Research Article

Long-Term Traditional Chinese Medicine Combined with NA Antiviral Therapy on Cirrhosis Incidence in Chronic Hepatitis B Patients in the Real-World Setting: A Retrospective Study

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Received 25 February 2020; Revised 19 June 2020; Accepted 16 July 2020; Published 4 August 2020

Academic Editor: Maria G. Miguel

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Objective. To evaluate the impact of long-term Traditional Chinese Medicine (TCM) syndrome differentiation combined with antiviral therapy with Nucleos (t) ide analogues (NAs) on the incidence of cirrhosis in patients with chronic hepatitis B. Methods. This retrospective cohort study included 521 patients with chronic hepatitis B who underwent a treatment course of ≥3 years from 1998–2019. Of the 521 patients, 261 were defined as TCM users while 260 were TCM nonusers (control group). All the enrolled subjects were followed up until February 2019 to measure the incidence and hazard ratio (HR) of cirrhosis, and the Cox proportional hazards regression model was used to analyze the independent factors affecting the occurrence of cirrhosis. Results. The cumulative incidence of TCM users and nonusers was 6.9% and 13.5%, respectively (P = 0.013). Results of the Kaplan–Meier analysis demonstrated that TCM users had a significantly lower cumulative incidence of cirrhosis than TCM nonusers (P = 0.011), and TCM users had a significantly lower liver cirrhosis risk than TCM nonusers (adjusted HR = 0.416, 95% CI, 0.231–0.749). The histological evaluation revealed improved fibrosis in 45.0% of TCM users and 11.1% of TCM nonusers (P = 0.033). The analysis of the prescriptions including total 119 single Chinese herbs medicinal demonstrated that “replenish qi and fortify the spleen,” “clear heat and dispel dampness,” and “soothe the liver and regulate qi” are the main treatment methods of TCM for CHB. Conclusions. Our study demonstrated that long-term TCM use may attenuate liver cirrhosis risk in patients with chronic hepatitis B (CHB).

1. Introduction

Hepatitis B virus (HBV) infection is a public health concern worldwide, and persistent HBV infection can cause chronic hepatitis B (CHB), which is a serious threat to human health, leading to a huge social burden. According to the latest data in 2018, there were approximately 292 million hepatitis B carriers worldwide in 2016, and the rate of hepatitis B surface antigen positivity in China was 6.1%, indicating that there were approximately 81 million people with HBV infection [1]. HBV-related cirrhosis is a major clinical outcome of chronic HBV infection. According to the World Health
Organization (WHO) estimates, approximately 30% of patients worldwide who died of cirrhosis in 2002 were HBV-associated cirrhosis [2]. The annual incidence of cirrhosis in patients with CHB is 2–10% [3]. The 5-year cumulative cirrhosis rate of patients with chronic HBV infection is 8–20%.

Several guidelines for hepatitis B prevention emphasize that the long-term treatment goal of CHB is to continuously inhibit and eliminate HBV DNA and to prevent and delay complications such as cirrhosis and HCC [4–6]. The value and status of antiviral treatment for CHB has been widely recognized. Nucleos (t) ide analogues (NAs) and poly-pegylated interferon (PEG-IFN) can inhibit HBV replication, thereby reducing the pathological damage to the liver. However, the persistence of covalently closed circular DNA (cccDNA) in hepatocytes is the main cause of prolonged relapse and easy recurrence of CHB after treatment is discontinued [7]. Numerous clinical trials have demonstrated that although long-term NA therapy has been shown to improve long-term clinical outcomes, long-term antiviral therapy alone does not completely prevent the occurrence of cirrhosis and HCC and face the challenge of patients’ poor compliance.

Traditional Chinese medicine (TCM) is based on syndrome differentiation, and it emphasizes an individualized treatment and holistic view. According to the diagnosis and treatment guidelines of TCM for CHB, the main mechanism and syndrome type of CHB are liver-gallbladder dampness-heat, liver depression, and spleen deficiency [8, 9]. TCM can play complementary roles in antiviral treatment, facilitate disease recovery, and delay its progress. Thus far, there are no specific antifibrotic chemical or biological drugs that can be used in clinical practice, and Chinese medicine has clear advantages in this field [10]. The combination of traditional Chinese and Western medicine has gradually become the main way to treat CHB in China. This approach can delay the progress of CHB and improve clinical prognosis. However, no long-term follow-up observations have been performed to verify the efficacy of this approach.

Thus, the present study aimed to determine the impact of long-term TCM syndrome differentiation combined with NA antiviral therapy on the incidence of cirrhosis in CHB patients.

2. Methods

2.1. Study Subjects. This retrospective cohort study was generated from real-world practice using the registration and inpatient and outpatient datasets from 1998–2019 from the Shanghai Shuguang Hospital and the Shanghai Pudong New District Infectious Diseases Hospital, China. Data obtained for analysis included data from electronic medical records (EMR), laboratory information system (LIS), and picture archiving and communication systems (PACS), as well as the detailed procedures, medication, and TCM prescriptions from paper-based medical records.

This cohort study screened 1288 patients with CHB between 18 and 65 years of age. Of these, 767 patients were excluded from the study based on the following exclusion criteria: presence of cirrhosis, nucleoside (acid) antiviral therapy for <3 years or single antiviral therapy with interferon; concomitant nonalcoholic fatty liver disease or alcoholic liver disease, presence of other severe systemic diseases, presence of schistosomiasis, and other types of viral hepatitis infection. The remaining 521 patients were diagnosed with CHB and had been receiving continuous treatment with NA for more than three years either with or without TCM; these patients were deemed eligible and were included in the data analysis. Figure 1 presents the flowchart of patient recruitment for this study.

According to the different degrees of exposure to TCM syndrome differentiation in clinical practice, patients were divided into two cohorts based on whether they received a combination of Chinese and Western medicine or whether they received Western medicine alone. Patients who received Chinese herbal medicine for HBV infection-related diseases for more than 3 months per year for ≥3 years were designated as TCM users. Patients who were exposed to TCM for less than 3 months per year regardless of the total treatment course were designated as TCM nonusers.

2.2. Efficacy Measures. The primary outcome was the cumulative incidence of cirrhosis in patients with CHB. Cirrhosis was mainly defined by ultrasonography, computed tomography (CT), or magnetic resonance imaging (MRI) of the liver. The second outcome was histological response during follow-up, and a positive response was defined as an improvement in necroinflammation by at least one grade in the Scheuer system for necroinflammation or improvement in fibrosis by at least one grade according to the Scheuer fibrosis staging classification.

2.3. Statistical Analyses. Statistical data were described using means ± standard deviation, medians, or percentages. Differences in the baseline characteristics between the two groups were assessed by Student’s t-test or chi-square test. The cumulative incidence of cirrhosis was determined using the Kaplan–Meier algorithm, and the log-rank test was used to evaluate the difference between the two groups. The independent factors affecting the incidence of cirrhosis were analyzed by multivariate analysis using the LR forward method of the Cox proportional hazards regression model. Data were statistically analyzed using SPSS v21.0 (IBM, Armonk, NY, USA). A P value of <0.05 (two-sided) was considered statistically significant.

3. Results

Table 1 describes the baseline characteristics of study patients. There were no significant differences between the two groups. The follow-up duration of patients who received a combination of traditional Chinese and Western medicine ranged from 3–21 years with a median follow-up of 8 years; that of patients who received Western medicine alone was 3–19 years with a median follow-up of 8 years (P > 0.05). The antiviral treatment duration with antiviral therapy was 3.1–20.6 years (median, 6.6 years) in the cohort that received...
Chinese and Western medicine and 3–18 years (median, 7 years) in the cohort that received Western medicine alone ($P > 0.05$).

A total of 53 liver cirrhosis events (18 in TCM users and 35 in TCM nonusers) occurred during the follow-up period. The cumulative incidence of cirrhosis was 6.9% in TCM users versus 13.5% in TCM nonusers ($\chi^2 = 6.143, P = 0.013$). Kaplan–Meier analysis demonstrated that TCM users had a lower cumulative incidence of cirrhosis than TCM nonusers ($\chi^2$ test $= 6.534, P = 0.011$; Figure 2). Table 2 illustrates the incidence of cirrhosis among patients with different baseline characteristics. Figure 3 illustrates the results of multivariable predictor of liver cirrhosis incidence with respect to the TCM utilization status along with the baseline characteristics in Tables 1 and 2. TCM users had a significantly lower liver cirrhosis risk than TCM nonusers (adjusted HR $= 0.416$, 95% CI, 0.231–0.749).

A total of 144 patients underwent liver biopsy at baseline (71 and 74 in the TCM user and nonuser groups, respectively). During the follow-up, 38 patients underwent paired liver biopsies (20 and 18 in the TCM user and nonuser groups, respectively). The results of the histological evaluation (Figure 4) revealed that 35.0% of TCM users and 22.2% of TCM nonusers achieved improvements in necroinflammation ($P = 0.627$) and 45.0% of TCM users and 11.1% of TCM nonusers achieved improvements in fibrosis symptoms ($P = 0.033$).

A total of 119 single Chinese herbs were collected in all the TCM prescriptions for CHB patients. The names and major functions of the mostly used single Chinese herbs are shown in Table 3, which were mainly heat-clearing medicinal, dampness-resolving medicinal, spleen-fortifying medicinal, and qi-regulating medicinal. It was demonstrated that “replenish qi and fortify the spleen,” “clear heat and dispel dampness,” and “soothe the liver and regulate qi” are the main treatment methods of TCM for CHB.

**4. Discussion**

CHB is an immune-mediated disease, and the interaction of the HBV with innate and acquired immune responses determines the clinical outcome of HBV infection. There are three obstacles for curing CHB: the persistence of the virus cccDNA, dysfunction of the immune response to HBV, and pathological changes in liver tissue (inflammation, necrosis, fibrosis, and cirrhosis). The development of HBV infection from chronic hepatitis to cirrhosis and even to liver cancer is a multistage pathogenic process and is affected by many factors, including virus (such as the HBV genotype, viral mutation, viral load, and viral proteins levels), host (such as age, gender, genetic polymorphism, presence or absence of other viral infections, and intestinal microecology), and other exogenous factors, and also whether or not patients receive antiviral therapy in a timely and standardized manner [11]. The HBV DNA load, serum HBsAg level, HBeAg statue, and ALT levels have been shown to be closely related to the clinical outcome of CHB [12].

Antiviral therapy is the key to preventing disease progression and improving the prognosis of patients with CHB. A number of studies have shown that long-term antiviral therapy can substantially improve the degree of liver fibrosis in patients with CHB. In a long-term follow-up study (mean follow-up duration 89.9 months) of CHB patients, the incidence of cirrhosis and HCC significantly reduced in CHB.
patients who received lamivudine compared with patients who did not receive antiviral therapy [13]. It was reported that, among the CHB patients treated with entecavir for 5 years, 88% showed improvement in liver fibrosis and 40% achieved cirrhosis reversal [14]. A 5-year open-label follow-up study of tenofovir in patients with CHB revealed that long-term inhibition of HBV can improve histology and reverse liver fibrosis and cirrhosis [15]. However, long-term use of NA antiviral therapy has gradually exposed problems such as drug resistance, viral relapse, and adverse reactions. Some studies also showed that even with active antiviral therapy, patients have varying degrees of liver histological lesions [16–19], and the long-term prognosis of CHB patients with nonantiviral indications is not optimistic.

TCM has a long history of treating CHB. TCM has over time formed a unique diagnosis and is associated with treatment advantages in long-term clinical practice. TCM plays an important role in regulating immune function, improving liver function, exerting antifibrosis and antiviral effects, and eliminating the clinical signs and symptoms of

| Characteristics                        | TCM users (n = 261) | TCM nonusers (n = 260) | P value |
|----------------------------------------|--------------------|------------------------|---------|
| Age (years)                            |                    |                        |         |
| <40                                    | 164 (62.8)         | 152 (58.5)             | 0.307   |
| ≥40                                    | 97 (37.2)          | 108 (41.9)             |         |
| Gender (n, %)                          |                    |                        |         |
| Male                                   | 167 (64.0)         | 171 (65.8)             | 0.670   |
| Female                                 | 94 (36.0)          | 89 (34.2)              |         |
| Duration of illness (n, %)             |                    |                        |         |
| 0–5                                    | 62 (23.8)          | 70 (26.9)              |         |
| 5–10                                   | 55 (21.2)          | 48 (18.5)              | 0.757   |
| 10–15                                  | 59 (22.6)          | 62 (23.8)              |         |
| 15–                                   | 85 (32.6)          | 80 (30.8)              |         |
| Smoking history (n, %)                 |                    |                        |         |
| Yes                                    | 33 (12.6)          | 28 (10.8)              | 0.506   |
| No                                     | 228 (87.4)         | 232 (89.2)             |         |
| Drinking history (n, %)                |                    |                        |         |
| Yes                                    | 45 (17.2)          | 39 (15.0)              | 0.487   |
| No                                     | 216 (82.8)         | 221 (85.0)             |         |
| Family history of CHB-related diseases  |                    |                        |         |
| Yes                                    | 96 (36.8)          | 101 (39.8)             | 0.549   |
| No                                     | 165 (63.2)         | 159 (61.2)             |         |
| HBeAg                                  |                    |                        |         |
| Positive                               | 153 (58.6)         | 156 (60.0)             | 0.749   |
| Negative                               | 108 (41.4)         | 104 (40.0)             |         |
| HBV DNA level (copies/mL)              |                    |                        |         |
| $10^3$–$10^4$                          | 39 (14.9)          | 35 (13.5)              |         |
| $10^4$–$10^5$                          | 31 (11.9)          | 29 (11.2)              |         |
| $10^5$–$10^6$                          | 34 (13.0)          | 46 (17.7)              | 0.644   |
| $10^6$–$10^7$                          | 45 (17.2)          | 49 (18.8)              |         |
| $10^7$                                 | 112 (42.9)         | 101 (38.8)             |         |
| ALT (IU/ml ULN)                        |                    |                        |         |
| 1–2                                    | 48 (18.4)          | 54 (20.8)              | 0.714   |
| 2–3                                    | 137 (52.5)         | 129 (49.6)             |         |
| 3–                                    | 76 (29.1)          | 77 (29.6)              |         |

**Figure 2:** Cumulative incidence of cirrhosis in CHB patients with and without TCM syndrome differentiation. The incidence of liver cirrhosis was significantly lower ($P = 0.011$) in TCM users (n = 261) than in TCM nonusers (n = 260).
A systematic review of TCM with NA therapy for treating CHB revealed that it is better than NA therapy alone in terms of improving liver function, HBeAg seroconversion rate, HBV DNA negative conversion rate, and the incidence of YMDD mutation [20]. According to TCM theory, hepatitis B-related cirrhosis locations include the liver, spleen, and kidney, and the pathological products of hepatitis B-related cirrhosis include blood stasis, water, stagnation of qi, and dampness of phlegm. The basic pathogenesis is qi deficiency and blood stasis and phlegm.

**Table 2: Incidence of cirrhosis for CHB patients with different baseline characteristics.**

| Characteristics | Cirrhosis (n = 53) | Noncirrhosis (n = 468) | P value |
|-----------------|--------------------|------------------------|---------|
| Age (years)     |                    |                        |         |
| <40             | 32 (10.1)          | 284 (89.9)             | 0.965   |
| ≥40             | 21 (10.2)          | 184 (89.8)             |         |
| Gender (n, %)   |                    |                        |         |
| Male            | 42 (12.4)          | 296 (65.8)             | 0.021   |
| Female          | 11 (6.0)           | 172 (94.0)             |         |
| Duration of illness (n, %) |        |                        |         |
| 0~5             | 5 (3.8)            | 127 (96.2)             | 0.158   |
| 5~10            | 9 (8.7)            | 94 (91.3)              |         |
| 10~15           | 14 (11.6)          | 107 (88.4)             |         |
| ≥15             | 25 (15.2)          | 140 (84.8)             |         |
| Smoking history (n, %) |            |                        |         |
| Yes             | 8 (12.6)           | 53 (10.8)              | 0.419   |
| No              | 45 (87.4)          | 415 (89.2)             |         |
| Drinking history (n, %) |            |                        |         |
| Yes             | 14 (16.7)          | 70 (83.3)              | 0.032   |
| No              | 39 (82.8)          | 398 (91.1)             |         |
| Family history of CHB-related diseases (n, %) | |                      |         |
| Yes             | 27 (13.7)          | 170 (86.3)             | 0.016   |
| No              | 26 (8.0)           | 298 (92.0)             |         |
| HBeAg Positive  | 39 (12.5)          | 270 (87.5)             | 0.027   |
| Negative        | 14 (6.6)           | 198 (93.4)             |         |
| HBV DNA level (copies/mL) |        |                        |         |
| 10^3~10^4       | 3 (4.1)            | 71 (95.9)              | 0.022   |
| 10^4~10^5       | 4 (3.3)            | 56 (96.7)              |         |
| 10^5~10^6       | 6 (7.5)            | 74 (92.5)              |         |
| 10^6~10^7       | 11 (11.7)          | 83 (88.3)              |         |
| 10^7            | 29 (14.6)          | 184 (85.4)             |         |
| ALT (IU/ml ULN) |                    |                        |         |
| 1~2             | 8 (7.8)            | 94 (92.2)              | 0.473   |
| 2~3             | 26 (9.8)           | 240 (90.2)             |         |
| 3~               | 19 (12.4)          | 134 (87.6)             |         |

**Figure 3:** Multivariable predictor of liver cirrhosis. Results of the multivariable predictor of liver cirrhosis according to TCM utilization status along with the baseline characteristics, as shown in Tables 1 and 2.
Figure 4: Histological response of CHB patients with and without TCM use. A total of 38 patients had paired liver biopsy (20 and 18 in TCM users and nonusers, respectively). Histological evaluation showed that 45.0% of TCM users and 11.1% of TCM nonusers achieved improved fibrosis ($P < 0.05$) (a) and 35.0% of TCM users and 22.2% of TCM nonusers achieved improvement in necroinflammation ($P > 0.05$) (b).

Table 3: The names and major functions of single Chinese herb medicinal mostly used in TCM prescriptions for CHB patients.

| Chinese herb medicinal | Major functions | No. of users | %
|------------------------|----------------|-------------|--------------------------|
| Atractylodes macrocephala Koidz. (Bai Shu) | Fortify the spleen and replenish qi, dry dampness, and induce diuresis | 180 | 68.97 |
| Atractylodes lancea (Thunb.) DC. (Cang Shu) | Dry dampness and fortify the spleen, dispel wind, and disperse cold | 153 | 58.62 |
| Corydalis yanhusuo W. T. Wang ex Z. Y. Su et C.Y. (Yan Hu Suo) | Active blood and resolve stasis and move qi to relieve pain | 129 | 49.43 |
| Artemisia capillaris Thunb. (Yin Chen) | Induce diuresis to dry dampness, fortify spleen, and tranquilize heart | 94 | 36.02 |
| Gardenia jasminoides Ellis (Zhi Zi) | Purge fire and except vexed, clear heat and drain dampness, and cool blood and detoxify | 82 | 31.42 |
| Serissa japonica (EU_hunb.) EU_hunb. (Liu Yue Xue) | Disperse wind and release the exterior, clear heat and drain dampness, stimulate the circulation of the blood, and cause the muscles and joints to relax | 79 | 30.72 |
| Cleistocactus sepium (Wu Zei Gu) | Stop bleeding with astringency, inhibit acidity to relieve pain, drain dampness, and disperse abscesses | 72 | 27.59 |
| Pinellia ternata (Ban Xia) | Dry dampness to resolve phlegm, direct qi downward to relieve emesis, and disperse stuffiness and nodules | 67 | 25.67 |
| Scutellaria barbata D. Don (Ban Zhi Lian) | Clear heat and detoxify and resolve stasis to induce diuresis | 64 | 24.52 |
| Spreading Hedysotis Herb (She She Cao) | Clear heat and drain dampness and detoxify | 57 | 21.84 |
| Citrus reticulata Blanco (Qing Pi) | Soothe the liver and regulate qi and disperse accumulation and stagnation | 55 | 21.07 |
| Citrus reticulata Blanco (Chen Pi) | Regulate qi and fortify the spleen and dry dampness to resolve phlegm | 45 | 17.24 |
| Lysimachia christinae Hance (Jin Qian Cao) | Drain dampness and anti-icteric and induce diuresis to relieve strangury | 45 | 17.24 |
| Bupleurum chinense (Chai Hu) | Harmonize and release the exterior and interior, soothe the liver to relieve depression, raise yang and lift prolapsed organs, and interrupt malaria | 44 | 16.86 |
| Glycyrrhiza uralensis Fisch (Gan Cao) | Fortify the spleen and replenish qi, clear heat and detoxify, dispel phlegm to suppress cough, relieve spasm and pain, and moderate herbs | 43 | 16.48 |
| Setaria italica (L.) Beauv. (Gu Ya) | Promote digestion, invigorate the stomach, and increase the appetite | 41 | 15.71 |
| Plantago depressa Willd. (Che Qian Cao) | Clear heat, induce diuresis, dispel phlegm, and cool blood and detoxify | 40 | 15.33 |
Infectious Diseases Hospital (Grade II Level A). Further (Grade III Level A) and the Shanghai Pudong New District study only operated in the Shanghai Shuguang Hospital each herb. Finally, due to the limited access to data, our let alone identifying the specific effective components in even determine a common prescription that is effective, a small proportion of each prescription, and it is difficult to a long-term follow-up period. EU_his could result in using very and one patient could visit different physicians during the physician may prescribe different prescriptions at each visit, primarily on the physicians' individual experiences: one cine; thus, the prevalence of TCM may be underestimated. medical records to reduce this bias as much as possible.

and outpatient claims datasets, as well as paper-based be incomplete and there could be biases in data collection ficiencies of a retrospective study, patient information may must be included in further studies. Second, given the de- tion. As these factors are more likely to affect TCM use, they bias due to unrecognized confounding factors such as household income, educational background, and occupa- As these factors are more likely to affect TCM use, they must be included in further studies. Second, given the de- ficiencies of a retrospective study, patient information may be incomplete and there could be biases in data collection and on follow-up; thus, we used registration and inpatient and outpatient claims datasets, as well as paper-based medical records to reduce this bias as much as possible. Third, this study did not consider over-the-counter TCM use for herbal medicines and traditional Chinese patent medicine; thus, the prevalence of TCM may be underestimated. Fourth, syndrome differentiation according to TCM is based primarily on the physicians' individual experiences: one physician may prescribe different prescriptions at each visit, and one patient could visit different physicians during the long-term follow-up period. This could result in using very small proportion of each prescription, and it is difficult to even determine a common prescription that is effective, let alone identifying the specific effective components in each herb. Finally, due to the limited access to data, our study only operated in the Shanghai Shuguang Hospital (Grade III Level A) and the Shanghai Pudong New District Infectious Diseases Hospital (Grade II Level A). Further multicenter studies need to be carried out to verify the effect of TCM.

5. Conclusions
In conclusion, this retrospective study demonstrated that the long-term use of syndrome differentiation according to TCM could attenuate the liver cirrhosis risk in patients with CHB. The findings of this study should be further validated by randomized controlled trails.

Abbreviations

| Abbreviation | Description |
|--------------|-------------|
| ALT | Alanine aminotransferase |
| CHB | Chronic hepatitis B |
| CT | Computed tomography |
| CCCDNA | Covalently closed circular DNA |
| EMR | Electronic medical record |
| HBV | Hepatitis B virus |
| HBeAg | Hepatitis B e antigen |
| HBsAg | Hepatitis B surface antigen |
| HCC | Hepatocellular carcinoma |
| HR | Hazard ratios |
| LIS | Laboratory information system |
| MRI | Magnetic resonance imaging |
| NAs | Nucleos (t) ide analogues |
| PACS | Picture archiving and communication systems |
| PEG-IFN | Poly-pegylated interferon |
| TCM | Traditional Chinese medicine |
| WHO | World Health Organization |

Data Availability

The datasets used and analyzed during this study are available from the corresponding authors upon reasonable request.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

Authors' Contributions

Zhi-Jun Hou performed the study and collected the data. Jing-Hao Zhang and Xin Zhang analyzed the data and wrote the manuscript. Chao Zheng and Xiao-jun Zhu collected and assembled the data. Qi-hua Ling and Zhen-hua Zhou supervised the study and helped design the study. Xiao-ni Kong and Man Li proofread the manuscript. Jian-jie Chen, Yue-Qiu Gao, and Xue-Hua Sun conceived and designed the study. Zhi-Jun Hou, Jing-Hao Zhang, and Xin Zhang contributed equally to this work.

Acknowledgments

This work was supported by the National Natural Science Foundation of China (81673938, 81673935, and 81774256), training plan of outstanding young medical talents, Shanghai Municipal Health Bureau (2018YQ43), the Three-
References

[1] Polaris Observatory Collaborators, “Global prevalence, treatment, and prevention of hepatitis B virus infection in 2016: a modelling study,” *The Lancet Gastroenterology and Hepatology*, vol. 3, no. 6, pp. 383–403, 2018.

[2] J. F. Perz, G. L. Armstrong, L. A. Farrington, Y. J. F. Hutin, and B. P. Bell, “The contributions of hepatitis B virus and hepatitis C virus infections to cirrhosis and primary liver cancer worldwide,” *Journal of Hepatology*, vol. 45, no. 4, pp. 529–538, 2006.

[3] G. Fattovich, F. Bortolotti, and F. Donato, “Natural history of chronic hepatitis B: special emphasis on disease progression and prognostic factors,” *Journal of Hepatology*, vol. 48, no. 2, pp. 335–352, 2008.

[4] J. Hou, G. Wang, F. Wang et al., “Guideline of prevention and treatment for chronic hepatitis B (2015 update),” *Journal of Clinical and Translational Hepatology*, vol. 5, no. 4, pp. 297–318, 2017.

[5] European Association for the Study of the Liver, P. Lampertico, K. Agarwal et al., “EASL 2017 Clinical Practice Guidelines on the management of hepatitis B virus infection,” *Journal of Hepatology*, vol. 67, no. 2, pp. 370–398, 2017.

[6] N. A. Terrault, A. S. F. Lok, B. J. McMahon et al., “Update on prevention, diagnosis, and treatment of chronic hepatitis B: AASLD 2018 hepatitis B guidance,” *Hepatology*, vol. 67, 2018.

[7] P. Revill and S. Locarnini, “Antiviral strategies to eliminate hepatitis B virus covalently closed circular DNA (cccDNA),” *Current Opinion in Pharmacology*, vol. 30, pp. 144–150, 2016.

[8] Hepatobiliary Disease Group, Internal Medicine of Traditional Chinese, and China Association of Traditional Chinese Medicine, “Expert consensus of hepatology, World federation of Chinese medicine societies; hepatology group, Chinese association of the integration of traditional and western medicine. Guidelines for traditional Chinese medical diagnosis of chronic hepatitis B,” *Journal of Clinical and Experimental Hepatology*, vol. 28, no. 03, pp. 164–168, 2012, in Chinese.

[9] Hepatobiliary Specialized Committee of China Association of Chinese Medicine and Liver Diseases Specialized Committee of China Medical Association of Minorities, “The clinical guidelines of diagnosis and treatment of chronic hepatitis B with traditional Chinese medicine (2018),” *Journal of Clinical and Experimental Hepatology*, vol. 34, no. 12, pp. 2520–2525, 2018, in Chinese.

[10] Liver Disease Committee, Chinese Association of Integrative Medicine, “Guidelines for the diagnosis and treatment of liver fibrosis in integrative medicine practice (2019),” *Journal of Clinical and Experimental Hepatology*, vol. 35, no. 7, pp. 1444–1449, 2019, in Chinese.

[11] Y.-F. Liaw, “Natural history of chronic hepatitis B virus infection and long-term outcome under treatment,” *Liver International*, vol. 29, no. 1, pp. 100–107, 2009.

[12] C.-J. Liu and J.-H. Kao, “Global perspective on the natural history of chronic hepatitis B: role of hepatitis B virus genotypes A to J,” *Seminars in Liver Disease*, vol. 33, no. 2, pp. 097–102, 2013.

[13] M. F. Yuen, W. K. Seto, D. H. Chow et al., “Long-term lamivudine therapy reduces the risk of long-term complications of chronic hepatitis B infection even in patients without advanced disease,” *Antiviral Therapy*, vol. 12, no. 8, pp. 1295–1303, 2007.

[14] A. S.-F. Lok, “Long-term therapy of chronic hepatitis B reverses cirrhosis,” *Nature Reviews Gastroenterology & Hepatology*, vol. 10, no. 4, pp. 199–200, 2013.

[15] P. Marcellin, E. Gane, M. Buti et al., “Regression of cirrhosis during treatment with tenofovir disoproxil fumarate for chronic hepatitis B: a 5-year open-label follow-up study,” *The Lancet*, vol. 381, no. 9865, pp. 468–475, 2013.

[16] Y. L. Zhang and J. W. Zhou, “Liver histopathological features of mildly abnormal ALT in patients with CHB,” *Chinese Journal of Health Laboratory Technology*, vol. 21, no. 1, pp. 26–27, 2011, in Chinese.

[17] J. J. Liu, B. Ding, Z. H. Xu et al., “Histological characters and clinical relationship between chronic HBV and inactive HBsAg carriers,” *Shenzhen Journal of Integrated Traditional Chinese and Western Medicine*, vol. 19, no. 4, pp. 216–219, 2009, in Chinese.

[18] X. H. Gong, F. Z. Wang, H. Li et al., “Investigation on transformation of 197 HBsAg carriers,” *Chinese Journal of Public Health*, vol. 22, no. 10, pp. 1251–1253, 2006, in Chinese.

[19] W. Y. Deng, “Analysis of the relationship between age and outcome in 654 patients infected with HBV,” *Chinese Journal of Integrated Traditional and Western Medicine on Liver Disease*, no. S1, p. 122, 2003, in Chinese.

[20] S. Ren, H. Zhang, X. C. Hu et al., “Meta-analysis of the clinical efficacy of combination therapy with traditional Chinese medicine and nucleoside analogue antiviral drugs for chronic hepatitis B,” *Journal of Clinical and Experimental Hepatology*, vol. 29, no. 4, pp. 264–269+275, 2013, in Chinese.

[21] P. Liu, Y. Y. Hu, C. Liu et al., “Multicenter clinical study on Fushehuayu capsule against liver fibrosis due to chronic hepatitis B,” *World Journal of Gastroenterology*, vol. 11, no. 19, pp. 2892–2899, 2005.

[22] F.-R. Yang, B. W. Fang, and J. S. Lou, “Effects of Fuwanga pill on hepatic fibrosis in vivo and vitro,” *World Journal of Gastroenterology*, vol. 19, no. 32, pp. 5326–5333, 2013.

[23] H. B. Cai, X. G. Sun, and F. Liu et al., “Effects of dahuanghechang pills on cytokines and mitogen activated protein kinase activation in rats with hepatic fibrosis,” *Journal of Ethnopharmacology*, vol. 132, no. 1, pp. 157–164, 2010.

[24] L. Miao, W. N. Yang, X. Q. Dong et al., “Combined anluoxuangan and entecavir treatment significantly improve the improvement rate of liver fibrosis in patients with chronic hepatitis B virus infection,” *Zhonghua Gan Zang Bing Za Zhi*, vol. 27, no. 7, pp. 521–526, 2019, in Chinese.

[25] L. Yao, Z. M. Yao, H. Weng et al., “Effect of rat serum containing Biejiajian oral liquid on proliferation of rat hepatic stellate cells,” *World Journal of Gastroenterology*, vol. 10, no. 13, pp. 1911–1913, 2004.