tr>
<tr>
<td>AB_{8-15}</td>
<td>IGP-2119</td>
</tr>
<tr>
<td>AB_{9-16}</td>
<td>IGP-2122</td>
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<tr>
<td>AB_{8-15} DG</td>
<td>IGP-2405</td>
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<tr>
<td>AB_{8-15} PG</td>
<td>IGP-2406</td>
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<tr>
<td>AB_{8-is} branched</td>
<td>IGP-2407</td>
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<tr>
<td>AB_{8-is-bio}</td>
<td>IGP-2258</td>
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<tr>
<td>AB_{6-13-bio}</td>
<td>IGP-2259</td>
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</tbody>
</table>

Table 3: Characterization of the N-truncated 8-specific AB, TeiA (Truncated eight amyloid), antibodies during cloning. The isotype was determined, reactivity in bridging assay (high OD
is indicative for a high affinity) and specificity in a Luminex format on bio-peptides captured on avidin beads. The final subclone that has been used for further characterization is also indicated.

<table>
<thead>
<tr>
<th>Clone</th>
<th>Ig subtype</th>
<th>Bridging assay (OD450)</th>
<th>Luminex assay (MFI)</th>
<th>Subclone</th>
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<tr>
<td></td>
<td></td>
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<td>IGP-2258</td>
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<tr>
<td>2.13.E5</td>
<td>IgG2b</td>
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<td>IgG1</td>
<td>0.1478</td>
<td>2451.5</td>
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<td></td>
</tr>
</tbody>
</table>

**Example 4: Characterization of N-Truncated eight specific (TeiA) antibodies**

In order to further substantiate the specificity of these TeiA antibodies on Aβ, two approaches were taken: (1) 2D gelanalysis of a formic acid extract of a human Alzheimer brain and (2) a mixture of 'full-size' synthetic Aβ peptides (Anaspec) differing in their N-terminus used on a SELDI approach (Merchant et al., 2000).

The results of these approaches are shown in figures 6&7. Brain tissue sampling and 2D analysis have been performed essentially as described in Sergeant et al (2003).

To reveal the position of the Aβ42 peptides, a new 42-C-terminal specific antibody was used 7G12H1 (equivalent to 21F12 as described in Sergeant et al (2003)). The different spots have been characterized with mass-spectrometry to correspond to the different N-truncations as presented on the Figure 6.
Example 5: mAb TeiAl.6 (Aβ N-trunc8) is specific for parenchymal amyloid deposits and does not recognize vascular amyloid deposits

One of the side-effects of the passive immunization is the frequency of microhemorrhages. Such increase in the number of microhemorrhages may be explained by the fixation of injected antibodies to the aggregated AB peptides within vessel walls (Paris et al., 2000; Pfeifer et al., Cerebral Hemorrhage After Passive Anti-AB Immunotherapy, Science 15 November 2002; Vol. 298. no. 5597, p. 1379). Thus, truncated AB species are also original targets since they are not mainly found in amyloid angiopathy. As demonstrated in figure 8A and 8B, on adjacent human AD brain sections, a classical AB antibody labels both parenchymal and vascular amyloid deposits (A, arrows and arrowheads, respectively, 6E10 antibody).

Using a truncated 8 antibody (B, here TeiA1.6), only parenchymal amyloid deposits are labelled (B, arrows) but not vascular amyloid deposits (B, arrowheads).

Altogether, these data indicate that amino 8-truncated AB antibodies specifically target parenchymal amyloid deposits and not interact with vascular amyloid deposits which have been suggested to be responsible for the perivascula effects (hemorrhages, encephalopathies) observed with other anti-Abeta immune approaches.

Example 6: Intracranial administration of N-Truncated eight specific (TeiA) antibodies in transgenic mice lead to a decrease in amyloid plaque burden

In order to demonstrate therapeutic efficacy of TeiA antibodies, they were injected in the hippocampus of transgenic mice bearing amyloid plaques in the brain and 7day after administration, cerebral amyloid peptide plaque load was quantified by immunohistochemistry. Briefly, under stereotaxic conditions, 1 or 2 μg of antibody were injected in the right hippocampus (unilateral injection) in ThyAPP S-LxPSiMi46L mice (Blanchard et al., 2003 Exp Neurology 184:247; WO0120977). The antibodies injected were: two commercial classical AB antibodies (4G8 and 6E10) and the TeiA antibodies TeiA 1.1, 1.6, 1.8 and 2b6.

Seven days after injection, animals were euthanized and brains treated for immunohistochemistry. After brain postfixation, 40 μm coronal cryosections were performed and sections 400 μm apart were stained with biotinylated 4G8 anti-Abeta as a "revealing" antibody to evaluate the amyloid load present in the brain. In the case where
4G8 antibody had been injected in the brain, the revealing antibody used was biotinylated 6E10 to avoid masking of epitopes. Biotinylated antibodies were detected with a standard avidin-peroxidase detection kit (Vectastain® ABC kit Vector Laboratories).

In each brain section, the amyloid peptide load was calculated in each hemisphere (injected and non-injected) in five different brain subregions [hippocampus, cortical region 1 (dorsal), cortical region 2 (lateral), cortical region 3 (latero-ventral) and thalamic]. After acquisition of images on an Olympus scanner system, quantification was performed semi-automatically with the Mercator ExploraNova system. For each animal, three brain sections were quantified, positioned quite similarly with regard to the injection point: one next to the injection point, one rostral and one caudal to the injection point. As previously described (Wilcock et al, 2003, J Neurosci 23:3745; Oddo et al, 2004, Neuron 43:321), 4G8 injection lead to a significant decrease in the amyloid peptide deposits in the injected hemisphere when compared to the non-injected hemisphere (Figure 9). This effect was variable between brain sections as might be expected from this local injection of the antibody. TeiAl.6 antibody also led to a significant decrease in brain amyloid in the injected side that was more pronounced in this series of experiments in the cortical region 3 (Figure 10). Analysis of 4 mice (aged 7 months) indicated a significant decrease (Figure 11).

Similarly TeiA1.8 led to a significant decrease in brain amyloid in the injected side that was more pronounced in this series of experiments in the cortical region 2 (Figure 12). Analysis of 4 mice (age 7 months) indicated a significant decrease (Figure 13).

These data indicate that TeiA antibodies 1.6 and 1.8 decreased brain amyloid load even after short term administration and compare well with classical anti-AB antibodies. It is of interest to note that animals already presented a significant deposition of amyloid at the time of administration, therefore suggesting a therapeutic rather than solely preventive potential for TeiA antibodies.

TeiA antibodies could therefore provide a good therapeutic effect against amyloid load in Alzheimer Disease patients.
CLAIMS

1. An antibody which specifically binds to the N-terminal region of $\text{A} \beta_{9-x}$ peptide, $x$ being comprised from 11 to 42, and recognises neither $\text{A} \beta_{5-40}$ nor $\text{A} \beta_{5-42}$.

2. Antibody according to claim 1, wherein said antibody presents a high specificity for the free N-terminal end of $\text{A} \beta_{5-X}$ peptide.

3. Antibody according to claim 1 or 2, wherein said antibody presents a high affinity with respect to $\text{A} \beta_{5-X}$ peptide.

4. Antibody according to any of claims 1 to 3, wherein said antibody specifically target parenchymal amyloid deposits of $\text{A} \beta_{8-x}$ peptide in the brain and does not interact with vascular amyloid deposits.

5. Antibody according to any of claims 1 to 4, wherein $x$ is comprised from 15 to 42, in particular a monoclonal antibody.

6. Antibody according to any of claims 1 to 5, wherein the variable region comprises one of the following pairs of amino acid sequences, corresponding respectively to the light and heavy chain:

- Antibody TeiA 1.6 (secreted by hybridoma IGH521)

**Light chain variable region:**

CDR-L1

$$\text{SSLTVTAGEKVTSCKSSQSLAGEYQKNYLTYWQYQQKPGQPPKLIIYWAST}$$

CDR-L2

$$\text{RDSGPDRFTGSGTDLITISVQAEDLAVYYCNDTYPYLTFA}$$

(SEQ ID NO : 1)

**Heavy chain variable region:**

CDR-H1

$$\text{GGLVQPGSSLRLSCAISOFTFSDFYMEDWVRQPPKRLIEWIASENKAKDYTf}$$
CDR-H3
BYSASVKGRFIVSRDTSQSILYLQMNLRAEDTAIYYCATYHDYAMDYWGQ

GTSVTSS (SEQ ID NO: 2)

• Antibody TeiA 1.7 (secreted by hybridoma IGH522)

Light chain variable region:

CDR-L1  CDR-L2
SSLTVTAGEKVTMNCKSSQNLNSGNQVNYLTWFQQKPGQQPKLLIYWAST

CDR-L3
RESGVPRFIGSGSGTDTLTINSVQAEDLAVYYCQNDYRYPLTFGAG
(SEQ ID NO:3)

Heavy chain variable region:

CDR-H1  CDR-H2
GGLVQPGGSLRLSCATSGFTFSDFYMEWVRQPPGRREWIAASRDICAIYBT

CDR-H3
EYSASVKGRFIVSRDTSQSILYLQMNLRAEDTAIYYCATYHDYAMDYWGQ

GTSVTSS (SEQ ID NO: 4)

• Antibody TeiA 1.8 (secreted by hybridoma IGH523)

Light chain variable region:

CDR-L1  CDR-L2
SSLAVTAGERVTMSCKSSLTLNSGSQNYLTWYQQKPGQQPKLLIYWAST

CDR-L3
ISGVPRFTGSGTGDTLTISSVQAEDLAVYYCQKDYSYPLTFGAG
(SEQ ID NO:5)

Heavy chain variable region:

CDR-H1  CDR-H2
GGLVQPGGSLRLSCATAGFTFTQYMSWVRQPPGKALEWLATIRNKAKGFT
CDR-H3
TEYSASVKGRFTISRDNQSILYLQMSTLRAGDSATYYCAVYGNAYMDYW
QGTSVNVSS (SEQ ID NO : 6)

• Antibody TeiA 2b.6 (secreted by hybridoma IGH524)

Light chain variable region:
CDR-L1 CDR-L2
SSLTVTAGEKVTMSCKSSQSLFNSGRQTNLYLTWFQQRPGQAPKLIIYWASTR
CDR-L3
GSGVPDRFTGSGTEFTLTISSVQAEDLAVYCYCNQDYTYPLTFGAG
(SEQ ID NO : 7)

Heavy chain variable region:
CDR-H1 CDR-H2
GGLVQPGGSLRLSCATSGFTFTDFYMEWVRQPPGKRWIAASRNKAKGYT
CDR-H3
T1YSASYKGRFIVSRDTSQGILYLQMSALRAEDTAIYYCAIYRYYAMDYWGQ
GTSVTVSS (SEQ ID NO : 8)

• Antibody TeiA 1.1 (secreted by hybridoma IGH525)

Light chain variable region:
CDR-L1 CDR-L2
SSLTVTAGEKVTMSCTSSQLFKSGTQLiNYLTWYYQQKPGQPPKLLIIYWASTR
CDR-L3
ISGVPRFTGSGTDFTLTISSVQAEDLAVYCYCNQDYTYPLTFGAG
(SEQ ID NO:9)

Heavy chain variable region:
CDR-H1 CDR-H2
GGLVQPGGSLRLSCATSGFTFSDFEWEVRQPPGKRWIAASRNKAKGYT
CDR-H3

TEYSASYKGRFIVSRDTSQSILYLQMNALRAEDTAIYYCAIYRYAMOYWGQ

GTSVTVSS (SEQ ID NO: 10)

7. Antibody according to any of claims 1 to 6, wherein the CDR of the light and heavy chain of the variable region comprises one of the following amino acid sequences:
   • Antibody TeiA 1.6 (IGH521 sequence)
     CDR of the light chain variable region:
     CDR-L1:
     KSSQSLLAGRYQKNYLT (SEQ ID NO: 11)
     CDR-L2:
     WASTRDSG (SEQ ID NO: 12)
     CDR-L3:
     QNDTYTYFLT (SEQ ID NO: 13)

     CDR of the heavy chain variable region:
     CDR-H1:
     GPTFSDFYMI (SEQ ID NO: 14)
     CDR-H2:
     ASRNKANDYTEYSASVKG (SEQ ID NO: 15)
     CDR-H3
     YHDYAMDY (SEQ ID NO: 16)

   • Antibody TeiA 1.7 (IGH522 sequence)
     CDR of the light chain variable region:
     CDR-L1:
     KSSQNLNLNSGNQNYLT (SEQ ID NO: 17)
     CDR-L2:
     WASTRESG (SEQ ID NO: 18)
     CDR-L3:
     QKDYRYPLT (SEQ ID NO: 19)
CDR of the heavy chain variable region:

CDR-H1:
GFTFSDFYME (SEQ ID NO: 14)

CDR-H2:
ASRDKAKDYTTEYSASVKG (SEQ ID NO: 20)

CDR-H3:
YFSYAMDY (SEQ ID NO: 21)

- Antibody TeiA 1.8 (IGH523 sequence)

CDR of the light chain variable region:

CDR-L1:
KSSLTLLNSGSQTNYLTD (SEQ ID NO: 22)

CDR-L2:
WASTRESG (SEQ ID NO: 18)

CDR-L3:
QKDYSYPLT (SEQ ID NO: 23)

CDR of the heavy chain variable region:

CDR-H1:
GFTFTDQYMS (SEQ ID NO: 24)

CDR-H2:
TIRNKAKGFTTEYSASVKG (SEQ ID NO: 25)

CDR-H3:
YGNYAMDY (SEQ ID NO: 26)

- Antibody TeiA 2b.6 (IGH524 sequence)

CDR of the light chain variable region:

CDR-L1:
KSSQSLFNSGEQTYNLTD (SEQ ID NO: 27)

CDR-L2:
WASTRGS (SEQ ID NO: 28)
CDR-L3:
QNDYTYFLT (SEQ ID NO : 13)

CDR of the heavy chain variable region:
CDR-H1:
OFTFTDFYME (SEQ ID NO : 29)
CDR-H2:
ASRNKANGYTYTEYSASVKG (SEQ ID NO : 30)
CDR-H3:
YRYYAMDY (SEQ ID NO : 31)

• Antibody TeiA 1.1 (IGH525 sequence)

CDR of the light chain variable region:
CDR-L1:
TSSQSLFNSOTQTNYLT (SEQ ID NO : 32)
CDR-L2:
WASTRESG (SEQ ID NO :18)
CDR-L3:
QNDYTYFLT (SEQ ID NO : 13)

CDR of the heavy chain variable region:
CDR-H1:
OFTFSDFIE (SEQ ID NO : 33)
CDR-H2:
ASRNKNYDYKTEYSASYKO (SEQ ID NO : 34)
CDR-H3:
YRHYAMDY (SEQ ID NO : 35)

8. Antibody according to any of claims 1 to 7, wherein said antibody is labelled with a compound chosen from the group comprising: a radionuclide, a fluor, an enzyme label, an enzyme substrate, an enzyme co-factor, enzyme inhibitor and a hapten.
9. Antibody according to any of claims 1 to 8, which is a humanised antibody.

10. Hybridoma producing an antibody according to any of claims 1 to 7.

11. Hybridoma according to claim 10, wherein said hybridoma have been deposited on August 23, 2007, at:

   BCCM / LMBP Plasmid Collection under the following Accession No:
   - TeiA 1.6 or 2.6F4C2 (IGH521) - > LMBP 6594CB
   - TeiA 1.7 or 2.8A3F8 (IGH522) - > LMBP 6595CB
   - TeiA 1.8 or 1.3B12H3 (IGH523) - > LMBP 6596CB
   - TeiA 2b.6 or 2.13E5E4 (IGH524) -- > LMBP 6597CB
   - TeiA 1.1 or 3.46B10E7 (IGH525) - > LMBP 6598CB

12. Peptide preparation to generate an immune response giving rise to antibody production which is efficient to reduce the amyloid deposits and to isolate an antibody and consisting of:

   Aβ 8-x mimicking peptide: SGYGVHH GC-KLH

13. Process of preparation of an antibody of claims 1 to 9, which specifically binds to the N-terminal region of Aβ_x peptide and does not recognise Aβ_{42}, x being comprised from 11 to 42, in particular 15 to 42, which presents a high specificity, and comprising an immunisation step of an appropriate animal with a Aβ_{8-x} peptide and a T-helper epitope, in particular with a Aβ_{8-x} peptide fused with a T-helper epitope, or a Aβ_{8-x} branched peptide, in particular a Aβ_{8,i5} peptide.

14. Process of preparation of an antibody according to claim 13, wherein said antibody binds specifically to the N-terminal region of Aβ_{8-i5} peptide, does not recognise Aβ_{42} and which presents a high specificity with respect to Aβ_{8-i5} peptide, such as determined on Western Blot.

15. Antibody, which binds specifically to the N-terminal region of Aβ_{8-x} peptide, such as obtained by a process as defined in claim 13 or 14.
16. A method for determining *in vitro* amyloid burden in a mammal, comprising the following steps:
   (i) quantifying the level of N-terminal truncated \( \text{A} \beta_{3-4} \) in a body fluid of said mammal, using the antibody of any of claims 1 to 9 or 15,
   (ii) comparing the level of antibody of said mammal to those obtained with a control mammal, and
   (iii) deducing from step (ii) if said mammal is suffering from a neurological disease provided the biomarker level is modified with respect to the level measured in the control mammal, in particular is higher than the level measured in the control mammal.

17. The method according to claim 16, wherein the mammal is a human.

18. The method according to claim 16 or 17, wherein specificity and sensitivity of said antibody toward \( \text{A} \beta_{3-42} \) is higher than 63%, preferably comprised from about 63 to about 100%, more preferably comprised from about 75% to 85% and more preferably comprised from 85% to 100%.

19. The method according to any of claims 16 to 18, wherein said body fluid is cerebrospinal fluid (CSF) or blood.

20. A method for determining, in a mammal, the susceptibility to a disease associated with \( \beta \)-amyloid formation and/or aggregation such as Alzheimer's disease, for determining, in a mammal, the risk of developing a disease associated with \( \beta \)-amyloid formation and/or aggregation such as Alzheimer's disease, for screening of the clearance of \( \beta \)-amyloid deposition in a mammal, or for predicting the level of \( \beta \)-amyloid burden in a mammal, said method comprising the following steps:
   (i) determining, in said mammal, the amount of peptide \( \text{A} \beta_{3-4} \) with an antibody according to any of claims 1 to 9 or 15,
   (ii) comparing the amount determined in step (i) with the amount of antibody specific of said N-terminal region of \( \text{A} \beta_{3-4} \) peptide in a control mammal, and
   (iii) concluding from the comparison in step (ii), whether the mammal is susceptible
to a disease associated with β-amyloid formation and/or aggregation such as Alzheimer's disease, whether the mammal is at risk of developing a disease associated with β-amyloid formation and/or aggregation such as Alzheimer's disease, whether the β-amyloid deposition in a mammal is cleared, or what the level of β-amyloid is in said mammal.

21. The method according to claim 20, wherein the amount of antibody specific of the N-terminal region of Aβ₈₋ₓ peptide is determined on a tissue sample obtained from said mammal.

22. A kit comprising at least one buffer, and at least one detection compound, at least one N-truncated Aβ₈₋ₓ specific antibody as defined in any of claims 1 to 9 or 15.

23. The kit of claim 22, further comprising a preferably labelled second antibody which binds to the antibody of any of claims 1 to 9 or 15.

24. A therapeutic composition comprising as active ingredient an antibody of any of claims 1 to 9 or 15, or comprising synthetic peptides with a free N-terminal-end mimicking the free N-terminal-end of N-truncated Aβ peptides, in association with a pharmaceutically acceptable vehicle.

25. The therapeutic composition of claim 24 being suitable for the administration to an individual of a dose of an antibody from 1 mg/kg/day to 200 mg/kg/day.

26. A vaccine composition comprising as active ingredient an antibody of any of claims 1 to 9 or 15, fragments or derivatives thereof, or comprising synthetic peptides with a free N-terminal-end mimicking the free N-terminal-end of N-truncated Aβ peptides, in association with a pharmaceutically acceptable vehicle.

27. The vaccine composition of claim 26 being suitable for the administration to an individual of a dose of an antibody from 1 mg/kg/day to 200 mg/kg/day.
28. Use of at least one of the antibodies of any of claims 1 to 9 or 15, for the preparation of a drug or a vaccine intended for the prevention or the treatment of Alzheimer disease.

29. Use of at least one of the antibodies of any of claims 1 to 9 or 15, for the preparation of a drug or a vaccine intended for the clearance of β-amyloid burden.

30. A method of clearance of β-amyloid burden in a mammal comprising the administration of a composition of claim 24 to 27 to the said mammal.

31. Use of a therapeutic or vaccine composition according to claim 24 or 26, for the induction of an immune response in a mammal being affected by or susceptible to develop an Alzheimer disease.
Figure 1

NH2-...SEWYMDAEFRHDSGSTVYEHVHLKLYLFFAEDVGSNGKAGIGGLMLGMGQVHAT

657

APP

β

α

γ
Figure 2
- Brain formic acid extract, SDS-PAGE, and Western-blotting with 21F12

Figure 4
Figure 6
Figure 12
Figure 13
INTERNATIONAL SEARCH REPORT

A. CLASSIFICATION OF SUBJECT MATTER

INV. C07K16/18 A61K39/395 G01N33/577 C12N5/18 C07K14/47
A61P25/28

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)
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 practical, search terms used)
EPO-Internal

C. DOCUMENTS CONSIDERED TO BE RELEVANT

<table>
<thead>
<tr>
<th>Category</th>
<th>Citation of document with indication, where appropriate, of the relevant passages</th>
<th>Relevant to claim No</th>
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<td>X</td>
<td>WO 2004/013172 A (INNOGENETICS NV [BE]); DELACOURTE ANDRE [FR]; SERGEANT NICOLAS [FR]) 12 February 2004 (2004-02-12) cited in the application the whole document, especially example 2, table 4, figures 6, 7</td>
<td>1-4, 8, 13-31</td>
</tr>
</tbody>
</table>

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 document 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

Date of the actual completion of the international search

10 December 2008

Date of mailing of the international search report

05/03/2009

Name and mailing address of the ISA/
European Patent Office, P B. 5818 Patentliana 2 NL - 2280 HV Rijswijk
Tel (+31-70) 340-2040, Fax (+31-70) 340-3016

Authorized officer

Bayer, Annette
<table>
<thead>
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<th>Citation of document, with indication, where appropriate, of the relevant passages</th>
<th>Relevant to claim No</th>
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<tbody>
<tr>
<td>A</td>
<td>SERGEANT N ET AL: &quot;Truncated beta-amyloid peptide species in pre-clinical Alzheimer's disease as new targets for the vaccination approach&quot; JOURNAL OF NEUROCHEMISTRY, NEW YORK, NY, US, vol. 85, no. 6, June 2003 (2003-06), pages 1581-1591, XP002257887 ISSN: 0022-3042 cited in the application the whole document, especially page 1589, last complete paragraph</td>
<td></td>
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</tbody>
</table>
**INTERNATIONAL SEARCH REPORT**

**Box No. II** Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. □ Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:

2. □ Claims Nos.: because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

3. □ Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

**Box No. III** Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see additional sheet

1. □ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.

2. □ As all searchable claims could be searched without effort justifying an additional fees, this Authority did not invite payment of additional fees.

3. □ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4. □ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

   see annex

**Remark on Protest**

- □ The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.

- □ The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.

- □ No protest accompanied the payment of additional search fees.

Form PCT/ISA/210 (continuation of first sheet (2)) (April 2005)
This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

<table>
<thead>
<tr>
<th>Claim Numbers</th>
<th>Description</th>
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<tr>
<td>1. 1-5, 8-10, 13-31 (all partially)</td>
<td>An antibody which specifically binds to the N-terminal region of Abeta8-x peptide, x being comprised from 11 to 42, and recognises neither Abetal-40 nor Abetal-42 and related subject-matter.</td>
</tr>
<tr>
<td>2. 1-11, 13-31 (all partially)</td>
<td>Monoclonal antibody TeiA 1.6, hybridoma producing said antibody and related subject-matter.</td>
</tr>
<tr>
<td>4. 1-11, 13-31 (all partially)</td>
<td>Monoclonal antibody TeiA 1.8, hybridoma producing said antibody and related subject-matter.</td>
</tr>
<tr>
<td>5. 1-11, 13-31 (all partially)</td>
<td>Monoclonal antibody TeiA 2b.6, hybridoma producing said antibody and related subject-matter.</td>
</tr>
<tr>
<td>7. 12 (complete)</td>
<td>Peptide preparation consisting of A-beta8-x mimicking peptide: SGYGVHHGC-KLH.</td>
</tr>
<tr>
<td>8. 24-27, 30, 31 (all partially)</td>
<td>A therapeutic composition comprising synthetic peptides with a free N-terminal-end mimicking the free N-terminal end of N-truncated A-beta peptides and related subject-matter.</td>
</tr>
<tr>
<td>Patent document cited in search report</td>
<td>Publication date</td>
</tr>
<tr>
<td>----------------------------------------</td>
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<tr>
<td>WO 2004013172 A</td>
<td>12-02-2004</td>
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