Compatibility of Achyranthes bidentata components in reducing inflammatory response through Arachidonic acid pathway for treatment of Osteoarthritis

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\textbf{ABSTRACT}

Achyranthes bidentata is a common traditional Chinese medicine (TCM) used in treating osteoarthritis (OA). The compatibility between effective components has now become a breakthrough in understanding the mechanism of TCM. This study aimed at determining the optimal compatibility and possible mechanism of \textit{Achyranthes bidentata} for OA treatment. Results showed that the adhesion score of the OA group is higher than NC group, and showed a trend of down-regulation in the intervention group. The CHI3L1 and IL-1\textbeta in joint fluid of the OA group was significantly increased compared to the sham operation group (NC group). Group G, I, and L exhibited significantly down-regulated CHI3L1, while groups C, F, I, K, and L exhibited reduced IL-1\textbeta. Joint adhesion, damage in cartilage, and synovial tissue was found in the OA model, cartilage tissue was found recovered in groups I, J, and L, and synovial tissue was recovered in group G, I, and L. Thus, group I and L were chosen for metabolite analysis, and indole-3-propionic acid was slightly up-regulated, while koeiginequineone A, prostaglandin H2, and 1-hydroxy-3-methoxy-10-methylacridonewere down-regulated in group I and L. According to functional analysis, the arachidonic acid (AA) metabolic pathway is enriched. Down-regulated expression of vital proteins in the AA metabolism pathway, such as PGE2 and COX2 in group I and L were verified. In conclusion, Hydroxyecdysone, Oleanolic acid, Achyranthes bidentata polysaccharide at a compatibility of 0.03-\textmu g/mg, 2.0-\textmu g/mg, 20.0-\textmu g/mg or 0.03-\textmu g/mg, 2.0-\textmu g/mg, 10.0-\textmu g/mg, respectively, may be the optimal compatibility of \textit{Achyranthes bidentata}.

\textbf{1. Introduction}

Osteoarthritis (OA) is a common chronic joint disorder involving pathological changes in the articular cartilage, synovium, subchondral bone, and other joint tissues [1]. Increasing age, obesity, nutritional factors, and genetic susceptibility were considered the primary risk factors contributing to the prevalence of OA [2,3]. The OA-involved inflammatory response is now well recognized, mainly manifested in joint pain, swelling, and stiffness [4]. Phenotypic shift, apoptosis, and aberrant expression of inflammation-related genes, including catabolic genes, such as NOS, COX, and MMPs were reported to be involved in inflammation-induced injuries [5,6]. The CHI3L1 is a glycoprotein secreted by articular chondrocytes, synoviocytes, and macrophages [7]. Studies have stated that chondrocytes of human osteoarthritic cartilage secrete the inflammation associated chitinecin CHI3L1 [8]. CHI3L1 was also reported to be functionally related to the development of osteoarthritis and could be used as markers [9]. IL-1\textbeta is considered one of the key cytokines involved in OA’s pathogenesis. Additionally, patients with OA have an elevated level of IL-1\textbeta in both synovial fluid, synovial membrane, cartilage, and subchondral bone layer [10].

Generally, NSAIDs and acetaminophen are considered first-line anti-inflammatory therapies for treating OA, cyclooxygenase-2 (COX-2) inhibitors and intra-articular steroid injections are also considered treatment options [11,12]. However, side effects, including gastrointestinal ulcers and perforations, cardiovascular complications, and opportunistic infections caused by immunosuppressive agents, limited the use of these anti-inflammatory...
therapies [13, 14]. Achyranthes bidentata, derived from natural plants, is a common traditional Chinese medicine (TCM) used in treating orthopedics and traumatology, has an anti-inflammatory effect on the articular cartilage of animal models of osteoarthritis [15]. Reports indicate that Achyranthes bidentata can reduce inflammatory responses by targeting TNF, IL-6, and TP53, suppressing glycolysis and apoptosis by MAPK/AKT signaling pathway, and decreasing osteoclastogenesis and bone resorption by inhibiting RANKL signaling [16–18]. Achyranthes bidentata contains three active components, including Hydroxyecdysone, Oleanolic acid, and Achyranthes bidentata polysaccharide, the concentration of these three components varies depending on the production area, processing, and testing methods [19]. To meet the criteria of precise medication, the compatibility between the effective components has now become a breakthrough in understanding the pharmacological mechanism of Chinese medicine [12]. However, no optimal therapeutic compatibility of the three components has been determined. Therefore, exploring the optimal therapeutic compatibility of the three active components, including Hydroxyecdysone, Oleanolic acid, and Achyranthes bidentata polysaccharide, and the possible pharmacological mechanism of Achyranthes bidentata is necessary.

Metabolomics are widely used in the prediction, diagnosis, and pharmacology in disease research, and play an important role in characterizing, efficacy evaluation, and pharmacological mechanism research in the application of TCM [20, 21]. Differential expressed metabolites (such as phosphatidylcholine, lysophosphatidylcholine, ceramides, myristate derivatives, and carnitine derivatives) as possible biomarkers related to the pathogenesis of OA have been found via metabolomics analysis by detecting various specimens of OA (cartilage, synovium, serum, plasma, urine, and synovial fluid, etc.) [22, 23]. Various metabolic pathways have affected amino acid metabolism in OA, including branched chain amino acids and arginine, and phospholipid metabolism involving conversion of phosphatidylcholine to lysophosphatidylcholine [24]. However, there is still a lack of relevant research focusing on metabolite changes during the treatment of OA using TCM.

In this study, a metabolomics strategy based on UPLC-TOF/MS for detecting the therapeutic compatibility and exploring the pharmacological mechanism of Achyranthes bidentata components was conducted, aiming to provide new ideas for OA treatment.

2. Material and methods

2.1 Construction of osteoarthritis rat model

Ninety male Wistar rats (250–300 g) were purchased from Guangdong Medical Laboratory Animal Center and fed in forevergen biosciences Co., Ltd (Guangzhou, China). The rats were randomly divided into an OA model group and a sham operation group after one week of adaptation. The OA rat model (N = 82) was constructed using injected 50-μl (3-mg) Monomer sodium iodoacetate (MIA) (sigma, USA) at a concentration of 60-mg /ml to unilateral knee joint, then stretched and bent for 30 seconds to fully spread the MIA to the entire joint. The sham operation group (N = 8) was injected with 50-μl normal saline [25]. Two weeks after the operation, we randomly sacrificed four rats from the OA model group and two rats from the sham operation group to determine whether the OA model was successfully constructed by conducting a pathological examination. The remaining 84 rats were divided into 14 groups: sham operation group (NC group) (N = 6), OA group (N = 6), and 12 intervention groups (A-L group) (N = 6 per group). The NC group and OA model group were treated with 0.9% saline, and the intervention groups were treated with three Achyranthes Bidentata components in 12 different ratios once daily for six weeks. The serum, cartilage tissue, and synovial tissue were collected and stored at −80°C until use. Our animal experiment was approved by the Animal Care and Usage Committee of Guangzhou Forevergen Biosciences Center (IACUC-G16052).

2.2 Preparation of 3 Achyranthes bidentata components in 12 different ratios

The three components of Achyranthes bidentata are: Achyranthes bidentata polysaccharide, Oleanolic acid (C27H44O7) (Yangling Ciyuan
Biotechnology, Xi’an, China), and Hydroxyecdyson (C30H48O3) (Zhenqiang Biotechnology, Shenzhen, China). The concentration gradient of these three components is as follows: high, medium, and low concentrations of Hydroxyecdyson were 0.12-μg/mg, 0.06-μg/mg, and 0.03-μg/mg. High and low concentrations of Oleanolic acid were 2.0-μg/mg and 1.0-μg/mg, 0.03-μg/mg. High and low concentrations of Achyranthes bidentata polysaccharide were 20.0-μg/mg and 10.0-μg/mg. Twelve intervention groups with different compatibility of Achyranthes bidentata are shown in Table 1.

### 2.3 Assessment of cartilage and synovial tissue damage

Adhesion rating scale of joint: Rats were sacrificed, incised the lateral side of the right patella, and fully exposed the knee joint, the adhesion in the joint was then evaluated using a semi-quantitative score. According to Rothkopf’s scoring standard: zero points for no adhesion; one point for film-like adhesion; two points for moderate adhesion; three points for dense fibrous adhesion [26]. Cartilage tissue damage was assessed following the Mankin’s score [27]. Synovitis was assessed according to KRENN score: by evaluating synovial cell proliferation, vascular proliferation, and inflammatory cell infiltration (0–3 levels, respectively), the sum of the three indices is the synovitis score. Synovitis was divided into three levels, namely, none (0–1 points), mild (2–3 points), moderate (4–6 points), and high (7–9 points). The higher the score, the more severe the inflammation [2].

### 2.4 Hematoxylin-eosin (HE) staining

Cartilage and synovial tissue were fixed >24 h, and then dehydrated in gradient alcohol with 75% ethanol for four hours, 85% ethanol for two hours, 90% ethanol for two hours, 95% alcohol for one hour, and 100% ethanol for one hour. The samples were embedded with paraffin at −20°C and cut into slices of 4-μm. We then used xylene to dewax the slices and dehydrated them in gradient ethanol vice versa. HE staining was conducted as follows: first, we stained the slices with hematoxylin for 3–5 minutes. Second, the slices were incubated with eosin for five minutes. Last, we dehydrated the slices in gradient ethanol (85% alcohol and 95% alcohol, each for five minutes). The samples were observed under a light microscope.

### 2.5 Enzyme-linked immune sorbent assay (ELISA)

The expression level of synovial fluid cartilage glycoprotein 39 (YKL-40) and interleukin-1beta was detected using Chitinase-3-like protein 1 (CHI3L1) ELISA kit (Wuhan Huamei, China), rat interleukin 1β (IL-1β) ELISA Kit (Wuhan Huamei, China), and Rat PGE2 Elisa Kit (Solarbio, China) according to the manufacturer’s instructions. A Thermo Scientific™ Multiskan™ FC microplate reader (ThermoFish, the USA) was used for detecting OD values at 450-nm.

### 2.6 Western blot analysis

Total proteins were isolated from rat cartilage tissue using RIPA lysis buffer supplemented with PMSF and proteinase inhibitor, the concentration of all proteins was determined using a BCA protein assay kit (Beyotime, China). SDS-polyacrylamide gel (4%–15%) electrophoresis was then used to separate 25-μg protein, and then transferred to PVDF membranes (Millipore, USA). A 5% BSA (BioFroxx, Germany) was used for PVDF membrane blocking. The PVDF membranes were incubated with primary antibodies against COX2 (1:1000, 4872, CST, USA), GAPDH(1:10,000, ab8245, Abcam, USA) at °C overnight, and secondary antibody for one hour after TBST washes at room temperature. The

| Group | A | B | C | D | E | F | G | H | I | J | K | L |
|-------|---|---|---|---|---|---|---|---|---|---|---|---|
| Hydroxyecdyson (μg/mg) | 0.12 | 0.12 | 0.12 | 0.12 | 0.06 | 0.06 | 0.06 | 0.06 | 0.03 | 0.03 | 0.03 | 0.03 |
| Oleanolic acid(μg/mg) | 2.0 | 2.0 | 1.0 | 1.0 | 2.0 | 2.0 | 1.0 | 1.0 | 2.0 | 2.0 | 1.0 | 1.0 |
| Achyranthes bidentata polysaccharide(μg/mg) | 20.0 | 10.0 | 20.0 | 10.0 | 20.0 | 10.0 | 20.0 | 10.0 | 20.0 | 20.0 | 20.0 | 10.0 |
membranes were incubated with enhanced chemiluminescence reagents before scanning. The gray values were analyzed using Image J software, and GAPDH was used as a reference gene as the control for normalization.

2.7 High-throughput metabolome detection

Metabolites were extracted from the serum samples with extraction solution (methanol: acetonitrile = 1:1 (V/V), containing the isotope-labeled internal standard mixture). LC-MS/MS analysis was conducted using the UHPLC system (Vanqish, Thermo Fisher Scientific), chromatographic separation of target compounds was conducted using a Waters ACQUITY UPLC BEH amide column (2.1 mm × 100 mm, 1.7-μm). The auto-sampler temperature was 4°C, with an injection volume of 3-μL. Phase A was the mobile phase consisting of 25-mmol/L ammonium acetate and 25-mmol/L ammonia hydroxide (pH = 9.75), and phase B was acetonitrile. Using gradient distillation: 0 ~ 0.5 min, 95% B; 0.5 ~ 7 min, 95%~65% B; 7 ~ 8 min, 65%~40% B; 8 ~ 9 min, 40% B; 9 ~ 9.1 min, 40%~95% B; 9.1 ~ 12 min, 95% B. Mobile phase flow rate: 0.5-mL/min, column temperature: 30°C. The Thermo Q Exactive HFX mass spectrometer can conduct primary and secondary mass spectrometry data acquisition in the control of acquisition software (Xcalibur, Thermo), with parameters as follows: Sheath gas flow rate: 30 Arb, Aux gas flow rate: 25 Arb, Capillary temperature: 350°C, Full ms resolution: 60,000, MS/MS resolution: 7500, Collision energy: 10/30/60 in NCE mode, Spray Voltage: 3.6 kV (positive) or −3.2 kV (negative). The raw data were converted to the mzXML format using ProteoWizard. R program package based on XCMS was used for peak identification, peak extraction, peak alignment, and integration, etc. Then, an in-house MS2 database (Biotree DB) was applied in metabolite annotation with a cutoff value at 0.3.

2.8 Orthogonal partial least squares method-discriminant analysis (OPLS-DA)

SIMCA software (V16.0.2, Sartorius Stedim Data Analytics AB, Umea, Sweden) was used to perform logarithmic (LOG) conversion and UV formatting. First, we conducted Orthogonal partial least squares discriminant analysis (OPLS-DA) based on the first principal component and verified the quality of the model with 7-fold cross-validation. Then, we used the cross-validated R2Y (the interpretability of the model to categorical variable Y) and Q2 (model’s predictability) to evaluate the effectiveness of the model. Finally, different random Q2 values were obtained by randomly changing the order of the categorical variable Y to further test the validity of the model.

2.9 Differential metabolite screening and functional analysis

Multivariate statistical analysis was used to analyze the metabolite data. Metabolites with a p-value of Student’s t-test ≤ 0.05, and a Variable Importance in the Projection (VIP) value of the OPLS-DA model ≥ 1 were considered significantly different. The differential metabolites were then analyzed using a volcano plot. We calculated the Euclidean distance matrix for the quantitative values of the differential metabolites and clustered the differential metabolites with a complete linkage method. Key pathways related to differential expressed metabolites were analyzed using the Kyoto Encyclopedia of Genes and Genomes (KEGG) and PubChem (http://pubchem.ncbi.nlm.nih.gov). Database for Annotation, Visualization, and Integrated Discovery (DAVID: https://david.ncifcrf.gov/) was applied for analyzing the enrichment of the KEGG pathway of the differentially expressed metabolites [28].

2.10 Statistical analysis

The data were presented as mean ± SD. One-way ANOVA was used to analyze the expression levels of inflammatory cytokines, semi-quantitative analysis of the pathological test, and the proteins of target metabolic pathways. The Bonferroni multiple-comparison method was used for mean comparison. Statistical significance was calculated using IBM SPSS Statistics (version 22.0, IBM SPSS Inc., Chicago, IL).
3. Results

Achyranthes bidentate has an anti-inflammatory effect on the articular cartilage of animal models of osteoarthritis, it contains three active components, including Hydroxyecdysone, Oleanolic acid, and Achyranthes bidentata polysaccharide, no optimal therapeutic compatibility of the three components have been determined. Our study aimed at determining the optimal compatibility and possible mechanism of Achyranthes bidentate for OA treatment, and finally, it found that Hydroxyecdysone, Oleanolic acid, and Achyranthes bidentata polysaccharide at compatibility of 0.03-μg/mg, 2.0-μg/mg, 20.0-μg/mg, and 0.03-μg/mg, 2.0-μg/mg, 10.0-μg/mg may be the optimal compatibility of Achyranthes bidentate.

3.1 Achyranthes Bidentata component information

Three components of Achyranthes bidentata were purchased and identified by the m/z value. MS detection of the monomer components of Achyranthes bidentata showed that the molecular formula of Hydroxyecdysone was C30H48O3, and the m/z value was 11.7 (Figure 1a); the molecular formula of Oleanolic acid was C27H44O7, and the m/z value was 25.19 (Figure 1b). Achyranthes bidentata polysaccharide is a compound component with no molecular formula. It is a high-content active ingredient in the forage plant Achyranthes bidentate, mainly composed of three monosaccharide components: glucose, mannose, and fructose.

3.2 Joint adhesion and inflammatory level after treatment

Joint adhesion and inflammatory levels of 14 groups, including the NC group, the OA group, and 12 intervention groups (A-L group) were detected on Day 56 after model construction (Figure 2a). Rothkopf adhesion semi-quantitative scoring showed that the adhesion score of the OA group is higher compared to the NC group, and showed a trend of down-regulation in the intervention groups. Group H and I have the most obvious down-regulation; however, no statistical significance has been identified (Figure 2b). The CHI3L1 in the joint fluid of the OA group was significantly increased compared to the NC group and was significantly down-regulated in groups G, I, and L (Figure 2c). The expression of IL-1β in the OA group was significantly up-regulated compared with the NC group, and the expression in groups C, F, I, K, and L was significantly reduced (Figure 2d).

3.3 Pathological changes of cartilage tissue and synovial tissue

The HE staining of cartilage tissue showed that the NC group had intact cartilage tissue, and the cells were evenly distributed. Compared to the NC group, the OA group showed destroyed perichondrium, with shifted and thinned fiber tissue, as well as disordered chondrocytes, with partial calcifications. The A-L group, the intervention groups of Achyranthes bidentate, can reverse the cartilage tissue damage to a certain extent, in which groups I, J, and L have the most significant recovery (Figure 3a).
The Mankin’s score for cartilage tissue assessment of the OA group was significantly increased compared to the NC group, and showed significant reductions in the intervention groups, especially in groups I and L (Figure 3b).

The HE staining of synovial tissue showed aligned synovial tissue cells in the NC group, while synovial cell proliferation, fibrous tissue proliferation, and inflammatory cell infiltration were seen in the OA group and showed alleviation in the intervention groups (Figure 3c). KRENN score of synovitis tissue evaluation was reduced in intervention groups, especially in groups G, I, and L (Figure 3d).

3.4 Hierarchical cluster analysis of differentially expressed metabolites in group I and L

Metabolomics can discover significantly changed metabolites and reveal the possible mechanism of combability of Achyranthes bidentata in OA treatment. Based on the above results, group I and group L showed the most effective efficacy and were chosen for further metabolic analysis. The OPLS-DA score showed that sample distinction is significant, and the samples are all within the 99% confidence interval (Hotelling’s T-squared ellipse) (Figure 4a).

The metabolites whose abundance is down-regulated or up-regulated in the NC group and I group compared with the OA group was chosen, the hierarchical cluster analysis of the differential expressed metabolites between the I group, NC group, and OA group was divided into four clusters with one up-regulated and three down-regulated (Figure 4b). The metabolites whose abundance is down-regulated or up-regulated in the NC group and L group compared with the OA group were chosen, the hierarchical cluster analysis of differential expressed metabolites between the L group, NC group, and OA group was divided into three clusters with one up-regulated and two down-regulated (Figure 4c). Indole-3-propionic acid was up-regulated, while 1-hydroxy-3-methoxy-10-methylacridone, koeiginequinone A, and prostaglandin H2 were down-regulated as determined in groups I and L.

3.5 Arachidonic acid metabolic pathway involved in the compatibility of Achyranthes bidentata

Using the differentially expressed metabolites found in this study, we predicted the metabolic pathways. The enrichment analysis showed that

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**Figure 2.** Efficacy evaluation of osteoarthritis after Achyranthes Bidentata treatment. (a) The research flow chart of our study: Rat OA model was constructed by injecting MIA, 12 intervention groups was given with 12 different compatibility of the Achyranthes bidentata for six weeks (A-L group, N = 6 per group). The osteoarthritis was detected on Day 56. (b) The Achyranthes Bidentata intervention group showed reduced Rothkopf adhesion score compared with the OA group. (c) ELISA analysis showed that group G, I, and L showed significantly down regulated CHI3L1. (d) ELISA analysis showed that the expression of IL-1β in group C, F, I, K, and L were significantly reduced compared with the OA group. Compared with NC group, *p < 0.05; compared with OA group, **p < 0.05.
the arachidonic acid (AA) metabolic pathway was significantly enriched in group I (Figure 5a).

AA pathway plays a major role in the inflammation metabolic network, including two main metabolic pathways: (1) Through the modulation of cyclooxygenase (COX) and generating several prostaglandins (PEGs); (2) Through the action of lipoxygenase (LOX) in the production of leukotrienes and lipid peroxides. Considering that prostaglandin H2 was up-regulated in the OA group, and down-regulated in both groups I and L in our study, we further verified the expression of key

Figure 3. Pathological changes of cartilage tissue and synovial tissue. (a) HE staining showed cartilage tissue repair in Achyranthes Bidentata intervention group compared with the OA group. (b) Group I and L showed the most significant reduction in Mankin’s score of cartilage tissue assessment. (c) HE staining showed synovial tissue repair in Achyranthes Bidentata intervention group compared with the OA group. (d) Group G, I, and L showed the most significant reductions in the KREEN score of synovitis tissue evaluation. Compared with the NC group, \(^*p < 0.05\); compared with OA group, \(^b p < 0.05\).
enzymes in COX metabolic pathway, including PEG2 and COX2. ELISA analysis indicated that the expression of PGE2 was significantly reduced in groups I and L (Figure 5b). The WB analysis showed that the expression of COX2 was also reduced in groups I and L (Figure 5c and d).

4. Discussion

Achyranthes bidentata is widely used in OA treatment, however, its complex mixture makes it difficult to meet the standards of precision medicine during the past decade. Current research has suggested that Hydroxyecdysone can enhance protein synthesis and promote cell regeneration, while Achyranthes bidentata polysaccharide can enhance the host immune response [29,30]. Thus, we believed that research on the compatibility of Achyranthes bidentata was the key to clarifying this herbal medicine’s pharmacological and pharmacodynamic mechanisms.

OA can be induced by cytokines, which leads to various matrix-degrading enzymes to then be produced, and the cytokines and chemokines secretion through signal transduction pathways were further amplified, causing joint pain, swelling, and synovitis [4,31]. The main elevated cytokines in patients with OA, include IL-1β, TNF-α, and IL-6, while other cytokines (including IL-15, IL-17, IL-18, IL-21, and LIF), Chemokines (including CCL5, IL-8, GRO-α, and MCP-1), and anti-inflammatory factors (IL-4, IL-10, and IL-13) also showed an up-regulation [32]. It is worth noting that CHI3L1, an important inflammatory factor, is reported to be significantly increased in the serum, synovial fluid, and cartilage tissue of patients with OA [33]. In this study, we found that IL-1β and CHI3L1 were significantly higher in the OA group compared to the NC group. The expression of IL-1β was significantly reduced in groups C, F, I, K, and L with different compatibility of Achyranthes bidentata, while CHI3L1 was significantly reduced only in groups G, I, and L, which is consistent with the previous report. Alternatively, joint adhesion, damage in cartilage and synovial tissue found in the OA model also showed significant recovery in the compatibility of Achyranthes bidentata...
intervention groups, especially in groups I and L. The pharmacological and pharmacodynamic mechanisms of *Achyranthes bidentata* are difficult to clarify due to their complex structure, which is not enough to reach the criteria of precise medication. As we know, the optimal compatibility of TCM components, including the compatibility of different drugs and the compatibility of different isolates from one drug, can improve its efficacy and reduce its toxicity [34]. Previous studies used rat chondrocytes pretreated with *Achyranthes bidentata Blume* at 3-μg/mL, 10-μg/mL, and 30-μg/mL, and subsequently stimulated with IL-1β (10-ng/mL) and discovered that *Achyranthes bidentata Blume* might be a potential drug for treating osteoarthritis [35]. β-Ecdysterone was reported to suppress interleukin-1β-induced apoptosis and inflammation in rat chondrocytes through inhibition of NF-κB signaling pathway [36]. Oleanolic acid also showed peripheral antinociceptive and anti-inflammatory effects in a rat model of OA [37]. While no optimal therapeutic compatibility of the three components of *Achyranthes bidentata* has been determined. Thus, we conducted 12 differential compatibility of *Achyranthes bidentata* to determine the optimal compatibility of three *Achyranthes bidentata* components, so that each component can play a synergistic effect to each other, to reach the maximum efficacy and safety. Our results indicated that the compatibility of *Hydroxyecdysone*, *Oleanolic acid*, and *Achyranthes bidentata polysaccharide* in groups I and L at the level of 0.03-μg/mg, 2.0-μg/mg, 20.0-μg/mg and 0.03-μg/mg, 0.03-μg/mg, 10.0-μg/mg can play an anti-inflammatory role and effective reverse of joint damage in OA treatment. We further conducted metabolite analysis for revealing the pharmacological mechanism of the compatibility of *Achyranthes bidentata* in OA treatment.

Metabolomics can discover significantly changed metabolites, and may reveal the possible

**Figure 5.** Arachidonic acid metabolic pathway may participate in *Achyranthes bidentata* treatment. (a) Bubble chart of enrichment analysis of differential metabolite pathways in group I. The arachidonic acid metabolic pathway was significantly enriched. (b) ELISA analysis of PGE2 was significantly reduced in group I and L. (c and d) The expression of COX2 was verified to be reduced in groups I and L using WB analysis. Compared with NC group, *a* p < 0.05; compared with OA group, *b* p < 0.05.
mechanism of combability of Achyranthes bidentata in OA treatment [38]. In this study, hierarchical cluster analysis of the differential expressed metabolites showed that indole-3-propionic acid (IPA) was up-regulated, while 1-hydroxy-3-methoxy-10-methylacridonew, koeiginequinone A, and prostaglandin (PG) H2 were down-regulated as determined in groups I and L. IPA is secreted by the commensal bacteria located in the intestines, absorbed by intestinal epithelial cells, diffused into the bloodstream, and play an anti-inflammation role in the intestine, brain, and liver, indicating the possible modulation of IPA on the compatibility of Achyranthes bidentata as group I and L in OA treatment. Additionally, PGs have been recognized to play an important role in articular diseases, indicating the possibility of PGs as therapeutic biomarkers of Achyranthes bidentata with compatibility as group I and L in OA treatment.

Further enrichment analysis showed that AA metabolic pathway is significantly enriched in both groups I and L, indicating that AA pathway participates in the pharmacologic mechanisms of compatibility of Achyranthes bidentata as group I and L in OA treatment. Arachidonic acid (AA) is a long-chain unsaturated fatty acid, which generates unstable PGG2 and PGH2 through the cyclooxygenase (COX) pathway, and generates two types of essential mediators, including prostaglandins (PGEs) (such as PGD and PGE2) and thromboxanes (TXAs) [41]. Among the PGs, PGE2 is widely distributed in multiple cells and tissues and acts as a major mediator of inflammation in rheumatoid arthritis (RA) [42]. PGE2 was mainly produced by synovial cells, chondrocytes, and macrophages/monocytes, accompanied by up-regulation of COX-2 level in activated articular cells [43,44]. We then verified the down-regulated expression of PEG2 and COX2 in the compatibility of Achyranthes bidentata as groups I and L, indicating that PEG2 and COX2 could be used as therapeutic biomarkers and that AA metabolic pathway may participate in the optimal compatibility of Achyranthes bidentata in the OA treatment. Thus, we suggested that the optimal compatibility of Achyranthes bidentata may be regarded as an anti-inflammatory agent in OA treatment by interfering with the activation of inflammation or blocking the inflammatory response through the AA pathway.

5. Conclusion

In summary, Hydroxyecdysone, Oleanolic acid, and Achyranthes bidentata polysaccharide at a compatibility of 0.03-μg/mg, 2.0-μg/mg, 20.0-μg/mg and 0.03-μg/mg, 2.0-μg/mg, 10.0-μg/mg may be the optimal compatibility of Achyranthes bidentata. The pathological mechanism of compatibility of Achyranthes bidentata in OA treatment is related to the inhibition of the AA pathway. These findings expand our understanding of the optimal compatibility and pharmacological mechanism of Achyranthes bidentata in OA treatment.

Availability of Data and Material

The data in this study are available from the corresponding author by reasonable request.

Ethical approval statement

Our animal experiment was approved by the Animal Care and Usage committee of Guangzhou Forevergen Biosciences Center (IACUC-G16052).

Authors’ contributions

All authors contributed to the study conception and design. Material preparation, data collection, and analysis were performed by Dujun Ma, Liping Peng, Yuan Li, Zhouwei Liao, and Tian Yu. The first draft of the manuscript was written by Zanzhu Li and all authors commented on previous versions of the manuscript. All authors read and approved the final manuscript.

Disclosure statement

No potential conflict of interest was reported by the author(s).

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