

# World Journal of *Gastrointestinal Pharmacology and Therapeutics*

*World J Gastrointest Pharmacol Ther* 2021 March 18; 12(2): 32-39





**ORIGINAL ARTICLE**

**Retrospective Study**

- 32 Histopathological evaluation of long-term tenofovir disoproxil fumarate treatment in patients with hepatitis be antigen-negative chronic hepatitis B

*Abayli B, Abaylı C, Gencdal G*

## Contents

*World Journal of Gastrointestinal Pharmacology and Therapeutics*

**Bimonthly Volume 12 Number 2 March 18, 2021**

### ABOUT COVER

Editorial Board Member, Ugo Cioffi, MD, PhD, Full Professor, Surgeon, Department of Surgery, University of Milan, Via Festa del Perdono 7, Milan 20122, Italy. [ugocioffi5@gmail.com](mailto:ugocioffi5@gmail.com)

### AIMS AND SCOPE

The primary aim of the *World Journal of Gastrointestinal Pharmacology and Therapeutics* (WJGPT, *World J Gastrointest Pharmacol Ther*) is to provide scholars and readers from various fields of gastrointestinal pharmacology and therapeutics with a platform to publish high-quality basic and clinical research articles and communicate their research findings online.

WJGPT mainly publishes articles reporting research results obtained in the field of gastrointestinal pharmacology and therapeutics and covering a wide range of topics including acid-related disorders, functional gastrointestinal disorders, fundamentals of gastrointestinal pharmacology, *etc.*

### INDEXING/ABSTRACTING

The WJGPT is now abstracted and indexed in PubMed, PubMed Central, China National Knowledge Infrastructure (CNKI), China Science and Technology Journal Database (CSTJ), and Superstar Journals Database.

### RESPONSIBLE EDITORS FOR THIS ISSUE

Production Editor: *Ji-Hong Lin*; Production Department Director: *Yun-Xiaoqian Wu*; Editorial Office Director: *Jin-Lei Wang*.

#### NAME OF JOURNAL

*World Journal of Gastrointestinal Pharmacology and Therapeutics*

#### ISSN

ISSN 2150-5349 (online)

#### LAUNCH DATE

May 6, 2010

#### FREQUENCY

Bimonthly

#### EDITORS-IN-CHIEF

Emanuele Sinagra, Sin-Hyeog Im

#### EDITORIAL BOARD MEMBERS

<https://www.wjgnet.com/2150-5349/editorialboard.htm>

#### PUBLICATION DATE

March 18, 2021

#### COPYRIGHT

© 2021 Baishideng Publishing Group Inc

#### INSTRUCTIONS TO AUTHORS

<https://www.wjgnet.com/bpg/gerinfo/204>

#### GUIDELINES FOR ETHICS DOCUMENTS

<https://www.wjgnet.com/bpg/GerInfo/287>

#### GUIDELINES FOR NON-NATIVE SPEAKERS OF ENGLISH

<https://www.wjgnet.com/bpg/gerinfo/240>

#### PUBLICATION ETHICS

<https://www.wjgnet.com/bpg/GerInfo/288>

#### PUBLICATION MISCONDUCT

<https://www.wjgnet.com/bpg/gerinfo/208>

#### ARTICLE PROCESSING CHARGE

<https://www.wjgnet.com/bpg/gerinfo/242>

#### STEPS FOR SUBMITTING MANUSCRIPTS

<https://www.wjgnet.com/bpg/GerInfo/239>

#### ONLINE SUBMISSION

<https://www.f6publishing.com>

© 2021 Baishideng Publishing Group Inc. All rights reserved. 7041 Koll Center Parkway, Suite 160, Pleasanton, CA 94566, USA

E-mail: [bpgoffice@wjgnet.com](mailto:bpgoffice@wjgnet.com) <https://www.wjgnet.com>

# Retrospective Study

## Histopathological evaluation of long-term tenofovir disoproxil fumarate treatment in patients with hepatitis be antigen-negative chronic hepatitis B

Bahri Abayli, Cansu Abaylı, Genco Gencdal

**ORCID number:** Bahri Abayli 0000-0001-5583-8642; Cansu Abaylı 0000-0001-8510-1748; Genco Gencdal 0000-0002-5856-5384.

**Author contributions:** Abayli B and Abaylı C were responsible for the concept, clinical follow-up of patients, writing of article, supervision; Gencdal G was responsible for the concept, study design, organization and coordination of the trial, data analysis, writing of article, supervision; all members of the study team contributed to the management or administration of the trial.

### Institutional review board

**statement:** This retrospective study was organised in accordance with the Helsinki Declaration. Local ethics committee approval was obtained (No. 11.03.2020-52/756).

### Informed consent statement:

Patients were not required to give informed consent to the study because the analysis used anonymous clinical data that were obtained after each patient agreed to treatment by written consent.

**Conflict-of-interest statement:** For all of authors of this manuscript,

**Bahri Abayli, Cansu Abaylı,** Seyhan Devlet Hastanesi, Adana 090, Turkey

**Genco Gencdal,** Department of Gastroenterology, Hepatology and Liver Transplantation, Koc University, School of Medicine, İstanbul 34300, Turkey

**Corresponding author:** Genco Gencdal, MD, Associate Professor, Department of Gastroenterology, Hepatology and Liver Transplantation, Koc University, School of Medicine, özel GOP hastanesi endoskopi birimi gop istanbul, İstanbul 34300, Turkey.  
[gencogencdal@yahoo.co.uk](mailto:gencogencdal@yahoo.co.uk)

## Abstract

### BACKGROUND

Hepatitis B virus is a universal health problem. There are approximately 250 million people living with hepatitis B worldwide, and approximately 600000 of these people die every year due to the virus.

### AIM

To compare the pretreatment and post-treatment histopathological results of patients with hepatitis be antigen (HBeAg)-negative chronic hepatitis B (CHB) who had been receiving tenofovir disoproxil fumarate (TDF) treatment at our clinic for at least 5 years.

### METHODS

Patients with HBeAg-negative CHB who were being treated with TDF (245 mg/d) were included in the study. Liver biopsies of patients before TDF treatment and liver biopsies after 5 years of TDF treatment were retrospectively compared.

### RESULTS

A total of 50 HBeAg-negative CHB patients were included in the study (mean age:  $47.9 \pm 10.4$  years, men: 27.54%). Histological improvement was observed in 78% (39) of the patients after 5 years of treatment. After the 5 years of treatment, the mean Ishak score of the patients was  $1.3 \pm 1.3$ , and the mean histologic activity index score was  $4.1 \pm 2.8$ . A 1.53 point reduction in Ishak fibrosis score was detected after long-term TDF treatment.

### CONCLUSION

there is no potential, personal, financial arrangement (e.g., employment, consultancies, honoraria, stock ownership or options, expert testimony, grants/patents received, and royalties) with a company whose product figures prominently in the submitted manuscript or with a company that makes a competing product.

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

**Open-Access:** This article is an open-access article that was selected by an in-house editor and fully peer-reviewed by external reviewers. It is distributed in accordance with the Creative Commons Attribution NonCommercial (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:** Invited manuscript

**Specialty type:** Gastroenterology and hepatology

**Country/Territory of origin:** Turkey

**Peer-review report's scientific quality classification**

Grade A (Excellent): 0  
Grade B (Very good): 0  
Grade C (Good): C, B  
Grade D (Fair): 0  
Grade E (Poor): 0

**Received:** December 18, 2020

**Peer-review started:** December 18, 2020

**First decision:** January 10, 2021

**Revised:** January 10, 2021

**Accepted:** March 10, 2021

**Article in press:** March 10, 2021

**Published online:** March 18, 2021

**P-Reviewer:** Manrai M

**S-Editor:** Zhang L

**L-Editor:** A

**P-Editor:** Liu JH

Liver biopsies after 5 years of TDF treatment revealed a significant histological response and a regression of the necroinflammatory score compared to pretreatment liver biopsies. To better understand the effects of antiviral treatments on the improvement of liver histology, long-term studies involving larger numbers of patients are needed.

**Key Words:** Hepatitis B; Tenofovir; Chronic hepatitis; Biopsy; Liver; Fibrosis

©The Author(s) 2021. Published by Baishideng Publishing Group Inc. All rights reserved.

**Core Tip:** Hepatitis B virus is a universal health problem. There are approximately 250 million people living with hepatitis B worldwide, and approximately 600000 of these people die every year due to the virus. Viral suppression with treatment can also lead to histological healing. In the current study, we aimed to compare the pretreatment and post-treatment histopathological results of patients with hepatitis be antigen-negative chronic hepatitis B who have been receiving tenofovir disoproxil fumarate (TDF) treatment at our clinic for at least 5 years. Liver biopsies after 5 years of TDF treatment revealed a significant histological response and a regression of the necroinflammatory score compared to pretreatment liver biopsies.

**Citation:** Abayli B, Abaylı C, Gencdal G. Histopathological evaluation of long-term tenofovir disoproxil fumarate treatment in patients with hepatitis be antigen-negative chronic hepatitis B. *World J Gastrointest Pharmacol Ther* 2021; 12(2): 32-39

**URL:** <https://www.wjgnet.com/2150-5349/full/v12/i2/32.htm>

**DOI:** <https://dx.doi.org/10.4292/wjgpt.v12.i2.32>

## INTRODUCTION

Hepatitis B virus (HBV) is a universal health problem. There are approximately 250 million people living with hepatitis B worldwide, and approximately 600000 of these people die every year due to the virus. The treatment of chronic HBV infection depends on many factors including clinical variables (e.g., liver inflammation and/or the presence or absence of cirrhosis), the patient's immunological response to infection [e.g., hepatitis be antigen (HBeAg) status], risk factors for the rapid progression of the disease (e.g., age > 40 and family history of hepatocellular carcinoma), and virological factors (e.g., HBV viral load and genotype). Treatment strategies for chronic hepatitis B (CHB) typically include pegylated interferon or nucleos(t)ide analogues (e.g., entecavir and tenofovir). Interferon is recommended primarily for young patients with compensated liver disease who do not want to undergo long-term treatment<sup>[1-4]</sup>.

For HBeAg-negative CHB patients, the predicted response to treatment is less accurate. Treatment should be started immediately after the diagnosis of HBeAg-negative CHB, because untreated spontaneous remission is rarely seen in this group. The aim of treatment in patients with CHB is to reduce the mortality and morbidity associated with the disease and to increase the quality of life and the lifetime of the patient by preventing complications such as cirrhosis, liver failure and hepatocellular carcinoma that may occur with the progression of the disease. The main goal in therapy is to achieve the long-term suppression of HBV deoxyribonucleic acid (DNA) replication. Viral suppression with treatment can also lead to histological healing. Recent studies have found histological improvements in sexually transmitted disease patients who received antiviral therapy<sup>[2-9]</sup>.

In the current study, we aimed to compare the pretreatment and post-treatment histopathological results of patients with HBeAg-negative CHB who have been receiving tenofovir disoproxil fumarate (TDF) treatment at our clinic for at least 5 years.



## MATERIALS AND METHODS

### *Study design and patient population*

Patients with HBeAg-negative CHB who received TDF treatment (245 mg/d) were included in this study. Patients were included in the study if they were over 18 years, had received antiviral treatment for at least 5 years, came to the outpatient control regularly, underwent liver biopsy before and after treatment, and had available laboratory parameters. Patients were excluded if they were HBeAg-positive, were previously treated with interferon or another antiviral, had a treatment incompatibility, or had been receiving treatment for less than 5 years. Patients were also excluded in cases of human immunodeficiency virus, hepatitis C or hepatitis D co-infection were detected. Elecsys instrument (Roche Diagnostics, Italy) was used to detect HBsAg, anti-HBs, HBeAg and anti-HBe. The real-time polymerase chain reaction AmpliPrep/COBAS TaqMan HBV test 2.0 (Roche Molecular Systems, NJ, United States) was used to quantify HBV DNA. Liver biopsies of patients were evaluated by an experienced pathologist.

### *Definitions of treatment responses*

In our hospital, patients are followed up in the hepatology outpatient clinic according to the 2017 hepatitis B European Association for the Study of the Liver guidelines. According to these guidelines, a decrease in necroinflammatory activity [indicated by a  $\geq 2$  point decrease in the histologic activity index (HAI) or in the Ishak system] without worsening fibrosis compared to the pretreatment histological findings is determined as histological response. The virological response in patients who receive NA treatment is defined as undetectable by a sensitive polymerase chain reaction assay when HBV DNA is below the 10 IU/mL limit of detection. Serological responses for HBsAg are HBsAg loss and the development of anti-HBs. Normalisation of alanine aminotransferase (ALT) levels based on the ULN (40 IU/L) is determined as biochemical response.

### *Statistical analysis*

The data are presented as the mean, median, standard deviation and percentage. All analysis was performed using IBM Statistic Package for Social Science Statistics, V.20.0 (IBM Corp., Armonk, NY, United States). The Kolmogorov-Smirnov test was used to assess normality of quantitative variables. Differences in the variables pre and post treatment were analysed by the Wilcoxon test within groups. All tests were two-tailed, and  $P < 0.05$  was considered to be statistically significant.

### *Ethical approval*

This retrospective study was organised in accordance with the Helsinki Declaration. Local ethics committee approval was obtained (No. 11.03.2020-52/756).

## RESULTS

### *Demographic characteristics of the study population*

A total of 50 HBeAg-negative chronic HBV patients were included in the study. The demographic characteristics of the patients are presented in [Table 1](#). The baseline mean HAI score of the patients was  $7.2 \pm 3.2$ , and the mean baseline Ishak fibrosis score was  $2.2 \pm 1.4$ . The average time from the start of treatment to liver biopsy was  $60.8 \pm 9.7$  wk. All patients were treated with tenofovir disoproxil fumarate (245 mg/d).

### *Histological response*

All biopsy samples were evaluated by a pathologist specialising in liver diseases. Histological improvement was observed in 78% (39) of the patients after 5 years of treatment. After 5 years of treatment, the mean Ishak score of the patients was  $1.3 \pm 1.3$ , and the mean HAI score was  $4.1 \pm 2.8$ . A 1.53 point reduction in Ishak fibrosis scores after long-term treatment was obtained. The long term treatment (60 wk) resulted in most of patients with no or minimal necroinflammation, as assessed by HAI score ([Figure 1](#)), and no or minimal fibrosis, as defined by Ishak classification ([Figure 2](#)).

Eleven patients had advanced fibrosis or cirrhosis (Ishak score  $\geq 4$ ) before the treatment. After long-term treatment, all patients demonstrated at least a 1 point reduction in the Ishak fibrosis score, with a median reduction of 2.9 points from pre-

Table 1 Demographics of the patients before and after treatment

		mean $\pm$ SD	P value
Age		47.9 $\pm$ 10.4	
Male		27 (54%)	
Treatment period (wk)		60.8 $\pm$ 9.7	
AST	BT	82.3 $\pm$ 218.6	< 0.001
	AT	23.7 $\pm$ 14.1	
ALT	BT	74.3 $\pm$ 118.1	< 0.001
	AT	23.7 $\pm$ 14.1	
T.Bil.	BT	0.8 $\pm$ 0.4	0.024
	AT	0.8 $\pm$ 1	
Albumin	BT	3.9 $\pm$ 0.7	> 0.5
	AT	4 $\pm$ 0.8	
Creatine	BT	1 $\pm$ 1.98	> 0.5
	AT	0.8 $\pm$ 0.2	
PLT	BT	208000 $\pm$ 55000	0.007
	AT	238000 $\pm$ 78000	
Ishak score	BT	2.2 $\pm$ 1.4	0.002
	AT	1.3 $\pm$ 1.3	
HAI score	BT	7.2 $\pm$ 3.2	< 0.001
	AT	4.1 $\pm$ 2.8	

ALT: Alanine aminotransferase; AST: Aspartate aminotransferase; TBIL: Total bilirubin; PLT: Platelets; HAI: Histology activity index; BT: Before treatment; AT: After treatment.

treatment values

### Virological response

At the time of post treatment biopsy, 100% of the patients (50/50) had an HBV DNA level < 300 copies/mL; therefore, genotypic testing for resistance was not performed.

### Serological response

All patients were HBeAg-negative. After long-term tenofovir treatment, two of the 50 patients showed HBsAg seroconversion.

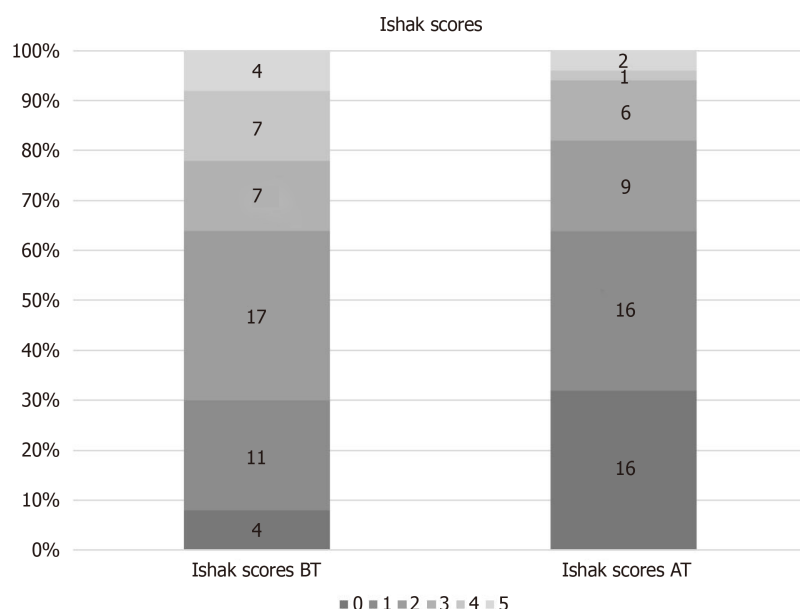
## DISCUSSION

Many studies have shown a statistically significant and consistent correlation between the viral load level or viral load change and the histological grade and biochemical and serological responses over the course of chronic HBV.

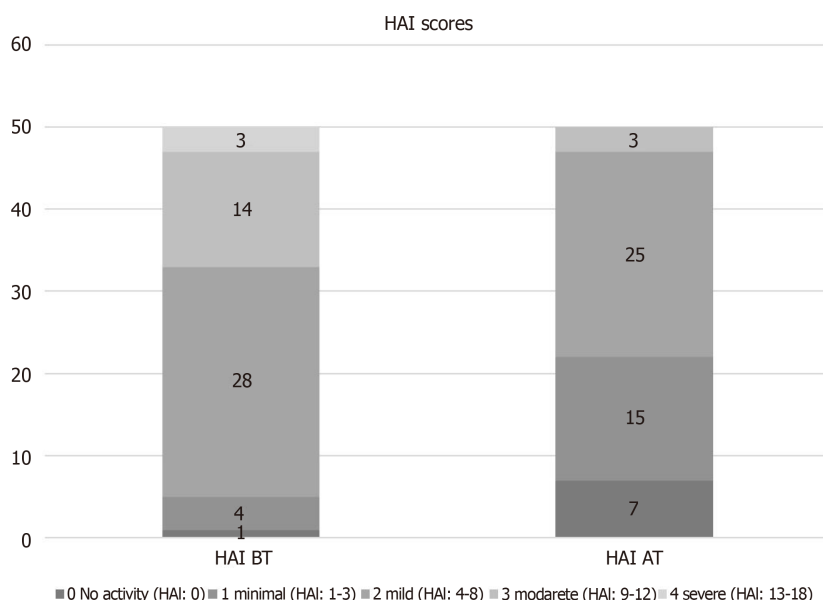
With antiviral therapy, viral replication is brought under control, thereby suppressing necroinflammatory activity in the vast majority of patients. In this way, progressive liver damage is avoided, and there is a decreased risk of hepatocellular carcinoma. Untreated spontaneous recovery is rare in patients with HBeAg-negative CHB. Therefore, after the diagnosis is made, non-cirrhotic patients who meet the criteria should immediately begin treatment and continue it until HBsAg seroconversion occurs. Entecavir and tenofovir are frequently used for this treatment because of their low resistance profile. In patients with hepatitis B with cirrhosis, lifetime treatment is recommended<sup>[2-9]</sup>.

According to recent studies, tenofovir and entecavir are frequently used in cirrhotic and non-cirrhotic patients with CHB, and these drugs have been shown to be safe and effective. These two molecules are frequently used owing to their low resistance





**Figure 1** Ishak scores of the patients before and after treatment. BT: Before treatment; AT: After treatment.



**Figure 2** Histology activity index scores of the patients before and after treatment. HAI: Histologic activity index; BT: Before treatment; AT: After treatment.

profiles<sup>[9-11]</sup>. According to initial studies, tenofovir treatment results in fibrosis improvement, including cirrhosis regression, in the majority of patients. In one study, 51% of the 348 patients with paired biopsies (at baseline and at week 240) showed fibrosis regression in their follow-up biopsy. Interestingly, 71 of 96 patients (74%) with Ishak stage 5 or 6 were found to have no cirrhosis at week 240<sup>[12]</sup>. In their review, Pol *et al*<sup>[13]</sup> reported the safety and efficacy data from two real-world cohorts in the United Kingdom and Europe (362 NA-naïve patients, follow up time: 9-28 mo). In this report, virological suppression was detected in 80%-89% of patients; breakthrough was detected in 2% of patients without any corresponding resistance mutations. HBeAg seroconversion was seen in 7%-18% of patients, and HBsAg loss occurred in 2% of the European cohort. ALT normalisation was detected in 87% of patients by week 30 wk. Pan *et al*<sup>[14]</sup> reported the real-world safety and efficacy of TDF (90 Asian-American patients, 48 wk period). Ten percent of the patients had a prior treatment history with lamivudine or adefovir. The authors detected virological suppression in 82% of patients, HBeAg seroconversion was detected in 12% and ALT normalisation was



detected in 66% of the patients by the end of follow up. No TDF resistance was detected, and the treatment was considered well-tolerated. In the study conducted by Buti *et al*<sup>[15]</sup>, after 5 years of tenofovir treatment, improvement in the Knodell score ( $\geq 1$  point decrease) was found in 93.8% of cirrhotic patients and in 90.8% of non-cirrhotic patients, and no difference was found between the groups. A similar histological response with tenofovir treatment, regardless of the presence of cirrhosis, has been reported<sup>[15]</sup>. In a study, Tatar *et al*<sup>[16]</sup> reported remarkably good HBV DNA suppression, good biochemical response rate and improvement of liver necroinflammation in 52 CHB patients who were treated with TDF (The mean follow-up:  $33 \pm 11$  mo). In accordance with these studies, our results showed HBV DNA to be negative in 100% of the patients after treatment; 38 (76%) had an improved Ishak fibrosis score, 34 (68%) had an improved necroinflammatory score ( $\geq 2$  point improvement of HAI score), and HBsAg seroconversion was detected in 2 patients.

Some studies have reported a strong histological response and fibrosis regression in patients with advanced fibrosis/cirrhosis. In a study with patients who received tenofovir treatment for 5 years, a decline in fibrosis score was observed in 51%. The authors stated that this rate increased to 74% in patients with Ishak fibrosis scores of 5 and 6 before treatment, and histological improvement was greater than 91% in those with Ishak fibrosis scores  $>2$ <sup>[12]</sup>. In our study, 11 of the 50 patients had advanced fibrosis or cirrhosis (Ishak score  $\geq 4$ ) at baseline. After long-term tenofovir therapy, all 11 patients demonstrated at least a 1 point reduction in the Ishak fibrosis score, with a median reduction from baseline of 2.9 points.

This study is limited by its retrospective nature, which did not allow for the initial data to be diversified. Many pretreatment demographic characteristics that could affect the response to treatment could not be verified, and the effect of these factors on the treatment response could not be investigated.

## CONCLUSION

Compared to the initial liver biopsies, the liver biopsies performed at least 5 years after the initiation of TDF treatment revealed a significant histological response and regression of the necroinflammatory score. These promising findings should be verified in a larger population by conducting a multicentre, prospective study.

## ARTICLE HIGHLIGHTS

### Research background

Hepatitis B virus is a universal health problem. There are approximately 250 million people living with hepatitis B worldwide, and approximately 600000 of these people die every year due to the virus. Viral suppression with treatment can also lead to histological healing.

### Research motivation

Recent studies have found histological improvements in sexually transmitted disease patients who received antiviral therapy.

### Research objectives

In the current study, we aimed to compare the pretreatment and post-treatment histopathological results of patients with hepatitis be antigen (HBeAg)-negative chronic hepatitis B (CHB) who have been receiving tenofovir disoproxil fumarate (TDF) treatment at our clinic for at least 5 years.

### Research methods

Patients with HBeAg-negative CHB who were being treated with TDF (245 mg/d) were included in the study. Liver biopsies of patients before TDF treatment and liver biopsies after 5 years of TDF treatment were retrospectively compared.

### Research results

A total of 50 HBeAg-negative CHB patients were included in the study (mean age:  $47.9 \pm 10.4$  years, men: 27.54%). Histological improvement was observed in 78% (39) of the patients after 5 years of treatment. After the 5 years of treatment, the mean Ishak score

of the patients was  $1.3 \pm 1.3$ , and the mean histologic activity index score was  $4.1 \pm 2.8$ . A 1.53 point reduction in Ishak fibrosis score was detected after long-term TDF treatment.

### Research conclusions

Liver biopsies after 5 years of TDF treatment revealed a significant histological response and a regression of the necroinflammatory score compared to pretreatment liver biopsies.

### Research perspectives

These promising findings should be verified in a larger population by conducting a multicentre, prospective study.

## REFERENCES

- 1 **World Health Organization.** Hepatitis B fact sheet. [cited 4 December 2020]. Available from: <http://www.who.int/mediacentre/factsheets/fs204/en/>
- 2 **Terrault NA, Lok ASF, McMahon BJ, Chang KM, Hwang JP, Jonas MM, Brown RS Jr, Bzowej NH, Wong JB.** Update on Prevention, Diagnosis, and Treatment of Chronic Hepatitis B: AASLD 2018 Hepatitis B Guidance. *Clin Liver Dis (Hoboken)* 2018; **12**: 33-34 [PMID: [30988907](#) DOI: [10.1002/cld.728](#)]
- 3 **Sarin SK, Kumar M, Lau GK, Abbas Z, Chan HL, Chen CJ, Chen DS, Chen HL, Chen PJ, Chien RN, Dokmeci AK, Gane E, Hou JL, Jafri W, Jia J, Kim JH, Lai CL, Lee HC, Lim SG, Liu CJ, Locarnini S, Al Mahtab M, Mohamed R, Omata M, Park J, Piratvisuth T, Sharma BC, Sollano J, Wang FS, Wei L, Yuen MF, Zheng SS, Kao JH.** Asian-Pacific clinical practice guidelines on the management of hepatitis B: a 2015 update. *Hepatol Int* 2016; **10**: 1-98 [PMID: [26563120](#) DOI: [10.1007/s12072-015-9675-4](#)]
- 4 **European Association for the Study of the Liver.** EASL 2017 Clinical Practice Guidelines on the management of hepatitis B virus infection. *J Hepatol* 2017; **67**: 370-398 [PMID: [28427875](#) DOI: [10.1016/j.jhep.2017.03.021](#)]
- 5 **Gish RG, Given BD, Lai CL, Locarnini SA, Lau JY, Lewis DL, Schluep T.** Chronic hepatitis B: Virology, natural history, current management and a glimpse at future opportunities. *Antiviral Res* 2015; **121**: 47-58 [PMID: [26092643](#) DOI: [10.1016/j.antiviral.2015.06.008](#)]
- 6 **Iloeje UH, Yang HI, Su J, Jen CL, You SL, Chen CJ;** Risk Evaluation of Viral Load Elevation and Associated Liver Disease/Cancer-In HBV (the REVEAL-HBV) Study Group. Predicting cirrhosis risk based on the level of circulating hepatitis B viral load. *Gastroenterology* 2006; **130**: 678-686 [PMID: [16530509](#) DOI: [10.1053/j.gastro.2005.11.016](#)]
- 7 **Sanai FM, Helmy A, Bzeizi KI, Babatin MA, Al-Qahtani A, Al-Ashgar HA, Al-Mdani AS, Al-Akwaa A, Almutharea S, Khan MQ, Alghamdi AS, Farah T, Al-Hamoudi W, Saadeh M, Abdo AA.** Discriminant value of serum HBV DNA levels as predictors of liver fibrosis in chronic hepatitis B. *J Viral Hepat* 2011; **18**: e217-e225 [PMID: [21692936](#) DOI: [10.1111/j.1365-2893.2011.01437.x](#)]
- 8 **Dienstag JL, Goldin RD, Heathcote EJ, Hann HW, Woessner SL, Stephenson SL, Gardner S, Gray DF, Schiff ER.** Histological outcome during long-term lamivudine therapy. *Gastroenterology* 2003; **124**: 105-117 [PMID: [12512035](#) DOI: [10.1053/gast.2003.50013](#)]
- 9 **Schiff ER, Lee SS, Chao YC, Kew Yoon S, Bessone F, Wu SS, Kryczka W, Lurie Y, Gadano A, Kitis G, Beebe S, Xu D, Tang H, Iloeje U.** Long-term treatment with entecavir induces reversal of advanced fibrosis or cirrhosis in patients with chronic hepatitis B. *Clin Gastroenterol Hepatol* 2011; **9**: 274-276 [PMID: [21145419](#) DOI: [10.1016/j.cgh.2010.11.040](#)]
- 10 **Chang TT, Liaw YF, Wu SS, Schiff E, Han KH, Lai CL, Safadi R, Lee SS, Halota W, Goodman Z, Chi YC, Zhang H, Hindes R, Iloeje U, Beebe S, Kreter B.** Long-term entecavir therapy results in the reversal of fibrosis/cirrhosis and continued histological improvement in patients with chronic hepatitis B. *Hepatology* 2010; **52**: 886-893 [PMID: [20683932](#) DOI: [10.1002/hep.23785](#)]
- 11 **Papachrysos N, Hytioglou P, Papalavrentios L, Sinakos E, Kouvelis I, Akriviadis E.** Antiviral therapy leads to histological improvement of HBeAg-negative chronic hepatitis B patients. *Ann Gastroenterol* 2015; **28**: 374-378 [PMID: [26126929](#)]
- 12 **Marcellin P, Gane E, Buti M, Afdhal N, Sievert W, Jacobson IM, Washington MK, Germanidis G, Flaherty JF, Aguilar Schall R, Bornstein JD, Kittrinos KM, Subramanian GM, McHutchison JG, Heathcote EJ.** Regression of cirrhosis during treatment with tenofovir disoproxil fumarate for chronic hepatitis B: a 5-year open-label follow-up study. *Lancet* 2013; **381**: 468-475 [PMID: [23234725](#) DOI: [10.1016/S0140-6736\(12\)61425-1](#)]
- 13 **Pol S, Lampertico P.** First-line treatment of chronic hepatitis B with entecavir or tenofovir in 'real-life' settings: from clinical trials to clinical practice. *J Viral Hepat* 2012; **19**: 377-386 [PMID: [22571899](#) DOI: [10.1111/j.1365-2893.2012.01602.x](#)]
- 14 **Pan CQ, Trinh H, Yao A, Bae H, Lou L, Chan S;** Study 123 Group. Efficacy and safety of tenofovir disoproxil fumarate in Asian-Americans with chronic hepatitis B in community settings. *PLoS One* 2014; **9**: e89789 [PMID: [24594870](#) DOI: [10.1371/journal.pone.0089789](#)]

- 15 **Buti M**, Fung S, Gane E, Afdhal NH, Flisiak R, Gurel S, Flaherty JF, Martins EB, Yee LJ, Dinh P, Bornstein JD, Mani Subramanian G, Janssen HL, George J, Marcellin P. Long-term clinical outcomes in cirrhotic chronic hepatitis B patients treated with tenofovir disoproxil fumarate for up to 5 years. *Hepatol Int* 2015; **9**: 243-250 [PMID: [25788199](#) DOI: [10.1007/s12072-015-9614-4](#)]
- 16 **Tatar B**, Gül S, Köse Ş, Pala E. Long-Term Effects of Tenofovir on Liver Histopathology in Patients with Chronic Viral Hepatitis B Infection. *Turk Patoloji Derg* 2020; **1**: 154-158 [PMID: [32149362](#) DOI: [10.5146/tjpath.2020.01478](#)]



Published by **Baishideng Publishing Group Inc**  
7041 Koll Center Parkway, Suite 160, Pleasanton, CA 94566, USA

**Telephone:** +1-925-3991568

**E-mail:** [bpgoffice@wjgnet.com](mailto:bpgoffice@wjgnet.com)

**Help Desk:** <https://www.f6publishing.com/helpdesk>

<https://www.wjgnet.com>

