

# Efficacy of Initial Treatment with Clevudine in Naïve Patients with Chronic Hepatitis B

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**Background/Aims:** Clevudine, a pyrimidine nucleoside analogue, has potent antiviral effects in patients with chronic viral hepatitis B (CHB). We report the efficacy of initial treatment with clevudine in naïve patients with CHB living in Daejeon and Chungcheong Province, South Korea.

**Methods:** One hundred five adults with CHB were administered 30 mg of clevudine per day for an average of 51 weeks. We evaluated viral markers and liver biochemistry retrospectively every 3 months.

**Results:** Alanine aminotransferase (ALT), aspartate aminotransferase (AST), and hepatitis B virus (HBV) DNA before the treatment were  $184 \pm 188$  IU/L,  $150 \pm 138$  IU/L, and  $7.1 \pm 1.2$  log copies/mL, respectively. Undetectable rates ( $< 60$  IU/mL) of DNA were 36.2%, 68.9%, 83.6%, 76.2%, and 75.8% at 12, 24, 36, 48, and 60 weeks, respectively. Seroconversion rates were 9.1%, 13.6%, 24.6%, 26.5%, and 26.1% and ALT normalization rates were 64.5%, 78.1%, 87.9%, 90.0% at 12, 24, 36, and 48 weeks, respectively. Six patients (5.7%) had a viral breakthrough.

**Conclusions:** Clevudine is a useful drug in the initial treatment of patients with CHB, with a potent antiviral effect and low incidence of viral breakthrough. (*Korean J Intern Med* 2010;25:372-376)

**Keywords:** Clevudine; Drug resistance; HBV seroconversion; Hepatitis B, chronic

## INTRODUCTION

Hepatitis B virus (HBV) infection, which leads to chronic hepatitis and cirrhosis, may be responsible for life-threatening liver disease in some individuals, with 350 million carriers worldwide [1]. Substantial advances in oral nucleoside/nucleotide analogs have been made for treating chronic hepatitis B (CHB) in past decades. Nucleoside or nucleotide analogues competitively inhibit replication of HBV DNA through DNA polymerase. Lamivudine (LAM) is a first-generation oral nucleoside

analogue for treating CHB. Previous studies in naïve patients found a cumulative LAM resistance incidence of 70 to 80% after 4 to 5 years of therapy [2,3]. Entecavir shows very low resistance and a potent antiviral effect in naïve patients with CHB. Although there is no evidence of carcinogenesis in human studies, this drug has produced some tumors such as adenoma of the lung, neurogenic tumors, and hepatocellular carcinoma in mice [4]. Clevudine is a nucleoside analogue of the unnatural L-configuration, which has potent antiviral activity against HBV [1,5]. Studies about the efficacy of initial treatment

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with clevudine in naïve chronic hepatitis B patients have been reported [1,6-11] and indicate that clevudine is very useful for treating CHB in the short term. However, these studies had short-term follow-ups of 12 or 24 weeks [1,6-11]. More long-term follow-up period studies are needed for evaluating the efficacy of clevudine treatment in patients with CHB. We prescribed clevudine to 105 naïve patients with CHB living in Daejeon and Chungcheong Province in South Korea for an average of 51 weeks.

## METHODS

### Study design

This was a retrospective study and was conducted at six medical centers in Daejeon and Chungcheong Province, South Korea. Eligible patients were diagnosed with CHB with or without hepatitis B e antigen (HBeAg), HBV DNA levels above  $1.0 \times 10^5$  copies/mL,  $> 2.0 \times 10^4$  IU/mL or  $> 0.4$  pg/mL, and alanine aminotransferase (ALT) levels  $> 80$  IU/L. All of the patients were prescribed clevudine for more than 6 months. The exclusion criteria were malignancy, alcoholism ( $> 20$  g/day), severe fatty liver on ultrasonography, and a medication history of nucleoside or nucleotide analogues or interferon. The serum HBV DNA level was measured using the DNA PCR hybridization method (Roche, Indianapolis, IN, USA). The lower limit of HBV DNA detection was 300 copies/mL. Viral response was defined as  $< 300$  copies/mL of HBV DNA. We defined viral breakthrough as a ten times rebound of HBV DNA copies compared with previous copies during treatment. Enrolled patients

**Table 1. Baseline patient characteristics before treatment**

No.	105
Mean age, yr	42.7 ± 11.9
Male / Female	65 : 40
CHB	90
LC	15 (14.3)
Follow-up period, wk	51.1 (24 - 72)
AST, IU/L	150 ± 138
ALT, IU/L	184 ± 188
HBV DNA, log copies/mL	7.1 ± 1.2
eAg positive	71

Values are presented as number (%) or mean ± SD.

CHB, chronic hepatitis B; LC, liver cirrhosis; AST, aspartate aminotransferase; ALT, alanine aminotransferase; HBV, hepatitis B virus; eAg, e antigen.

were prescribed 30 mg clevudine daily. We checked liver biochemistry, HBeAg/Ab, and HBV DNA every 12 weeks.

### Statistical analysis

SPSS version 10.2 (SPSS Inc., Chicago, IL, USA) was used. Fisher's exact test and the chi-square test were used to compare viral response and serum ALT normalized rate according to HBeAg status.

**Table 2. Viral response rate of hepatitis B virus DNA and a comparison of HBeAg-positive and HBeAg-negative patients**

	Total virologic response	HBeAg (+)	HBeAg (-)	<i>p</i> value <sup>a</sup>
Initial DNA, log copies/mL	7.1 ± 1.2	7.4 ± 1.1	6.5 ± 1.0	
12 wk	25/69 (36.2)	10/43 (23.3)	15/26 (67.7)	0.004
24 wk	64/93 (68.9)	37/62 (59.7)	27/31 (87.1)	0.007
36 wk	61/73 (83.6)	38/48 (79.2)	23/25 (92.0)	0.160
48 wk	48/63 (76.2)	32/44 (72.3)	16/19 (84.2)	0.326
60 wk	25/33 (75.8)	17/26 (65.4)	8/9 (88.9)	0.179

Values are presented as number (%).

HBeAg, hepatitis B e antigen.

<sup>a</sup> *p* values are calculated for comparison of HBeAg-positive and HBeAg-negative patients.

**Table 3. Serum alanine aminotransferase (ALT) normalization rate**

Weeks	ALT normalization rate			p value
	Total	HBeAg (+)	HBeAg (-)	
12	49/76 (64.5)	45/71 (63.4)	4/5 (80.0)	NS
24	82/105 (78.1)	53/71 (74.6)	29/34 (85.3)	NS
36	80/91 (87.9)	55/63 (87.3)	25/28 (89.3)	NS
48	71/80 (90.0)	50/57 (87.7)	21/23 (91.3)	NS

Values are presented as number (%).

HBeAg, hepatitis B e antigen; NS, not significant.

## RESULTS

### Patient characteristics

A total of 105 patients including 15 (14.3%) patients with liver cirrhosis were enrolled at six medical centers from February 2007 to August 2008. Seventy-one patients were HBeAg positive, and 34 patients were negative. The mean age was 42.7 years, and the mean follow-up period was 51.1 weeks. Median ALT at baseline was 184 IU/L, and median baseline HBV DNA was 7.1 log<sub>10</sub> copies/mL (Table 1).

### Viral response

Viral response rates in all patients were 36.2%, 68.9%, 83.6%, 76.2%, and 75.8% at 12, 24, 36, 48, and 60 weeks, respectively. Viral response rates in HBeAg-positive patients were 23.3%, 59.7%, 79.2%, 72.3%, and 65.4%, and those in HBeAg-negative patients were 67.7%, 87.1%, 92.0%, 84.2%, and 88.9% at the corresponding weeks. The rates in HBeAg-positive patients were significantly lower at 12 and 24 weeks compared with those in HBeAg-negative patients. No significant difference was observed after 24 wks (Table 2).

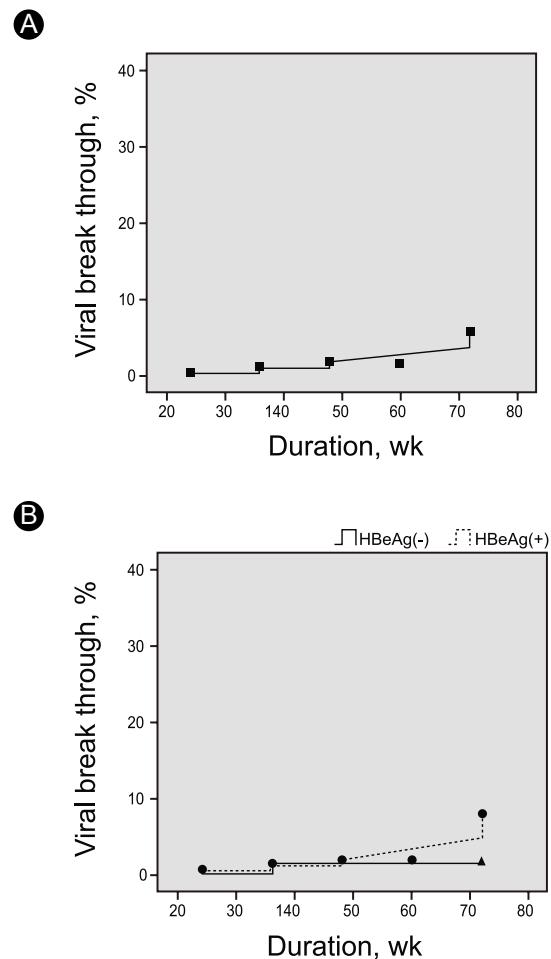
### Serum ALT normalization rate

The serum ALT normalization rates in all patients were 78.1% and 90.0% at 24 and 48 weeks, respectively. The serum ALT normalization rates in HBeAg-positive versus HBeAg-negative patients were 63.4% vs. 80%, 74.6% vs. 85.3%, 87.3% vs. 89.3%, and 87.7% vs. 91.3% at 12, 24, 36, and 48 weeks, respectively (Table 3).

### HBeAg loss and seroconversion rate

A total of 71e HBeAg-positive patients were enrolled. Among these patients, the HBeAg loss was 20.3%, 30.6%,

and 43.3% at 24, 48, and 60 weeks, respectively, and the HBeAg seroconversion rates were 13.6%, 26.5%, and 26.1%, at 24, 48, and 60 weeks (Table 4).



**Figure 1.** Incidence rate of viral breakthrough according to treatment period. (A) Viral breakthrough rate of total patients. (B) Viral breakthrough rate of HBeAg (+) and HBeAg (-) patients.

**Table 4. HBeAg loss rate and seroconversion rate in HBeAg-positive patients**

Weeks	No. of patients	HBeAg loss	Seroconversion
12	44	6 (13.6)	4 (9.1)
24	59	12 (20.3)	8 (13.6)
36	57	18 (31.6)	14 (24.6)
48	49	15 (30.6)	13 (26.5)
60	23	10 (43.3)	6 (26.1)

Values are presented as number (%).  
HBeAg, hepatitis B e antigen.

### Viral breakthrough

The cumulative incidence rate of viral breakthrough in the HBeAg-positive versus HBeAg-negative patients was 1.4% vs. 0%, 3.3% vs. 3.4%, and 6.7% vs. 4.1% at 24, 36, and 48 weeks, respectively (Fig. 1). Six patients underwent treatment for viral breakthrough. Five cases were identified at 48 weeks, and one case at week 60. Five of these six patients underwent a tyrosine-methionine-aspartate-aspartate mutation (YMDD) examination, and four patients were positive (Table 5).

## DISCUSSION

Clevudine is a nucleoside analog with an unnatural L-configuration. The mechanism by which clevudine exerts its antiviral effects is believed to be by the intracellular formation of the monophosphate, diphosphate, and triphosphate forms. Clevudine enters cells through both

facilitated nucleoside transport and nonfacilitated passive diffusion and a substrate of three intracellular kinases is responsible for its phosphorylation [12].

When 21 patients received 30 mg clevudine for 24 weeks and were followed up for another 24 weeks off therapy, the clevudine treatment was well tolerated and resulted in more potent antiviral activity and a higher ALT normalization rate than a 12-week treatment, with durable efficacy at week 24 of therapy [1]. However, the experimental period in this study was less than 1 year. In another larger-scale study, 243 patients with HBeAg-positive CHB were randomized (3 : 1) to receive 30 mg clevudine once daily (n = 182) or placebo (n = 61) for 24 weeks. Patients were followed for a further 24 weeks off therapy. In that study, the 24-week clevudine therapy was well tolerated and showed a potent and sustained antiviral effect without evidence of viral resistance during the treatment period [9]. Here, we administered clevudine to 105 patients with CHB for about 1 year. As in other studies, clevudine showed powerful antiviral effects in this study [13]. With a limit of detection of < 300 copies/mL, the viral response rate was 76.2% (HBeAg-positive patients, 72.3%; HBeAg-negative patients, 84.2%) at 48 weeks. This result was remarkable enough to be compared with entecavir and tenofovir, which have negative seroconversion rates of 67% and 76%, respectively, after a 1-year treatment, but a direct comparison is difficult because of differences in the study methods and patient characteristics [9,10].

Viral response was highest at 36 weeks and subsequently decreased because some patients experienced viral breakthrough, and the HBV DNA PCR test at 48 or 60 weeks was not performed in some patients who had a

**Table 5. Clinical features of patients with viral breakthrough**

Age/Gender	DNA at baseline, copies/mL	eAg at baseline	ALT at baseline, U/L	Time until VBT, wk	YMDD mutation
31/M	$5.3 \times 10^8$	+	262	36	Negative
50/M	$7.0 \times 10^6$	+	78	48	Positive
60/M	$3.1 \times 10^7$	+	80	24	Positive
53/M	$4.0 \times 10^5$	-	114	36	Positive
48/F	$1.3 \times 10^7$	+	311	60	Positive
50/M	$1.5 \times 10^8$	+	636	40	Not done

eAg, e antigen; ALT, alanine aminotransferase; VBT, viral breakthrough; YMDD, tyrosine-methionine-aspartate-aspartate mutation.

good viral response.

The serum ALT normalization rate of all patients was 78.1% (HBeAg-positive patients, 74.6%; HBeAg-negative patients, 85.3%) and 90.0% (HBeAg-positive patients, 87.7%; HBeAg-negative patients, 91.3%) at 24 and 48 weeks, respectively. This result also shows similar efficacy to other antiviral agents. Approximately 6% of all patients with CHB showed viral breakthrough at 48 weeks. The viral breakthrough incidence rates were 2.9% and 7.0% for HBeAg-negative and -positive patients respectively; however, the difference was not statistically significant.

The cumulative viral breakthrough rate until week 48 was 6.3% (HBeAg-positive group, 7.0% [4/57]; HBeAg-negative group, 4.5% [1/22]) (Fig. 1).

Clevudine had powerful antiviral effects and had relatively lower viral breakthrough incidence rate compared with lamivudine (16-32% after 12 months); it is expected to replace lamivudine therapy [14]. When we performed the YMDD mutation test on five of six patients with viral breakthrough, the YMDD mutation was found in four patients. This suggested that clevudine cross-reacted with lamivudine. In particular, although our enrolled number of patients was insufficient to reach a definitive conclusion, clevudine appears to have been more effective for the HBeAg-negative group. Noticeably, although this study was not controlled, the HBeAg seroconversion rate at 48 weeks was 26.5% (13/49), which was relatively high compared with that for lamivudine (16-18%) [15]. A limitation of this study was that the examinations were not performed at regular intervals at all times, as it was a retrospective study and had a relatively short-term follow-up period of 1 year compared with other studies on medications that have been more fully researched. Furthermore, although no patient stopped clevudine because of myositis or others side effects, we did not measure serum creatine phosphokinase regularly or assess side effects. Despite these limitations, the results suggest that clevudine has the potential to be an effective therapeutic medication for patients with CHB; however, longer-term research will be needed for further evaluation.

#### Conflict of interest

No potential conflict of interest relevant to this article was reported.

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