**Name of journal:** *World Journal of Gastroenterology*

**ESPS Manuscript NO: 28654**

**Manuscript Type:** **ORIGINAL ARTICLE**

***Clinical Trials Study***

**Phase IIb trial of *in vivo* electroporation mediated dual-plasmid hepatitis B virus DNA vaccine in chronic hepatitis B patients under lamivudine chemotherapy**

Yang FQ *et al.* Phase IIb trial of therapeutic HBV DNA vaccine

Fu-Qiang Yang, Gui-Rong Rao, Gui-Qiang Wang, Yue-Qi Li, Yao Xie, Zhan-Qing Zhang, Cun-Liang Deng, Qing Mao, Jun Li, Wei Zhao, Mao-Rong Wang, Tao Han, Shi-Jun Chen, Chen Pan, De-Ming Tan, Jia Shang, Ming-Xiang Zhang, Yue-Xin Zhang, Ji-Ming Yang, Guang-Ming Chen

**Fu-Qiang Yang, Gui-Rong Rao,** **Guang-Ming Chen,** Liver Disease Research Center, Guangzhou 458 Hospital, Guangzhou 510062, Guangdong Province, China

**Gui-Qiang Wang,** Department of Infectious Diseases, Peking University First Hospital, Beijing 100034, China

**Yue-Qi Li,** Beijing 302 Hospital, Beijing 100039, China

**Yao Xie,** Ninth Clinical Department, Beijing Ditan Hospital, Beijing 100011, China

**Zhan-Qing Zhang,** Liver Diseases Department of Public Health Center, Fudan University, Shanghai 201508, China

**Cun-Liang Deng,** Department of Infectious Diseases, Affiliated Hospital of Luzhou Medical College, Luzhou 646000, Sichuan Province, China

**Qing Mao,** PLA Institute of Infectious Diseases, Southwest Hospital, Third Military Medical University, Chongqing 400038, China

**Jun Li,** Jiangsu Province People's Hospital, Nanjing 210029, Jiangsu Province, China

**Wei Zhao,** Nanjing 2nd Hospital, Nanjing 210003, Jiangsu Province, China

**Mao-Rong Wang,** Nanjing 81 Hospital, Nanjing 210002, Jiangsu Province, China

**Tao Han,** Tianjin 3st Center Hospital, Tianjin 300170, China

**Shi-Jun Chen,** Jinan Infectious Desease Hospital, Jinan 250000, Shandong Province, China

**Chen Pan,** Fuzhou Infectious Desease Hospital, Fuzhou 350025, Fujian Province, China

**De-Ming Tan,** Xiangya Hospital, Central South University, Changsha 410008, Hunan Province, China

**Jia Shang,** Henan Province People's Hospital, Zhengzhou 450003, Henan Province, China

**Ming-Xiang Zhang,** The Sixth People's Hospital of Shenyang, Shenyang 110006, Liaoning Province, China

**Yue-Xin Zhang,** The First Affiliated Hospital of Xinjiang Medical University, Urumqi 830054, [Xinjiang](http://cn.bing.com/dict/clientsearch?mkt=zh-CN&setLang=zh&form=BDVEHC&ClientVer=BDDTV3.5.0.4311&q=%E6%96%B0%E7%96%86%E7%BB%B4%E5%90%BE%E5%B0%94%E8%87%AA%E6%B2%BB%E5%8C%BA) [Uygur](http://cn.bing.com/dict/clientsearch?mkt=zh-CN&setLang=zh&form=BDVEHC&ClientVer=BDDTV3.5.0.4311&q=%E6%96%B0%E7%96%86%E7%BB%B4%E5%90%BE%E5%B0%94%E8%87%AA%E6%B2%BB%E5%8C%BA) [Autonomous](http://cn.bing.com/dict/clientsearch?mkt=zh-CN&setLang=zh&form=BDVEHC&ClientVer=BDDTV3.5.0.4311&q=%E6%96%B0%E7%96%86%E7%BB%B4%E5%90%BE%E5%B0%94%E8%87%AA%E6%B2%BB%E5%8C%BA) [Region](http://cn.bing.com/dict/clientsearch?mkt=zh-CN&setLang=zh&form=BDVEHC&ClientVer=BDDTV3.5.0.4311&q=%E6%96%B0%E7%96%86%E7%BB%B4%E5%90%BE%E5%B0%94%E8%87%AA%E6%B2%BB%E5%8C%BA), China

**Ji-Ming Yang,** Infection Disease hospital of TianJin, Tianjin 300101, China

**Author contributions:** Yang FQ, Rao GR, Wang GQ and Chen GM contributed to conception and design; Wang GQ is the leader of multi-center and PI of the clinical research; Yang FQ, Rao GR, Wang GQ, Li YQ, Xie Y, Zhang ZQ, Deng CL, Mao Q, Li J, Zhao W, Wang MR, Han T, Chen SJ, Pan C, Tan DM, Shang J, Zhang MX, Zhang YX, Yang JM and Chen GM contributed to acquisition of data, or analysis and interpretation of data; Yang FQ, Rao GR, Wang GQ, Li YQ, Xie Y, Zhang ZQ, Deng CL, Mao Q, Li J, Zhao W, Wang MR, Han T, Chen SJ, Pan C, Tan DM, Shang J, Zhang MX, Zhang YX, Yang JM and Chen GM have been involved in drafting the manuscript or revising it critically for important intellectual content; all authors have given final approval of the version to be published.

**Supported by** Yigan Biological Products Co., Ltd. of Guangzhou Pharmaceutical Holdings Ltd. (GPC, Guangzhou, China); It was also funded by Guangdong Provincial Sci. & Tech. Project, No. 2012A080204009; Guangdong Provincial Natural Science Fund, No. 2014A030313770; Guangdong Provincial Public Benefit Foundation, No. 2015A010107011; and National Key Program Grant for Management of AIDS and Viral Hepatitis in China 11th 5-Year Plan, No. 2008ZX10002-003.

**Institutional review board statement:** The study was reviewed and approved by the ethics committee of the Peking University First Hospital, as well as by the committees of each of the participating centers.

**Clinical trial registration statement:** This trial was registered with ClinicalTrials.org (NCT01487876).

**Informed consent statement:** All study participants, or their legal guardian, provided informed written consent prior to study enrollment.

**Conflict-of-interest statement:** Not declared.

**Data sharing statement:** No additional data are available.

**Open-Access:** This article is an open-access article which was selected by an in-house editor and fully peer-reviewed by external reviewers. It is distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>

**Manuscript source:** Unsolicited manuscript

**Correspondence to: Guang-Ming Chen, MD,** Liver Disease Research Center, Guangzhou 458 Hospital, Guangzhou 510062, Guangdong Province, China. chgm0616@sina.com

**Telephone:** +86-20-87376240

**Fax:** +86-21-64085875

**Received:** July 22, 2016

**Peer-review started:** July 22, 2016

**First decision:** September 20, 2016

**Revised:** September 30, 2016

**Accepted:** October 27, 2016

**Article in press:**

**Published online:**

**Abstract**

***AIM***

To assess the efficacy and safety of *in vivo* electroporation (EP)-mediated dual-plasmid hepatitis B virus (HBV) DNA vaccine *vs* placebo for sequential combined therapy with lamivudine (LAM) in patients with chronic hepatitis B.

***METHODS***

Two hundred and twenty five patients were randomized to LAM + vaccine (vaccine group, *n =* 109) or LAM + placebo (control group, *n =* 116). LAM treatment lasted 72 wk. Patients received the DNA vaccine or placebo by intramuscular injection mediated by EP at weeks 12 (start of treatment with vaccine or placebo, SOT), 16, 24, and 36 (end of treatment with vaccine or placebo, EOT).

***RESULTS***

In the modified intent to treat population, more patients had a HBV DNA decrease of > 2 log10 IU/mL in the vaccine group at week 12 after EOT compared with the control group. A trend toward a difference in the number of patients with undetectable HBV DNA at week 28 after EOT was obtained. Adverse events were similar. In the dynamic per protocol set, which excluded adefovir (ADV) add-on cases at each time point instantly after ADV administration due to LAM antiviral failure, more patients had a HBV DNA decrease of > 2 log10 IU/mL in the vaccine group at week 12 and 28 after EOT compared with the control group. More patients with undetectable HBV DNA at week 28 after EOT in the vaccine group were also observed. Among patients with viral load < 1000 copies/mL at week 12, more patients achieved HBeAg seroconversion in the vaccine group than among controls at week 36 after EOT, as well as less virological breakthrough and YMDD mutations.

***CONCLUSION***

The primary endpoint was not achieved using the HBV DNA vaccine. The HBV DNA vaccine could only be beneficial in subjects that have achieved initial virological response under LAM chemotherapy.

**Key words:** Chronic hepatitis B; DNA vaccine; *In vivo* electroporation; Lamivudine -resistant mutants; Randomized placebo-controlled trial

**© The Author(s) 2016.** Published by Baishideng Publishing Group Inc. All rights reserved.

**Core tip:** The study aimed to assess the efficacy and safety of *in vivo* electroporation -mediated dual-plasmid HBV DNA vaccine *vs* placebo for sequential combined therapy with lamivudine (LAM) in patients with chronic hepatitis B. The results indicated that the primary endpoint was not achieved using the HBV DNA vaccine. The HBV DNA vaccine might only be beneficial in subjects that have achieved initial virological response under LAM chemotherapy.

Yang FQ, Rao GR, Wang GQ, Li YQ, Xie Y, Zhang ZQ, Deng CL, Mao Q, Li J, Zhao W, Wang MR, Han T, Chen SJ, Pan C, Tan DM, Shang J, Zhang MX, Zhang YX, Yang JM, Chen GM. Phase IIb trial of *in vivo* electroporation mediated dual-plasmid HBV DNA vaccine in chronic hepatitis B patients under lamivudine chemotherapy. *World J Gastroenterol* 2016; In press

**INTRODUCTION**

Chronic hepatitis B (CHB) is a chronic infection caused by the hepatitis B virus (HBV). There is no specific symptoms and the diagnosis is based on the clinical description accompanied by laboratory findings (IgM anti-HBc-negative and positive result for HBsAg, HBeAg, or HBV DNA)[[1](#_ENREF_1)]. The CHB burden is global, but more significant in Asia, Pacific Islands, sub-Saharan Africa, Amazon, and Eastern Europe[[2](#_ENREF_2)]. Hepatitis B virus (HBV) is endemic in China, with about 110 million HBV carriers and at least 300000 people dying from HBV-related diseases each year[[3](#_ENREF_3)].

Anti-viral therapy aiming at clearing or radically inhibiting virus replication is the most effective treatment for CHB, and consists of interferon and reverse transcriptase inhibitors of oral nucleos(t)ide analogues (NUC) such as lamivudine (LAM), adefovir (ADV), entecavir (ETV), and tenofovir (TDF). Impaired host immune responses in patients with CHB constitute an obstacle to virus clearance. Therefore, alternative therapeutic strategies using vaccines have been suggested[[4](#_ENREF_4),[5](#_ENREF_5)].

Compared with protein vaccines, DNA vaccines mediate the *de novo* intracellular viral protein expression and synthesis in host cells, consequently boosting host-specific T cell-mediated response against HBV[[6-8](#_ENREF_6)]. However, a recent study revealed that DNA vaccines have a limited efficacy[[9](#_ENREF_9)]. The humoral and cellular immune responses of a HBV PreS2/S DNA vaccine in chronic HBV carriers are lower than in healthy volunteers[[7](#_ENREF_7),[10](#_ENREF_10)]. Therefore, alternative gene delivery strategies are being considered, including *in vivo* electroporation (EP), which was studied in animal models and approved for a phase I trial among healthy individuals[[11](#_ENREF_11)].

In order to enhance the cellular immunity of the DNA vaccine, we developed a construct encoding an interleukin-2 and interferon-γ fusion protein to potentiate local antigen presentation by antigen-presenting cells (APCs), and consequently to increase immunogenicity against HBV[[12](#_ENREF_12),[13](#_ENREF_13)].

Therefore, we conducted a phase IIa study in which patients with CHB received LAM with or without EP-mediated dual-plasmid HBV DNA vaccine. The vaccine has been shown to be safe and immunologically effective, with the HBV-specific T-cell responses induced under LAM chemotherapy showing a correlation with suppression of viral replication[[14](#_ENREF_14)]. LAM is a well-tolerated suppressor of viral replication with a strong antiviral potential and its effects are similar in Asian and Western populations[[15](#_ENREF_15)]. Nevertheless, it was reported that a large number of CHB patients under LAM monotherapy develop drug resistance mutation and virological breakthrough (VBT)[[16](#_ENREF_16)]. As suggested by a previous study[[14](#_ENREF_14)], adding vaccine therapy to LAM could be effective to prevent VBT. The aim of the present study was to further assess the efficacy and safety of the dual-plasmid strategy with LAM in patients with CHB compared with placebo and LAM.

**MATERIALS AND METHODS**

***Study design***

This was a double blind, randomized, placebo-controlled study, approved by the Chinese State Food and Drug Administration (CFDA) (license number: 2006L03542). Patients with CHB on a 72-wk LAM treatment (100 mg daily; GlaxoSmithKline Pharmaceutical Corporation, Suzhou, Jiangsu Province, China) were enrolled at 18 centers. This trial was registered with ClinicalTrials.org (NCT01487876) and approved by the ethics committee of the Peking University First Hospital, as well as by the committee of each of participating center. Written informed consent was obtained from each participant prior to enrolment. The study was conducted between April 2011 and October 2013 and in compliance with the Declaration of Helsinki. The study was monitored by Beijing Guoxin Ze Ding International Pharmaceutical Co., Ltd. (China), an independent contract research organization.

***Participants***

CHB diagnosis was carried out based on clinical symptoms and liver function tests[[17](#_ENREF_17)]. Inclusion criteria were: 1) 18-55 years old; 2) HBsAg- and HBeAg-positive for at least 6 months with HBV DNA > 106 copies/mL (1.79 × 105 IU/mL) within 4 weeks prior to randomization; and 3) serum alanine aminotransferase (ALT) 2-10 times the upper limit of normal within 4 weeks before randomization. Exclusion criteria were: (1) suspected hepatic tumors or alpha-fetoprotein (AFP) > 100 ng/mL; (2) cirrhosis; (3) co-infection with hepatitis A, C, D, or E virus; (4) co-infection with HIV; (5) other causes of liver disease; (6) serious medical or psychiatric illness; (7) abnormal serum creatinine, thrombocyte count, hemoglobin, or serum total bilirubin; or (8) pregnancy. Previous treatments for CHB were allowed, but not within 6 mo before study participation.

Between April 2011 and October 2013, 229 patients were screened and randomly assigned to the LAM + vaccine (vaccine) and LAM + placebo (control) groups (Figure 2); 111 patients were randomized to the vaccine arm and two were excluded due to loss to follow-up or protocol violation; 118 patients were randomized to the control arm and two were excluded (Figure 1). Therefore, the 225 patients that received at least one dose of study medication were designated as the ITT population: 109 in the vaccine group and 116 in the control group; 222 patients were analyzed as the modified ITT (mITT) population, excluding two patients with HBV M204V mutant in the vaccine arm and one who discontinued the allocated study agent in the control arm; and finally 107 (96.3%) and 115 (97.5%) patients with good compliance and no additional dosing were included in the PPS analysis (Figure 1).

Baseline characteristics were comparable between the two groups (Table 1). At SOT, serum HBV DNA, HBeAg, and HBsAg were similar between the two groups (all *P* > 0.05) (Table 1).

The numbers of patients who needed ADV add-on treatment, as proposed by the protocol, could be determined by the physicians in their respective local clinical centers, and are presented in Table 2. During the 72-wk study period, a similar proportion of patients in the two arms needed ADV add-on therapy.

***Randomization and blinding***

Patients were randomly assigned to the LAM + vaccine (vaccine) and LAM + placebo (control) groups. Identical labels with computer-generated random unique numeric codes were used to identify all study drug vials. An independent biostatistician was in charge of randomization using the SAS program (SAS Institute Inc., Cary, NC, USA), as well as labeling. The block size and seed numbers were sealed with the randomization list and kept at the CFDA. Participants, investigators, and sponsors were blinded until the end of the trial.

***EP-mediated dual-plasmid HBV DNA vaccine administration***

As described previously[[18](#_ENREF_18)], the dual-plasmid consists of a pS2.S HBV DNA vaccine plasmid (pcDNA3.1+/S2.S) encoding the HBV envelope middle protein and a pFP adjuvant plasmid (pcDNA3.1-/IL2+IFN-γ) containing the fused sequence of hIL-2 and hIFN-γ. The S2.S gene was amplified from the plasmid pHBVα1 with the whole HBV s gene fragment (type ayw) by PCR, and then inserted in the eukaryotic expression vector pcDNA3.1+ after the use of the EcoR1 enzyme. Supplementary Supplementary Figure 1A shows a representative map of the vaccine plasmid pS2.S of the HBV DNA vaccine. Total RNA was extracted from human peripheral blood mononuclear cells (PBMC), and cDNA was synthesized by reverse transcription. PCRs were carried out to amplify the recombined gene fragments of human interleukin-2 (hIL-2) and human interferon-γ (hIFN-γ). The fusion gene was named hIIF. The hIIF gene was cloned conventionally into the eukaryotic expression vector pcDNA3.1- after digestion by *EcoR*I. Supplementary Figure 1B shows a representative map of the adjuvant plasmid pFP of the HBV DNA vaccine. The pharmaceutical grade dual-plasmid was re-constituted in phosphate-buffered saline and evaluated for efficacy, toxicity, and safety in animals as an investigative new drug (IND)[[19](#_ENREF_19),[20](#_ENREF_20)]. The plasmids were manufactured by Guangzhou Biotech Pharmaceutical Co., Ltd., China.

Patients in the vaccine group received four immunizations with EP-mediated dual-plasmid HBV DNA vaccine (ED-DNA) consisting of a primer (week 12/start of vaccine treatment, SOT) and three boosting doses (weeks 16, 24, and 36/end of vaccine treatment, EOT) (Figure 2)[[14](#_ENREF_14)]. Each dose of 4 mL of ED-DNA (2 mg of DNA vaccine plasmid and 2 mg of adjuvant plasmid) was intramuscularly injected into four sites of the deltoid muscle on both sides of the patient, immediately followed by *in vivo* EP. Placebo (4 mL of vaccine diluent) was administrated using the same method.

The custom designed electro-pulse generator can generate square wave signals with a voltage of 0-300 V and pulse duration of 0.01-100 ms in any specific combination sequences. The device uses a medical-grade sterilized two-needle electrode made of silver Chinese acupuncture needles (diameter: 0.3 mm; length: 12 mm, for single use only). The two needles are 1 cm apart.

***Endpoints***

Responses were assessed at 12, 28, and 36 wk after EOT. The primary endpoint was the rate of undetectable HBV DNA or HBeAg seroconversion (defined as loss of HBeAg). The secondary endpoints included: (1) HBV DNA load suppression (number of patients who had HBV DNA > 2 log10 IU/mL decrease); (2) occurrence of LAM-resistant mutations or VBT; and (3) normalization of ALT levels.

***Laboratory assays***

At the end of follow-up, all serum samples were assayed by a central laboratory or at each participating center, as previously described[[14](#_ENREF_14)]. HBV serological markers at each visit were assayed using the same lots of reagents. Serum HBsAg/HBeAg status and HBV DNA levels were detected at the central laboratory (AxSYM test, dynamic range of HBsAg was 0.05-250 IU/mL, Abbot, Chicago, IL, USA; COBAS AmpliPrep/COBAS TaqMan HBV Test, ranging from 20 to 1.7 × 108 IU/mL, Roche Diagnostics, Basel, Switzerland).

Venous blood samples were obtained and peripheral blood mononuclear cells (PBMC) were separated for immunologic response test. The number of IFN-γ-secreting T lymphocytes among PBMC was assessed using an enzyme-linked immunosorbent spot (ELISPOT) assay using the human IFN-γ ELISPOT kit (BD Biosciences, San Diego, CA, USA), as previously described[[14](#_ENREF_14)]. Peptides synthesized by GL Biochem Ltd (Shanghai, China), including HLA-2 restricted CTL epitope peptides and proteins in HBV (preS2+S) T cell antigen, were used to stimulate the cells during the ELISPOT assays. The number of IFN-γ spots was counted on an AID ELISPOT reader system (AID, Germany). Data were expressed as mean SFCs/106 PBMC. Otherwise, PBMC were cultured under simulation with HBsAg. After 9 d, supernatants were assessed for HBV antigen-specific IFN-γ secretion by enzyme-linked immunosorbent assay (ELISA).

PBMC were used for cytotoxicity analysis by flow cytometry (FACSCalibur; BD Biosciences, Franklin Lake, NJ, USA), as follows. PBMC were first stained with PerCP-conjugated α-CD3 and APC-conjugated α-CD8. Then, fixation and permeabilization were performed, followed by staining with PE-conjugated IFN-γ monoclonal antibodies (BD Biosciences, Franklin Lake, NJ, USA). Results were analyzed using the FlowJo software (version 5.7.2; Tree Star Inc., Ashland, OR, USA).

***Post-treatment immuno-memory tests***

Among all participants, 45 patients were recruited 50-70 weeks after the 72-wk LAM treatment period and before unblinding. Then, IFN-γ levels in PMBC supernatants were detected by ELISA after 9 d of *in vitro* culture after HBsAg stimulation. In this case, a positive T-cell response was defined as two times increase of control level interferon-γ (IFN-γ) secretion. Therefore, the threshold of <100 pg/mL upon 9 days of culture of PBMC stimulated by HBsAg was selected. A positive response of CD8+ intracellular INF-γ secreting T-cells in PMBC was defined as no less than 0.1% frequency of positive IFN-γ intracellular staining CD8 cells, measured by flow cytometry (FACS), upon stimulation by various HBV envelope antigens.

***Safety***

All health-related events were documented during the 72-wk treatment period. Physical examinations, vital signs, hematology tests, and urinalysis were performed. In addition, anti-nuclear antibodies and anti-DNA antibodies were measured in all subjects. Patients with a failure to LAM antiviral therapy or with VBT occurrence received adefovir (ADV, 10 mg/d) as add-on treatment.

The adverse events were evaluated by clinicians and defined as mild (transient, tolerable, and minor symptoms disappearing by themselves in the intervals of drug administration), moderate (relatively long-lasting symptoms affecting daily life of the patients and requiring reduction of drug dose or administration frequency), or severe (long-lasting symptoms causing disability to the patients, who only recovered after drug discontinuation and with appropriate treatments).

***Statistical analysis***

We assumed that 40% of patients would have HBV DNA loss in the DNA vaccine group compared with 20% in the placebo arm at EOT. Therefore, 106 patients per group (total of 212 patients) were needed to achieve a 90% power with a two-sided significance level of 0.05. Assuming a drop-out rate of 15%, 240 patients were required.

A database was established using Epidata 3.1 (http://www.epidata.dk/). SAS 9.2 (SAS institute Inc., Cary, NC, USA) was used for statistical analysis. Data double entry was performed, including range and consistency checks. Analysis was conducted on all eligible patients according to the intent-to-treat (ITT) principle. A dynamic per protocol set analysis (dPPS) was performed to take into account the patients who received adefovir as add-on treatment. All categorical data were analyzed using the chi-square test, Fisher’s exact test, or Mann-Whitney *U* test, as appropriate. Continuous data were analyzed using the Student *t* test or Wilcoxon Rank Sum test, as appropriate. *P* values < 0.05 were considered statistically significant.

**RESULTS**

***mITT analysis***

The mITT analysis revealed a modest and transient therapeutic effect. At week 12 after EOT, there were more patients with a HBV DNA decrease of > 2 log10 IU/mL in the vaccine group (10.3% *vs* 3.5%, *P =* 0.044) compared with the control group. In addition, there was a trend toward differences in the number of patients in the two groups with undetectable HBV DNA at 28 wk after EOT (28.0% *vs* 17.4%, *P =* 0.058) (Table 3) and a decrease in HBeAg > 1 log10 IU/mL (34.6% *vs* 23.5%, *P =* 0.068) 36 wk after EOT (Table 3). There was no difference in HBV DNA levels between the two groups at any time point.

Detection for drug-resistant mutants, as well as all other HBV serological markers, showed no remarkable differences between the two groups throughout the study.

***dPPS analysis***

Taking into account the patients with ADV add-on treatment (Table 2), a dPPS was performed: cases were excluded dynamically from the PPS at each visit when they had received ADV add-on treatment. Compared with the control group, more patients with a HBV DNA decrease of > 2 log10 IU/mL were found in the vaccine group at weeks 12 (11.2% *vs* 3.8%, *P =* 0.038) and 28 (17.3% *vs* 7.1%, *P =* 0.040) after EOT. In addition, there were more patients with undetectable HBV DNA at week 28 after EOT (38.7% *vs* 23.0%, *P =* 0.030) in the vaccine group compared with controls (Table 3).

***Stratified analysis***

A stratified analysis was performed, as previously described[[21](#_ENREF_21)]. At SOT, 30 patients (11 and 19 in the vaccine and control groups, respectively) with serum HBV DNA levels < 1000 copies/mL (1.79 × 102 IU/mL) were selected. More patients achieved HBeAg seroconversion in the vaccine group (6/11, 54.5%) than among controls (3/19, 15.8%) (*P =* 0.042) at week 36 after EOT (Table 3). At SOT, there were three cases with HBeAg seroconversion in the vaccine group compared to four in the control group. The three patients in the vaccine group remained in seroconversion status throughout the trial until the endpoint, while the seroconversion status among the four control patients was unstable due either to HBeAb loss or reappearance of HBeAg, resulting in only two patients remaining in seroconversion at week 36 after EOT. Two patients in the vaccine group (2/11) had HBsAg levels below 1000 IU/mL at SOT, and showing HBeAg seroconversion at week 36 after EOT. Meanwhile, eight patients in the control group (8/19) had HBsAg levels below 1000 IU/mL at SOT, with two showing HBeAg seroconversion at week 36 after EOT. In addition, less VBT (18.2% *vs* 47.4%, *P =* 0.140) and YMDD mutations (9.1% *vs* 42.1%, *P =* 0.100) were observed in the vaccine group at week 36 after EOT, but the difference was not statistically significant.

Furthermore, in the dPPS analysis of the 30 patients by excluding the ADV add-on cases at each time point instantly after ADV administration due to LAM antiviral failure, viral load decrease at weeks 28 (*P =* 0.017) and 36 (*P =* 0.023) after EOT were more remarkable in the LAM + vaccine group (Figure 3A), in which more patients (8/9 *vs* 8/18, *P =* 0.042) had serum HBV DNA maintained at undetectable levels at week 28 after EOT. Similarly, less VBT (0.0% *vs* 41.2%, *P =* 0.058) and YMDD mutations (0.0% *vs* 35.3%, *P =* 0.063) were observed in the vaccine group at week 36 after EOT.

Baseline serum HBsAg levels at SOT were higher in the vaccine group (mea*n =* 5,287 IU/mL) than in controls (1494 IU/mL) (Figure 3B), probably due to lower ALT levels at week 0 (mean of 148 U/L) compared with control values (mean of 195 U/L, Figure 3C). As shown in Figure 2B, throughout the trial, serum HBsAg tended to decrease slightly (4116 IU/mL at week 36 after EOT) in the vaccine group, while remaining at a similar level with some elevation in the control group (2991 IU/mL at week 36 after EOT).

In the present study, normalized ALT levels displayed a substantial decrease during the first 16 weeks of treatment and then stabilized (Figure 3C). No significant differences were observed between the two groups. However, two patients in the vaccine group experienced a flare-up with ALT elevation up to 593 U/L in one VBT patient and 424 U/L in another patient with well suppressed viral status at weeks 12 and 36 after EOT.

These analyses were also performed in the group of patients with >1000 copies/mL (1.79×102 IU/mL), but no difference could be observed in any parameter.

***Immune response***

Forty-five patients were recruited 50-70 weeks after EOT, including 22 and 23 in the vaccine and control groups, respectively, to assess immuno-memory responses in venous blood samples.

After 9 d of HBsAg stimulation, 19/22 (86.4%) patients showed positive immunological response with IFN-γ secretion in the vaccine group, compared with 13/23 (56.5%) patients in the control group (*P =* 0.027). In addition, intracellular IFN-γ secreting CD3+/CD8+ T cells among PMBC were detected using flow cytometry after 6-8 h of simulation with HBV-related antigens, and 17/22 (77.3%) versus 10/23 (43.5%) patients showed positive results in the vaccine and control groups, respectively (*P =* 0.021) (Table 4).

In either *ex vivo* or *in vitro* ELISPOT assay, the two groups showed similar levels of IFN-γ-secreting T-cells per million PBMC, after stimulation by HBV specific peptide pools for 48 h or 9 d.

***Safety***

Vaccine injections were well tolerated with limited mild adverse events observed in 14 of 111 (12.6%) DNA vaccine recipients compared with 15 of 118 (12.7%) patients receiving placebo. Adverse effects did not lead to any dropout during the study. Only two patients in the vaccine group (1.8%) and one in the control group (0.9%) reported non-drug-related serious adverse events during treatment (*P* > 0.05). In the vaccine group, a 38-year-old male patient had a severe CHB relapse due to LAM resistance at week 60. The other patient was a 46-year-old female who underwent surgery for breast fibroma and thyroid adenoma at weeks 54 and 55. In the control group, a 43-year-old female patient was diagnosed with primary liver cancer by ultrasound and computed tomography at week 36 after EOT. All three patients were admitted to the hospital for further treatments. These events were reported as severe adverse events. Four patients receiving the DNA vaccine (3.6%) and five patients receiving the placebo (4.2%) had increased total bilirubin levels at week 36 after EOT (*P* > 0.05). No clinically significant serological abnormalities were observed. No antinuclear antibodies against double-strand DNA were detected in any patients during and after treatment.

**Discussion**

The aim of this study was to assess the efficacy of the dual-plasmid strategy with LAM in patients with CHB compared with a placebo + LAM as a phase IIb trial. Our results indicated that the primary endpoint was not achieved using the EP-mediated dual-plasmid HBV DNA vaccine. The HBV DNA vaccine could only be beneficial in subjects that have achieved initial virological response under LAM chemotherapy.

The efficacy did not reach the pre-set primary endpoint as per the registered protocol, but the mITT analysis showed that serum HBV DNA and HBeAg levels decreased, and the secondary efficacy measures of the treatment were better in the DNA vaccine + LAM chemotherapy group than in the placebo + LAM group. There was a higher number of patients with a HBV DNA decrease of > 2 log10 IU/mL in the vaccine group at week 12 after EOT compared with the control group. There were tendencies toward a difference in the number of patients with undetectable HBV DNA at week 28 after EOT and that with a decrease in HBeAg of > 1 log10 IU/mL at week 36 after EOT. The vaccine was safe, well tolerated, and resulted in a rate of adverse events comparable to that of the control group. Again, the secondary efficacy measures of the vaccine was more obvious in the dPPS analysis, in which there was a higher number of patients with a HBV DNA decrease of > 2 log10 IU/mL in the vaccine group at weeks 12 and 28 after EOT, as well as that with undetectable HBV DNA at week 28 after EOT. The discrepancy between the data sets is that we took into account the patients who needed add-on therapy with ADV in the PPS population and thus suggests that the added value of the DNA vaccine, as a kind of HBV-specific immune modulatory therapy upon LAM antiviral therapy, might be achieved in easy-to-treat patients with CHB but without antiviral failure. Therefore, these results suggest that the effect of the vaccine could be better in highly selected patients, but additional studies are necessary to identify these patients adequately.

In the present study, the frequency of VBT was higher than that in our previous phase IIa study. Indeed, the occurrence of VBT in patients receiving LAM + vaccine was 50.0% versus 63.5% in the LAM + placebo at week 72, compared with 18.2% and 47.4%, respectively, which is higher than that in the previous trial[[14](#_ENREF_14)]. In addition, the numbers of patients achieving a decrease of HBV DNA > 2 log10 at week 72 were comparable between the two groups, as well as the numbers of patients with undetectable HBV DNA, in disagreement with our previous phase IIa trial[[14](#_ENREF_14)]. Many reasons could explain these discrepancies. There is a possibility that some patients were not naïve to NUC and some mutations could already have been present. The patients enrolled in the present study were different from the ones originally recruited in the phase IIa trial, which included only four centers in Beijing, while the present study involved many centers across China. Nevertheless, the results of the stratified analysis are similar to our previous findings[[14](#_ENREF_14)], and may suggest that the lower HBV DNA load at week 72 is probably due to lower rates of VBT and YMDD mutation, but this needs to be confirmed.

In the stratified analysis of 30 patients with a viral load < 1000 copies/mL (1.79 × 102 IU/mL) at SOT, more patients achieved HBeAg seroconversion in the vaccine group than among controls. In addition, less VBT and YMDD mutations were observed in the vaccine group. If HBeAg seroconversion is selected as trial endpoint for the DNA vaccine, the SOT baseline serum HBsAg should be concerning in this stratified analysis, as it favors HBeAg seroconversion at the end[[22](#_ENREF_22)]. Interestingly, baseline serum HBsAg levels at SOT were higher in the vaccine group than among controls, probably due to the higher ALT levels at week 0 among controls, which is an indicator of host cellular immune response and would have been favorable for viral clearance among the patients of this group. During the trial course or at the endpoint, serum HBsAg tended to decrease slightly in the vaccine group, while remaining at the relatively same level with some elevation in the control group. ALT flare-ups occurred only in the vaccine arm, but were not associated with VBT in one case. Such host-induced flare-ups have been reported in patients responding to pegylated interferon-alpha 2a (peg-IFN), and are thought to reflect the immune clearance of HBV[[21](#_ENREF_21),[22](#_ENREF_22)].

HBeAg seroconversion has been used as a reliable criterion for therapeutic efficacy of antiviral treatments and could be the impetus for HBsAg loss and seroconversion[[23](#_ENREF_23)]. It is considered a therapeutic endpoint and has been associated with immune restoration or control, and with reduced morbidity and mortality of CHB patients[[14](#_ENREF_14),[23](#_ENREF_23),[24](#_ENREF_24)]. In the present study, LAM showed good efficacy during the first stage of viral suppression, and the adjusted results provided clinical evidence that DNA vaccine can induce an immune control of HBV infection in CHB patients who had already achieved initial virological responses under LAM antiviral therapy. Indeed, treatment guidelines suggest that therapy may be discontinued after 2-3 years of consolidation therapy in HBeAg-positive non-cirrhotic CHB patients that achieve HBeAg seroconversion during an antiviral therapy[[25](#_ENREF_25)]. It was reported by recent studies that patients with NUC therapy either switching to or adding-on peg-IFN achieve higher rates of sustained response than those continuing NUC monotherapy[[21](#_ENREF_21),[26](#_ENREF_26),[27](#_ENREF_27)]. Combination therapy of antivirals and immunomodulators, either to improve response to NUC, IFN, or newly developed therapeutic vaccines, is considered crucial for investigators to achieve “functional” cure of HBV, defined as HBsAg loss or off-therapy immune control[[27](#_ENREF_27)]. The present study suggests that for patients receiving long-term therapy, it might be possible to enhance the chances of HBeAg seroconversion by adding-on a finite course of immunomodulatory therapy, especially a HBV-specific one.

Patients with CHB often present immune defects such as weak or undetectable helper and cytotoxic T cell-mediated immune responses[[22](#_ENREF_22),[28](#_ENREF_28)], and immune modulation therapy aims to induce a sustainable HBV-specific multi-epitope and polyclonal T cell response. Therapeutic HBV DNA vaccine, as an HBV-specific immunomodulator, might be superior to interferon, as a non-specific one, in terms of adverse effects and dosing regimens, given that vaccine therapy was well tolerated and only four injections were administered in our present and previous studies. A previous study in macaques revealed that DNA vaccine may induce some immune memory[[29](#_ENREF_29)]. Results from our Phase IIa trial showed that the patients receiving the DNA vaccine had a significant elevation of HBV-specific IFN-γ-secreting T cell count compared with baseline[[14](#_ENREF_14)], suggesting that the DNA vaccine plays a role in boosting the host cellular immune responses in favor of viral eradication among CHB patients. Because local laboratories were not equipped to assess T cells and because of logistics issues for shipping fresh samples to a central laboratory, we could not assess T cells in the present Phase IIb trial. Nevertheless, we attempted to evaluate immune response in this cohort by recalling the patients who completed the follow-up and before unblinding. Compared with the control group, there was a higher proportion of positive responses against HBsAg and HBV T-cell related peptides in terms of IFN-γ secretion and phenotypes of T cells from the PBMC of patients receiving DNA vaccination. Additional studies are necessary to determine the optimal doses and administration strategies.

This study is not without limitations, as the sample size was small in the stratified analysis, it potentially prevented us from reaching a conclusion on its efficacy. In addition, the follow-up was short, preventing the evaluation of the long-term efficacy and the beneficial outcome of the HBeAg seroconversion achieved by the vaccine. In the present study, only one vaccine administration schedule was tried, based on our previous phase IIa trial[[14](#_ENREF_14)], but additional studies could explore the best administration schedule. Last but not the least, we failed to assess the real-time HBV-specific T-cell immune responses of the subjects during the trial.

Even though the clinical efficacy as a whole was a minor outcome in this trial, our findings are still of relevance for future studies, in which DNA vaccine will sequentially be combined with more potent NUC or NUC less susceptible to drug resistance, like ETV or TDF. Since there were only four doses of the vaccine in the present trial, investigations on more appropriate strategies of vaccination including the number of injections and timing, either in animals or small-scale clinical trials, are crucial before large-scale trials. The results of positive memory T-cell responses in the present study would be implicative for these further studies, aiming to intensify and prolong the HBV-specific T-cell responses induced by the vaccine under NUC chemotherapy among CHB patients.

In conclusion, the primary endpoint was not achieved using the HBV DNA vaccine. The HBV DNA vaccine might only be beneficial in subjects that have achieved initial virological response under LAM chemotherapy. Additional studies are necessary to identify the subset of patients that could benefit the most from the EP-mediated DNA vaccination.

**COMMENTS**

***Background***

Chronic hepatitis B (CHB) infection leads to serious morbidity and mortality. Antivirals have some efficacy, but virological breakthrough will eventually occur.

***Research frontiers***

DNA vaccine are a novel type of vaccine that induce the production of antigen proteins by the cells.

***Innovations and breakthroughs***

The primary endpoint was not achieved using the HBV DNA vaccine. Nevertheless, the HBV DNA vaccine could only be beneficial in subjects that have achieved initial virological response under lamivudine chemotherapy.

***Applications***

Patients with a response to lamivudine could receive the DNA vaccine as an add-on therapy in order to prolong the response.

***Terminology***

A DNA vaccine is a vaccine that introduces a DNA sequence in host cells to make them produce the protein that will serve as the antigen for antibody production. A virological breakthrough occurs when an antiviral therapy fails.

***Peer-review***

This manuscript is well written and interesting.

**REFERENCES**

1 **Weinbaum CM**, Williams I, Mast EE, Wang SA, Finelli L, Wasley A, Neitzel SM, Ward JW. Recommendations for identification and public health management of persons with chronic hepatitis B virus infection. *MMWR Recomm Rep* 2008; **57**: 1-20 [PMID: 18802412]

2 **Sorrell MF**, Belongia EA, Costa J, Gareen IF, Grem JL, Inadomi JM, Kern ER, McHugh JA, Petersen GM, Rein MF, Strader DB, Trotter HT. National Institutes of Health Consensus Development Conference Statement: management of hepatitis B. *Ann Intern Med* 2009; **150**: 104-110 [PMID: 19124811]

3 **Sunbul M**. Hepatitis B virus genotypes: global distribution and clinical importance. *World J Gastroenterol* 2014; **20**: 5427-5434 [PMID: 24833873 DOI: 10.3748/wjg.v20.i18.5427]

4 **Gurunathan S**, Klinman DM, Seder RA. DNA vaccines: immunology, application, and optimization\*. *Annu Rev Immunol* 2000; **18**: 927-974 [PMID: 10837079 DOI: 10.1146/annurev.immunol.18.1.927]

5 **Shi TD**, Wu YZ, Jia ZC, Zou LY, Zhou W. Therapeutic polypeptides based on HBV core 18-27 epitope can induce CD8+ CTL-mediated cytotoxicity in HLA-A2+ human PBMCs. *World J Gastroenterol* 2004; **10**: 1902-1906 [PMID: 15222033]

6 **Kim CY**, Kang ES, Kim SB, Kim HE, Choi JH, Lee DS, Im SJ, Yang SH, Sung YC, Kim BM, Kim BG. Increased in vivo immunological potency of HB-110, a novel therapeutic HBV DNA vaccine, by electroporation. *Exp Mol Med* 2008; **40**: 669-676 [PMID: 19116452 DOI: 10.3858/emm.2008.40.6.669]

7 **Mancini-Bourgine M**, Fontaine H, Scott-Algara D, Pol S, Bréchot C, Michel ML. Induction or expansion of T-cell responses by a hepatitis B DNA vaccine administered to chronic HBV carriers. *Hepatology* 2004; **40**: 874-882 [PMID: 15382173 DOI: 10.1002/hep.20408]

8 **Chen M**, Jagya N, Bansal R, Frelin L, Sällberg M. Prospects and progress of DNA vaccines for treating hepatitis B. *Expert Rev Vaccines* 2016; **15**: 629-640 [PMID: 26652035 DOI: 10.1586/14760584.2016.1131615]

9 **Fontaine H**, Kahi S, Chazallon C, Bourgine M, Varaut A, Buffet C, Godon O, Meritet JF, Saïdi Y, Michel ML, Scott-Algara D, Aboulker JP, Pol S. Anti-HBV DNA vaccination does not prevent relapse after discontinuation of analogues in the treatment of chronic hepatitis B: a randomised trial--ANRS HB02 VAC-ADN. *Gut* 2015; **64**: 139-147 [PMID: 24555998 DOI: 10.1136/gutjnl-2013-305707]

10 **Roy MJ**, Wu MS, Barr LJ, Fuller JT, Tussey LG, Speller S, Culp J, Burkholder JK, Swain WF, Dixon RM, Widera G, Vessey R, King A, Ogg G, Gallimore A, Haynes JR, Heydenburg Fuller D. Induction of antigen-specific CD8+ T cells, T helper cells, and protective levels of antibody in humans by particle-mediated administration of a hepatitis B virus DNA vaccine. *Vaccine* 2000; **19**: 764-778 [PMID: 11115698]

11 **Yang HY**, Chen GM, Cui YM, Xia Z, Mei C, Rao GR, Guo-Yu MO, Yang RC. Clinical study on safety and immunogenicity of therapeutic dual-plasmid HBV DNA vaccine mediated by in vivo electroporation. *Jiefangjun Yixue Zazhi* 2013; **38**: 204-209

12 **Yang FQ**, Chen GM, Rao GR, Yu-Hong XU, Zhao YG, Xiao-Qiang HE, Sun XH, Hou JL. Interleukin-2 and Interferon-Fusion Protein Expression Plasmid Enhanced Potency of HBV DNA Vaccination in HBV Transgenic Mice. *Redai Yixue Zazhi* 2008

13 **Yang FQ**, Chen GM, Rao GR, Yu-Hong XU, Zhao YG, Xiao-Qiang HE, Sun XH, Hou JL. Adjuvant effect of IL-2 and IFN-γ fusion protein gene plasmid on HBV DNA vaccination in BALB/c mice. *Zhongguo Xiandai Yixue Zazhi* 2008: 838-842

14 **Yang FQ**, Yu YY, Wang GQ, Chen J, Li JH, Li YQ, Rao GR, Mo GY, Luo XR, Chen GM. A pilot randomized controlled trial of dual-plasmid HBV DNA vaccine mediated by in vivo electroporation in chronic hepatitis B patients under lamivudine chemotherapy. *J Viral Hepat* 2012; **19**: 581-593 [PMID: 22762143 DOI: 10.1111/j.1365-2893.2012.01589.x]

15 **Lai CL**, Chien RN, Leung NW, Chang TT, Guan R, Tai DI, Ng KY, Wu PC, Dent JC, Barber J, Stephenson SL, Gray DF. A one-year trial of lamivudine for chronic hepatitis B. Asia Hepatitis Lamivudine Study Group. *N Engl J Med* 1998; **339**: 61-68 [PMID: 9654535 DOI: 10.1056/NEJM199807093390201]

16 **Schalm SW**. Clinical implications of lamivudine resistance by HBV. *Lancet* 1997; **349**: 3-4 [PMID: 8988109 DOI: 10.1016/S0140-6736(05)62154-X]

17 **Desmet VJ**, Gerber M, Hoofnagle JH, Manns M, Scheuer PJ. Classification of chronic hepatitis: diagnosis, grading and staging. *Hepatology* 1994; **19**: 1513-1520 [PMID: 8188183]

18 **He X**, Chen G, Huang Y. Construction and identification of therapeutic double-plasmid hbv DNA vaccine. *Jiefangjun Yixue Zazhi* 2003; 28: 493-496

19 **Mo G,** Chen G, Yang F. Induction of immune responses and safety profiles in rhesus macaques immunized with the therapeutic DNA vaccine against hepatitis B virus. *Jiefangjun Yixue Zazhi* 2003; **28**: 508-510

20 **Mo G**, Chen G, Huang Z. A study of distribution in the tissues and long-term toxicity of therapeutic DNA vaccine in mice. *Jiefangjun Yixue Zazhi* 2003; **28**: 504-507

21 **Ning Q**, Han M, Sun Y, Jiang J, Tan D, Hou J, Tang H, Sheng J, Zhao M. Switching from entecavir to PegIFN alfa-2a in patients with HBeAg-positive chronic hepatitis B: a randomised open-label trial (OSST trial). *J Hepatol* 2014; **61**: 777-784 [PMID: 24915612 DOI: 10.1016/j.jhep.2014.05.044]

22 **Mukherjee R**, Reddy PB, Arava J, Rao P, Mitnala S, Gupta R, Reddy D. Relationship between serum HBsAg level, HBV DNA level, and peripheral immune cells in patients with chronic hepatitis B virus infection. *Hepat Med* 2010; **2**: 157-162 [PMID: 24367213 DOI: 10.2147/HMER.S13796]

23 **Xu DZ**, Wang XY, Shen XL, Gong GZ, Ren H, Guo LM, Sun AM, Xu M, Li LJ, Guo XH, Zhen Z, Wang HF, Gong HY, Xu C, Jiang N, Pan C, Gong ZJ, Zhang JM, Shang J, Xu J, Xie Q, Wu TF, Huang WX, Li YG, Xu J, Yuan ZH, Wang B, Zhao K, Wen YM. Results of a phase III clinical trial with an HBsAg-HBIG immunogenic complex therapeutic vaccine for chronic hepatitis B patients: experiences and findings. *J Hepatol* 2013; **59**: 450-456 [PMID: 23669281 DOI: 10.1016/j.jhep.2013.05.003]

24 **Zhou H**, Min J, Zhao Q, Gu Q, Cong H, Li Y, He S. Protective immune response against Toxoplasma gondii elicited by a recombinant DNA vaccine with a novel genetic adjuvant. *Vaccine* 2012; **30**: 1800-1806 [PMID: 22240340 DOI: 10.1016/j.vaccine.2012.01.004]

25 The guideline of prevention and treatment for chronic hepatitis B (2010 version). *Zhonghua Liu Xing Bing Xue Za Zhi* 2011; **32**: 405-415 [PMID: 21569677]

26 **Brouwer WP**, Xie Q, Sonneveld MJ, Zhang N, Zhang Q, Tabak F, Streinu-Cercel A, Wang JY, Idilman R, Reesink HW, Diculescu M, Simon K, Voiculescu M, Akdogan M, Mazur W, Reijnders JG, Verhey E, Hansen BE, Janssen HL. Adding pegylated interferon to entecavir for hepatitis B e antigen-positive chronic hepatitis B: A multicenter randomized trial (ARES study). *Hepatology* 2015; **61**: 1512-1522 [PMID: 25348661 DOI: 10.1002/hep.27586]

27 **Lampertico P**. The royal wedding in chronic hepatitis B: The haves and the have-nots for the combination of pegylated interferon and nucleos(t)ide therapy. *Hepatology* 2015; **61**: 1459-1461 [PMID: 25645624 DOI: 10.1002/hep.27731]

28 **Dimitropoulou D**, Karakantza M, Tsamandas AC, Mouzaki A, Theodorou G, Gogos CA. T-lymphocyte subsets in peripheral blood and liver tissue of patients with chronic hepatitis B and C. *In Vivo* 2011; **25**: 833-840 [PMID: 21753143]

29 **Kulkarni V**, Rosati M, Jalah R, Ganneru B, Alicea C, Yu L, Guan Y, LaBranche C, Montefiori DC, King AD, Valentin A, Pavlakis GN, Felber BK. DNA vaccination by intradermal electroporation induces long-lasting immune responses in rhesus macaques. *J Med Primatol* 2014; **43**: 329-340 [PMID: 24810337 DOI: 10.1111/jmp.12123]

**P-Reviewer:** Rasti M **S-Editor:** Yu J **L-Editor:** **E-Editor:**

**Specialty type:** Gastroenterology and hepatology

**Country of origin:** China

**Peer-review report classification**

Grade A (Excellent): 0

Grade B (Very good): B

Grade C (Good): 0

Grade D (Fair): 0

Grade E (Poor): 0



**Figure 1** **Patient disposition and different analysis populations.** M204V mutant: The lamivudine-resistant amino acid mutation of HBV; mITT: Modified intent to treat.



**Figure 2** **Study design.** Four intramuscular injections of either DNA vaccine (DNA Vacc.) or placebo mediated by *in vivo* electroporation were given to each patient. LAM: Lamivudine.



**A**

**B**

**C**

|  |
| --- |
|  |
| Vaccine / Placebo |
| LAM |
| No. of patients |
| Vaccine | 11 11 11 11 11 10 9 8 11 10 11 11 8 11 10 11 11 10 9 8 |
| Control | 19 19 19 19 19 18 18 16 19 19 19 19 16 19 19 18 19 19 17 17 |

**Figure 3 Serological kinetics of hepatitis B virus DNA (A), HBsAg (B) and alanine aminotransferase (C) in 30 patients with week 12 HBV DNA < 1000 copies/mL (1.79 × 102 IU/mL).** Data are presented as mean ± SEM. DNA Vacc.: Lamivudine (LAM) plus DNA vaccine arm; Ctr.: LAM + placebo arm.

**Table 1 Characteristics of the modified intent to treat population**

|  |  |  |  |
| --- | --- | --- | --- |
| **Characteristics** | **Vacc. (*n =* 107)** | **Ctr. (*n =* 115)** | ***P* value** |
| Demography |  |  |  |
|  Age, median (IQR) | 28.9 (23.1-37.1) | 29.3 (24.0-35.4) | 0.950 |
|  Male sex, *n* (%) | 86 (78.9) | 86 (74.1) | 0.400 |
|  Weight (kg), median (IQR) | 62.5 (56.0-70.0) | 63.0 (56.0-70.0) | 0.907 |
|  Genotype B, *n* (%) | 42/107 (39.3) | 39/115 (33.9) | 0.409 |
| Week 0 laboratory results |  |  |  |
|  ALT (IU/L), median (IQR) | 119.0 (94.0-167.5) | 136.0 (102.6-195.3) | 0.066 |
|  HBV DNA (log10 IU/mL), median (IQR) | 8.1 (7.7-8.3) | 8.1 (7.5-8.4) | 0.9532 |
|  HBeAg (COI), median (IQR) | 852.0 (223.5-1093.0) | 926.6 (508.5-1076.0) | 0.1972 |
|  HBsAg (IU/mL), median (IQR) | 10261.0 (3953.0-24089.0) | 8637.0 (2748.0-24436.0) | 0.7542 |
| Week 12 laboratory results1 |  |  |  |
| HBV DNA (log10 IU/mL), median (IQR) | 3.5 (2.6-4.5) | 3.4 (2.6-4.6) | 0.7542 |
|  HBeAg (COI), median (IQR) | 147.4 (15.6-650.0) | 45.9 (11.9-1011.0) | 0.6052 |
|  HBsAg (IU/mL) median (IQR) | 4072.0 (1608.0-8514.0) | 3023.5 (1167.0-11194.0) | 0.4602 |

1The time point when the injection of study agent started; 2Analyzed with Wilcoxon Rank Sum test. mITT: Modified intent to treat; LAM: Lamivudine; IQR: Interquartile ranges; ALT: Alanine aminotransferase; COI: Semi-quantification by S/CO [the signal value ratio of sample (S) to the control (CO)]; Vacc., LAM + vaccine arm; Ctr., LAM + placebo arm.

**Table 2 Cumulative number of patients in the per protocol set with adefovir add-on treatment due to lamivudine antiviral failure at each time point of the trial *n* (%)**

|  |  |  |
| --- | --- | --- |
| **Week** | **Vacc. (*n =* 102)** | **Ctr. (*n =* 113)** |
| 24 | 1 (0.98) | 0 (0.00) |
| 36 | 2 (1.96) | 1 (0.88) |
| 40 | 4 (3.92) | 5 (4.42) |
| 48 | 7 (6.86) | 8 (7.08) |
| 52 | 11 (10.78) | 14 (12.39) |
| 56 | 17 (16.67) | 18 (15.93) |
| 60 | 27 (26.47) | 26 (23.01) |
| 64 | 34 (33.33) | 32 (28.32) |

Vacc.: LAM + vaccine arm; Ctr.: LAM + placebo arm; ADV: Adefovir; LAM: Lamivudine.

**Table 3 Virological responses after the end of treatment with DNA vaccination or placebo injection in contrast to the baseline at week 12 in different analysis populations**

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **Population** | **Responses** | **12 wk after EOT (week 48)** | **28 wk after EOT (week 64)** | **36 wk after EOT (week 72)** |
| **Vacc.** | **Ctr.** | ***P* value** | **Vacc.** | **Ctr.** | ***P* value** | **Vacc.** | **Ctr.** | ***P* value** |
| mITT | No. of patients | 107 | 115 |  | 107 | 115 |  | 107 | 115 |  |
| HBV DNA (log10 IU/mL), median (IQR) | 3.06 (0-4.63) | 3.05 (1.43-5.02) | 0.742 | 2.18 (0-5.01) | 2.78 (1.41-4.86) | 0.164 | 1.88 (0-4.36) | 2.37 (0-5.38) | 0.329 |
| HBV DNA > 2 log10 IU/mL decrease, *n* (%) | 11 (10.3) | 4 (3.48) | **0.044** | 13 (12.2) | 6 (5.2) | 0.065 | 10 (9.4) | 7 (6.1) | 0.362 |
| HBV DNA undetectable, n (%) | 27 (25.2) | 23 (20.0) | 0.351 | 30 (28.0) | 20 (17.4) | 0.058 | 24 (22.4) | 26 (22.6) | 0.975 |
| HBeAg > 1 log10IU/mL decrease, *n* (%) | 22 (20.6) | 22 (19.1) | 0.789 | ND | ND |  | 37 (34.6) | 27 (23.5) | 0.068 |
| HBeAg seroconversion, *n* (%) | 16 (15.0) | 10 (8.7) | 0.147 | ND | ND |  | 18 (16.8) | 15 (13.0) | 0.429 |
| dPPS | No. of patients | 98 | 108 |  | 75 | 87 |  | 68 | 81 |  |
| HBV DNA (log10 IU/mL), median (IQR) | 2.99 (0-4.59) | 3.05 (1.43-5.02) | 0.593 | 2.16 (0-4.66) | 2.78 (1.41-4.86) | 0.104 | 1.85 (0-3.78) | 2.37 (0-5.38) | 0.262 |
| HBV DNA > 2 log10IU/mL decrease, *n* (%) | 11 (11.2) | 4 (3.8) | **0.038** | 13 (17.3) | 6 (7.1) | **0.040** | 10 (14.7) | 7 (8.9) | 0.246 |
| HBV DNA undetectable, *n* (%) | 27 (27.6) | 23 (21.3) | 0.296 | 29 (38.7) | 20 (23.0) | **0.030** | 24 (35.3) | 26 (32.1) | 0.681 |
| HBeAg > 1 log10IU/mL decrease, *n* (%) | 20 (20.4) | 21 (19.4) | 0.863 | ND | ND |  | 29 (42.7) | 25 (30.9) | 0.136 |
| HBeAg seroconversion, *n* (%) | 16 (16.3) | 10 (9.3) | 0.127 | ND | ND |  | 18 (26.5) | 15 (18.5) | 0.244 |
| Subgroup1 | No. of patients | 11 | 19 |  | 11 | 19 |  | 11 | 19 |  |
| HBV DNA (log10 IU/mL), median (IQR) | 0 (0-1.16) | 1.43 (0-2.64) | 0.268 | 0 (0-0) | 1.73 (0-3.6) | 0.053 | 0 (0-0) | 0 (0-4.32) | 0.172 |
| HBV DNA undetectable, *n* (%) | 8 (72.7) | 7 (36.8) | 0.128 | 8 (72.7) | 8 (44.1) | 0.142 | 7 (63.6) | 9 (47.4) | 0.466 |
| HBeAg seroconversion, *n* (%) | 5 (45.5) | 2 (10.5) | 0.068 | ND | ND |  | 6 (54.6) | 3 (15.8) | **0.042** |
| dPPS of subgroup2 | No. of patients | 11 | 19 |  | 9 | 18 |  | 9 | 17 |  |
| HBV DNA (log10 IU/mL), median (IQR) | 0 (0-1.16) | 1.43 (0-2.64) | 0.268 | 0 (0-0) | 1.73 (0-3.6) | 0.053 | 0 (0-0) | 0 (0-4.32) | 0.172 |
| HBV DNA undetectable, *n* (%) | 8 (72.7) | 7 (36.8) | 0.128 | 8 (88.9) | 8 (44.4) | **0.042** | 7 (77.8) | 9 (52.9) | 0.399 |
| HBeAg seroconversion, *n* (%) | 5 (45.5) | 2 (10.5) | 0.068 | ND | ND |  | 6 (66.7) | 3 (17.7) | **0.028** |

1Subgroup, 30 patients with week 12 HBV DNA < 1000 copies/mL(1.79 × 102 IU/mL); 2dPPS of subgroup, population for analysist of the 30 patients with week 12 HBV DNA < 1000 copies/mL (1.79 × 102 IU/mL) by ruling out the ADV add-on cases at each time point instantly after ADV administration due to lamivudine (LAM) antiviral failure. Data were analyzed with Fisher’s exact test. mITT: Modified intent to treat; dPPS: Dynamic Per Protocol Set by ruling out the adefovir (ADV) add-on cases at each time point instantly after ADV administration due to lamivudine (LAM) antiviral failure; ND: Not detected; Vacc.: LAM + DNA vaccine arm; Ctr., LAM + placebo arm.

**Table 4 Hepatitis B virus antigen-specific T-cell responses in 45 patients from week 50 to 70 after the trial*****n* (%)**

|  |  |  |  |
| --- | --- | --- | --- |
| **Positive responses** | **Vacc. (*n =* 22)** | **Ctr. (*n =* 23)** | ***P* value** |
| 1IFN-γ secretion  | 19 (86.4) | 13 (56.5) | 0.047 |
| 2CD3 +CD8 +IFN-γ | 17 (77.3) | 10 (43.5) | 0.021 |

1A positive T-cell response was defined as two times control level increase of interferon-γ (IFN-γ) secretion, tested by ELISA, that in itself was no less than 100 pg/ml upon 9 d cultivation of PBMC stimulated by HBsAg; 2A positive T-cell response was defined as no less than 0.1 percentage frequency of positive IFN-γ intracellular staining CD8 cells, tested by FACS, upon stimulation by various HBV envelope antigens. Vacc., LAM + DNA vaccine arm; Ctr., LAM + placebo arm.