Chiropractic Therapy Modulated Gut Microbiota and Attenuated Allergic Airway Inflammation in an Immature Rat Model

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Background: As a type of traditional Chinese massage, chiropractic therapy is applied to prevent and treat children with asthma in China. However, its mechanism of action is unclear. Allergic airway inflammation plays a key role in the occurrence and development of asthma, in which changes in gut microbiota are involved. The present study investigated the influence of chiropractic therapy on allergic airway inflammation (AAI) and gut microbiota in an immature rat model.

Material/Methods: Three-week-old male Sprague-Dawley rats were divided randomly into control (CN), AAI, and chiropractic (CP) groups. AAI and CP groups were sensitized and challenged with ovalbumin (OVA) to induce AAI. The CP group received chiropractic therapy during AAI modelling. AAI was assessed by cell counts in bronchoalveolar lavage fluid and HE staining of lung tissues. Plasma OVA-sIgE, IFN-γ, IL-4, and IL-10 levels were detected by ELISA. DNA extraction from feces samples was used for 16S rRNA gene sequencing and analyzed for gut microbiota by Quantitative Insights Into Microbial Ecology (QIIME).

Results: AAI group had significantly lower richness and diversity of gut microbiota along with Th2 response and allergic airway inflammation. Moreover, the AAI group had lower abundance of butyrate-producing bacterial taxa with more Lactobacillus. Chiropractic therapy significantly increased the richness and diversity of gut microbiota and increased butyrate-producing bacterial taxa and decreased Lactobacillus, along with attenuating Th2 response and allergic airway inflammation during AAI modelling.

Conclusions: Chiropractic therapy attenuated allergic airway inflammation and optimized gut microbiota in an immature rat model, which might promote the development of adult-like butyrogenic milieu, immunotolerance, and inflammation attenuation.

MeSH Keywords: Asthma • Massage • Microbiota

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Background

Asthma is recognized as a chronic inflammatory airway disease, ranking among the top 20 conditions worldwide for disability-adjusted life years in children [1]. Allergic airway inflammation plays a critical role in the development and progression of asthma [2]. Th2 helper 2 cell (Th2) response, characterized by activation of Th2 cytokines and increased production of immunoglobulin (Ig) E, is closely related with the initiation of allergic airway inflammation in asthma [2]. Interleukin-4 (IL-4) is the key effector in Th2 response and is indispensable to induce Th2 cell development and IgE production [3].

The host immune system coordinates the balance of effector and regulatory immune cells, as well as anti- and pro-inflammatory cytokines, through interaction with microbiota in a physical condition [4]. Mounting evidence links the development of asthma to the dysbiosis of gut microbiota [5,6]. A loss of microbial diversity induces skewed Th2 response and promotes asthma development [7,8]. The diversity of gut microbiota is essential to functional redundancy and diversity for consistent establishment of a healthy signature of short-chain fatty acids (SCFAs) [9,10]. Microbe-producing SCFAs can increase IL-10+ T cells, promote naïve T-cell differentiation into interferon (IFN)-γ-producing Th1, and provide protection against inflammatory response [11–13]. IFN-γ, as a key Th1 cytokine, can inhibit IL-4 production and regulate the differentiation and aggregation of eosinophils [14]. IL-10, produced by various T cells, can block Th2 response, induce and maintain immunotolerance, and inhibit inflammation [15,16]. Therefore, regulation of gut microbiota is an important strategy in the primary prevention of allergic diseases [17]. However, so far, the evidence that supports probiotics administration for reducing the risk of children asthma is insufficient [18].

Massage therapy is a traditional healing method and has long been applied to prevent and treat children with asthma in China and in Western countries [19]. A systemic review indicated that massage can significantly enhance the effect of current conventional therapies for children with asthma [19]. However, the underlying mechanism is still unclear. Chiropractic therapy (Nie Ji therapy in Chinese, literally means pinching-along-the-spine therapy) is an important and popular traditional Chinese massage for children [20,21]. Our previous clinical trials showed that chiropractic therapy attenuated the symptoms of allergic diseases, along with possible modulation of gut microbiota [22,23]. Ovalbumin (OVA) sensitization and aerosol challenge are widely used to induce allergic airway inflammation [24]. Therefore, we designed this study to investigate the influence of chiropractic therapy on gut microbiota and OVA-induced allergic airway inflammation (AAI) in an immature rat model.

Material and Methods

Animals

Male specific pathogen-free (SPF) Sprague-Dawley rats (aged 3 weeks, weight: 47.92±4.38 g, provided by Qinglongshan, Nanjing, China) were kept in Plexiglas cages in the animal experimental laboratory, and were provided with food and water ad libitum under a standard 12/12 h light/dark cycle. The animal protocol was approved by the Animal Care and Use Committee of Nanjing University of Chinese Medicine (No 201805A005). The experiment was conducted in accordance with the National Institutes of Health Guidelines for the Care and Use of Laboratory Animals.

Allergic airway inflammation (AAI) model and group treatments

After 4-day acclimatization, all rats were divided randomly into a control group (CN, n=6), an AAI group (n=8), and a chiropractic group (CP, n=8). The AAI model was induced in the AAI group and CP group, in which rats were sensitized on day 9 and day 16 with intraperitoneal (i.p.) injections of 1 mg OVA (OVA, Grade V, Sigma, No A5503) mixed into 1.5 mL 4% Al(OH)3 gel (BF040, Xi’an Heart company, China). From day 23 to day 26, the rats were challenged with 1% OVA aerosol (OVA, Grade V, Sigma, No A5503) for 30 min once a day in a plastic chamber connected to an atomizer (402AI type, Jiangsu Yuwell Medical Instruments Co., Ltd., Jiangsu, China) set at its highest volume. The CN group was treated in the same way, but was sensitized and challenged with saline. Chiropractic therapy was performed in the CP group once a day from day 5 to day 26 before and during OVA-induced modelling (Figure 1). No rats died before sacrifice.

The procedure of chiropractic therapy was performed as follows: The manipulator, who had been trained before the study, put on a disposal mask and gloves, and then held a rat gently in the left palm. First, he stroked the rat’s back slowly with the right hand from the neck to the bottom 20 times (strok- ing speed: ~5 cm/s, pressure: ~5 N detected by FingerTPS system, Pressure Profile System, Inc, USA). Second, he pinched the back skin with the fingers of right hand and pushed continuously from the bottom to the neck 25 repeats (~10 s/repeat, pressure: ~12 N) from day 5 to day 14 and for 30 repeats from day 15 to day 26 to avoid manipulation adaptation. Finally, he stroked the back up and down 3 times. The manipulating force was moderate to keep rats relatively calm without obvious discomfort or screaming during the whole procedure. The rats in the CN group and AAI group were only housed individually in clean cages for the same duration as the intervention.
On day 27, all rats were anaesthetized with i.p. injections of 5 mL/kg 25% Urethane (No. U2500, Sigma USA). Blood was collected from the abdominal aorta into tubes with ethylene diamine tetraacetic acid. Plasma was isolated by centrifugation at 3000 rpm for 20 min at 4.0°C. The supernatant was collected and stored at –80°C. Bronchoalveolar lavage fluid (BALF) was collected immediately after blood collection. The trachea was punctured with a 12-gauge gavage needle and 3 ml PBS was slowly injected into the left lung, and then withdrawn and injected again 5 times to collect BALF. The middle lobe of the right lung was collected and put into a flask containing 4% buffered formalin solution for hematoxylin-eosin (HE) staining. Two fresh fecal pellets from the colon were collected from rats and frozen immediately in liquid nitrogen and stored at –80°C for gut microbiota analysis.

**Inflammatory cell counts in BALF**

Total inflammatory cells were counted by automatic cell counter (Countstar IC1000, Shanghai China). The collected BALF was centrifuged at 5000 rpm for 10 min at 4°C and the remaining cell pellets were made into cell smears. Different types of inflammatory cells were observed by Wright-Giemsa staining following the manufacturer’s instructions (G1020, Solarbio, Beijing China). Cells were differentially counted under light microscopy (DP71/BX60, Olympus Corp., Tokyo, Japan).

**Histopathological observation of the lung by HE staining**

The samples were dehydrated, embedded in paraffin, and cut into 4-μm sections for routine HE staining (agents provided by Sinopharm Beijing, China; Baso, Zhuhai, China). Images were obtained under light microscopy (DP71/BX60, Olympus Corp., Tokyo, Japan).

**Measurement of plasma OVA-sIgE, IFN-γ, IL-4, and IL-10 levels by enzyme-linked immunosorbent assay (ELISA)**

The plasma levels of OVA-sIgE, IFN-γ, IL-4, and IL-10 were measured by ELISA according to the manufacturer’s protocol (JEB-15429, JEB-13737, JEB-13730, JEB-13734 JinYibai, Nanjing, China). Every sample was duplicated to mitigate biases.

**DNA extraction and 16S rRNA gene sequencing for gut microbiota**

Fecal pellets were homogenized using a beadbeating method (FastPrep bead matrix E, MP Biomedicals, Santa Ana, CA), and total DNA was extracted with the E.Z.N.A., R Stool DNA Kit (SKU: D4015-02 Omega Bio-tek, Norcross, GA) following the manual. Purity and quality of the genomic DNA were checked on 0.8% agarose gels. The V3-V4 hypervariable region of bacterial 16S rRNA gene were amplified as previously described [25]. PCR products were purified with a QIAGen Gel Extraction Kit (28704, QIAGEN, Germany), quantified by RT-PCR, and sequenced at Allwegenec, China. Every sample were processed in triplicate to mitigate reaction-level PCR biases. Deep sequencing was performed on the Miseq platform.

**Statistical analyses**

For gut microbiota sequencing, the dataset was analyzed by VSEARCH. The sequences were clustered into operational taxonomic units (OTUs) at 97% similarity level [25]. Alpha diversity indices were calculated at OTUs in Quantitative Insights Into Microbial Ecology (QIIME), including observed species, Chao1 richness index, and Shannon diversity index [26]. To compare differences in bacterial composition between the groups, the analysis of similarities (ANOSIM) was performed on unweighted UniFrac distance matrixes and presented as principal coordinate analysis (PCoA) [27,28]. Linear discriminant analysis effect size (LEfSe) was used to detect typical abundant taxa among groups [29]. Based on LEfSe, the comparison of the relative abundance of bacterial taxa between groups was analyzed by Metastats [30].

Data are expressed as mean±standard deviation (SD). Except for the analysis of gut microbiota mentioned above, the differences of other data among groups were analyzed by one-way analysis of variance (ANOVA) and differences between 2 groups were analyzed by two-tailed t test with SPSS 22.0 (SPSS, Inc., USA.) A P value <0.05 was considered statistically significant.

**Results**

**Chiropractic therapy attenuated allergic airway inflammation (AAI)**

Compared with the CN group, OVA sensitization and challenge significantly increased the total numbers of inflammatory cells and eosinophils in the AAI group (P<0.001, P<0.001; Figure 2), and increased the numbers of lymphocytes and neutrophils (P<0.001, P<0.01; Figure 2). To further investigate the inflammation status, the histopathological changes of the lungs were
assessed by HE staining. The AAI group showed obvious infiltration of inflammatory cells around the bronchioles and arteries and in the lungs. Moreover, the AAI group showed constricted bronchioles and thicker walls (Figure 3). These results indicated that OVA sensitization and challenge induced AAI in immature rats. Compared with the AAI group, chiropractic therapy remarkably decreased the counts of total inflammatory cells, eosinophils, lymphocytes, and neutrophils (P<0.01, P<0.001, P<0.001 vs. CN group; P<0.05, **P<0.01 vs. AAI group).

Chiropractic therapy increased the production of IFN-γ and IL-10 and attenuated Th2 response

Compared with the CN group, the AAI group showed significantly increased levels of OVA-sIgE and IL-4 (P<0.01, P<0.001; Figure 4A, 4B), along with significantly lower levels of IFN-γ (P<0.05; Figure 4C). IL-10 levels in the AAI group decreased but not significantly compared with the CN group (P>0.05; Figure 4D). Chiropractic therapy significantly decreased the production of OVA-sIgE and IL-4 (P<0.05, P<0.001; Figure 4A, 4B) and significantly enhanced the production of IFN-γ and IL-10 (P<0.001, P<0.05; Figure 4C, 4D) during AAI modelling compared with the AAI group.

Chiropractic therapy increased the richness and diversity of gut microbiota

A total of 12 phyla and 137 genera were identified among groups. Compared with the CN group, the number of observed species as shown by Chao 1 and Shannon indices were decreased significantly in the AAI group (P<0.01, P<0.001; Figure 5A–5C). Therefore, OVA-induced AAI led to significant loss of the richness and abundance of gut microbiota in immature rats. Chiropractic therapy significantly increased the number of observed species, Chao 1 and Shannon indices compared with the AAI group (P<0.05, P<0.05; Figure 5A–5C), indicating that chiropractic therapy attenuated the loss of richness and diversity of gut microbiota during AAI modelling.

Taxonomic characterization of the gut microbial profile among groups

To characterize the general differences in gut microbial communities among groups, PCoA was performed on unweighted UniFrac distances (Figure 6). The PCoA plots demonstrated significant separation between the CN and AAI groups (P<0.05) as well as between CP and AAI groups (P<0.01) and no significant separation between the CP and CN group (P>0.05). LEfSe was performed to determine typical abundant bacterial taxa as respective biomarkers among groups. The following were over-represented in the AAI group as Phylum Firmicutes, class Bacilli, family Lactobacilaceae, genus Lactobacillus, species Lactobacillus_vaginalis and Lactobacillus_gasseri, whereas order NB1_n and genus Ruminiclostridium_1 and Ruminococcaceae_UCG_014 were over-represented in the CN group. The higher
**Figure 4.** Plasma levels of OVA-sIgE, IL-4, IFN-, and IL-10. (A–D) All these indexes were measured in duplicate by ELISA (n=6–8).

* P<0.05, ** P<0.01, *** P<0.001 vs. CN group; # P<0.05 vs. AAI group.

**Figure 5.** The altered richness and diversity of gut microbiota represented by observed species, Chao1, and Shannon indexes. (A–C) These analyses were based on 16S rRNA gene sequencing. Every sample was processed in triplicate (n=5). * P<0.05, ** P<0.01 vs. CN group; # P<0.05, ### P<0.001 vs. AAI group.

**Figure 6.** PCoA of unweighted UniFrac distances among groups (PCoA plots with Principal Coordinates (PC) 1 and PC 2). Each dot stands for a sample. ANOSIM reveals significant separation between the CN and AAI groups (P<0.05) as well as between the CP and AAI groups (P<0.01) and no significant separation between the CP and CN groups (P>0.05).
abundance of *Lactobacillus* explained the higher abundance of phylum Firmicutes in AAI immature rats (Figure 7A, 7B). In the CP group, Phyla Bacteroidetes and Proteobacteria, class Bacteroidia, families *Bacteroidales_S24_7_group*, *Prevotellaceae*, *Ruminococcaceae* and *Staphylococcaceae* were over-represented (all LDA score >2, P<0.05; Figure 7A, 7B).

**Chiropractic therapy modulated bacterial taxonomic composition**

To know more about the differential taxonomic composition among groups, the comparison of relative abundance of selected bacterial taxa at phylum, family, genus, and species levels were performed in detail based on LEfSe results.

At the phylum level, gut microbiota was mainly composed of Firmicutes, Bacteroidetes, Proteobacteria, Actinobacteria, and Saccharibacteria among the groups. In the AAI group, the relative abundance of Bacteroidetes, Proteobacteria, and Saccharibacteria significantly decreased (P<0.05, P<0.01, P<0.01; Figure 8A, 8B) and the relative abundance of Firmicutes increased (P<0.01; Figure 8A, 8B) compared with the CN group. Compared with the AAI group, the CP group showed significantly decreased abundance of Firmicutes (P<0.05; Figure 8A, 8B) and increased abundance of Bacteroidetes, Proteobacteria, and Saccharibacteria (P<0.01, P<0.01, P<0.01; Figure 8A, 8B). At the family level, the AAI group had significantly lower abundance of *Ruminococcaceae* and *Bacteroidales_S24_7_group* (P<0.01, P<0.01; Figure 9A) as well as higher abundance of...
Lactobacillaceae, but without a significant difference (P=0.056 >0.05; Figure 9A). Compared with the AAI group, the CP group had higher abundance of Ruminococcaceae, Bacteroidales_S24_7_group and Prevotellaceae (P<0.01, P<0.01, P<0.001; Figure 9A), as well as lower Lactobacillaceae (P<0.01; Figure 9A).

At the genus level, the AAI group had lower abundance of several genera such as Roseburia, Ruminococcus_1, Ruminococcaceae_UCG-014 compared with the CN group (P<0.01, P<0.01, P<0.01; Figure 9A). At the genus level, the AAI group had lower abundance of Staphylococcaceae compared with the CN group (P<0.01, P<0.05; Figure 10A, 10B). The CP group had higher Roseburia, Ruminococcus_1 and lower abundance of Lactobacillus_10 (P<0.01; Figure 10A, 10B) compared with the AAI group. At the species level, the AAI group had higher abundance of Lactobacillus_vaginalis and Lactobacillus_gasseri compared with the CN group (P<0.01, P<0.05; Figure 9B). The CP group had lower abundance of Lactobacillus_vaginalis and Lactobacillus_gasseri compared with the AAI group (P<0.01, P<0.01; Figure 9B).

**Discussion**

We found that OVA sensitization and challenge caused skewed Th2 response and allergic airway inflammation in immature rats. Moreover, OVA sensitization and challenge caused loss
of gut microbiota richness and diversity, which is consistent with the results of a previous OVA-induced asthmatic animal study [31]. Meanwhile, except for the extreme high abundance of phylum Firmicutes, the abundances of other major phyla were generally lower in AAI immature rats.

Unexpectedly, AAI immature rats had a higher abundance of *Lactobacillus*, which are always considered as probiotic bacteria [32]. Lactate produced by *Lactobacillus* provides an unfavorable environment for the growth of many pathogenic bacteria and also acts as a permeabilizer of the Gram-negative bacterial outer membrane [32]. As a facultative anaerobe, *Lactobacillus* is typically abundant in the early colonization stage before the replacement with anaerobic bacteria [33,34]. Changes in gut bacteria composition precede the altered level of microbiota-derived metabolite in the bloodstream [35]. A typical example of bacterial metabolites is SCFA, among which butyrate, acetate, and propionate are the most abundant in adults as well as lactate in infants [34,35]. There are cross-feeding interactions between different species; therefore, a diverse microbiota is key to establishing a healthy SCFA signature [10].

In our study, AAI rats had less butyrate-producing bacteria such as *Bacteroidales S24-7* and *Ruminococcaceae* [37,38] and less other phyla with more *Lactobacillus*. This result indicated a dysbiotic gut microbiota and potentially unhealthy SCFAs signature in AAI immature rats, which might negatively affect the immunotolerance development and worsen AAI.

Massage is reported to improve the immunity [39] and have good effects on improving asthma [19,40]. As a Chinese traditional massage, chiropractic therapy was initially applied for indigestion among children by pinching the skin along the spine, and it demonstrates very good clinical effects in the treatment of various digestive problems among children [20,21]. It has now become a popular therapy for prevention and treatment of allergic diseases in pediatric massage departments. Clinical studies showed that chiropractic therapy significantly decreased nighttime symptom scores and acute attack times and enhanced clinical therapeutic effect among children with asthma [41,42]. However, its mechanism is still unclear. Our previous study indicated that chiropractic therapy protected intestinal epithelium barrier and helped with anti-inflammation via promoting the production of specialized proresolving mediators in an animal experiment [43] and helped with the improvement of allergic diseases in clinical studies [22,23].

The present study revealed that chiropractic therapy significantly increased the richness and diversity of gut microbiota.
during AAI modelling. Moreover, chiropractic therapy increased the main butyrate-producing bacterial taxa such as families Bacteroidales S24-7 and Ruminococaceae and Roseburia [37,38]. Ruminococaceae can also produce acetate, which Roseburia can use to produce butyrate [44]. In addition, chiropractic therapy increased Prevotellaceae, which have important roles in nutrient absorption and immune regulation [45]. Chiropractic therapy also significantly decreased Lactobacillus. Therefore, chiropractic therapy might help the transition from a lactate-rich environment to a more adult-like butyrogenic environment in AAI immature rats. A healthy signature of SCFAs, especially more effective butyrate [11], can enhance the production of IFN-γ and IL-10 as well as provide protection against inflammation [12–14]. Therefore, the regulation of gut microbiota by chiropractic therapy might contribute to the enhanced production of IFN-γ and IL-10. The coexistence of increased IFN-γ and IL-10 production is key to anergy of antigen-specific T-cell during exposure to allergens [46]. These results might explain the attenuation of Th2 response and allergic airway inflammation by chiropractic therapy. In addition, our study further shows the importance of a diverse gut microbiota for immune homeostasis.

Animal experiments demonstrated that chiropractic therapy can improve the gastrointestinal dysfunction by regulating abnormal levels of brain-gut peptides (BGPS), including cholecystokinin-8 (CCK-8), ghrelin, vasoactive intestinal peptide (VIP), and substance P (SP), either in plasma, in the colon, or in the hypothalamus [47–49]. BGPs are critical neurotransmitters, which not only modulate gastrointestinal function [50,51] but also participate in the interaction between gut microbiota and brain-gut axis [52]. It is well known that the brain can also influence gut microbiota indirectly by regulating gastrointestinal motility and secretion, or directly by releasing signaling molecules [52]. Results of the present study and previous studies indicate that chiropractic therapy can optimize gut microbiota, partially via regulating BGPs production and the brain-gut axis.

However, there is certain limitation in our study. Firstly, our study did not evaluate the production of SCFAs; therefore, there is no direct evidence for a causal relationship between SCFAs and allergic airway inflammation. Secondly, our study did not test BGPs level since previous studies have already focused on it. Thirdly, the sample size of animals was small in our study and thus limited the statistical power. Further investigations are warranted based on this preliminary study.

**Conclusions**

To the best of our knowledge, this is the first study to assess the effect of massage on gut microbiota. In brief, our study indicated that AAI coexisted with dysbiosis of gut microbiota in immature rats and chiropractic therapy attenuated allergic airway inflammation and significantly modulated the gut microbiota. Chiropractic therapy might facilitate the development of a more adult-like butyrogenic environment, which can promote immunotolerance and inflammation attenuation. Our study supports chiropractic therapy as a good adjunct therapy for children with asthma and other allergic diseases.

**Conflict of interests**

None.

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