



(86) **Date de dépôt PCT/PCT Filing Date:** 2005/09/29
(87) **Date publication PCT/PCT Publication Date:** 2006/04/06
(45) **Date de délivrance/Issue Date:** 2019/06/25
(85) **Entrée phase nationale/National Entry:** 2007/03/16
(86) **N° demande PCT/PCT Application No.:** GB 2005/003756
(87) **N° publication PCT/PCT Publication No.:** 2006/035237
(30) **Priorité/Priority:** 2004/09/29 (GB0421639.6)

(51) **Cl.Int./Int.Cl. G01N 33/68** (2006.01),
A61P 25/28 (2006.01)

(72) **Inventeurs/Inventors:**
WESTBROOK, JULES, GB;
BYERS, HELEN, GB;
WARD, MALCOLM, GB;
LOVESTONE, SIMON, GB;
HYE, ABDUL, GB;
LYNHAM, STEPHEN, GB;
...

(73) **Propriétaires/Owners:**
PROTEOME SCIENCES PLC, GB;
KING'S COLLEGE LONDON, GB

(74) **Agent:** BERESKIN & PARR LLP/S.E.N.C.R.L., S.R.L.

(54) **Titre : PROCÉDES ET COMPOSITIONS LIÉS À LA MALADIE D'ALZHEIMER**
(54) **Title: METHODS AND COMPOSITIONS RELATING TO ALZHEIMER'S DISEASE**

(57) **Abrégé/Abstract:**

Methods and compositions relating to Alzheimer's disease are provided. Specifically, proteins that are differentially expressed in the Alzheimer's disease state relative to their expression in the normal state are provided. Proteins associated with Alzheimer's disease are identified and described. Methods of diagnosis of Alzheimer's disease using the differentially expressed proteins are also provided, as are methods for the identification and therapeutic use of compounds for the prevention and treatment of Alzheimer's disease.

(72) **Inventeurs(suite)/Inventors(continued):** JOUBERT, RICHARD, DE; PREFOT, PETRA, DE; KUHN, KARSTEN, DE; BAUMANN, CHRISTIAN, DE; SCHAEFER, JUERGEN, DE; PRINZ, THORSTEN, DE; KIENTLE, STEFAN, DE

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
6 April 2006 (06.04.2006)

PCT

(10) International Publication Number
WO 2006/035237 A3

(51) International Patent Classification:

G01N 33/68 (2006.01) **A61P 25/28** (2006.01)

(21) International Application Number:

PCT/GB2005/003756

(22) International Filing Date:

29 September 2005 (29.09.2005)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

0421639.6 29 September 2004 (29.09.2004) GB

(71) **Applicants** (for all designated States except US): **PROTEOME SCIENCES PLC** [GB/GB]; Coveham House, Downside Bridge Road, Cobham Surrey KT11 3EP (GB). **KING'S COLLEGE LONDON** [GB/GB]; Strand, London Greater London WC2R 2LS (GB).

(72) Inventors; and

(75) **Inventors/Applicants** (for US only): **WESTBROOK, Jules** [GB/GB]; Proteome Sciences plc, Coveham House, Downside Bridge Road, Cobham Surrey KT11 3EP (GB). **BYERS, Helen** [GB/GB]; Proteome Sciences plc, Coveham House, Downside Bridge Road, Cobham Surrey KT11 3EP (GB). **WARD, Malcolm** [GB/GB]; Proteome Sciences plc, Coveham House, Downside Bridge Road, Cobham Surrey KT11 3EP (GB). **LOVESTONE, Simon** [GB/GB]; King's College London, Strand, London Greater London WC2R 2LS (GB). **HYE, Abdul** [GB/GB]; King's College London, Strand, London Greater London WC2R 2LS (GB). **LYNHAM, Stephen** [GB/GB]; King's College London, Strand, London Greater London WC2R 2LS (GB). **JOUBERT, Richard** [FR/DE]; Proteome Sciences R & D GmbH & Co. KG, Altenhöferallee 3, 60438 Frankfurt (DE). **PREFOT, Petra** [DE/DE]; Proteome Sciences R & D GmbH & Co. KG, Altenhöferallee 3, 60438 Frankfurt (DE). **KUHN, Karsten** [DE/DE]; Proteome Sciences R & D GmbH & Co. KG, Altenhöferallee 3, 60438 Frankfurt (DE). **BAUMANN, Christian** [DE/DE]; Proteome Sciences R & D GmbH & Co. KG, Altenhöferallee 3, 60438 Frankfurt (DE). **SCHAEFER,**

Juergen [DE/DE]; Proteome Sciences R & D GmbH & Co. KG, Altenhöferallee 3, 60438 Frankfurt (DE). **PRINZ, Thorsten** [DE/DE]; Proteome Sciences R & D GmbH & Co. KG, Altenhöferallee 3, 60438 Frankfurt (DE). **KIENLE, Stefan** [DE/DE]; Proteome Sciences R & D GmbH & Co. KG, Altenhöferallee 3, 60438 Frankfurt (DE).

(74) **Agents:** **KIDDLE, Simon** et al.; Mewburn Ellis LLP, York House, 23 Kingsway, London Greater, London WC2B 6HP (GB).

(81) **Designated States** (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.

(84) **Designated States** (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

- with international search report
- before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments

(88) Date of publication of the international search report:

3 August 2006

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) **Title:** METHODS AND COMPOSITIONS RELATING TO ALZHEIMER'S DISEASE

(57) **Abstract:** Methods and compositions relating to Alzheimer's disease are provided. Specifically, proteins that are differentially expressed in the Alzheimer's disease state relative to their expression in the normal state are provided. Proteins associated with Alzheimer's disease are identified and described. Methods of diagnosis of Alzheimer's disease using the differentially expressed proteins are also provided, as are methods for the identification and therapeutic use of compounds for the prevention and treatment of Alzheimer's disease.

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Methods and Compositions Relating to Alzheimer's Disease**Field of the Invention**

The present invention relates to methods and compositions relating to Alzheimer's disease. Specifically, the present invention identifies and describes proteins that are differentially expressed in the Alzheimer's disease state relative to their expression in the normal state and, in particular, identifies and describes proteins associated with Alzheimer's disease. Further, the present invention provides methods of diagnosis of Alzheimer's disease using the differentially expressed proteins. Still further, the present invention provides methods for the identification and therapeutic use of compounds for the prevention and treatment of Alzheimer's disease.

Background of the Invention

Dementia is one of the major public health problems of the elderly, and in our ageing populations the increasing numbers of patients with dementia is imposing a major financial burden on health systems around the world. More than half of the patients with dementia have Alzheimer's disease (AD). The prevalence and incidence of AD have been shown to increase exponentially. The prevalence for AD in Europe is 0.3% for ages 60-69 years, 3.2% for ages 70-79 years, and 10.8% for ages 80-89 years (Rocca, Hofman et al. 1991). The survival time after the onset of AD is approximately from 5 to 12 years (Friedland 1993).

Alzheimer's disease (AD), the most common cause of dementia in older individuals, is a debilitating

neurodegenerative disease for which there is currently no cure. It destroys neurons in parts of the brain, chiefly the hippocampus, which is a region involved in coding memories. Alzheimer's disease gives rise to an irreversible progressive loss of cognitive functions and of functional autonomy. The earliest signs of AD may be mistaken for simple forgetfulness, but in those who are eventually diagnosed with the disease, these initial signs inexorably progress to more severe symptoms of mental deterioration. While the time it takes for AD to develop will vary from person to person, advanced signs include severe memory impairment, confusion, language disturbances, personality and behaviour changes, and impaired judgement. Persons with AD may become non-communicative and hostile. As the disease ends its course in profound dementia, patients are unable to care for themselves and often require institutionalisation or professional care in the home setting. While some patients may live for years after being diagnosed with AD, the average life expectancy after diagnosis is eight years.

In the past, AD could only be definitively diagnosed by brain biopsy or upon autopsy after a patient died. These methods, which demonstrate the presence of the characteristic plaque and tangle lesions in the brain, are still considered the gold standard for the pathological diagnoses of AD. However, in the clinical setting brain biopsy is rarely performed and diagnosis depends on a battery of neurological, psychometric and biochemical tests, including the measurement of biochemical markers such as the ApoE and tau proteins or

the beta-amyloid peptide in cerebrospinal fluid and blood.

Biomarkers may possibly possess the key in the next step for diagnosing AD and other dementias. A biological marker that fulfils the requirements for the diagnostic test for AD would have several advantages. An ideal biological marker would be one that identifies AD cases at a very early stage of the disease, before there is degeneration observed in the brain imaging and neuropathological tests. A biomarker could be the first indicator for starting treatment as early as possible, and also very valuable in screening the effectiveness of new therapies, particularly those that are focussed on preventing the development of neuropathological changes. A biological marker would also be useful in the follow-up of the development of the disease.

Markers related to pathological characteristics of AD; plaques and tangles (A β and tau) have been the most extensively studied. The most promising has been from studies of CSF concentration of A β (1-40), A β (1-42) and tau or the combination of both proteins in AD. Many studies have reported a decrease in A β (1-42) in CSF, while the total A β protein or A β (1-40) concentration remain unchanged (Ida, Hartmann et al. 1996; Kanai, Matsubara et al. 1998; Andreasen, Hesse et al. 1999).

Summary of the Invention

Broadly, the present invention relates to methods and compositions for the diagnosis of Alzheimer's disease. More specifically, the present invention identifies and

describes proteins that are differentially expressed in the Alzheimer's disease state relative to their expression in the normal state.

In a first aspect, the invention provides a method of diagnosing Alzheimer's disease in a subject, the method comprising detecting one or more of a differentially expressed protein identified by the methods described herein in a tissue sample or body fluid sample from said subject. Preferably, the method is an in vitro method.

In all aspects, the methods of the present invention may also be used in relation to pre-Alzheimer's stages such as mild cognitive impairment (MCI) as well as advanced Alzheimer's disease.

In another aspect, the present invention provides a method of determining the nature or degree of Alzheimer's disease in a human or animal subject, the method comprising detecting one or more of a differentially expressed protein identified by the methods described herein in a tissue sample or body fluid sample from said subject. Thus, the methods of the present invention encompass methods of monitoring the progress of Alzheimer's disease or of disease progression from MCI to Alzheimer's disease. Also encompassed are prognostic methods, for example prognosis of likely progression from MCI to Alzheimer's disease, or prognosis of likely duration or severity of Alzheimer's disease.

In a preferred embodiment the method comprises:

(a) establishing a paradigm in which at least one protein is differentially expressed in relevant tissue or body fluid sample from, or representative of, subjects having differential levels of Alzheimer's disease;

(b) obtaining a sample of the tissue or body fluid sample from the subject;

(c) determining the presence, absence or degree of expression of the differentially expressed protein or proteins in the sample; and

(d) relating the determination to the nature or degree of the Alzheimer's disease by reference to a previous correlation between such a determination and clinical information.

In one embodiment, the progression of the disorder may be tracked by using the methods of the invention to determine the severity of the disorder, e.g. global dementia severity). In another embodiment, the duration of the disorder up to the point of assessment may be determined using the methods of the invention.

For example, expression of an Ig lambda chain C region (see spot 177, Fig 6) may correlate with global dementia severity. Expression of a serum albumin precursor (see spot 165, Fig 6) may show a negative correlation with the duration of the disease.

This method allows the type of Alzheimer's disease of a patient to be correlated to different types to prophylactic or therapeutic treatment available in the art, thereby enhancing the likely response of the patient to the therapy.

In some embodiments, more than one protein is differentially expressed, providing a multi-protein fingerprint of the nature or degree of the Alzheimer's disease. Preferably, at least four proteins are differentially expressed.

Conveniently, the patient sample used in the methods of the invention can be a tissue sample or body fluid sample such as a blood, plasma, serum or urine sample. Use of body fluids such as those listed is preferred because they can be more readily obtained from a subject. This has clear advantages in terms of cost, ease, speed and subject wellbeing. Blood, blood products such as plasma, and urine are particularly preferred.

The step of detecting the differentially expressed protein may be preceded by a depletion step to remove the most abundant proteins from the sample, as described below.

Preferably, at least one of the differentially expressed proteins is a protein shown in Fig 6, Fig 7 or Fig 12. In preferred embodiments, the differentially expressed protein is apolipoprotein A-IV precursor, apolipoprotein C-III precursor, transthyretin, galectin 7, complement C4 precursor, alpha-2-macroglobulin precursor, Ig alpha-1 chain C, histone 2B, Ig lambda chain C region, fibrinogen gamma chain precursor, complement factor H, inter-alpha-trypsin heavy chain H4 precursor, complement C3 precursor, clusterin precursor, gamma or beta actin, haptoglobin precursor or the serum albumin precursor isoform found in spot ID no 2, 14, 15, 123, 165, 176 or

184 of Fig 6 or fragments thereof. Preferred fragments are a C-terminal fragment of Apo-AIV or a C4 alpha region of complement C4 precursor lacking the anaphylatoxin domain. For example, the fragment may comprise amino acid residues 270-309 of apolipoprotein A-IV; residues 1446-1744 of complement C4, or may be an N-terminal fragment of apolipoprotein A-IV which migrates as a polypeptide of 10-16kD or a polypeptide of 28 kD in SDS-PAGE, or a fragment of any of the proteins in Fig 7 with a molecular weight of 6430, 14640, 27147 or 14646 Da. Other preferred fragments comprise the areas indicated in bold in figures 9, 10, and 13 to 19.

Preferred fragments are less than 50, less than 100, less than 150 less than 200, less than 250, less than 300, less than 350, less than 400, less than 500, less than 600, less than 700, less than 800, less than 900, less than 1000, less than 1100, less than 1200, less than 1300, less than 1400, less than 1500, less than 1600, less than 1700, less than 1800, less than 1900 or less than 2000 amino acids in length.

The expression of certain differentially expressed proteins may be increased in subjects with Alzheimer's disease as compared to control subjects. The expression of other differentially expressed proteins may be decreased in subjects with Alzheimer's disease as compared to control subjects. Figs 6, 8 and 12 indicate whether the expression of the proteins disclosed therein is increased or decreased in Alzheimer's versus control subjects. It is thus clear from the figures whether an increase or decrease in expression is indicative of the

disease state for all the proteins listed therein.
Including the preferred proteins listed above.

Preferably, a differentially expressed protein shows a fold difference in expression of at least 1.5, at least 1.6, at least 1.7, at least 1.8, at least 1.9, at least 2.0, at least 2.5, at least 3, at least 3.5, at least 4, at least 5, at least 10 or more between the level found in patients with Alzheimer's versus control subjects.

The differentially expressed protein may be detected using an antibody specific to that protein, for example in an ELISA assay or Western blotting. Alternatively, the differentially expressed protein may be detected by, amongst others, 2D gel electrophoresis or mass spectrometry techniques including LS/MS/MS, MALDI-TOF or SELDI-TOF. The sample may be immobilised on a solid support for analysis.

In one embodiment, a diagnosis may be made solely on the basis of the pattern of spots on a 2D gel prepared from a subject sample. The pattern of spots obtained from Alzheimer's disease or MCI subjects may be compared directly with the pattern obtained from control subject samples, without the need for identifying individual proteins.

In one embodiment, an antibody sandwich technique where antibodies specific for one or more of the biomarkers is added and the immobilised antibodies capture the biomarker protein. The captured proteins are then detected using a second antibody that may be directly

labelled with a signal generating agent (enzyme, fluorescent tag, radiolabel etc.) or may be detected using further amplification (labelled secondary antibody, streptavidin/biotin systems with enzyme, fluorophore, radiolabel etc.). Other immunological methods may include one-dimensional or two-dimensional gel electrophoresis of patient samples followed by transfer to a solid surface using techniques such as Western blotting and subsequent detection using antibodies specific for the AD biomarkers.

In an alternative embodiment, autoantibodies to the biomarkers may be detected by using the Western blotting approach described above using either samples from a patient or representative of AD and then detecting the presence of antibodies specific for the biomarker that are present in the blood of AD patients but not in controls.

The method may further comprise determining an effective therapy for treating the Alzheimer's disease.

In a further aspect, the present invention provides a method of treatment by the use of an agent that will restore the expression of one or more differentially expressed proteins in the Alzheimer's disease state towards that found in the normal state in order to prevent the development or progression of Alzheimer's disease. Preferably, the expression of the protein is restored to that of the normal state.

In a further aspect, the present invention provides a method whereby the pattern of differentially expressed proteins in a tissue sample or body fluid sample or urine of an individual with Alzheimer's disease is used to predict the most appropriate and effective therapy to alleviate the Alzheimer's disease.

Also provided is a method of screening an agent to determine its usefulness in treating a Alzheimer's disease, the method comprising:

- (a) obtaining a sample of relevant tissue taken from, or representative of, a subject having Alzheimer's disease symptoms, who or which has been treated with the agent being screened;
- (b) determining the presence, absence or degree of expression of the differentially expressed protein or proteins in the tissue from, or representative of, the treated subject; and,
- (c) selecting or rejecting the agent according to the extent to which it changes the expression, activity or amount of the differentially expressed protein or proteins in the treated subject having Alzheimer's disease symptoms.

Optionally, the method may further comprise, prior to step (a), the step of establishing a paradigm in which at least one protein is differentially expressed in relevant tissue from, or representative of, subjects having Alzheimer's disease symptoms and normal subjects.

Preferably, the agent is selected if it converts the expression of the differentially expressed protein

towards that of a normal subject. More preferably, the agent is selected if it converts the expression of the protein or proteins to that of the normal subject.

Also provided is a method of screening an agent to determine its usefulness in treating Alzheimer's disease, the method comprising:

- (a) obtaining over time samples of relevant tissue or body fluid taken from, or representative of, a subject having Alzheimer's disease symptoms, who or which has been treated with the agent being screened;
- (b) determining the presence, absence or degree of expression of a differentially expressed protein or proteins in said samples; and,
- (c) determining whether the agent affects the change over time in the expression of the differentially expressed protein in the treated subject having Alzheimer's disease symptoms.

Optionally, the method may further comprise, prior to step (a), the step of establishing a paradigm in which at least one protein is differentially expressed in relevant tissue or body fluid from, or representative of, subjects having Alzheimer's disease symptoms and normal subjects; and/or

establishing that expression of said differentially expressed protein diverges over time in subjects having Alzheimer's disease symptoms and normal subjects.

Samples taken over time may be taken at intervals of weeks, months or years. For example, samples may be taken at monthly, two-monthly, three-monthly, four-

monthly, six-monthly, eight-monthly or twelve-monthly intervals.

A change in expression over time may be an increase or decrease in expression, compared to the initial level of expression in samples from the subject and/or compared to the level of expression in samples from normal subjects. The agent is selected if it slows or stops the change of expression over time.

In the screening methods described above, subjects having differential levels of protein expression comprise:

- (a) normal subjects and subjects having Alzheimer's disease symptoms; and,

- (b) subjects having Alzheimer's disease symptoms which have not been treated with the agent and subjects Alzheimer's disease which have been treated with the agent.

In alternative embodiments, the subjects having differential levels of protein expression comprise:

- (a) normal subjects who have and have not been treated with the agent; and one or both of

- (b) subjects having mild cognitive impairment who have and have not been treated with the agent; and

- (c) subjects having Alzheimer's disease symptoms who have and have not been treated with the agent.

Preferably, the differential levels of protein expression are not observed in normal subjects who have and have not been treated with the agent.

The subjects having Alzheimer's disease symptoms are preferably human subjects with Alzheimer's disease.

Alternatively, the subjects having Alzheimer's disease symptoms may be an animal model such as mutant amyloid precursor protein (APP) transgenic mice, presenilin-1 (PS-1) transgenic mice, and/or double transgenic APP/PS-1 transgenic mice.

The tissue or body fluid samples may be, for example, brain tissue, blood, plasma, serum, saliva or cerebrospinal fluid samples.

In one embodiment, the paradigm is established using two-dimensional (2D) gel electrophoresis carried out on the relevant tissue or a protein-containing extract thereof.

In another embodiment, the paradigm is established using SELDI analysis of the relevant tissue or a protein-containing extract thereof. Preferably, the tissue or extract is immobilised on a solid support, for example a chip.

Conveniently, a depletion step may be performed prior to 2D gel electrophoresis or SELDI analysis, to remove the most abundant proteins from the samples and reduce background.

The method may further comprise the step of isolating a differentially expressed protein identified in the method, and optionally the step of characterising the isolated protein.

Preferably, at least one of the differentially expressed proteins is a protein shown in Fig 6, Fig 7, Fig 8 or Fig 12 or a rodent equivalent thereof. In preferred embodiments, the differentially expressed protein is apolipoprotein A-IV precursor, apolipoprotein C-III precursor, transthyretin, galectin 7, complement C4 precursor, complement factor H, S100 calcium binding protein or ceruloplasmin, inter-alpha-trypsin heavy chain H4 precursor, complement C3 precursor, clusterin precursor, gamma or beta actin, haptoglobin precursor or fragments thereof. Preferred fragments are a C-terminal fragment of Apo-AIV or a C4 alpha region of complement C4 precursor lacking the anaphylatoxin domain. For example, the fragment may comprise amino acid residues 270-309 of apolipoprotein A-IV; residues 1446-1744 of complement C4.

Preferred fragments will comprise one or more of the sequences highlighted in Figs 9, 10 and 13-19.

In a further aspect, the invention provides a method of making a pharmaceutical composition which comprises having identified an agent using the method described above, the further step of manufacturing the agent and formulating it with an acceptable carrier to provide the pharmaceutical composition.

In a further aspect, the invention provides a method of identifying a protein which is differentially expressed in relevant tissue or body fluid sample from subjects

with mild cognitive impairment and/or subjects with Alzheimer's disease and normal subjects, comprising:

- i) immobilising a tissue sample or body fluid sample or protein-containing extract thereof on a solid support
- ii) analysing the immobilised proteins by surface enhanced laser desorption time of flight mass spectroscopy
- iii) comparing the spectra obtained to detect differences in protein expression between Alzheimer's subjects and normal subjects.

Also provided is protein which is differentially expressed in relevant tissue from, or representative of subjects having differential levels of Alzheimer's disease symptoms and which is as obtainable by the methods described herein or by two-dimensional gel electrophoresis carried out on said tissue or a protein-containing extract thereof, the method comprising:

- (a) providing non-linear immobilized pH gradient (IPG) strips of acrylamide polymer 3 mm x 180 mm;

- (b) rehydrating the IPG strips in a cassette containing 25 ml. of an aqueous solution of urea (8M), 3-[(Cholamidopropyl)dimethylammonio]-1-propanesulphonate (CHAPS, 2% w/v), 0.5% IPG Pharmalyte and a trace of Bromophenol Blue;

- (c) emptying the cassette of liquid, transferring the strips to an electrophoretic tray fitted with humid electrode wicks, electrodes and sample cups, covering the strips and cups with low viscosity paraffin oil;

- (d) applying 200 micrograms of an aqueous solution of dried, powdered material of the relevant body tissue in urea (8M), CHAPS (4% w/v), Tris (40 mM), 0.5% IPG

Pharmalyte and a trace of Bromophenol Blue to the sample cups, at the cathodic end of the IPG strips;

(e) carrying out isoelectric focusing on the gel at S1 500V step-n-hold (s/h) for 1h; S2 500V s/h for 2h; S3 1000V gradient (G) for 1h; S4 1000V s/h for 2h; S5 8000V G for 2h and S6 8000V s/h for a time effective to enable the proteins to migrate in the strips to their pI-dependent final positions;

(f) equilibrating the strips within the tray with 100 ml of an aqueous solution containing Tris-HCl (50 mM) pH 6.8, urea (6M), glycerol (30% v/v), SDS (2% w/v) and DTT (10mg/ml) ;

(g) replacing this solution by 100 ml. of an aqueous solution containing Tris-HCl (50 mM) pH 8.8, urea (6M), glycerol (30% v/v), SDS (2% w/v), iodoacetamide (25mg/ml) and a trace of Bromophenol Blue and incubating for 20 minutes;

(h) providing a vertical gradient slab gel 160 x 200 x 1.5 mm of acrylamide/piperazine-diacrylyl cross-linker(9-16%T/2.6%C), polymerised in the presence of TEMED (0.5% w/v), ammonium persulphate (0.1% w/v) and sodium thiosulphate (5 mM), in Tris-HCl (0.375M) pH 8.8 as leading buffer;

(i) over-layering the gel with sec-butanol for about 2 hours, removing the overlay and replacing it with water;

(j) cutting the IPG gel strips to a size suitable for the second dimensional electrophoresis, removing 6 mm from the anode end and 14 mm from the cathode end;

(k) over-layering the slab gel with an aqueous solution of agarose (0.5% w/v) and Tris-glycine-SDS (25 mM-198 mM-0.1% w/v) as leading buffer, heated to 70°C and loading

the IPG gel strips onto the slab gel through this overlaid solution;

(l) running the second dimensional electrophoresis at a constant current of 40 mA at 8-12°C for 5 hours; and
(m) washing the gel.

This invention is based, in part, on systematic search strategies involving sensitive detection of proteins by 2D-electrophoresis. To aid the identification of differentially expressed protein a standard marker set of proteins such as those available from Genomic Solutions may be run on an extra lane to 2D electrophoresis.

The examples presented below demonstrate the successful use of the experimental paradigms of the invention to identify target proteins associated with Alzheimer's disease.

Definitions

"Differential expression", as used herein, refers to at least one recognisable difference in tissue or body fluid protein expression. It may be a quantitatively measurable, semi-quantitatively estimatable or qualitatively detectable difference in tissue protein expression. Thus, a differentially expressed protein (herein DEP) may be strongly expressed in tissue in the normal state and less strongly expressed or not expressed at all in tissue in the Alzheimer's disease state. Conversely, it may be strongly expressed in tissue in the Alzheimer's disease state and less strongly expressed or not expressed at all in the normal state. Further, expression may be regarded as differential if the protein

undergoes any recognisable change between the two states under comparison.

The term "paradigm" means a prototype example, test model or standard.

Wherever a differentially expressible protein is used in the screening procedure, it follows that there must have been at some time in the past a preliminary step of establishing a paradigm by which the differential expressibility of the protein was pre-determined. Once the paradigm has been established, it need not be re-established on every occasion that a screening procedure is carried out. The term "establishing a paradigm" is to be construed accordingly.

"Relevant tissue" means any tissue involved in brain function, in particular tissue involved in Alzheimer's disease.

"Tissue/Body fluid... ..representative of... ..subjects" means any tissue or body fluid in which the above-mentioned biological change can be simulated for laboratory purposes and includes, for example, a primary cell culture or cell line derived ultimately from relevant tissue.

The term "subjects" includes human and animal subjects.

The treatments referred to above can comprise the administration of one or more drugs or foodstuffs, and/or other factors such as diet or exercise.

The differentially expressed proteins (DEPs) include "fingerprint proteins", "target proteins" or "pathway proteins".

The term "fingerprint protein", as used herein, means a DEP, the expression of which can be used, alone or together with other DEPs, to monitor or assess the condition of a patient suspected of suffering from Alzheimer's disease. Since these proteins will normally be used in combination, especially a combination of four or more, they are conveniently termed "fingerprint proteins", without prejudice to the possibility that on occasions they may be used singly or along with only one or two other proteins for this purpose. Such a fingerprint protein or proteins can be used, for example, to diagnose a particular type of Alzheimer's disease and thence to suggest a specific treatment for it.

The term "diagnosis", as used herein, includes the provision of any information concerning the existence, non-existence or probability of the disorder in a patient. It further includes the provision of information concerning the type or classification of the disorder or of symptoms which are or may be experienced in connection with it. This may include, for example, diagnosis of the severity of the disorder. It encompasses prognosis of the medical course of the disorder, for example its duration, severity and the course of progression from MCI to Alzheimer's disease.

Currently disease status is assessed by duration of disease from inception to present (longer duration equals more severe disease) and clinical assessment measures. These assessment measures include clinical tests for memory and other cognitions, clinical tests for function (abilities of daily living) and clinical assessments of global severity. Trials of potential therapies in AD are currently evaluated against these measures. The FDA and other medicines approval bodies require as part of these assessments measures of both cognition and global function. The Global Dementia Scale is one such measure of global function. It is assessed by rater assessment of severity including cognition and function against a standardised set of severity criteria.

The term "target protein", as used herein, means a DEP, the level or activity of which can be modulated by treatment to alleviate Alzheimer's disease. Modulation of the level or activity of the target protein in a patient may be achieved, for example, by administering the target protein, another protein or gene which interacts with it or an agent which counteracts or reduces it, for example an antibody to the protein, competitive inhibitor of the protein or an agent which acts in the process of transcription or translation of the corresponding gene.

The term "alleviate", as used herein, in relation to Alzheimer's disease means any form of reducing one or more undesired symptoms or effects thereof. Any amelioration of Alzheimer's disease of the patient falls

within the term "alleviation". Amelioration may also include slowing down the progression of the disease.

Alternatively or additionally, the DEPs can interact with at least one other protein or with a gene involved in the regulation of brain function. Such other proteins are termed herein "pathway proteins" (PPs). The term is applied to the protein with which the DEP interacts, not to the DEP itself, although a pathway protein can be another DEP.

By way of example, embodiments of the present invention will now be described in more detail with reference to the accompanying figures.

Brief Description of the Figures

Fig 1 shows spectra for the 6430 Da peak identified by SELDI analysis in normal (top) and Alzheimer's disease (bottom) subjects.

Fig 2 shows spectra for the 14640 Da peak identified by SELDI analysis in normal (top) and Alzheimer's disease (bottom) subjects.

Fig 3 shows spectra for the 27147 Da peak identified by SELDI analysis in normal (top) and Alzheimer's disease (bottom) subjects.

Fig 4 shows spectra for the 14646 Da peak identified by SELDI analysis in pooled normal (top) and Alzheimer's disease (bottom) subjects.

Fig 5 shows a silver stained gel obtained from material extracted from the chips used for SELDI analysis. The bands (1-6) excised and analysed by LC/MS/MS are indicated by arrows.

Fig 6 shows differentially expressed proteins identified by 2D gel analysis and mass spectroscopy.

Fig 7 shows differentially expressed proteins identified by SELDI and LC/MS/MS.

Fig 8 shows differentially expressed proteins identified by qPST.

Fig 9 shows the sequence coverage (indicated in bold) obtained for apolipoprotein A-IV (P06727) in the 14.6kDa band isolated on the Q10 SAX2 SELDI chip. C-terminal residues 270 - 396 are underlined.

Fig 10 shows sequence coverage (indicated in bold) obtained for Complement C4 precursor (P01028) in 2DE spot 164.

Fig 11 shows a 2D gel obtained from the pre-depletion analysis. The differentially expressed spots are circled.

Fig 12 lists the differentially expressed spots identified by the pre-depletion analysis. Column 3 gives the accession number for the human protein, column 4 the mean normalised spot volume in the control samples, column 6 the mean normalised spot volume in the disease

samples, column 8 the mean normalised spot volume in the disease sample divided by that in the control sample, column 9 the significance (p-value) of the difference in spot volumes compared by Student's t-test, column 10 the number of gels in the control group in which the spot was detected. CV is coefficient of variation.

Fig 13 shows sequence coverage (indicated in bold) obtained for alpha-2 macroglobulin (P01023) in the pre-depletion analysis. The signal sequence is underlined.

Fig 14 shows sequence coverage (indicated in bold) obtained for inter-alpha trypsin inhibitor heavy chain H4 precursor (Q14624) in the pre-depletion analysis. The signal sequence is underlined.

Fig 15 shows sequence coverage (indicated in bold) obtained for complement C3 precursor (P01024) in the pre-depletion analysis. The signal sequence is underlined.

Fig 16 shows sequence coverage (indicated in bold) obtained for clusterin precursor (P10909) in the pre-depletion analysis. The signal sequence is underlined.

Fig 17 shows sequence coverage (indicated in bold for spot 832 and bold italic for spot 652) obtained for complement C4 precursor (P01028) in the pre-depletion analysis. The signal sequence is underlined.

Fig 18 shows sequence coverage (indicated in bold) obtained for gamma actin (P63261) in the pre-depletion analysis. The signal sequence is underlined.

Fig 19 shows sequence coverage (indicated in bold) obtained for haptoglobin precursor (P00738) in the pre-depletion analysis. The signal sequence is underlined.

Detailed Description

Methods and compositions for the treatment of Alzheimer's disease. Proteins termed 'target proteins' and/or fingerprint proteins are described which are differentially expressed in Alzheimer's disease states relative to their expression in normal states. Methods for the identification of such fingerprint and target proteins are also described.

'Differential expression' as used herein indicates that a protein is present at different levels in samples from normal and diseased subjects.

Also described below are methods for prognostic and diagnostic evaluation of Alzheimer's disease states and for the identification of subjects exhibiting a predisposition to Alzheimer's disease.

1. Identification of differentially expressed and pathway proteins

In one embodiment, the present invention concerns methods for the identification of proteins which are involved in Alzheimer's disease. Such proteins may represent proteins which are differentially expressed in Alzheimer's disease states relative to their expression in normal states. Such differentially expressed proteins may represent 'target' or 'fingerprint' proteins.

Methods for the identification of such proteins are described in Section 1. Methods for the further characterisation of such differentially expressed proteins and for their identification as target and/or fingerprint proteins are presented below in Section 1.1.

'Differential expression', as used herein, refers to both qualitative as well as quantitative differences in protein expression. Thus a differentially expressed protein may qualitatively have its expression activated or completely inactivated in normal versus Alzheimer's disease state. Such a qualitatively regulated protein will exhibit an expression pattern within a given tissue, cell type or body fluid sample which is detectable in either control or Alzheimer's disease subject, but not detectable in both. Alternatively, such a qualitatively regulated protein will exhibit an expression pattern within a given tissue, cell type or body fluid sample, which is detectable in either control or Alzheimer's disease subjects but not detectable in both.

'Detectable', as used herein, refers to a protein expression pattern, which are detectable using techniques described herein.

Alternatively, a differentially expressed protein may have its expression modulated, i.e. quantitatively increased or decreased, in normal versus Alzheimer's disease states. The degree to which expression differs in normal versus Alzheimer's disease states need only be large enough to be visualised via standard characterisation techniques, such as silver staining of 2D-electrophoretic gels. Other such standard

characterisation techniques by which expression differences may be visualised are well known to those skilled in the art. These include successive chromatographic separations of fractions and comparisons of the peaks, capillary electrophoresis, separations using micro-channel networks, including on a micro-chip, SELDI analysis and qPST analysis.

Chromatographic separations can be carried out by high performance liquid chromatography as described in Pharmacia literature, the chromatogram being obtained in the form of a plot of absorbance of light at 280 nm against time of separation. The material giving incompletely resolved peaks is then re-chromatographed and so on.

Capillary electrophoresis is a technique described in many publications, for example in the literature "Total CE Solutions" supplied by Beckman with their P/ACE 5000 system. The technique depends on applying an electric potential across the sample contained in a small capillary tube. The tube has a charged surface, such as negatively charged silicate glass. Oppositely charged ions (in this instance, positive ions) are attracted to the surface and then migrate to the appropriate electrode of the same polarity as the surface (in this instance, the cathode). In this electroosmotic flow (EOF) of the sample, the positive ions move fastest, followed by uncharged material and negatively charged ions. Thus, proteins are separated essentially according to charge on them.

Micro-channel networks function somewhat like capillaries and can be formed by photoablation of a polymeric material. In this technique, a UV laser is used to generate high energy light pulses that are fired in bursts onto polymers having suitable UV absorption characteristics, for example polyethylene terephthalate or polycarbonate. The incident photons break chemical bonds with a confined space, leading to a rise in internal pressure, mini-explosions and ejection of the ablated material, leaving behind voids which form micro-channels. The micro-channel material achieves a separation based on EOF, as for capillary electrophoresis. It is adaptable to micro-chip form, each chip having its own sample injector, separation column and electrochemical detector: see J.S.Rossier et al., 1999, Electrophoresis 20: pages 727-731.

Surface enhanced laser desorption ionisation time of flight mass spectrometry (SELDI-TOF-MS) combined with ProteinChip technology can also provide a rapid and sensitive means of profiling proteins and is used as an alternative to 2D gel electrophoresis in a complementary fashion. The ProteinChip system consists of aluminium chips to which protein samples can be selectively bound on the surface chemistry of the chip (eg. anionic, cationic, hydrophobic, hydrophilic etc). Bound proteins are then co-crystallised with a molar excess of small energy-absorbing molecules. The chip is then analysed by short intense pulses of N2 320nm UV laser with protein separation and detection being by time of flight mass spectrometry. Spectral profiles of each group within an experiment are compared and any peaks of interest can be

further analysed using techniques as described below to establish the identity of the protein.

Quantitative protein sequence tag (qPST) technology may also be used to detect differentially expressed proteins. Briefly, the proteins in the samples for comparison are labelled with a stable isotope tag. A different isotope is used for each sample. The proteins are enzymatically cleaved and the labelled peptides in each sample are quantified by mass spectrometry. In this way, expression of equivalent proteins in the different samples can be compared directly by comparing the intensities of their respective isotopic peaks.

Detection of differentially expressed proteins may be preceded by a depletion step to remove the most abundant proteins from the sample. The large majority of the protein composition of serum/plasma consists of just a few proteins. For example, albumin, which is present at a concentration of 35-50 mg/ml, represents approximately 54% of the total protein content with IgG adding other 16%. In contrast, proteins changing in response to disease, for example as a result of tissue leakage, may circulate at 10ng/ml. This vast dynamic range of protein concentrations represents a major analytical challenge and to overcome the problem, a multiple affinity depletion column can be used to remove the most highly abundant proteins (eg the 5, 6, 7, 8, 9 or 10 most highly abundant proteins). This enables the detection of changes in lower abundance ranges because more starting material can be used and there is less interference from

the highly abundant molecules. Such a depletion strategy can be applied before any detection method.

Differentially expressed proteins may be further described as target proteins and/or fingerprint proteins. 'Fingerprint proteins', as used herein, refer to a differentially expressed protein whose expression pattern may be utilised as part of a prognostic or diagnostic Alzheimer's disease evaluation or which, alternatively, may be used in methods for identifying compounds useful for the treatment of Alzheimer's disease. A fingerprint protein may also have characteristics of a target protein or a pathway protein.

'Target protein', as used herein, refers to a differentially expressed protein involved in Alzheimer's disease such that modulation of the level or activity of the protein may act to prevent the development of Alzheimer's disease. A target protein may also have the characteristics of a fingerprint protein or a pathway protein.

1.1 Characterisation of differentially expressed proteins

Differentially expressed proteins, such as those identified via the methods discussed above, may be further characterised by using, for example, methods such as those discussed herein. Such proteins will be referred to herein as 'identified proteins'.

Analyses such as those described herein, yield information regarding the biological function of the

identified proteins. An assessment of the biological function of the differentially expressed proteins, in addition, will allow for their designation as target and/or fingerprint proteins.

Specifically, any of the differentially expressed proteins whose further characterisation indicates that a modulation of the proteins expressed or a modulation of the proteins activity may ameliorate Alzheimer's disease will be designated 'target proteins', as defined above, in Section 1.

Any of the differentially expressed proteins whose expression pattern contributes to a protein 'fingerprint' profile correlative of Alzheimer's disease, will be designated a 'fingerprint protein'. 'Fingerprint profiles' will be more fully discussed below. It should be noted that each of the target proteins may also function as fingerprint proteins.

A variety of techniques can be utilised to further characterise the identified proteins. First, the corresponding nucleotide sequence of the identified protein may be obtained by utilising standard techniques well known to those of skill in the art, may, for example, be used to reveal homologies to one or more known sequence motifs which may yield information regarding the biological function of the identified protein.

Secondly, the biological function of the identified proteins may be more directly assessed by utilising

relevant in vivo and in vitro systems. In vivo systems may include, but are not limited to, animal systems which naturally exhibit Alzheimer's disease-like symptoms and/or pathology, or ones which have been engineered to exhibit such symptom and/or pathology. Further, such systems may include systems for the further characterisation of Alzheimer's disease, and may include, but are not limited to, naturally occurring and transgenic animal systems.

In vitro systems may include cell lines derived from such animals or Alzheimer's disease subjects. Animal models may be used to generate cell lines, containing one or more cell types involved in Alzheimer's disease, that can be used as cell culture models for this disorder. While primary cultures derived from the transgenic animals of the invention may be utilised, the generation of continuous cell lines is preferred. For examples of techniques which may be used to derive a continuous cell line from the transgenic animals, see Small, et al., 1985, Mol. Cell Biol. 5: 642-648.

Preferred transgenic animal models of Alzheimer's disease include mice overexpressing glycogen synthase kinase (GSK) (see Lucas et al (2001) EMBO J. 20, p27-39), mice overexpressing mutant alleles of APP or PS1 and double (APP/PS1) transgenic mouse models overexpressing mutant alleles of both APP and PS1. Double transgenic mice resulting from a cross between a mutant APP line Tg2576 and a mutant PS1M146L transgenic line is reported in Holcomb et al., Nat Med. 1998 Jan;4(1):97-100).

In further characterising the biological function of the identified proteins, the expression of these proteins may be modulated within the in vivo and/or in vitro systems, i.e. either overexpressed or underexpressed in, for example, transgenic animals and/or cell lines, and its subsequent effect on the system then assayed.

Alternatively, the activity of the identified protein may be modulated by either increasing or decreasing the level of activity in the in vivo and/or in vitro system of interest, and its subsequent effect then assayed.

The information obtained through such characterisations may suggest relevant methods for the treatment of Alzheimer's disease using the protein of interest. For example, treatment may include a modulation of protein expression and/or protein activity. Characterisation procedures such as those described herein may indicate where such modulation should involve an increase or a decrease in the expression or activity of the protein of interest. Such methods of treatment are discussed below in Section 4.

2. Differentially expressed proteins

Identified proteins, which include differentially expressed proteins such as those identified in Section 1 above, are described herein. Specifically, the amino acid sequences of such identified proteins are described. Further, antibodies directed against the identified protein, and cell- and animal-based models by which the identified proteins may be further characterised and utilised are also discussed in this Section.

2.1 Antibodies specific for differentially expressed or pathway proteins

The present invention also relates to methods for the production of antibodies capable of specifically recognising one or more differentially expressed or pathway protein epitopes. Such antibodies may include, but are not limited to, polyclonal antibodies, monoclonal antibodies (mAbs), humanised or chimeric antibodies, single chain antibodies, Fab fragments, F(ab')₂ fragments, fragments produced by a Fab expression library, anti-idiotypic (anti-Id) antibodies, and epitope-binding fragments of any of the above. Such antibodies may be utilised as part of Alzheimer's disease treatment methods, and/or may be used as part of diagnostic techniques whereby patients may be tested for abnormal levels of fingerprint, target, or pathway gene proteins, or for the presence of abnormal forms of such proteins.

For the production of antibodies to a differentially expressed or pathway protein, various host animals may be immunised by injection with a differentially expressed or pathway protein, or a portion thereof. Such host animals may include, but are not limited to, rabbits, mice and rats, to name but a few. Various adjuvants may be used to increase the immunological response, depending on the host species, including active substances such as lysolecithin, Pluronic polyols, polyanions, peptides, oil emulsions, keyhole limpet hemocyanin, dinitrophenol, and potentially useful human adjuvant such as BCG bacille Calmette-Fuerin) and *Corynebacterium parvum*.

Polyclonal antibodies are heterogeneous populations of antibody molecules derived from the sera of animals immunised with an antigen, such as target proteins, or an antigenic functional derivative thereof. For the production of polyclonal antibodies, host animals such as those described above, may be immunised by injection with differentially expressed or pathway protein supplemented with adjuvants as also described above.

Monoclonal antibodies, which are homogeneous populations of antibodies to a particular antigen, may be obtained by any technique, which provides for the production of antibody molecules by continuous cell lines in culture. These include, but are not limited to, the hybridoma technique of Kohler and Milstein (1975, Nature 256; 495-497; and US Patent No: 4,376,110), the human β -cell hybridoma technique (Kosbor, et al., 1983, Immunology Today 4: 72; Cole, et al., 1983, Proc. Natl. Acad. Sci. USA 80: 2026-2030), and the EBV-hybridoma technique (Cole, et al., 1985, Monoclonal Antibodies and Cancer Therapy, Alan R. Liss Inc., pp. 77-96). Such antibodies may be of any immunoglobulin class including IgG, IgM, IgE, IgA, IgD and any subclass thereof. The hybridoma producing the mAb of this invention may be cultivated in vitro or in vivo. Production of high titers of mAbs in vivo makes this the presently preferred method of production.

In addition, techniques developed for the production of 'chimeric antibodies' (Morrison, et al., 1984, Proc. Natl. Acad. Sci. 81: 6851-6855; Neuberger, et al., 1984, Nature 312: 604-608; Takeda, et al., 1985, Nature 314:

452-454) by splicing the genes from a mouse antibody molecule of appropriate antigen specificity together with genes from a human antibody molecule of appropriate biological activity can be used. A chimeric antibody is a molecule in which different portions are derived from different animal species, such as those having a variable region derived from a murine mAb and a human immunoglobulin constant region.

Alternatively, techniques described for the production of single chain antibodies (US Patent No: 4,946,778; Bird, 1988, Science 242: 423-426; Huston, et al., 1988, Proc. Natl. Acad. Sci. USA 85: 5879-5883; and Ward, et al., 1989, Nature 334: 544-546) can be adapted to produce differentially expressed or pathway protein-single chain antibodies. Single chain antibodies are formed by linking the heavy and light chain fragments of the Fv region via an amino acid bridge, resulting in a single chain polypeptide.

Antibody fragments, which recognise specific epitopes, may be generated by known techniques. For example, such fragments include, but are not limited to, the F(ab')₂ fragments which can be produced by pepsin digestion of the antibody molecule and the Fab fragments which can be generated by reducing the disulfide bridges of the F(ab')₂ fragments. Alternative, Fab expression libraries may be constructed (Huse, et al., 1989, Science 246: 1275-1281) to allow rapid and easy identification of monoclonal Fab fragments with the desired specificity.

3 Assays for amelioration of Alzheimer's disease symptoms

The differentially expressed proteins described herein may used to test compounds for the ability to prevent or ameliorate Alzheimer's disease.

Such compounds may be tested in human subjects in clinical trials. Any compound which restores the expression of a differentially expressed protein or proteins towards the normal level may be of potential use in treating Alzheimer's disease, i.e. reducing Alzheimer's disease symptoms or slowing the progression of Alzheimer's disease.

With regard to intervention, any treatments that restore or partially restore marker protein expression to normal levels should be considered as candidates for Alzheimer's disease therapeutic intervention. Dosages of test agents may be determined by deriving dose-response curves, as discussed in Section 6 below.

Similarly, any treatments that can prevent the development of Alzheimer's disease or prevent progression to levels of more advanced Alzheimer's disease should be considered as candidates for the Alzheimer's disease therapeutic intervention.

In addition, animal models of Alzheimer's disease, such as those described above, may be used to identify compounds capable of treating Alzheimer's disease symptoms. Such animal models may be used as test substrates for the identification of drugs,

pharmaceuticals, therapies and interventions which may be effective in treating such disorders. The response of the animals to the exposure may be monitored by assessing the expression of the marker proteins and comparing it to that of wild-type mice.

Protein expression patterns may be utilised in conjunction with animal model systems to assess the ability of a compound to ameliorate Alzheimer's disease symptoms, or prevent the progression of Alzheimer's disease. For example, the expression pattern of one or more fingerprint proteins may form part of a fingerprint profile, which may then be used in such an assessment. Fingerprint profiles may be characterised for Alzheimer's disease states within the animal model systems. Subsequently, these known fingerprint profiles may be compared to ascertain the effect a test compound has to modify such fingerprint profiles, and to cause the profile to more closely resemble that of a more desirable fingerprint. For example, administration of a compound may cause the fingerprint profile of an Alzheimer's disease model system to more closely resemble the control system, or may prevent further changes in fingerprint profile. Administration of a compound may, alternatively, cause the fingerprint profile of a control system to begin to mimic an Alzheimer's disease state, which may, for example, be used in further characterising the compound of interest, or may be used in the generation of additional animal models.

4. Compounds and methods for treatment of Alzheimer's disease

Described below are methods and compositions whereby Alzheimer's disease symptoms may be ameliorated or the progression of Alzheimer's disease slowed or halted. It is possible that Alzheimer's disease symptoms may be brought about, at least in part, by an abnormal level of target protein, or by the presence of a target protein exhibiting an abnormal activity. As such, the reduction in the level and/or activity of such target protein would bring about the amelioration Alzheimer's disease symptoms. Techniques for the reduction of target protein gene expression levels or target protein activity levels are discussed in Section 4.1.

Alternatively, it is possible that Alzheimer's disease symptoms may be brought about, at least in part, by the absence or reduction of the level of target protein expression, or a reduction in the level of a target protein's activity. As such, an increase in the level of target protein gene expression and/or the activity of such proteins would bring about the amelioration Alzheimer's disease symptoms. Techniques for increasing target protein gene expression levels or target protein activity levels are discussed in Section 4.2.

4.1 Compounds that inhibit expression, synthesis or activity of target proteins

As discussed above, target proteins involved in Alzheimer's disease may cause such disorders via an increased level of target protein activity. A variety of techniques may be utilised to inhibit the expression,

synthesis, or activity of such target genes and/or proteins.

For example, compounds which exhibit inhibitory activity, may be used in accordance with the invention to prevent mild cognitive impairment or Alzheimer's disease symptoms. Such molecules may include, but are not limited to, peptides (such as, for example, peptides representing soluble extracellular portions of target protein transmembrane receptors), phosphopeptides, small organic or inorganic molecules, or antibodies (including, for example, polyclonal, monoclonal, humanised, anti-idiotypic, chimeric or single chain antibodies, and FAb, F(ab')₂ and FAb expression library fragments, and epitope-binding fragments thereof). Techniques for determination of effective doses and administration of such compounds are described below, in Section 6.1. Inhibitory antibody techniques are further described below, in Section 4.1.2.

Further, antisense, siRNA and ribozyme molecules, which inhibit expression of the target protein gene, may also be used in accordance with the invention to inhibit the aberrant target protein gene activity. Such techniques are described below, in Section 4.1.1; triple helix molecules may be utilised in inhibiting the aberrant target protein gene activity.

4.1.1 Inhibitory antisense, ribozyme and triple helix approaches

Antisense, ribozyme and triple helix molecules may be designed to reduce or inhibit either wild type, or if appropriate, mutant target protein gene activity.

Techniques for the production and use of such molecules are well known to those of skill in the art.

Antisense RNA and DNA molecules act to directly block the translation of mRNA by hybridising to targeted mRNA and preventing protein translation. With respect to antisense DNA, oligodeoxy-ribonucleotides derived from the translation initiation site, e.g. between the -10 and +10 regions of the target gene nucleotide sequence of interest, are preferred.

Ribozymes are enzymatic RNA molecules capable of catalysing the specific cleavage of RNA. (For a review, see Rossi, J., 1994, Current Biology 4: 469-471). The mechanism of ribozyme action involves sequence specific hybridisation of the ribozyme molecule to complementary target RNA, followed by an endonucleolytic cleavage. The composition of ribozyme molecules must include one or more sequences complementary to the target protein mRNA, and must include the well known catalytic sequence responsible for mRNA cleavage. For this sequence, see US Patent No:5,093,246. As such, within the scope of the invention are engineered hammerhead motif ribozyme molecules that specifically and efficiently catalyse endonucleolytic cleavage of RNA sequences encoding target proteins.

Specific ribozyme cleavage sites within any potential RNA target are initially identified by scanning the molecule of interest for ribozyme cleavage sites which include the following sequences, GUA, GUU and GUC. Once identified, short TNA sequences of between 15 and 20 ribonucleotides

corresponding to the region of the target protein gene, containing the cleavage site may be evaluated for predicted structural features, such as secondary structure, that may render the oligonucleotide sequence unsuitable. The suitability of candidate sequences may also be evaluated by testing their accessibility to hybridise with complementary oligonucleotides, using ribonuclease protection assays.

RNA interference (RNAi) is a process of sequence-specific, post-transcriptional gene silencing in animals and plants, initiated by double-stranded RNA (dsRNA) that is homologous in sequence to the silenced gene. RNAi is mediated by short double-stranded RNA molecules (small interfering RNAs or siRNAs). siRNAs may be introduced into a cell as short RNA oligonucleotides of 10-15bp, or as longer dsRNAs which are subsequently cleaved to produce siRNAs. The RNA may be introduced into the cell as RNA, or may be transcribed from a DNA or RNA vector.

siRNA molecules may be synthesized using standard solid or solution phase synthesis techniques which are known in the art. Alternatively, siRNA molecules or longer dsRNA molecules may be made recombinantly by transcription of a nucleic acid sequence, preferably contained within a vector as described below.

Another alternative is the expression of a short hairpin RNA molecule (shRNA) in the cell. shRNAs are more stable than synthetic siRNAs. A shRNA consists of short inverted repeats separated by a small loop sequence. One inverted repeat is complementary to the gene target. The

shRNA is then processed into an siRNA which degrades the target gene mRNA and suppresses expression. shRNAs can be produced within a cell by transfecting the cell with a DNA construct encoding the shRNA sequence under control of a RNA polymerase III promoter, such as the human H1 or 7SK promoter. Alternatively, the shRNA may be synthesised exogenously and introduced directly into the cell. Preferably, the shRNA sequence is between 40 and 100 bases in length, more preferably between 40 and 70 bases in length. The stem of the hairpin is preferably between 19 and 30 base pairs in length. The stem may contain G-U pairings to stabilise the hairpin structure.

Nucleic acid molecules to be used in triplex helix formation for the inhibition of transcription should be single stranded and composed of deoxynucleotides. The base composition of these oligonucleotides must be designed to promote triple helix formation via Hoogsteen base pairing rules, which generally require sizeable stretches of either purines or pyrimidines to be present on one strand of a duplex. Nucleotide sequences may be pyrimidine-based, which will result in TAT and CGC⁺ triplets across the three associated strands of the resulting triple helix. The pyrimidine-rich molecules provide base complementary to a purine-rich region of a single strand of the duplex in a parallel orientation to that strand. In addition, nucleic acid molecules may be chosen that are purine-rich, for example, containing a stretch of G residues. These molecules will form a triple helix with a DNA duplex that is rich in GC pairs, in which the majority of the purine residues are located

on a single strand of the targeted duplex, resulting in GGC triplets across the three strands in the triplex.

Alternatively, the potential sequences that can be targeted for triple helix formation may be increased by creating a so-called "switchback" nucleic acid molecule. Switchback molecules are synthesised in an alternating 5'-3', 3'-5' manner, such that they base pair with first one strand of a duplex and then the other, eliminating the necessity for a sizeable stretch of either purines or pyrimidines to be present on one strand of a duplex.

Anti-sense RNA and DNA, siRNAs, ribozyme and triple helix molecules of the invention may be prepared by any method known in the art for the synthesis of DNA and RNA molecules. They include techniques for chemically synthesising oligodeoxyribonucleotides and oligoribonucleotides well known in the art such as, for example, solid phase phosphoramidite chemical synthesis. Alternatively, RNA molecules may be generated by in vitro and in vivo transcription of DNA sequences encoding the antisense RNA molecule. Such DNA sequences may be incorporated into a wide variety of vectors, which incorporate suitable RNA polymerase promoters such as the T7 or SP6 polymerase promoters. Alternatively, antisense cDNA constructs that synthesise antisense RNA constitutively inducibly, depending on the promoter used, can be introduced stably into cell lines.

4.1.2 Antibodies for the inhibition of target protein

Antibodies that are both specific for target protein and interfere with its activity may be used to inhibit target protein function. Where desirable, antibodies specific for mutant target protein, which interferes with the activity of such mutant target product, may also be used. Such antibodies may be generated, using standard techniques described in Section 2. above, against the proteins themselves or against peptides corresponding to portions of the proteins. The antibodies include, but are not limited to, polyclonal, monoclonal, Fab fragments, single chain antibodies, chimeric antibodies, etc.

In instances where the target gene protein is intracellular and whole antibodies are used, internalising antibodies may be preferred. However, lipofectin or liposomes may be used to deliver the antibody or a fragment of the Fab region, which binds to the target protein epitope into cells. Where fragments of the antibody are used, the smallest inhibitory fragment, which binds to the target protein's binding domain, is preferred. For example, peptides having an amino acid sequence corresponding to the domain of the variable region of the antibody that binds to the target protein may be used. Such peptides may be synthesised chemically or produced via recombinant DNA technology using methods well known in the art (e.g. see Creighton, 1983, *supra*; and Sambrook et al, 1989, *supra*).

Alternatively, single chain neutralising antibodies, which bind to intracellular target protein epitopes, may also be administered. Such single chain antibodies may be administered, for example, by expressing nucleotide sequences encoding single-chain antibodies within the target cell populating by utilising, for example, techniques such as those described in Marasco et al (Marasco, W. et al, 1993, Proc. Natl. Acad. Sci. USA, 90: 7889-7893).

In instances where the target protein is extracellular, or is a transmembrane protein, any of the administration techniques described below, in Section 6, which are appropriate for peptide administration may be utilised to effectively administer inhibitory target protein antibodies to their site of action.

4.2 Methods for restoring target protein activity

Target proteins that cause Alzheimer's disease may be underexpressed in Alzheimer's disease disorder situations. Alternatively, the activity of target protein may be diminished, leading to the development of Alzheimer's disease symptoms. Described in this Section are methods whereby the level of target protein may be increased to levels wherein Alzheimer's disease symptoms are prevented or ameliorated. The level of target protein activity may be increased, for example, by either increasing the level of target protein present or by increasing the level of active target protein which is present.

For example, a target protein, at a level sufficient to ameliorate Alzheimer's disease symptoms may be administered to a patient exhibiting such symptoms. Any of the techniques discussed below, in Section 6, may be utilised for such administration. One of skill in the art will readily know how to determine the concentration of effective, non-toxic doses of the normal target protein, utilising techniques such as those described below.

Further, patients may be treated by gene replacement therapy. One or more copies of a normal target protein gene or a portion of the gene that directs the production of a normal target protein with target protein gene function, may be inserted into cells, using vectors which include, but are not limited to, adenovirus, adeno-associated virus, and retrovirus vectors, in addition to other particles that introduce DNA into cells, such as liposomes. Additionally, techniques such as those described above may be utilised for the introduction of normal target protein gene sequences into human cells.

Cells, preferably autologous cells, containing normal target protein gene sequences may then be introduced or reintroduced into the patient at positions which allow for the prevention or amelioration of Alzheimer's disease symptoms. Such cell replacement techniques may be preferred, for example, when the target protein is a secreted, extracellular protein.

Additionally, antibodies may be administered which specifically bind to a target protein and by binding,

serve to, either directly or indirectly, activate the target protein function. Such antibodies can include, but are not limited to, polyclonal, monoclonal, FAb fragments, single chain antibodies, chimeric antibodies and the like. The antibodies may be generated using standard techniques such as those described above, in Section 2.3, and may be generated against the protein themselves or against proteins corresponding to portions of the proteins. The antibodies may be administered, for example, according to the techniques described above.

5. Pharmaceutical preparations and methods of administration

The identified compounds, nucleic acid molecules and cells that affect target protein expression, synthesis and/or activity can be administered to a patient at therapeutically effective doses to prevent or to treat or to ameliorate Alzheimer's disease. A therapeutically effective dose refers to that amount of the compound sufficient to result in amelioration of symptoms Alzheimer's disease, or alternatively, to that amount of a nucleic acid molecule sufficient to express a concentration of protein which results in the amelioration of such symptoms.

5.1 Effective dose

Toxicity and therapeutic efficacy of such compounds can be determined by standard pharmaceutical procedures in cell cultures or experimental animals, e.g. for determining by ED₅₀ (the dose therapeutically effective in 50% of the population) and by determining the ED₅₀ of any side-effects (toxicity - TD₅₀). The dose ratio between

toxic and therapeutic effects is the therapeutic index and it can be expressed as the ratio TD_{50}/ED_{50} . Compounds, which exhibit large therapeutic indices, are preferred. While compounds that exhibit toxic side effects may be used, care should be taken to design a delivery system that targets such compounds to the site of affected tissue in order to minimise potential damage to uninfected cells and, thereby, reduce side effects.

The data obtained from the animal studies can be used in formulating a range of dosage for use in humans. The dosage of such compounds lies preferably within a range of circulating concentrations that include the ED_{50} with little or no toxicity. The dosage may vary within this range depending upon the dosage form employed and the route of administration utilised.

5.2 Formulations and use

Pharmaceutical compositions for use in accordance with the present invention may be formulated in conventional manner using one or more physiologically acceptable carriers or excipients.

Thus, the compounds and their physiologically acceptable salts and solvates may be formulated for administration by inhalation or insufflation (either through the mouth or the nose) or oral, buccal, parenteral and rectal administration.

For oral administration, the pharmaceutical compositions may take the form of, for example, tablets or capsules prepared by conventional means with pharmaceutically

acceptable excipients such as binding agents (e.g. pre-gelatinised maize starch, polyvinylpyrrolidone or hydroxypropyl methyl-cellulose); fillers (e.g. lactose, microcrystalline cellulose or calcium hydrogen phosphate); lubricants (e.g. magnesium stearate, talc or silica); disintegrants (e.g. potato starch or sodium starch glycollate); or wetting agents (e.g. sodium lauryl sulphate). The tablets may be coated by methods well known in the art. Liquid preparations for oral administration may take the form of, for example, solutions, syrups or suspensions, or they may be presented as a dry product for constitution with water or other suitable vehicle before use. Such liquid preparations may be prepared by conventional means with pharmaceutically acceptable additives such as suspending agents (e.g. sorbitol syrup, cellulose derivatives or hydrogenated edible fats); emulsifying agents (e.g. lecithin or acacia); and preservatives (e.g. methyl or propyl-p-hydroxybenzoates or sorbic acid). The preparations may also contain buffer salts, flavouring, colouring and sweetening agents as appropriate.

Preparations for oral administration may be suitably formulated to give controlled release of the active compound.

For buccal administration the compositions may take the form of tablets or lozenges formulated in conventional manner.

For administration by inhalation, the compounds for use according to the present invention are conveniently

delivered in the form of an aerosol spray presentation from pressurised packs or a nebuliser, with the use of a suitable propellant, e.g. dichlorodifluoromethane, trichlorofluoromethane, dichloro-tetrafluoroethane, carbon dioxide or other suitable gas. In the case of a pressurised aerosol the dosage unit may be determined by providing a valve to deliver a metered amount. Capsules and cartridges of, e.g. gelatin, for use in an inhaler or insufflator may be formulated containing a powder mix of the compound and a suitable powder base such as lactose or starch.

The compounds may be formulated for parenteral administration by injection, e.g. by bolus injection or continuous infusion. Formulations for injection may be presented in unit dosage form, e.g. in ampoules or in multi-dose containers, with an added preservative. The compositions may take such forms as suspensions, solutions or emulsions in oily or aqueous vehicles, and may contain formulatory agents such as suspending, stabilising and/or dispersing agents. Alternatively, the active ingredient may be in powder form for constitution with a suitable vehicle, e.g. sterile pyrogen-free water, before use.

The compounds may also be formulated in rectal compositions such as suppositories or retention enemas, e.g. containing conventional suppository bases such as cocoa butter or other glycerides.

In addition to the formulations described previously, the compounds may also be formulated as a depot preparation.

Such long acting formulations may be administered by implantation, for example, subcutaneously or intramuscularly) or by intramuscular injection. Thus, for example, the compounds may be formulated with suitable polymeric or hydrophobic materials (for example, as an emulsion in an acceptable oil) or ion exchange resins, or as sparingly soluble derivatives, for example, as a sparingly soluble salt.

The compositions may, if desired, be presented in a pack or dispenser device, which may contain one or more unit dosage forms containing the active ingredient. The pack may, for example, comprise metal or plastic foil, such as blister pack. The pack or dispenser device may be accompanied by instructions for administration.

6. Diagnosis of Alzheimer's disease

A variety of methods may be employed for the diagnosis of Alzheimer's disease, monitoring progression of mild cognitive impairment and Alzheimer's disease, the predisposition to Alzheimer's disease, and for monitoring the efficacy of any Alzheimer's disease compounds during, for example, clinical trials and for monitoring patients undergoing clinical evaluation for the treatment of Alzheimer's disease. The differentially expressed and fingerprint proteins can also be used to define the nature or degree of Alzheimer's disease to aid in the identification and/or selection of treatments for the disorder.

Alzheimer's disease is characterised by a progressive, insidious onset, two or more deficits in cognitive

function, and the absence of any other illnesses that could account for the dementia

In addition to memory loss, there may be disorientation, poor attention span, and language impairment. There is likely to be a decline in the activity of daily living, and possibly also impaired perception and personality changes. Behavioural symptoms include delusions, aggression, agitation, anger, wandering, hallucinations, and sleep disturbance.

A simple test assessing orientation, registration, calculations and attention, recall, language, and visual-spatial function may be used for an initial diagnosis.

Structural imaging by standard CT or MRI may also be used. Typically a non-contrast head CT scan suffices, but MRI is preferred for those who have hypertension or diabetes, who are at risk for cerebral vascular disease.

Alzheimer's disease may be confirmed histologically by autopsy or brain biopsy showing neurofibrillary tangles and senile plaques.

Identifying individuals at risk from Alzheimer's disease may involve diagnosis of mild cognitive impairment (MCI). (MCI) may be a transitional state between normal aging and dementia. There are different types of MCI. There may be cognitive impairment in multiple areas of cognitive function, in addition to memory. In some cases, memory is normal but some other domain of cognitive function is abnormal.

Amnesic MCI appears to be a risk state for the development of Alzheimer's disease. Amnesic impairment is defined by subjective memory complaints. These patients have poor memory performance for their age and education on formal testing when compared to age-matched peers. General cognitive functions and the ability to perform the activities of daily living should be entirely normal. The amnesic type of MCI is associated with hippocampal atrophy, neurofibrillary tangles in the medial temporal lobes, and elevated levels of Tau in the cerebrospinal fluid (CSF).

Methods for diagnosing Alzheimer's disease or predisposition to Alzheimer's disease may also, for example, utilise reagents such as the differentially expressed and fingerprint proteins described above, and antibodies directed against differentially expressed, as described above. Specifically, such reagents may be used for the detection of either an over- or an under-abundance of target protein relative to the normal state.

The methods described herein may be performed, for example, by utilising pre-packaged diagnostic kits comprising at least one specific differentially expressed/finger print protein or anti-differentially expressed/fingerprint protein antibody reagent described herein, which may be conveniently used, e.g. in clinical settings, to diagnose patients exhibiting Alzheimer's disease symptoms.

Any cell type, tissue or body fluid in which the fingerprint protein is expressed may be utilised in the diagnostics described herein. Examples of suitable samples types include cell samples, tissue samples, and fluid samples such as blood, urine, serum, saliva, cerebrospinal fluid or plasma.

Among the methods which can be utilised herein, are methods for monitoring the efficacy of compounds in clinical trials for the treatment of Alzheimer's disease. Such compounds can, for example, be compounds such as those described above, in Section 4. Such a method comprises detecting, in a patient sample, a protein, which is differentially expressed in the Alzheimer's disease state relative to its expression in a normal state.

During clinical trials, for example, the expression of a single differentially expressed protein, or alternatively, a fingerprint pattern of a cell involved in Alzheimer's disease can be determined in the presence or absence of the compound being tested. The efficacy of the compound can be followed by comparing the expression data obtained to the corresponding known expression patterns in a normal state. Compounds exhibiting efficacy are those which alter the protein expression and/or the fingerprint pattern to more closely resemble that of the normal state, or which stabilise protein expression and/or the fingerprint pattern i.e. prevent progression of the disease.

The detection of the protein differentially expressed in the Alzheimer's disease state relative to their expression in a normal state can also be used for monitoring the efficacy of potential compounds for the treatment of Alzheimer's disease during clinical trials. During clinical trials, for example, the level and/or activity of the differentially expressed protein can be determined in relevant cells and/or tissues and/or body fluids in the presence or absence of the compound being tested. The efficacy of the compound can be followed by comparing the protein level and/or activity data obtained to the corresponding known levels/activities for the cells and/or tissues and/or body fluids in a normal state. Compounds exhibiting efficacy are those which alter the pattern of the cell and/or tissue sample and/or body fluid from an Alzheimer's disease subject to more closely resemble that of the normal state or which stabilise the pattern i.e. prevent progression of the disease.

Experimental

Subjects

The study population is derived from a large, longitudinally assessed, community based population of people with AD (NINCDS-ADRDA probable), other dementias and normal elderly persons. Samples are available on over 1000 subjects, all whom have detailed clinical assessment. Clinical research data includes systematic diagnostic, cognitive and behavioural assessments. Approximately 50ml blood (4x10ml in BD vacutainer K3E 15% tubes and 1x10ml in exetainer) is drawn from each subject. Subjects have had no food or fluid intake for

more than 2 hours prior to collection. One BD vacutainer K3E (plasma) and exetainer (serum) is used for proteomics study. The serum/plasma samples collected for proteomics are spun at 3000rpm for 8min within 2h of collection.

Analysis

Serum/plasma samples were lysed and rehydrated in a 2D lysis buffer consisting of 8M Urea, 2% w/v CHAPS, 0.5% IPG Pharmalyte (pH 3-10; Amersham Biotech, UK. The lysed samples were then subjected to isoelectrofocusing using 18cm 3-10NL Immobiline pH gradient strips. IPG electrofocusing of the rehydrated strips was carried out for 16h using the following protocol: S1 500V step-n-hold (s/h; i.e. the electric current applied to the strip is gradually increased in steps holding at particular settings for the times indicated) for 1h; S2 500V s/h for 2h; S3 1000V gradient (G) for 1h; S4 1000V s/h for 2h; S5 8000V G for 2h and a final step S6 8000V s/h for 8h with the IPGphorTM.

Electrofocused IPG strips were then equilibrated in a SDS equilibration buffer (50mM Tris-HCl pH8.8, 6M urea, 30%(v/v) glycerol, 2% SDS, and trace amount of bromophenol blue) with 10mg/ml dithiothreitol (DTT) for 20min, followed by 20min step with 25mg/ml Iodoacetamide. The equilibrated strips were then separated on a 10% acrylamide second dimension electrophoresis gel using the Ettan Dalt II system.

Following the electrophoresis the gels were placed in separate staining boxes and fixed using 40% ethanol/10% acetic acid for 1h at room temperature and then stained

according to Hochstrasse protocol (Table 1). Gel analysis was performed using the Melanie 3 software and Mann and Whitney rank sum test and False Discovery Rate statistical analysis was carried out to compare subject groups.

Table 1 - Hochstrasse staining protocol

Staining step	Time	Sample preparation In-gel reduction, alkylation and digestion (with trypsin) were performed prior to subsequent analysis by mass spectrometry. Cysteine
Fix 40% ethanol/10% acetic acid	1 h	
Soak in 5% ethanol/5% acetic acid	3hr or overnight	
Wash in water	5min	
Soak in 0.5M Sodium acetate, 1% gluteraldehyde	1.5 h	
Wash	4x15min	
Soak in 0.05% Naphthalene sulphonic acid	2x30min	
Rinse in water	4x15min	
Silver stain (12g silver, 20ml ammonium hydroxide and 3ml 10M sodium hydroxide)	25min	
Wash	4x4min	
Develop (0.005% citric acid and 0.1% formaldehyde)	As required	
Stop solution (5% tris and 2% acetic acid)	1-2h	
Storage solution (35% ethanol and 5% glycerol)		

residues were reduced with DTT and derivatized by treatment with iodoacetamide to form stable carbamidomethyl (CAM) derivatives. Trypsin digestion was carried out overnight at room temperature after an initial 1hr incubation at 37°C.

MALDI-TOF Mass spectrometry

The digested sample (3 µl) was desalted and concentrated using ZipTipC18 microtips (Millipore). Peptides were

eluted in 4 μ l 50% acetonitrile/ 0.1% trifluoroacetic acid. 0.5 μ l was then loaded onto a target plate with 0.5 μ l matrix (α -Cyano-4-hydroxy-cinnamic acid). Peptide mass fingerprints were acquired using a Voyager De-Pro, MALDI-TOF mass spectrometer (Applied Biosystems). The mass spectra were acquired in reflectron mode with delayed extraction. An autolytic tryptic peptide of mass 2163.0569 Da was then used to lock mass the acquired spectra, to achieve a mass accuracy of better than 30ppm.

LC/MS/MS

Peptides were extracted from the gel pieces by a series of acetonitrile and aqueous washes. The extract was pooled with the initial supernatant and lyophilised. Each sample was then resuspended in 6 μ l of 50mM ammonium bicarbonate and analysed by LC/MS/MS. Chromatographic separations were performed using an Ultimate LC system (Dionex, UK). Peptides were resolved by reverse phase chromatography on a 75 μ m C18 PepMap column. A gradient of acetonitrile in 0.05% formic acid was delivered to elute the peptides at a flow rate of 200 nl/min. Peptides were ionised by electrospray ionisation using a Z-spray source fitted to a QTOFmicro (Waters Corporation). The instrument was set to run in automated switching mode, selecting precursor ions based on their intensity, for sequencing by collision-induced fragmentation. The MS/MS analyses were conducted using collision energy profiles that were chosen based on the m/z and the charge state of the peptide.

Results

Analysis of all control group (n=50) and case group (n=50) 2D gel images and subjecting them to statistical analysis. A total of 16 protein spots show a significant result ($p < 0.05$) (see Fig 6).

The results shown in Fig 6 are unambiguous matches as they are based on exact matching of multiple MS/MS spectra. The sequence of selected proteins showing the peptide coverage obtained is given in Figs 8 to 10.

Class prediction using peptide fingerprinting

A class prediction analysis was performed in order to determine whether the pattern of peptide spots on 2DGE could predict caseness as determined clinically. Support Vector Machines (SVM), a supervised machine learning algorithm for prediction of class set in a group based upon a training set of data¹³, was used. SVM is most typically used in microarray analyses. However the statistical challenges are similar for proteomics and SVM has previously been used as a class prediction model for various proteomic studies^{14,15}. Using GeneSpring (Silicon Genetics) the original 25 cases and 25 controls were designated as a training set and then the replication 25 cases and 25 controls designated as a test set. All identified proteins were used as possible identifiers and with the parameters Polynomial Dot Product Order 1 and Diagonal Scaling Factor 1; 34 of the 50 test-samples were correctly identified as being either cases or controls. Sensitivity was 56% and specificity 80% using SVM analysis of 2DGE data alone.

Identification of peptides that differentiate between cases and controls

The normalised spot optical density in both the initial set of cases and controls and the replication set was compared. Mean differences between patients and controls at each spot were compared using the Wilcoxon rank-sum (Mann-Whitney) tests. The p-values for the null hypothesis of no mean differences were saved, sorted by increasing value and ranked. A false discovery rate index (FDR) was computed as the ratio of the rank number and the theoretical probability (which is the rank number divided by the total number of spots). Fifteen spots were identified to have a FDR of less than .50. These were then identified using LC-MS/MS (Fig 6).

Correlation of peptide spots with clinical parameters

Although the cases and controls were similarly aged it was possible that the observed peptide or spot differences were due to an association with age, gender or APOE genotype. A correlation analysis was thus performed for the 15 spots that differed between cases and controls in all 100 subjects with age, gender and APOE genotype. Data was first scaled to unit variance so as to standardise the scales upon which the variables were compared (i.e. each value was divided by the standard deviation of all the values for that particular variable). The Pearson correlation coefficient was then calculated. Cases with missing values were excluded pairwise. There were no strong correlations of any spot

with age, gender or APOE. Two spots weakly correlated with age, two with gender and one with APOE genotype.

An ideal biomarker would not only be different between cases and controls but would be a marker of disease progression. The 15 spots showing case-control differences in all 50 cases were thus correlated with duration of dementia and severity as measured by MMSE and GDS. Two spots correlated moderately and significantly with measures of disease progression and global dementia severity ($r^2 = 0.29$ with spot 177) and duration of disease ($r^2 = -0.29$ with spot 166). Thus, one peptide - an Ig lambda chain C region (spot 177) correlates with global dementia severity. The other marker of disease progression examined, duration, shows a negative correlation with albumin (Spot 165).

Pre-depletion analysis

In these experiments, human plasma samples were depleted to remove the 6 most abundant proteins before the 2D gel electrophoresis step.

Methods

60 human plasma samples (30 Controls and 30 disease subjects) were depleted using a removal column from Agilent. The samples were separated by 2D electrophoresis (pH 3-10NL, 10% SDS-PAGE, 75µg protein load). Gels were silver-stained, scanned (8bit, 200 dpi) and quantitatively analysed with Progenesis. To pick gel plugs from preparative gels, several control samples were mixed together and 3 gels run (2 gels with 205µg protein

load and 1 gel with 350µg protein load). The same strategy was used with disease samples to run preparative gels. Protein spots were then de-stained, trypsinated and polypeptides were spotted onto MALDI target with Spot handling workstations (GE Amersham Biosciences). Peptide profiles generated were analysed with Ms-Fit programme in combination with the Swiss-Prot database.

Results

Gel images of proteins extracted from control and disease samples were analysed with Progenesis (v2005). Each group (Control and Disease) were based on 29 analytical gels. Spot detection, matching were performed with Progenesis, then the spot data were exported to Excel and a macro developed in-house was used to calculate coefficient of variation (CV%), T-Test and Regulation facto or change.

11 spots were selected for analysis based on the following selection criteria: spots have to be found within at least 60% of gels, 2-fold up/down regulation and $p \text{ value} < 0.005$. Fig 11 displays the location of these 11 spots in the reference gel. This image corresponds to the 2D profile of proteins extracted from a control sample. The normalised volumes of the 11 spots detected in gels was analysed and is given in Fig 12.

All protein spots were picked from 3 to 6 different preparative gels and submitted to MS analyses. All protein spots were successfully identified as shown in Fig 12. In the down regulated spots, we found two spots of alpha-2-macroglobulin precursor (174; 178), one spot

of inter-alpha-trypsin inhibitor heavy chain H4 precursor (232), one spot of a mix of complement C3 precursor with clusterin precursor (712) as indicated in grey in Fig 12 and one spot of complement C3 precursor (713). In the up regulated spots, we found two spots of complement C4 precursor (652; 832), one spot of actin (675) and three spots of haptoglobin precursor (702; 703; 706).

To estimate the coverage of proteins identified and to discriminate the different chains or isoforms, for each spot, a common list of peptide masses was established. This list regroups all peptide masses matched corresponding to the same spot picked in 3 to 6 preparative gels. The amino acids belonging to the peptides matched are underlined in Figs 13 to 19.

Discussion

The 11 spots analysed identified 7 regulated proteins between control- and disease samples, namely alpha-2-macroglobulin, inter-alpha-trypsin inhibitor heavy chain H4, complement C3, complement C4, actin cytoplasmic and haptoglobin.

Alpha-2-macroglobulin protein is able to inhibit all four classes of proteinases by a unique "trapping" mechanism. The observed molecular weight of the gel spots (~100 kDa, Fig. 11, spots 174 ; 178), matched by PMF, cover mainly the N-terminus of the protein (Fig. 14). The protein identified may thus correspond to a fragment of the full-length sequence of alpha-2-macroglobulin. As spots identified as alpha-2-macroglobulin belong to the same chain of spots (Fig. 11), it is possible that the

difference between the two spots may be due to a post-translation modification.

There are two isoforms of inter-alpha-trypsin inhibitor heavy chain H4. Isoform 1 has 930 amino-acids and isoform 2 has 914 amino-acids. This protein is cleaved by plasma kallikrein to yield 100 kDa and a 35 kDa fragments. The resulting 100 kDa fragment is further converted to a 70 kDa fragment. The masses matched by PMF cover the sequence up to amino acid (aa) 688. This sequence corresponds to isoform 1 and may include the 70 kDa fragment and a short potentially active peptide. In this case, there is good agreement between the theoretical molecular weight and pI (74 kDa and 6.04 respectively) and the observed ones from the gel spot (see Fig. 11, spot 232).

Complement C3 precursor plays a central role in the activation of the complement system. This protein contains two chains (alpha and beta). We identified peptide masses covering the sequence from aa 714 to aa 1360 (Fig. 15), which corresponds to the alpha chain of complement C3. The theoretical molecular weight and pI of the alpha chain (115 kDa and 5.55 respectively) are not in agreement with the observed ones from the gel spots (see Fig. 11, spots 712, 713). The alpha chain is processed into different fragments. It appears that a temporary peptide appearing during the activation of complement system. As spots identified as complement C3 belong to the same chain of spots (Fig. 11), it is possible that the difference is due to a post-translational modification.

Complement C4 plays a central role in the activation of the classical pathway of the complement system. This protein contains three chains (alpha, beta and gamma). We identified peptide masses covering the alpha and beta chains for spot 832 and only alpha chain for spot 652 (Fig. 17). The theoretical molecular weights and pIs of these chains differ from the observed ones from the gel spots (see Fig. 11, spots 652 ; 832). As for complement C3, clusterin precursor protein contains two chains (alpha and beta). We identified peptide masses covering the alpha and beta chains (Fig. 16). The theoretical molecular weight and pI of clusterin (50 kDa and 5.89 respectively) are in agreement with those from the gel (Fig. 11, spot 712). It appears the full-length protein was identified.

Surface enhanced laser desorption ionisation time of flight mass spectrometry [SELDI-TOF-MS].

SELDI-TOF-MS and ProteinChip technology were combined to identify protein peaks differing between Alzheimer's and control subjects, followed by extraction of material from the chips to allow further characterisation of the material and identification of the components present.

METHOD (SELDI analysis)

The SELDI analysis comprises of a comparison of AD cases and control samples and data has been obtained for both a set of individual samples as well as a pooled set of samples. In each case spectral profiles of sera from control and AD cases were compared.

A). Analysis of a set of Individuals

Control and AD sera from individuals were run on Q10-SAX2chips:

n = 4 control

n = 4 AD

Serum samples were prepared fresh by diluting 20µl serum with 30µl SELDI lysis buffer. Five microlitres of sample were spotted onto each spot as necessary.

The chips were processed using the following protocol:

Chip Preparation

A hydrophobic ring is drawn around each spot using a PAP pen and the PAP allowed to dry thoroughly by placing chip on the SELDI machine for up to 25 minutes.

Sample Preparation

Serum diluted in SELDI lysis buffer using a 40:60 ratio (40 µL serum + 60 µL lysis buffer). Typically, this dilution will render the sample at a 20 mg/mL to 30 mg/mL concentration. Therefore, using a 5 µL aliquot of the lysis buffer sample will enable between 100 µg to 150 µg protein to be loaded on each spot.

Samples are vortexed and incubated on ice until ready to use, then briefly centrifuged samples immediately before use (30 secs, 14, 000 rpm).

Chip Equilibration

The chip is placed in a 15 mL Falcon tube and 10 - 15 mL 100 mM Tris buffer pH 9 at room temperature added, then

mixed on a rotary mixer for 5 minutes. The procedure is repeated twice.

Sample Application

After the last equilibration step, the chip is removed and dried carefully with soft tissue. 5 μ L of sample is pipetted onto each spot, the chip is placed in a sealed humidity chamber and placed on a shaker for 30 minutes.

Chip Washing

After incubation, the sample is carefully removed from each spot and the chip replaced in the Falcon tube. 10 - 15 mL 100 mM Tris buffer pH 9 is added, and the Falcon tube mixed on a rotary mixer and for 5 minutes. This is repeat four more times, then the chip washed twice in double distilled water.

Chip Drying

After the last wash step, the chip is removed and dried carefully with soft tissue, then left to air-dry at room temperature for 25 minutes.

SPA Application

2 x 0.6 μ L saturated SPA matrix (freshly made) is pipette onto each spot. The first application is allowed to dry before applying the second 0.6 μ L aliquot. The SPA is then left to dry for up to 10 minutes on the SELDI machine.

The chips are then read on the SELDI machine.

The following criteria were applied for data analysis:

Clustering criteria: 5 s/n; 100 % spectra; 0.3 % mass; 2 s/n; add est. peaks.

Normalisation: Total ion count between 3,000 and 30,000 Da only.

RESULTS

Spot to spot reproducibility between loadings of the same sample was very good. Good correlation was achieved. Patient to patient variability in both control and dementia groups was observed. This may be due to differences in protein amount as well as idiosyncratic differences. Using very stringent clustering, 3 peaks were found to be statistically significant ($p = 0.05$) and these were visually verified to check validity. The three peaks of interest (see Figs 1-3) are as follows:

Mr 6,430 Da abundance in AD	1.62 fold increase in $p = 0.027$
Mr 14,640 Da abundance in AD	2.29 fold increase in $p = 0.036$
Mr 27,147 Da abundance in AD	2.82 fold increase in $p = 0.004574$

B) Analysis of Pooled sets

A set of pooled samples were analysed using exactly the same methods and criteria as described above. Here, however, we analysed 3 pooled AD samples versus 3 pooled controls where each pool contains serum from at least 25 individuals. In this manner we have encompassed samples

from over 75 individuals with AD and compared them against a control cohort representing 75 number of individuals. Pooled groups are described as: AD Pool 1, 2 and 3 comprising of 25, 25 and 25 unique individuals respectively. Similarly, the pooled controls are described as: Control Pool 1, 2 and 3 comprising of 25, 25 and 25 unique individuals respectively.

RESULTS

Using very stringent clustering, 1 peak was found to be statistically significant ($p = 0.05$) and this was visually verified to check validity.

The peak of interest (see Fig 4) is as follows:

Mr 14,646 Da	1.72 fold increase in
abundance in AD	$p = 0.037$

SELDI Peak identification strategy

The differentially expressed proteins identified by SELDI analysis were further analysed by SDS-PAGE. Bands corresponding to the MW of differentially expressed proteins were excised for analysis by mass spectroscopy.

Material was extracted from chips Q10 854 & 855 ("individual" samples) by boiling for 10 minutes in Laemmli buffer and control and disease spots were pooled into separate Eppendorf tubes. Extracted material was separated using SDS-PAGE (18 %, tris-glycine, Novex) and the gel was initially stained with Colloidal Coomassie Blue (CCB) but no bands were visualised. Subsequently the same gel was re-stained using modified (MS-compatible) silver stain (Fig 5).

Six bands, between 11 and 20 kDa, were visualised and these were excised from the 1st control lane for analysis by LC/MS/MS as described above.

Identified proteins are shown in Fig 7.

Further analysis of identified proteins

Apolipoprotein A-IV

Sequence coverage obtained for apolipoprotein A-IV (P06727) is shown in Fig 9 for the 14.6kDa band isolated on the Q10 SAX2 SELDI chip

The molecular weight of the biomarker of interest observed within the SELDI profiling experiments was determined to be 14640 +/- 6 Da. The 14.6kDa species is thought to be a fragment of ApoA-IV based on the facts that the intact protein should be observed at higher mass (45kDa) and that the peptides observed in the LC/MS/MS analysis only represented the C-terminal region of the protein. The observed molecular weight is in good agreement with the average molecular weight of 14636 Da predicted for residues 270 - 396 of the sequence defined for apolipoprotein A-IV within the Swiss Prot database entry P06727.

Both authentic full length apolipoprotein A-IV and a C-terminal fragment of apolipoprotein A-IV comprising of residues 270 - 396 may thus represent serum biomarkers of Alzheimer's disease.

Complement C4 precursor

Sequence coverage obtained for Complement C4 precursor (P01028) in 2DE spot 164 is shown in Fig 10. Spot 164 was identified on the basis of several peptides indicated in underlined bold and this defines the protein in Spot 164 as a C-Terminal fragment extending from residues 1466 - 1744..

Quantitative Protein Sequence Tag (qPST) analysis

10 disease samples and 10 control samples were individually immunodepleted for the 6 most highly abundant proteins. 2 pools consisting of either the disease or the control samples were generated and applied to the qPST procedure (precleavage with CNBr, labelling with dimethylglycine, trypsination and fractionation by strong cation exchange). The obtained SCX fractions were analysed by LC-MS and LC-MSMS using the QTOF-II instrument following the standard approach (LC-MS and LC-MSMS by three different data acquisition methods).

Results*Identification of proteins*

As stated above, three different MSMS acquisition strategies were employed:

1. Data Dependent Analysis to obtain as many as possible peptide ID's (1 mass window).
2. Data acquisition by an 'include list' containing regulated pairs, ie peptides whose intensity varied between disease and control samples (regulation criteria: ≥ 2 / $\leq 0,5$)
3. Data acquisition by an 'include list' containing non-paired MS-signals.

Taking all results from these three approaches into account and correcting them for redundancy, 88 protein IDs were obtained.

Directed searching for regulated proteins by include list (pairs with a regulation ≥ 2.0 / ≤ 0.5) MSMS strategy and crossmatching:

8 peptides were identified which could be crossmatched to regulations. These 8 peptides represent five proteins (the peptide grouping to obtain protein ID's was achieved by the ProteinProphet algorithm).

The ID's of these five proteins are shown in Figure 8.

The 2 peptides which represent protein 1 also occur in Ig alpha-1 chain C region, so that the protein ID's 1 and 2 in fact represent one ID (Ig alpha-1 chain C region).

The hypothetical protein DKFZP686C02220 is a unique one (in fact, one peptide is unique, the second one can occur in several proteins). This protein has typical signatures of immunoglobulins (regarding InterPro entries), and the second peptide also occurs in Ig alpha-2 chain C region.

The proteins 4 and 5 represent one protein ID (haptoglobin precursor) because both peptides occur also in haptoglobin precursor, but the corresponding peptides were grouped as individual proteins by the algorithm used.

Validation of APO-AIV fragments using Western blotting

Western blotting has been undertaken to confirm that the 14.6 kDa species was a fragment of APO-AIV.

Plasma samples were diluted 1:10 with double distilled water and assayed using a Bradford dye-binding method (diluted samples permit handling of suitably sized aliquot volumes).

SDS-PAGE was carried out using 20 µg sample per lane (2 µg if sample is a denatured primary or secondary antibody) on 16 % acrylamide gels, 1.5 mm thick, 10 wells (NOVEX) for 1 hr 80 V; 1 ½ hrs 125 V. This was followed by Western Blotting onto nitrocellulose membrane at 50 V

for 1 ½ hrs. The blots were probed with the following antibodies:

Anti-ApoA-IV (N-terminal specific), Santa Cruz Biotechnology, Inc.

Anti-ApoA-IV (C-terminal specific), Santa Cruz Biotechnology, Inc.

Both antibodies are affinity purified goat polyclonals raised against a peptide mapping near the amino (N-terminal) or carboxy (C-terminal) terminus of ApoA-IV of human origin. These antibodies were chosen since probing for the N- and C-terminals should increase the chance of detection of the ApoA-IV protein and/or fragments.

Several bands were found that appear to be ApoA-IV specific and also discriminatory for AD. These bands do not appear in the secondary antibody-only control blot for control or AD samples.

Bands 3-6 which are observed in the 10-16 kDa region are discriminatory for AD bands 3 - 6, but also appear to align with bands in the denatured ApoA-IV antibody lanes. It has also been observed that bands 3 - 6 are much stronger on blots where the N-terminal specific anti-ApoA-IV antibody has been used.

Two other key bands are observed. Band 1 is observed at approximately 45kDa and appears to correspond to the full length mature APO-AIV protein. Band 2 is observed at approximately 28kDa and appears to be an N-terminal fragment of APO-AIV.

Complement factor H validation.Methods*Sample dilution*

Plasma samples were diluted to 1 in 8 in Phosphate buffered saline (PBS). An equal volume of Laemmli 2x sample buffer was added and then boiled for 10min until use.

Western blot

SDS gel electrophoresis was performed using the Fisher Scientific 36 well, 1.5mm gels (all solutions were purchased from National Diagnostics). Samples were separated on a 10% resolving gel with a 4% stacking gel (all solutions were purchased from National Diagnostics). Samples (20 μ l) were separated initially for 30min at 110V and then for 60min at 150V until the dye front just began to enter the running buffer.

The gel was transferred to PVDF (Amersham Biosciences) using a Semi-dry transblot (Bio-Rad) for 45min at 15V. The membrane was then blocked in 5% milk made in PBS-Tween and probed with Complement factor H primary antibody (Abcam, UK) overnight at 4°C. The bands were detected with a chemiluminescence Western detection kit (ECL+, Amersham Biosciences) and the membranes were scanned using Storm fluorescence scanner (Amersham Biosciences).

An immunoreactive band was observed at 139kDa (CfH) and the optical density was quantified using the Image Quant (Amersham Biosciences) software. Analysis was by non-parametric Mann-Whitney using the SPSS package.

Results

Western blot data was acquired from plasma from 128 people with NINCDS-ADRDA probable AD and 78 normal healthy elderly controls. Cases with AD had a 32% increase in CFH (Mann-Whitney; table 2)

Table 2

Diagnosis	Number	Mean CFH	SD	SEM
Controls	128	65.6	65.5	5.8
Probable AD	78	96.0	96.8	11.0

There was a gender difference with a relatively higher CFH value in females overall relative to males ($p=0.05$). However CFH was higher in cases with AD relative to controls even when considering genders separately ($p<0.01$; table 3)

Table 3

Females only	Number	Mean CFH	SEM
Controls	78	73.0	8.9
Probable AD	64	102.7	13.0
Total	142	86.4	7.7

A receiver operator curve (ROC) analysis showed that CFH performs better than chance as a diagnostic test.

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Claims

1. An *in vitro* method of diagnosing Alzheimer's disease in a subject, the method comprising:
 - (a) providing a body fluid sample from said subject;
 - (b) detecting the expression of alpha-2-macroglobulin precursor, clusterin precursor and fibrinogen gamma chain precursor in the body fluid sample, wherein alpha-2-macroglobulin precursor, clusterin precursor and fibrinogen gamma chain precursor are detected by mass spectrometry; and
 - (c) diagnosing the subject as having Alzheimer's disease when the expression of alpha-2-macroglobulin precursor, clusterin precursor and fibrinogen gamma chain precursor in the body fluid sample differs from the expression of alpha-2-macroglobulin precursor, clusterin precursor and fibrinogen gamma chain precursor in a control sample representative of normal subjects.
2. The method of claim 1, wherein the method further comprises detecting the expression of complement C3 precursor in the body fluid sample, wherein complement C3 precursor is detected by mass spectrometry, and diagnosing the subject as having Alzheimer's disease when the expression of complement C3 precursor in the body fluid sample differs from the expression of complement C3 precursor in a control sample representative of normal subjects.
3. The method of claim 1 or 2, wherein the method further comprises detecting the expression of one or more proteins selected from the group consisting of transthyretin, complement C4 precursor, complement factor H, and haptoglobin precursor in the body fluid sample.
4. The method of any one of claims 1 to 3, wherein the body fluid sample is plasma, serum, blood, saliva, urine or CSF.
5. The method of claim 4, wherein the body fluid sample is plasma.

6. The method of any one of claims 1 to 5, wherein the sample is immobilized on a solid support.

7. A method of monitoring the progression of Alzheimer's disease in a subject, the method comprising:

detecting the expression of alpha-2-macroglobulin precursor, clusterin precursor and fibrinogen gamma chain precursor in a body fluid sample from said subject at a first time point;

detecting the expression of alpha-2-macroglobulin precursor, clusterin precursor and fibrinogen gamma chain precursor in a body fluid sample from said subject at a later time point;

whereby differing expression of the alpha-2-macroglobulin precursor, clusterin precursor and fibrinogen gamma chain precursor in the sample from the subject at the later time point compared to the first time point indicates an increase in the severity of Alzheimer's disease.

8. The method of claim 7, wherein the method further comprises:

detecting the expression of complement C3 precursor in the body fluid sample from said subject at the first time point and in the body fluid sample at the later time point, whereby differing expression of the complement C3 precursor in the sample from the subject at the later time point compared to the first time point indicates an increase in the severity of Alzheimer's disease.

9. The method of claim 7 or 8, wherein the method further comprises:

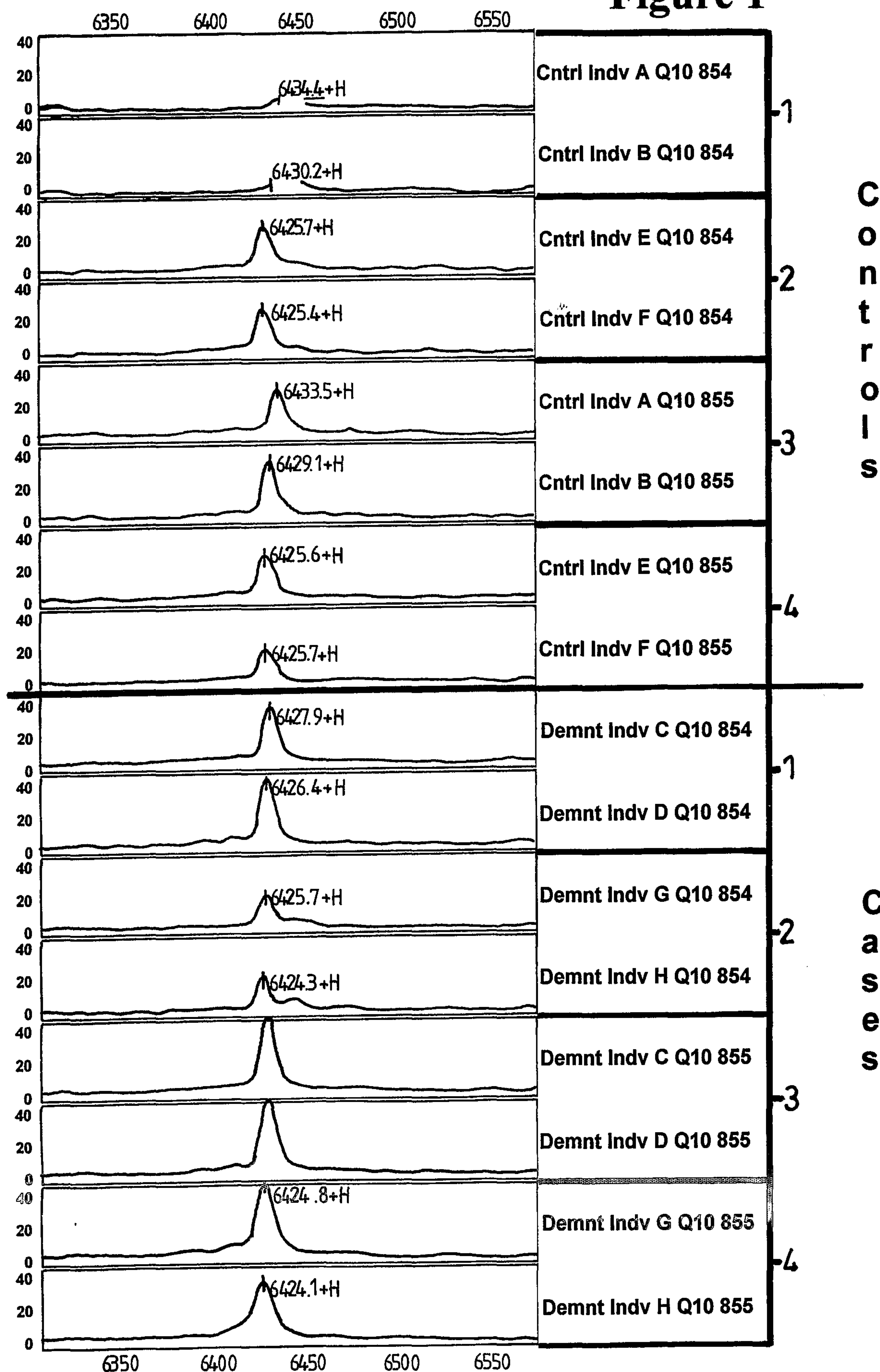
detecting the expression of one or more proteins selected from the group consisting of transthyretin, complement C4 precursor, complement factor H, and haptoglobin precursor in the body fluid sample from said subject in the body fluid sample at the first time point and in the body fluid sample at the later time point, whereby differing expression of the one or more proteins selected from the group consisting of transthyretin, complement C4 precursor, complement factor H, and haptoglobin precursor at the later time point compared to the first time point indicates an increase in the severity of Alzheimer's disease.

10. The method of any one of claims 7 to 9, wherein the body fluid sample is plasma, serum, blood, saliva, urine or CSF.
11. The method of claim 10, wherein the body fluid sample is plasma.
12. The method of any one of claims 7 to 11, wherein the alpha-2-macroglobulin precursor, the clusterin precursor and fibrinogen gamma chain precursor are detected by mass spectrometry.
13. The method of any one of claims 7 to 12, wherein the sample is immobilized on a solid support.
14. A kit comprising:
 - a reagent for detecting alpha-2-macroglobulin precursor;
 - a reagent for detecting clusterin precursor, and
 - a reagent for detecting fibrinogen gamma chain precursor.
15. The kit of claim 14, further comprising a reagent for detecting complement C3 precursor.
16. The kit of any one of claims 14 or 15, further comprising a reagent for detecting one or more proteins selected from the group consisting of transthyretin, complement C4 precursor, complement factor H, and haptoglobin precursor.
17. The kit of any one of claims 14 to 16, wherein the reagent for detecting alpha-2-macroglobulin precursor is an antibody specific to alpha-2-macroglobulin precursor, the reagent for detecting clusterin precursor is an antibody specific to clusterin precursor, and the reagent for detecting fibrinogen gamma chain precursor is an antibody specific to fibrinogen gamma chain precursor.
18. The kit of any one of claims 14 to 17, for use in the diagnosis of Alzheimer's disease.

19. Use of a kit according to any one of claims 14 to 18 for diagnosing Alzheimer's disease in a subject.

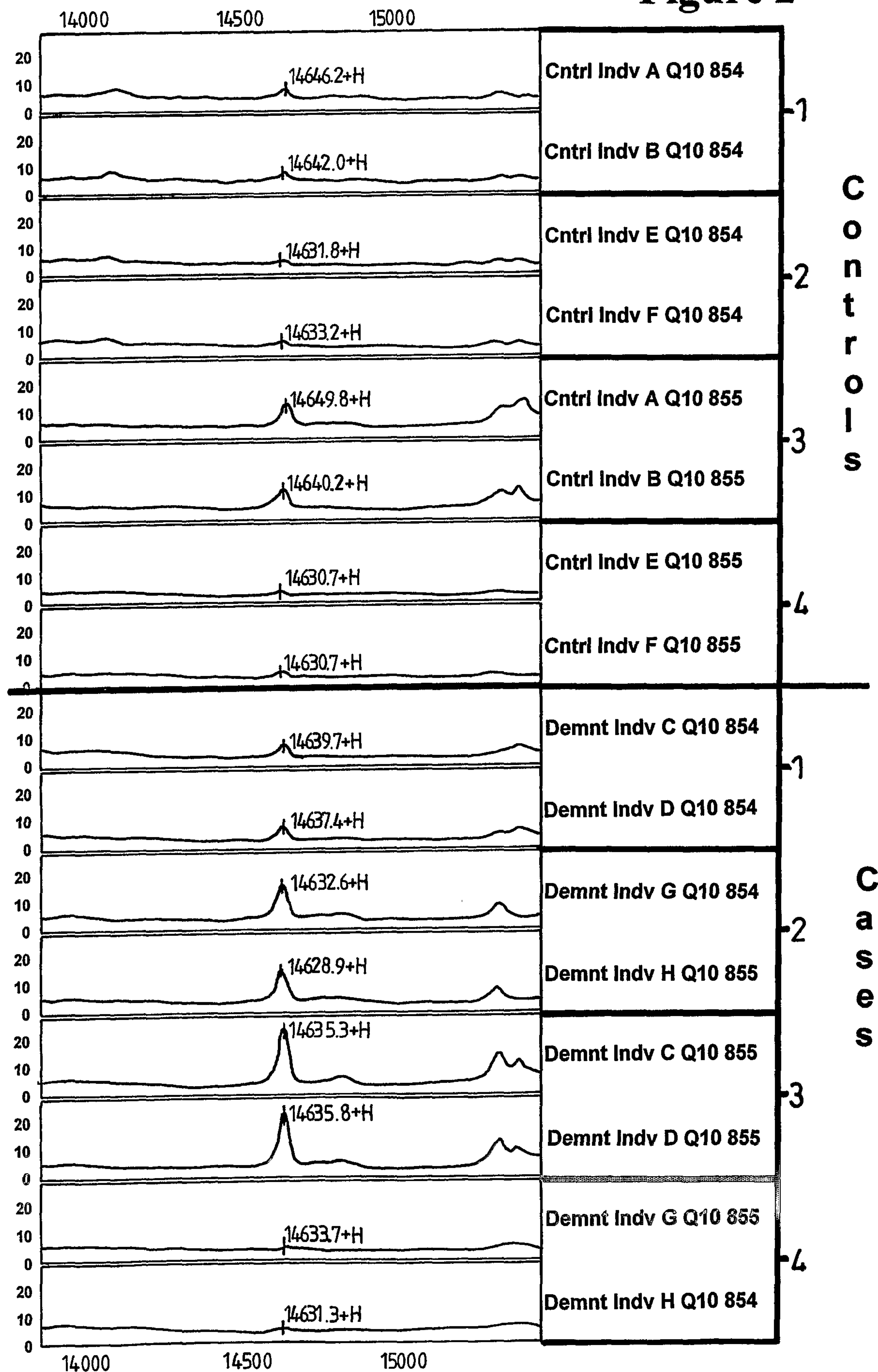
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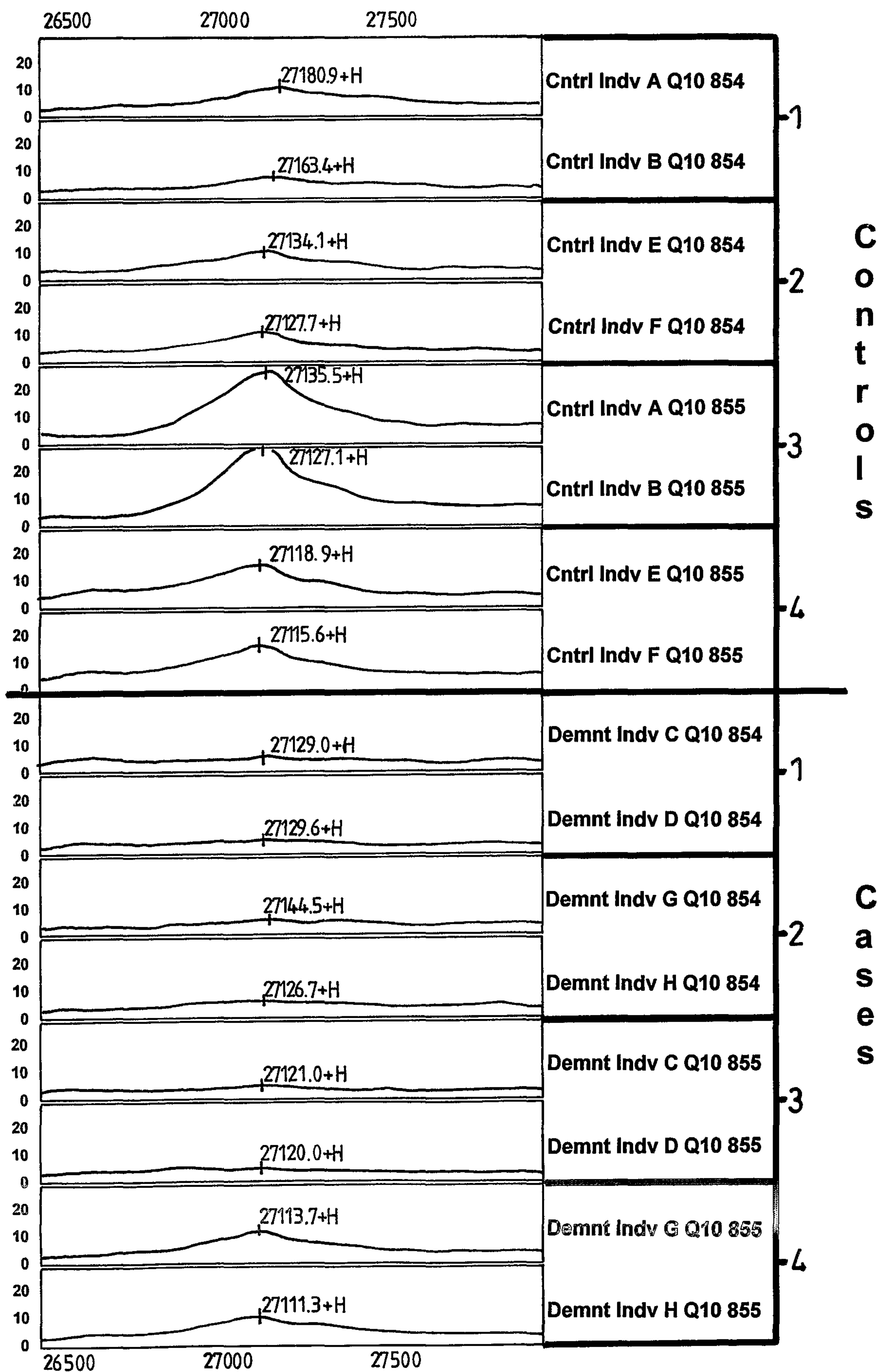
Figure 1



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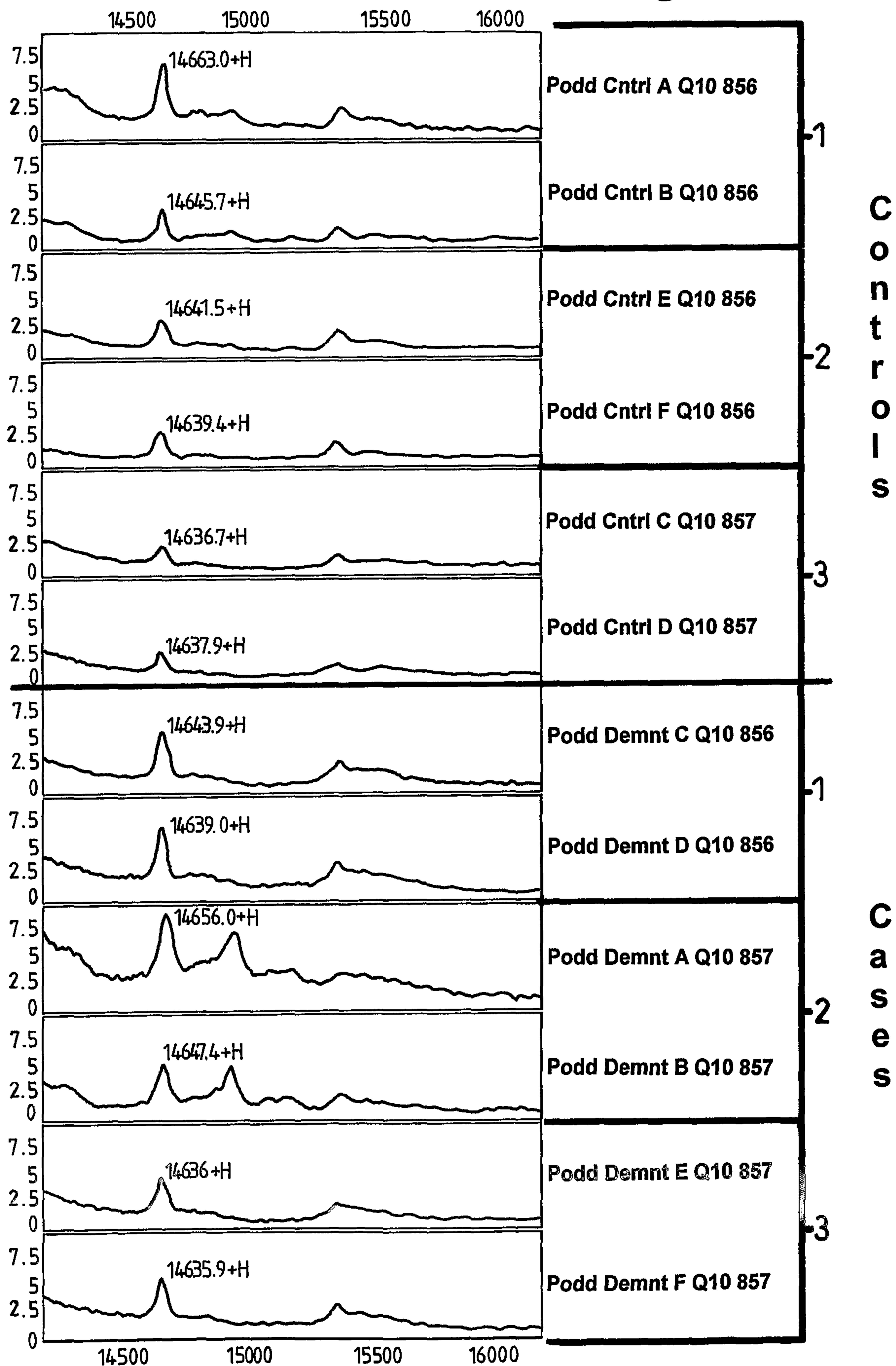
Figure 2



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Figure 4



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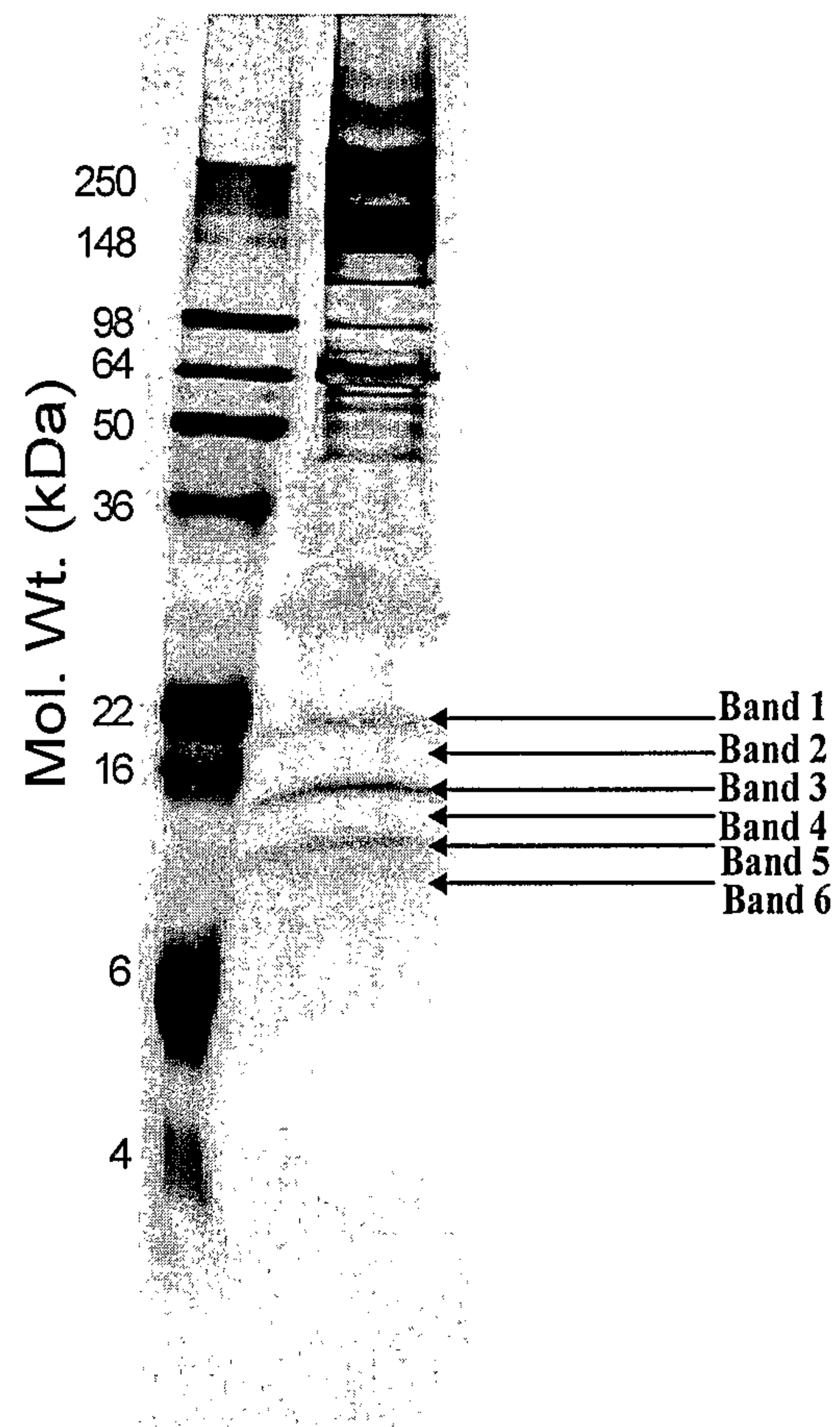


Figure 5

Spot No.	Rank	p Value	Fold Difference	State Change	Protein I.D.	Accession No.	Search Log No.
196	1	0.00030199	1.78	↑ AD	Desmoplakin (DP) (250/210 kDa paraneoplastic pemphigus antigen) Ig kappa chain C region Ig kappa chain V-II region TEW Serum amyloid P-component precursor (SAP) (9.5S alpha-1-glycoprotein)	P15924 P01834 P01617 P02743	7495 7542 7542 7951
171	2	0.001255545	2.11	↑ AD	Ig kappa chain C region Serum albumin precursor Galectin-7 (Gal-7) (HKL-14) (PI7) (p53-induced protein 1)	P01834 P02768 P47929	5623 7954 5623
2 (old)	3	0.001447694	13.75	↑ AD	Complement factor H precursor (H factor 1) Serum albumin precursor Alpha-2-macroglobulin precursor (Alpha-2-M) Ceruloplasmin precursor (EC 1.16.3.1) (Ferroxidase)	P08603 P02768 P01023 P00450	6672 6672 6672 6672
184	4	0.005360087	2.43	↑ AD	Ig lambda chain C regions Ig lambda chain V-III region LOI Serum albumin precursor Complement factor H-related protein 2 precursor (FHR-2)	P01842 P80748 P02768 P36980	7818 7818 7818 7818
177 (old)	5	0.005382883	1.92	↑ AD	Ig lambda chain C regions Serum albumin precursor Ig lambda chain V-III region LOI Ig kappa chain C region	P01842 P02768 P80748 P01834	5627 7955 7955 7955
4	6	0.005985336	8.83	↑ AD	Alpha-2-macroglobulin precursor (Alpha-2-M)	P01023	7827
170	7	0.01167553	1.98	↑ AD			
13	8	0.015500401	4.23	↓ AD	Inter-alpha-trypsin inhibitor heavy chain H4 precursor (ITI heavy chain H4) Ceruloplasmin precursor (EC 1.16.3.1) (Ferroxidase)	Q14624 P00450	7829 7829
165 (old)	9	0.018305158	1.58	↓ AD	Serum albumin precursor	P02768	5625
164	10	0.020647469	2.03	↓ AD	Complement C4 precursor [Contains: C4a anaphylatoxin; C4b]	P01028	7821

Figure 6

14	(old)	11	0.025004429				Ig gamma-1 chain C region	P01857	7821
					↓ AD	10.82	Serum albumin precursor	P02768	6227
							Histone H2B.a/g/h/k/l (H2B.1 A)	P62807	6227
							(H2B/a) (H2B/g) (H2B/h) (H2B/k) (H2B/l)		
126		12	0.028979402		↓ AD	1.6	CD5 antigen-like precursor (SP-alpha) (CT-2) (IgM-associated peptide)	O43866	7493
							Serum albumin precursor	P02768	7952
							Ig mu chain C region	P01871	7952
176		13	0.029106689		↑ AD	1.75	Ig lambda chain C regions	P01842	7816
							Serum albumin precursor	P02768	7816
							Ig lambda chain V-III region LOI	P80748	7816
123		14	0.031441346		↑ AD	1.36	Serum albumin precursor	P02768	7462
1		15	0.034723104		↑ AD	3.32	Alpha-2-macroglobulin precursor (Alpha-2-M)	P01023	7823
							Ig alpha-1 chain C region	P01876	7823

Band No.	Protein I.D.	Species	Accession No.	MW (Da)	Gel	MW (Da)	pI	No. Peptides Matched	Percentage Coverage	Error (ppm)	Search Log No.	Peptide Matched
SP1_1C	Haptoglobin precursor	Human	P00738	19300		45177	6.13	5	9%	16	5288	
SP1_2C	Transferrin	Human	g1339885	18200		12835	5.33	8	87%	223	5275	
	Serum albumin precursor	Human	P02768	18200		69248	5.82	3	6%	236	5274	
	Complement C4 precursor	Human	P01028	18200		192650	6.65	1	0%	236	5274	GLEELQFLGSK
	Fibrinogen alpha/alpha-E chain precursor	Human	P02671	18200		94914	5.7	1	3%	224	5274	EVTISEDGSDCPEAMDGLTSGIGTLDGFR
SP1_3C	Chain A, Transferrin	Human	g143295	14900		13753	5.35	10	92%	102	5251	
	Apolipoprotein A-IV precursor (Apo-AIV)	Human	P06727	14900		45343	5.28	2	10%	100	5247	
	Serum albumin precursor	Human	P02768	14900		69321	5.92	1	1%	109	5247	
SP1_4C	Transferrin precursor	Human	P02766	14200		15877	5.52	5	60%	170	5260	
	Hemoglobin beta chain	Human	P02023	14200		15857	6.81	4	28%	12	5260	
	Serum albumin precursor	Human	P02768	14200		69321	5.92	4	7%	7	5260	
SP1_5C	Haptoglobin-related protein precursor	Human	P00739	12600		38983	6.42	5	10%	102	5294	
	Transferrin precursor	Human	P02766	12600		15877	5.52	2	16%	129	5294	
	Serum albumin precursor	Human	P02768	12600		69321	5.92	4	10%	120	5294	
	Apolipoprotein C-III precursor (Apo-CIII)	Human	P02656	12600		10845	5.23	1	16%	122	5294	DALSSVQESQVAQAR
	Hemoglobin alpha	Human	P01922	12600		15227	9.84	2	17%	118	5294	
	Hemoglobin beta chain	Human	P02023	12600		15857	6.81	1	15%	131	5294	SAVTALWCKYNVDEVGGEALGR
SP1_6C	Serum albumin precursor	Human	P02768	11600		69321	5.92	6	11%	244	5280	
	Apolipoprotein C-III precursor (Apo-CIII)	Human	P02656	11600		10845	5.23	3	37%	244	5280	
	Haptoglobin precursor	Human	P00738	11600		45177	6.13	2	6%	233	5280	
	Vitronectin precursor (Serum spreading factor) (S-protein)	Human	P04004	11600		54271	5.55	1	3%	249	5280	SIAQYWLGCFAFGHL

Figure 7

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No	IPI Accession no	SWISS- PROT Accession no	Name	No of matched peptides	regulation (control / disease)	CV (%)
1	IPI00166866	P01876	MGC27165 PROTEIN	2	0,38	7
2	IPI00336074	P01876	IG ALPHA-1 CHAIN C REGION	2	0,35	4
3	IPI00423461	P01842	HYPOTHETICAL PROTEIN DKFZP686C0222 0 (FRAGMENT)	2	0,35	24
4	IPI00431645	P00738	HAPTOGLOBIN PRECURSOR	1	0,33	-
5	IPI00478493	P00738	HAPTOGLOBIN PRECURSOR	1	0,34	-

Figure 8

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1 MFLKAVVLTALVAVAGARA EVSADQVATV MWDYFSQLSN NAKEAVEHLQ
51 KSELTQQLNALFQDKLGEVN TYAGDLQKKL VPFATELHER LAKDSEKLKE
101 EIGKELEELR ARLLPHANEV SQKIGDNLRE LQQRLEPYAD QLRTQVNTQA
151 EQLRRQLTPY AQRMERVLRE NADSLQASLR PHADELKAKI DQNVEELKGR
201 LTPYADEFKV KIDQTVHEELR RSLAPYAQDT QEKLNHQLEG LTFQMKKNAE
251 ELKARISASA EELRQRLAPL AEDVRGNLKG NTEGLQKSLA ELGGHLDQQV
301 EEFRRRVEPY GENFNKALVQ QMEQLRQKLG PHAGDVEGHL SFLEKDLRDK
351 VNSFFSTFKE KESQDKTSL PELEQQQEQQ QEQQQEQVQM LAPLES

Figure 9

1 MRLLWGLIWA SSFFTLSLQK PRLLLFSPSV VHLGVPLSVG VQLQDVPRGQ
 51 VVKGSVFLRN PSRNNVPCSP KVDFTLSSER DFALLSLQVP LKDAKSCGLH
 101 QLLRGPEVQL VAHSPWLKDS LSRTTNIQGI NLLFS SRRGH LFLQTDQPIY
 151 NPGQVRVYRV FALDQKMRPS TDTITVMVEN SHGLRVRKKE VYMPSSIFQD
 201 DFVIPDISEP GTWKISARFS DGLESNSSTQ FEVKKYVLPN FEVKITPGKP
 251 YILTVPGHLD EMQLDIQARY IYGKPVQGVA YVRFGLLDED GKKTFFRGLE
 301 SQTCLVNGQS HISLSKAEFQ DALEKLNMG I TDLQGLRLYV AAAIIESPGG
 351 EMEEAELTSW YFVSSPFLD LSKTKRHLVP GAPFL LQALV REMSGSPASG
 401 IPVKVSATVS SPGSVPEVQD IQQNTDGSQ VSIPI IIPQT ISELQLSVSA
 451 GSHPHAIARL TVAAPPSGGP GFLSIERPDS RPPRVGDTLN LNLRAVGSGA
 501 TFSHYYMIL SRGQIVFMNR EPKRTLTSVS VFVDHHLAPS FYFVAFYYHG
 551 DHPVANSLRV DVQAGACEGK LELSVDGAKQ YRNGE SVKLH LETDSLALVA
 601 LGALDTALYA AGSKSHKPLN MGKVFEAMNS YDLGC GPGGG DSALQVFQAA
 651 GLAFSDGDQW TLSRKRLSCP KEKTTRKKRN VNFQK AINEK LGQYASPTAK
 701 RCCQDGVTRL PMMRSCEQRA ARVQQPDCRE PFLSC CQFAE SLRKKSRDKG
 751 QAGLQRALEI LQEEDLIDED DIPVRSFFPE NWLWRVETVD RFQILTLWLP
 801 DSLTTWEIHG LSLSKTKGLC VATPVQLRVF REFHLHLRLP MSVRRFEQLE
 851 LRPVLYNYLD KNLTVSVHVS PVEGLCLAGG GGLAQ QVLVP AGSARPVAFS
 901 VVPTAAAVS LKVVARGSFE FPGDAVSKV LQIEKEGAIH REELVYELNP
 951 LDHRGRTLEI PGNSDPNMIP DGDFNSYVRV TASDP LDTLG SEGALSPGGV
 1001 ASLLRLPRGC GEQTMIIYLA TLAASRYLDK TEQWS TLPPE TKDHAVDLIQ
 1051 KGYMRIQQFR KADGSYAAWL SRDSSTWLTA FVLKVLSLAQ EQVGG SPEKL
 1101 QETSNWLLSQ QQADGSFQDP CPVLDRSMQG GLVGNDETVA LTAFVTIALH
 1151 HGLAVFQDEG AEPLKQORVEA SISKANSFLG EKASAGLLGA HAAAITAYAL
 1201 SLTKAPVDLL GVAHNNLMAM AQETGDNLYW GSVTG SQSNA VSPTPAPRNP
 1251 SDPMPQAPAL WIETTAYALL HLLLHEGKAE MADQA SAWLT RQGSFQGGFR
 1301 STQDTVIALD ALSAYWIASH TTEERGLNVT LSSTGRNGFK SHALQLNNRQ
 1351 IRGLEEELQF SLGSKINVKV GGNSKGTLKV LRTYN VLDMK NTTCQDLQIE
 1401 VTVKGHVEYT MEANEDYEDY EYDELPKDD PDAPL QPVTP LQLFEGRNR
 1451 RRREAPKVVE EQESRVHYTV CIWRNGKVGL SGMAI ADVTL LSGFHALRAD
 1501 LEKLTSLSDR YVSHFETEGP HVLLYFDSVP TSREC VGF EA VQEV PVGLVQ
 1551 PASATLYDYY NPERRCSVFY GAPSKSRLLA TLCSA EVCQC AEGKCPRQRR
 1601 ALERGLQDED GYRMKFACY Y PRVEYGFQVK VLRED SRAAF RLFETKITQV
 1651 LHFTKDVKAA ANQMRNFLVR ASCRLRLPEG KEYLIMGLDG ATYDLEGHPQ
 1701 YLLDSNSWIE EMPSERLCRS TRQRAACAQL NDFLQ EYGTQ GCQV

Figure 10

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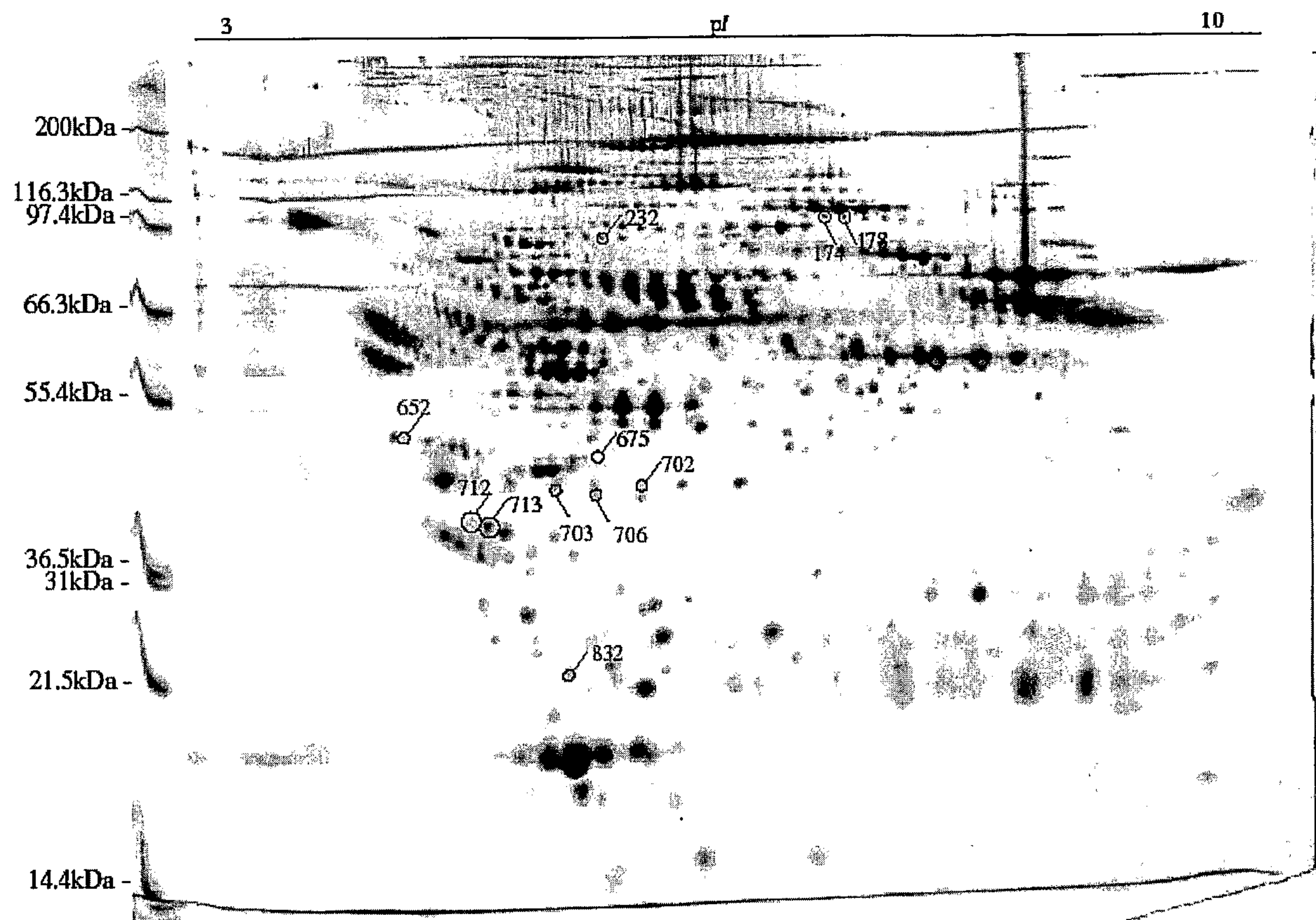


Figure 11

Spot no.	Protein Name	Acc. No.	Norm. Vol	CV (%)	Norm. Vol	CV (%)	Expressio n ratio	T-test (p)	Detection ratio	Theoretical Mr	pl	Cover -age (%)
174	alpha-2-macroglobulin precursor	P01023	0,36604	52	0,14510	57	0,40	2,50852E-06	28/21	160796 Da	5,95	13,8
178	alpha-2-macroglobulin precursor	P01023	0,29348	53	0,12475	57	0,43	1,25521E-05	27/20	160796 Da	5,95	11,7
232	Inter-alpha-trypsin inhibitor heavy chain H4 precursor	Q14624	0,32468	81	0,14481	64	0,45	0,001793752	28/27	103358 Da	6,51	25,6
712	Complement C3 precursor	P01024	0,81225	74	0,36446	62	0,45	0,001371446	28/24	184967 Da	6,00	14,9
712	Clusterin precursor	P10909	0,81225	74	0,36446	62	0,45	0,001371446	28/24	50062 Da	5,89	22,9
713	Complement C3 precursor	P01024	3,45803	67	1,44590	63	0,42	9,67927E-05	29/29	184967 Da	6,00	16,9
652	Complement C4 precursor	P01028	0,18067	122	0,39844	69	2,21	0,003595593	24/25	192771 Da	6,60	5,9
675	Actin cytoplasmic 2 (Gamma/beta actin)	P63261	0,21268	89	0,45443	74	2,14	0,002107165	25/28	41793 Da	5,31	49,1
702	Haptoglobin precursor	P00738	0,09210	86	0,40084	115	4,35	0,002920373	20/25	43349 Da	6,13	21,2
703	Haptoglobin precursor	P00738	1,54479	95	4,64500	76	3,01	0,000171685	24/28	43349 Da	6,13	23,9
706	Haptoglobin precursor	P00738	1,00814	112	3,26743	89	3,24	0,000607583	21/28	43349 Da	6,13	23,6
832	Complement C4 precursor	P01028	0,24743	126	0,61914	88	2,50	0,003005256	28/28	192771 Da	6,60	4,5

Figure 12

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1 MGKNKLLHPS LVLLLLLVLLP TDASVSGKPQ YMLVLPSSLH TETTEKGCVL LSYLNETHVT
 61 SASLESVRGN RSLFTDLEAE NDVLHCVAFA VPKSSSNEEV MELTVQVKGP TQEFKKRTTV
 121 MVKNEDSLVF VQTDKSIYKP GQTVKFRVVS MDENFHPLNE LIPLVYIQDP KGNRIAQWQS
 181 FQLEGGLKQF SFPLSSEPFQ GSYKVVVQKK **SGGRTEHPFT** **VEEFVLPKFE** VQVTVPKIIT
 241 ILEEEMNVSV CGLYTYGKPV PGHVTVSICR KYSDASDCHG EDSQAFCEKF **SGQLNSHGCF**
 301 **YQQVKTKVFQ** LKRKEYEMKL **HTEAQIQEEG** **TVVELTGRQS** SEITRTITKL **SFVKVDSHFR**
 361 **QGIPFFGQVR** **LVDGKGVPIP** **NKVIFIRGNE** ANYYSNATTD EHGLVQFSIN TTNVMGTSLT
 421 VRVNYKDRSP CYGYQWVSEE HEEAHTAYL VFSPSKSEFVH LEPMSHELPC GHTQTVQAHY
 481 ILNGGTLLGL KKLSFYYLIM AKGGIVRTGT HGLLVKQEDM **KGHFSISIPV** **KSDIAPVARL**
 541 LIYAVLPTGD VIGDSAKYDV **ENCLANKVDL** **SFSPSQSLPA** **SHHLRVTA** **PQSVCALRAV**
 601 DQSVLLMKPD AELSASSVYN LLPEKDLTGF PGPLNDQDDE DCINRHNVI NGITYTPVSS
 661 TNEKDMYSFL EDMGLKAFTN SKIRKPK**MCP** **QLQQYEMHGP** **EGLRVGFYES** DVMGRGHARL
 721 VHVEEPHTET VRKYFPETWI WDLVVVNSAG VAEVGVTVPD TITEWKAGAF CLSEDAGLGI
 781 SSTASLRAFQ PFFVELTMPY SVIRGEAFTL KATVLNLYPK CIRVSVQLEA SPAFLAVPVE
 841 KEQAPHCICA NGRQTVSWAV TPKSLGNVNF TVSAEALESQ ELCGTEVPSV PEHGRKDTVI
 901 KPLLVEPEGL EKETTFNSLL CPSGGEVSEE LSLKLPPNVV EESARASVSV LGDILGSAMQ
 961 NTQNLLQMPY GCGEQNMVLF APNIYVLDYL NETQQLTPEV KSKAIGYLNT GYQRQLNYKH
 1021 YDGSYSTFGE RYGRNQGNTW LTAFVLKTFA QARAYIFIDE AHITQALIWL SQRQKDNGCF
 1081 RSSGSLNNA IKGGVEDEV LSAYITIAL EIPLTVTHPV VRNALFCLES AWKTAQEGDH
 1141 GSHVYTKALL AYAFALAGNQ DKRKEVLKSL NEEAVKKDNS VHWERPQKPK APVGHFYEPQ
 1201 APSAEVEMTS YVLLAYLTAQ PAPTSEDLT ATNIVKWITK QQNAQGGFSS TQDTVVALHA
 1261 LSKYGAATFT RTGKAAQVTI QSSGTFSSKF QVDNNNRLLL QQVSLPELPG EYSMKVTGEG
 1321 CVYLQTSKY NILPEKEEFP FALGVQTLPO TCDEPKAHTS FQISLSVSYT GSRASNMAI
 1381 VDVKMVSGFI PLKPTVKMLE RSNHVSRTEV SSNHVLIYLD KVSNQTLSLF FTVLQDVPVR
 1441 DLKPAIVKVY DYYETDEFAI AEYNAPCSKD LGNA

Figure 13

15/20

1 MKPPRPVRTC SKVLVLLSLL AIHQTTTAEK NGIDIYSLTV DSRVSSR**FAH** **TVVTSRVVNR**
61 **ANTVQEATFQ** **MELPKKAFIT** NFSMNIDGMT YPGIIKEKAE AQAQYSAAVA KGKSAGLVKA
121 TGRNME**QFQV** **SVSVAPNAKI** **TFELVYEELL** **KRRLGVYELL** **LKVRPQQLVK** HLQMDIHIFE
181 PQGISFLETE STFMTNQLVD ALTTWQNKTK AHIRFKPTLS **QQQKSPEQQE** **TVLDGNLIIR**
241 YDVDRAIS**GG** **SIQIENGYFV** **HYFAPEGLTT** **MPKNVVFVID** KSGSMSGRKI QQTREALIKI
301 LDDLSPRDQF NLIVFSTEAT QWRPSLVPAS AENVNKARSE AAGIQALGGT NINDAMLMAV
361 QLLDSSNQEE RLPEGSVSLI **ILLTDGDPTV** **GETNPRSIQN** NVREAVSGRY SLFCLGFGFD
421 VSYAFLEKLA **LDNGGLARRI** HEDSDSALQL QDFYQEVANP LLTAVTFEYP SNAVEEVTQN
481 NFRLLFKGSE MVVAGKLQDR **GPDVLTATVS** **GKLPTQNITE** QTESSVAEQE AEFQSPKY**IF**
541 **HNFMERLWAY** LTIQQLLEQT VSASDADQQA LRNQALNLSL AYSFVTPLTS MVVTKPDDQE
601 QSQVAEKPME GESRNRNVHS **GSTFFKYYLQ** GAKIPKPEAS **FSPRRGWNRQ** AGAAGSRM**NF**
661 **RPGVLSSRQL** **GLPGPPDVPD** **HAAYHPFRRL** AILPASAPPA TSNPDPAVSR VMNMKIEETT
721 MTTQTPAPIQ APSAILPLPG QSVERLCVDP RHRQGPVNLL SDPEQGVEVT GQYEREKAGF
781 SWIEVTFKNP LVVWHASPEH VVVTRNRRSS AYKWKETLFS VMPGLKMTMD KTGLLLLSDP
841 DKVTIGLLFW DGRGEGRLRL LRDTDRFSSH VGGTLGQFYQ EVLWGSPAAS DDGRRTL RVQ
901 GNDHSATRER RLDYQEGPPG VEISCWSVEL

Figure 14

16/20

1 MGPTSGPSLL LLLLTHLPLA LGSPMYSIIT PNILRLESEE TMVLEAHDAQ GDVPVTVTVH
 61 DFPGKKLVLS SEKTVLTPAT NHMGNVTFIT PANREFKSEK GRNKFVTVQA TFGTQVVEKV
 121 VLVSLQSGYL FIQTDKTIYT PGSTVLYRIF TVNHKLLPVG RTVMVNIENP EGIPVKQDSL
 181 SSQNQLGVLP LSWDIPELVN MGQWKIRAYY ENSPQQVFST EFEVKEYVLP SFEVIVEPTE
 241 KFYIYNEKG LEVTITARFL YGKKVEGTAF VIFGIQDGEQ RISLPESLKR IPIEDGSGEV
 301 VLSRKVLLDG VQNLRAEDLV GKSLYVSATV ILHSGSDMVQ AERSGIPIVT SPYQIHFTKT
 361 PKYFKPGMPF DLMVFVTNPD GSPAYRVPVA VQGEDTVQSL TQGDGVAKLS INTHPSQKPL
 421 SITVRTKKQE LSEAEQATRT MQALPYSTVG NSNNYLHLSV LRTEL RPGET LNVNELLRMD
 481 RAHEAKIRYY TYLIMNKGRL LKAGRQVREP GQDLVVLPLS ITTDFIPSFR LVAYYTLIGA
 541 SGQREV VADS VWVDVKDSCV GSLVVKSGQS EDRQPVPGQQ MTLKIEGDHG ARVVLVAVDK
 601 GV FVLNKKNK LTQSKIWDVV EKADIGCTPG SGKDYAGVFS DAGLTFTSSS GQQT AQRAEL
 661 QCPQPAARRR RSVQLTEKRM DKVGKYPKEL RKCCEDGMRE NPMRFSCQRR **TRFISLGEAC**
 721 **KKVFLDCCNY** ITELRRQHAR ASHLGLARSN LDEDIAEEN IVSRSEFPES WLWNVEDLKE
 781 PPKNGISTKL MNIFLKDSIT TWEILAVSMS DKKGICVADP FEVTVMQDFF IDLRLPYSVV
 841 RNEQVEIRAV LYN YRQNQEL KVRVELLHNP AFCSLATTKR RHQQT VTI PP KSSL SVPYVI
 901 VPLKTGLQEV EVKAAVYHHF ISDGV RKS LK VVPEGIR MNK TVAVRTLDPE RLGREG**VQKE**
 961 **DIPPADLSDQ** **VPDTESETRI** **LLQGTPVAQM** **TEDA VDAERL** KHLIVTPSGC GEQNMIGMTP
 1021 TVIAVHYLDE TEQWEKFGLE KRQGALELIK **KGYTQQLAFR** **QPSSAFAAFV** **KRAPSTWLTA**
 1081 YVVKV**FSLAV** **NLIAIDSQVL** **CGAVKWLILE** KQKPDGV**FQE** DAPVIHQEMI GGLRNNNEKD
 1141 MALTAFLIS **LQEAKDICEE** **QVNSLP GSIT** **KAGDFLEANY** **MNLQRSYTVA** **IAGYALA QMG**
 1201 RLKGPLL NKF LTTAKDKNRW EDPGKQLYNV **EATSYALLAL** **LQLKDFDFVP** **PVVRWLNEQR**
 1261 **YYGGGYGSTQ** **ATEMV FQALA** **QYQKDAPDHQ** **ELNLDVSLQL** **PSRSSKITHR** **IHWESASLLR**
 1321 SEETKENE**GF** **TVTAEGKGQG** **TL SVVTMYHA** **KAKDQLTCNK** FDLKVTIKPA PETEKRPQDA
 1381 KNTMILEICT RYRGDQDATM SILDISM MTG FAPDTDDLKQ LANGVDRYIS KYELDKAFSD
 1441 RNTLIIYLDK VSHSEDDCLA FKVHQYFNVE LIQPGAVKVY AYYNLEESCT RFYHPEKEDG
 1501 KLNKLCRDEL CRCAEENC FI QKSDDKV TLE ERLDKACEPG VDYVYKTRLV KVQLSND FDE
 1561 YIMAIEQTIK SGSDEVQVGQ QRTFISPIKC REALKLEEK H YLMWGLSSD FWGEKPNLSY
 1621 IIGKDTWVEH WPEEDECQDE ENQKQCQDLG AFTESMVVFG CPN

Figure 15

17/20

1 MMKTLLLFVG LLLTWESQV LGDQTVSDNE LQEMSNQGSK **YVNKEIQNAV** NGVKQIKTLI
61 EKTNEERKTL LSNLEEAKKK KEDALNETRE SETKLKELPG **VCNETMMALW** **EECKPCLKQT**
121 CMKFYARVCR SGSGLVGRQL EEFLNQSSPF YFWMNGDRID SLENDRQQT HMLDVMQDHF
181 **SRASSIIDEL** **FQDRFFTREP** **QDTYHYLPFS** **LPHRRPHFFF** **PKSRIVRSLM** PFSPYEPLNF
241 HAMFQPFLEM IHEAQQAMDI HFHSPAFOHP PTEFIREGDD DRTVCREIRH NSTGCLRMKD
301 QCDKCREILS VDCSTNNPSQ AKLRRELD**ES** **LQVAERLTRK** YNELLKSYQW KMLNTSSLLE
361 QLNEQFNWVS RLANLTQGED QYYLRVTTVA SHTSDSDVPS GVTEVVVKLF **DSDPITVTVP**
421 **VEVSRKNPKF** METVAEKALQ EYRKKHREE

Figure 16

18/20

1 MRLLLWGLIWA SSFFTLSLQK PRLLLFSPSV VHLGVPLSVG VQLQDVPRGQ VVKGSVFLRN
 61 PSRNNVPCSP KVDFTLSSER DFALLSLQVP LKDAKSCGLH QLLRGPEVQL VAHSPWLKDS
 121 LSRTTNIQGI NLLFSSRRGH LFLQTDQPIY NPGQVRVRYRV FALDQKMRPS TDTITVMVEN
 181 SHGLRVRKKE VYMPSSIFQD DFVIPDISEP GTWKISARFS DGLESNSSTQ FEVKKYVLPN
 241 FEVKITPGKP YILTVPGHLD EMQLDIQARY IYGKPVQGVA YVRFGLLDED GKKTFFRGLE
 301 SQTCLVNGQS HISLSKAEFQ DALEKLNMGIT DLQGLRLYV AAATIESPGG EMEEAELTSW
 361 YFVSSPFSLD LSKTKRHLVP GAPFLLQALV REMSGSPASG IPVKVSATVS SPGSVPEVQD
 421 IQQNTDGSGQ VSIPIIIPQT ISELQLSVSA GSPHPAIARL TVAAPPSGGP GFLSIERPDS
 481 RPPRVGDTLN LNLRAVGSGA TFSHYYYMIL SRGQIVFMNR EPKRTLTSVS VFVDHHLAPS
 541 FYFVAFYYHG DHPVANSLRV DVQAGACEGK LELSVDGAKQ YRNGESVKLH LETDSLALVA
 601 LGALDTALYA AGSKSHKPLN MGKVFEAMNS YDLGCGPGGG DSALQVFQAA GLAFSDGDQW
 661 TL~~SRKRLSCP~~ **KEKT**TRKKRN VNFQKAIN**EK** **LGQYASPTAK** RCCQDGVTRL PMMRSCEQRA
 721 ARVQQPDCRE PFLSCCQFAE SLRKKSRDKG QAGLQRALEI LQEEIDLDED DIPVRSFFPE
 781 NWLWRVETVD RFQILTLWLP DSLTTWEIHG LSLSK**TKGLC** **VATPVQLRVF** **REFHLHLRLP**
 841 **MSVRR**FEQLE LRPVLYNYLD KNLTVSVHVS PVEGLCLAGG GGLAQQVLVP AGSARPVAFS
 901 VVPTAAAVS LKVVAR**GSFE** **FPVGDAVSKV** **LQIEKEGAIH** **REELVYELNP** **LDHRGR****TLEI**
 961 **PGNSDPNMIP** **DGDFNSYVRV** **TASDPLDTLG** **SEGALSPGGV** **ASLLRLPRGC** **GEQ****TM****IYLAP**
 1021 **TLAAS**RYLDK TEQWSTLPPE TKDHAVDLIQ KGYMRIQQFR KADGSYAAWL SRDSSTWLTA
 1081 FVLKVLSLAQ EQVGGSPEKL QETSNWLLSQ QQADGSFQDP CPVLDRSMQG GLVGNDETVA
 1141 LTAFVTIALH HGLAVFQDEG AEPLKQORVEA SISKANSFLG EKASAGLLGA HAAAITAYAL
 1201 SLTKAPVDLL GVAHNNLMAM AQETGDNLYW GSVTGSQSNA VSPTPAPRNP SDPMPQAPAL
 1261 WIETTAYALL HLLLHEGKAE MADQASAWLT **RQGSFQGGFR** STQDTVIALD ALSAYWIASH
 1321 TTEERGLNVT LSSTGRNGFK SHALQLNNRQ IRGLEEELQF SLGSKINVKV GGNSKGTLKV
 1381 LRTYNVLDK NTTCQDLQIE VTVKGHVEYT MEANEDYEDY EYDELPKDD PDAPLQPVTP
 1441 LQLFEGRRNR RRREAPKVVE EQESRVHYTV CIWRNGKVGL SGMAIADVTL LSGFHALRAD
 1501 LEKLTSLSDR YVSHFETEGP HVLLYFDSVP TSRECVGFEA VQEVFVGLVQ PASATLYDYY
 1561 NPERRCSVYF GAPSKSRLLA TLCSAEVCQC AEGKCPRQRR ALERGLQDED GYRMKFACY
 1621 PRVEYGFQVK VLREDSRAAF RLFETKITQV LHFTKDVCAA ANQMRNFLVR ASCRLRLEPG
 1681 KEYLIMGLDG ATYDLEGHPQ YLLDSNSWIE EMPSERLCRS TRQRAACAQL NDFLQEYGTQ
 1741 GCQV

Figure 17

19/20

1 MEEEEIAALVI DNGSGMCKAG FAGDDAPRAV **FPSIVGRPRH** QGVMVGMGQK **DSYVGDEAQS**
61 **KRGILTLKYP** IEHGIVTNWD DMEKIWHHTF YNELRVAPEE **HPVLLTEAPL** **NPKANREKMT**
121 QIMFETFNTP AMYVAIQAVL SLYASGR**TTG** **IVMDSGDGVT** **HTVPIYEGYA** **LPHAILRLDL**
181 **AGRDLTDYLM** KILTERGYSF TTTAEREIVR DIKEKLCYVA LDFEQEMATA ASSSSLEKSY
241 ELPDGQVITI GNERFRCPEA LFQPSFLGME SCGIHETTFN SIMKCDVDIR **KDLYANTVLS**
301 **GGTTMYPGIA** DRMQKEITAL APSTMKIKII **APPERKYSVW** IGGASILASLS TFQQMWISKQ
361 **EYDESGPSIV** HRKCF

Figure 18

20/20

1 MSALGAVIAL LLWGQLEAVD SGNDVTDIAD DGCPKPPEIA HGYVEHSVRY QCKNYKLRRT
61 EGDGVYTLND KKQWINKAVG DKLPECEADD GCPKPPEIAH GYVEHSVRYQ CKNYYKL RTE
121 GDGVYTLNNE KQWINKAVGD KLPECEAVCG KPKNPANPVQ RILGGHLD AK GSFPWQAKMV
181 SHHNLT TGAT LINEQWLLTT AKNLFLNHSE NATAKDIAPT **LTLYVGKKQL** VEIEKVVLHP
241 NYSQVDIGLI KLKQKVS VNE RVMPICLPSK **DYAEVGRVGY** VSGWGRNANF **KFTDHLKYVM**
301 **LPVADQDQCI** **RHYEGSTVPE** **KKTPKSPVGV** **QPILNEHTFC** **AGMSKYQEDT** CYGDAGSAFA
361 VHDLEEDTWY **ATGILSFDKS** **CAVAEYGVYV** KVTSIQDWVQ KTIAEN

Figure 19