Chinese Herbal Medicine Treatment Improves the Overall Survival Rate of Individuals with Hypertension among Type 2 Diabetes Patients and Modulates In Vitro Smooth Muscle Cell Contractility

Ying-Ju Lin¹,²*, Tsung-Jung Ho¹,³,⁴*, Yi-Chun Yeh⁵, Chi-Fung Cheng⁵, Yi-Tzone Shiao⁶, Chang-Bi Wang⁶, Wen-Kuei Chien⁷, Jin-Hua Chen⁷,⁸, Xiang Liu⁹, Hsinyi Tsang⁹, Ting-Hsu Lin², Chiu-Chu Liao², Shao-Mei Huang², Ju-Pi Li¹,¹⁰, Cheng-Wen Lin¹¹, Hao-Yu Pang¹¹, Jaung-Geng Lin¹, Yu-Ching Lan¹², Yu-Huei Liu²,¹³, Shih-Yin Chen¹,², Fuu-Jen Tsai¹,²,¹⁴*, Wen-Miin Liang⁵*

¹ School of Chinese Medicine, China Medical University, Taichung, Taiwan, ² Genetic Center, Department of Medical Research, China Medical University Hospital, Taichung, Taiwan, ³ Division of Chinese Medicine, China Medical University Beigang Hospital, Yunlin, Taiwan, ⁴ Division of Chinese Medicine, Tainan Municipal An-Nan Hospital-China Medical University, Tainan, Taiwan, ⁵ Graduate Institute of Biostatistics, School of Public Health, China Medical University, Taichung, Taiwan, ⁶ Heart Center, China Medical University Hospital, Taichung, Taiwan, ⁷ Biostatistics Center, College of Management, Taipei Medical University, Taipei, Taiwan, ⁸ School of Health Care Administration, College of Management, Taipei Medical University, Taipei, Taiwan, ⁹ National Institute of Allergy and Infectious Diseases, National Institutes of Health, Bethesda, Maryland, United States of America, ¹⁰ Rheumatism Research Center, China Medical University Hospital, Taichung, Taiwan, ¹¹ Department of Medical Laboratory Science and Biotechnology, China Medical University, Taichung, Taiwan, ¹² Department of Health Risk Management, China Medical University, Taichung, Taiwan, ¹³ Graduate Institute of Integrated Medicine, China Medical University, Taichung, Taiwan, ¹⁴ Asia University, Taichung, Taiwan

* These authors contributed equally to this work.
* d0704@mail.cmuh.org.tw (FJT); wmliang.cmu@gmail.com (WML)

Abstract

Type 2 diabetes (T2D) is a chronic, multifactorial, and metabolic disorder accounting for 90% diabetes cases worldwide. Among them, almost half of T2D have hypertension, which is responsible for cardiovascular disease, morbidity, and mortality in these patients. The Chinese herbal medicine (CHM) prescription patterns of hypertension individuals among T2D patients have yet to be characterized. This study, therefore, aimed to determine their prescription patterns and evaluate the CHM effect. A cohort of one million randomly sampled cases from the National Health Insurance Research Database (NHIRD) was used to investigate the overall survival rate of CHM users, and prescription patterns. After matching CHM and non-CHM users for age, gender and date of diagnosis of hypertension, 980 subjects for each group were selected. The CHM users were characterized with slightly longer duration time from diabetes to hypertension, and more cases for hyperlipidaemia. The cumulative survival probabilities were higher in CHM users than in non-CHM users. Among these top 12 herbs, Liu-Wei-Di-Huang-Wan, Jia-Wei-Xiao-Yao-San, Dan-Shen, and Ge-
Gen were the most common herbs and inhibited in vitro smooth muscle cell contractility. Our study also provides a CHM comprehensive list that may be useful in future investigation of the safety and efficacy for individuals with hypertension among type 2 diabetes patients.

Introduction

Type 2 diabetes (T2D) is a chronic, multifactorial, and metabolic disorder and accounts for 90% of those with diabetes worldwide [1]. In Asia and the eastern Pacific region, China was home to the largest number of adults with diabetes (i.e. 90.0 million, or 9% of the population), followed by India (61.3 million, or 8% of the population) and Bangladesh (8.4 million, or 10% of the population) [2–4]. In Taiwan, T2D is one of the top 10 leading causes of death, suggesting that this disease is one of the most important health problems today. T2D is characterized by abnormally high levels of blood glucose resulting from impaired pancreatic β cell function, decreased insulin sensitivity in target tissues, and increased glucose output from the liver [5, 6]. Chronic hyperglycemia causes multiple organ damage and failure, affecting sites including the blood vessels and heart, eyes, kidneys, and nerves. Diabetes related cardiovascular disease, retinopathy, nephropathy, neuropathy, and peripheral circulatory disorders are believed to be responsible for the symptoms, signs, ill-defined secondary conditions, and mortality observed in patients with diabetes.

Hypertension (high blood pressure, usually > 130/80 mmHg) is very common among T2D patients. Almost half of T2D patients have high blood pressure, which doubles their risk of cardiovascular disease [7, 8]. Diabetes related cardiovascular diseases are believed to be responsible for the high morbidity and mortality rates of this condition [9]. In prospective studies, blood pressure control was twice as effective as glucose control in preventing diabetes related cardiovascular disease [10–13]. Therefore, both control of blood pressure and glucose levels in order to prevent substantial diabetic related complications and mortality continues to be an important public health concern.

In diabetic patients, significant improvements can be achieved by lifestyle modification [14] and treatment with hypoglycemic or anti-hyperglycemic, insulin sensitizing, and insulin secretion enhancing agents [15–17]. However, side effects are still frequently reported when using these therapeutic regimes. Meta-analyses show increased cardiovascular and mortality risk when using metformin, sulfonylurea, and thiazolidinediones [15–18]. Long-term thiazolidinedione use increases the risk of fracture, lower respiratory tract infection, and bladder cancer among those with diabetes [17, 19, 20]. These reports have prompted the search for alternative and complementary therapies for better management of diabetes and its related complications. Chinese herbal medicine (CHM) has been used in clinical practice for clinical, chronic, and irreversible diseases for hundreds of years. It has also been used in the management of diabetes, as well as diabetes related complications and mortality [21–23].

CHM is an important aspect of health care in Taiwan and is provided by licensed traditional Chinese medicine (TCM) doctors. It has also been covered under the National Health Insurance (NHI) program since 1996 [24, 25]. Residents in Taiwan are able to choose regular medical treatments, CHM, or both. All claims are collected by the National Health Insurance Research Database (NHIRD). Therefore, this claim database can be used as a platform to explore the utilization and therapeutic effects of Chinese herbal therapies prescribed by these TCM doctors in Taiwan. The characteristics of TCM use in Taiwan have been investigated by population-based studies for several diseases including childhood asthma [26], breast cancer
[27], chronic kidney disease [28], diabetes [29], endometriosis [30], primary dysmenorrhea [31], schizophrenia [32], and Sjögren's syndrome [33] etc.

In this study, we also used a population-based database to investigate the demographic characteristics, the overall survival analysis and prescription patterns of individuals with hypertension among type 2 diabetes patients according to CHM usage. In addition, we also evaluate the effect of selected herbal formulas and single herbs on smooth muscle cell contractility.

Results
Demographic characteristics and overall survival analysis of individuals with hypertension among type 2 diabetes patients according to CHM usage

In this study, the database claims identified 984 CHM users and 2,434 non-CHM users with hypertension among type 2 diabetes patients from a cohort of one million randomly sampled cases from the National Health Insurance Research Database (NHIRD) [34] (Fig 1A). The demographic characteristics of CHM users versus non-CHM users (total subjects) are shown in the left side of Table 1. There were significant different frequency distributions for age, gender, duration from diabetes to hypertension, comorbidity (cardiovascular disease and hyperlipidaemia), and income for these two groups \( p < 0.05 \). The CHM users (total subjects) were characterized with younger age, more females, longer duration time from diabetes to hypertension, lesser cases for cardiovascular disease, more cases for hyperlipidaemia, and higher incomes. The one-to-one match method was used to match CHM users and non-CHM users. After matching these two groups for age, gender and date of diagnosis of hypertension, CHM and non-CHM users were selected (Fig 1B and Table 1 right side). There were significant different frequency distributions for duration from diabetes to hypertension, and hyperlipidaemia \( p < 0.05 \). The CHM users were characterized with slightly longer duration time from diabetes to hypertension, and more cases for hyperlipidaemia.

The cumulative survival probability of individuals with hypertension among type 2 diabetes patients according to CHM usage were shown in Fig 2. The overall survival rate was different between CHM users and matched non-CHM users \( p < 0.001 \). The cumulative survival probabilities were higher in CHM users than in matched non-CHM users suggesting that CHM may be beneficial for longer survival of hypertension individuals among type 2 diabetes patients.

Twelve most common herbal formulas and single herbs prescribed by TCM doctors for the treatment of hypertension individuals among type 2 diabetes patients

The 12 most common Chinese herbal formulas and single herbs prescribed for the CHM users analyzed in this study are listed in Table 2. The follow-up person-years were from hypertension to the study end as shown in Fig 1B. The herbal composition of these herbal formulas and single herbs were also shown in Table A in S1 File. Shu-Jing-Huo-Xue-Tang was the most commonly prescribed herbal formula, followed by Liu-Wei-Di-Huang-Wan and Jia-Wei-Xiao-Yao-San. Among the top 12 herbal formulas, Ji-Sheng-Shen-Qi-Wan and Zi-Bai-Di-Huang-Wan are 2 derivative formulas of Liu-Wei-Di-Huang-Wan. Therefore, Liu-Wei-Di-Huang-Wan and its various derivatives were the most common herbal formulas prescribed by Chinese medical doctors for the CHM users in this study. Of the 12 most common single herbs, Yan-Hu-Suo was the most commonly prescribed, followed by Dan-Shen and Ge-Gen.
Fig 1. Flow recruitment diagram. A: Chart showing the protocol for enrollment of study subjects. B: Follow-up time for CHM and matched non-CHM users.

doi:10.1371/journal.pone.0145109.g001
Table 1. Demographic characteristics of total subjects and frequency matched subjects with hypertension among type 2 diabetes patients according to CHM usage.

| Characteristics                  | Total subjects | Frequency matched subjects |
|----------------------------------|----------------|---------------------------|
|                                  | Number         | Number                    |
|                                  | N = 2,436      | N = 984                   |
|                                  | N %            | N %                       |
|                                  | p value        |                           |
|                                  | N = 980        | N = 980                   |
|                                  | N %            | N %                       |
|                                  | p value        |                           |
| **Age**                          |                |                           |
| <60 yrs                          | 1,406          | 965                       | 441 | 44.82 | <0.0001 |
| 60–70 yrs                        | 978            | 662                       | 316 | 32.11 |          |
| 70–80 yrs                        | 767            | 584                       | 183 | 18.6  |          |
| >= 80 yrs                        | 269            | 225                       | 44  | 4.47  |          |
| **Gender**                       |                |                           |
| Male                             | 2,042          | 1,566                     | 476 | 48.37 | <0.0001 |
| Female                           | 1,378          | 870                       | 508 | 51.63 |          |
| **Duration from diabetes to hypertension** |            |                           |
| 1–2 years                        | 919            | 694                       | 225 | 22.87 | 0.0006  |
| 2–4 years                        | 1,512          | 1,074                     | 438 | 44.51 |          |
| >= 5 years                       | 989            | 668                       | 321 | 32.62 |          |
| **Cardiovascular disease**       |                |                           |
| No                               | 2,852          | 1,999                     | 853 | 86.69 | 0.001   |
| Yes                              | 568            | 437                       | 131 | 13.31 |          |
| **Ischaemic heart disease**      |                |                           |
| No                               | 2,664          | 1,889                     | 775 | 78.76 | 0.4382  |
| Yes                              | 756            | 547                       | 209 | 21.24 |          |
| **Chronic kidney disease**       |                |                           |
| No                               | 3,225          | 2,288                     | 937 | 95.22 | 0.1692  |
| Yes                              | 195            | 148                       | 60  | 6.08  |          |
| **Hyperlipidaemia**              |                |                           |
| No                               | 1,964          | 1,468                     | 496 | 50.41 | <0.0001 |
| Yes                              | 1,456          | 968                       | 488 | 49.59 | 0.0145  |
| **Obesity**                      |                |                           |
| No                               | 3,374          | 2,409                     | 965 | 98.07 | 0.0587  |
| Yes                              | 46             | 27                        | 19  | 1.93  |          |
| **Alcohol-related illness**      |                |                           |
| No                               | 3,369          | 2,397                     | 972 | 98.78 | 0.4047  |
| Yes                              | 51             | 39                        | 12  | 1.22  |          |
| **Tobacco use**                  |                |                           |
| No                               | 3,401          | 2,426                     | 975 | 99.09 | 0.0726  |
| Yes                              | 19             | 10                        | 9   | 0.91  |          |
| **INCOME**                       |                |                           |
| <NT20000                         | 739            | 582                       | 157 | 15.96 | <0.0001 |
| NT20000–NT30000                  | 677            | 491                       | 186 | 18.9  |          |
| NT30000–NT40000                  | 1,438          | 982                       | 456 | 46.34 |          |
| >= NT40000                       | 566            | 381                       | 185 | 18.8  |          |
| **Urbanization level**           |                |                           |
| 1                                | 817            | 574                       | 243 | 24.7  | 0.3331  |
| (Continued)                      |                |                           |

Continued.
Effect of most common herbal formulas and single herbs on smooth muscle cell contractility

Smooth muscle cell contractility can be monitored by measuring the phosphorylation of myosin light chain protein [35]. The Y27632 compound (Rho kinase inhibitor) was used as the control for decreased myosin light chain phosphorylation [36]. And the calyculin compound was used as the control for increased myosin light chain phosphorylation [35]. We chose most common used two herbal formulas and two single herbs from these top 12 herbs (herbal Table 1. (Continued))

| Characteristics | Total subjects | Frequency matched subjects |
|-----------------|----------------|---------------------------|
|                 | Total Number   | non-CHM user | CHM user | p value | Total Number | non-CHM user | CHM user | p value |
|                 | N = 2,436      | N = 984       |          |          | N = 980      | N = 980      |          |          |
|                 | N %            | N %          |          |          | N %          | N %          |          |          |
| 2               | 1,033          | 739          | 30.34    | 294      | 29.88        | 292          | 29.8      |          |
| 3               | 497            | 340          | 13.96    | 157      | 15.96        | 157          | 16.02     |          |
| 4               | 585            | 421          | 17.28    | 164      | 16.67        | 163          | 16.63     |          |
| 5               | 488            | 362          | 14.86    | 126      | 12.8         | 125          | 12.76     |          |

CHM, Chinese herbal medicine; N, number; NT, new Taiwan dollars.
Urbanization level: 1 indicates the highest level of urbanization and 5 is the lowest level.
p values were obtained by chi-square test.
p value (p < 0.05) was highlighted in bold italic.
doi:10.1371/journal.pone.0145109.t001

Fig 2. Cumulative survival curves of individuals with hypertension among type 2 diabetes patients according to Chinese herbal medicine (CHM) usage.
doi:10.1371/journal.pone.0145109.g002
formulas: Liu-Wei-Di-Huang-Wan and Jia-Wei-Xiao-Yao-San; single herbs: Dan-Shen and Ge-Gen) according to their frequencies of prescriptions and average duration for prescription. A10 cells (rat smooth muscle cells) were treated with these herbs at the concentrations as indicated (Fig 3A and 3B; S3A and S3B Fig). As shown, these four herbs reduced phosphorylation of myosin light chain. These herbs inhibited the phosphorylation of myosin light chain protein, suggesting that these most common herbs may be beneficial for smooth muscle cell contractility.

Table 2. Twelve most common herbal formulas and single herbs prescribed by TCM doctors for the treatment of hypertension individuals among type 2 diabetes patients.

| Number of Person — years | Frequency of prescriptions | Percentage of usage person | Average daily dose (g) | Average duration for prescription (days) |
|--------------------------|---------------------------|-----------------------------|------------------------|-----------------------------------------|
| **Total**                | 4,875                     | 38,140                      | 100                    | 11.9                                    | 7.6                                     |
| **Herbal formula**       |                           |                             |                        |                                         |                                         |
| Shu-Jing-Huo-Xue-Tang    | 2,153                     | 1,995                       | 39.2                   | 3.9                                     | 6.8                                     |
| Liu-Wei-Di-Huang-Wan     | 1,906                     | 2,137                       | 34.3                   | 4                                       | 8.3                                     |
| Jia-Wei-Xiao-Yao-San     | 1,719                     | 1,625                       | 32.4                   | 4.1                                     | 8.5                                     |
| Ge-Gen-Tang              | 1,674                     | 1,190                       | 31.2                   | 4.3                                     | 6.7                                     |
| Shao-Yao-Gan-Cao-Tang    | 1,643                     | 1,150                       | 30.7                   | 3.3                                     | 7.2                                     |
| Ma-Xing-Shi-Gan-Tang     | 1,549                     | 1,349                       | 29.1                   | 3.9                                     | 6.2                                     |
| Xue-Fu-Zhu-Yu-Tang       | 1,557                     | 1,308                       | 28.4                   | 4                                       | 8.9                                     |
| Du-Huo-Ji-Sheng-Tang     | 1,479                     | 1,380                       | 27.4                   | 4.7                                     | 7.8                                     |
| Chuan-Xiong-Cha-Tiao-San | 1,404                     | 1,062                       | 26.9                   | 4                                       | 6.2                                     |
| Ji-Sheng Shen-Qi-Wan     | 1,438                     | 1,576                       | 26                     | 4.1                                     | 9.6                                     |
| Gan-Lu-Yin               | 1,387                     | 1,391                       | 25.9                   | 3.7                                     | 7.5                                     |
| Zhi-Bai-Di-huang-Wan     | 1,437                     | 1,364                       | 25.8                   | 4                                       | 10.1                                    |
| **Single herb**          |                           |                             |                        |                                         |                                         |
| Yan-Hu-Suo               | 4,732                     | 29,455                      | 97.2                   | 4                                       | 7.8                                     |
| Ge-Gen                   | 1,830                     | 1,665                       | 35.2                   | 1.1                                     | 7.6                                     |
| Dan-Shen                 | 1,716                     | 1,596                       | 34.4                   | 1.4                                     | 8.2                                     |
| Tian-Hua-Fen             | 1,798                     | 2,179                       | 34.3                   | 1.3                                     | 10.3                                    |
| Jie-Geng                 | 1,751                     | 1,697                       | 33.2                   | 1.1                                     | 9.2                                     |
| Bei-Mu                   | 1,618                     | 1,436                       | 31.5                   | 1                                       | 6.6                                     |
| Huang-Qin                | 1,607                     | 1,409                       | 31.5                   | 1.1                                     | 8                                       |
| Niu-Xi                   | 1,635                     | 1,345                       | 30.6                   | 0.9                                     | 7.9                                     |
| Mai-Men-Dong             | 1,516                     | 1,287                       | 28.6                   | 1.2                                     | 8.9                                     |
| Huang-Qi                 | 1,473                     | 1,807                       | 28.3                   | 1.4                                     | 8.8                                     |
| Xuan-Shen                | 1,496                     | 1,157                       | 28.2                   | 1.2                                     | 9                                       |
| Xing-Ren                 | 1,472                     | 1,057                       | 27.9                   | 1.1                                     | 6.7                                     |

TCM, traditional Chinese medicine.

Follow-up time was from hypertension to the study end (Fig 1B).

doi:10.1371/journal.pone.0145109.t002
In this study, we used a population-based database to investigate the demographic characteristics, the overall survival analysis and prescription patterns of individuals with hypertension.

**Discussion**

In this study, we used a population-based database to investigate the demographic characteristics, the overall survival analysis and prescription patterns of individuals with hypertension.
among type 2 diabetes patients according to CHM usage. In addition, we also evaluate the effect of two herbal formulas and two single herbs from these top 12 herbs on smooth muscle cell contractility. We found that the cumulative survival probabilities were higher in CHM users than in non-CHM users. We also described the most common prescribed CHMs. The single herbs and the herbal formulas inhibited smooth muscle cell contractility. Our results suggest that adjunctive CHM therapy treatment may improve the overall survival rate of individuals with hypertension among type 2 diabetes patients and some of them modulate smooth muscle cell contractility.

Our results showed that the overall survival rate was higher in CHM users than in non-CHM users from hypertension individuals among type 2 diabetes patients. And we also found that patients treated with any CHM, herbal formulas or single herbs had the trend of lower risks of the death, macrovascular and microvascular diseases as the endpoints after adjusted for age, duration from diabetes to hypertension, and comorbidities by the conditional logistic analysis (Tables B–G in S1 File).

The regular medical treatments (other than CHM) between CHM and non-CHM users showed that there were more patients in non-CHM users who have used anti-diabetes drugs from diabetes to the index date (Table H in S1 File). However, there were no anti-diabetes drug usage differences between these two groups from index date to index date + 365 (Table I in S1 File). As for the anti-hypertension drugs from diabetes to the index day (Table H in S1 File), there were more patients in non-CHM users who have used anti-hypertension drugs- ACEI or ARB (p < 0.05). However, there were more patients in CHM users who have used anti-hypertension drugs- beta blocking agents (p < 0.05). From index date to index date + 365 (Table I in S1 File), there were more patients in non-CHM users who have used anti-hypertension drugs- ACEI or ARB (p < 0.05). CHM has been reported to reduce progression from impaired glucose tolerance to diabetes [37, 38]. Furthermore, CHMs have been used to successfully treat diabetes via increased insulin secretion and sensitivity, enhanced glucose uptake by adipose and muscle tissues, inhibition of glucose absorption by the intestine, inhibition of glucose production by hepatocytes, and anti-inflammatory activity [21–23, 39].

The most common herbal formulas were Liu-Wei-Di-Huang-Wan and Jia-Wei-Xiao-Yao-San. Liu-Wei-Di-Huang-Wan were the most common herbal formula in this study. Its various derivatives (Ji-Sheng-Shen-Qi-Wan, and Zi-Bai-Di-Huang-Wan) were also noted in our herbal formula list. Liu-Wei-Di-Huang-Wan are composed of Rx. Rehmanniae Preparata, Fr. Corni, Rx. Dioscoreae, Poria, Cx. Moutan, and Rz. Alismatis. Liu-Wei-Di-Huang-Wan has been used to treat diabetes, pre-diabetes, fatigue, and metabolic syndrome [29, 40]. Furthermore, scientific evidence has suggested that Liu-Wei-Di-Huang-Wan can decrease visceral fat deposition [41], increase plasma levels of adiponectin and improve insulin resistance [42], and improve the lipid profile indicating a reduction of cardiovascular risk [43]. And Liu-Wei-Di-Huang-Wan combined with antihypertensive drugs appears to be effective in improving blood pressure and symptoms in patients with essential hypertension [44]. We have found that there were also no significant differences in the osmolarity and cell survival rate of cells among these herbal formulas and single herbs as compared with the cell only control, suggesting that the osmolarity of the Chinese herbal medicine are suitable for the cells in culture (S1A and S1B Fig). Furthermore, our functional analysis by measuring the phosphorylation of myosin light chain protein and the collage contraction assay (S2A–S2D Fig) first showed that smooth muscle cell contractility was inhibited by treatment with Liu-Wei-Di-Huang-Wan, which was in agreement with previous clinical observations [44]. Jia-Wei-Xiao-Yao-San are composed of Rx. Angelicae Sinensis, Rx. Paeoniae Alba, Poria, Rz. Atractylodis Macrocephalae, Rx. Bupleuri, Cx. Moutan, Fr. Gardeniae, and Rx. Gly. Jia-Wei-Xiao-Yao-San is used to treat symptoms including nervousness, palpitations, headache, anorexia, night sweating, dry eyes, hot flashes,
and irregular menstruation; it also has hepatoprotective effects [45–49]. However, there were no related literatures related to the effect of Jia-Wei-Xiao-Yao-San on diabetes or hypertension. To our knowledge, this is the first study to show that Jia-Wei-Xiao-Yao-San can inhibit smooth muscle cell contractility by measuring the phosphorylation of myosin light chain protein and the collage contraction assay (S2A–S2D Fig).

Dan-Shen was the most common single herb and composed of Radix Salviae Miltiorrhizae. In previous studies, Dan-Shen has been shown to have protective effects on the cardiovascular system [50–54] and pulmonary arteries [55, 56]. Furthermore, active component (SalB) from Dan-Shen can exhibit antidiabetic activity and inhibit symptoms of diabetes mellitus in rats and these effects may partially be correlated with its insulin sensitivity, glycogen synthesis and antioxidant activities [57–60]. Ge-Gen was composed of Radix Puerariae and contains an isoflavonoid glycoside with hypotensive effects, with excellent clinical results in the treatment of hypertension [61]. Furthermore, puerarin is a major active ingredient of Ge-Gen and exerts significant protective effects against diabetic retinopathy in rats via regulating angiogenesis factors expressions [62]. Interestingly, our results suggest that patients treated with the single herb- Ge-Gen had the statistical significance of lower risk of acute myocardial infarction and nephropathy (Table B and F in S1 File). There were no trends or statistical significance observed from the ischemic stroke, hemorrhagic stroke and amputation as the endpoint (Table C–E in S1 File). We are the first to suggest that Dan-Shen and Ge-Gen were the most common single herbs for individuals with hypertension among type 2 diabetes patients and in vitro functional analysis suggested that smooth muscle cell contraction was inhibited by treatment with these herbs.

By integrating the National Health Insurance Research Database (NHIRD) review with our in vitro functional data, we were able to investigate the mechanism of action of CHM in the treatment of disease. Limitations of this study included a lack of blood physiological and biochemical measures in this database, such as blood pressure or blood sugar. The NHIRD limitations also include lacks of genetic factors, environmental factors (including levels of job stress and exercise), personal histories (including education and body mass index), and potential disease misclassifications [63–67]. The usage of CHM improves the overall survival rate of individuals with hypertension among type 2 diabetes patients and also these CHM treatment modulates smooth muscle cell contractility. Our study provides a CHM comprehensive list that may be useful in future investigation of the safety and efficacy for individuals with hypertension among type 2 diabetes patients.

Materials and Methods

Ethical statement

This study was evaluated and approved for the purchase of the National Health Insurance Research Database (NHIRD) by the Human Studies Committee of China Medical University Hospital, Taichung, Taiwan. No informed consent was required because the data were analyzed anonymously. The cell line rat aortic smooth muscle cell line A10 cells (BCRC number:60127; used in Fig 3A and 3B) were purchased from Food Industry Research and Development Institute in Taiwan (https://catalog.bcrifirdi.org.tw/BSAS_cart/controller?event=SEARCH&bcrc_no=60127&type_id=4&keyword=smooth;•muscle;•cells). These cells were derived from the thoracic aorta of rats and served as a commonly used model of vascular smooth muscle cells [35] and were approved by the Animal Care and Use Committee (IACUC) of China Medical University, Taichung, Taiwan.
National Health Insurance Research Database (NHIRD) resource in Taiwan

The national health insurance (NHI) program in Taiwan was started in 1995 to make health care available for all residents of Taiwan. As of 2010, over 99% of residents were enrolled in the program [68]. The NHI program provides the National Health Insurance Research Database (NHIRD) resource (http://nhird.nhri.org.tw/en/index.htm) for scientists in Taiwan and only for research purposes. Data for this study were retrieved from the “Longitudinal Health Insurance Database (LHID2000)”, which includes all the original claim data and registration files for 1,000,000 beneficiaries, randomly sampled from the year 2000 Registry for Beneficiaries (n = 23.72 million) under the NHI program. This database contains information on patient demographics, diagnoses, prescriptions, records of clinical visits and hospitalizations, inpatient orders, ambulatory care, and socio-demographic factors. Disease diagnoses are coded using the International Classification of Disease, 9th Revision, Clinical Modification (ICD-9-CM). This database also include traditional Chinese medicine services (Chinese herbal medicine (CHM), acupuncture, and manipulative therapies; http://www.nhi.gov.tw/English/webdata/webdata.aspx?menu=11&menu_id=592&WD_ID=592&webdata_id=3161). The data are from the National Health Insurance Research Database, Taiwan (NHIRD) http://nhird.nhri.org.tw/en/index.html. Contact nhird@nhri.org.tw for details and data access.

Study population

This study was designed as a population-based retrospective cohort study. In this study, a cohort of one million individuals randomized selected from NHI (Taiwan) was used. The sampled population was representative of all NHI beneficiaries. The study subjects were selected from this cohort and were shown in Fig 1A. There were 84,032 individuals with diabetes (ICD-9-CM: 250) between 1998 and 2010. The hypertension ICD-9-CM used in this study was from 401–405. Individuals under the age of 20 were excluded. Individuals without hypertension (ICD-9-CM: 401–405), who had hypertension before diabetes, who had hypertension within 1 year after diabetes, and who had hypertension after 2009 were also excluded. In addition, at least one of the following enrollment criteria had to be met for identifying patients with hypertension in the study: (1) one or more inpatient admissions with diagnosis of hypertension, or (2) three or more outpatient visits within one-year period, each with a diagnosis of hypertension. The first date which satisfied the above (1) or (2) criteria was defined as the date of diagnosis of hypertension. After all of these criteria were applied, 10,664 study subjects were included in the study cohort.

Definition of CHM and non-CHM users

Study subjects with a record of cumulative CHM drug days more than 28 within first year after hypertension were defined as CHM users (N = 984, Fig 1B). Study subjects with no recorded of CHM usage were defined as non-CHM users (N = 2,434). The date of satisfying the criterion of cumulative 28 drug days of CHM prescription was designated as the index date (Fig 1B). The one-to-one match method was used to match CHM users and non-CHM users. After matching these two groups for age, gender and date of diagnosis of hypertension, CHM and non-CHM users were selected (Fig 1B and Table 1 right side). A total of 980 subjects for each group were selected. The study endpoint was the following: date of death, date of withdrawal from the NHI program, or date of follow-up termination (31 Dec. 2010). This study was designed as a population-based retrospective cohort study and to explore the effect of Chinese Herbal Medicine treatment on the overall survival rate of individuals with hypertension among type 2 diabetes patients.
Chinese herbal medicine (CHM)

All drug codes for CHM (herbal formulas and single herbs) were collected and grouped according to their name. The frequencies of prescriptions, cumulative drug doses, average durations of per prescription, and follow-up person years were calculated from hypertension to the study end for the CHM users. Herbal formulas usually constituted a combination of 2 to 17 herbs (Table A in S1 File), created by experienced TCM doctors; these formulas have been used for thousands of years. Single herbs were obtained from plant, animal, or mineral material and can be mixed with other herbs to create a formula. The herbal formulas and single herbs in the NHIRD database were all produced by Good Manufacturing Practice (GMP) certified traditional Chinese medicine manufacturers based in Taiwan. These manufacturers included Sun Ten Pharmaceutical Co. Ltd., Shang Chang Pharmaceutical Co. Ltd., Chuang Song Zong Pharmaceutical Co. Ltd., KO DA Pharmaceutical Co. Ltd., and Kaiser Pharmaceutical Co. Ltd.

Study covariates

We collected demographic data such as age, gender, income and urbanization levels. Urbanization levels in Taiwan are divided into five strata according to the Taiwan National Health Research Institute publications, with level 1 referring to the most urbanized communities and level 5 referring to the least urbanized communities. We identified the diagnoses of comorbidities which defined by following diagnoses recorded before the diagnosis date of hypertension: cardiovascular disease (ICD-9-CM: 430–437), ischaemic heart disease (ICD-9-CM: 410–414), chronic kidney disease (ICD-9-CM: 582–583) and hyperlipidaemia (ICD-9-CM: 272) (Table 1). The diagnosis criteria for each comorbidity were similar to those for hypertension.

We also applied the conditional multivariable logistic regression adjusted for all variables in the Table 1 to assess the effect of CHM on the occurrence of blood pressure related disease such as acute myocardial infarction, ischemic stroke, hemorrhagic stroke, amputation, and nephropathy (the results were shown in Tables B–F in S1 File). The covariates included CHM user, age, income, duration from diabetes to hypertension, and comorbidities before hypertension including cardiovascular disease (ICD-9-CM: 430–437), ischaemic heart disease (ICD-9-CM: 410–414), chronic kidney disease (ICD-9-CM: 582–583) and hyperlipidaemia (ICD-9-CM: 272).

Cell culture, reagents, and Western blotting

Rat aortic smooth muscle cells (A10 cell line) were cultured in Dulbecco’s Modified Eagle's Medium (DMEM) supplemented with 10% fetal bovine serum (FBS), 100 U/mL penicillin, 100 U/mL streptomycin, and 2 mM L-glutamine (Gibco). Y27632, and calyculin A were purchased from Sigma (St. Louis, MO, USA). A10 cells were treated with Y27632 (10 μM), and calyculin A (50 μg/ml) for 10 min. The treated cells were lysed in RIPA buffer (Thermo Scientific ™) and then were applied to Western blot analysis and staining with anti-phospho-myosin light chain (MLC) (1:1,000 dilution), anti-total-MLC (1:1,000 dilution), and anti-beta actin (1:1,000 dilution) antibodies (Fig 3; S3 Fig). The monoclonal anti-phospho-MLC (phospho-myosin light chain 2 [Ser19] mouse mAb; catalog number: 3675) and polyclonal anti-total-MLC (myosin light chain 2 antibody; catalog number: 3672) rabbit antibodies were from Cell Signaling Technology, Inc. The anti-beta actin (actin antibody [mAbGEa]; catalog number: NB100-74340) mouse monoclonal antibody was obtained from Novus Biologicals. The experimental protocol used for Western blotting has been described previously [69, 70]. Briefly, cells were harvested, washed, and lysed in lysis buffer (50 mM Tris-HCl [pH 7.5], 150 mM NaCl, 5 mM EDTA, 1% Triton X-100, 0.1% SDS) supplemented with protease inhibitor cocktail (Roche). The lysates were resolved by 12% SDS-PAGE and transferred to polyvinylidene fluoride membranes.
The membranes were incubated with primary antibodies overnight at 4°C and then incubated with alkaline phosphatase-conjugated secondary antibodies (Goat anti-Mouse IgG (H+L) Polyclonal Secondary Antibody, HRP conjugate; 1:5000 dilutions; catalog number: A16072; Thermo Fisher Scientific). Signals were visualized using a SuperSignal West Femto Maximum Sensitivity Substrate Detection Kit; catalog number: 34096; Thermo Fisher Scientific) in accordance with the manufacturer’s instructions.

Statistical analysis

We presented demographic data such as age, gender, duration from diabetes to hypertension, comorbidities (cardiovascular disease, ischaemic heart disease, chronic kidney disease, and hyperlipidaemia), income, and urbanization level for both groups (CHM and non-CHM users) using count and percentage for categorical variables, and used chi-squared tests to assess their differences (Table 1). We sorted the cumulative person-years for each herbal formula and single herb and listed the top 12 most common herbal formulas and single herbs (Table 2). We employed Kaplan-Meier method to estimate the cumulative survival probabilities and used the log-rank test to explore the effect of Chinese Herbal Medicine treatment on the overall survival rate of individuals with hypertension among type 2 diabetes patients (CHM and non-CHM users; Fig 2). We also used conditional logistic analysis to explore the effect of CHM therapy as well as these most commonly used herbs on the reduction of macrovascular and microvascular diseases, and death as the endpoints during follow up (Tables B–G in S1 File). All p-values less than 0.05 were considered significant. All data management and statistical analyses were performed using Statistical Analysis System (SAS) software (version 9.3; SAS Institute, Cary, NC, USA).

Supporting Information

S1 Fig. Osmolarity and cell survival rate of cells treated with Chinese herbal medicine. (A) Detection of the osmolarity from the cell culture medium as shown above by using Vapro TM Osmometer, Model 5520. The standard 290 and the concentrations of NaCl (0%, 0.5%, 1%, and 2%) were used as the controls. Cells were in the presence of cells only, Y10 (Y27632 at 10 μM), single herbs- Dan-Shen and Ge-Gen, and herbal formula- Liu-Wei-Di-Huang-Wan and Jia-Wei-Xiao-Yao-San (5 and 10 μg/ml). Similar results were obtained in three independent experiments. Values represent the mean± S.D. (B) % of cell survival rate of cells treated with Chinese herbal medicine. Cells were in the presence of cells only, Y10 (Y27632 at 10 μM), single herbs- Dan-Shen and Ge-Gen, and herbal formula- Liu-Wei-Di-Huang-Wan and Jia-Wei-Xiao-Yao-San (5 and 10 μg/ml) for 24 h and were detected by using the WST-1 assay. Similar results were obtained in three independent experiments. Values represent the mean± S. D.

S2 Fig. Effect of Chinese herbs on contraction of collagen gels. Cell-embedded collagen gels were prepared according to the manufacturer’s instructions (CELL BIOLABS, INC., cell contraction assay (catalog number CBA-201-T). (A) The surface area of collagen gels was calculated at 24 h, 48 h, and 120 h in the presence of cells only, Y10 (Y27632 at 10 μM), 1X BDM-contraction inhibitor as the controls. The surface area of collagen gels of single herbs- Dan-Shen and Ge-Gen, and herbal formula- Liu-Wei-Di-Huang-Wan and Jia-Wei-Xiao-Yao-San (5 and 10 μg/ml) was also calculated. The contraction of collagen gel was expressed in a percentage, with the surface area of the cells only serving as 100%. Similar results were obtained in three independent experiments. Values represent the mean± S.D. (B) The surface area of
collagen gels was shown at 24 h in the presence of cells only (No. 1), Y10 (No. 2; Y27632 at 10 μM), 1X BDM-contraction inhibitor (No. 3) as the controls. The surface area of collagen gels of single herbs- Dan-Shen (No. 4; 5 μg/ml) and Ge-Gen (No. 5; 5 μg/ml), and herbal formula- Liu-Wei-Di-Huang-Wan (No. 6; 5 μg/ml) and Jia-Wei-Xiao-Yao-San (No. 7; 5 μg/ml) was also shown. (C) The surface area of collagen gels was shown at 48 h in the presence of cells only (No. 1), Y10 (No. 2; Y27632 at 10 μM), 1X BDM-contraction inhibitor (No. 3) as the controls. The surface area of collagen gels of single herbs- Dan-Shen (No. 4; 5 μg/ml) and Ge-Gen (No. 5; 5 μg/ml), and herbal formula- Liu-Wei-Di-Huang-Wan (No. 6; 5 μg/ml) and Jia-Wei-Xiao-Yao-San (No. 7; 5 μg/ml) was also shown. (D) The surface area of collagen gels was shown at 120 h in the presence of cells only (No. 1), Y10 (No. 2; Y27632 at 10 μM), 1X BDM-contraction inhibitor (No. 3) as the controls. The surface area of collagen gels of single herbs- Dan-Shen (No. 4; 5 μg/ml) and Ge-Gen (No. 5; 5 μg/ml), and herbal formula- Liu-Wei-Di-Huang-Wan (No. 6; 5 μg/ml) and Jia-Wei-Xiao-Yao-San (No. 7; 5 μg/ml) was also shown. (PPTX)

S3 Fig. Original uncropped and unadjusted blots of Fig 3. (A) herbal formulas; (B) single herbs. The antibodies (anti-Phospho-MLC, anti- Total-MLC, and anti-β-actin) used here were shown in the left of the S3 Fig. (PPTX)

S1 File. Supporting tables for Chinese herbal medicine treatment in hypertension individuals among type 2 diabetes patients. Herbal composition of twelve most common herbal formulas and single herbs prescribed by TCM doctors for the treatment of hypertension individuals among type 2 diabetes patients (Table A). Results of conditional multivariable logistic regression on the occurrence of acute myocardial infarction (Table B). Results of conditional multivariable logistic regression on the occurrence of ischemic stroke (Table C). Results of conditional multivariable logistic regression on the occurrence of hemorrhagic stroke (Table D). Results of conditional multivariable logistic regression on the occurrence of amputation (Table E). Results of conditional multivariable logistic regression on the occurrence of nephropathy (Table F). Results of conditional multivariable logistic regression on the occurrence of death (Table G). Regular medical treatment (from diabetes to index day) among type 2 diabetes patients according to CHM usage (Table H). Regular medical treatment (from index day to index day +365) among type 2 diabetes patients according to CHM usage (Table I). (DOCX)

Acknowledgments
The authors wish to thank the Division of Chinese Medicine, China Medical University Beigang Hospital for administrative assistance and consultation and the Aim for Top University Plan of the Ministry of Education, Taiwan at the China Medical University. We also thank Drs. Ya-Hui Chi, Kuan-Teh Jeang, Yuan-Chia Chang, and Willy W.L. Hong for technical help and suggestions.

Author Contributions
Conceived and designed the experiments: YJL TJH FJT WML. Performed the experiments: THL CCL SMH. Analyzed the data: CBW WKC JHC YCY YTS YCL CFC. Contributed reagents/materials/analysis tools: XL HT JPL CWL HYP JGL YHL SYC WML. Wrote the paper: YJL WML.
References

1. Geneva WHO. Definition, diagnosis and classification of diabetes mellitus and its complications. Part 1: Diagnosis and classification of diabetes mellitus. WHO/NCD/NCS/992. 1999.

2. Yang W, Lu J, Weng J, Jia W, Ji L, Xiao J, et al. Prevalence of diabetes among men and women in China. N Engl J Med. 2010; 362(12):1090–101. Epub 2010/03/26. doi: 10.1056/NEJMoa0908292 362/12/1090 [pii]. PMID: 20335585.

3. Saquib N, Saquib J, Ahmed T, Khanam MA, Cullen MR. Cardiovascular diseases and type 2 diabetes in Bangladesh: a systematic review and meta-analysis of studies between 1995 and 2010. BMC Public Health. 2012; 12:434. Epub 2012/06/15. doi: 10.1186/1471-2458-12-434 1471-2458-12-434 [pii]. PMID: 22694884; PubMed Central PMCID: PMC3487781.

4. Shera AS, Rafique G, Khawaja IA, Baqui S, King H. Pakistan National Diabetes Survey: prevalence of glucose intolerance and associated factors in Baluchistan province. Diabetes Res Clin Pract. 1999; 44 (1):49–58. Epub 1999/07/22. S0168-8227(99)00017-0 [pii]. PMID: 10414940.

5. Stumvoll M, Goldstein BJ, van Haeften TW. Type 2 diabetes: principles of pathogenesis and therapy. Lancet. 2005; 365(9467):1333–46. Epub 2005/04/13. S0140-6736(05)61032-X [pii] doi: 10.1016/S0140-6736(05)61032-X PMID: 15823385.

6. Kahn SE, Hull RL, Utzschneider KM. Mechanisms linking obesity to insulin resistance and type 2 diabetes. Nature. 2006; 444(7121):840–4. Epub 2006/12/15. nature05482 [pii] doi: 10.1038/nature05482 PMID: 17167471.

7. Wang B, Ni Y, Zhong J, Sun F. Effects of incretins on blood pressure: a promising therapy for type 2 diabetes with hypertension. J Diabetes. 2012; 4(1):22–9. Epub 2011/11/02. doi: 10.1111/j.1753-0407.2011.00167.x PMID: 22040104.

8. Lin K, Lloyd-Jones DM, Li D, Carr JC. Quantitative imaging biomarkers for the evaluation of cardiovascular complications in type 2 diabetes mellitus. J Diabetes Complications. 2014; 28(2):234–42. Epub 2013/12/07. S15517138(13)00211-0 [pii] doi: 10.1016/j.jdiacomp.2013.09.008 PMID: 24309215.

9. Lebovitz HE. Etiology and pathogenesis of diabetes mellitus. Pediatr Clin North Am. 1984; 31(3):521–30. Epub 1984/06/01. PMID: 6375587.

10. Intensive blood-glucose control with sulphonylureas or insulin compared with conventional treatment and risk of complications in patients with type 2 diabetes (UKPDS 33). UK Prospective Diabetes Study (UKPDS) Group. Lancet. 1998; 352(9131):857–65. Epub 1998/09/22. S0140673698070196 [pii] PMID: 9742976.

11. Tight blood pressure control and risk of macrovascular and microvascular complications in type 2 diabetes: UKPDS 38. UK Prospective Diabetes Study Group. BMJ. 1998; 317(7160):703–13. Epub 1998/09/11. PMID: 9732337; PubMed Central PMCID: PMC28659.

12. Mehler PS, Coll JR, Estacio R, Esler A, Schrier RW, Hiatt WR. Intensive blood pressure control reduces the risk of cardiovascular events in patients with peripheral arterial disease and type 2 diabetes. Circulation. 2003; 107(5):753–6. Epub 2003/02/13. PMID: 12578880.

13. Estacio RO, Jeffers BW, Gifford N, Schrier RW. Effect of blood pressure control on diabetic microvascular complications in patients with hypertension and type 2 diabetes. Diabetes Care. 2000; 23 Suppl 2: B54–6. Epub 2000/06/22. PMID: 10860192.

14. Li S, Culver B, Ren J. Benefit and risk of exercise on myocardial function in diabetes. Pharmacol Res. 2003; 48(2):127–32. Epub 2003/06/01. S1043661803000999 [pii] PMID: 12798664.

15. Rao AD, Kuhadiya N, Reynolds K, Fonseca VA. Is the combination of sulfonylureas and metformin associated with an increased risk of cardiovascular disease or all-cause mortality?: a meta-analysis of observational studies. Diabetes Care. 2008; 31(8):1672–8. Epub 2008/05/07. doi: 10.2337/dc08-0167 dc08-0167 [pii] PMID: 18458139; PubMed Central PMCID: PMC2494623.

16. Roumie CL, Hung AM, Greervy RA, Grijalva CG, Liu X, Murff HJ, et al. Comparative effectiveness of sulfonylurea and metformin monotherapy on cardiovascular events in type 2 diabetes mellitus: a cohort study. Ann Intern Med. 2012; 157(9):601–10. Epub 2012/11/07. doi: 10.7326/0003-4819-157-9-20121106-00003 1389945 [pii] PMID: 23128859.

17. Loke YK, Singh S, Furberg CD. Long-term use of thiazolidinediones and fractures in type 2 diabetes: a meta-analysis. CMAJ. 2009; 181(9):32–9. Epub 2008/12/17. doi: 10.1503/cmaj.080486 cmaj.080486 [pii] PMID: 19073651; PubMed Central PMCID: PMC261065.

18. Lincoff AM, Wolski K, Nicholls SJ, Nissen SE. Pioglitazone and risk of cardiovascular events in patients with type 2 diabetes mellitus: a meta-analysis of randomized trials. JAMA. 2007; 298(10):1180–8. Epub 2007/09/13. 298/10/1180 [pii] doi: 10.1001/jama.298.10.1180 PMID: 17848652.

19. Singh S, Loke YK, Furberg CD. Long-term use of thiazolidinediones and the associated risk of pneumonia or lower respiratory tract infection: systematic review and meta-analysis. Thorax. 2011; 66(5):383–8. Epub 2011/02/18. doi: 10.1136/thx.2010.152777 thx.2010.152777 [pii] PMID: 21325145.
20. Turner RM, Kwok CS, Chen-Turner C, Maduakor CA, Singh S, Loke YK. Thiazolidinediones and associated risk of Bladder Cancer: a Systematic Review and Meta-analysis. Br J Clin Pharmacol. 2013; 72(1):39–43. Epub 2012/10/12. doi: 10.1111/bcp.12306 PMID: 23425197.

21. Li GG, Kam A, Wong KH, Zhou X, Omar EA, Alqahtani A, et al. Herbal medicines for the management of diabetes. Adv Exp Med Biol. 2012; 771:396–413. Epub 2013/02/09. PMID: 23393692.

22. Ceylan-Isik AF, Fliethman RM, Wold LE, Ren J. Herbal and traditional Chinese medicine for the treatment of cardiovascular complications in diabetes mellitus. Curr Diabetes Rev. 2008; 4(4):320–3. Epub 2008/11/11. PMID: 18991600.

23. Li WL, Zheng HC, Bukuru J, De Kimpe N. Natural medicines used in the traditional Chinese medical system for therapy of diabetes mellitus. J Ethnopharmacol. 2004; 92(1):1–21. Epub 2004/04/22. doi: 10.1016/j.jep.2003.12.031 S0378874104000315 [pii]. PMID: 15099842.

24. Fang RC, Tsai YT, Lai JN, Yeh CH, Wu CT. The traditional Chinese medicine prescription pattern of diabetes mellitus patients in Taiwan: A population-based study. J Ethnopharmacol. 2012; 142(1):17–26. Epub 2012/04/10. doi: 10.1016/j.jep.2012.04.005 S0378-8741(12)00233-4 [pii]. PMID: 22577985; PubMed Central PMCID: PMC3502954.

25. Chen FP, Chen TJ, Kung YY, Chen YC, Chou LF, Chen FJ, et al. Use frequency of traditional Chinese medicine in Taiwan. BMC Health Serv Res. 2007; 7:26. Epub 2007/02/27. 1472-6963-7-26 [pii] doi: 10.1186/1472-6963-7-26 PMID: 17319950; PubMed Central PMCID: PMC1810531.

26. Huang TP, Liu PH, Lien AS, Yang SL, Chang HH, Yen HR. Characteristics of traditional Chinese medicine use in children with asthma: a nationwide population-based study. Allergy. 2013; 68(12):1610–3. Epub 2013/10/15. doi: 10.1111/all.12273 PMID: 24117783.

27. Lai JN, Wu CT, Wang JD. Prescription pattern of Chinese herbal products for breast cancer in Taiwan: a population-based study. Evid Based Complement Alternat Med. 2012; 2012:891893. Epub 2012/06/12. doi: 10.1155/2012/891893 PMID: 22685488; PubMed Central PMCID: PMC3368194.

28. Hsieh CF, Huang SL, Chen CL, Chen WT, Chang HC, Yang CC. Non-aristolochic acid prescribed Chinese herbal medicines and the risk of mortality in patients with chronic kidney disease: results from a population-based follow-up study. BMJ Open. 2014; 4(2):e004033. Epub 2014/02/25. doi: 10.1136/bmjopen-2013-004033 bmjopen-2013-004033 [pii]. PMID: 24581496; PubMed Central PMCID: PMC3991999.

29. Huang CY, Tsai YT, Lai JN, Hsu FL. Prescription pattern of Chinese herbal products for diabetes mellitus in Taiwan: a population-based study. Evid Based Complement Alternat Med. 2013; 2013:201329. Epub 2013/07/12. doi: 10.1155/2013/201329 PMID: 23643864; PubMed Central PMCID: PMC3703417.

30. Fang RC, Tsai YT, Lai JN, Yeh CH, Wu CT. The traditional Chinese medicine prescription pattern of endometriosis patients in Taiwan: a population-based study. Evid Based Complement Alternat Med. 2012; 2012:591391. Epub 2012/10/12. doi: 10.1155/2012/591391 PMID: 23056141; PubMed Central PMCID: PMC3463977.

31. Pan JC, Tsai YT, Lai JN, Fang RC, Yeh CH. The traditional Chinese medicine prescription pattern of patients with primary dysmenorrhea in Taiwan: a large-scale cross sectional survey. J Ethnopharmacol. 2014; 152(2):314–9. Epub 2014/01/15. doi: 10.1016/j.jep.2014.01.002 S0378-8741(14)00011-7 [pii]. PMID: 24417866.

32. Lin HC, Yang WC, Lee HC. Traditional Chinese medicine usage among schizophrenia patients. Complement Ther Med. 2008; 16(6):336–42. Epub 2008/11/26. doi: 10.1016/j.ctim.2007.11.001 S0965-2299(07)00113-6 [pii]. PMID: 19028334.

33. Yu MC, Lin SK, Lai JN, Wei JC, Cheng CY. The traditional Chinese medicine prescription patterns of Sjogrens patients in Taiwan: A population-based study. J Ethnopharmacol. 2014. Epub 2014/06/07. S0378-8741(14)00418-8 [pii] doi: 10.1016/j.jep.2014.05.049 PMID: 24905866.

34. Lin CH, Sheu WH. Hypoglycaemic episodes and risk of dementia in diabetes mellitus: 7-year follow-up study. J Intern Med. 2013; 273(1):102–10. Epub 2012/09/26. doi: 10.1111/j.1460-6732.2012.00116.x PMID: 23003116.

35. Bhadriraju K, Elliott JT, Nguyen M, Plant AL. Quantifying myosin light chain phosphorylation in single adherent cells with automated fluorescence microscopy. BMC Cell Biol. 2007; 8:43. Epub 2007/10/19. 1471-2121-8-43 [pii] doi: 10.1186/1471-2121-8-43 PMID: 17941977; PubMed Central PMCID: PMC2213650.

36. Uehata M, Ishizaki T, Satoh H, Ono T, Kawahara T, Morishita T, et al. Calcium sensitization of smooth muscle mediated by a Rho-associated protein kinase in hypertension. Nature. 1997; 389(6654):990–4. Epub 1997/11/14. doi: 10.1038/42017 PMID: 9353125.

37. Lian F, Li G, Chen X, Wang X, Piao C, Wang J, et al. Chinese herbal medicine Tianqi reduces progression from impaired glucose tolerance to diabetes: a double-blind, randomized, placebo-controlled, multicenter trial. J Clin Endocrinol Metab. 2014; 99(2):648–55. Epub 2014/01/18. doi: 10.1210/jc.2013-3276 PMID: 24432995.
38. Grant SJ, Bensoussan A, Chang D, Kiat H, Klupp NL, Liu JP, et al. Chinese herbal medicines for people with impaired glucose tolerance or impaired fasting blood glucose. Cochrane Database Syst Rev. 2009;(4):CD006690. Epub 2009/10/13. doi: 10.1002/14651858.CD006690.pub2 PMID: 19821382; PubMed Central PMCID: PMC3191296.

39. Hui H, Tang G, Go VL. Hypoglycemic herbs and their action mechanisms. Chin Med. 2009; 4:11. Epub 2009/06/16. doi: 10.1186/1749-8546-4-11 1749-8546-4-11 [pii]. PMID: 19523223; PubMed Central PMCID: PMC2704217.

40. Qian Y, Xue YM, Li J, Zhu B, Pan YH, Zhang Y. [Effect of Liuweidihuang pills in preventing diabetes mellitus in OLETF rats]. Nan Fang Yi Ke Da Xue Xue Bao. 2010; 30(1):21–4. Epub 2010/02/02. PMID: 20117976.

41. Xue YM, Luo R, Zhu B, Zhang Y, Pan YH, Li CZ. [Liuweidihuang pills reduces visceral fat deposition in Otsuka Long-Evans Tokushima Fatty rats]. Nan Fang Yi Ke Da Xue Xue Bao. 2006; 26(10):1446–8. Epub 2006/10/26. PMID: 17062348.

42. Qian Y, Xue YM, Li J. [Effects of Liuweidihuang pills on plasma adiponectin level in OLETF rats]. Nan Fang Yi Ke Da Xue Xue Bao. 2008; 28(1):34–6. Epub 2008/01/30. PMID: 18227021.

43. van Wietmarschen HA, van der Greef J, Schroen Y, Wang M. Evaluation of symptom, clinical chemistry and metabolomics profiles during Rehmannia six formula (R6) treatment: an integrated and personalized data analysis approach. J Ethnopharmacol. 2013; 150(3):851–9. Epub 2013/10/15. doi: 10.1016/j.jep.2013.09.041 S0378-8741(13)00688-0 [pii]. PMID: 24120517.

44. Wang J, Yao K, Yang X, Liu W, Feng B, Ma J, et al. Chinese patent medicine liu wei di huang wan combined with antihypertensive drugs, a new integrative medicine therapy, for the treatment of essential hypertension: a systematic review of randomized controlled trials. Evid Based Complement Alternat Med. 2012; 2012:714805. Epub 2012/12/22. doi: 10.1155/2012/714805 PMID: 23258998; PubMed Central PMCID: PMC3520441.

45. Qu Y, Gan HQ, Mei QB, Liu L. Study on the effect of Jia-Wei-Xiao-Yao-San decoction on patients with functional dyspepsia. Phytother Res. 2010; 24(2):245–8. Epub 2009/07/18. doi: 10.1002/ptr.2920 PMID: 19610026.

46. Chiao TH, Fu PK, Chang CH, Chang SN, Chiahung Mao F, Lin CH. Prescription patterns of Chinese herbal products for post-surgery colon cancer patients in Taiwan. J Ethnopharmacol. 2014; 155(1):702–8. Epub 2014/06/20. doi: 10.1016/j.jep.2014.06.012 S0378-8741(14)00460-7 [pii]. PMID: 24945402.

47. Wang BR, Chang YL, Chen TJ, Chiu JH, Wu JC, Wu MS, et al. Coprescription of Chinese herbal medicine Jiu Wei Di Huang Wan and Western medication among female patients with breast cancer in Taiwan: analysis of national insurance claims. Patient Prefer Adherence. 2014; 8:671–82. Epub 2014/05/24. doi: 10.2147/PPA.S61280 ppa-8-671 [pii]. PMID: 24855343; PubMed Central PMCID: PMC4065731.

48. Chen LC, Tsao YT, Yen KY, Chen YF, Chou MH, Lin MF. A pilot study comparing the clinical effects of Jia-Wey Shiau-Yau San, a traditional Chinese herbal prescription, and a continuous combined hormone replacement therapy in postmenopausal women with climacteric symptoms. Maturitas. 2003; 44(1):55–62. Epub 2003/02/06. S0378113902003146 [pii]. PMID: 12568736.

49. Chien SC, Chang WC, Lin PH, Chang WP, Hsu SC, Chang JC, et al. A Chinese herbal medicine, jia-wei-xiao-yao-san, prevents dimethylnitrosamine-induced hepatic fibrosis in rats. ScientificWorldJournal. 2014; 2014:217525. Epub 2014/07/06. doi: 10.1155/2014/217525 PMID: 24995353; PubMed Central PMCID: PMC40465731.

50. Zhang T, Xu J, Li D, Chen J, Shen X, Xu F, et al. Salvinolic acid A, a matrix metalloproteinase-9 inhibitor of Salvia miltiorrhiza, attenuates aortic aneurysm formation in apolipoprotein E-deficient mice. Phytomedicine. 2014; 21(10):1137–45. Epub 2014/06/12. doi: 10.1016/j.phymed.2014.05.003 S0944-7113(14)00224-4 [pii]. PMID: 24916705.

51. Jiang B, Li D, Deng Y, Teng F, Chen J, Xue S, et al. Salvinolic acid A, a novel matrix metalloproteinase-9 inhibitor, prevents cardiac remodeling in spontaneously hypertensive rats. PLoS One. 2013; 8(3):e59621. Epub 2013/03/28. doi: 10.1371/journal.pone.0059621 PONE-D-12-35832 [pii]. PMID: 23533637; PubMed Central PMCID: PMC3606118.

52. Woo KS, Yip TW, Chook P, Kwong SK, Szeto CC, Li JK, et al. Cardiovascular Protective Effects of Adjunctive Alternative Medicine (Salvia miltiorrhiza and Pueraria lobata) in High-Risk Hypertension. Evid Based Complement Alternat Med. 2013; 2013:132912. Epub 2013/03/28. doi: 10.1155/2013/132912 PMID: 23533460; PubMed Central PMCID: PMC3606734.

53. Hu F, Koon CM, Chan JY, Lau KM, Kwan YW, Fung KP. Involvements of calcium channel and potassium channel in Danshen and Gegen decoction induced vasodilation in porcine coronary LAD artery. Phytomedicine. 2012; 19(12):1051–8. Epub 2012/08/15. doi: 10.1016/j.phymed.2012.07.007 S0944-7113(12)00217-6 [pii]. PMID: 22989578.
54. Yang TY, Wei JC, Lee MY, Chen CM, Ueng KC. A randomized, double-blind, placebo-controlled study to evaluate the efficacy and tolerability of Fufang Danshen (Salvia miltiorrhiza) as add-on antihypertensive therapy in Taiwanese patients with uncontrolled hypertension. Phytother Res. 2012; 26(2):291–8. Epub 2011/09/03. doi: 10.1002/ptr.3548 PMID: 21887804.

55. Wang J, Lu W, Wang W, Zhang N, Wu H, Liu C, et al. Promising therapeutic effects of sodium tanshinone IIA sulfonate towards pulmonary arterial hypertension in patients. J Thorac Dis. 2013; 5(2):169–72. Epub 2013/04/16. doi: 10.3978/j.issn.2072-1439.2013.02.04 [pii]. PMID: 23585945; PubMed Central PMCID: PMC3621926.

56. Wang J, Jiang Q, Wan L, Yang K, Zhang Y, Chen Y, et al. Sodium tanshinone IIA sulfonate inhibits canonical transient receptor potential expression in pulmonary arterial smooth muscle from pulmonary hypertensive rats. Am J Respir Cell Mol Biol. 2013; 48(1):125–34. Epub 2012/10/10. doi: 10.1165/rcmb.2012-0071OC PMID: 23065131; PubMed Central PMCID: PMC3547081.

57. Huang M, Wang P, Xu S, Xu W, Chu K, Lu J. Biological activities of salvianolic acid B from Salvia miltiorrhiza on type 2 diabetes induced by high-fat diet and streptozotocin. Pharm Biol. 2015; 53(7):1058–65. Epub 2015/01/24. doi: 10.3109/13880209.2014.959611 PMID: 25612777.

58. Raoufi S, Baluchnejadmojarad T, Roghani M, Ghazanfari T, Khojasteh F, Mansouri M. Anti-diabetic potential of salvianolic acid B in multiple low-dose streptozotocin-induced diabetes. Pharm Biol. 2015; 53(12):1803–9. Epub 2015/04/18. doi: 10.3109/13880209.2015.1008148 PMID: 25885938.

59. Lian F, Wu L, Tian J, Jin M, Zhou S, Zhao M, et al. The effectiveness and safety of a danshen-containing Chinese herbal medicine for diabetic retinopathy: a randomized, double-blind, placebo-controlled multicenter clinical trial. J Ethnopharmacol. 2015; 164:71–7. Epub 2015/02/11. doi: 10.1016/j.jep.2015.01.048 S0378-8741(15)00063-X [pii]. PMID: 25666427.

60. Cai H, Liu L, Wang Y, Yu Y, Liu W. Protective effects of injection against learning and memory impairments in streptozotocin-induced diabetic rats. Exp Ther Med. 2014; 8(4):1127–30. Epub 2014/09/05. doi: 10.3892/etm.2014.1919 etm-08-04-1127 [pii]. PMID: 25187809; PubMed Central PMCID: PMC4151631.

61. Qicheng F. Some current studies and research approaches relating to the use of plants in the traditional Chinese medicine. J Ethnopharmacol. 1980; 2(1):57–63. Epub 1980/03/01. PMID: 7464185.

62. Teng Y, Cui H, Yang M, Song H, Zhang Q, Su Y, et al. Protective effect of puerarin on diabetic retinopathy in rats. Mol Biol Rep. 2009; 36(5):1129–33. Epub 2008/07/01. doi: 10.1007/s11033-008-9288-2 PMID: 18587665.

63. Koo M, Chen CH, Tsai KW, Lu MC, Lin SC. Ambulatory medical services utilization for menstrual disorders among female personnel of different medical professions in Taiwan: a nationwide retrospective cohort study. BMC Womens Health. 2015; 15(1):66. doi: 10.1186/s12905-015-0220-3 PMID: 26306618; PubMed Central PMCID: PMC4450071.

64. Chen MJ, Tsai YT, Liu JM, Lee YC, Wu MS, Chiu HM, et al. Statins and the risk of pancreatic cancer in Type 2 diabetic patients-A population-based cohort study. Int J Cancer. 2015. doi: 10.1002/ijc.29813 PMID: 26296262.

65. Wang SH, Chen DY, Lin YS, Mao CT, Tsai ML, Hsieh MJ, et al. Cardiovascular Outcomes of Sitagliptin in Type 2 Diabetic Patients with Acute Myocardial Infarction, a Population-Based Cohort Study in Taiwan. PLoS One. 2015; 10(6):e0131122. doi: 10.1371/journal.pone.0131122 PMID: 26115092; PubMed Central PMCID: PMC44382692.

66. Chang CH, Chen SJ, Liu CY. Risk of Developing Depressive Disorders following Hepatocellular Carcinoma: A Nationwide Population-Based Study. PLoS One. 2015; 10(8):e0135417. doi: 10.1371/journal.pone.0135417 PMID: 26295711; PubMed Central PMCID: PMC4546687.

67. Chen YC, Kok VC, Chien CH, Horng JT, Tsai JJ. Cancer risk in patients aged 30 years and above with type 2 diabetes receiving antidiabetic monotherapy: a cohort study using metformin as the comparator. Ther Clin Risk Manag. 2015; 11:1315–23. doi: 10.2147/TCRM.S91513 PMID: 26357479; PubMed Central PMCID: PMC4559233.

68. Wild S, Roglic G, Green A, Sicree R, King H. Global prevalence of diabetes: estimates for the year 2000 and projections for 2030. Diabetes Care. 2004; 27(5):1047–53. Epub 2004/04/28. PMID: 15111519.

69. Liu X, Feng R. Inhibition of epithelial to mesenchymal transition in metastatic breast carcinoma cells by c-Src suppression. Acta Biochim Biophys Sin (Shanghai). 2010; 42(7):496–501. Epub 2010/08/14. doi: 10.1093/abb/gmq043 gmq043 [pii]. PMID: 20705589.

70. Liu X, Feng R, Du L. The role of enoyl-CoA hydratase short chain 1 and peroxiredoxin 3 in PP2-induced apoptosis in human breast cancer MCF-7 cells. FEBS Lett. 2010; 584(14):3185–92. Epub 2010/06/15. doi: 10.1016/j.febslet.2010.06.002 S0014-5793(10)00483-7 [pii]. PMID: 20541551.