<table>
<thead>
<tr>
<th>Title</th>
<th>Current antiviral therapy of chronic hepatitis B: Efficacy and safety</th>
</tr>
</thead>
<tbody>
<tr>
<td>Author(s)</td>
<td>Lam, YF; Yuen, MF; Seto, WK; Lai, CL</td>
</tr>
<tr>
<td>Citation</td>
<td>Current Hepatitis Reports, 2011, v. 10 n. 4, p. 235-243</td>
</tr>
<tr>
<td>Issued Date</td>
<td>2011</td>
</tr>
<tr>
<td>URL</td>
<td><a href="http://hdl.handle.net/10722/144892">http://hdl.handle.net/10722/144892</a></td>
</tr>
<tr>
<td>Rights</td>
<td>The Author(s); This work is licensed under a Creative Commons Attribution-NonCommercial-NoDerivatives 4.0 International License.</td>
</tr>
</tbody>
</table>
Current Antiviral Therapy of Chronic Hepatitis B: Efficacy and Safety

Yuk-Fai Lam · Man-Fung Yuen · Wai-Kay Seto · Ching-Lung Lai

Published online: 9 August 2011
© The Author(s) 2011. This article is published with open access at Springerlink.com

Abstract The treatment of chronic hepatitis B is in constant evolution. Interferon, the first agent licensed for chronic hepatitis B treatment, has been superseded by the growing popularity of nucleoside/nucleotide analogues (NA). However, resistance to these agents is a major challenge. Newer NAs, such as entecavir and tenofovir dipivoxil fumarate, have very low resistance rates and favorable safety profiles. Long-term use of these agents can effectively suppress hepatitis B virus DNA, leading to decrease in incidence of hepatic flares, as well as in the development of cirrhosis and hepatocellular carcinoma. The efficacy and safety of various antiviral agents is discussed in this review.

Keywords Hepatitis B · Nucleos(t)ide Analogs · Resistance · HBV DNA

Introduction

Chronic hepatitis B (CHB) infection is a major health burden worldwide, with around 400 million people affected [1]. Patients with CHB have a 15% to 40% chance of developing cirrhosis, hepatic decompensation and hepatocellular carcinoma in their lifetime [2, 3]. The risk of having hepatocellular carcinoma is increased by more than 100–200 fold compared to healthy subjects [4]. Currently two classes of agents are available for treatment of chronic hepatitis B, namely immunomodulatory therapy (conventional interferon and pegylated interferon) and nucleoside/nucleotide analogs (NA).

Conventional Interferon and Pegylated Interferon

The first agent approved for treatment of chronic hepatitis B was conventional interferon-α [5]. Pegylated-interferon, which was licensed in 2005, has the advantage of ease of administration when compared to conventional interferon. They mainly work through immunomodulation.

These agents decrease viral loads and increase rates of hepatitis B e antigen (HBeAg) seroconversion to antibody against HBeAg (anti-HBe) [6]. They also have the advantage of use for a finite duration. However, the course of treatment has been extended from 16 weeks with the conventional interferon to 48 weeks with pegylated interferon.

Interferons are poorly tolerated because of their severe side effects. The side effects include influenza-like illness, anorexia, flares of autoimmune disease, thyroid dysfunction, myelosuppression, hepatitis flare, hepatic decompensation, and neuropsychiatric adverse events like depression, irritability, and even suicidal tendency. High relapse rate and costs are also drawbacks of immunomodulatory therapy [7, 8].

On long-term follow-up, a majority of patients still have detectable hepatitis B virus (HBV) DNA after interferon treatment [9, 10]. In addition, most studies fail to demonstrate a reduction in incidence of hepatocellular carcinoma with conventional interferon on long term follow-up [11–13].

Nucleoside/Nucleotide Analogue

Currently five NAs are approved for the treatment of CHB. They are lamivudine, adefovir dipivoxil (ADV),
Telbivudine, entecavir and tenofovir disoproxil fumarate (TDF) (Tables 1 and 2)

They are effective in suppressing viral loads, facilitating HBeAg seroconversion, achieving alanine aminotransferase (ALT) normalization, and improving liver fibrosis. They are generally well tolerated and serious adverse events are rarely encountered [14–17].

There is growing evidence that prolonged and effective suppression of HBV DNA can decrease the risk of cirrhosis and hepatocellular carcinoma. Therefore the current trend of treatment for chronic hepatitis B is to use long term NA therapy to achieve permanent virologic suppression, with loss of hepatitis B surface antigen (HBsAg) as the ideal end-point [18].

The comparative efficacy and resistance rates of the five nucleoside and nucleotide analogues are shown in Table 1 and 2 respectively.

**Lamivudine**

Lamivudine was originally used for HIV infection. It was approved by the United States Food and Drug Administration (FDA) in 1998 as the first nucleoside analogue for the treatment of CHB. Lamivudine is the (−) enantiomer of 2′,3′-dideoxy 3′-thiacytidine. It is phosphorylated into the triphosphate form (3TC-TP) and is incorporated into the growing chain of DNA during reverse transcription of the first strand of HBV DNA and synthesis of the second strand of HBV DNA, resulting in chain termination and inhibiting HBV DNA synthesis [15, 19].

**Efficacy and Resistance**

Lamivudine is effective in suppressing viral replication in both HBeAg-positive and HBeAg-negative patients.

The Asia Hepatitis Lamivudine Study Group concluded that lamivudine 100 mg, when compared to placebo, achieved a higher rate of HBeAg seroconversion (16% versus 4%) and sustained ALT normalization (72% versus 24%) at 1 year. Greater proportion of patients in the group receiving lamivudine 100 mg daily, when compared to placebo, achieved improvement of necroinflammatory activity by 2 points or more at 1 year (56% versus 25%) [20]. The efficacy of lamivudine is confirmed by another multicentre trial performed in the United States. HBeAg-positive patients on lamivudine 100 mg daily, as compared to placebo, have higher rates of histological improvement (52% versus 23%), HBeAg seroconversion (32% versus 11%), undetectable HBV DNA (44% versus 16%) and alanine aminotransferase normalization (41% versus 16%) at 48 weeks [21]. For extended lamivudine therapy to 3 years, more patients achieved HBeAg seroconversion, alanine aminotransferase normalization and sustained HBV DNA suppression [22].

Lamivudine is also efficacious in HBeAg-negative patients. After 1 year of treatment, 96% of patients achieved alanine aminotransferase normalization and 68% achieved undetectable HBV DNA [23].

Lamivudine is effective in preventing progression of cirrhosis and developing of hepatocellular carcinoma [24–26]. A large prospective study for patients with established cirrhosis [27] showed that patients on lamivudine 100 mg daily, compared to placebo, were less likely to have increase in Child-Turcotte- Pugh (CTP) score (3.4% versus 8.8%) and development of hepatocellular carcinoma (3.9% versus 7.4%). In fact, this study, originally planned for a follow-up period of 5 years, was terminated at median of 32.4 months because of the significant differences observed between the treatment group and the placebo group.

**Table 1 Efficacy of nucleoside/nucleotide analogs treatment for chronic hepatitis B (HBeAg-positive/HBeAg-negative)**

<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>1 year</td>
<td>16/−</td>
<td>12/−</td>
<td>23/−</td>
<td>21/−</td>
<td>21/−</td>
</tr>
<tr>
<td>2 year</td>
<td>29/−</td>
<td>29/−</td>
<td>30/−</td>
<td>31/−</td>
<td>27/−</td>
</tr>
<tr>
<td>4 (or 5*) year</td>
<td>47/−</td>
<td>48/−</td>
<td>NA/−</td>
<td>44/− a</td>
<td>29/−</td>
</tr>
<tr>
<td>ALT normalization (%)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1 year</td>
<td>72/96</td>
<td>48/72</td>
<td>77/74</td>
<td>68/78</td>
<td>68/76</td>
</tr>
<tr>
<td>2 year</td>
<td>NA/60</td>
<td>74/73</td>
<td>70/77</td>
<td>87/89</td>
<td>NA/NA</td>
</tr>
<tr>
<td>4 (or 5*) year</td>
<td>69/NA</td>
<td>NA/69 a</td>
<td>NA/NA</td>
<td>80/NA a</td>
<td>NA/NA</td>
</tr>
<tr>
<td>Undetectable HBV DNA by PCR (%)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1 year</td>
<td>36/68</td>
<td>21/51</td>
<td>60/88</td>
<td>67/90</td>
<td>76/93</td>
</tr>
<tr>
<td>2 year</td>
<td>NA/42</td>
<td>40/71</td>
<td>56/82</td>
<td>80/94</td>
<td>89/91</td>
</tr>
<tr>
<td>4 (or 5*) year</td>
<td>NA/NA</td>
<td>NA/67 a</td>
<td>NA/NA</td>
<td>94/NA a</td>
<td>96/100</td>
</tr>
</tbody>
</table>

ALT alanine aminotransferase, HBeAg hepatitis B e-antigen, NA not available

a denotes 5 year data
The major problem with long-term lamivudine therapy is drug resistance. Lamivudine is known to have a low genetic barrier. The classical mutation associated with lamivudine is rtM204V/I and rtL180M located in the YMDD locus of hepatitis B virus polymerase [20, 21, 28, 29].

The rate of undetectable HBV DNA decreased from 77% at 12 months of lamivudine treatment, to 52% at 24 months and 42% at 36 months, mainly due to development of drug resistance [23, 30]. For cirrhotic patients on lamivudine, those who have drug resistance are more likely to have increase CTP score when compared to patients without lamivudine resistance [27]. Hepatitic flares and even liver failure were described in patients with lamivudine resistance. Hepatitic flare was noted in 10% of lamivudine-treated patients at 1 year and 18% at 2 years [30, 31].

Safety

Lamivudine has excellent safety profile. Long term study up to 6 years showed no major adverse events and complications associated with lamivudine treatment [30, 32].

Current Roles of Lamivudine Therapy

In view of the low genetic barrier of lamivudine, it is not recommended as the first line treatment for CHB [33•, 34•]. However, lamivudine may have a role as monotherapy for those with these favorable parameters: Baseline HBV DNA less than 9 log, ALT higher than or equal to 2 times upper limit of normal and good viral suppression of less than 4 log at week 4. The rate of lamivudine resistance for patients who have these favorable parameters is low [35].

Lamivudine also has a role as preemptive treatment for CHB patients receiving a finite course of immunosuppression or chemotherapy [36, 37]. The American Association for the Study of Liver Disease guidelines recommends that lamivudine can be used as prophylactic treatment if the anticipated duration of immunosuppression or chemotherapy is less than 12 months and the baseline HBV DNA is undetectable [33•].

Although lamivudine is classified as a pregnancy Category C drug, lamivudine has long term safety data in pregnancy. Pregnant women on lamivudine have the same rate of birth defect compared to the general population [38]. Lamivudine is also useful for patients who have resistance toward adefovir dipivoxil as they have different pattern in viral resistance [33•]. However, entecavir is superior to lamivudine for patients with adefovir resistance.

Adefovir Dipivoxil

ADV was approved by the United States FDA for treatment of chronic hepatitis B in 2002. ADV is a nucleotide analog. It is converted into adefovir and phosphorylated into its active form in the body. Adefovir diphosphonate is incorporated into the viral DNA and thus inhibits HBV DNA reverse transcription and viral replication [14].

Efficacy and Resistance

In a phase III trial involving 515 HBeAg-positive patients who were treated with ADV or placebo, more patients in the adefovir group achieved undetectable HBV DNA (21% versus 0%), alanine aminotransferase normalization (48% versus 16%), histological improvement (53% versus 25%) and HBeAg seroconversion (12% versus 6%) at 48 weeks of treatment [39]. A second study confirmed the efficacy of ADV in HBeAg-negative patients. Patients receiving ADV 10 mg daily when compared with placebo, achieved greater response in terms of histologic improvement (64% versus 33%), undetectable HBV DNA (51% versus 0%) and normalization of alanine aminotransfasc (72% versus 29%) at 48 weeks [40]. Further clinical studies show that extended ADV monotherapy up to 5 years can achieve better histological, virologic and biochemical outcomes [41].

For patients who have lamivudine resistance, combination therapy of lamivudine and adefovir or adefovir monotherapy can achieve alanine aminotransfaser normalization and HBV DNA suppression [42]. However recent studies favor the approach of adding adefovir to lamivudine in lamivudine resistant patients instead of switching to adefovir monotherapy because the latter approach may lead to a higher risk of adefovir resistance (7% versus 18% at 2 years) [43].

ADV is efficacious in decompensated cirrhosis. In a study involving 226 lamivudine-resistant patients with

<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Resistance (%)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1 year</td>
<td>14</td>
<td>0</td>
<td>2.2–5</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>2 year</td>
<td>39</td>
<td>3</td>
<td>10.8–25.1</td>
<td>&lt;1</td>
<td>0</td>
</tr>
<tr>
<td>5 year</td>
<td>60–70</td>
<td>20–29</td>
<td>NA</td>
<td>1.2</td>
<td>NA</td>
</tr>
</tbody>
</table>

Table 2 Resistance rate of nucleoside/nucleotide analogs for treatment of chronic hepatitis B

NA not available
decompensated cirrhosis awaiting liver transplant, HBV DNA became undetectable in 59% and 65% at week 48 and 96 with ADV treatment [44]. In the same study with 241 post-liver transplantation lamivudine-resistant CHB patients, HBV DNA became undetectable in 40% and 65% at weeks 48 and 96 with ADV treatment. There is also improvement of CTP score with adefovir treatment in lamivudine-resistant chronic hepatitis B cirrhosis.

Cumulative rate of genotypic resistance to ADV at 5 years for HBsAg-positive and HBsAg-negative patients are 20% and 29%, respectively [41, 45]. The mutations associated with adefovir are N236T and A181V/T [46]. Viruses with these 2 mutations remain susceptible to entecavir whereas viruses with A181V/T mutations are cross-resistant to lamivudine [47].

Safety

The major side effect of ADV is nephrotoxicity. Studies using ADV 10 mg versus ADV 30 mg found that the higher dose is associated with greater impairment of renal function [39]. ADV at 10 mg is well tolerated and renal side effects are reported in 3% of patients with compensated liver disease after 5 years of treatment [41]. The renal impairment is reversible with dose reduction or drug withdrawal. For patients receiving ADV, renal function should be monitored every 3 months or more frequently if there is pre-existing renal impairment [33].

Current Role of Adefovir Therapy

With the availability of newer agents, such as entecavir and TDF with high antiviral efficacy and high genetic barrier, ADV monotherapy is no longer recommended as the first option for treatment-naive patients [33, 34].

In the case of lamivudine or telpivudine resistance, adefovir therapy is shown to be useful. It is recommended that ADV is added to lamivudine/telbivudine instead of switching to ADV monotherapy [43]. However, TDF is superior to ADV for these patients. In the case of renal impairment, ADV is best avoided. If it is used, dosage should be adjusted and close monitoring of renal function is necessary.

Telbivudine

Telbivudine was approved by the United States FDA for treatment of CHB in 2006. Telbivudine is the L-enantiomer of thymidine. After phosphorylation into active triphosphate form, it is incorporated into HBV viral DNA and causes termination of chain synthesis, thereby suppressing HBV DNA replication.

Efficacy and Resistance

In the GLOBE trial [48], 1367 patients including both HBsAg-positive and HBsAg-negative patients were randomized to telbivudine 600 mg daily and lamivudine 100 mg daily for 1 year. In HBsAg-positive patients, patients on telbivudine, when compared to lamivudine, are more likely to achieve undetectable HBV DNA (<300 copies/mL) (60% versus 40.4%) and HBsAg seroconversion (22.5% versus 21.5%). For HBsAg-negative patients, there was also a higher rate of undetectable HBV DNA for patients on telbivudine compared with lamivudine (88.3% versus 77.4%). Less patients on telbivudine developed drug resistance when compared to lamivudine in both HBsAg-positive (5.0% versus 11.0%) and HBsAg-negative patients (2.2% versus 10.7%). HBsAg-positive patients on telbivudine had better histologic response than those on lamivudine at 1 year (64.7% versus 56.3%). Another similar study performed in China also demonstrated superiority of telbivudine to lamivudine in terms of undetectable HBV DNA [49]. For chronic hepatitis B patients with cirrhosis, telbivudine can improve CTP score at 48 weeks of treatment [49, 50].

One of the major drawbacks with long-term telbivudine treatment is the development of drug resistance. The main mutation is rtM204I in the YMDD motif. Secondary mutations rtL80I/V and L80I/V + L180M may be associated with the rtM208I mutation. Unlike lamivudine, rtM204V is not associated with telbivudine resistance [48, 51]. After 2 years of therapy, the resistance is found in 25.1% of HBsAg-positive and 10.8% of HBsAg-negative patients [52].

Safety

Telbivudine is a pregnancy category B drug. In a study involving 61 pregnant women in China to determine the efficacy of telbivudine in decreasing maternal transmission of the hepatitis B virus to their infants, no adverse effects
were observed in both the mothers and the newborns in telbivudine-treated patients [55].

Current Role of Telbivudine Therapy

Both the American Association for the Study of Liver Disease and European Association for the Study of the Liver guidelines do not recommend telbivudine as the first line of treatment due to the high rate of drug resistance with long-term use [33, 34]. In multivariate analyses of the GLOBE study data, low baseline HBV DNA (<9 log10 for HBeAg positive patients, <7 log10 for HBe-negative patients) and undetectable HBV DNA at 24 weeks (<300 copies per mL) are associated with low rate of resistance at 2 years (HBeAg positive: 1.8%; HBeAg negative: 2.3%) [56]. A recent study showed that HBV DNA level of less than 200 IU/mL at week 12 of telbivudine therapy was also predictive of higher chance of undetectable HBV DNA (78.6%) and lower chance of resistance (0%) at year 3 [57]. Telbivudine may have a role in patients with these favorable characteristics.

Entecavir

Entecavir was approved by the FDA as the third nucleoside analog for the treatment of chronic hepatitis B in 2005. Entecavir is a guanosine analogue. It undergoes phosphorylation into di-phosphate and tri-phosphate metabolites, which is then incorporated into HBV DNA polymerase to inhibit viral replication. It also inhibits the priming of HBV DNA polymerase, a step which involves guanosine [58]. The inhibition of the priming of HBV DNA polymerase is unique among the currently licensed nucleoside/nucleotides.

Efficacy and Resistance

Entecavir has significant antiviral activity against chronic hepatitis B. In a double blinded prospective study, 715 HBeAg-positive patients were randomized to receive the standard doses of entecavir or lamivudine. More patients in the entecavir group achieve undetectable HBV DNA levels (67% versus 36%), alanine aminotransferase normalization (68% versus 60%) and histological improvement (72% versus 62%) at 48 weeks of treatment. The reduction of HBV DNA from baseline to week 48 was greater with entecavir than with lamivudine (6.9 versus 5.4 log copies per mL). HBeAg seroconversion rate at 48 weeks with entecavir group was 21%, while that for lamivudine group was 18% [59]. Entecavir is also effective for HBeAg-negative patients. In a parallel prospective study involving 648 HBeAg-negative CHB patients, who were randomized to the standard doses of entecavir or lamivudine for 52 weeks, more patients in the entecavir group achieved undetectable HBV DNA (90% versus 72%), normalization of ALT (78% versus 71%) and histological improvement (70% versus 61%). The mean reduction of HBV DNA was higher in entecavir group (5.0 versus 4.5 log copies per mL) [60].

While the efficacy of earlier generation nucleoside/nucleotide analogs decreases with time due to development of resistance, entecavir demonstrates potent antiviral efficacy for up to 5 years [61–63]. For patients taking entecavir up to 5 years, 94% achieved undetectable HBV DNA and 80% had normal ALT levels. After 96 weeks of treatment, 31% had HBeAg seroconversion; 5.1% also had HBsAg seroconversion. After 5 years, an additional 23% had HBeAg seroconversion; and 1.4% had HBsAg seroconversion [63].

Long term monitoring showed entecavir has a low resistance rate in treatment-naïve patients up to 5 years. The cumulative probability of genotypic entecavir resistance was 1.2% [64]. Entecavir resistance only occurs when, in addition to the two substitutions associated with lamivudine resistance (M204I/V +/− L180M), a third “signature” substitution develops at positions T184, S202 or M250 [65]. Therefore, for patients who have lamivudine resistance, there is lower genetic barrier for development of entecavir resistance. For this group of patients, the cumulative probability of genotypic entecavir resistance and genotypic resistance associated with virological breakthrough was 51% and 43%, respectively at 5 years. Because of this, entecavir is not recommended in patients with lamivudine resistance [64*].

Safety

Entecavir has similar safety profile with lamivudine. Serious adverse events are rarely seen [59, 60]. In a preclinical animal study, there is a higher incidence of species-specific solid tumors with doses of entecavir above the therapeutic range compared to placebo [66]. However, post-market surveillance up to 2009 failed to show any increase in incidence of malignancy with entecavir use [67]. Five cases of lactic acidosis on entecavir for decompensated HBV cirrhosis were reported [68]. However, the lactic acidosis was reversible in 4 of the patients. It is not certain whether the lactic acidosis was related to the multiorgan failure of the patients or to entecavir.

Current Role of Entecavir Therapy

Sustained effective virologic suppression is essential in prevention of complications of chronic hepatitis B. Entecavir has high antiviral potency, low resistance rate, and excellent safety profile with prolonged use. Entecavir is therefore one of the “ideal” treatment options for CHB, especially for treatment-naïve individuals. Both the
AASLD and EASL guidelines recommend entecavir as one of the first line agents for treatment of chronic hepatitis B [33•, 34•].

**Tenofvir Disoproxil Fumarate**

TDF is a nucleotide analogue initially approved for the treatment of HIV infection. It was approved by the United States FDA in 2008 for treatment of CHB. TDF is a prodrug of tenofovir. Tenofovir is phosphorylated into its active form and binds directly with viral polymerase and thereby suppressing viral replication [69, 70].

**Efficacy and Resistance**

Tenofovir and adefovir share similar molecular structure; tenofovir is found to be as potent as adefovir in an equal molar basis in in-vitro study. As tenofovir is less nephrotoxic, the approved daily dose of tenofovir is 300 mg which is more potent than adefovir at 10 mg. In a pivotal trial comparing tenofovir and adefovir, more patients achieved undetectable HBV DNA (HBV DNA <400 copies per mL) at 48 weeks in tenofovir group in both HBeAg-positive (76% versus 13%) and HBeAg-negative (93% versus 63%) patients [71]. All patients were put on open-label tenofovir after 48 weeks. At week 144 of tenofovir treatment, 93% of HBeAg-positive and 99% of HBeAg-negative patients have undetectable HBV DNA [72]. Furthermore, no resistance to tenofovir was detected up to week 192 [73•, 74, 75].

Several studies have demonstrated antiviral efficacy of tenofovir in lamivudine resistant patients. An Australian prospective study involving 60 patients with incomplete virologic response to both lamivudine and adefovir were given TDF or combination of TDF and lamivudine, 64% of patients achieved undetectable HBV DNA (<15 IU/mL) [76]. HBV mutation associated with adefovir resistant rtN236T and rtA181V/T are associated with decrease in response to tenofovir in in-vitro studies [77, 78]. However, clinical studies on tenofovir showed mixed results in adefovir resistant patients [76, 79, 80]. A retrospective multicenter study involving patients with treatment failure with lamivudine, adefovir, entecavir, either as sequential or add-on therapy, showed that TDF was less efficacious in patients with adefovir-associated mutations compared to lamivudine-associated mutations (52% versus 100%) in achieving undetectable HBV DNA (<15 IU/mL) [79]. However, no patients had virological breakthrough in the observed period of the study. In another German study involving 105 adefovir treated patients, 81% of patients have undetectable HBV DNA (<400 copies per ml) at week 48. The treatment response was not affected by baseline ADV- resistance or lamivudine resistance [80].

In decompensated liver disease, TDF has comparable improvement of virological, biochemical and clinical parameters compared to entecavir. Both agents achieved similar rates of HBV DNA undetectability (HBV DNA <400 copies per mL) at week 48, normal alanine aminotransferase and improvement of CTP score [81].

There is no report of virologic resistance of TDF in chronic hepatitis B monoinfection [73•]. In a study involving patients with HBV-HIV co-infection, rtA194T is found to be associated with reduced susceptibility to TDF in-vitro when lamivudine mutations rtM204V and rtL180m were present [82]. However, this is not confirmed by another study [83]. The role of rtA194T in TDF resistance is still unknown.

**Safety**

From the experience in HIV infection, tenofovir is associated with a small decline in renal function of 9.8 mL/min/1.37 m2 in 5 years [84]. However, large scale clinical studies showed there is no evidence of compromised renal function with tenofovir in patients with chronic hepatitis B monoinfection [72]. TDF is also associated with renal proximal tubular disfunction and renal phosphate wasting [85]. Cases of acute renal failure and nephrogenic diabetes insipidus have been reported [86]. Renal toxicity is usually reversible when tenofovir is stopped.

TDF is a pregnancy category B drug. There is no increase in rate of birth defects in HIV infected mother taking tenofovir compared to baseline [34].

**Current Role of TDF Therapy**

TDF is recommended as the one of the first line agents for treatment naïve patients. Given the potent antiviral efficacy, low resistance rate and minimal toxicity, it can achieve long term effective HBV DNA suppression and has the potential to become one of the “ideal” treatment options for chronic hepatitis B. Tenofovir is also useful in patients with lamivudine, telbivudine and entecavir resistance.

**Conclusions**

Over the past 10 years, there has been considerable improvement in the treatment of chronic hepatitis B. Interferon-based immunomodulatory therapy is gradually being replaced by nucleoside/nucleotide analogs which have better antiviral potency and safety profiles. Newer nucleoside/nucleotide analogs including entecavir and TDF have potent antiviral potency and high genetic barrier to resistance, and thus are superior to agents such as lamivudine, telbivudine, and adefovir.
Current treatment strategy of chronic hepatitis B is to achieve prolonged and effective viral suppression [18]. Due to the presence of intracellular covalently closed circular DNA in hepatocytes, there is a chance of viral rebound if NA is discontinued even when the serum HBV DNA have declined to levels below the detectable limits of polymerase chain reaction assays.

With the availability of entecavir and more recently TDF, prolonged effective viral suppression can be achieved with long-term treatment. It is likely that the complications with CHB including cirrhosis and hepatocellular carcinoma will decrease in the future with these newer agents.

However, more studies on the safety profiles and efficacy on long term use of these newer agents are needed.

Disclosure  Ching-Lung Lai acted as speaker for GlaxoSmithKline, Bristol Myers-Squibb and Gilead; Man-Fung Yuen acted as consultant/speaker and received research grants from Bristol Myers-Squibb, GlaxoSmithKline, Novartis and Roche Diagnostics; Yuk-Fai Lam and Wai-Kay Seto reported no potential conflicts of interest relevant to this article.

Open Access This article is distributed under the terms of the Creative Commons Attribution Noncommercial License which permits any noncommercial use, distribution, and reproduction in any medium, provided the original author(s) and source are credited.

References

Papers of particular interest, published recently, have been highlighted as:

- Of importance

18. Lai CL, Yuen MF. Chronic hepatitis B—new goals, new treatment. N Engl J Med. 2008;359:2488–91. This article suggests the new goal of chronic hepatitis B treatment should be prolonged and effective viral suppression. The authors also provide evidence to support this.
33. • Lok AS, McMahon BJ. Chronic hepatitis B: update 2009. Hepatology. 2009;50:661–2. This is the American Association for the Study of the Liver Diseases Guideline, which gives a comprehensive overview on the management of chronic hepatitis B.
34. • EASL Clinical Practice Guidelines: management of chronic hepatitis. B. J Hepatol. 2009;50:227–42. This is the latest European Association for the Study of the Liver Guideline on the management of chronic hepatitis B.


74. Heathcote EJ, Gane EJ, DdeMan RA, et al. Long term (4 year) efficacy and safety of tenofovir disoproxil fumarate (TDF) treatment in HBeAg-positive patients (HBeAg+) with chronic hepatitis B (study 103); preliminary analysis (abstract). Hepatology. 2010;52(4(Suppl)):556A.


