Increased MANF Expression in the Inferior Temporal Gyrus in Patients With Alzheimer Disease

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Alzheimer disease (AD) is an aging-related disorder linked to endoplasmic reticulum (ER) stress. The main pathologic feature of AD is the presence of extracellular senile plaques and intraneuronal neurofibrillary tangles (NFTs) in the brain. In neurodegenerative diseases, the unfolded protein response (UPR) induced by ER stress ensures cell survival. Mesencephalic astrocyte-derived neurotrophic factor (MANF) protects against ER stress and has been implicated in the pathogenesis of AD. MANF is expressed in neurons of the brain and spinal cord. However, there have been no investigations on MANF expression in the brain of AD patients. This was addressed in the present study by immunohistochemistry, western blotting, and quantitative analyses of postmortem brain specimens. We examined the localization and expression levels of MANF in the inferior temporal gyrus of the cortex (ITGC) in AD patients (n = 5), preclinical (pre-)AD patients (n = 5), and age-matched non-dementia controls (n = 5) by double immunofluorescence labeling with antibodies against the neuron-specific nuclear protein neuronal nuclei (NeuN), ER chaperone protein 78-kDa glucose-regulated protein (GRP78), and MANF. The results showed that MANF was mainly expressed in neurons of the ITGC in all 3 groups; However, the number of MANF-positive neurons was significantly higher in pre-AD (Braak stage III/IV) and AD (Braak stage V/VI) patients than that in the control group. Thus, MANF is overexpressed in AD and pre-AD, suggesting that it can serve as a diagnostic marker for early stage disease.

Keywords: Alzheimer disease, cerebral cortex, endoplasmic reticulum stress, MANF, hyperphosphorylated tau, senile plaque

INTRODUCTION

Alzheimer disease (AD) is a progressive neurodegenerative disease with insidious onset (Mathys et al., 2019) that constitutes a major public health burden (Scheltens et al., 2016; Alzheimer’s Association, 2020). The prevalence of AD among people over the age of 65 years is estimated to be 10–30%, which is increasing with the aging of the global population (Masters et al., 2015). AD is characterized by a progressive decline in cognitive function and neuronal loss (Pietronigro et al., 2016). There are two types of brain lesion that are the pathologic hallmarks of AD: extracellular...
TABLE 1 | Demographic and clinicopathologic information of the samples.

| NBB no. | Autopsy  | Braak stage | Group  | Age (years) | Sex | PMD (h) | BW (g) | CSF pH |
|---------|----------|-------------|--------|-------------|-----|---------|--------|--------|
| 15033   | S15/033  | 0           | A      | CON         | 93  | M       | 07:40  | 1155   | 6.20  |
| 14043   | S14/043  | 0           | O      | CON         | 60  | F       | 08:10  | 1310   | 6.58  |
| 01045   | S01/115  | I           | B      | CON         | 83  | M       | 04:35  | 1367   | 6.49  |
| 02018   | S02/043  | I           | B      | CON         | 92  | F       | 07:00  | 1193   | 6.45  |
| 04026   | S04/074  | I           | B      | CON         | 91  | F       | 07:45  | 1054   | 6.90  |
| 14014   | S14/014  | III         | B      | Pre-AD      | 90  | F       | 06:05  | 1255   | 6.12  |
| 14020   | S14/020  | III         | O      | Pre-AD      | 92  | F       | 06:35  | 1305   | 6.12  |
| 95059   | S95/140  | IV          | NA     | Pre-AD      | 86  | F       | 03:20  | 995    | 7.14  |
| 08075   | S08/241  | IV          | C      | Pre-AD      | 88  | M       | 05:00  | 1296   | 6.45  |
| 09096   | S09/301  | IV          | C      | Pre-AD      | 92  | M       | 08:25  | 1117   | 6.14  |
| 97015   | S97/045  | V           | NA     | AD          | 85  | F       | 03:10  | 1044   | 6.90  |
| 00054   | S00/115  | V           | C      | AD          | 59  | M       | 07:45  | 1171   | 6.29  |
| 00119   | S00/264  | V           | C      | AD          | 85  | F       | 06:10  | 1003   | 6.65  |
| 11121   | S11/121  | V           | NA     | AD          | 95  | M       | 07:00  | 1143   | 6.18  |
| 02069   | S02/203  | VI          | C      | AD          | 70  | F       | 08:15  | 876    | 6.30  |

AD, Alzheimer disease; BW, brain weight; CON, non-dementia control; CSF, cerebrospinal fluid; F, female; M, male; NA, not available; NBB, Netherlands Brain Bank; NFT, neurofibrillary tangles; PMD, postmortem delay; pre-AD, preclinical Alzheimer disease.

senile plaques composed of amyloid β-peptide (Aβ), and intraneuronal neurofibrillary tangles (NFTs) consisting of paired helical filaments of hyperphosphorylated tau protein (Braak and Del Tredici, 2018). AD patients exhibit different degrees of nucleolar pyknosis in neurons or even the disappearance of neurons. The pathogenesis of AD has been linked to dysfunction of the endoplasmic reticulum (ER) (Mukherjee and Soto, 2011; Cabral-Miranda and Hetz, 2018), which is the site of protein folding and secretion in eukaryotic cells (Gerakis and Hetz, 2018). ER stress, which is induced by the accumulation of unfolded or misfolded proteins in the ER (Ghemrawi and Khair, 2020), has been proposed as a mechanism underlying Aβ-induced Alzheimer-like neuropathology (Goswami et al., 2020). Although homeostatic mechanisms such as the unfolded protein response (UPR) can restore normal ER function (Rahman et al., 2018), prolonged ER stress can lead to cell dysfunction and death (Bravo et al., 2013; Lu et al., 2014).

Mesencephalic astrocyte-derived neurotrophic factor (MANF) – originally named arginine-rich protein (ARP) or arginine-rich mutated in early tumors (ARMET) – is an evolutionarily conserved secreted protein expressed in the rodent brain that has been shown to play a protective role in the response to ER stress (Apostolou et al., 2008; Wang et al., 2014). Like cerebral dopamine neurotrophic factor (CDNF), MANF is classified as a neurotrophic factor (Lindholm and Saarma, 2010; Lindahl et al., 2017) that participates in the UPR (Apostolou et al., 2008). MANF was shown to rescue neurons from apoptosis and ER stress (Hellman et al., 2011), and knocking down MANF expression induced the UPR and increased the neurotoxic effects of Aβ (Xu et al., 2019). Additionally, chronic activation of the UPR in the brain has been reported in MANF-deficient mice (Pakarinen et al., 2020). Glucose-regulated protein (GRP)78 is an ER stress-associated marker (Sakono and Kidani, 2017) and UPR-regulated chaperone that interacts with MANF (Yan et al., 2019). In general, MANF is upregulated and plays a protective role in the response to ER stress (Xu et al., 2019). MANF exerted neuroprotective effects against ethanol-induced neurodegeneration by alleviating ER stress, which may be relevant to other ER stress-related neurodegenerative diseases (Wang Y. et al., 2021).

MANF has shown protective effects in animal models of AD (Xu et al., 2019), Parkinson disease (Voutilainen et al., 2009), spinocerebellar ataxia (Yang et al., 2014), ischemic brain damage (Airavaara et al., 2010), retinal degeneration (Lu et al., 2018), cardiac ischemia (Arrieta et al., 2020), and liver injury (Sousa-Victor et al., 2019). Given the diverse pathologies that MANF can alleviate, its involvement in diseases related to the activation of the UPR is expected. However, to date there have been no reports on the expression of MANF in the brain of AD patients.

The inferior temporal gyrus of the cortex (ITGC) is a key brain area involved in cognitive functions including memory, auditory cognition, and semantics (Meunier and Barbeau, 2013). The ITGC plays an important role in verbal fluency, which is affected soon after the onset of AD (Scheff et al., 2011). The hippocampus, a region severely affected in AD, is connected to the ITGC (Mégevand et al., 2017). To address the above point, the present study examined the subcellular localization and expression of MANF in the ITGC of human brain specimens from pre-AD and AD patients in order to clarify its role in AD pathogenesis.

MATERIALS AND METHODS

Brain Specimens

Paraffin-embedded postmortem human brain tissue specimens were provided by the Netherlands Brain Bank (NBB) (Amsterdam, Netherlands). The brains were donated to
FIGURE 1 | p-tau and Aβ17-24 expressions in the ITGC detected by immunohistochemistry. (A–H) p-tau and Aβ17-24 immunoreactivity in control subjects (left) and AD patients (right). Patients with AD showed extensive accumulation of Aβ plaques [arrowheads in panels (F,H)] and NFTs [arrowheads in panels (B,D)] in the ITGC. Scale bar, 50 µm (A,B,E,F), 20 µm (C,D,G,H).

research after patients or their closest relatives provided written, informed consent. AD pathology was evaluated according to Braak stage (Braak and Braak, 1991). Ten AD cases (age range, 59–95 years; male-to-female ratio, 2:3) and 5 sex- and age-matched control cases (age range, 60–93 years; male-to-female ratio, 2:3) were included in the analysis. The AD cases were further classified into pre-AD (Braak stages: III/IV) and AD (Braak stages: V/VI) (n = 5 each). Non-dementia control cases (Braak stage: 0/I) had no known clinical history of dementia, and the cause of death was unrelated to the central nervous system. Demographic and clinicopathologic data for the samples are shown in Table 1.
FIGURE 2 | Mesencephalic astrocyte-derived neurotrophic factor levels detected by western blotting with a polyclonal antibody. Cell lysis solutions and human brain temporal cortex protein extracts were probed with a polyclonal anti-MANF antibody to evaluate specificity. The immunoblot shows a clear single band of ∼18 kDa in all protein extracts and cell lysis solutions.

Tissue Preparation
Paraffin blocks containing the ITGC were stored at room temperature under protection from light. Serial coronal sections were cut on a microtome (RM2235; Leica, Wetzlar, Germany) at a thickness of 6 µm and stored at room temperature. We selected 1 section every 150 µm (or approximately every 25 sections) for a total of 45 sections from the ITGC, as well as 1 section each from the anterior, middle, and posterior cortices. Thus, 45 brain tissue sections were used for MANF immunohistochemistry from pre-AD, AD, and non-dementia control specimens for quantification of MANF-immunoreactive neurons in the ITGC. Additionally, in order to detect pathologic lesions (Aβ and tau) and determine whether MANF is primarily expressed in neurons and determine its subcellular localization, several adjacent sections were randomly selected from pre-AD, AD, and non-dementia control.

Specific Assessment of MANF Antibodies
To ensure that the rabbit polyclonal anti-MANF antibody used in this study (ARMET/ARP; cat. no. Ab67271; Abcam, Cambridge, United Kingdom) could detect MANF in human brain specimens, we tested its specificity in cell lysates of HEK 293T and SHSY5Y cell line and homogenates of human brain temporal cortex tissue by western blotting under denaturing conditions as previously described (Huang et al., 2020). The SHSY5Y (Catalog number TCHu97), HEK293T (Catalog number GNHu17) cells were purchased from the typical culture preservation committee of Chinese academy of sciences. To further confirm the specificity of the anti-MANF antibody, the levels of MANF expression in hepatocellular tissues of hepatocyte-specific MANF-knockout (HKO) control mice and wild-type (WT) control mice were measured by immunohistochemistry. In previous studies, the efficiency of MANF knockout from the HKO control mice have been detected with the use of western blotting (Yang et al., 2021). The hepatocellular tissue sections of the HKO control mice and WT control mice were provided by Prof. Yuxian Shen of the Anhui Medical University.

Immunohistochemistry
Brain sections were deparaffinized to water according to standard procedures. After washing with phosphate-buffered saline (PBS; pH value 7.2–7.4; 3 × 10 min), antigen retrieval was performed by boiling the sections in citric acid buffer and steaming for 1 min 15 s. The sections were left to cool at room temperature for about 30 min, then flushed with warm water (50°C) for 5 min and washed with PBS (3 × 10 min). Endogenous peroxidase activity was quenched by incubating the sections in a solution of 3% H2O2 and methanol for 10 min at 25°C. The tissue was blocked in 10% sheep serum albumin solution at 37°C for 1 h, followed by overnight incubation at 4°C with rabbit polyclonal anti-MANF antibody at 1:400 dilution. The following day, the sections were warmed at room temperature for 15 min in a covered humid chamber, then washed in PBS (3 × 10 min) before incubation with biotin-conjugated goat anti-rabbit IgG for 15 min at room temperature (≥25°C). After washing in PBS

TABLE 2 | Mesencephalic astrocyte-derived neurotrophic factor expression in the inferior temporal gyrus cortex in pre-AD and AD patients and non-dementia controls.

| Variable                        | AD               | Pre-AD          | Control          |
|---------------------------------|------------------|-----------------|------------------|
| Diameter of nucleoli (µm)       | 3.68 ± 0.77**    | 3.70 ± 0.82**   | 4.16 ± 1.22      |
| Cytoplasmic MANF-positive neurons/1 mm² | 115.81 ± 21.24** | 116.58 ± 21.99** | 100.85 ± 19.91  |

Values represent mean ± standard deviation; **P < 0.05 vs. control group (two-way analysis of variance). AD, Alzheimer disease; MANF, mesencephalic astrocyte-derived neurotrophic factor; pre-AD, preclinical Alzheimer disease.
FIGURE 3 | Mesencephalic astrocyte-derived neurotrophic factor levels in the HKO control mice and WT control mice detected by immunohistochemistry with polyclonal antibodies. No positive staining was detected in the hepatocellular tissues of the HKO control mice (A,B). MANF was detected in the hepatocellular tissues of the WT control mice (C,D). Scale bar: 50 µm (A,C) and 20 µm (B,D).

(3 × 10 min), diaminobenzidine reagent was added dropwise, and the colorimetric reaction was allowed to proceed for 1–1.5 min. The sections were washed and then stained with hematoxylin, dehydrated through a graded series of alcohol and xylene, and mounted with neutral gum for observation under a light microscope (CX43; Olympus, Tokyo, Japan). Immunohistochemical detection of tau and Aβ was performed using monoclonal antibodies against phosphorylated (p-)tau (Ser202, Thr205) (AT8; Thermo Fisher Scientific, Waltham, MA, United States; cat. no. MN1020) and Aβ17–24 (4G8; Biolegend, San Diego, CA, United States; cat. no. 800701), both used at 1:200 dilution. Immunohistochemistry was performed on liver sections as previously described (Wang P. et al., 2021).

To determine whether MANF is mainly expressed in neurons and to determine its subcellular localization, we performed double immunofluorescence labeling of MANF/neuronal nuclei (NeuN) and MANF/GRP78 as previously described (Shen et al., 2012; Gao et al., 2018; Herranen et al., 2020), with minor modifications to the protocol. The following antibodies were used: rabbit polyclonal anti-MANF (1:400 dilution); mouse monoclonal anti-NeuN (clone A60, cat. no. MAB377; Abcam) (1:400 dilution); and rabbit polyclonal anti-GRP78 (GRP78 BiP, cat. no. ab21685; Abcam) (1:200 dilution). Nuclei were stained with 4',6-diamidino-2-phenylindole (cat. no. C1005; Beyotime, Shanghai, China). The specificity of these antibodies has been reported in previous studies (Zhu et al., 2007; Shen et al., 2012; Braak and Del Tredici, 2018; Adaikkan et al., 2019; Xu et al., 2019).

Measurements
Images of the brain sections were obtained with a digital slide scanner (Pannoramic MIDI; 3DHISTECH, Budapest, Hungary) and Case Viewer software (3DHISTECH). Quantitative analysis of the density of neurons expressing MANF in the cytoplasm was carried out using ImageJ software (Schneider et al., 2012). The diameters of the region of interests (ROIs) in nucleoli were measured with MetaMorph software (Molecular Devices, San Jose, CA, United States). The methods used for region selection and measurement have been described elsewhere (Hu et al., 2002, 2003; Thangavel et al., 2008). Briefly, the cortical selection in each tissue section was defined in the 5 ROIs (size: 600 × 317 µm) on the digitized autoradiograms using Case Viewer software, and the area was measured. Each selection was drawn from the cortical surface extending perpendicularly to the gray and white matter boundary. Up to 5 selections were defined for each ROI but in some ROIs, the number of selections was limited by loss of tissue integrity. In each image of the ITGC acquired under high magnification (40× objective), the average optical density of immunopositive areas was calculated for each visual field; the density of neurons expressing MANF in the cytoplasm was determined. Additionally, the diameter of nucleoli of nucleolated neuronal profiles from 5 ROIs per ITGC tissue section under high magnification (40× objective) was measured by an investigator who was blinded to the clinicopathologic data of the subjects.

Statistical Analysis
Statistical analyses were performed using SPSS v22.0 (SPSS Inc., Chicago, IL, United States). Data are expressed as mean ± standard deviation. Based on Shapiro–Wilk test, almost all data were skewed, so non-parametric tests were used. Differences between groups were evaluated with the Kruskal–Wallis H test for multiple comparisons. Two-way analysis of variance was used to compare MANF expression in the ITGC...
among pre-AD, AD, and non-dementia control cases. \( P < 0.05 \) was considered significant.

### RESULTS

#### Clinicopathologic Information of pre-AD, AD, and Control Cases

The clinicopathologic information for the study population is summarized in Table 1. There were no differences in age \( (P = 0.428) \), postmortem delay \( (P = 0.650) \), body weight \( (P = 0.125) \), and cerebrospinal fluid pH \( (P = 0.404) \) among pre-AD and AD patients and control subjects. Immunohistochemical analysis revealed that p-tau immunoreactivity was predominantly in the ITGC and was higher in AD patients (NBB no. 00119; Autopsy S00/264) compared to control subjects (NBB no. 01045; Autopsy S01/115). A\( \beta_{17−24} \) (4G8) expression was also higher in AD cases (NBB no. 11121; Autopsy S11/121) than in controls (NBB no. 15033; Autopsy S15/033) (Figure 1).

#### Specificity of the Anti-MANF Antibody

We evaluated the specificity of the polyclonal anti-MANF antibody by western blotting using cell cultures and human brain temporal cortex extracts. The antibody recognized a single band at \( \sim 18 \text{ kDa} \) – which is the known molecular weight of MANF protein (Figure 2). Furthermore, we have not detected MANF expression in hepatocellular tissues of HKO control mice.
FIGURE 5 | Representative images of the expression of MANF and GRP78 in the ITGC. (A–H) The pictures above are representative images of immunofluorescence labeling for MANF (red) and GRP78 (green). The nuclei were stained with DAPI (blue). Picture D is a merged image of panels (A–C). The magnified images show the expression and cellular distribution of MANF and GRP78, respectively. Scale bar: 50µm (A–D) and 10µm (E–H).

by immunohistochemical staining, compared with WT control mice (Figure 3).

Nucleoli of ITGC Neurons Are Reduced in Size in Pre-AD and AD
The diameter of neuronal nucleoli of ITGC neurons was significantly smaller in pre-AD and AD patients than in control subjects (P < 0.05; Table 2).

Distribution of MANF-Positive Neurons in the ITGC and Subcellular Localization of MANF
MANF expression was detected in the ITGC of human brain specimens (NBB no. 02018; Autopsy S02/043). Neurons in layer IV and V had especially strong MANF immunoreactivity (Figure 4). Double immunofluorescence labeling showed partial superimposition of GRP78 and MANF, which were mainly distributed throughout the ER (NBB no. 97015; Autopsy S97/045) (Figure 5). A large number of neurons in the ITGC express both MANF and NeuN, indicating that MANF is a protein expressed primarily in neurons (NBB no. 00119; Autopsy S00/264) (Figure 6).

Mesencephalic Astrocyte-Derived Neurotrophic Factor Is Overexpressed in the ITGC in Pre-AD and AD
Mesencephalic astrocyte-derived neurotrophic factor expression in the ITGC of pre-AD, AD, and non-dementia control cases was evaluated by immunohistochemistry. MANF immunoreactivity was observed in all three groups, mainly in neurons (Figure 7). The rank order of expression level was pre-AD > AD >> control (Figure 8). Additionally, the number of neurons in the ITGC with cytoplasmic MANF expression per unit area (1 mm²) was higher in pre-AD and AD patients than non-dementia control cases. MANF was mainly expressed in the cytoplasm of neurons (Table 2).

DISCUSSION
The results of this study demonstrate that proteins associated with AD pathogenesis were more highly expressed in the ITGC of patients with AD than in control subjects, which is consistent with previous reports (Zhu et al., 2007; Lacosta et al., 2017). These AD biomarkers include p-tau and Aβ17–24, which are components of NFTs and amyloid plaques, respectively. Mesencephalic astrocyte-derived neurotrophic factor is an evolutionarily conserved protein with both cytoprotective and immunomodulatory effects (Neves et al., 2016) that is highly expressed in the developing mammalian cortex and is involved in neurite extension and the regulation of ER homeostasis in neurons (Adaikkan et al., 2019). MANF was shown to play a protective role in cell survival by attenuating the neurotoxicity resulting from ER stress (Xu et al., 2019). Moreover, treatment with recombinant MANF or MANF overexpression alleviated Aβ-induced UPR activation caused by ER stress, while knocking down MANF promoted UPR activation and enhanced the toxicity of Aβ (Xu et al., 2019). Exogenous MANF stimulated nerve repair in dopaminergic neurons.
MANF Expression in Alzheimer’s Disease

FIGURE 6 | Representative images of MANF and NeuN expression in the ITGC. (A–F) Representative micrographs of double immunolabeling with antibodies against MANF (red) and NeuN (green). MANF was mainly detected in neurons. Scale bar: 100 µm (A–F).

(Petrova et al., 2003; Voutilainen et al., 2009; Hao et al., 2017; Liu et al., 2018) and although it was not essential for neuron survival in mouse embryo, the endogenous protein was shown to be necessary for maintaining neuronal ER homeostasis both in vitro and in vivo (Pakarinen et al., 2020). As an ER stress-associated protein, MANF has been implicated in chronic stress and multiple neurodegenerative diseases including AD (Zhu et al., 2017).

In our study, we examined the expression of MANF in the ITGC of human brain specimens from pre-AD and AD patients and non-dementia control cases by immunohistochemistry. A representative Western blot result shows that the MANF protein was stably expressed in HEK 293T and SHSY5Y cells. However, compared with the WT control mice, MANF did not express in the hepatic cells of the HKO control mice. The reported specificity of the anti-MANF antibody (Lindholm et al., 2008; Yang et al., 2017) was confirmed by western blotting with the detection of a single protein band at ∼18 kDa. MANF was mainly present in neurons of the ITGC, consistent with the known expression pattern of MANF in the aging human cerebral cortex and adult mouse brain (Lindholm et al., 2008; Tseng et al., 2017; Yang et al., 2017; Danilova et al., 2019). This was also confirmed by MANF/NeuN double immunolabeling experiments.

The size of nucleoli of ITGC neurons was significantly smaller in pre-AD and AD than in control brains, in line with previous observations (Hu et al., 2003); however, MANF immunoreactivity in neurons was higher in patients than in controls. GRP78 and MANF interact as part of a macromolecular complex in the ER (Glembotski et al., 2012). AD is related to ER calcium deficiency (Trychta et al., 2018); depletion of ER calcium leads to dissociation of the MANF/GRP78 complex and MANF secretion (Apostolou et al., 2008; Glembotski et al., 2012). ER calcium depletion also causes ER stress, which further results in the upregulation of MANF. In the present study, double immunolabeling of MANF/GRP78 showed that the MANF seems localized within ER. Based on these findings, we speculate that in pre-AD and AD, calcium depletion and severe chronic ER stress leads to the upregulation of MANF and activation of the apoptosis signaling pathway, resulting in the shrinkage of neuronal nucleoli and neuronal death.

A large number of neurons in the ITGC express both MANF and NeuN, indicating that MANF is a protein expressed primarily in neurons. MANF expressed by adeno-associated virus (AAV) was predominantly detected

https://www.proteinatlas.org/ENSG00000145050-MANF/tissue
in the cytoplasm of infected cells (Yang et al., 2017), which is consistent with the localization of endogenous MANF. AAV-induced MANF expression was also observed in neurons and glia of the cerebral cortex following ischemia (Airavaara et al., 2010). MANF is upregulated in neurons under pathologic conditions such as focal cerebral ischemia, and a larger infarct area was observed in MANF-deficient brains (Shen et al., 2012), suggesting that MANF exerts a protective effect against ischemic injury in cortical neurons (Mätlik et al., 2018). However, a larger sample size and more detailed cytologic analyses are needed to confirm our results and to determine whether the increased level of MANF in the pre-AD and AD brain is related to a perturbation of ER homeostasis.

There were some limitations to our study, we did not examine the expression of ER stress or apoptosis markers to clarify the function of MANF in AD. Nonetheless, our results demonstrate that MANF is overexpressed in neurons in the brain of pre-AD and AD patients, suggesting that it can serve as a diagnostic marker for early stage disease.

![FIGURE 7](image1.png)

**FIGURE 7** | Mesencephalic astrocyte-derived neurotrophic factor expression in the ITGC of AD, pre-AD, and control cases detected by immunohistochemistry. (A–F) MANF expression in non-dementia control (A,B), pre-AD (C,D), and AD (E,F) samples shown at low (20×) (A,C,E) and high (40×) (B,D,F) magnification.

![FIGURE 8](image2.png)

**FIGURE 8** | Increased MANF expression in the ITGC of pre-AD and AD groups. Overall expression in both groups was higher than in the control group, while the expression level was lower in the AD group than in the pre-AD group, although the difference was non-significant. **P < 0.05 vs. control group.**
DATA AVAILABILITY STATEMENT

The original contributions presented in the study are included in the article-supplementary material, further inquiries can be directed to the corresponding authors.

ETHICS STATEMENT

Paraffin-embedded postmortem human brain specimens were obtained via the Netherlands Brain Bank with informed written consent from the patients or their next of kin for the autopsy and use of brain material and clinical files for research purposes. The patients/participants provided their written informed consent to participate in this study.

AUTHOR CONTRIBUTIONS

X-CL designed the study, performed the immunohistochemistry and statistical analysis, and drafted the manuscript. X-HQ conducted the tissue preparation and parts of the experiments. HF and K-QZ participated in the study design and the statistical analysis. G-HC and Q-SW conceived, designed, and supervised all aspects of the study and revised the manuscript. All authors read and approved the final manuscript.

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FUNDING

This work was supported by grants from the National Key R&D Program of China (no. 2016YFC1305900) and the National Natural Science Foundation of China (no. 81671316).

ACKNOWLEDGMENTS

We are grateful to the Netherlands Brain Bank for providing human brain samples, and thank Jiang-Ning Zhou (USTC) for help with the experiment. We thank Prof. Yu-Xian Shen (Anhui Medical University) for supplying HKO control mice and WT control mice tissue sections.

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**Conflict of Interest:** The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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