Early and sustained altered expression of aging-related genes in young 3xTg-AD mice

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Alzheimer’s disease (AD) is a multifactorial neurological condition associated with a genetic profile that is still not completely understood. In this study, using a whole gene microarray approach, we investigated age-dependent gene expression profile changes occurring in the hippocampus of young and old transgenic AD (3xTg-AD) and wild-type (WT) mice. The aim of the study was to assess similarities between aging- and AD-related modifications of gene expression and investigate possible interactions between the two processes. Global gene expression profiles of hippocampal tissue obtained from 3xTg-AD and WT mice at 3 and 12 months of age (m.o.a.) were analyzed by hierarchical clustering. Interaction among transcripts was then studied with the Ingenuity Pathway Analysis (IPA) software, a tool that discloses functional networks and/or pathways associated with sets of specific genes of interest. Cluster analysis revealed the selective presence of hundreds of upregulated and downregulated transcripts. Functional analysis showed transcript involvement mainly in neuronal death and autophagy, mitochondrial functioning, intracellular calcium homeostasis, inflammatory response, dendritic spine formation, modulation of synaptic functioning, and cognitive decline. Thus, overexpression of AD-related genes (such as mutant APP, PS1, and hyper-phosphorylated tau, the three genes that characterize our model) appears to favor modifications of additional genes that are involved in AD development and progression. The study also showed overlapping changes in 3xTg-AD at 3 m.o.a. and WT mice at 12 m.o.a., thereby suggesting altered expression of aging-related genes that occurs earlier in 3xTg-AD mice.

Keywords: Alzheimer’s disease; aging; microarray; inflammation; synaptic plasticity; mitochondria

Abbreviations: AD, Alzheimer’s disease; m.o.a., months of age; Aβ, β-amyloid; WT, wild type; qRT-PCR, quantitative real-time PCR; ER, endoplasmic reticulum; LTP, long-term potentiation; NFT, neurofibrillary tangle; GABA, γ-aminobutyric acid; IPA, Ingenuity Pathway Analysis; SNP, single-nucleotide polymorphism; LOWESS, locally weighted scatterplot smoothing

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Results

Cluster analysis of the investigated profiles revealed four different gene clusters (Supplementary Table 1), described in the following sections.

Cluster A. Cluster A consists of 35 transcripts that, compared with age-matched WT mice, were found downregulated in 3 m.o. and 12 m.o. 3xTg-AD mice. The same gene set was found to be upregulated in 12 m.o. WT mice when compared with 3 m.o. WT (Figure 1a). The main biological functions associated with these genes are shown in Figure 2a and Table 1.

Cluster B. Cluster B consists of 59 transcripts that, compared with 3 m.o. WT mice, were upregulated in 3 m.o. 3xTg-AD mice and 12 m.o. WT mice. The same set was downregulated in 12 m.o. 3xTg-AD mice when compared with 12 m.o. WT mice (Figure 1b). These genes are implicated in key biological functions depicted in Figure 2b and Table 2.

Cluster C. Cluster C consists of 193 transcripts that, compared with age-matched WT mice, were upregulated in 3 m.o. and 12 m.o. 3xTg-AD mice. The same set was upregulated in 12 m.o. WT mice compared with 3 m.o. WT mice (Figure 1c). Functional analysis revealed that the overexpressed transcripts are associated with functions depicted in Figure 2c and Table 3.

Cluster D. Cluster D consists of 76 transcripts that, compared with 3 m.o. WT mice, were downregulated in 3 m.o. 3xTg-AD mice and 12 m.o. WT mice. These transcripts were upregulated in 12 m.o. 3xTg-AD mice when compared with 3 m.o. WT mice (Figure 1d). The main biological functions associated with these genes are provided in Figure 2d and Table 4.

Figure 1 Unsupervised hierarchical clustering analysis. Transcripts that are clustered according to their expression values (log ratios) are shown. Each row indicates a transcript. The nine columns depict three replicates for each of the three experimental conditions listed on the top of the figure. Quantitative changes in gene expression are shown in colors. Red and green indicate upregulated and downregulated transcripts, respectively. Black indicates no changes in expression. Missing data points are shown in gray. (a) Cluster of 35 transcripts that were found downregulated in 3 m.o. 3xTg-AD versus 3 m.o. WT mice and 12 m.o. 3xTg-AD versus 12 m.o. WT mice. The same genes were instead upregulated in 12 m.o. 3xTg-AD versus 3 m.o. WT mice. (b) Cluster of 59 transcripts that were found to be specifically upregulated in 3 m.o. 3xTg-AD versus 3 m.o. WT mice and in 12 m.o. WT versus 3 m.o. WT mice. The same genes were instead downregulated in 12 m.o. 3xTg-AD versus 12 m.o. WT mice. (c) Cluster of 193 transcripts that were upregulated in all the experimental conditions (3 m.o. 3xTg-AD versus 3 m.o. WT mice and 12 m.o. 3xTg-AD versus 12 m.o. WT mice as well as in 12 m.o. WT versus 3 m.o. WT mice). (d) Cluster of 76 transcripts that were found downregulated in 3 m.o. 3xTg-AD versus 3 m.o. WT mice and in 12 m.o. WT versus 3 m.o. WT mice. The same genes were instead upregulated in 12 m.o. 3xTg-AD versus 12 m.o. WT mice.

TaqMan qRT-PCR: microarray data validation. In order to validate the microarray results, quantitative real-time PCR (qRT-PCR) analysis was performed on RNA extracted from the same hippocampal samples employed for microarray experiments. Analysis was performed on three upregulated genes (BECN1, CST3, and GABRA5) belonging to cluster C. In all cases, qRT-PCR confirmed the microarray results (Figure 3). Moreover, expression of the employed housekeeping gene (GAPDH) was stable among all the samples.

Interestingly, with the limitation and interpretative caution dictated by the low number of genes investigated by qRT-PCR (n = 3), we observed that young 3xTg-AD animals did already reach maximal expression that was not further changed, in a statistically significant manner, by aging (see 12 m.o. 3xTg-AD animals). On the contrary, in WT mice, these changes were occurring in an age-dependent way. Interestingly, young 3xTg-AD mice actually showed expression values that were higher compared with old WT mice.
Discussion

The main purpose of the study was to compare gene effects promoted by the pro-AD environment offered by the 3xTg-AD mice with those triggered, in WT mice, by senescence.

The most relevant finding is the identification of profile modifications detected in young 3xTg-AD mice that closely resemble those occurring upon physiological aging in WT animals, thereby suggesting that 3 m.o. 3xTg-AD mice...
undergo premature modulation of aging-related genes (Figure 4). For clarity and to improve readability of the section, we are discussing results of the three experimental conditions as follows:

- 3 m.o. 3xTg-AD versus 3 m.o. WT mice;
- 12 m.o. 3xTg-AD versus 12 m.o. WT mice;
- 12 m.o. WT versus 3 m.o. WT mice.

### 3 m.o. 3xTg-AD versus 3 m.o. WT mice

Comparisons between expression profile changes in young 3xTg-AD mice versus 3 m.o. WT mice revealed overexpression of several genes that are depicted in clusters B and C (Figures 1 and 4). These genes modulate several mechanisms (discussed below) that are serving a pathogenic role in AD.

**[Ca²⁺], homeostasis.** Deregulation of [Ca²⁺], is a key contributing factor for AD development and progression. Perturbations in Ca²⁺ handling by the endoplasmic reticulum (ER) and mitochondria as well as impaired intra/extracellular cation exchange contribute to the neuronal dysfunction and degeneration occurring in brain aging and AD. In cluster C, we found five genes involved in [Ca²⁺], homeostasis (ATP2B1, CACNB4, RYR2, PKD2, and NDUFA9). ATP2B1 encodes for the plasma membrane Ca²⁺-ATPase, a key system controlling high-capacitance/low-affinity Ca²⁺ extrusion, whereas CACNB4 encodes for the voltage-dependent L-type calcium channel subunit β-4, a critical route for Ca²⁺ influx. Altered expression and functionality of these proteins correlate with deregulation of [Ca²⁺], as well as with development of a seizure-prone phenotype. CACNB4 is also associated with neuroprotective mechanisms in animal models of ischemic brain injury. We also found changes in the expression of RYR2. RYR2 encodes for the ryanodine receptor 2, a major system controlling Ca²⁺ release in pre-symptomatic 3xTg-AD mice. This data set suggests a complex scenario in which enhanced [Ca²⁺] extrusion, mediated by ATP2B1, may compensate for impaired [Ca²⁺] buffering and excessive ER-Ca²⁺ release operated by RYR2.

**Mitochondrial functioning.** Alterations of mitochondrial functioning are crucial in physiological aging as well as in neurodegenerative conditions. In clusters B and C, we found transcripts involved in the modulation of mitochondrial morphology and metabolism (TRAK1 and Ndufs1 (cluster B); Hax1, Ndufa9, Gadap1, Kif1b, Hadbh, Pgam1, Ppia, and Dnm1l (cluster C)). Gadap1 and Dnm1l encode for proteins that promote mitochondrial fission; Pgam1 encodes for an enzyme of the glycolytic pathway that is involved in mitochondrial-mediated protective effects against Aβ-induced cell death; TRAK1 plays a critical role in axonal transport of mitochondria. Finally, Ndufs1 encodes for the NADH-ubiquinone oxidoreductase 75 kD, a subunit that regulates the activity of mitochondrial complex I. The early and sustained overexpression of transcripts involved in...
mitochondrial fission (GDAP1, and DNML1) and neuro-protection (PGAM1) is in line with recent findings indicating that mitochondrial fission is neuroprotective through reduction of oxidative damages.15

**Inflammatory response.** Neuroinflammation is detrimental in brain aging and age-related cognitive decline.16 The AD brain shows chronic, discrete, and microlocalized areas of inflammation as well as the appearance of activated microglia and reactive astrocytes in the vicinity of senile plaques.1 Moreover, inflammatory-related genes and proteins (i.e., CD33, NLRP3) play a substantial role in Aβ-driven neurodegeneration and AD progression.17,18 Cluster C revealed the presence of some transcripts (APP, IRF1, PPIA, VCAM1, TIMP2, HMGB1L1, PGAM1, and JAK1) that encode for proteins involved in the inflammatory response. It is noteworthy that in analogy with what was found for mitochondrial functioning, 3xTg-AD mice showed an early

### Table 3 IPA functional analysis of cluster C genes

| Category                                      | P-value   | Molecules                                                                 |
|-----------------------------------------------|-----------|---------------------------------------------------------------------------|
| Cell-to-cell signaling and interaction        | 4.94E-05-2.9E-02 | B2M, APBA2, VCAM1, GABRA5, USP14, JAK1, NRP2, MAPK1, KIF1B, CRK, HES1 (includes EG:15205), VDAC3, App, IRF1 (includes EG:16362), SYBU, PMP22, DCC, RSU1 |
| Cellular assembly and organization            | 4.94E-05-2.9E-02 | B2M, APBA2, FHL1 (includes EG:14199), YWHAH, GPR12, MAPK1, KIF1B, HAX1, LPPR4, BOK, SNX3, CRK, BECN1, App, WASL, MYRIP, STAR13, DCC, MAP1LC3B, NRN1, VCAM1, NRP2, MED1 (includes EG:19014), RYR2, SEPT7, SMARC5, YWHAZ, STX38L, MAPK9, GDAP1, DNM1L, VDAC3, App, TSG101, PPIA, S100B, TRRAP, Hmg1l1, HES1 (includes EG:15205), B2M, APBA2, VCAM1, MAPK1, GDAP1, HES1 (includes EG:15205), App, RCN1, VDAC3, App, TSG101, PPIA, Hmg1l1, DNM1L, NRN1 |
| Cell morphology                               | 6.66E-05-3.45E-02 | B2M, APBA2, VCAM1, KIF1B, HES1 (includes EG:15205), YWHAH, GPR12, MAPK1, GDAP1, PMP22, DCC, MAP1LC3B, NRN1, VCAM1, NRP2, MED1 (includes EG:19014), RYR2, SEPT7, SMARC5, YWHAZ, STX38L, MAPK9, GDAP1, DNM1L, VDAC3, App, TSG101, PPIA, S100B, TRRAP, Hmg1l1, HES1 (includes EG:15205), B2M, APBA2, VCAM1, MAPK1, GDAP1, HES1 (includes EG:15205), App, RCN1, VDAC3, App, TSG101, PPIA, Hmg1l1, DNM1L, NRN1 |
| Organismal development                        | 9.48E-05-2.9E-02 | PAFAH1B2, VCAM1, JAK1, MAPK1, NRP2, MED1 (includes EG:19014), PKD2 (includes EG:18764), MAPK9, BECN1, App, H2AFZ, TSG101, WASL, Hmg1l1, CACNB4, TIMP2 |
| Nervous system development and function       | 1.75E-04-3.24E-02 | B2M, APBA2, USP14, GABRA5, MAPK1, YWHAH, KIF1B, ITM2B, LPPR4, CRK, HES1 (includes EG:15205), App, SOX2, CACNB4, B2M, APBA2, VCAM1, MAPK1, GDAP1, HES1 (includes EG:15205), App, RCN1, VDAC3, App, TSG101, PPIA, Hmg1l1, DNM1L, NRN1 |
| Neurological disease                          | 2.83E-04-3.34E-02 | GABRA5, YWHAH, MAPK1, ATTP6V1C1, ATTP8B1, KIF1B, LPPR4, SLMAP, SOX2, PDHA1, TTB2, PGAM1, GABRB1, PKIA, PGRMC1, G3BP2, IPTP1, HIAT1, YWHAZ, SPARC1L, KIFAP3, CT3, TIMP2 |
| Cell signaling                                | 1.23E-03-3.34E-02 | G3BP2, JAK1, MAPK1, ATP2B1, PKD2 (includes EG:18764), RYR2, CACNB4, STX38L, MAPK9, CRK, KIFAP3 (includes EG:16363), SOX2, CLK1, TRIM13, TTB2, PPM1A, RSU1, TIMM8B, TIMP2 |
| Cellular compromise                           | 1.23E-03-3.45E-02 | VCAM1, MAPK1, SEPT7, PGAM1, CRK, GDAP1, DNM1L, TSG101, App, IRF1 (includes EG:16362) |
| Cell death                                    | 1.93E-03-3.45E-02 | PAFAH1B2, CA2, VCAM1, JAK1, MAPK1, KIF1B, MED1 (includes EG:19014), TSG101, PPIA, S100B, TRRAP, CACNB4, TIMP2 |
| Inflammatory response                         | 5.57E-03-3.34E-02 | VCAM1, JAK1, Hmg1l1, PPIA, PGAM1, App, IRF1 (includes EG:16362), TIMP2 |

### Table 4 IPA functional analysis of cluster D genes

| Category                                      | P-value   | Molecules                                                                 |
|-----------------------------------------------|-----------|---------------------------------------------------------------------------|
| Cell-to-cell signaling and interaction        | 9.74E-05-4.89E-02 | ST5, EPHB1, ARID1A, CCN3D, BAIAp2, RGS4, STAT3 |
| Nervous system development and function       | 9.74E-05-4.99E-02 | ISLR2, NEUROD2, EPB1, PRPF19, CCN3D, ACTB, S100B, BAIAp2, PDE4A, RGS4, STAT3, DIO2, KLK6, RN6, HIP1R |
| Organ development                             | 3.98E-04-4.59E-02 | EPHB1, CCN3D, STAT3, DIO2 |
| Cellular development                          | 3.87E-04-4.78E-02 | PRPF19, PAG24, ELAVL3, PDE4A, RGS4, STAT3, DIO2, B11, GPC1, NEUROD2, LMD1, CCN3D, S100B, BAIAp2, BRD4, UBF, TRRAP, SCAND1 |
| Cell cycle                                    | 8.76E-04-4.78E-02 | GPC1, MAP4, ARID1A, null, CCN3D, PAG24, EHMT2 (includes EG:10919), LAS1L, BRD4, STAT3, TRRAP |
| Lipid metabolism                              | 2.39E-03-3.47E-02 | GPC1, MAP4, null, COTL1, HIP1R |
| Cell death                                    | 3.14E-03-3.6E-02 | PRPF19, ARID1A, PAG24, ACTB, PDE4A, RGS4, STAT3, CAPN10, KLK8, B11, ITPK1, GPC1, MAP4, LMD1, CCN3D, S100B, VCP, UBF, TEOX2 |
| Cell morphology                               | 4.07E-03-4.39E-02 | ISLR2, MAP4, ACTB, BAIAp2, VCP, S100B, PDE4A, BRD4, RGS4, STAT3, TRRAP, RNF6, PRPF19, PAG24, ACTB, RGS4, STAT3, NAA10, B11, GPC1, LMD1, CCN3D, EIF4A1, S100B, BRD4, PTPRS, LASL1, UBF, TRRAP |
| Cell growth and proliferation                 | 4.07E-03-4.99E-02 | ISLR2, MAP4, ACTB, BAIAp2, VCP, S100B, PDE4A, BRD4, RGS4, STAT3, TRRAP, RNF6, PRPF19, PAG24, ACTB, RGS4, STAT3, NAA10, B11, GPC1, LMD1, CCN3D, EIF4A1, S100B, BRD4, PTPRS, LASL1, UBF, TRRAP |
| Neurological disease                          | 8.12E-03-3.21E-02 | S100B, STAT3 |
activation of upstream (IRF1) and downstream (JAK1) proinflammatory transcripts.

Additional genes involved in the inflammatory response (Hsp70, IL10 receptor, and CD200) were also found (cluster D). However, these transcripts were downregulated (instead of upregulated when comparing 3 m.o. 3xTg-AD mice with 3 m.o. WT mice). Among these, Hsp70 and CD200 are known to play an anti-inflammatory role. Thus, the overexpression of inflammatory genes appears to be coupled with simultaneous decrease in the expression of anti-inflammatory transcripts, thereby suggesting an overall proinflammatory status.

Figure 3 Validation of microarray gene expression data by qRT-PCR. Bar graph shows mRNA levels of CST3, GABRA5, and BECN1 measured by real-time PCR in young and old WT mice and 3xTg-AD mice as well as in old WT mice and 3xTg-AD mice. Data are expressed as mean values of relative fold changes ± S.D. of three independent experiments performed in triplicate. The symbol * indicates that CST3, GABRA5, and BECN1 are significantly upregulated when compared with gene expression in 3 m.o. WT mice (P<0.05). The symbol # indicates that CST3, GABRA5, and BECN1 are significantly upregulated when compared with expression of these genes in 12 m.o. WT mice (P<0.01).

Dendrite formation, synaptic plasticity, and memory. Age- and AD-related cognitive decline is associated with synaptic dysfunction. In that respect, we found overexpression of several transcripts (CD200, Neurod1, SOX9, SNAP25, EGR1, OPHN1, ET1, CDH2, GABRA1, and Spond1 (cluster B): APBA2, APP, GABRA5, MAPK1, PMP22, SYBU, USP14, VDAC3, CST3, DCC, HES1, HEY1, HPRT1, SOX2, VWA5H, CRK, GPR12, HMGB1L1, STK38L, and VCAM1 (cluster C)) that are implicated in dendritic spine formation, modulation of synaptic functioning, and cognitive decline. Overexpression of transcripts encoding for proteins involved in synaptic activity and neurotransmission may be interpreted as a protective event against AD-related synaptic dysfunctions.

Overexpression of transcripts present in cluster C suggests a bimodal set of activities. One set of actions may counteract neuronal damage and promote protective effects aimed at maintaining cognitive abilities. For instance, APBA2 is known to bind to APP and reduce Aβ production, thereby counteracting memory deficits and promoting long-term potentiation (LTP) in Tg AD mice. Similarly, CST3 up-regulation is protective against the neurotoxic effects of Aβ oligomers and oxidative stress damage. This phenomenon has been reported in Tg AD models, in the brains of AD patients, and in those of elderly individuals. The GABRA5 transcript increase found in our 3xTg-AD mice lends support to the compensatory hypothesis as GABRA5 has been reported to be downregulated with aging and this process has been linked to defective spatial memory performance.

These protective actions are counteracted by the concurrent up-regulation of transcripts that, on the contrary, are associated with neuronal dysfunction. This is the case of APP and ERK2 transcripts. Given the aminoacidic composition of murine Aβ, murine APP is marginally involved in the formation of toxic Aβ species, but some studies have also proposed that the protein can favor ERK1,2 activation, tau phosphorylation, and the formation of NFTs. Chronic ERK2 activation by Aβ oligomers can be detrimental as early and sustained ERK2 stimulation has been shown to promote cognitive decline in AD mice and induce neuronal toxicity through caspase-3 activation.

Figure 4 Graphical synopsis of cluster analysis results. Synoptic view of all the transcriptome profiles that were found altered in the three experimental conditions. The main findings associated with observed gene expression changes are summarized. Note that, as depicted by the dark gray boxes, 3 m.o. 3xTg-AD versus 3 m.o. WT mice as well as 12 m.o. WT versus 3 m.o. WT mice show a similar pattern of expression changes in clusters B, C, and D. Also note that, only in the case of genes of cluster C, a common pattern of expression profiles is observed across all the three experimental conditions.
It is interesting to note that in clusters A and D, we found downregulation of transcripts (PEBP1 and Complexin III (cluster A) and BIN, ACTB, BAIAP2, EphB, KLK8, ARD1, PDE4, Ehmt2, Neurod2, STAT3, S100B, and CCND3 (cluster D)) that are involved in dendritic spine formation, modulation of synaptic functioning, and cognitive decline. In this respect, PEBP1 and Complexin III downregulation is associated with impaired cholinergic neurotransmission (PEBP1) and synaptic transmission deficits (Complexin III), the two mechanisms largely impaired in AD and in mild cognitive impairment. BAIAP2, EphB1, BIN1, and KLK8 are involved in synaptic plasticity. BAIAP2 (also known as IRSp53) modulates NMDA receptor-related synaptic transmission, LTP, as well as learning and memory. BAIAP2 knockout mice show impaired spatial learning and novel object recognition. Thus, BAIAP2 downregulation might be involved in the development of cognitive deficits affecting this model that has been also shown to suffer from significant decrease in BDNF levels. NMDAR deregulation has been associated with EphB (a family of ligand proteins) receptor deficits and linked to the appearance of neurological disorders including AD. Still, in the set of synaptic plasticity-related genes that we found underexpressed, KLK8 seems important as the gene is involved in brain development, neuronal plasticity, and stress response. Thus, KLK8 downregulation can favor aberrant CNS development. S100B is a calcium-binding protein released by astroglial cells. Compelling evidence indicates that transgenic animals, overexpressing S100B, show neuronal loss and increased expression of the pro-apoptotic protein clusterin, a protein that is increased in hippocampi and frontal cortices of AD patients. In summary, our findings suggest that 3xTg-AD mice not only show early modulation of transcripts that exert detrimental effects on synaptic functioning but also develop changes that counteract this impairment. The process appears to go on chronically as the same pattern is present in 12 m.o. 3xTg-AD mice.

**Neuronal death, cell cycle, and autophagy.** Aberrant expression of cell cycle- and autophagy-related molecules plays a pivotal role in the neuronal loss associated with AD and brain aging. Our analysis revealed overexpression of a group of transcripts (SPIN1, JAK3, Neurod1, SOX9, YWHAE, EGR1, and HSPA1A/HSPA1B (cluster B); Beclin-1, MAP1LC3-B, HADHB, AK5, CyC1, SIRT7 and ENC1 (cluster C)) involved in autophagy, cell cycle, and cell death, thereby suggesting the potential activation of a complex network of neuronal death-related responses occurring in young 3xTg-AD mice. Early EGR1 upregulation has been reported in AD patients and can contribute, along with other cell cycle-related genes, to NFT formation, a phenomenon that is also mediated by the increased expression of cell cycle proteins. Furthermore, overexpression of SPIN1 (a gene involved in cell cycle impairment) promotes chromosome instability, a process leading to cellular senescence and apoptosis. Sirt7, a less investigated member of the sirtuin family, is a nuclear protein expressed in the brain with a cellular distribution and functions that are still unknown. Experiments employing Sirt7 knockout mice or Sirt7-overexpressing cells have indicated the antiproliferative role of the protein and suggested that Sirt7 activity can improve tissue integrity in aging animals. Beclin-1 and MAP1LC3-B are two autophagy-related genes whose overexpression might counteract pathology development in 3 m.o. 3xTg-AD mice. This hypothesis is consistent with recent findings indicating that early induction of autophagy reduces cognitive decline in 3xTg-AD mice. Autophagy, in fact, clears out damaged proteins that are prone to aggregation, and the process can be neuroprotective by mediating the degradation of misfolded proteins. Intriguingly, in AD, autophagy can play a dual role. On one hand, it is protective by promoting the removal of Aβ and tau aggregates and, on the other hand, autophagosomes are nevertheless sites of choice for accumulation and production of Aβ. Analysis of networks of genes involved in neuronal death also showed several transcripts that, compared with young WT mice, were found downregulated in young 3xTg-AD mice (ARID1A, BRD4, CCND3, EHMT2, GPC1, LASS1L, MAP4, PA2G4, PTGES3, STAT3, and TRRAP (cluster D)). Interestingly, aberrant expression of CCND3 has been reported to be closely associated with AD-related neuronal death occurring in specific hippocampal regions. Intriguingly, in AD, autophagy can play a dual role. On one hand, it is protective by promoting the removal of Aβ and tau aggregates and, on the other hand, autophagosomes are nevertheless sites of choice for accumulation and production of Aβ. Analysis of networks of genes involved in neuronal death also showed several transcripts that, compared with young WT mice, were found downregulated in young 3xTg-AD mice (ARID1A, BRD4, CCND3, EHMT2, GPC1, LASS1L, MAP4, PA2G4, PTGES3, STAT3, and TRRAP (cluster D)). Interestingly, aberrant expression of CCND3 has been reported to be closely associated with AD-related neuronal death occurring in specific hippocampal regions.

**12 m.o. 3xTg-AD versus 12 m.o. WT mice.** In order to evaluate how AD-like progression affects gene expression, we investigated profiles of 12 m.o. 3xTg-AD mice compared with 12 m.o. WT mice and matched this set of data with results of 3 m.o. 3xTg-AD mice that were compared with age-matched WT mice. The analysis revealed a similar overexpression of all transcripts of cluster C and downexpression of those of cluster A. Interesting differences appeared when analyzing transcripts belonging to clusters B and D (Figures 1 and 4).

Cluster B transcripts, overexpressed in young 3xTg-AD mice, were downregulated in 12 m.o. 3xTg-AD mice when compared with age-matched WT mice, finding that supports the hypothesis that aged 3xTg-AD mice may show deficits in counteracting disease progression. Underexpression of many transcripts found in cluster B has been previously reported in several models of aging and AD as well as in AD patients. This is the case for SNAP25 (encoding for a key protein involved in synaptic vesicle cycle and neurotransmission modulation), GABRA1 (encoding for a γ-aminobutyric acid (GABA) receptor subunit downregulated in AD brains), CDH2 (encoding for N-cadherin, whose deregulation triggers apoptotic pathway activation), and OPHN1 (encoding for a Rho-GTPase involved in controlling excitatory synapses maturation and plasticity). Cluster D transcripts, downexpressed in 3 m.o. 3xTg-AD, were instead overexpressed in aged 3xTg-AD. The upregulation of transcripts that are involved in the modulation of synaptic plasticity, neuronal proliferation, survival, differentiation, and regulation of cell division process might reflect a compensatory neuroprotective response (BAIAP2), although the net result suggests a more complex scenario (see, for instance, the role of KLK8). BAIAP2 downregulation is related to altered synaptic plasticity as observed in
BAIAP2-deficient mice, whereas its overexpression increases dendritic spine density and postsynaptic potentiation, ultimately enhancing cognition. KLK8 deficiency promotes decreased synaptic plasticity, whereas, on the contrary, gain in KLK8 mRNA levels has been observed in AD brains and equally associated with detrimental effects on hippocampal functioning. Inhibition of S100B has positive effects on the Aβ load and glosis observed in AD patients, whereas S100B overexpression has been associated with exacerbation of AD clinical signs. Finally, altered expression of cell cycle proteins (i.e., CCND3) in postmitotic neurons has been observed both in aging and in AD.

Collectively, over- and underexpression of the aforementioned transcripts confirm a pattern of responses in which some genes exert detrimental effects that are in line with the development of AD-like pathology. At the same time, other transcripts appear to counteract these effects. Interestingly, this process appears to progress chronically as a similar pattern has been reported in 3 m.o. and in 12 m.o. 3xTg-AD mice.

12 m.o. WT versus 3 m.o. WT mice. Finally, we compared expression profiles of 12 m.o. WT mice with 3 m.o. WT mice to analyze the effects of physiological aging.

Interestingly, changes driven by aging were largely overlapping with those observed in 3 m.o. 3xTg-AD mice when compared with 3 m.o. WT mice. We, in fact, observed overexpression of several transcripts belonging to clusters B and C and downregulation of a gene data set present in cluster D (Figures 1 and 4). The functional correlates of these changes have been discussed above.

Overall, these data support the hypothesis that the pro-AD environment offered by the 3xTg-AD mice accelerates and promotes early modifications of genes that are otherwise changed by aging in old WT mice. Moreover, comparison between 12 m.o. WT mice and age-matched 3xTg-AD mice showed expression changes that were similar only for genes belonging to cluster C.

Interestingly, Ingenuity Pathway Analysis (IPA) analysis of this cluster revealed transcripts (HSPA12A, HIAT1, KIAA1409, NRP2, STARD13, UBB, and PKIA) that genome-wide association studies have recently shown to be involved in AD-related single-nucleotide polymorphisms (SNPs) as well as transcripts (APP, ATP6V1C1, BECN1, CST3, DCC, DNM1L, GABRA5, GABRB1, HIAT1, HSPA12A, KIAA1409, LPPR4, MAPK9, NRP2, PKIA, RYR2, STARD13, UBB, and YWHAZ) that are closely related to AD development.

Finally, cluster A is the only one showing a unique behavior in 3xTg-AD mice, indicating changes that are specifically triggered by the pro-AD genetic background of the mice. It is intriguing to hypothesize that these genes have a functional role in AD progression and represent potential AD biomarkers.

Among these, we observed a downregulation of Homer3 and Dysferlin. Homer3 interacts with APP and inhibits Aβ production. Thus, Homer3 downregulation can lead to enhanced Aβ production. Accumulation in senile plaques of Dysferlin, a protein involved in membrane repair, leads to defective neuronal repair. Thus, it is conceivable that the Dysferlin downregulation that we found in the 3xTg-AD mice can exert a similar detrimental effect.

Conclusions

With the limitation of any Tg-AD model, our data support the hypothesis that aging and AD share some mechanistic similarities. In our Tg-AD mouse, the pro-AD environment promotes and accelerates changes that, upon aging, are otherwise occurring in WT animals. Although the 3xTg-AD mice harbor three mutations that do not occur, at the same time, in AD human brains, this cumulative pro-AD background modulates in Tg mice the early on expression of genes, the so-called BioAge genes (associated with neuronal loss, glial activation, and lipid dysmetabolism) that are in humans biomarkers for brain aging and found prematurely expressed in AD patients. The same accelerated expression of aging-related genes in AD has been recently reported in a study that analyzed AD brains, thereby providing support for the heuristic value of 3xTg-AD mice as preclinical model of AD.

Our data support and integrate the age-based hypothesis for AD, a concept that postulates that aging is required to set in motion the AD-related pathogenic cascade. Our findings suggest the presence of a feed-forward mechanism by which pro-AD factors can promote aging-related gene profile changes, thereby boosting and accelerating the disease process. Furthermore, this is the first study that offers an extensive analysis of the transcriptomic profile of the 3xTg-AD model, one of the most comprehensive preclinical models of the disease. The novel set of AD-related genes found in our analysis will, likely, further studies investigating the molecular roadmap of the disease and its modulation by aging.

Materials and Methods

Animal models. All the procedures involving animals were approved by the institutional ethics committee (CeSI protocol n.: AD-301) and performed according to institutional guidelines and in compliance with national (DL no. 116, GU, Suppl. 40, 18 February 1992) and international laws and policies. The 3xTg-AD (harboring AD-related human mutation APP (Swe), PS1 (M146V), and tau (P301L) transgenes) female mice (n = 2 at 3 m.o.a.; n = 2 at 12 m.o.a) and age-matched female control WT mice (129SV × C57BL/6, n = 2 at 3 m.o.a.; n = 2 at 12 m.o.a.) were kept in a temperature-controlled room at 25°C under a 12-h light/dark cycle and fed ad libitum with tap water and commercial chow. Mice at 3 or 12 m.o.a. were anesthetized and killed by decapitation, hippocampi excised, tissue transferred into RNA-later solution, and stored at −80°C for RNA processing.

Microarray analysis. Hippocampi were homogenized using a hand glass potter and total RNA extracted using the SVTotal RNA Isolation System kit following the manufacturer’s instructions (Promega, Madison, WI, USA). RNA purity and quantity were estimated using an Agilent 8453 spectrophotometer (Agilent, Santa Clara, CA, USA). RNA samples were not pooled. Using the Amino Allyl MessageAmp II aRNA Amplification Kit (Ambion, Grand Island, NY, USA) RNA was amplified and fluorescently labeled with Cy3-Cy5 fluorescent signals were detected with a confocal laser scanner ‘ScanArray Express’ (Packard BioScience, Waltham, MA, USA) and analyzed with the ‘ScanArray Express–MicroArray Analysis System 3.0’ software (Perkin Elmer, Waltham, MA, USA). Raw data are stored in the GEO public database (accession number: GSE35210). Raw data were normalized using locally weighted scatterplot smoothing (LOWESS) algorithm. A transcript was considered differentially expressed when showing an absolute log-ratio value of ≥ ±0.5, that is, 1.4 fold-change in transcript quantity. The transcript data set
was analyzed with gene clustering analysis using Cluster 3.0 (Stanford University Labs, Stanford, CA, USA). We evaluated only spots showing significant expression in at least 80% of the experiments. Data were centered to the mean of each mRNA and unsupervised average linkage hierarchical clustering of both the mRNAs and samples was performed using uncentered correlation metrics. Dendrograms and selected transcripts were visualized by employing TreeView software (Stanford University Labs). Gene biological functions were inferred using IPA software (Ingenuity Systems, Redwood City, CA, USA).

Real-time PCR. Microarray results were validated by performing qRT-PCR analysis on RNA extracted from hippocampi obtained from the same samples used for microarray analysis. Gene expression of three genes found upregulated by microarray (BECN1, CST3, and GABRA5) was evaluated. The GAPDH housekeeping gene was employed as internal control. qRT-PCR was performed in a total volume of 50 μl containing: 1 × TaqMan Universal PCR Master Mix, no AmpErase UNG, and 2 μl of cDNA using TaqMan assay on an ABI 7900HT Sequencing Detection System (Applied Biosystems, Paisley, UK). Transcript primers and probe sets employed were: GAPDH: primers FW: CTTTGGCAAGCTTACCTGGG, probe: CACCTGGTTGCTGTAAGCGGGGCCATTGATTA; BECN1: primers FW: GTACCAGCTTGGCTCATTAGG, probe: ACACAGTCGAGAAACTACG, probe: CCCCCAGAAGATATCCGAGCCTACCC; CST3: primers FW: CTGTGACCTCTGGTCTCATGAC, probe: CCTTGCTCAGTGTCCTTGC, probe CACCCTGTTGCTG. Gene biological functions were inferred using IPA software (Ingenuity Systems, Redwood City, CA, USA).

Conflict of Interest

The authors declare no conflict of interest.

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