Enhanced Glial Reaction and Altered Neuronal Nitric Oxide Synthase are Implicated in Attention Deficit Hyperactivity Disorder

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Attention deficit hyperactivity disorder (ADHD) has a complex etiology, and its specific causal factors remain to be elucidated. Aberration of nitric oxide synthase (nNOS) and inflammation, together with astrocytic and microglial cells have been continually associated with several neurological disorders, including ADHD. Using spontaneously hypertensive rat (SHR), we investigated the changes in nNOS, inflammatory, microglial and astrocytic markers in the frontal cortex and hippocampus at three different ages: onset of hypertension stage (i.e., 6 weeks after birth of SHR), established hypertension stage (i.e., 12 weeks after birth of SHR) and senescent stage (i.e., 12 months after birth of SHR), and compared with its age-matched normotensive control, Wistar-Kyoto (WKY) rats. A significant upregulation of Iba-1 expression in the senescent stage of SHR was observed. Further, we observed an upregulated nNOS expression in both onset and established stages of SHR, and a downregulated nNOS in the senescent stage. Our study showed an age-related increment of astrogliosis in the cortex and hippocampi of aged SHR. On the basis of our results, alterations in the nNOS and Iba-1 expressions, as well as age-related astrogliosis, may contribute to ADHD pathogenesis.

Keywords: attention deficit hyperactivity disorder (ADHD), microglia, neuronal nitric oxide synthase, astrogliosis, spontaneously hypertensive rat

1 INTRODUCTION

Attention deficit hyperactivity disorder (ADHD) is a common neurobehavioral disorder accompanied by brain abnormalities, and is characterized by marked attention deficit, short attention span, hyperactivity, or impulsiveness. It is more common in children, with an estimated prevalence of 3%–5% (Polanczyk et al., 2007). However, children are not the only ones who suffer from ADHD. The disorder can progress into adulthood and lead to unstable relationships, poor work or school performance, and low self-esteem (Okie, 2006). The spontaneously hypertensive rat (SHR) is a genetic model and bred from progenitor Wistar Kyoto rat (Aparicio et al., 2019) and is the most commonly used and studied animal model of ADHD (Bucci et al., 2008). SHRs exhibit behaviors similar to those of ADHD patients in a variety of
testing paradigms, including increased motor activity, impulsivity, and inattention. Evidences including candidate genes, neurotransmitter dysfunction, neuropathology and pharmacology supports construct validity of SHR (Song et al., 2020). Meanwhile, the dopaminergic and (-)-noradrenaline systems of spontaneously hypertensive rats are hypofunctional, which is one of the important mechanisms in the pathogenesis of ADHD (Bucci et al., 2008).

Nitric oxide is a widely distributed gaseous chemical messenger generated by nitric oxide synthase (Mittleman et al.) from L-arginine (Calabrese et al., 2007), and can be synthesized by three NOS isoforms; neuronal NOS (nNOS), inducible NOS (iNOS), and endothelial NOS (eNOS) (Lind et al., 2017). Nitric oxide acts as a non-conventional neurotransmitter in the CNS, and is involved in the regulation of autonomic functions (Qadri et al., 2003). Furthermore, nitric oxide acts as a vasodilator in the cardiovascular system; thus, alterations in nitric oxide production can affect blood pressure and behavior (Kluknsvsky et al., 2016). The neuronal isoforms encoded by NOS1 are the main source of nitric oxide in the CNS. nNOS has been detected in different types of neurons in the cerebellum, hypothalamus, striatum, cerebral cortex, and hippocampus (Domek-Łopacińska and Strosznajder, 2010; Spiers et al., 2019), having a close association with behavioral, memory, cognition, and learning ability. Previous studies have demonstrated that blocking nNOS in the brain may lead augmented systemic mean arterial pressure (MAP) in SHRs in comparison to age-matched WKY rats (Ferrari and Fior-Chadi, 2005). Nitric oxide synthase has been linked to a variety of behavioral disorders, including ADHD. In particular, NOS1 knocked-out mice are more aggressive and impulsive (Volke et al., 1997).

Oxidative stress plays crucial role in the pathophysiology of ADHD. Oxidative stress known as an im-balance between oxidants and antioxidants in the cells. This imbalance due to malfunction of the antioxidant system or by an uncontrolled level of reactive oxygen species (ROS) (Corona, 2020). Brain is the most vul-nerable organ in the body by the oxidative stress which can harm the integrity of neurons. The in-ammation is characterized by changes in microglia, astrocytes, cytokines, chemokines, and related molecular processes within the brain. Neuroinflammation is a key player in psychiatric disorders, such as bipolar disorder, depression, and schizophrenia (Dunn et al., 2019). In the cerebrospinal fluid of ADHD patients, increased pro-inflammatory factor (i.e., TNF-β) and decreased anti-inflammatory cytokine (i.e., IL-4) were detected (Mittleman et al., 1997). The microglia are the central nervous system’s resident immune cells (Réus et al., 2015). Microglia plays an important role in brain damage, neuronal phagocytosis, network connectivity during development, and secretion of inflammatory and anti-inflammatory cytokines (Corona, 2020). Dopamine (specifically, D1R) agonists inhibit microglial activation, concomitant with increased neuronal activity and improved cognitive impairment in ADHD patients (Yokokura et al., 2021). Therefore, targeting the microglia could be an interventional strategy for ADHD.

Astrocytes have unique functional and morphological characteristics. In ADHD, lack of lactic acid produced by the brain’s astrocyte can cause ADHD hyperactivity (Russell et al., 2006; Medin et al., 2019). Moreover, astrocyte can promote the induction and progression of inflammatory states, which are significantly associated with disease status or severity (Corona, 2020). TNF-α, ROS and other inflammatory stimuli induce expression of NF-kB in the astrocytic nucleus and aggravate neuroinflammation (Linnerbauer et al., 2020). Also, the MAPK/NF-kB signaling minimizes the activation of astrocyte and microglia, and lowers the expression of IL-1, TNF-α and other inflammatory factors in ADHD (Song et al., 2020).

The SHR is the most commonly used and studied animal model of ADHD. This is because SHRs exhibit behaviors similar to those of ADHD patients, including increased motor activity, impulsivity, and inattention (Bucci et al., 2008). More importantly, the hypofunctionality of dopaminergic and noradrenaline systems, which are key mechanisms in ADHD pathogenesis, are evident in SHRs (Bucci et al., 2008). In this study, SHRs were divided into three groups based on phases: the onset of hypertension (6 weeks), established hypertension (12 weeks), and the senescent stage (12 months). Alterations in nNOS expression, activation of microglia, and astroglisis in the brain at each stage were investigated and analogized with its age-matched normotensive control, Wistar-Kyoto (WKY) rats, from which the SHRs were originally derived (Ganten et al., 1991).

2 MATERIALS AND METHODS

2.1 Animals
Six-week-old, twelve-week-old, and twelve-month-old SHR, along with age-matched Wistar Kyoto rats (Beijing Vital River Laboratory Animal Technology Co. Ltd., Beijing) were used. Rats were housed under a 12 h light/dark cycle with free availability to food and water. Experiments were carried out following the NIH Guide for the Care and Use of Laboratory Animals.

2.2 Open Field Test
Sixty rats were used, with ten rats from each age group of SHR and WKY. The animals were placed in a square apparatus (100 cm × 100 cm × 45 cm) with black walls on all sides for 5 min. The apparatus was placed under a camera and adjusted. Using the ANY-maze software, set-up programs were calibrated to the recorded areas. Each rat was placed in the center and allowed to move freely. The movement and total distance of rats were recorded during the test time. The arena was cleaned with 75% ethanol after every trial to avoid affecting the test of the subsequent rats.
2.3 Tissue Preparation
Before their death, six rats from each age group of SHR and WKY were anesthetized with pentobarbital sodium (50 mg/kg, i.p.), and perfused intracardially with 300 ml of cool normal saline, followed by 500 ml of 4% paraformaldehyde in 0.1 M phosphate buffer (PBS, pH 7.4). The whole brain was carefully removed and dissected. Brain tissues were post-fixed at 4°C in the same fixative overnight and then transferred to 30% sucrose in PBS until the tissue dropped to the bottom of the container. Brain tissues were cut into five sets of serial coronal sections with a cryostat at −20°C. Each fifth section was collected in each group for free-floating immunohistochemistry and immunofluorescence labeling. 20-μm-thick sections were mounted on gel-coated slides and cover-slipped with Permount (Fisher Scientific, Pittsburgh, PA).

Twelve rats from each age group of SHR and WKY were sacrificed by decapitation, and brains were carefully harvested. The frontal cortex and hippocampus were dissected and immediately frozen in liquid nitrogen for Western blotting and reverse transcriptase-polymerase chain reaction (n = 6 per age group).

2.4 Immunohistochemistry and Immunofluorescence Labeling
All manipulations were performed in 24-well culture plates at room temperature. Sections were incubated in 0.01 M PBS with 3% H2O2 and 10% methanol for 5 min to quench the endogenous peroxidase, followed by blocking in 5% normal goat serum in 0.01 M PBS. Sections were then incubated in primary rabbit polyclonal anti-nNOS (1:500; Santa Cruz Biotechnology, Santa Cruz, CA) overnight at 4°C. After a thorough wash in PBS, sections were incubated with biotinylated donkey biotinylated goat anti-rabbit IgG (1:400; Vector Laboratories, Burlingame, CA) for 1 h at room temperature, followed by an avidin-biotin-peroxidase complex (Vectastain Elite ABC Kit, 1:300; Vector Laboratories, Burlingame, CA) for 1.5 h at room temperature. The reaction product was developed with 0.025% 3,3′-diaminobenzidine (DAB) and 0.0033% H2O2 in 0.2 M Tris-HCl (pH 7.6), and the reaction stopped by rinsing the slides with 0.2 M Tris-HCl. Sections were mounted onto 0.02% poly-L-lysine-coated slides, allowed to dry at room temperature, dehydrated through graded series of alcohol, cleared in xylene, and finally cover-slipped.

For immunofluorescence labeling, sections were permeabilized and blocked with 0.3% Triton X-100 and 10% normal goat serum in 0.01 M PBS, incubated in primary antibodies against nNOS overnight at 4°C, incubated with 1:500 Alexa Fluor 488 Donkey anti-rabbit IgG (Molecular Probes) secondary antibodies for 1 h at 37°C, and finally covered with anti-fade gel/mount aqueous mounting media (Southern Biotech, Al, United States). All negative control sections were incubated in PBS without primary antibodies. Immunofluorescence labeling neurons were photographed with a confocal imaging system (the Bio-Rad MRC-1024UV system, United Kingdom) connected to a Zeiss Axiohot microscope. The digital images were processed using confocal assistant software (Bio-Rad, Richmond, CA).

2.5 Immunofluorescence Staining
After anesthesia, rats were perfused with 1% PBS and 4% PFA. The brain was quickly removed from the skull, soaked in 4% PFA, and fixed at 4°C for 24 h. Then remove paraformaldehyde and add 30% sucrose solution for dehydration until the brain sinks into the bottom of the solution. OCT was used to embed and make continuous frozen sections at 20 μm. Serial consecutive sections were processed for immunofluorescence staining to detect GFAP as a marker of astroglial reaction, Iba-1 as a microglia marker. Collect and pick out the complete brain slices, wash off the embedding agent with PBS solution, 3 times / 5 min. Then, after blocking with blocking solution for 1 h, primary antibodies Iba-1 (Abcam, 1:500), GFAP (Santa Cruz, 1:100) and TNF-α (Abmart, 1:400) were added and incubated overnight at 4°C; Then wash with PBS 3 times for 5 min each time. Add secondary antibody solution and incubate at 37°C away from light for 1 h; Wash 3 times with PBS for 5 min each time and seal with DAPI (VECTASHIELD, United States) containing sealant. Then it was observed with a fluorescence microscope (Olympus BX51, Tokyo, Japan) at an excitation wavelength from 547 to 570 nm (Cy3, red), 494–520 nm (FITC, green), and 360–460 nm (DAPI, blue). Images were taken at 200x magnification.

2.6 Reverse Transcriptase-Polymerase Chain Reaction
Total RNA was extracted with an RNA isolation reagent, Trizol TM (Invitrogen, Carlsbad, CA). Quantitation of RNA was performed by measuring the absorbance of each sample at 260 nm. Quality check was done by gel electrophoresis. RNA samples (2 μl) from each age group were used to generate cDNA. Reverse transcription was carried out using Super-Script™ III Reverse Transcriptase (Invitrogen, Carlsbad, CA). Thermocycling was performed with a gradient thermocycler (Takara, Japan) using GoTaq® Flexi DNA Polymerase (Promega, Madison, WI). β-actin was used as the internal control. The primers used were: nNOS, 5′-acc acc tgt ctc ctc tag cc-3′ (forward), 5′-cct tgt tgt ggc gac tt-3′ (reverse), size of amplicon was 304 bp; β-actin, 5′-tgc tac caa ctt gga cga ca-3′ (forward), 5′-aag gaa ggc tgt aag gc-3′ (reverse), size of amplicon was 573 bp.

The PCR reaction started at first by denaturing cycle at 94°C for 5 min, followed by an amplification profile of 35 reaction cycles with denaturation at 94°C for 30 s, primer annealing at 55°C for 30 s, and elongation at 72°C for 1 min. PCR was performed simultaneously on SHR and WKY samples, with internal controls running in parallel. All RT-PCR experiments included negative controls, in which template RNA or reverse transcriptase was omitted. Each PCR product (10 μl) was electrophoresed on a 2% agarose gel containing Ethidium bromide. Resultant gel bands were visualized in a UV-transilluminator, with the optical density of the bands determined with a MultilmageTM II light cabinet (DE-500) that was equipped with Fluochrome Beta 1.1 software (Alpha Innotech Corp., CA). The relative amount of the
2.7 Western Blot Analysis

Total proteins were extracted from brains with 2 mM PMSF in 1 ml ice-cold RIPA buffer added protease inhibitor cocktail EDTA-free. The protein concentrations were determined using the Bradford protein assay. SDS-PAGE was performed on 12% polyacrylamide slab gel, electrophoresed in Tris-glycine buffer under denaturing conditions, and transferred to PVDF membrane at 70 V for 1.5 h at 4°C in a Bio-Rad TransBlot apparatus. Membranes were blocked for 1 h at room temperature with 5% (wt/vol) nonfat milk in Tris-buffered saline 0.05% (vol/vol) Tween 20 (TBS-Tween; 50 mM Tris base, 200 mM NaCl, 0.05% Tween 20). Membranes were then incubated for 12 h at room temperature with primary polyclonal rabbit anti-GFAP (1:500, Thermo Fisher Scientific, Waltham, MA).

Membranes were rinsed thrice (each wash lasting 10 min) with TBS 0.05% Tween 20 (TBST), incubated at room temperature for 1 h with secondary horseradish peroxidase (HRP)-conjugated goat anti-rabbit IgG (1:5,000, Santa Cruz Biotechnology, CA) and then washed again with TBST. Proteins were detected by enhanced chemiluminescence (ECL reagents; Amersham, Piscataway, NJ) exposed to radiographic film (Hyperfilm ECL; Amersham) as described in the technical manual provided by Amersham Pharmacia Biotech, Inc. Densitometry of the immunoreactive bands of interest was performed by acquiring digital images of the radiographic film (Hyperfilm ECL; Amersham) using Bio-Rad Quantity One 1-D software. To normalize protein bands to a gel loading control, membranes were washed in TBST and re-probed with rabbit anti-β-actin (1:5000, Abcam, Cambridge, MA) followed by incubation with peroxidase-conjugated goat anti-rabbit (1:5000, Santa Cruz Biotechnology, CA) and ECL detection. For the negative control, the primary antibody was omitted.

2.8 Statistical Analysis

Data from morphometric studies of immunohistochemistry, Western blot analysis, and RT-PCR were presented as mean ± SD. Data from the open field test were presented as mean ± SEM. One-way analysis of variance (ANOVA) with post hoc Tukey tests was used to determine statistical significance. p value of <0.05 was considered statistically significant. All statistical tests and graphs were performed or generated using GraphPad Prism Version 8.0 (GraphPad Prism Software, Inc., CA, United States).

3 RESULTS

3.1 Spontaneously Hypertensive Rat Display Behavior Paradigm of Attention Deficit Hyperactivity Disorder

The open-field test showed that the locomotor trajectory, total distance and mean speed of SHRs were significantly different from WKY rats. Regarding the activity trajectory (Figure 1A), SHR rats had a wider range of activity and more complex trajectories, whereas WKY rats had a simple and single route with a smaller range and more reciprocal movements. The total distance was significantly higher in SHRs at all ages compared with WKY rats (**p < 0.0001**) in each age group. Nonetheless, this difference was insignificant (**p = 0.0913**, **p = 0.0912**). At 12 months, the distance and speed were significantly decreased when compared with 6- and 12 week-old rats (**p < 0.0001**, **p < 0.0001**).
3.2 mRNA Expression of Neuronal Nitric Oxide Synthase is Increased in the Frontal Cortex and Hippocampus of Spontaneously Hypertensive Rat

At weeks 6 and 12 post-natal stage we found higher levels of nNOS mRNA levels in frontal cortex in SHRs than WKY rats (Figures 2, 3, p < 0.05). The expression of nNOS in 12 weeks SHR was higher than that in 6 weeks SHR. WKY rats also showed the same trend. However, the mRNA level of nNOS was considerably increased in the aforementioned brain regions of aged WKY (12 months old) but declined substantially in the age-matched SHRs (Figures 2, 3G,H, p < 0.01).

3.3 GFAP Expression Increases Gradually With Age

Immunoblotting analysis showed the protein level of GFAP was increased with age in the frontal cortex and hippocampus of SHR and WKY rats (Figure 4). Higher GFAP protein level was detected in the brain of SHR when compared with WKY rats at 6 and 12 weeks after birth (p < 0.05, Figures 4B,C). Additionally, there was a significant difference in GFAP level at 12 months (p < 0.01, Figure 4).

3.4 The Expression of GFAP Increases in the Frontal Cortex and Hippocampus of Spontaneously Hypertensive Rat

Immunofluorescence staining (Figures 5A,B) was used to evaluate the expression of astrocyte and the immunofluorescence intensity of GFAP positive cells (Figure 5C). The GFAP expression was increased in the hippocampus and prefrontal cortex of SHR and WKY rats. Also, the GFAP expression was significantly higher in the CA1 region of SHR hippocampus compared with the WKY group at 6 weeks (p < 0.001, Figure 5C). Further, the GFAP expression was increased in other periods when compared with the WKY group (p < 0.01, Figure 5C). In addition, the GFAP expression was significantly increased in the SHR group at 12 months (aging) compared with the WKY group in the prefrontal cortex (p < 0.05, Figure 5C).
3.5 The Expression of Iba-1 Increases Gradually in the Frontal Cortex and Hippocampus of Spontaneously Hypertensive Rat

Immunofluorescence staining (Figures 6A,B) was used to detect microglial activation in the hippocampal CA1 region and the prefrontal cortex. Iba-1 expression was significantly higher in the hippocampal CA1 region and prefrontal cortex region of SHR compared with WKY rats at 12 months ($p < 0.001$, Figure 6C). In comparison with WKY group at 6 and 12 weeks, Iba-1 expression was increased in SHR at the same time ($p < 0.01$, Figure 6C).

3.6 The Expression of TNF-α Increases Gradually in the Frontal Cortex and Hippocampus of Spontaneously Hypertensive Rat

Immunofluorescence staining (Figures 7A,B) was used to evaluate the expression of TNF-α and the immunofluorescence intensity of TNF-α positive cells (Figure 7C). The TNF-α expression was increased in the hippocampus and prefrontal cortex of SHR rats when compared with the age-matched WKY rats. Moreover, TNF-α expression in the CA1 region of the hippocampus was significantly higher than the WKY group at 6 weeks ($p < 0.01$). The TNF-α expression also increased in other periods compared with WKY group ($p < 0.05$). In addition, the TNF-α expression was significantly upregulated in the SHR group compared with the WKY group in the prefrontal cortex of each periods ($p < 0.05$, Figure 7C).

4 DISCUSSION

In the present study, SHRs (an animal model of ADHD) were obtained by selective breeding of WKY rats (control animals) with the highest blood pressure. These animal strains are normotensive at birth, and gradually develop stable hypertension during the first weeks of life (Rubattu et al., 2018). The blood pressure of SHR rises around 5–6 weeks of...
age, and steadily increases to reach systolic blood pressure of 180–200 mmHg. SHR has several features of hypertensive organ damage, including cardiac hypertrophy, cardiac failure, and renal dysfunction in the latter stages of their life. As such, this strain has been employed for evaluating genetic factors in hypertension (Doris, 2017). Besides, SHRs have been used as an animal model for studying vascular brain damage and related pathogenic development of learning and memory deficits (Hussein et al., 2005; Li et al., 2009; Tayebati et al., 2012; Leffa et al., 2019).

The open field test (OFT) is a common experimental method for evaluating the ability of animal self-motion, and is used in multidisciplinary experimental studies. In behavioral studies of ADHD, the distance and speed of movement of rats in the open field test are often used as an important basis for evaluating the two core symptoms of ADHD: hyperactivity and impulsivity. Generally, locomotion is used to assess hyperactive behavior, and the speed of movement is used to evaluate the state of agitated impulsivity in animals.

In this study, SHRs showed marked differences in locomotion when compared to the WKY rats. After 6 weeks, SHRs demonstrated considerable long motor distance and fast motor speed, which are typical of hyperactive and impulsive behaviors, and are behavioral features of ADHD.

Centrally produced nitric oxide affects autonomic responses and participates in the regulation of sympathetic tone to the periphery (Han et al., 2010). Our study found upregulated nNOS expression in the frontal cortex and hippocampus at the onset and established stages of hypertension. This was consistent with a previous study that established gene expression of nNOS was increased in central autonomic centers in animals with increased sympathetic activity. The above statement supports the hypothesis that nitric oxide plays an important role in the maintenance of homeostatic balance through the modulation of sympathetic activity (Hojná et al., 2007). Thus, during over-activity of the sympathetic system in hypertensive condition, the sustained increase in blood pressure and related functional alteration in the autonomic system might act as a stimulus for exacerbate nNOS activity, and nitric oxide production in spontaneously hypertensive rats might be enhanced in CNS to compensate for the elevated blood pressure (Hojná et al., 2007; Sun et al., 2019). However, the excess amount of nitric oxide, if not balanced by antioxidants, will react with superoxide (O$_2^-$) and other reactive oxygen species to produce peroxynitrite, and trigger oxidative stress. Noteworthy is that oxidative stress is involved in neurodegenerative diseases owing to its free-radical properties (Colas et al., 2006; Török, 2008). Besides, it has been correlated with the pathophysiology of ADHD (Álvarez-Arellano et al., 2020).

In the aged animals (i.e., 12 month-old), the brain nNOS expression of WKY rats was significantly higher. This was in line with a previous study that showed that aged animals exhibit increased in cortical NOS activity regardless of their genetic background (Law et al., 2003). Nitric oxide has been implicated in neurodegenerative diseases due to its oxidative properties. The upregulation of nNOS expression in aged WKY might be due to the abatement of free radical scavengers and diminished anti-oxidative capacity in old animals. Thus, elevated nitric oxide release would increase oxidative stress, and coincide with cognitive impairments owing to the accumulation of oxidative damages. Concerning aged SHR, since considerable upregulated nNOS in the early stages of the aging process affects the microenvironment in each sub-regions of the CNS, the secondary change in oxidative stress, excitotoxicity, energy depletion, neuroinflammation together with apoptosis could lead to significant loss of nNOS expressing neurons. The diminution of nNOS positive neuron number in the brain of aged SHR, as evidenced from our experiments, was consistent with this hypothesis, which might, at least in part, explain the decline tendency of nNOS expression in the older SHR.

Oxidative stress in ADHD causes an inflammatory response that affects ADHD attention deficit and hyperactivity disorder (Corona, 2020). Under different pathological conditions, abnormal induction of nNOS leads to excessive production of NO, and results in oxidative stress. ROS can cause microglia activation (Kim et al., 2019). Iba-1 expression was significantly higher in 12 months SHR than the WKY rats. This suggests significant microglia activation in aged hypertensive rats, which may be attributed to the accumulation of oxidative damage. The nNOS expression in the 12 month-old SHRs confirmed this theory. Activated

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**FIGURE 4** The level of GFAP protein expressed in the frontal cortex. (A) The Western blot result of GFAP in the frontal cortex of SHR and WKY. (B) The decreased expression of GFAP in the frontal cortex was more severe in the SHR than in WKY. *p < 0.05 versus age-matched WKY control, **p < 0.01 versus age-matched WKY control. (C) The change of GFAP expression in the frontal cortex in aged process of SHR and WKY. The values represent mean ± SD.
Microglia release superoxide anion free radicals, nitric oxide, TNF-α, IL-1 and other inflammatory factors that exacerbate inflammatory damage (Rèus et al., 2015). TNF-α expression was higher in 12 months SHR than WKY rats. Compared with the WKY rats, SHR rats at 6 and 12 weeks showed the same trend. High levels of proinflammatory cytokine can affect synaptic plasticity and neurogenesis (Anand et al., 2017; Leffa et al., 2018). Therefore, cytokines can affect cognitive function. This was consistent with dementia behavior in aged hypertensive rats. In addition, the TNF-α is up-regulated in tryptophan metabolism, which is associated with the severity of ADHD symptoms (Aarsland et al., 2015; Anand et al., 2017).

Astrogliosis is an increase in the size of specific cytoskeletal protein GFAP-immunoreactive astrocytes. It is common in several brain damage (Tomassoni et al., 2004). A higher GFAP

**FIGURE 5** GFAP immunofluorescence showed that the expression of GFAP in the cortex and hippocampus of SHR rats was increased compared with that of WKY rats. (A) the expression of GFAP in prefrontal cortex of WKY and SHR, (B) the expression of GFAP in hippocampal CA1 of WKY and SHR, bar = 20 μm *p < 0.05 versus age-matched WKY control. **p < 0.01, age-matched WKY control. ***p < 0.001 age-matched WKY control. The values represent mean ± SD. (C) The mean intensity of GFAP was upregulated in the SHR at each age-matched group in the hippocampus and frontal cortex.
expression has been reported in several neurodegenerative disorders and autogenic senescence-accelerated mouse (SAM) model (Tomassoni et al., 2004; Wu et al., 2005; Tayebati et al., 2012). The increased astrocyte-neuron contacts could restrict or reduce synaptic plasticity by decreasing the potential interneuronal contacts (Finch, 2003). Our results showed GFAP-positive astrocytes increased with age in both the frontal cortex and hippocampus of SHR and WKY, suggesting it is an age-related pathological change of CNS. We also speculated that an increase in astrocyte volume might lead to a corresponding decrement in neuronal volume. Our results from the present study showed GFAP-positive astrocytes to increase with age in the frontal cortex and hippocampus of SHR and WKY rats, suggesting it is an age-related pathological change of the CNS. Also, we speculated that an increase in astrocyte volume might lead to a corresponding decrement in neuronal volume. Our results showed the brains of the SHR model had more severe astrogliosis conditions than the WKY rats, especially at the senescence stage of 12 months, which was incongruity with marked loss of nNOS neurons at this stage. This implies that elevated nitric oxide production in the early stage and early onset of astrogliosis occurring in the brain of animals with established hypertension could hinder the normal survival conditions of functional neurons, and lead to abnormalities, such as impaired learning and memory.

It is worth noting that there are some limitations regarding the use of SHR as a disease model for ADHD. With reference to our present study, the model was selectively bred, and so is difficult to separate the effects of hypertension from hyperactivity (Majdak et al., 2016). In addition, ADHD has been associated with

![FIGURE 6](A) The expression of iba-1 in the prefrontal cortex of WKY and SHR. (B) the expression of iba-1 in the hippocampal CA1 region of WKY and SHR. (C) Iba-1 immunofluorescence showed that the expression of iba-1 protein in the cortex and hippocampus of SHR was increased compared with that of WKY Group, bar = 100 μm. **p < 0.01 versus age-matched WKY control. ***p < 0.001, vs. age-matched WKY control. The values represent mean ± SD.
dysfunction in monoamine neurotransmitters pathways, such as 5-hydroxytryptamine (5-HT), dopamine (DA), and norepinephrine (NE) (Klein et al., 2017). Particularly, 1 month after the birth of SHRs, brain expressions of tyrosine hydroxylase and DAT genes are decreased, dopamine and norepinephrine levels are abnormal, and decrease in function exhibits signs of ADHD (Russell, 2007). In their adulthood, expression of tyrosine hydroxylase and DAT genes and monoamine neurotransmitter levels increase, causing a decrease in ADHD symptoms, such as curtailed hyperactive and impulsive behaviors. The specific reasons for the decrease in the distance and speed of senile SHR need to be further studied.

5 CONCLUSION

The imbalanced expression of nNOS, abnormal production of nitric oxide, microglia activation, and age-related astrogliosis that are present in specific brain sub-regions related to learning ability and memory formation may contribute to pathological changes in the brain of hypertensive animal model, ultimately leading to memory and cognitive impairments in ADHD. Investigations targeting the neuronal toxic effects of excess nitric oxide, alleviating age-related astrogliosis processes, and preventing the loss of

FIGURE 7 | TNF-α immunofluorescence showed that the expression of TNF-α protein in the hippocampus and frontal cortex of SHR was increased compared with that of WKY group, bar = 100 μm. (A) The expression of TNF-α in the hippocampus CA1 region of WKY and SHR. (B) the expression of TNF-α in the frontal cortex of WKY and SHR. (C) The mean intensity of TNF-α was upregulated in the SHR at each age-matched group in the hippocampus and frontal cortex. *p < 0.05 versus age-matched WKY control. **p < 0.01, versus age-matched WKY control. The values represent mean ± SD.
cholinergic neurons could potentially be an effective interventional medium for ADHD patients.

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 author.

ETHICS STATEMENT
The animal study was reviewed and approved by the Ethics Committee of Zhejiang University.

AUTHOR CONTRIBUTIONS
Conceptualization, PZ and HF; methodology, HF, and SC; software, HF, SC, and NA; validation, PZ and HF; formal analysis, HF, and SC; investigation, HF, and GS; resources, HF, SY and CI; data curation, HF; writing original draft preparation, HF, SC, and PZ writing and editing, PZ and NA visualization, HF, SC and GS; supervision, MF; project administration, MF; funding acquisition, MF All authors have read and agreed to the published version of the manuscript.

FUNDING
This research was funded by the National Natural Science Foundation of China, grant number 81871063.

ACKNOWLEDGMENTS
We are grateful to the Core Facilities of Hangzhou Normal University Affiliated Xiaoshan Hospital. Institute of Neuroscience of Zhejiang University for their technical assistance. Laboratory animal center of Zhejiang University for animal breeding and behavioral testing.
