The Naturally Occurring YMDD Mutation among Patients Chronically Infected HBV and Untreated with Lamivudine: A Systematic Review and Meta-Analysis

Youwen Tan1,2,*, Kegin Ding1,*, Jing Su1,*, Xuan Trinh3, Zhihang Peng1, Yuhua Gong2, Li Chen2, Qian Cui1, Na Lei1, Xin Chen1, Rongbin Yu1,*

1 Department of Epidemiology and Biostatistics, School of Public Health, Nanjing Medical University, Nanjing, China, 2 Department of Hepatosis, The Third People’s Hospital of Zhenjiang City, Zhenjiang, China, 3 Department of Pharmacology, University of Gothenburg, Göteborg, Sweden

Abstract

**Background:** Several recent reports have demonstrated that tyrosine (Y)-methionine (M)-aspartic acid (D)-aspartic acid (D) (YMDD) motif mutations can naturally occur in chronic HBV patients without antiviral treatment such as lamivudine therapy. This paper aims to assess the overall spontaneous incidence and related risk factors of YMDD-motif mutations among lamivudine-naive chronic HBV carriers, so as to provide some clue for clinical treatment of hepatitis B.

**Methodology/Principal Findings:** Chinese and English literatures were searched for studies reporting natural YMDD mutations among untreated chronic HBV patients from 2001 to 2010. The incidence estimates were summarized and analyzed by meta-analyses. Forty-seven eligible articles from eight countries were selected in this review (13 in English and 34 in Chinese). The pooled incidence of YMDD-motif mutation among untreated chronic HBV patients from eight countries was 12.21% (95% CI: 9.69%–14.95%). China had an incidence of 13.38% (95% CI: 10.90%–16.07%) and seven other countries had an incidence of 9.90% (95% CI: 3.28%–19.55%), respectively. Lamivudine therapy would increase the risk of mutations 5.23 times higher than the untreated patients. A higher HBV DNA copy number was associated with increased incidence of natural YMDD mutation. No significant difference was found in YMDD mutation incidence between groups of different gender, age, HBeAg status, patients’ ALT (alanine aminotransferase) level, and between the groups of HBV genotype B and C.

**Conclusions:** The YMDD-motif mutations can occur spontaneously with a relatively high incidence in CHB patients untreated with lamivudine. These mutations might be the consequence of accumulated base mismatch due to the nature of viral polymerase. More fundamental and clinical studies are needed to clarify the influence of YMDD mutations in hepatitis B progression and antiviral treatment.

Citation: Tan Y, Ding K, Su J, Trinh X, Peng Z, et al. (2012) The Naturally Occurring YMDD Mutation among Patients Chronically Infected HBV and Untreated with Lamivudine: A Systematic Review and Meta-Analysis. PLoS ONE 7(3): e32789. doi:10.1371/journal.pone.0032789

Editor: Joerg F. Schlaak, University Hospital of Essen, Germany

Received October 19, 2011; Accepted January 31, 2012; Published March 27, 2012

Copyright: © 2012 Tan et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

Funding: This work was supported by the National Science and Technology Major Project (2012ZX10001-001, 2011ZX10004-902) (http://www.nmp.gov.cn/), the Priority Academic Program Development of Jiangsu Higher Education Institutions (PAPD) (http://202.119.175.131/pages/index.aspx), the Natural Science Foundation Key Project of Jiangsu Province, China (BK2010079) (http://www.jspdf.cn/) and the Social Development Project of Zhenjiang City (SH2009016) (http://kjj.zhenjiang.gov.cn/). The funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.

Competing Interests: The authors have declared that no competing interests exist.
* E-mail: rongbinyu@njmu.edu.cn

These authors contributed equally to this work.

Introduction

Hepatitis B virus (HBV) infection has long been a serious public health problem around the world. HBV is one of the main causes of acute and chronic hepatitis in humans. Nearly 400 million people are infected chronically with HBV worldwide [1]. It has been demonstrated that chronic infection with this virus is linked to the development of cirrhosis and hepatocellular carcinoma, accounting for 0.5–0.75 million deaths per year.

Antiretroviral treatment is the main clinical treatment of chronic hepatitis B (CHB). Lamivudine, a nucleoside analogue, has been widely used because of its high effectiveness. However, long-term use of lamivudine may lead to the emergence of lamivudine-resistance in some HBV cases. The resistance generation is closely associated with mutations in the highly conserved YMDD motif, which is in the catalytic domain C of viral DNA polymerase. The YMDD motif has an amino acid sequence of tyrosine (Y)-methionine (M)-aspartic acid (D)-aspartic acid (D) [2] and is both the binding and functional site of lamivudine.

Recent reports revealed that besides the mutations caused by lamivudine therapy, some YMDD mutations can occur spontaneously in lamivudine-untreated CHB patients. Because HBV reverse transcriptase has no proofreading activity during the replication process, mutations can naturally occur due to random nucleotide misincorporation [3]. The most common substitutions are methionine at amino acid position-204 to either isoleucine (rtM204I, YIDD mutant) or valine (rtM204V, YVDD mutant) [4,5]. Many studies have reported the incidence and characteristic of spontaneous YMDD-motif mutation in lamivudine-naive CHB patients. However, the reported incidences were quite different.
(0% to 31.58%) or even contradict with each other. This paper reviews the available publications, aiming to comprehensively provide not only the incidence but also associated factors of natural YMDD-motif mutation among lamivudine-untreated CHB patients. The results should provide scientific evidences for clinical treatment and future research of YMDD mutation.

Materials and Methods

Search strategy

A number of sources were used to access data for this review. Published articles were searched in PubMed [http://www.ncbi.nlm.nih.gov/pubmed] and Embase [http://www.embase.com/search] database from January 2001 to December 2010 using the search terms: YMDD, lamivudine, HBV OR hepatitis B, spontaneous mutation OR natural mutation, untreated, and naive. The two Chinese databases: China National Knowledge Infrastructure (CNKI) [http://epub.edu.cnki.net/] and WanFang [http://g.wanfangdata.com.cn/] were also searched for articles related to HBV and YMDD motif mutation from January 2001 to December 2010. The reference lists of relevant articles were also searched manually for complement. This review was conducted and reported according to the PRISMA (Preferred Reporting Items for Systematic Reviews and Meta-Analyses) Statement issued in 2009 (Table S1) [6].

Selection criteria

All the potentially relevant papers were reviewed independently by two investigators through assessing the eligibility of each article and useful data were extracted with standardized data-extraction forms. Studies published in English and Chinese were eligible if they fulfilled the following criteria: (1) studies in the mentioned four databases with full text, or with available data in abstract; (2) study objects contain CHB patients with NO treatment of lamivudine or other antiviral drugs; (3) studies report the rates of YMDD mutation among lamivudine-untreated CHB cases. The exclusion criteria were: (1) studies without specific sample origins; (2) studies with overlapping time intervals of the same sample collection; (3) study objects were treated with lamivudine; (4) study objects were patients with acute hepatitis B; (5) subset of a published article by the same authors; (6) studies present unclear data or obviously paradoxical data.

Data extraction and quality assessment

The following information, though some studies did not contain all of them, were extracted from the literatures: first author’s name, publication date, journal title, study area, sample size (the number of CHB patients untreated with lamivudine), the number of cases with YMDD mutation, the method of detecting YMDD mutation, gender and age of study objects, the status of HBsAg (positive or negative), the amount of serum HBV DNA, ALT (alanine aminotransferase) level, and HBV genotype. All the numerical values of HBV DNA copies were converted to logarithm. Two investigators (Keqin Ding and Yuhua Gong) extracted and collected data independently using a standardized data-extraction protocol. Any discrepancy was settled by consensus of the whole team.

Statistical analysis

In this study, random effect model or fixed effect model was used for meta-analysis, according to the heterogeneity between studies which was tested with the Q test (P<0.10 was considered indicative of statistically significant heterogeneity) and the I² statistic (values of 25%, 50% and 75% are considered to represent low, medium and high heterogeneity respectively). The fixed effect model was used when there was no significant heterogeneity (I²<50%); otherwise the random effect model was used. P values were calculated by χ² tests. All the reported P values are two-sided and P values less than 0.05 were regarded as statistically significant for all included studies.

Publication bias was examined by a funnel plot of log OR (odds ratio) against its standard error using Begg’s test, and the degree of asymmetry was tested statistically using Egger’s unweighed regression asymmetry test. Publication bias exists if the rank correlation coefficient from Begg’s test is close to 1, or the slope parameter from Egger regression is high.

Data manipulation and statistical analyses were undertaken using the Statistical Software Package (STATA) 10.0 (STATA Corporation, College Station, TX, USA, 2009) and Review Manager 4.2 (The Cochrane Collaboration, Oxford, United Kingdom, 2003).

Results

According to the literature search strategies, a total of 202 studies (54 studies in PubMed and Embase, and 148 studies in CNKI and Wanfang database) were identified, and 155 studies were excluded based on the inclusion and exclusion criteria. Finally there were 47 studies adopted in this study (13 studies in English and 34 studies in Chinese) as showed in Figure 1 and Figure S1 [6].

The studies of our review were involved in the following 18 provinces and municipalities in China and seven other countries: Central China (Hubei, Hunan), North China (Beijing, Liaoning, Jilin, Shan‘xi), South China (Hainan, Guangdong, Guangxi), East China (Shanghai, Shandong, Jiangsu, Zhejiang), Northwest China (Shaanxi, Xinjiang), and Southwest China (Chongqing, Sichuan, Yunnan); Japan, Korea, Iran, Italy, Turkey, Jordan, and South Africa. A total of 4,555 CHB patients untreated with lamivudine and other antiviral drugs were involved in this review. A great majority of these cases were collected from hospitals.

Difference in the incidence of YMDD mutation between lamivudine-treated cases and lamivudine-untreated cases

Twelve studies compared YMDD mutation incidence between 611 lamivudine-treated CHB patients and 551 untreated CHB patients. There was no significant statistical heterogeneity (χ² = 15.18, P = 0.17, I²= 27.5%). The meta-analysis with fixed effects models demonstrated higher YMDD mutation rate in lamivudine-treated CHB cases than in untreated CHB cases (OR = 5.23, 95% CI: 3.55–7.17; Z = 8.37, P<0.01) (Figure 2). Begg’s test and Egger’s test showed that publication bias existed.

Incidence of YMDD motif mutation among lamivudine-untreated CHB patients

As showed in Table 1, the pooled incidence of all the eight countries in YMDD mutation among lamivudine-untreated cases is 12.21% (95% CI: 9.69%–14.95%), while the pooled incidence in China is 13.38% (95% CI: 10.90%–16.07%). We also calculated the incidence in each sub-region of China. The naturally occurring YMDD mutation was relatively more severe in Northwest China and Southwest China, where the incidences were 20.19% (95% CI: 14.03%–27.10%) and 17.23% (95% CI: 6.07%–32.58%), respectively. The lowest incidence was in North China with the rate of 6.71% (95% CI: 3.87%–10.26%). The incidences of naturally occurring YMDD motif mutation in Central China, East China, and South China were 12.77% (95%
Figure 1. Flow diagram of the systematic literature research.
doi:10.1371/journal.pone.0032789.g001

Figure 2. Difference in the incidence of YMDD mutation between lamivudine-treated patients and untreated patients.
doi:10.1371/journal.pone.0032789.g002
| Region            | Province       | Author | Year | Untreated patients | Cases of YMDD mutation | Incidence of YMDD mutation (95%CI) (%) | Pooled incidence of YMDD mutation (95%CI) (%) | Weight (%) |
|-------------------|----------------|--------|------|--------------------|------------------------|---------------------------------------|-----------------------------------------------|------------|
| China             | Central China  | Xiang T [7] | 2010 | 40                 | 3                      | 8.51 (2.02, 18.84)                     | 12.77 (7.06, 19.87)                           | 7.10       |
|                   |                | Wei J [8]  | 2010 | 449                | 46                     | 10.35 (7.69, 13.31)                    |                                               |            |
|                   | Hunan          | Tang XL [9] | 2009 | 119                | 23                     | 19.59 (13.01, 27.14)                   |                                               |            |
|                   | North China    | Shi M [10] | 2006 | 98                 | 5                      | 5.54 (1.92, 10.87)                     | 6.71 (3.87, 10.26)                           | 8.34       |
|                   | Shan’xi         | QI QS [11] | 2005 | 36                 | 2                      | 6.68 (1.01, 16.85)                     |                                               |            |
|                   | Jinlin          | Xiong Y [12] | 2008 | 30                 | 1                      | 4.70 (0.18, 14.77)                     |                                               |            |
|                   | Beijing         | Ren YQ [13] | 2004 | 65                 | 6                      | 9.84 (3.89, 18.10)                     |                                               |            |
|                   | East China      | Wu C [14]  | 2007 | 38                 | 3                      | 8.93 (2.13, 19.75)                     | 11.18 (8.34, 14.35)                          | 30.30      |
|                   |                | Min XC [16] | 2009 | 196                | 21                     | 10.90 (6.96, 15.63)                    |                                               |            |
|                   |                | Huo H [17]  | 2006 | 183                | 40                     | 22.02 (16.33, 28.26)                   |                                               |            |
|                   | Jiangsu         | Yu HZ [18]  | 2009 | 20                 | 2                      | 11.79 (1.87, 28.62)                    |                                               |            |
|                   |                | Qian JB [19] | 2002 | 86                 | 3                      | 4.01 (0.93, 9.10)                      |                                               |            |
|                   |                | Qiu YW [20]  | 2008 | 380                | 63                     | 16.66 (13.11, 20.55)                   |                                               |            |
|                   |                | Cheng NL [21]| 2008 | 170                | 14                     | 8.48 (4.78, 13.11)                     |                                               |            |
|                   | Zhejiang        | Zhang XM [22]| 2004 | 122                | 12                     | 10.17 (5.47, 16.11)                    |                                               |            |
|                   |                | Zheng XM [23]| 2003 | 40                 | 3                      | 8.51 (2.02, 18.84)                     |                                               |            |
|                   |                | Cheng FJ [24]| 2006 | 89                 | 12                     | 13.90 (7.56, 21.74)                    |                                               |            |
|                   |                | Zheng NH [25]| 2007 | 229                | 16                     | 7.17 (4.20, 10.84)                     |                                               |            |
|                   |                | Yan MH [26]  | 2003 | 110                | 19                     | 17.57 (11.09, 25.16)                   |                                               |            |
|                   | South China     | WU J [27]  | 2009 | 49                 | 7                      | 14.99 (6.56, 26.08)                    | 14.91 (8.04, 23.41)                          | 16.03      |
|                   |                | Huang ZM [28]| 2005 | 104                | 28                     | 27.14 (19.12, 36.00)                   |                                               |            |
|                   |                | Qian YQ [29] | 2005 | 125                | 30                     | 24.22 (17.15, 32.02)                   |                                               |            |
|                   | Guangdong       | QU YY [30]  | 2006 | 140                | 20                     | 14.52 (9.22, 20.79)                    |                                               |            |
|                   |                | Liu JH [31]  | 2008 | 176                | 7                      | 4.22 (1.77, 7.69)                      |                                               |            |
|                   |                | Guo HB [32]  | 2004 | 29                 | 6                      | 21.65 (9.02, 37.88)                    |                                               |            |
|                   |                | Yu J [33]    | 2007 | 71                 | 4                      | 6.24 (1.87, 12.94)                     |                                               |            |
|                   | NorthWest       | Li CM [34]  | 2003 | 39                 | 3                      | 8.71 (2.07, 19.27)                     | 20.19 (14.03, 27.10)                        | 13.80      |
|                   |                | Zhang XH [35]| 2003 | 150                | 32                     | 21.53 (15.34, 28.40)                   |                                               |            |
|                   | Shaanxi         | Qu JX [36]  | 2010 | 82                 | 21                     | 25.91 (17.11, 35.81)                   |                                               |            |
|                   |                | Qu JX [37]   | 2008 | 106                | 28                     | 26.65 (18.72, 35.38)                   |                                               |            |
|                   |                | Pan HQ [38]  | 2010 | 110                | 31                     | 28.40 (20.43, 37.11)                   |                                               |            |
|                   | SouthWest       | Zhang Y [39] | 2005 | 60                 | 5                      | 8.99 (3.17, 17.38)                     |                                               |            |
|                   |                | Li L [40]    | 2005 | 54                 | 11                     | 20.92 (11.28, 32.54)                   | 17.23 (6.07, 32.58)                          | 8.06       |
|                   |                | Liu TH [41]  | 2004 | 55                 | 3                      | 6.22 (1.46, 13.97)                     |                                               |            |
|                   | Sichuan         | He X [42]   | 2007 | 12                 | 1                      | 11.25 (0.49, 33.19)                    |                                               |            |
|                   |                | Wang YQ [43] | 2009 | 190                | 60                     | 31.70 (25.30, 38.42)                   |                                               |            |
|                   | Yunnan          | Wang YQ [43] | 2009 | 190                | 60                     | 31.70 (25.30, 38.42)                   |                                               |            |
|                   | Pooled incidence of YMDD mutation in China |                |      |                    |                                       | 13.38 (10.90, 16.07)                        | 83.63      |
|                   | Other countries | Matsuda M [44]| 2004 | 20                 | 1                      | 6.96 (0.28, 21.36)                     | 9.90 (3.28, 19.55)                           | 16.37      |
|                   | Japan           | Seta T [45] | 2000 | 10                 | 0                      | 2.32 (2.02, 18.80)                     |                                               |            |
|                   | Korea           | Heo J [47]  | 2004 | 40                 | 3                      | 8.51 (2.01, 18.84)                     |                                               |            |
|                   | Iran            | AB-Olyaee S [48]| 2008 | 147                | 0                      | 0.17 (0.16, 1.47)                      |                                               |            |
|                   | Turkey          | Akarsu M [51]| 2006 | 71                 | 13                     | 18.76 (10.65, 28.53)                   |                                               |            |
|                   | Jordan          | Masaadeh HA [52]| 2008 | 4                  | 1                      | 29.48 (18.4, 71.94)                    |                                               |            |
|                   | South Africa    | Selabe SG [53]| 2007 | 15                 | 3                      | 21.78 (5.67, 44.52)                    |                                               |            |
|                   | Pooled incidence of YMDD mutation in these eight countries |                |      |                    |                                       | 12.21 (9.69, 14.95)                        | 100.00     |
CI: 7.06%–19.87%), 11.18% (95% CI: 8.34%–14.35%), and 14.91% (95% CI: 8.04%–23.41%), respectively. In addition, the pooled incidence of the seven foreign countries was 9.90% (95% CI: 3.28%–19.55%).

Incidence of natural YMDD mutation between gender and age

Stratified analyses showed that natural YMDD mutation was not associated with gender (Figure 3) and age (Figure 4).

Eleven studies contained the available information of gender distribution, including 967 males and 252 females. There was no significant statistical heterogeneity ($\chi^2 = 5.52, P = 0.85, I^2 = 0$). Males and females had equal chances of natural YMDD mutation (OR = 1.12, 95% CI: 0.76–1.64; $Z = 0.57, P = 0.57$) (Figure 3). No significant publication bias was observed in this analysis.

The average age was mentioned in six studies including 177 cases with YMDD mutations and 702 cases without YMDD mutation. Statistical heterogeneity was observed between groups ($\chi^2 = 66.00, P < 0.00001, I^2 = 92.4\%$). By using the random effects models, no significant difference of average age was showed between groups (WMD (weighted mean difference) = −3.73, 95% CI: −11.49–4.03; $Z = 0.94, P = 0.35$) (Figure 4). Begg’s test and Egger’s test indicated that publication bias existed.

Relation of natural YMDD mutation with pathogen-related characteristics

Statistical analyses also showed that the occurrence of YMDD mutation was not associated with HBeAg status (Figure 5), HBV genotype (Figure 6), and ALT level (Figure 7). However, viral load was showed to be associated with YMDD mutation. Neither statistical heterogeneity nor publication bias was observed in those

Figure 3. Difference in the incidence of YMDD mutation between male and female patients.
doi:10.1371/journal.pone.0032789.g003

Figure 4. Difference in average ages between the patients with YMDD mutations and the patients without mutation.
doi:10.1371/journal.pone.0032789.g004
analyses, except that publication bias existed in articles of HBV genotype analysis.

Twelve studies including 1194 HBeAg-positive cases and 639 HBeAg-negative cases showed that the incidence of natural YMDD-motif mutation was the same in different HBeAg status groups (OR = 1.03, 95% CI: 0.67–1.58; Z = 0.15, P = 0.88). It also seems that ALT level did not play a role in occurrence of natural YMDD mutation among patients with or without lamivudine therapy (WMD = 0.04, 95% CI: −3.23–3.14; Z = 0.03, P = 0.98). Six studies reported 149 genotype-B HBV cases and 382 genotype-C HBV cases. No significant difference was found in rates of natural YMDD mutation between genotype B and genotype C (OR = 1.62, 95% CI: 0.99–2.64; Z = 1.92, P = 0.05).

Eight studies (including 129 mutants and 774 cases without mutation) referred to the relation of HBV-DNA viral load with YMDD mutation. As showed in Figure 8, the HBV DNA copy number was higher in the group with YMDD mutation than in the group without mutation (WMD = 0.27, 95% CI: 0.10–0.45; Z = 3.09, P = 0.002).

**Discussion**

The application of lamivudine is a breakthrough in viral treatment. Most patients with chronic hepatitis B had excellent response to lamivudine, but development of YMDD mutants attenuated efficacy of lamivudine treatment. Based on the studies included in this analysis, lamivudine treatment increased the risk of YMDD mutation to 5 fold of untreated controls (OR = 5.23, 95% CI: 3.55–7.71). Although mutations in YMDD motif increased during lamivudine treatment, many studies reported that this mutation could also be spontaneous. As showed in this report, the pooled incidence of natural YMDD mutation was as high as 12.21% (95% CI: 9.69%–14.95%). This result mostly represented the current situation in East Asia, especially China,
since in this research 37 studies were from China and only 10 were from other 7 countries including Japan and Korea. It is not strange because China has a high prevalence of HBV and numbers of reports have been accumulated with clinical observation of YMDD mutations. However, the incidence of spontaneous YMDD mutations did not differ among countries (Table 1). Gender and age were not related with spontaneous YMDD mutations either (Figure 3 or Figure 4). These results suggest that pathogen factors rather than host genetic factors are the determinants of these natural YMDD mutants.

Of all the pathogen-related factors studied in this review, viral load was correlated with the incidence rate of natural YMDD mutation, while patients’ HBeAg status and ALT level had no influence. A cohort study by Atkins M et al. reported that mutant virus strains were more likely to appear in lamivudine-treated patients with larger virus amount and longer therapy duration [54]. It seems like the YMDD mutation might occur at certain rate and accumulate during HBV replication. The relation between YMDD mutation and infection duration was not conclusive because it is difficult to determine the accurate infection duration. It is also hard to define a proper study interval. Min XC et al [16] found no difference in YMDD natural mutation between the group with 30-year or more infection duration and the group with less than 30-year duration. The issue is that 30-year duration may not be an appropriate study interval to detect accumulation effect of mutations.

Hepatitis B virus has eight genotypes classified from A to H, which variant in geographic distribution and characteristics. Genotypes B and C are predominant in East Asia such as China, Korea, and Japan [55]. A few reports showed an association between emergence rate of YMDD mutants and HBV genotypes. Huang ZM et al [28] demonstrated that the mutations mostly occur in genotype C and its mixed genotypes. Controversially Min XC et al [16] and Huo H et al [17] indicated these mutants emerge more frequently in genotype B virus than in genotype C virus. By this meta-analysis, no significant difference was observed in incidence of YMDD mutation between genotype B and genotype C. Due to inadequate information in the adopted literatures we failed to analyze the other genotypes in this review. Multinational studies would be valuable to clarify the relationship between the occurrence of YMDD mutants and HBV genotypes.

It has been commonly recognized that YMDD mutations can occur spontaneously. There are two major concerns following the

---

**Table 1. Study results of natural YMDD mutation among patients with HBV.**

| Study or sub-category | N  | With mutation Mean(SD) | WO mutation Mean(SD) | WMD(fixed) 95%CI | Weight % | WMD(fixed) 95%CI |
|----------------------|----|------------------------|----------------------|------------------|----------|-----------------|
| Akarsu M 2006        | 13 | 21.00(5.00)            | 21.00(7.00)          | 95.45            | 0.00     | [-3.26,3.26]   |
| Git YW 2008          | 63 | 175.30(75.10)          | 317                  | 2.43             | 3.60     | [16.85,24.05]  |
| Git YY 2006          | 20 | 53.90(61.30)           | 120                  | 1.35             | -13.90   | [41.33,13.53]  |
| Ren YQ 2004          | 6  | 121.75(82.67)          | 59                   | 0.38             | 12.38    | [39.55,64.31]  |
| Zhang Y 2005         | 5  | 113.23(56.32)          | 55                   | 0.39             | 2.79     | [47.97,53.55]  |
| Total(95%CI)         | 107| 609                    |
| Test for heterogeneity | ch^2=1.33, df=4(P=0.86), P=0% |
| Test for overall effect: Z=0.03(P=0.98) |

---

**Figure 7. Difference in ALT level between the patients with YMDD mutations and the patients without mutation.**

doi:10.1371/journal.pone.0032789.g007

**Figure 8. Difference in HBV DNA copies between the patients with YMDD mutations and the patients without mutation.**

doi:10.1371/journal.pone.0032789.g008
finding. Firstly, do naturally occurring YMDD mutations influence the nature course of hepatitis B and worsen patients’ condition? Secondly, since lamivudine treatment increases YMDD mutations, is efficacy of lamivudine reduced during treatment? Currently there is no evidence that YMDD mutations are associated with ALT level. The result indicates that at least naturally existing YMDD mutants may not worsen patients’ liver function. Allen MJ et al [4] demonstrated that the sensitivity of YMDD mutant strains to lamivudine declined and viral titers rebounded during lamivudine treatment in vitro. This finding does not mean necessarily that patients with pre-existing mutations have to be identified for special care or other therapy, because YMDD mutants were found to be attenuated in replication capacity and pathogenicity [44,46]. In that case, YMDD mutants have little possibility to be dominant during infection and lamivudine should still be effective to treat hepatitis. All these opinions are not convincing enough to manipulate strategy of current lamivudine treatment and more fundamental and clinical evidences are needed to reveal the ultimate impact of YMDD mutations.

Considering the YMDD mutations could be acquired from other people instead of occur spontaneously, the natural mutation rate might be overestimated. However, there has been no laboratory evidence clarifying this possibility. Qu JX et al [36] indicated that children had higher mutation prevalence if YMDD mutants were detected in their parents. The question is whether these mutants are transmitted from parents or genetic background increases mutation rate. Plus the conclusion was drawn from fourteen families and might just be a coincidence. Further studies are needed to distinguish acquired mutation from spontaneous mutation, including comparison of infectivity between the wild type and mutant virus strains, transfection assay of cells, and animal experiments.

In conclusion, YMDD-motif mutation can occur naturally with a rather high rate about 12.21% among untreated CHB patients. Patients’ viral load is the factor that is admitted by majority to be correlated with mutation rate. Further mechanism researches are required to clarify the behaviors of mutants in addition to attenuated replication ability. Large-scale population studies in multi-countries are also necessary to evaluate the influence of YMDD mutations in hepatitis B progression and antiviral treatment.

Supporting Information
Figure S1 PRISMA Flowchart.

Table S1 PRISMA Checklist.

Author Contributions
Conceived and designed the experiments: RBY YWT KQD. Performed the experiments: KQD YHG LC QL XC. Analyzed the data: KQD ZHP. Contributed reagents/materials/analysis tools: JS ZHP. Wrote the paper: KQD YWT. Revised the manuscript: JS XT RBY.

References
1. Kao JH, Chen DS (2002) Global control of hepatitis B virus infection. Lancet Infect Dis 2: 395–403.
2. Tipples GA, Ma MM, Fischer KP, Bain VG, Ketneten NM, et al. (1996) Mutation in HBV RNA-dependent DNA polymerase confers resistance to lamivudine in vitro. Hepatology 24: 714–717.
3. Tsutoda A (2006) How do naturally occurring YMDD-motif mutants influence the clinical course of lamivudine-naive patients with chronic hepatitis B virus infection? J Gastroenterol Hepatol 21: 1783–1788.
4. Allen MI, Deslauriers M, Andrews CW, Tipples GA, Walters KA, et al. (1998) Identification and characterization of mutations in hepatitis B virus resistant to lamivudine. Hepatology 27: 1670–1677.
5. Stuyver IJ, Locarnini SA, Lok A, Richman DD, Carman WF, et al. (2001) Nomenclature for antiviral-resistant human hepatitis B virus mutations in the polymerase region. Hepatology 33: 751–757.
6. Moher D, Liberati A, Tetzlaff J, Altman DG, Grp P (2009) Preferred Reporting Items for Systematic Reviews and Meta-Analyses: The PRISMA Statement. Plos Medicine 6.
7. Xiang T, Liu X, Zhang LS, Dai WA, Li QZ, et al. (2010) The detection of YMDD mutation in CHB patients in Enshi Autonomous Prefecture. Journal of Hubei University for Nationalities 27: 47–49.
8. Wei J, Wang H, Li SL, Zeng LL (2010) Experimental research for YMDD motif natural mutations in the serum of patients with chronic hepatitis B. Chinese Journal of Hepatology 18: 855–856.
9. Tang XL (2009) Detection of spontaneous hepatitis B virus YMDD motif mutation by fluorescent hybridization double-probe PCR, and melting curve assay. Chinese Journal of Modern Medicine 19: 1544–1546.
10. Shi M, Yang JZ, Wang RS, Zhang H, Zhu YF, et al. (2006) Rapid quantitation of lamivudine-resistant mutants in lamivudine treated and untreated patients with chronic hepatitis B virus infection. Clin Chim Acta 373: 172–175.
11. Qi QS, Cheng SB, Yang Q, Yi DY (2005) Study on the therapeutic efficacy of lamivudine in chronic hepatitis B patients with YMDD mutations. Acta Academiae Medicinae Jiangsu 45: 42–49.
12. Xiong Y, Song YG (2005) Detection of YMDD mutants in patients with chronic hepatitis B during lamivudine treatment. Journal of Beijing University 6: 421–424.
13. Ren YQ, Hu DR, Fan GR, Wu YF (2004) Detection of YMDD mutation in CHB patients untreated with antiviral medicine. Journal of Clinical Hepatology 18: 85–88.
14. Wu C, Ren LY, Li Q (2007) Detection and analysis of HBV YMDD-motif mutants before and during lamivudine treatment. Medical Laboratory Science and Clinics 18: 46–48.
15. Wang JF, Zhang ZQ, Du MH, Gao T, Gu SM (2003) Detection of hepatitis B virus YMDD mutants and its clinical significance in patients with chronic hepatitis B. World Journal of infection 3: 405–406.
16. Min XC, Miao XH, Zhao SM, Zhao KK, Yang DG (2009) The spontaneous YMDD mutation rate in chronic hepatitis B patients. Chinese Journal of Hepatology 17: 887–890.
17. Huo H, Zhao SM, Cai X, Miu XH (2006) Epidemiological survey of spontaneous HBV YMDD motif mutation. Chinese Hepatology 11: 232–234.
18. Yu HZ, Yuan JF, Zhang XJ, Cheng Y (2009) Detection of YMDD mutation in patients with hepatitis B by double probe FQ-PCR assay. Medical Information 1: 19–20.
19. Qian JB, Sun NX, Du SC, Fan XF, Zhang YX (2002) Detection of YMDD motif mutation in lamivudine-treated and untreated CHB patients. Journal of Jiangsu Medicine 28: 408–410.
20. Qiu YW, Huang LH, Hu TH, Cheng SB, Yan Q (2008) Study of the disease progress and clinical treatment in the hepatitis B patients with YMDD mutation. Clinical Focus 23: 350–355.
21. Chen NL, Zhao SL, Liu XR, Wei YF, Wu HY, et al. (2008) Relationship of YMDD mutations and genotypes of HBV in antiviral-untreated patients with chronic hepatitis B. Chinese Journal of Clinical Physican 2: 1125–1127.
22. Zhang JM (2004) Clinical analysis on YMDD-motif spontaneous mutation in patients with chronic hepatitis B. Journal of Shaodong Medicine 44: 22–23.
23. Zheng JM, Ni PH (2003) Detection of YMDD mutation in HBV polymerase gene region with nested PCR. Shanghaji. Journal of Medical Laboratory Science 18: 261–264.
24. Cheng FJ, Cheng SL, Lu J, Pang WY (2006) Analysis of YMDD mutation in antiviral-untreated patients with chronic hepatitis B. Zejhuan Journal of Preventive Medicine 11: 11–12.
25. Zheng NH, Ji CB, Zhou WH, Ying H, Hu JD, et al. (2007) Detection of naturally emerging HBV YMDD mutation. Chinese Journal of Clinical Hepatology 23: 215–214.
26. Yan MH, Zhang C, Lin Q, Zhou RF (2003) Detection of YMDD motif mutation in lamivudine-untreated patients with chronic hepatitis B. Chinese Journal of Hepatology 11: 430–431.
27. Wu J, Qian X (2009) Correlation between hepatitis B virus genotypes and natural mutation of YMDD in Hainan. Journal of Hainan Medical College 15: 714–716.
28. Huang ZM, Huang QW, Qin YQ, He YZ, Qin HJ, et al. (2005) YMDD mutations in patients with chronic hepatitis B untreated with antiviral medicines. World J Gastroenterol 11: 867–870.
29. Qian YQ, Zhao DY, Qian HJ, Lin GC, Huang ZM (2003) The study of natural HBV YMDD mutation in 125 Zhuang patients with chronic hepatitis B in Guannxi province. Journal of Chinese Physican 7: 1531–1532.
30. Qu YY, Wei JJ, Zeng WT, Zhu KL, Ma PQ (2006) Clinical analysis of HBV YMDD mutation in patients untreated with lamivudine. Journal of Tropical Medicine 6: 791–793.
31. Liu JH, Zeng H, Cao BH, Li XY, Wen FY, et al. (2008) Study on the relationship between lamivudine treatment and YMDD motif mutation. Journal of Chinese Tropical Medicine 8: 1483–1487.

32. Guo HR, Li L, Ye XG, Wang H (2004) HBV genotyping and detecting of YMDD mutation in patients chronically infected HBV and untreated with antiviral drugs. Journal of Chinese Biotechnology 24: 107–109.

33. Yu J, Huang JM, Zhang YH, Zou DX, Peng LM (2007) Detection of HBV gene mutation and analyzing for the correlation factors of the mutations. International Medicine and Health Guidance News 13: 63–66.

34. Li CM, Zhang YX (2003) Detection of lamivudine-resistant HBV YMDD mutation and analyzing the relative factors of YMDD mutation. Chinese Journal of Integrated Traditional and Western Medicine on Liver Diseases 13: 137–139.

35. Zhang XH, Zhang YX, Sun LR, Wen Q, Zhou LQ, et al. (2003) The study of gene chips in detecting YMDD mutations in the region of HBV polymerase. National Medical Journal of China 83: 459–462.

36. Qu JX, Zeng QL, Liu ZG, Xu GH, Fen JH, et al. (2010) Spontaneous HBV YMDD motif mutations in patients with familial clustered chronic HBV infection. World Chinese Journal of Digestology 18: 2604–2607.

37. Qu JX, Liu R, Ren YQ, Fen JH (2008) Study in naturally occurring HBV YMDD mutations in patients with hepatic cirrhosis. Chinese Journal of Clinical Hepatology 24: 116–117.

38. Pan HQ, Zeng QL, Qu JX, Xu GH, Li YJ, et al. (2010) Epidemiological survey for spontaneous HBV YMDD motif mutation of CHB patients whom under 20 years old and in immune tolerant phase. Medical Information 23: 2094–2095.

39. Zhang Y, Wang PZ, Zhang Y, Li Y, Wei X, et al. (2005) The study of natural HBV YMDD-motif mutation in patients with chronic hepatitis B. Military Medical Journal of South China 19: 13–15.

40. Li L, Mao Q, Huang YP, Wang YM (2005) Detection of HBV YMDD mutation in patients with chronic hepatitis B and its significance. Acta Academiae Medicinarum Militaris Tertiae 27: 167–169.

41. Liu TH, Lang XJ (2004) Detection of YMDD mutation in patients with chronic hepatitis B after lamivudine treatment. Chinese Pharmaceuticals 13: 29.

42. He X, Jiang N (2007) Analysis on the HBV polymerase gene mutation in lamivudine-resistant patients. Journal of Sichuan University 38: 805–806.

43. Wang YQ, Pu D (2009) Clinical analysis of natural mutation of YMDD in HBV infection patients. Journal of China Tropical Medicine 9: 652–653.

44. Matsuda M, Suzuki F, Suzuki Y, Tsubota A, Akata N, et al. (2004) YMDD mutants in patients with chronic hepatitis B before treatment are not selected by lamivudine. J Med Virol 74: 361–366.

45. Seto T, Yokosuka O, Imazeki F, Tagawa M, Saijo H (2000) Emergence of YMDD motif mutants of hepatitis B virus during lamivudine treatment of immunocompetent type B hepatitis patients. J Med Virol 60: 8–16.

46. Kobayashi S, Ide T, Sata M (2001) Detection of YMDD motif mutations in some lamivudine-untreated asymptomatic hepatitis B virus carriers. J Hepatol 34: 394–398.

47. Heo J, Cho M, Kim HH, Shin YM, Jiang HJ, et al. (2004) Detection of YMDD motif mutants by oligonucleotide chips in lamivudine-untreated patients with chronic hepatitis B virus infection. J Korean Med Sci 19: 541–546.

48. Amini-Bavil-Olyaee S, Hosseini SY, Sahibi F, Alavian SM (2008) Hepatitis B virus (HBV) genotype and YMDD motif mutation profile among patients infected with HBV and untreated with lamivudine. Int J Infect Dis 12: 83–87.

49. Ramezanian A, Velayati AA, Rashian MR, Gachkar L, Bamizad M, et al. (2006) Rate of YMDD motif mutants in lamivudine-untreated Iranian patients with chronic hepatitis B virus infection. Int J Infect Dis 12: 252–255.

50. Villa E, Lei B, Taliani G, Grazioni A, Critelli R, et al. (2009) Pretreatment with pegylated interferon prevents emergence of lamivudine mutants in lamivudine-naive patients: a pilot study. Antivir Ther 14: 1081–1087.

51. Akarsu M, Sengorul A, Tankurt E, Sayiner AA, Topalak O, et al. (2006) Detection of YMDD motif variants in inactive hepatitis B carrier by Inno-Lipa HBV DR assay. J Gastroenterol and Hepatol 21: 1783–1788.

52. Masaadeh HA, Hayajneh WA, Alqudah EA (2008) Hepatitis B virus genotypes and lamivudine resistance mutations in Jordan. World J Gastroenterol 14: 7231–7234.

53. Selabe SG, Lukhwareni A, Song E, Leeuw YG, Burnett RJ, et al. (2007) Mutations associated with lamivudine-resistance in therapy-naive hepatitis B virus (HBV) infected patients with and without HIV co-infection: implications for antiretroviral therapy in HBV and HIV co-infected South African patients. J Med Virol 79: 1630–1634.

54. Atkins M, Hunt MC, Brown N, Gray F, Sanathanan L, et al. (1998) Clinical significance of YMDD mutant hepatitis B virus (HBV) in a large cohort of lamivudine-treated hepatitis B patients. Hepatology 28: 319A.

55. Nordier H, Couroucé AM, Couraget P, Echevarria JM, Lee SD, et al. (2004) Genetic diversity of hepatitis B virus strains derived worldwide: genotypes, subgenotypes, and HBsAg subtypes. Intervirology 47: 289–309.