Potential Molecular Mechanisms of Ephedra Herb in the Treatment of Nephrotic Syndrome Based on Network Pharmacology and Molecular Docking

Tianwen Yao, Qingliang Wang, Shisheng Han, Yan Lu, Yanqiu Xu, and Yi Wang

1Department of Nephrology, Yueyang Hospital of Integrated Traditional Chinese and Western Medicine, Shanghai University of Traditional Chinese Medicine, Shanghai 200437, China
2Department of Emergency, Yueyang Hospital of Integrated Traditional Chinese and Western Medicine, Shanghai University of Traditional Chinese Medicine, Shanghai 200437, China

Correspondence should be addressed to Yi Wang; drwangyi0110@126.com

Received 26 December 2021; Revised 30 April 2022; Accepted 11 June 2022; Published 5 July 2022

Objective. To explore the possible mechanisms of Ephedra herb (EH) in the treatment of nephrotic syndrome (NS) by using network pharmacology and molecular docking in this study.

Methods. Active ingredients and related targets of EH were obtained from the Traditional Chinese Medicine Systems Pharmacology (TCMSP) database, and the gene names corresponding to the proteins were found through the UniProt database. Then, target genes related to NS were screened out from GeneCards, PharmGKB, and OMIM databases. Next, the intersection targets were obtained successfully through Venn diagram, which were also seen as key target genes of EH and NS. Cytoscape 3.9.0 software was used to construct the effective "active ingredient-target" network diagram, and "drug-ingredient-target-disease (D-I-T-D)" network diagram. After that, the STRING database was used to construct a protein-protein interaction (PPI) network. Furthermore, Gene Ontology (GO) and Kyoto Encyclopedia of Genes and Genomes (KEGG) pathway enrichment involved in the targets were performed by the DAVID database and ClueGO plugin in Cytoscape. Finally, AutoDockTools software was used for molecular docking to verify the binding strength between main active ingredients and key target proteins.

Results. A total of 22 main active ingredients such as quercetin, kaempferol, luteolin, and naringenin were obtained, which could act on 105 targets related to NS. Through PPI network, 53 core targets such as AKT1, TNF, IL6, VEGFA, and IL1B were found, which might play a crucial role in the treatment of NS. Meanwhile, these targets were significantly involved in PI3K-Akt signaling pathway, TNF signaling pathway, AGE-RAGE signaling pathway, hepatitis B, and pathways in cancer through GO and KEGG enrichment analysis. The docking results indicated that active ingredients such as kaempferol, luteolin, quercetin, and naringenin all had good binding to the target protein AKT1 or TNF. Among them, luteolin and naringenin binding with AKT1 showed the best binding energy (-6.2 kcal/mol).

Conclusion. This study indicated that the potential mechanism of EH in treating NS may be related to PI3K-Akt signaling pathway, TNF signaling pathway, and AGE-RAGE signaling pathway, which provided better approaches for exploring the mechanism in treating NS and new ideas for further in vivo and in vitro experimental verifications.

1. Introduction

Nephrotic syndrome (NS) is characterized by massive proteinuria (greater than 3.5 g per day), hypoalbuminemia (less than 30 g/L), hyperlipidemia, systemic edema, and various complications [1]. As one of the most frequent glomerular diseases affecting more than millions of people worldwide, NS is caused by the dysfunctional glomerular filtration barrier [2]. In the last decades, NS has become a prevalent disease associated with high morbidity in 20%-40% of children [3, 4]. Nowadays, steroids and immunosuppressive drugs are the most common methods in the treatment of NS at home and abroad [5]. Usually, NS responds well to steroids [6]. However, frequent recurrences are common, which will require multidrug therapy with long-term side effects [7]. Obviously, those problems not only increase the difficulty...
of treatment but also accelerate the development of end-stage renal disease (ESRD), which can be a catastrophic event for the patient, family, and even society [8]. Therefore, it is crucial to search for more advanced and natural drugs with less side effects for the treatment of NS.

Ephedra herb (EH), also named as mahuang in Chinese, belongs to the stems of Ephedra sinica Stapf, Ephedra intermedi Schrenk & C.A.Mey., and Ephedra equisetina Bunge [9]. EH is one of the most ancient herbs recorded in Sheng Nong’s Herbal Classic, which plays an important role in terms of relieving exterior syndrome by diaphoresis, as well as inducing diuresis to alleviate edema. Chemical analyses demonstrate that a variety of specific ingredients have been extracted from EH, including flavonoids, alkaloids, organic acids, polysaccharides, tannins, volatile oils, and other active ingredients [10]. EH has been used for medicinal purposes for thousands of years with its biological and pharmacological properties, such as antioxidative, antimicrobial, anti-inflammatory, antiallergic, and antiproliferative effects [11–13]. Recently, it is widely used to treat bronchial asthma, cardiovascular related diseases, immune system diseases, cancer, and other diseases [14–16]. Moreover, it is reported that EH has positive effects on the treatment of NS through reducing protein in the urine, alleviating kidney functional and structural impairment [17, 18]. However, the mechanism of EH to treat NS is unclear, and the targets and pathways of its effective chemical ingredients are still unknown, which need to be further studied.

As a new technology to analyze the potential mechanism of compounds or single herbs, network pharmacology has been widely used in the field of TCM. It is known that network pharmacology is an emerging discipline that designs drugs based on systems biology theory and biological system network analysis [19]. So, network pharmacology is aimed at clarifying the molecular mechanism of drugs, providing guidance for the prediction of TCM, and exploring the relationships between drugs and diseases from a macro perspective [20], which is especially suitable for conducting multicomponent, multitarget, and synergistic studies of TCM [21]. Molecular docking is a computational method that can be used to study the binding interaction between molecule ligands and target proteins [22]. This helps not only in understanding the mechanism of their biological action but also accelerating the process of drug discovery [23].

In this study, network pharmacology and molecular docking methods were both used to screen the possible targets of EH intervention in NS and establish the network of EH active ingredients-targets-pathways-diseases, which may provide a basis for relevant experimental researches (Figure 1).

2. Materials and Methods

2.1. Screening Active Ingredients of EH. Traditional Chinese Medicine Systems Pharmacology (TCMSP, http://tcmspw.com/tcmspsearch.php) database is an open and accessible database resource [24]. Furthermore, it acts as a unique systems pharmacology platform of Chinese herbal medicines to explore the relationships between drugs, targets, and diseases. The keyword “Mahuang” was inputted into the search box to retrieve all the active ingredient data of EH. Because oral bioavailability (OB) and drug-likeness (DL) are important evaluation indexes for drugs to participate in the process of ADME (absorption, distribution, metabolism, and excretion), we took OB and DL as screening conditions [25]. In order to obtain active ingredients of EH more comprehensively, we set the condition of OB ≥ 30% and DL ≥ 0.18 to select them [26].

2.2. Screening Active Ingredient Targets of EH. We took “mahuang” as the keyword and searched “Related Targets” in the TCMSP database to find target proteins corresponding to the active ingredients of EH [27]. Then, the gene names corresponding to those target proteins were specified through UniProt (https://www.uniprot.org/) with Homo sapiens as the selected species. The Cytoscape software (version 3.9.0) was used to construct a network between active ingredients of EH and their corresponding target points. In the network, each node represented a target gene or molecule, and the connecting line between two nodes represented the relationship between them. It meant the stronger the degree of the node, the greater the role of the target in the network [28].

2.3. Collection of Target Genes Related to NS. With the keyword of “nephrotic syndrome”, target genes related to NS were collected from GeneCards (https://www.genecards.org), PharmGKB (https://www.pharmgkb.org), and OMIM (https://www.omim.org) databases. The three databases are recognized as good references for the collection of disease targets. Results accessed from GeneCards were screened for the relevance score ≥ 1.00 as the screening index. The repeated target genes corresponding to NS and the active ingredients of EH were deleted by Excel. Finally, the intersection targets were obtained successfully through Venn diagram in the website http://bioinformatics.psb.ugent.be/webtools/Venn/.

2.4. Construction of Drug-Ingredient-Target-Disease (D-I-T-D) Network. We obtained intersection target genes of EH and NS by Venn diagram. Then, a visual comprehensive network (D-I-T-D) was built by Cytoscape software (version 3.9.0) based on the interaction between drug (EH), active ingredients, intersection target genes, and diseases (NS).

2.5. Construction of Protein-Protein Interaction Network. The STRING database (https://string-db.org/) is an online protein-protein interaction (PPI) analysis database. It can collect and integrate data about known and predicted protein-protein associations from many organisms, including both direct and indirect interactions [29]. Therefore, the intersection targets of EH and NS were inputted into the STRING database to construct the PPI network. In order to ensure the high confidence of information, the minimum interaction score was set as 0.4, the species was set as Homo sapiens, and the isolated proteins were hidden.
Figure 1: A comprehensive strategy diagram for the study of the mechanism of EH acting on NS.
Table 1: Potential active ingredients of EH.

| Number | Mol ID   | Molecule name         | OB (%) | DL  |
|--------|----------|-----------------------|--------|-----|
| 1      | MOL010788| Leucopelargonidin     | 57.97  | 0.24|
| 2      | MOL002823| Herbacetin            | 36.07  | 0.27|
| 3      | MOL010489| Resivit               | 30.84  | 0.27|
| 4      | MOL000422| Kaempferol            | 41.88  | 0.24|
| 5      | MOL004798| Delphinidin           | 40.63  | 0.28|
| 6      | MOL000098| Quercetin             | 46.43  | 0.28|
| 7      | MOL000006| Luteolin              | 36.16  | 0.25|
| 8      | MOL00358 | Beta-sitosterol       | 36.91  | 0.75|
| 9      | MOL00449 | Stigmasterol          | 43.83  | 0.76|
| 10     | MOL00492 | (+)-Catechin          | 54.83  | 0.24|
| 11     | MOL001494| Mandenol              | 42.00  | 0.19|
| 12     | MOL001506| Supraene              | 33.55  | 0.42|
| 13     | MOL001755| 24-Ethylcholdest-4-en-3-one | 36.08 | 0.76|
| 14     | MOL001771| Poriferast-5-en-3beta-ol | 36.91 | 0.75|
| 15     | MOL002881| Diosmetin             | 31.14  | 0.27|
| 16     | MOL004328| Naringenin            | 59.29  | 0.21|
| 17     | MOL004576| Taxifolin             | 57.84  | 0.27|
| 18     | MOL005043| Campest-5-en-3beta-ol | 37.58  | 0.71|
| 19     | MOL005190| Eriodictyol           | 71.79  | 0.24|
| 20     | MOL005573| Genkwanin             | 37.13  | 0.24|
| 21     | MOL005842| Pectolinarigenin      | 41.17  | 0.30|
| 22     | MOL007214| (+)-Leucoyanidin      | 37.61  | 0.27|
| 23     | MOL011319| Truflex OBP           | 43.74  | 0.24|

2.6. Gene Ontology (GO) and Kyoto Encyclopedia of Genes and Genomes (KEGG) Pathway Enrichment. To further explore the pathways of the disease, the intersection targets of EH and NS were entered in the DAVID database (https://david.ncifcrf.gov/).

It is an online biological knowledge base and analytic tool, which is mainly used to obtain various biological information, including gene functional classification, functional annotation, and enriched pathways [30]. GO enrichment analysis is composed of biological process (BP), cellular component (CC), and molecular function (MF). KEGG is a knowledge base used to describe molecular interactions and reveal known metabolic pathways. The species was defined as Homo sapiens to perform GO and KEGG pathway enrichment analysis. According to the gene number and P value, the top 10 GO and KEGG items were selected for further analysis, respectively. Meanwhile, KEGG pathway enrichment analysis was also performed using ClueGO plugin in Cytoscape.

2.7. Molecular Docking of Active Ingredients with Key Targets. We selected two targets with the largest value in the PPI network as the receptors and chose the corresponding active ingredients as the ligands for molecular docking verification. Firstly, the PubChem database (https://pubchem.ncbi.nlm.nih.gov) was used to obtain the 2D structure diagrams of active ingredients, and we transformed them into 3D structure diagrams through Chem3D software (version 14.0). Then, protein crystal forms corresponding to the targets were obtained from the PDB database (https://www.rcsb.org/). After that, we imported them into PyMOL software (version 2.5) to remove water molecules. Next, the ingredient and corresponding protein were both processed by AutoDockTools 1.5.7 software and saved in pdbqt format as a ligand and a receptor, respectively. Finally, docking work was performed to analyze the results using PyMOL software and AutoDock Vina software (version 1.1.2).

3. Results

3.1. Screening Active Ingredients of EH. We used “mahuang” as the keyword to search active ingredients in the TCMSP database and found 363 kinds of chemical ingredients. With the screening conditions of OB ≥ 30% and DL ≥ 0.18, a total of 23 potential active ingredients satisfying the conditions were obtained in Table 1. Then, we screened out ingredients without corresponding targets and confirmed 22 potential active ingredients of EH, respectively: leucopelargonidin, herbacetin, resivit, kaempferol, delphinidin, quercetin, luteolin, beta-sitosterol, stigmasterol, (+)-catechin, mandenol, 24-ethylchololest-4-en-3-one, poriferast-5-en-3beta-ol, diosme- tin, naringenin, taxifolin, campest-5-en-3beta-ol, eriodictyol, genkwanin, pectolinarigenin, (+)-leucoyanidin, and truflex OBP. They were the main active ingredients of EH in the treatment of NS. Finally, their chemical abstracts service (CAS) number, chemical structure, molecular formula, and molecular weight are shown in Table 2.

3.2. Screening Active Ingredient Targets of EH and Construction of Active Ingredient-Target Network. 22 effective chemical ingredients of EH were inputted into the TCMSP platform, and 226 corresponding proteins were obtained. Then, target gene names corresponding to those proteins were found by UniProt. Finally, the active ingredients and targets were imported into Cytoscape 3.9.0 software to construct active ingredient-target network. As shown in Figure 2, the network between EH active ingredients and corresponding targets consisted of 249 nodes (1 herbal medicine node, 22 active ingredient nodes, and 226 corresponding target nodes) and 515 edges. The yellow inverted triangle node represented EH, the pink diamond nodes represented the active ingredients of EH, and the blue oval nodes represented the target genes. It was obvious that quercetin has the most number of targets. It could also be concluded from the figure that each active ingredient of EH had multiple targets, and each target could correspond to multiple active ingredients, which deeply showed the characteristics of multicomponent and multitarget actions of EH.

3.3. Collection of Gene Targets Related to NS. The keyword “nephrotic syndrome” was used to search relevant genes in GeneCards, PharmGKB, and OMIM databases. Then, a total of 2089 genes were obtained after deletion. Finally, intersection targets were acquired by overlapping the above targets

BioMed Research International
Table 2: Information table of active ingredients of EH.

| CAS number | Ingredient name | Chemical structure | Molecular formula | Molecular weight |
|------------|----------------|--------------------|-------------------|-----------------|
| 520-17-2   | Leucopelargonidin | ![Chemical structure](image1.png) | C_{15}H_{14}O_{6}  | 290.27          |
| 527-95-7   | Herbacetin      | ![Chemical structure](image2.png) | C_{15}H_{10}O_{7} | 302.25          |
| 480-17-1   | Resivit         | ![Chemical structure](image3.png) | C_{15}H_{14}O_{7} | 306.27          |
| 520-18-3   | Kaempferol      | ![Chemical structure](image4.png) | C_{15}H_{10}O_{6} | 286.24          |
| 13270-61-6 | Delphinidin     | ![Chemical structure](image5.png) | C_{15}H_{10}O_{7} | 303.24          |
| 117-39-5   | Quercetin       | ![Chemical structure](image6.png) | C_{15}H_{10}O_{7} | 302.24          |
| 491-70-3   | Luteolin        | ![Chemical structure](image7.png) | C_{15}H_{10}O_{6} | 286.24          |
| 83-46-5    | Beta-sitosterol | ![Chemical structure](image8.png) | C_{29}H_{50}O     | 414.71          |
| 83-48-7    | Stigmasterol    | ![Chemical structure](image9.png) | C_{29}H_{46}O     | 412.69          |
| CAS number | Ingredient name           | Chemical structure | Molecular formula | Molecular weight |
|------------|---------------------------|--------------------|-------------------|------------------|
| 154-23-4   | (+)-Catechin              | ![Chemical Structure](image1.png) | C_{15}H_{14}O_{6} | 290.27           |
| 544-35-4   | Mandenol                  | ![Chemical Structure](image2.png) | C_{20}H_{36}O_{2} | 308.50           |
| 67392-96-5 | 24-Ethylcholest-4-en-3-one | ![Chemical Structure](image3.png) | C_{29}H_{48}O | 412.69           |
| 18525-35-4 | Poriferast-5-en-3beta-ol | ![Chemical Structure](image4.png) | C_{29}H_{50}O | 414.71           |
| 520-34-3   | Diosmetin                 | ![Chemical Structure](image5.png) | C_{16}H_{12}O_{6} | 300.26           |
| 480-41-1   | Naringenin                | ![Chemical Structure](image6.png) | C_{15}H_{12}O_{5} | 272.25           |
| 480-18-2   | Taxifolin                 | ![Chemical Structure](image7.png) | C_{15}H_{12}O_{7} | 304.25           |
| 474-62-4   | Campest-5-en-3beta-ol     | ![Chemical Structure](image8.png) | C_{20}H_{44}O | 400.68           |
of EH and NS via the Venn diagram. As shown in Figure 3, there were 105 intersection targets of EH and NS.

3.4. Construction of Drug-Ingredient-Target-Disease (D-I-T-D) Network. The drug (EH), active ingredients, target genes, and disease (NS) were imported into Cytoscape 3.9.0 software to construct the D-I-T-D network. As shown in Figure 4, the yellow inverted triangle node represented EH, the purple octagon node represented NS, the pink diamond nodes represented the active ingredients of EH, and the 105 blue oval nodes represented the overlapping genes between disease and drug. It could be seen from the figure that querctin had the most number of overlapping genes. Furthermore, it clearly showed how EH treated NS through active ingredients and target genes.

3.5. Construction of Protein-Protein Interaction Network. We imported 105 overlapping targets into the STRING database to obtain the PPI relationship. As shown in Figure 5(a), the network had a total of 104 nodes and 2064 edges after removing the free one. In addition, the mean degree value of those overlapping targets was 39.70. According to the degree value, they were ranked from high to low. As shown in Figure 5(b), we selected targets with value greater than 39.70 and finally obtained 53 core targets. In terms of degree value, the top 10 key target proteins were RAC-alpha serine/threonine-protein kinase (AKT1), tumor necrosis factor (TNF), interleukin 6 (IL6), vascular endothelial growth factor a (VEGFA), interleukin 1 beta (IL1B), tumor protein P53 (TP53), mitogen-activated protein kinase 3 (MAPK3), caspase 3 (CASP3), jun proto-oncogene, AP-1 transcription

| CAS number | Ingredient name       | Chemical structure | Molecular formula | Molecular weight |
|------------|-----------------------|--------------------|-------------------|------------------|
| 552-58-9   | Eriodictyol           | ![Eriodictyol](image) | C_{15}H_{12}O_{6}  | 288.25           |
| 437-64-9   | Genkwanin             | ![Genkwanin](image) | C_{16}H_{12}O_{5}  | 284.26           |
| 520-12-7   | Pectolinarigenin      | ![Pectolinarigenin](image) | C_{17}H_{14}O_{6}  | 314.29           |
| 69256-15-1 | (+)-Leucocyanidin     | ![(+)-Leucocyanidin](image) | C_{15}H_{14}O_{7}  | 306.29           |
| 84-78-6    | Truflex OBP           | ![Truflex OBP](image) | C_{20}H_{30}O_{4}  | 334.45           |

Table 2: Continued.
factor subunit (JUN), and matrix metallopeptidase 9 (MMP9). It was believed that these targets may be significant for EH in the treatment of NS.

3.6. GO and KEGG Pathway Enrichment. To further explore the mechanism of EH on NS more systematically, the intersection target genes were inputted into DAVID database for GO analysis and KEGG pathway analysis ($P < 0.01$). It showed that the predicted target genes of EH were mainly enriched in 540 biological processes (BP), 64 cellular components (CC), and 93 molecular functions (MF). According to the number of target gens, the top 10 enriched conditions of GO analysis were listed, respectively, in Figure 6. It could be seen that green represented BP, orange represented CC, and purple represented MF. BP mainly included positive regulation of transcription from RNA polymerase II promoter, response to drug, negative regulation of apoptotic process, positive regulation of transcription, DNA-templated, positive regulation of cell proliferation, inflammatory response, apoptotic process, aging, positive regulation of gene expression, and signal transduction. CC mainly included cytosol, extracellular space, cytoplasm, nucleus, plasma membrane, extracellular exosome, extracellular region, nucleoplasm, mitochondrion, and membrane. MF mainly included protein binding, identical protein binding, enzyme binding, protein homodimerization activity, protein heterodimerization activity, transcription factor binding, zinc ion binding,
cytokine activity, protein kinase binding, transcription factor activity, and sequence-specific DNA binding.

The KEGG pathway enrichment was, respectively, completed via DAVID database and ClueGO plugin in Cytoscape. Through DAVID database, the KEGG pathway enrichment obtained a total of 114 pathways, of which the top 20 pathways were shown according to the size of the $P$ value in Figure 7. The $y$-axis represented the name of the pathway, and the $x$-axis represented the ratio of target genes to background genes. Besides, the size of dot represented the count of genes on different pathways, and the color of dot represented the significance of enrichment. Obviously, the data showed that EH mainly regulated pathways in cancer, hepatitis B, PI3K-Akt signaling pathway, HTLV-I infection, and TNF signaling pathway to treat NS. Through ClueGO plugin in Cytoscape, we obtained 29 representative pathways ($P < 0.01$) as shown in Figure 8. These pathways were divided into 11 categories based on signaling pathway

![Figure 4: Drug-ingredient-target-disease (D-I-T-D) network.](image)

![Figure 5: (a) PPI network diagram of the intersection targets of EH and NS. (b) 53 core targets arranged according to degree value.](image)
3.7. Molecular Docking of Active Ingredients with Key Targets. We selected AKT1 and TNF as the protein receptors because they had the highest degree value in the PPI network. Accordingly, the active ingredients including kaempferol, luteolin, quercetin, and naringenin were used as ligands for molecular docking verification. As shown in Figure 6 and Figure 7, the docking results indicated a good binding affinity between the active ingredients and the protein receptors, suggesting potential therapeutic effects against the diseases associated with these target genes.
Table 3: Molecular docking scores of major active ingredients and targets.

| Number | Ingredient | Target | Affinity value (kcal/mol) |
|--------|------------|--------|--------------------------|
| 1      | Kaempferol | AKT1   | -6.1                     |
| 2      | Luteolin   | AKT1   | -6.2                     |
| 3      | Quercetin  | AKT1   | -5.7                     |
| 4      | Naringenin | AKT1   | -6.2                     |
| 5      | Kaempferol | TNF    | -5.1                     |
| 6      | Luteolin   | TNF    | -5.3                     |
| 7      | Quercetin  | TNF    | -5.0                     |

Table 3, the affinities between these ingredients and targets were lower than -5.0 kcal/mol, which demonstrated that the main active ingredients of EH had a strong binding activity with targets. According to the docking results, the binding energy of kaempferol to AKT1 and TNF was -6.1 and -5.1 kcal/mol, the binding energy of luteolin to AKT1 and TNF was -6.2 and -5.3 kcal/mol, the binding energy of quercetin to AKT1 and TNF was -5.7 and -5.0 kcal/mol, and the binding energy of naringenin to AKT1 was -6.2 kcal/mol. Among them, luteolin and naringenin binding with AKT1 showed the best binding energy (-6.2 kcal/mol). Besides, four amino acid residues in AKT1, LYS-8, TRP-99, GLU-9, and HIS-89, formed multiple binding locations with luteolin (Figure 9(b)), while LEU-52 also bound to kaempferol through hydrogen bonding (Figure 9(a)). Two amino acid residues in AKT1, ARG-15 and GLU-85, formed multiple hydrogen bonds with quercetin (Figure 9(c)), and another two amino acid residues, GLN-47 and ALA-50, formed hydrogen bonds with naringenin (Figure 9(d)).

Three amino acid residues in TNF, HIS-52, ARG-60, and GLN-59, formed multiple binding locations with luteolin (Figure 9(f)), while HIS-52 and ARG-60 also bound to kaempferol and quercetin, respectively, through hydrogen bonding (Figures 9(e) and 9(g)).

4. Discussion

NS is a disease with considerable morbidity and high risk for relapse [31]. Despite developed medical conditions and increasingly diverse ways of reducing proteinuria, NS still cannot be effectively treated [32]. As a famous Chinese herbal medicine with a long history, EH has a wide range of pharmacological effects in the treatment of bronchial asthma, immune system diseases, cardiovascular related diseases, and cancer [33, 34]. Recently, EH is frequently used alone or in conjunction with other herbs to treat NS, but the relevant mechanism is not yet clear.

As is known, network pharmacology is a rapidly emerging discipline, which has the same thinking with TCM. In this study, we used network pharmacology theory and related tools to select the active ingredients and related targets of EH and linked them with NS. A total of 23 active ingredients with OB ≥ 30% and DL ≥ 0.18 were screened in the TCMSP database. From the “active ingredient-target” network diagram and “drug-ingredient-target-disease” network diagram, it could be seen that active ingredients including quercetin, kaempferol, luteolin, and naringenin were important nodes in the network. Besides, the network also intuitively displayed the specific relationship between EH and NS through 105 consensus genes.
Among them, quercetin is a natural flavonoid extracted from EH, other herbs, and numerous fruits and vegetables, which is known as a powerful herbal antioxidant. Pharmacological studies have shown that quercetin has several biological activities, such as antioxidant, anti-inflammatory, and anticarcinogenic effects [35]. More and more studies have proved that quercetin exerts renal protective effects through a variety of ways, including inhibiting ferroptosis, preventing

![Figure 9: Molecular docking of active ingredients with key targets: (a) kaempferol with AKT1, (b) luteolin with AKT1, (c) quercetin with AKT1, (d) naringenin with AKT1, (e) kaempferol with TNF, (f) luteolin with TNF, and (g) quercetin with TNF. The molecule was represented in a ball-stick model with atoms C and O in green and red, respectively.](image-url)
the aging of the kidney cells, alleviating renal fibrosis and apoptosis, and repressing oxidative stress and inflammation [36, 37]. As a natural anti-inflammatory compound, kaempferol has been reported to exert curative effects on kidney inflammatory damage [38]. Recently, numerous studies have shown that kaempferol has salutary effects for anti-inflammatory, antioxidant, analgesic, and anticancer properties [39]. With those properties, kaempferol has been demonstrated to exert beneficial therapeutic applications in nephrolithiasis, edema, and hyperlipidemia [40, 41]. Luteolin is a nontoxic flavonoid, which has become a hot area of research in an alternative medicine for several years [42]. Numerous studies have confirmed that luteolin has a wide range of biological activities, including antiapoptosis, antioxidative, anti-inflammatory, and anticancer properties [43]. It is generally accepted that luteolin exerts therapeutic effects on NS and other kidney injury [44, 45]. Therefore, it is speculated that these active ingredients may be important material basis of EH to treat NS.

The PPI network of the intersection targets of EH and NS clearly showed that AKT1, TNF, IL6, VEGFA, IL1B, TP53, MAPK3, CASP3, JUN, and MMP9 were the top 10 nodes with the highest degree value, which interacted with multiple active ingredients and played a key role in the network diagram. AKT family exists in three different isoforms, and they are named AKT1, AKT2, and AKT3, respectively. These unique isoforms are differentially expressed in many signaling pathways according to the developmental stage. Among them, AKT1 is involved in the proliferation, activation, apoptosis, and cell survival of interstitial fibroblasts and glomerular mesangial cells [46]. Kim et al. found that the deletion of AKT1 resulted in the attenuation of renal fibrosis and tubular dedifferentiation, which proved that AKT1 might serve as a therapeutic target in AKI-to-CKD progression [47]. Lin et al. reported that uncoupled mitochondrial respiration and increased oxidative stress were both found when mitochondria AKT1 was inhibited, which suggested that AKT1 signaling could be a novel target to develop new strategies for better treatment of various kidney diseases [48]. TNF is primarily produced by mononuclear macrophages and can prevent the body from the infringement by killing tumor cells. It is also proved to be an important cytokine with a wide range of biological effects, which is connected with the susceptibility and development of immune diseases, including NS [49].

GO analysis can be used to make a significant annotation of gene products. Among them, BP enrichment showed the coupling effect and transport mode of proteins in biological pathways. The results of CC analysis proved that intersection proteins participated in the cellular environment. Through MF analysis, we could demonstrate that some protein receptor activities were regulated by drugs, such as protein binding, which was closely related to proteinuria [50]. KEGG pathway enrichment analysis guides us to combine genomes with cellular and species to figure out the potential pathways of EH against NS. Our results showed that EH could be applied in the treatment of NS through PI3K-Akt signaling pathway, TNF signaling pathway, AGE-RAGE signaling pathway, hepatitis B, and pathways in cancer. In those pathways, multiple targets such as AKT1 and TNF were simultaneously regulated by active ingredients such as quercetin, kaempferol, and luteolin.

The PI3K-Akt signaling pathway has been proposed to modulate diverse biological processes and plays a key role in podocyte apoptosis, which is connected with the progress of proteinuria and renal function [51]. It has been reported that EH exerts anti-inflammatory and antioxidative stress effects through promoting the phosphorylation of PI3K and AKT proteins [52]. As the main component of EH, quercetin has been proved to improve renal fibrosis and apoptosis in chronic renal failure rats, as well as reducing water retention and toxin accumulation by inhibiting the PI3k/Akt pathway through targeting PIK3R [53]. Thus, regulating the PI3K/AKT pathway in podocytes through EH may be an important potential targeted therapy for NS in the future [54]. As one of important inflammatory pathways, the TNF signaling pathway plays a central role in the regulation of different cellular events such as immunity, apoptosis, cell proliferation, and differentiation [55]. It has been proved that the TNF signaling pathway is connected with cholesterol-dependent podocyte apoptosis and albuminuria, which may be primarily activated immediately in the process of various NS progression, including diabetic nephropathy and focal segmental glomerulosclerosis [56, 57]. It has been demonstrated that EH exhibits antioxidant, anti-inflammatory, and antiarthritic properties by decreasing proinflammatory cytokines, including inhibiting the TNF signaling pathway [58]. Besides, EH has been proved to exert anti-inflammatory potential via downregulation of TNF-α [59]. As one of the active ingredients extracted from EH, kaempferol can reduce the intensity of inflammatory processes by inhibiting the secretion of proinflammatory cytokine TNF-α and additionally increasing the expression of anti-inflammatory cytokine IL-10 [60]. Furthermore, the protective effect of kaempferol against streptozotocin-induced diabetic nephropathy has been proved to be associated with the downregulation of TNF-α and IL-6 [61]. The AGE-RAGE signaling pathway is connected with the regulation of oxidative stress and inflammation. When it is activated, nicotinamide adenine dinucleotide phosphate oxidase-induced reactive oxygen is produced, which will give rise to oxidative stress [62]. Therefore, targeting at the AGE-RAGE signaling pathway could be the promising potential approach for improving NS and ageing kidney.

The results of molecular docking were consistent with the results of network pharmacology because active ingredients such as kaempferol, luteolin, quercetin, and naringenin all had good binding to the target protein AKT1 or TNF, which provided a strong basis for the results of network pharmacology. So, the feasibility of TCM such as EH can be affirmed in the treatment of NS.

5. Conclusions

On the basis of network pharmacology and molecular docking, this study focused on the active ingredients of EH and its multitarget and multipathway characteristics in the process of treating NS. The results indicated that the potential
mechanism of EH in treating NS may be related to PI3K-Akt signaling pathway, TNF signaling pathway, and AGE-RAGE signaling pathway. It may provide new ideas and better approaches for the treatment of NS. However, further in vivo and in vitro experimental verifications are expected to be conducted in the future.

Data Availability
The data used to support the findings of this study are included within the article.

Conflicts of Interest
The authors declare that they have no conflicts of interest.

Authors’ Contributions
Tianwen Yao designed the experiment and drafted the manuscript. Yi Wang contributed to the study design and gave the theoretical guidance. Qingliang Wang and Shisheng Han contributed to statistical analyses. Yan Lu helped to draft the manuscript. Yanqiu Xu was responsible for the revision of the final version. All authors read and approved the final manuscript.

Acknowledgments
This work was supported by the Shanghai Sailing Program (21YF1448300) and Project of Inheritance and Scientific Innovation of Traditional Chinese Medicine of Shanghai (ZYCC2019005).

References
[1] K. Tsuji, S. Kitamura, and J. Wada, “MicroRNAs as biomarkers for nephrotic syndrome,” International Journal of Molecular Sciences, vol. 22, no. 1, p. 88, 2021.
[2] S. Veissi, B. Smeets, L. P. Heuvel, M. F. Schreuder, and J. Jansen, “Nephrotic syndrome in a dish: recent developments in modeling in vitro,” Pediatric Nephrology, vol. 35, no. 8, pp. 1363–1372, 2020.
[3] K. L. Gibson, P. Hansrivijit, and M. E. Farris, “Emerging agents for the management of nephrotic syndrome: progress to date,” Paediatric Drugs, vol. 18, no. 1, pp. 25–29, 2016.
[4] S. Agrawal, J. J. Zaritsky, A. Fornoni, and W. E. Smoyer, “Dyslipidaemia in nephrotic syndrome: mechanisms and treatment,” Nature Reviews. Nephrology, vol. 14, no. 1, pp. 57–70, 2018.
[5] H. Tamura, “Trends in pediatric nephrotic syndrome,” World Journal of Nephrology, vol. 10, no. 5, pp. 88–100, 2021.
[6] H. Fujisawa, Y. Nakayama, S. Nakao et al., “Effectiveness of immunosuppressive therapy for nephrotic syndrome in a patient with late-onset Fabry disease: a case report and literature review,” BMC Nephrology, vol. 20, no. 1, 2019.
[7] S. E. D. A. Hackl, P. Zed, J. Diefenhardt, R. E. Binz-Lotter, and L. T. Weber, “The role of the immune system in idiopathic nephrotic syndrome,” Molecular and Cellular Pediatrics, vol. 8, no. 1, 2021.
[8] M. L. Downie, C. Gallibois, R. S. Parekh, and D. G. Noone, “Nephrotic syndrome in infants and children: pathophysiology and management,” Paediatrics and International Child Health, vol. 37, no. 4, pp. 248–258, 2017.
[9] H. Odaguchi, S. Hyuga, M. Sekine et al., “The adverse effects of Ephedra herb and the safety of ephedrine alkaloids-free Ephedra herb extract (EEF),” Yakugaku Zasshi, vol. 139, no. 11, pp. 1417–1425, 2019.
[10] S. M. Miao, Q. Zhang, X. B. Bi, J. L. Cai, and M. L. Wang, “A review of the phytochemistry and pharmacological activities of Ephedra herb,” Chinese Journal of Natural Medicines, vol. 18, no. 5, pp. 321–344, 2020.
[11] S. Ibragic, S. Barbini, J. T. Oberlerchner, A. Potthast, T. Rosenau, and S. Böhmder, “Antioxidant properties and qualitative analysis of phenolic constituents in Ephedra spp. by HPTLC together with injection port derivatization GC-MS,” Journal of Chromatography B, vol. 1180, article 1112277, 2021.
[12] Y. Nagata, H. Ando, Y. Sasaki, and R. Suzuki, “Ephedra herb, Mao, inhibits antigen-induced mast cell degranulation by induction of the affinity receptor for IgE internalization,” Pharmaceutical Research, vol. 38, no. 4, pp. 569–581, 2021.
[13] S. Nakamori, J. Takahashi, S. Hyuga et al., “Analogic effects of Ephedra herb extract, ephedrine alkaloids-free Ephedra herb extract, ephedrine, and pseudoephedrine on formalin-induced pain,” Biological & Pharmaceutical Bulletin, vol. 42, no. 9, pp. 1538–1544, 2019.
[14] N. Oshima, “Efficient preparation of ephedrine alkaloids-free Ephedra herb extract and its antitumor effect and putative marker compound,” Yakugaku Zasshi, vol. 137, no. 2, pp. 173–177, 2017.
[15] N. Oshima, T. Yamashita, S. Hyuga et al., “Efficiently prepared ephedrine alkaloids-free Ephedra herb extract: a putative marker and antiproliferative effects,” Journal of Natural Medicines, vol. 70, no. 3, pp. 554–562, 2016.
[16] E. Z. Mpingirika, A. E. Hosseiny, S. M. S. Bakheit, R. Arafah, and A. Amleh, “Potential anticancer activity of crude ethanol, ethyl acetate, and water extracts of Ephedra foeminea on human osteosarcoma U2OS cell viability and migration,” BioMed Research International, vol. 2020, Article ID 3837693, 16 pages, 2020.
[17] Y. Ming, S. Cheng, W. Long et al., “The herbal formula granule prescription Mahuang decoction ameliorated chronic kidney disease which was associated with restoration of dysbiosis of intestinal microbiota in rats,” Evidence-Based Complementary and Alternative Medicine, vol. 2021, Article ID 4602612, 12 pages, 2021.
[18] Z. Dong, H. Dai, Y. Gao et al., “Effect of Mahuang Fuzi and Shenzhuo decoction on idiopathic membranous nephropathy: a multicenter, nonrandomized, single-arm clinical trial,” Frontiers in Pharmacology, vol. 12, 2021.
[19] H. Lu, J. Xu, B. Xie et al., “The multi-target mechanism of cyclosporin A in the treatment of vitiligo based on network pharmacology,” Dermatologic Therapy, vol. 34, no. 4, article e15023, 2021.
[20] L. Jiang, Z. Shi, and Y. Yang, “Network pharmacology-based approach to investigate the molecular targets of Rhubarb for treating cancer,” Evidence-Based Complementary and Alternative Medicine, vol. 2021, Article ID 9945633, 8 pages, 2021.
[21] G. Zhou, X. Feng, and A. Tao, “Explore the lipid-lowering and weight-reducing mechanism of lotus leaf based on network pharmacology and molecular docking,” Evidence-Based Complementary and Alternative Medicine, vol. 2021, Article ID 1464027, 7 pages, 2021.
[22] S. Abdelsattar, A. Dawoud, and M. A. Helal, “Interaction of nanoparticles with biological macromolecules: a review of molecular docking studies,” Nanotoxicology, vol. 15, no. 1, pp. 66–95, 2021.

[23] F. Stanzione, I. Giangreco, and J. C. Cole, “Use of molecular docking computational tools in drug discovery,” Progress in Medicinal Chemistry, vol. 60, pp. 273–343, 2021.

[24] J. Lu, J. Yan, J. Yan et al., “Network pharmacology based research into the effect and mechanism of Xijiao Dihuang decoction against sepsis,” Biomedicine & Pharmacotherapy, vol. 122, article 109777, 2020.

[25] G. Feipeng, X. Luxin, C. Beili et al., “Exploration of Ziziphi Spinosae semen in treating insomnia based on network pharmacology strategy,” Evidence-Based Complementary and Alternative Medicine, vol. 2021, Article ID 19888607, 12 pages, 2021.

[26] C. Jia, X. Pan, B. Wang, P. Wang, Y. Wang, and R. Chen, “Mechanism prediction of Astragalus membranaceus against cisplatin-induced kidney damage by network pharmacology and molecular docking,” Evidence-Based Complementary and Alternative Medicine, vol. 2021, Article ID 9516726, 15 pages, 2021.

[27] J. X. Ma, M. Ye, K. Ma et al., “Network pharmacology-based strategy for predicting active ingredients and potential targets of Coptis chinensis Franchin polycystic ovary syndrome,” Evidence-Based Complementary and Alternative Medicine, vol. 2021, Article ID 6651307, 15 pages, 2021.

[28] Z. Zeng, J. Hu, J. Jiang et al., “Network pharmacology and molecular docking-based prediction of the mechanism of Qianghuo Shengshi decoction against rheumatoid arthritis,” BioMed Research International, vol. 2021, Article ID 6623912, 12 pages, 2021.

[29] K. T. B. Crosara, E. B. Moda, Y. Xiao, and W. L. Siqueira, “Merging in-silico and in vitro salivary protein complex partners using the STRING database: a tutorial,” Journal of Proteomics, vol. 171, pp. 87–94, 2018.

[30] T. Han, Y. Zhou, and D. Li, “Relationship between hepatocellular carcinoma and depression via online database analysis,” Bioengineered, vol. 12, no. 1, pp. 1689–1697, 2021.

[31] F. Veltkamp, L. R. Rensma, A. H. M. Bouts, and Learns consortium, “Incidence and relapse of idiopathic nephrotic syndrome: meta-analysis,” Pediatrics, vol. 148, no. 1, 2021.

[32] C. Xue, B. Yang, J. Xu et al., “Efficacy and safety of rituximab in adult frequent-relapsing or steroid-dependent minimal change disease or focal segmental glomerulosclerosis: a systematic review and meta-analysis,” Clinical Kidney Journal, vol. 14, no. 4, pp. 1042–1054, 2021.

[33] F. Sioud, S. Amor, I. B. Tounia et al., “A new highlight of Ephedra alata deca properties as potential adjuvant in combination with cisplatin to induce cell death of 4T1 breast cancer cells in vitro and in vivo,” Cell, vol. 9, no. 2, p. 362, 2020.

[34] N. Oshima, T. Yamashita, N. Uchiyama et al., “Non-alkaloidal composition of Ephedra herb is influenced by differences in habitats,” Journal of Natural Medicines, vol. 73, no. 1, pp. 303–311, 2019.

[35] K. T. Huang, C. T. Wu, Y. Chang, F. M. Ho, C. K. Chiang, and S. H. Liu, “Therapeutic effect of quercetin polymeric nanoparticles on ischemia/reperfusion-induced acute kidney injury in mice,” Biochemical and Biophysical Research Communications, vol. 608, pp. 122–127, 2022.

[36] Y. Wang, F. Quan, Q. Cao et al., “Quercetin alleviates acute kidney injury by inhibiting ferropotosis,” Journal of Advanced Research, vol. 28, pp. 231–243, 2021.

[37] F. Abharzanjani and M. Hemmati, “Protective effects of quercetin and resveratrol on aging markers in kidney under high glucose condition: in vivo and in vitro analysis,” Molecular Biology Reports, vol. 48, no. 7, pp. 5435–5442, 2021.

[38] T. M. Yeh, C. D. Chang, S. S. Liu, C. I. Chang, and W. L. Shih, “Tea seed kaempferol triglycoside attenuates LPS-induced systemic inflammation and ameliorates cognitive impairments in a mouse model,” Molecules, vol. 27, no. 7, p. 2055, 2022.

[39] S. Chang, X. Li, Y. Zheng et al., “Kaempferol exerts a neuroprotective effect to reduce neuropathic pain through TLR4/NF-κB signaling pathway,” Phytotherapy Research, vol. 36, no. 4, pp. 1678–1691, 2022.

[40] P. Yuan, X. Sun, X. Liu et al., “Kaempferol alleviates calcium oxalate crystal-induced renal injury and crystal deposition via regulation of the AR/NOX2 signaling pathway,” Phytomedicine, vol. 86, article 153555, 2021.

[41] A. Ochiai, M. B. Othman, and K. Sakamoto, “Kaempferol ameliorates symptoms of metabolic syndrome by improving blood lipid profile and glucose tolerance,” Bioscience, Biotechnology, and Biochemistry, vol. 85, no. 10, pp. 2169–2176, 2021.

[42] R. Reudhabibadh, T. Binlathe, P. Chinthapothumikuntal et al., “Suppressing Cdk5 activity by luteolin inhibits MMP-13-induced apoptotic neuroblasta through Erk1/2 and Fak/Akt/GSK3β pathways,” Molecules, vol. 26, no. 5, p. 1307, 2021.

[43] Y. J. Lee, T. Lim, M. S. Han et al., “Anticancer effect of luteolin is mediated by downregulation of TAM receptor tyrosine kinases, but not interleukin-8, in non-small cell lung cancer cells,” Oncology Reports, vol. 37, no. 2, pp. 1219–1226, 2017.

[44] L. Tang, B. Deng, L. Shi, B. Wei, B. Ren, and X. Fu, “Effect of luteolin on 11beta-hydroxysteroid dehydrogenase in rat liver and kidney,” Evidence-Based Complementary and Alternative Medicine, vol. 2015, Article ID 834124, 7 pages, 2015.

[45] X. Xu, Z. Yu, B. Han et al., “Luteolin alleviates inorganic mercury-induced kidney injury via activation of the AMPK/mTOR autophagy pathway,” Journal of Inorganic Biochemistry, vol. 224, article 111583, 2021.

[46] M. Palma, C. Leroy, S. Salomé-Desnoulez et al., “A role for AKT1 in nonsense-mediated mRNA decay,” Nucleic Acids Research, vol. 49, no. 19, pp. 11022–11037, 2021.

[47] Y. Kim, Y. K. Park, S. H. Song et al., “Role of Akt1 in renal fibrosis and tubular dedifferentiation during the progression of acute kidney injury to chronic kidney disease,” The Korean Journal of Internal Medicine, vol. 36, no. 4, pp. 962–974, 2021.

[48] H. Y. Lin, Y. Chen, Y. Chen et al., “Tubular mitochondrial AKT1 is activated during ischemia reperfusion injury and has a critical role in predisposition to chronic kidney disease,” Kidney International, vol. 99, no. 4, pp. 870–884, 2021.

[49] M. Xiao, S. Bai, J. Chen et al., “Correlation of TNF-α and IL-10 gene polymorphisms with primary nephrotic syndrome,” Experimental and Therapeutic Medicine, vol. 20, no. 5, 2020.

[50] C. Zheng, X. Hu, L. Zhao, M. Hu, and F. Gao, “Clinical and pharmacological hallmarks of rifapentine&rsquo; use in diabetes patients with active and latent tuberculosis: do we know enough?,” Drug Design, Development and Therapy, vol. 11, pp. 2957–2968, 2017.

[51] C. Zhao, Y. Gu, L. Chen, and X. Su, “Upregulation of FoxO3a expression through PI3K/Akt pathway attenuates the
progression of lupus nephritis in MRL/lpr mice,” *International Immunopharmacology*, vol. 89, no. Part A, article 107027, 2020.

[52] L. Huang, B. Zhao, Q. Li, J. Wu, H. Jiang, and Q. Li, “Ephedrine alleviates middle cerebral artery occlusion-induced neurological deficits and hippocampal neuronal damage in rats by activating PI3K/AKT signaling pathway,” *Bioengineering*, vol. 12, no. 1, pp. 4136–4149, 2021.

[53] H. Tu, D. Ma, Y. Luo et al., “Quercetin alleviates chronic renal failure by targeting the PI3k/Akt pathway,” *Bioengineering*, vol. 12, no. 1, pp. 6538–6558, 2021.

[54] F.-q. Cui, Y.-F. Wang, Y.-b. Gao et al., “Effects of BSF on podocyte apoptosis via regulating the ROS-mediated PI3K/AKT pathway in DN,” *Journal of Diabetes Research*, vol. 2019, Article ID 9512406, 10 pages, 2019.

[55] N. Genov, A. Basti, M. Abreu, and A. Relógio, “Temporal splicing switches in elements of the TNF-pathway identified by computational analysis of transcriptome data for human cell lines,” *International Journal of Molecular Sciences*, vol. 20, no. 5, p. 1182, 2019.

[56] G. Wang, J. Ouyang, S. Li et al., “The analysis of risk factors for diabetic nephropathy progression and the construction of a prognostic database for chronic kidney diseases,” *Journal of Translational Medicine*, vol. 17, no. 1, 2019.

[57] L. Otalora, E. Chavez, D. Watford et al., “Identification of glomerular and podocyte-specific genes and pathways activated by sera of patients with focal segmental glomerulosclerosis,” *PLoS One*, vol. 14, no. 10, article e0222948, 2019.

[58] H. Ahsan, H. M. Irfan, M. H. Asim et al., “Therapeutic appraisal of ephedrine against rheumatoid arthritis: a new indication,” *Pakistan Journal of Pharmaceutical Sciences*, vol. 34, no. 4, pp. 1549–1554, 2021.

[59] H. S. Yaseen, M. Asif, M. Saadullah et al., “Methanolic extract of Ephedra ciliata promotes wound healing and arrests inflammatory cascade in vivo through downregulation of TNF-α,” *Inflammmopharmacology*, vol. 28, no. 6, pp. 1691–1704, 2020.

[60] M. Palacz-Wrobel, P. Borkowska, M. Paul-Samojedny et al., “Effect of apigenin, kaempferol and resveratrol on the gene expression and protein secretion of tumor necrosis factor alpha (TNF-α) and interleukin-10 (IL-10) in RAW-264.7 macrophages,” *Biomedicine & Pharmacotherapy*, vol. 93, pp. 1205–1212, 2017.

[61] S. Alshehri, “Kaempferol attenuates diabetic nephropathy in streptozotocin-induced diabetic rats by a hypoglycaemic effect and concomitant activation of the Nrf-2/Ho-1/antioxidants axis,” *Archives of Physiology and Biochemistry*, vol. 24, pp. 1–14, 2021.

[62] X. Q. Wu, D. D. Zhang, Y. N. Wang, Y. Q. Tan, X. Y. Yu, and Y. Y. Zhao, “AGE/RAGE in diabetic kidney disease and ageing kidney,” *Free Radical Biology & Medicine*, vol. 171, no. 8, pp. 260–271, 2021.