



Study on the mechanism of the Modified Ginseng-Schisandra Decoction (MGSD) in the treatment of recurrent respiratory tract infection (RRTI) based on network pharmacology

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Background: The aim of this study was to investigate the mechanism of Modified Ginseng-Schisandra Decoction in the treatment of recurrent respiratory tract infection (RRTI) using network pharmacology.

Methods: To screen the active ingredients of A Modified Ginseng-Schisandra Decoction, TCMSP, TCMID, Batman-TCM and PubChem database were applied. To predict the targets of active ingredients on RRTI, TCMSP, Pubmed, OMIM, Drug Bank, GAD and TTD database were used. The compounds-therapeutic target network was constructed with Cytoscape 3.7.2 software. The STRING database was used to construct a protein-protein interaction (PPI) network, and Kyoto Encyclopedia of Genes and Genomes (KEGG) pathway enrichment analysis was used to identify potential signal pathways.

Results: The 3 main active ingredients of Modified Ginseng-Schisandra Decoction obtained by screening were quercetin, kaempferol, and isoflavone; the main therapeutic targets were *PTGS2*, *ESR1*, *AR*, *PPARG*, *NOS2*, and others. Based on the PPI network, we found that the targets of Modified Ginseng-Schisandra Decoction were significantly enriched in (FDR <0.01) cancer pathway, tumor necrosis factor (TNF) signaling pathway, hypoxia-inducible factor (HIF-1) signaling pathway, and others.

Conclusions: Modified Ginseng-Schisandra Decoction can treat RRTI primarily through acting in the signal transduction of some key nodes of cancer pathway and TNF pathway. It exerts a direct or indirect influence on multiple signaling pathways, and has the characteristics of multicomponent, multitarget, and multichannel action.

Keywords: Modified Ginseng-Schisandra Decoction; recurrent respiratory tract infection (RRTI); network pharmacology; signal pathway; molecular mechanisms

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Introduction

Recurrent respiratory tract infection (RRTI) is a common and frequently occurring disease in children. It refers to the frequent occurrence of upper and lower respiratory tract infections within 1 year that exceed the normal range (1,2). According to the location of infection, RRTI can be divided into recurrent upper respiratory tract infection and recurrent lower respiratory tract infection (bronchitis and pneumonia) (1). According to statistics, the incidence of RRTI among children under the age of 1 in developed countries is as high as 25%, while the incidence in children between 1 and 4 years of age is 18% (3). In developing countries, the annual number of child deaths from RRTI is as high as 2 million (mainly pneumonia) (4). Relevant investigations show that this disease accounts for about 10–20% of respiratory diseases in China, with 70–80% of these patients being under 5 years old (5). As conditions in the environment change, the incidence of RRTI is increasing (6). Compared with lower respiratory tract infection, repeated upper respiratory tract infection is more common (7). RRTI is characterized by a long duration and a high likelihood of multiple recurrences. Children with severe disease can have several episodes per month. If the disease is not treated in timely and effective fashion, it can develop into chronic disease, with asthma, nephritis, myocarditis being common complications (8) that pose a considerable threat to the life and health of children (7). Traditional Chinese medicine (TCM) has attracted much attention in the prevention and treatment of RRTI in recent years.

RRTI can be classified as “body deficiency cold”, “cough and asthma”, “long cough”, “deficiency syndrome” and “spontaneous sweating” in the discourse of traditional Chinese medicine (9). The key pathogenesis lies in “deficiency” (10), and clinical treatment is based on stage treatment and syndrome differentiation.

Modified Ginseng-Schisandra Decoction is often used in the treatment of RRTI (“lung and spleen deficiency syndrome”), Ginseng-Schisandra decoction comes from a book named “Youyou Jicheng” written by Fuzheng Chen during the Qing dynasty. The original record is as follows: “The cough lasts for a long time, and there is no other evidence. It is due to lung deficiency. Only the spleen should be supplemented. Modified Ginseng-Schisandra Decoction. It’s also a magical prescription to treat long-term cough and deficiency of qi and weak qi, white face, and white lips.” The original prescription consists of ginseng, *Atractylodes macrocephala* Koidz., *Poria cocos*, *Schisandra*, *Ophiopogon*

japonicus, *Radix Glycyrrhizae* Preparete, ginger, and jujube. Later doctors often used this prescription to treat cough due to deficiency of lung and spleen qi. Based on clinical experience, we use Modified Ginseng-Schisandra Decoction to treat spleen deficiency syndrome after RRTI. The basic prescription is composed of *Radix astragali* (6 g), *Codonopsis pilosula* (6 g), *Atractylodes macrocephala* Koidz. (6 g), *Poria cocos* (9 g), *Schisandra* (3 g), *Ophiopogon japonicus* (6 g), oyster (15 g), *Radix glycyrrhizae* Preparete (3 g), and *Saposhnikovia divaricata* (10 g), with the ingredients being added or subtracted based on clinical symptoms. As for Children with RRTI, Modified Ginseng-Schisandra Decoction is used according to the following rules: 1 to 2 years old: 2 times/day, 1/3 doses/time; 3 to 6 years old; 2 times/day, 0.5 doses/time; 6 years old and above, 2 times/day, 1 dose/time. The treatment course lasted 4 weeks. Preliminary clinical studies have found that the Modified Ginseng-Schisandra Decoction can reduce TCM syndrome scores, and promote white blood cell (WBC), hemoglobin (Hb), C-reactive protein (CRP), and eosinophil (EOS) and other indicators to return to normal levels, with the clinical efficacy being superior to that of single-use *Bacillus subtilis* dual-live bacteria particles for treatment(11,12).

TCM prescriptions typically have multiple components, multiple targets, and multiple pathways. This study used network pharmacology to examine the potential pharmacological effects and molecular mechanisms of Modified Ginseng-Schisandra Decoction on the treatment of RRTI, so as to lay a theoretical foundation for its treatment of RRTI. We present the following article in accordance with the MDAR reporting checklist (available at <https://dx.doi.org/10.21037/tp-21-240>).

Methods

Research design

The general experimental design for this pharmacology study of Modified Ginseng-Schisandra Decoction in the treatment of RRTI included the following principal steps: the screening of active ingredients of Modified Ginseng-Schisandra Decoction, the establishment of a target database of active ingredients related to RRTI, and the establishment of the Modified Ginseng-Schisandra Decoction-Active compounds-therapeutic target network with subsequent KEGG enrichment analysis. The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013).

Database establishment

Screening the main active ingredients of the Modified Ginseng-Schisandra Decoction

We searched the Traditional Chinese Medicine Systems Pharmacology Database and Analysis Platform (TCMSP), Traditional Chinese Medicines Integrated Database (TCMID), and Batman-TCM (<http://bionet.ncpsb.org/batman-tcm/>) databases to find the main chemical components of *Radix astragali*, *Codonopsis pilosula*, *Atractylodes macrocephala* Koidz., *Poria cocos*, *Schisandra*, *Ophiopogon japonicus*, oyster, *Radix glycyrrhizae* Preparete, and *Saposhnikovia divaricata*, and used the PubChem database (<https://pubchem.ncbi.nlm.nih.gov/>) to check their compound names and molecular structures. With oral bioavailability (OB) $\geq 30\%$ and drug-likeness (DL) ≥ 0.18 as screening conditions, the compounds were screened and the active compound targets were obtained.

Prediction of targets of active ingredients in Modified Ginseng-Schisandra Decoction

The TCMSP database revealed the targets corresponding to the active ingredients of the Modified Ginseng-Schisandra Decoction. The targets from different sources were entered into UniProt (<http://www.uniprot.org/>), PubMed, and other databases with the search limited to the human species. In this way, the gene symbols corresponding to the targets were acquired for subsequent network pharmacological analysis.

Target mining of RRTIs

The keyword phrase “Recurrent respiratory tract infection” was used to search the Online Mendelian Inheritance in Man (OMIM) database (<https://www.omim.org/>), Drug Bank (<http://www.drugbank.ca/>), Gene Association Database (GAD; <http://geneticassociation-db.nih.gov/>), and Therapeutic Targets Database (TTD <http://db.idrblab.net/ttd/>) for all RRTI targets.

Construction and visualization of Modified Ginseng-Schisandra Decoction network diagram

Construction and visualization of Modified Ginseng-Schisandra decoction-Active compounds-therapeutic target

Drug-disease targets (DDTs) were obtained from identifying the intersection of Modified Ginseng-Schisandra Decoction target and the RRTI target with Venn (<http://bioinformatics.psb.ugent.be/webtools/Venn/>). The Modified

Ginseng-Schisandra Decoction-Active compounds-therapeutic target network was constructed with Cytoscape 3.7.2 software, and topology analysis and visualization were performed. The Network Analyzer tool in the software was used to calculate the degree value to evaluate the importance of the components and targets in the network.

Protein-protein interaction (PPI) network data

The Modified Ginseng-Schisandra decoction PPI network was constructed using the Search Tool for the Retrieval of Interacting Genes/Proteins (STRING) database (<https://string-db.org/>). The conditions were set to “minimum required interaction score =0.9” and “hide disconnected nodes in the network”.

Kyoto Encyclopedia of Genes and Genomes enrichment analysis

To further elucidate the different scientific connotations of Modified Ginseng-Schisandra decoction from a biological perspective, we used the KOBAS online server (<http://kobas.cbi.pku.edu.cn/kobas3/?T=1>) with the species set to human to conduct Kyoto Encyclopedia of Genes and Genomes (KEGG) enrichment analysis and Gene Ontology (GO) function enrichment analysis for the DDTs of the Modified Ginseng-Schisandra Decoction. This was followed by visualization using the KEGG Mapper tool (<https://www.genome.jp/kegg/mapper.html>) to analyze the specific role of related targets in important signaling pathways and to obtain a schematic diagram of Modified Ginseng-Schisandra Decoction regulatory signaling pathways (13).

Statistical analysis

The relative data were recorded with EXCEL. In KEGG pathway enrichment analysis, the pathways with $P < 0.05$ were considered statistical different and selected.

Results

Identifying the common targets of the Modified Ginseng-Schisandra Decoction and RRTI

A total of 156 active compounds of the Modified Ginseng-Schisandra Decoction (excluding duplicate compounds) were found by searching the three databases. Among them, 15, 16, 4, 6, 7, 3, 14, 88, and 16 corresponding compounds of *Radix astragali*, *Codonopsis pilosula*, *Atractylodes macrocephala*

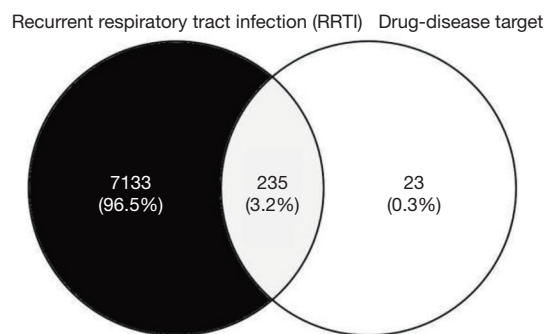


Figure 1 Drug-disease target Venn diagram of Modified Ginseng-Schisandra Decoction.

Koidz., *Poria cocos*, *Schisandra*, *Ophiopogon japonicus*, oyster, *Radix glycyrrhizae* Preparete, and *Saposhnikovia divaricata* were found, respectively.

After Uniprot corrected all target names, the number of potential targets of Modified Ginseng-Schisandra Decoction was 258. Using the GeneCards database, we searched and downloaded all 7,168 target genes related to RRTI. The integration of the potential targets of Modified Ginseng-Schisandra Decoction active chemical components with the targets of RRTI, resulted in 235 DDTs of the Modified Ginseng-Schisandra Decoction (Figure 1).

Analysis and visualization of the DDT network diagram based on the Modified Ginseng-Schisandra decoction

Cytoscape software was used to construct the DDT network graph. It was composed of 415 nodes and 2,130 edges. In Figure 2, the green represents the prescription drugs of Modified Ginseng-Schisandra Decoction [9], the red represents the active compounds of the Modified Ginseng-Schisandra Decoction [156], and the blue represents the DDT of Modified Ginseng-Schisandra Decoction [235].

The top 10 therapeutic ingredients with degree value were quercetin, kaempferol, isoflavones (among others) (degrees of 268, 102, and 62, respectively; see Table 1 for details). The top 10 treatment targets by degree were *PTGS2*, *ESR1*, *AR*, *PPARG*, *NOS2* (among others) (degrees of 108, 93, 88, 81, and 74, respectively; see Table 2 for details).

Analysis of the DDT PPI network of the modified Ginseng-Schisandra Decoction

A total of 235 DDTs of the Modified Ginseng-Schisandra

Decoction were imported into the STRING database (<https://string-db.org/>). With the filter condition set to “Homo sapiens” and the conditions set to “minimum required interaction score =0.9” and “hide disconnected nodes” in the network”, 216 target proteins were obtained. According to the degree value of the network topology index, a bar graph of the top 30 related protein nodes was drawn (Figure 3).

Enrichment analysis of the KEGG signal pathway of the core PPI

KEGG pathway analysis was performed on 216 target proteins in the PPI network with the condition of a false discovery rate (FDR) set as <0.01, and the top 30 signal pathways were visualized according to the FDR value. The specific results are shown in Figure 4. According to the correlation between the size of FDR, the size of enrichment factors, and RRTI, it was found that that important signal pathways included cancer signal pathway, tumor necrosis factor (TNF) signal pathway, hypoxia-inducible factor (HIF-1) signal pathway, among others. The KEGG Mapper tool was used to analyze the specific role of the Modified Ginseng-Schisandra Decoction in the cancer pathway, TNF signaling pathway, and HIF-1 signaling pathway, and the model diagrams of Modified Ginseng-Schisandra Decoction regulating the above signal pathways were constructed (Figures 5,6).

Discussion

The Modified Ginseng-Schisandra Decoction is composed of *Radix astragali*, *Codonopsis pilosula*, *Atractylodes macrocephala* Koidz., *Poria cocos*, *Schisandra*, *Ophiopogon japonicus*, oyster, *Radix glycyrrhizae* Preparete, *Saposhnikovia divaricata*, and is derived from the combination of Chinese classical prescriptions including the Ginseng-Schisandra Decoction, Yupingfeng Powder, Sijunzi Decoction, and Shengmai Yin. Chinese medicine believes that children are pure Yang bodies, and if ginseng is not used properly, it is easy to develop Yang heat and “excessive internal heat” (14). *Codonopsis pilosula* is sweet and flat, and is more suitable for children. Therefore, the original ginseng is exchanged for *Codonopsis*. *Codonopsis pilosula* and *Radix astragali* are principal drugs, which correspond to the spleen and lung meridian. *Codonopsis pilosula* has the effects of tonifying the middle-Jiao and Qi, invigorating the spleen and lungs, invigorating the blood and activating fluid, and eliminating dysphoria.

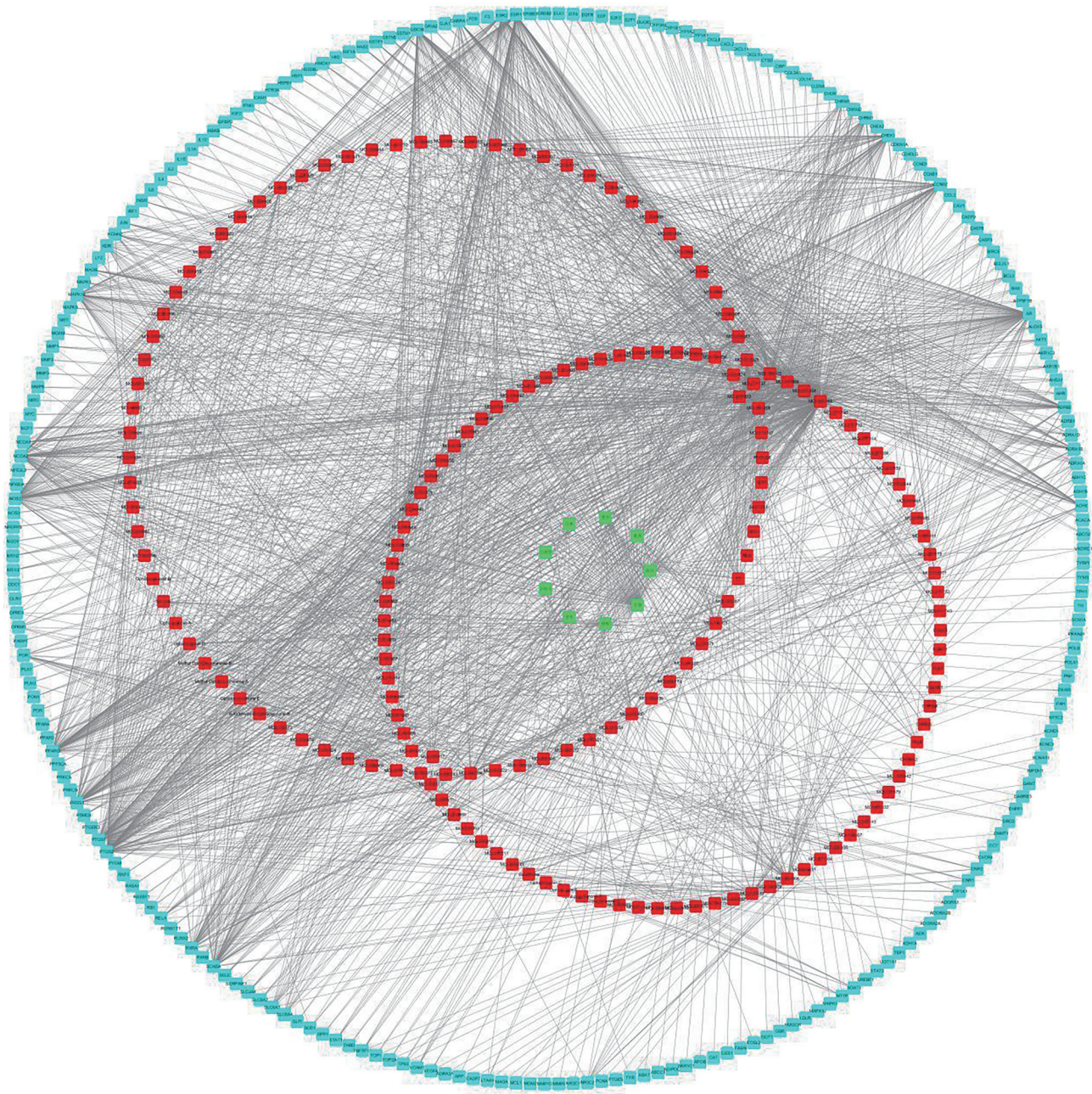


Figure 2 Drug-disease target network diagram of the Modified Ginseng-Schisandra Decoction.

It can enhance immune function, regulate the intestinal tract, reduce the absorption of inflammatory mediators and toxins, and has a good blood-replenishing effect (15). *Radix astragali* is good at entering the spleen and stomach. It is an important medicine for tonifying the middle-Jiao and Qi. It can tonify the lung and spleen, replenish qi for strengthening superficies. *Atractylodes macrocephala* Koidz

combined with *Poria cocos* can strengthen the spleen, dry dampness, and replenish qi; *Schisandra* with *Ophiopogon japonicus* has the functions of nourishing Yin and moistening lung, benefiting stomach and promoting body fluid, clearing away the heart fire and eliminating irritability; *Saposhnikovia divaricata* has the functions of dispelling wind, relieving the superficies, and eliminating dampness. According to

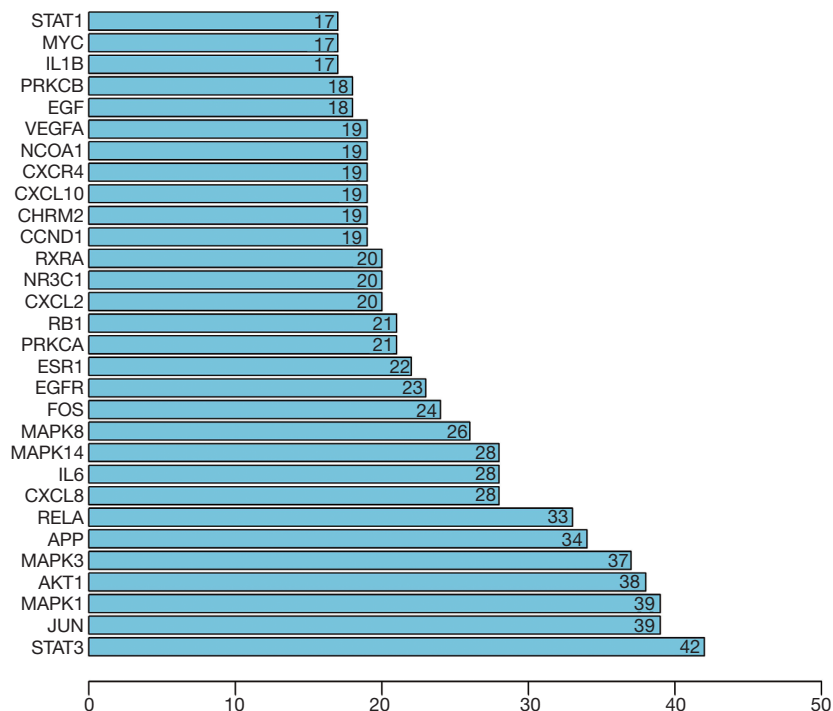
Table 1 Information of the top 10 active ingredients by degree value of Modified Ginseng-Schisandra Decoction

Mol ID	Molecule name	OB (%)	DL
MOL000442	1,7-Dihydroxy-3,9-dimethoxy pterocarpene	39.05	0.48
MOL000098	Quercetin	46.43	0.28
MOL003896	7-Methoxy-2-methyl isoflavone	42.56	0.2
MOL000392	Formononetin	69.67	0.21
MOL000354	Isorhamnetin	49.6	0.31
MOL000006	Luteolin	36.16	0.25
MOL000296	Hederagenin	36.91	0.75
MOL004328	Naringenin	59.29	0.21
MOL000417	Calycosin	47.75	0.24
MOL000378	7-O-methylisomucronulatol	74.69	0.3

OB, oral bioavailability; DL, drug-likeness.

Table 2 Top 10 drug-disease targets with degree of the Modified Ginseng-Schisandra Decoction

Uniprot ID	Target	Degree
P35354	Prostaglandin G/H synthase 2, PTGS2	108
Q9SAD4	Ethylene-responsive transcription factor, ESR1	93
P10275	Androgen receptor, AR	88
P37231	Peroxisome proliferator-activated receptor gamma, PPARG	81
P35228	Nitric oxide synthase, NOS2	74
P07477	Trypsin-1, PRSS1	73
Q15596	Nuclear receptor coactivator 2, NCOA2	69
P49841	Glycogen synthase kinase-3 beta, GSK3B	66
Q9FYK5	Ethylene-responsive transcription factor, ESR2	64
P23219	Prostaglandin G/H synthase 1, PTGS1	63

**Figure 3** Degree bar chart of the PPI network target protein of the Modified Ginseng-Schisandra Decoction. PPI, protein-protein interaction network.

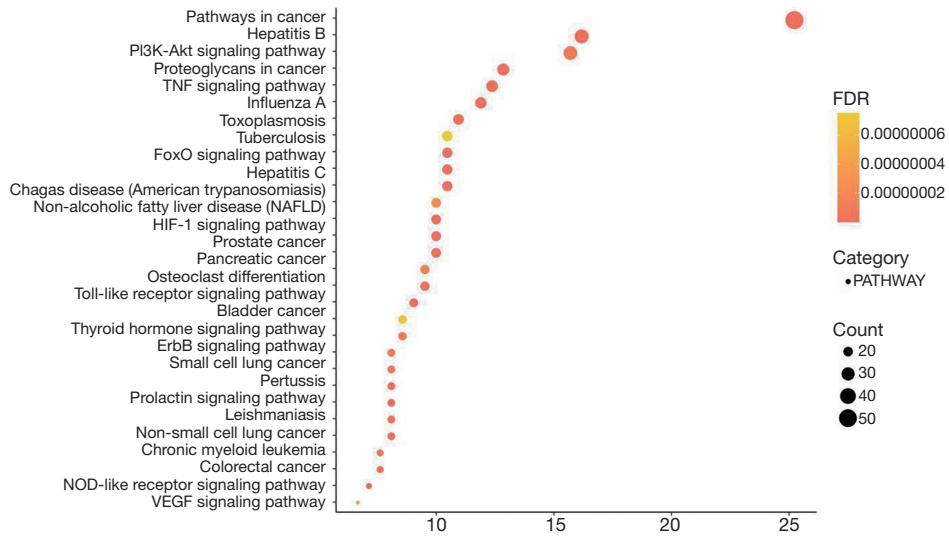


Figure 4 The main signaling pathways involved and mediated by the Modified Ginseng-Schisandra Decoction (top 30).

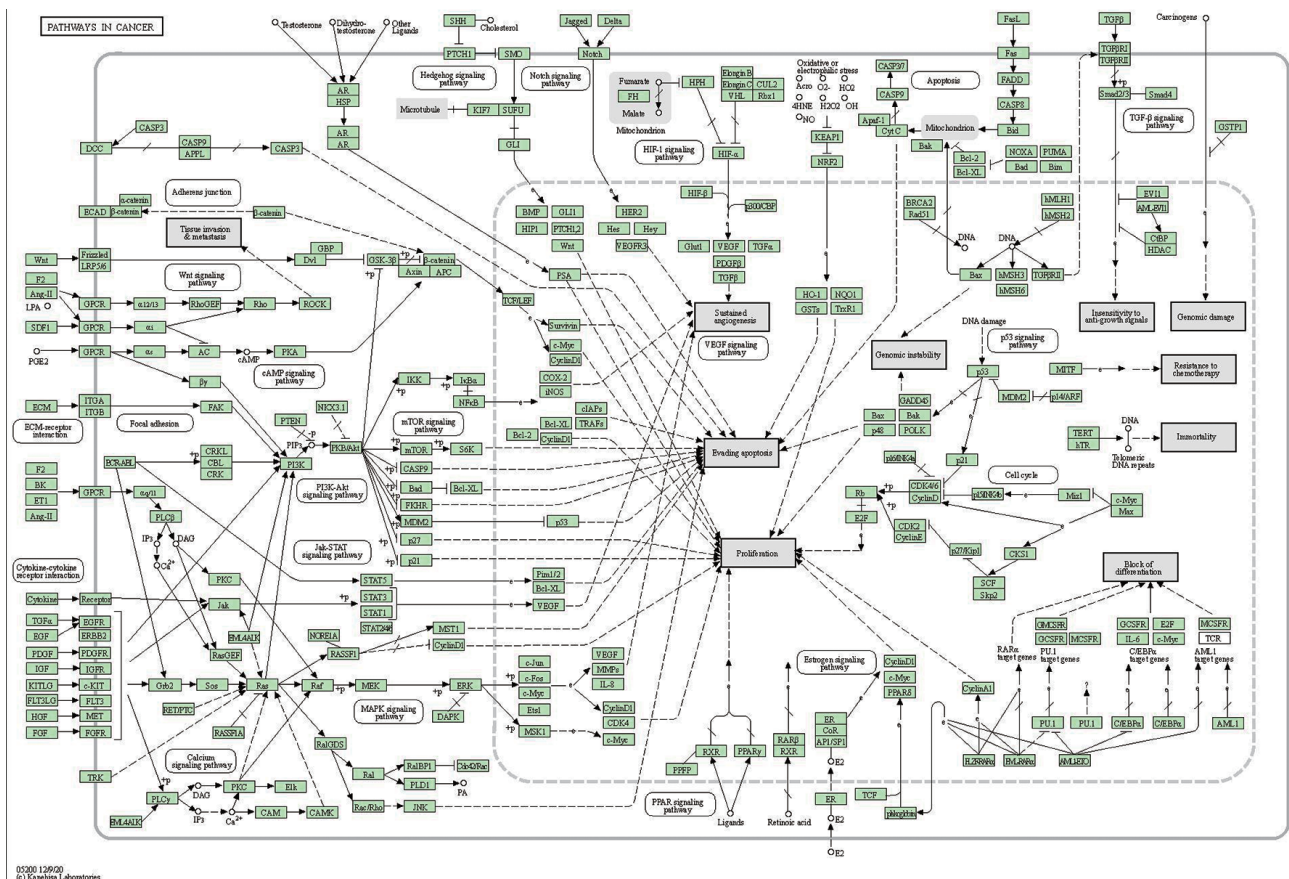


Figure 5 Cancer signaling pathway (Obtained from KEGG database) (13).

TNF SIGNALING PATHWAY

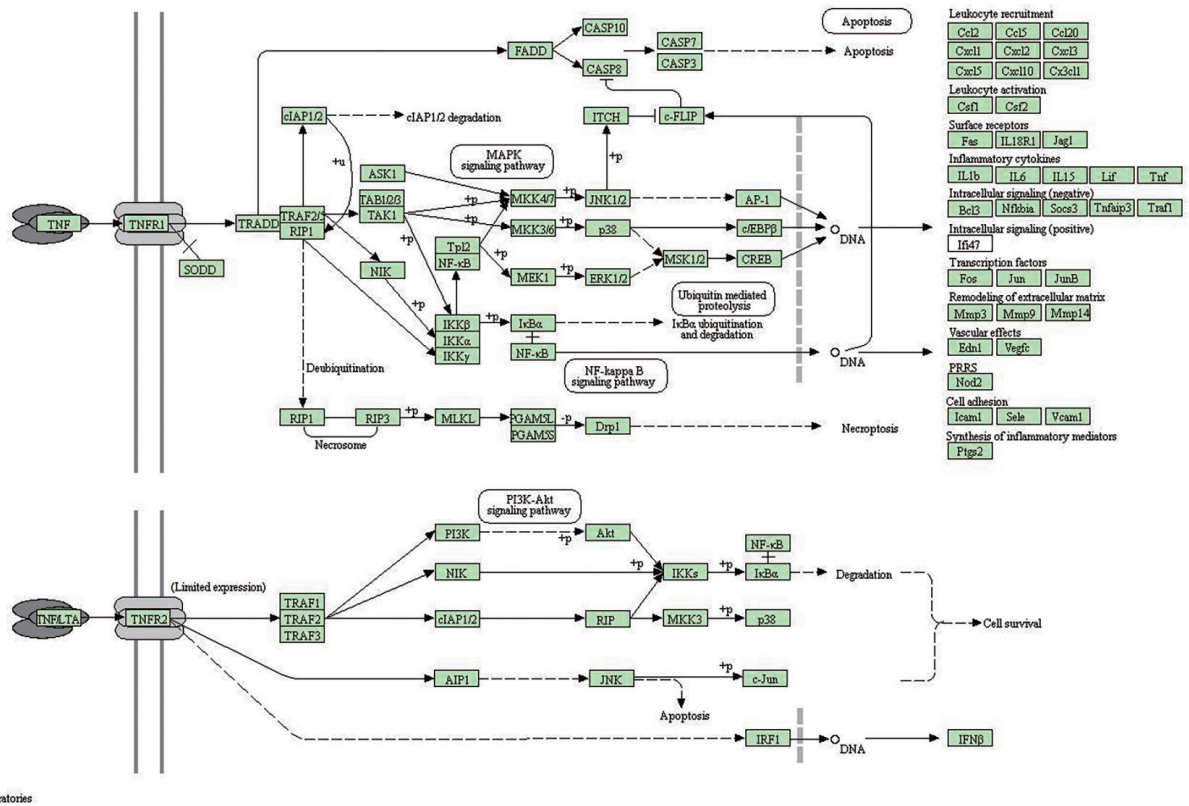


Figure 6 Tumor necrosis factor signaling pathway (Obtained from KEGG database) (13).

modern pharmacological research, *Saposhnikovia divaricata* has antipyretic, analgesic, anti-inflammatory, antibacterial, antiallergic, and regulating immune function effects (16). Traditional Chinese medicine believes that oysters have the functions of submerging Yang and replenishing Yin, reconciling Yin and Yang, and the combination of Yin and Yang means that “Righteousness is stored in the body, and evil cannot be dried.” Modern research has found that oysters’ chemical composition has calcium carbonate, iron, zinc, and other trace elements, and is rich in protein zinc and has a wide range of pharmacological activities (17), which can enhance the body’s defense against nonspecific stimuli. *Radix glycyrrhizae* Preparete can tonify the spleen, replenish qi and harmonize various medicines. The Modified Ginseng-Schisandra Decoction has the effects of nourishing the spleen and qi, nourishing Yin, and moisturizing the lungs.

The DDT network diagram of the Modified Ginseng Decoction in this study shows that, from the point of view of a single medicine, *Radix glycyrrhizae* Preparete can have up to

88 compounds, *Codonopsis pilosula* and *Saposhnikovia divaricata* have 16, and *Radix astragali* has 15. This suggests that *Radix glycyrrhizae* Preparete, *Codonopsis pilosula*, *Radix astragali*, and *Saposhnikovia divaricata* may play a more important role in the Modified Ginseng-Schisandra Decoction treatment of RRTI than previously thought. The results may be related to the single drug quantity and chemical composition. In regard to the compounds, the top three compounds in degree were MOL000098-quercetin, MOL000422-kaempferol, and MOL003896-isoflavone. Quercetin, kaempferol, and isoflavones are all flavonoids. Modern pharmacology has confirmed that quercetin has anti-inflammatory, antibacterial, antioxidant, antiviral, antihistamine, vasodilation, regulation of immune function, and other effects (16), and can interact with viral surface proteins to block viral adhesion (17). Kaempferol has anti-infection, anti-inflammatory, anti-cancer, and other effects (18). Isoflavones have significant antioxidant activity and can scavenge free radicals (19). These findings indicate that Modified Ginseng-Schisandra Decoction may exert a

synergistic effect through its main components, which jointly treat RRTI. In terms of the target, the top five degree values were PTGS2, ESR1, AR, PPARG, and NOS2. PTGS2/COX-2 is a key inducible enzyme that catalyzes prostaglandin synthesis in inflammation, which is enhanced in inflammatory stimulation, tissue damage, and other situations, and participates in the inflammatory response (20). Modern pharmacology has further confirmed that the anti-inflammatory activity of quercetin (21) and kaempferol (22) can act by mediating the expression of COX-2. Kaempferol could inhibit COX-2 expression in MCF-7 cell (23). Kaempferol-3-O-rutinoside inhibits macrophages from secreting inflammatory factors, such as TNF, like IL-6, TNF- α , COX-2 and iNOS (24). Quercetin reduces COX-2 and MPO expression in intestine ischemia reperfusion injury (25). The NF- κ B pathway is inhibited by Quercetin (26). Isoflavone could also inflammation via regulating COX-2 and NF- κ B pathway (27,28). Since Kaempferol, Quercetin and Isoflavone are the main components of Modified Ginseng-Schisandra Decoction, we speculated that the anti-inflammatory effect and treating repeated respiratory infection of Modified Ginseng-Schisandra Decoction might be mediated by the regulation of NF- κ B and COX-2. However, there is no report on Modified Ginseng-Schisandra Decoction targeting COX-2 in the treatment of RRTI, and this should be investigated further. The results of the PPI network shows that STAT3, JUN, MAPK1, AKT1, MAPK3, etc. may be important target groups for the treatment of RRTI with Modified Ginseng Decoction. Among them, the STAT3 degree value was the largest, indicating that it may occupy a major position in the DDT network in this prescription. This is not consistent with PTGS2, ESR1, AR, PPARG and NOS2 being in the core position in the DDT network diagram of Modified Ginseng-Schisandra decoction. This discrepancy be due to the manner in which the Chinese herbal medicine was processed, whether it was decocted, and the specificity of target action, and thus warrants further study. STAT3 is an important node in the development of inflammation. It can regulate the biological behavior of tumor cells and immune cells by mediating extracellular signals of inflammatory mediators, and it is thus a central molecule in the process of inflammation (29). In summary, the Modified Ginseng-Schisandra Decoction may be used to target STAT3 to affect its normal physiological and pathological process in treating RRTI.

The results of KEGG enrichment analysis show that cancer signaling pathway, TNF signaling pathway, and HIF-1 signaling pathway were the leading signaling pathways for the treatment of RRTI by Modified Ginseng-Schisandra Decoction. The cancer signaling pathway contains 15 signal transduction pathway-specific marker genes related to tumor formation. Among them, the necrosis factor kappa beta (NF- κ B) pathway, COX-2 pathway, and STAT pathway may be closely related to the occurrence and development of repeated respiratory infections. The NF- κ B/COX-2 pathway is an important part the important mechanism that leads to inflammatory disease, and it plays a critical role in the regulation of the body's inflammatory immune response (30). Therefore, it can be surmised that the Modified Ginseng-Schisandra Decoction may operate through signal transduction at some key nodes of the NF- κ B/COX-2 pathway to effectuate RRTI treatment. Tumor necrosis factor alpha (TNF- α) is an important mediator and regulator of mammalian immune response in both healthy and diseased conditions. It can control the development of the immune system, cell survival signaling pathways, proliferation, and metabolic processes. Studies have confirmed that gene blockade of TNF receptor 1 (TNFR 1) expression can lead to a phenotype similar to that found in mice that lack TNF-expression, which is characterized by a lack of B cell germinal centers, resistance to liposaccharide (LPS)-mediated systemic toxicity, and increased susceptibility to bacterial infections (31). The results of this study show that the Modified Ginseng-Schisandra Decoction can treat RRTI through TNF signaling pathway. In conclusion, the Modified Ginseng-Schisandra Decoction may play a role in the treatment of RRTI through the cancer signaling pathway, TNF signaling pathway, and HIF-1 signaling pathway.

In summary, this study used the network pharmacology to conduct a preliminary theoretical investigation into the mechanism of the Modified Ginseng-Schisandra Decoction in the treatment of RRTI. The research results clarify the relationship between the main potential active ingredients and target genes of Modified Ginseng-Schisandra Decoction and the pathway of RRTI, providing new ideas for further in-depth exploration of the mechanism of action. The limited information in current databases and the lack of support of experimental results should be considered the limitations of our study. Despite this, the results can be used to direct future research, with further in-depth experimental

investigation being needed in the future.

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Footnote

Reporting Checklist: The authors have completed the MDAR reporting checklist. Available at <https://dx.doi.org/10.21037/tp-21-240>

Conflicts of Interest: All authors have completed the ICMJE uniform disclosure form (available at <https://dx.doi.org/10.21037/tp-21-240>). The authors have no conflicts of interest to declare.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013).

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