

Authorship:

Cheng-Yuan Peng, M.D., Ph.D.

Division of Hepatogastroenterology, Department of Internal Medicine,
China Medical University Hospital
No. 2, Yuh-Der Rd, Taichung, 40447, Taiwan

Rong-Nan Chien, M.D.

Liver Research Unit
Chang Gung Memorial Hospital and University College of Medicine, Keelung
No.222, Maijin Rd., Anle Dist., Keelung City, Taiwan 204

Jia-Horng Kao, M.D., Ph.D.

Graduate Institute of Clinical Medicine, National Taiwan University College of Medicine
No. 7, Chung-Shan South Road, Taipei 100 Taiwan

Tsung-Hui Hu, M.D., Ph.D.

Division of Hepato-Gastroenterology, Department of Internal Medicine,
Kaohsiung Chang Gung Memorial Hospital and Chang Gung University College of Medicine
No 123, Ta-Pei Road, Niao Sung Dist., Kaohsiung 833, Taiwan

Chun-Che Lin, M.D., Ph.D.

Division of Gastroenterology and Hepatology, Department of Internal Medicine,
Chung Shan Medical University Hospital
N0.110, Sec. 1, Chien-Kuo N.Road, Taichung 402, Taiwan

Chi-Tan Hu, M.D., Ph.D.

Division of Gastroenterology, Department of Internal Medicine and Research Center for Hepatology,
Buddhist Tzu Chi General Hospital
Graduate Institute of Clinical Medicine, School of Medicine, Tzu Chi University, Hualien, Taiwan
No. 707, Sec. 3, Chung-Yang Rd., Hualien 970, Taiwan

Chi-Yi Chen, M.D.

Liver Unit of Internal Medical Department, Chia-Yi Christian Hospital
No. 539, Chung-Hsiao Rd., Chia-Yi City 539, Taiwan

Tsai-Yuan Hsieh, M.D., Ph.D.

Division of Gastroenterology, Department of Internal Medicine,
Tri-Service General Hospital, National Defense Medical Center
No.325, Sec.2, Cheng-Kung Rd., Neihu District, Taipei City 114, Taiwan

Han-Chieh Lin, M.D., Ph.D.

Division of Gastroenterology, Department of Medicine,
Taipei Veterans General Hospital
No.201, Sec. 2, Shipai Rd., Beitou District, Taipei City 112, Taiwan

Wan-Long Chuang, M.D. (Correspondence)

Affiliation: Department of Internal Medicine, Kaohsiung Medical University Hospital
Address: No.100, Tzyou 1st Road Kaohsiung 807, Taiwan
Tel: +886-7-3121101 #7475
Email: waloeh@kmu.edu.tw

Taiwan NA – Registry Group

Title:

Interim analysis of initial NUC treatment among NUC-naïve chronic hepatitis B patients in Taiwan

Background/Aims: Current guidelines recommend potent nucleos(t)ide analogues (NUCs) with a high genetic barrier to resistance as the initial treatment for chronic hepatitis B (CHB). This study aims to evaluate the rate of sustained NUC treatment (entecavir [ETV], lamivudine [LVD], or telbivudine [LdT]) and assess treatment compliance in the outpatient setting.

Methods: NUC-naïve CHB patients who received ETV, LVD or LdT monotherapy from August 2008 to July 2009 were randomly selected from 33 hospitals across Taiwan. Demographic and clinical data were retrieved from medical records. Enrolled patients were retro-prospectively followed for 144 weeks and herein we presented the interim results as of September 2012.

Results: There were 417 male and 161 female patients (mean age, 43.5 years). The proportion of patients who had sustained treatment with ETV, LVD and LdT for 144 weeks without any modification was: 98.9%, 80.0% and 88.9%, respectively. Table summarizes the rates of modification of initial treatments and reasons for modification. Figure shows the time to modification. Modified compliance rates (mean days/year [%] on therapy) from Weeks 96 to 144 were 96.8% (ETV), 97.9% (LVD) and 97.1% (LdT).

Conclusions: Our data show a high rate of sustained treatment and an acceptable rate of compliance in Taiwanese CHB patients treated with 144-week ETV.

Table: Rates of Modification of Initial Antiviral Treatment and Reasons for Modification

	Initial Antiviral Treatment			Total (N=578)
	ETV (N=463)	LVD (N=49)	LdT (N=66)	
Patients with initial treatment modification	41 (8.9%)	25 (51.0%)	26 (39.4%)	92 (15.9%)
Reasons for modification*				
Antiviral resistance	5 (12.2%)	10 (40.0%)	14 (53.8%)	29 (31.5%)
Intolerance to initial treatment	2 (4.9%)	3 (12.0%)	0	5 (5.4%)
Financial embarrassment	3 (7.3%)	1 (4.0%)	0	4 (4.3%)
Others	31 (75.6%)	11 (44.0%)	12 (46.2%)	54 (58.7%)

*Percentages were calculated based on the number of patients with initial treatment modification.

Figure: Time to initial treatment modification in Taiwanese CHB patients treated with ETV, LVD or LdT

