Mutation screening of patients with Alzheimer disease identifies APP locus duplication in a Swedish patient

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Abstract

Background: Missense mutations in three different genes encoding amyloid-β precursor protein, presenilin 1 and presenilin 2 are recognized to cause familial early-onset Alzheimer disease. Also duplications of the amyloid precursor protein gene have been shown to cause the disease. At the Dept. of Geriatric Medicine, Karolinska University Hospital, Sweden, patients are referred for mutation screening for the identification of nucleotide variations and for determining copy-number of the APP locus.

Methods: We combined the method of microsatellite marker genotyping with a quantitative real-time PCR analysis to detect duplications in patients with Alzheimer disease.

Results: In 22 DNA samples from individuals diagnosed with clinical Alzheimer disease, we identified one patient carrying a duplication on chromosome 21 which included the APP locus. Further mapping of the chromosomal region by array-comparative genome hybridization showed that the duplication spanned a maximal region of 1.09 Mb.

Conclusions: This is the first report of an APP duplication in a Swedish Alzheimer patient and describes the use of quantitative real-time PCR as a tool for determining copy-number of the APP locus.

Background

Missense mutations in three different genes encoding amyloid-β precursor protein [1] (APP, [MIM 104760]), presenilin 1 [2] (PSEN1, [MIM 104311]) and presenilin 2 [3] (PSEN2, [MIM 600759]) are recognized to cause familial early-onset Alzheimer disease (EO-AD; [MIM 104300]). The mutations are known to functionally change the proteolytic processing of the APP protein, which leads to an increased Aβ42/Aβ40 ratio and increased Aβ-deposition in the brain. In addition to nucleotide variations, an increased dosage of the APP gene is also known to cause Alzheimer disease (AD) [4]. This is in agreement with previous observations that patients with Down syndrome (DS, trisomy 21, [MIM 190685]), who carry an extra copy of the APP gene, develop AD early in age [5-7]. Duplication of the APP locus has been detected in six unrelated French families [4,8], two Dutch families [9], one Finnish family [10], five families from a UK screen [11], and in two Japanese families [12]. The reported size of APP duplications range from 0.5 to 6.5 Mb, with the exception of an extended discontinuous duplication identified to be 15.5 Mb plus 1.5 Mb found in the UK screen [11]. Notably, a study on EO-AD cases from Finland and Sweden failed to identify duplications of the APP locus in a total of 141 AD patient, and concluded that the prevalence in these populations are low [13].

At the Dept. of Geriatric Medicine, Karolinska University Hospital, Sweden, patients are referred for mutation screening at the Genetic Unit for the identification of nucleotide variations and for determining copy-number of the APP locus. During the period between April 2008 and June 2010, a total number of 22 patients with clinical AD were referred for mutation screening, and the first finding of an APP duplication in a Swedish Alzheimer patient was made. The duplication was identified by analysis of microsatellite

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markers and quantitative real-time PCR, while confirmed by array-based comparative genome hybridization (aCGH). The duplication covers 1.09 Mb on chromosome 21q21, including the entire gene for APP. In this cohort of patients referred for mutation screening, the genomic duplication occurs at a frequency of 4.5% (1/22). This shows the importance of continued screening for APP locus duplication in Swedish AD patients, in parallel with sequencing efforts for the detection of nucleotide variations.

Methods
Patient and family history
Twenty-two patients were referred for mutation screening in the genes APP, PSEN1, PSEN2 at the Genetics Unit as part of the clinical investigation at the Dept of Geriatric Medicine, Karolinska University Hospital, Sweden. We made a subjective classification of the 22 subjects into four groups: early onset familial AD (EO-FAD); familial AD (FAD); possible familial AD (poss. FAD) and AD (Additional file 1). Ten cases were classified as “EO-FAD”, where an autosomal dominant history of early onset (≤65 years of age) in at least three affected family members in two generations were found. Four of the cases were classified as “FAD”, where there was at least three affected family members in two generations but where the onset in one of the family members was >65 years of age, and thus not fulfilling the criteria for early-onset. Six of the subjects were categorized as “possible FAD”, since the number of identified family members with AD were not sufficient to definitively conclude the nature of inheritance. Finally, two patients were referred for mutation screening on the basis of autopsy-confirmed severe cerebral amyloid angiopathy (CAA) or because of an extreme early-onset, <30 years of age and classified as “AD.”

The study was performed in accordance with the Helsinki Declaration, with informed consent and approval from the local ethics committee (Stockholm).

DNA samples
Genomic DNA was extracted from peripheral blood using the manufacturer’s protocol (Genta Purgene Blood kit, Qiagen, Sweden). A DNA sample from an individual with trisomy 21 (T21) was used as a positive control for the microsatellite marker analyses and for the APP copy-number assay (Coriell Cell Repositories Camden, New Jersey, USA, (Catalog ID: GM02767).

DNA sequencing
Sequencing reactions were performed using BigDye® Terminator v3.1 Cycle sequencing kit (Applied Biosystems, Carlsbad, California, USA) followed by capillary electrophoresis on a ABI 3100 Genetic Analyzer (Applied Biosystems, Carlsbad, California, USA). Primers and PCR conditions are available on request.

Microsatellite marker analysis
DNA from patients, D01-D22, were amplified with primers for microsatellite marker GDB:188463 (D21S265), located 330 kb centromeric to the APP gene, and for GBD:196999 (APP-dint), located in intron 1 of APP [4]. Fluorescent PCR-products were separated by capillary electrophoresis on an ABI 3100 instrument (Applied Biosystems, Carlsbad, California, USA) and the electropherograms were analyzed by using the GeneMapper 3.7 software (Applied Biosystems, Carlsbad, California, USA). The area under each allele’s major peak was determined and the ratio for the two different allele-areas for a marker was calculated to evaluate copy-number variation. A balanced copy-number of two was scored for ratios between 0.8 and 1.4, whereas ratios less than 0.65 or greater than 1.8 were used to define samples with three alleles.

Amplification with Taq-polymerase can result in so-called stutter-peaks when two alleles only differ by two nucleotides in size [14]. Therefore, markers with alleles that differ by two nucleotides were allowed to have ratios between 1.4 - 1.8 and still score as a normal copy-number. Mono-allelic markers yielding only one peak were not informative for indicating copy-number.

Copy-number assay
Quantitative real-time PCR was performed with the Taqman® copy-number assay method according to the manufacturer’s protocol (Applied Biosystems, Carlsbad, California, USA). The assays amplify target-regions within the APP locus in exon 1 (Assay ID: Hs00569527_cn), exon 7 (Assay ID: Hs02339796_cn), and in exon 17 (Assay ID: Hs00255904_cn). Additional assays amplify regions in closely located genes; MIR155HG (Assay ID: Hs04070445_cn), ATP5J (Assay ID: Hs02552822_cn), CYYR1 (Assay ID: Hs01271635_cn), and ADAMTS1 (Assay ID: Hs00609065_cn) (Additional file 2). Briefly, target assay-primers with a FAM-dye labeled probe and the calibrator RNase P-primers with a VIC-dye labeled probe, were used together in a duplex real-time PCR-reaction. The cycle threshold (Ct)-value at log-linear phase was used to calculate delta Ct-valuesample (dCt-sample) between the target assay and calibrator assay (CtAssay - CtCalibrator = dCt-sample). The mean from tetraplicates of delta Ct_samples from each sample DNA (D01-D22, and T21), was used for calculating relative copy-number normalized to the mean dCt-value of the trisomy 21 control DNA (T21) by the equation: 2^((dCtT21-dCtSample)*3).

Comparative genome hybridisation
The commercially available 180 K oligonucleotide catalog design (design ID 022060, Agilent Technologies, Palo Alto, CA, USA) with complete genome coverage and overall median probe spacing of 13 kb was used.
Experiments were performed according to the manufacturer’s protocol with minor modifications. Briefly, patient DNA and a gender-matched control (Promega, Wisconsin, USA) were labeled by random priming (Enzo Life Sciences, New York, USA) with Cy3-dUTP and Cy5-dUTP, respectively. After denaturing the probe and pre-annealing with 50 μg of Cot-1 DNA (Invitrogen, California, USA), hybridization was performed at 65 °C with rotation for 24 hours. The array was scanned with the Agilent microarray scanner (Palo Alto, CA, USA). Feature Extraction (v10.2) and DNA Analytics (v4.0) software packages (Agilent Technologies) were used for analysis and duplication breakpoints were determined by visual inspection of the numerical normalized log2 ratio analysis and duplication breakpoints were determined by the copy-number assay (Palo Alto, CA, USA). The genomic positions refer to the Human Genome Build 36.1 (UCSC Genome Browser, assembly March 2006).

Results
Twenty-two patients were referred for mutation screening in the genes APP, PSEN1 and PSEN2 during the period of April 2008 to June 2010 at the Dept. of Geriatric Medicine, Karolinska University Hospital, Sweden. The screening involved genomic sequencing of exons 16-17 in APP, exons 2-12 in PSEN1, exons 3-7 and exon 12 in PSEN2. No variation in APP was detected in any of the 22 DNA samples. In contrast we identified polymorphic SNPs in both PSEN1 and PSEN2. Polymorphic SNPs in PSEN2 were frequently detected and, in addition to the seven previously reported polymorphisms, we also found a novel variation at IVS6 +91 (G>C; GRCh37:1:227075950) (see Additional file 3 and Additional file 4). The functional implication on pathogenesis was tested by in silico methods. We were not able to detect a functional impact of the variation, neither by splice-prediction analysis nor by examining conservation between species (Additional file 4). Taken together, our sequencing effort did not find variations that could explain the diseased state of the patients.

To address the possibility of a duplication of APP, two microsatellite markers GDB:196999, (APP-dint), and GDB:188463, (D21S265) were PCR amplified with fluorescent primers. When analyzing the 22 DNA samples, 16 were heterozygous (polymorphic) for the marker GDB:196999 and 20 were heterozygous for marker GDB:188463. In our set-up, the size of alleles for marker GDB:196999 ranged from 163 bp to 191 bp, and the size of alleles for GDB:188463 ranged from 241 bp to 255 bp (Table 1). The allele-ratio calculated for marker GDB:196999 was between 0.8 - 1.4 indicating a balanced diploid copy-number, except for sample D08 (Table 1). In sample D08, the areas for alleles 174 bp and 189 bp for GDB:196999 had a ratio of 2.24 (17752/7932) indicating an unbalanced 2:1 copy-number (Figure 1). Six of the individuals were monoallelic for the marker. When calculating the allele-ratios for marker GDB:188463 in the sample-set, thirteen out of 22 DNA samples had ratios between 0.8 - 1.4. Seven samples had ratios between 1.4 - 1.8 and five of these samples had alleles that only differed by two nucleotides in size (D03, D18, D20 - 22); thereby leading to inflation of the area of the shorter allele (Table 1). The ratios for D13 and D15 were slightly higher than 1.4 and monoallelic for GDB:196999. Two samples were monoallelic for marker GDB:188463 (D08 and D16) and therefore did not generate any allele ratios. As an example, in Figure 1, DNA sample D07 carries two alleles for both markers, GDB:196999 and GDB:188463, and the calculated ratio for alleles GDB:196999-174 and GDB:196999-189 was 1.24, (3213/2582), indicating a balanced copy-number of two. Similarly, the ratio for the alleles of GDB:188463 is calculated to be 1.02 (902/883) for sample D07. Also illustrated, sample D08 was not informative for marker GDB:188463 since it only carried one allele, the 247 bp (Figure 1b), but an unbalanced copy-number are visible as the ratio, 2.24 (17752/7932), for the alleles of marker GDB:196999. Furthermore, sample D09 was not informative for marker GDB:196999 since it only carried one allele (189 bp), but could be examined at the GDB:188463 marker, with a ratio of 1.08 (6683/6190) (Figure 1b). The control DNA carrying trisomy 21 displayed three alleles for both examined markers (Figure 1a-b). The finding indicated that sample D08 carries a duplication of the genomic locus for marker GDB:196999.

To confirm the APP copy-number detected by microsatellite analysis, we used a quantitative real-time PCR method (The TaqMan Copy Number Assay, Applied Biosystems Carlsbad, California). The number of copies was determined by comparing patient DNA with a positive control DNA from a subject with trisomy 21 (copy-number of three). Three different exons of the APP gene were amplified by a duplex real-time PCR method for exon 1, exon 7, and exon 17. By calculating the relative copy-number, samples D07 and D09 were found to be diploid for all three amplicons (Figure 2, top). In contrast, sample D08 was found to have a copy-number value of three for all three amplicons. Thus, the copy-number assay confirmed the presence of three copies detected by the microsatellite analysis for D08. All the other DNA samples were determined by the copy-number assay to have relative copy-number values of 1.52-2.33, indicating a copy-number of two (Table 1). Noteworthy, the copy-number assay for samples D13 and D15 showed no variation in any of the exons examined and thus could be determined to carry normal number of the APP gene.

We wanted to address how far the duplication extended into the flanking chromosomal regions and if any neighboring genes were duplicated in addition to APP. For this purpose, copy-number assays targeting the genes MIR155HG,
ATP5J, CYYR1, and ADAMTS1 were analyzed (for locations see Additional file 2). The relative copy-number for samples D07 and D09 was two for all of the four investigated genes (Figure 2 lower). The relative copy-number values ranged from 1.78 (ATP5J, sample D07) to 2.15 (ATP5J, sample D09). The duplication established in sample D08 was determined to include the locus for the ATP5J gene located centromeric as well as the locus for the CYYR1 gene located telomeric of APP and the relative copy-number for these two assays were close to three, (3.42 and 3.49 respectively) (Figure 2 lower). However, the duplication did not include the more distal gene MIR155HG, located centromeric, or the gene for ADAMTS1 located telomeric (Figure 2 lower) as indicated by the relative copy-number values of 2.14 and 2.07 respectively. We could therefore limit the duplication to a maximal region of 1.27 Mb on chromosome 21 between genomic position 25 868029 bp and 27 138783 bp (Build 36.1).

To fine-map the duplicated region, array-CGH was used by hybridizing DNA sample D08 to a 180 k array covering the APP locus. The method compares hybridization signals from the D08 DNA with a diploid control DNA. The signal intensities were plotted against positions on the chromosome and shows that the duplication covers a region on chromosome 21 restricted to a maximum size of 1.09 Mb within coordinates 25 950727 bp - 27 043885 bp, and to a minimum size of 1.01 Mb within coordinates 25 957532 bp - 26 971 366 bp (Figure 3). Thus, the array-CGH confirms the presence of a genomic duplication including the APP gene in sample D08.

**Discussion**

The genetic causes for EO-FAD are strongly linked to three genes: APP, PSEN1, and PSEN2. The frequency in which mutations in these genes contribute to EO-FAD is reported to range from 17% to around 80% [15-18], depending on the population examined, the sample size, and the diagnostic criteria (for review see Avramopoulou 2009) [19]. At the Dept. of Geriatric Medicine, Karolinska University Hospital, Sweden, patients with a suspected genetic cause of AD are referred for mutation screening in the genes APP, PSEN1, and PSEN2. In addition to using sequencing for mutation screening, we have developed a protocol for analyzing the copy-number of APP. The protocol includes both a microsatellite analysis and a copy-number assay. Here we report on the results from

| GDB:196999 alleles | GDB:188463 alleles | CNA-APP ratio |
|-------------------|-------------------|---------------|
| D01 185/189       | 243/253           | 1.03          |
| D02 174/189       | 241/245           | 1.37          |
| D03 174/189       | 245/247           | 1.57          |
| D04 189/199       | 245/247           | 1.18          |
| D05 174/189       | 245/249           | 0.92          |
| D06 191           | 243/249           | 0.93          |
| D07 174/189       | 249/253           | 1.02          |
| D08 174/189       | 247              | -             |
| D09 189           | 241/253           | 1.07          |
| D10 163/174       | 245/255           | 1.04          |
| D11 174/189       | 247/255           | 1.19          |
| D12 163/174       | 249/255           | 0.97          |
| D13 189           | 247/251           | 1.45          |
| D14 174/178       | 245/249           | 1.38          |
| D15 189           | 249/253           | 1.53          |
| D16 163/189       | 247              | -             |
| D17 189           | 241/247           | 1.24          |
| D18 174/191       | 247/249           | 1.63          |
| D19 178/189       | 247/255           | 1.12          |
| D20 178/189       | 247/249           | 1.67          |
| D21 189           | 247/249           | 1.73          |
| D22 189/193       | 247/249           | 1.70          |

DNA samples are numbered D01-D22. From left; microsatellite-analysis for marker GDB:196999 (APPdint) and GDB:188463 (D21S265). The column “alleles” for each of the two markers gives the allele size in base-pairs. Also, for the two markers, and given in the column “ratio”, are the ratios between the peak-areas under each alleles major peak. Relative copy-number values from the copy-number assays (CNA-APP) targeting APP-exon 1 (E1), exon 7 (E7) and exon 17 (E17). The values in each column are calculated as relative to trisomy 21-DNA control and given as mean of tetraplicates. The abbreviation “nd” stands for not determined. Bold figures indicate ratio and copy-number of samples with suspected copy-number variation.
sequence (see Additional file 3 and Additional file 4) and copy-number analyses of 22 AD patients referred for mutation screening on the basis of their familial history with the detection of an APP duplication in one of the 22 patients. The calculated duplication frequency found in the present study depends on the number of samples considered, i.e. when only the 10 EO-FAD patients are included the frequency is 10%, and when including all 22 samples the frequency is 4.5%.

In contrast, our sequencing efforts did not identify any nucleotide variations in the three genes that could explain the diseased state. The low frequency of mutations in the Swedish population is also notable in earlier studies [20,21]. The mutations found so far in Swedish patients are six in total, and includes two in APP [22,23] and four in PSEN1 [20,24,25]. This finding strengthens the notion that mutations in APP, PSEN1 and PSEN2 are less frequent in Swedish families than in the reports on European EO-FAD. Significantly, 10 of the 22 patients had a family history with early-onset (< 65 year-old age), and an additional four of the patients came from families with an autosomal dominant history of AD.

The recognition of APP duplication as a genetic cause for AD has raised the issue of to what degree it explains familial EO-AD. Others have reported frequencies of APP-duplications between 2.7% and 8% in familial cases of AD [4,9,12]. This indicates that the contribution of APP duplications as a cause for the disease is one third to half of that from APP nucleotide missense mutations [15]. A report on screening for APP duplication in 77 Swedish and 64 Finnish patients did not succeed in finding any copy-number variations [13]. Again, this indicated that frequencies of genetic variations may differ between populations of different origin. However, in our study, the frequency of duplications is in the range of previous reports, with frequencies of 8% in French [4] and Japanese AD-families [12], and at 2.7% in Dutch familial AD-cases [9].

The duplication at chromosome 21 we reported is restricted to a maximum size of 1.09 Mb and to a minimum size of 1.01 Mb, which includes the genes for MRLP39, JAM2, ATPSJ, GABPA, APP, and CYYR1. All but two of the reported APP-duplications are of different sizes, and thereby include different genes. From the reports of different-sized duplications, it has also been
suggested that there is no single genomic architectural feature that serves as a recombination substrate, but that the APP locus may be a hotspot region for recombination due to multiple low copy repeats [9]. This is supported by our finding in a Swedish patient, since it adds a novel-sized duplication. It has been suggested that sequencing of the duplication break-points may shed further light on the molecular basis of the recombination in the region.

The two different-sized duplications found in Dutch families contain only the APP gene which strongly suggests that APP duplication is the only requirement for causing AD. The clinical characteristics in patient with APP-duplications are typically early-onset dementia, cerebral amyloid angiopathy (CAA) and intracerebral hemorrhage (ICH). When looking within families there are phenotypes of seizure and dementia with Lewy-bodies [8-10,26], but clearly these are not found in all diseased individuals. Rather, intra-familial diversity is

Figure 2 Bar graph showing copy-number from the copy-number assay targeting APP exons (top) and genes flanking APP gene (lower). On the y-axis is the copy-number calculated relative to trisomy 21-DNA control, and on the x-axis are three patient DNA samples, D07, D08, D09 and the trisomy-21 DNA (Tri21). For top graph, three different assays at the APP locus were used, positioned in Exon1, (blue), Exon7 (red), and in Exon17 (yellow)). For lower graph, four different assays were used targeting the genes of; MIR155HG, (blue), ATP5J (red), CYYR1 (yellow), and ADAMTS1 (light blue). Assays were performed in tetraplicate and bars are shown as mean ±standard deviation.
commonly found. In the Swedish family originating from proband III:2, (DNA sample D08), there was an early onset at 55 years in the cousin (III:4) (Figure 4). Individual III:4 had a medical history of syncope and a diagnosis of subcortical dementia. Similarly, the history of individual II:5 also contained remarks of syncope, as well as vertigo and the diagnosis of multi-infarct dementia. Individual II:4 had a history of mental illness and was hospitalized under the diagnosis of schizophrenia and psychosis. The duplication has only been confirmed in proband III:2, since there are no available DNA samples from other affected family members.

Conclusions
To conclude, we report the first Swedish EO-AD patient carrying an APP-duplication. We have successfully used the copy-number assay, showing that it is reliable and straightforward to use. It is also a good complement to the use of microsatellites to determine copy-number since these require the presence of heterozygous markers to be informative. We would like to emphasize the value of continued screening for APP-duplications in the
Swedish population of EO-AD, since we found it to be more frequent than nucleotide sequence variations in our cohort of patients referred for genetic mutation screening.

Additional material

Additional file 1: Table showing classification of the 22 cases of AD according to their family history with table legend.

Additional file 2: Schematic picture of the APP locus with surrounding genes with figure legend.

Additional file 3: Table showing nucleotide variations found in the 22 DNA samples for sequencing of APP, PSEN1, and PSEN2 with table legend.

Additional file 4: Method and result of on In-silico analysis of nucleotide variation IVS6 +9 and.

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Authors’ contributions

HT carried out the molecular genetic studies and made the in-silico analysis. JS carried out the genome hybridization study. MF and JB carried out the neuropathology. CG was responsible for referral of patients to genealogy of the patients as genetic counselors. IN judged clinical data and JS carried out the genome hybridization study. MF and JB carried out the molecular genetic studies and made the Cloning of a gene bearing missense mutations in early-onset familial Alzheimer’s disease. Nature 1995, 375(6534):754-760.

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