Why adefovir is not yet available in Canada

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The introduction of lamivudine for the management of chronic hepatitis B virus (HBV) infection heralded a new era in the management of the disease. Until then, the only available therapy was interferon-alpha (IFN-α), which was expensive and difficult to take. It required three injections a week for four months to two years for some forms of HBV. IFN-α was able to induce a response in a small proportion of patients. In hepatitis B e antigen (HBeAg)-positive patients, seroconversion from HBeAg to the antibody for HBV (anti-HBe) occurred in just over 30% of treated patients (1,2). Alanine aminotransferase (ALT) normalized and HBV DNA became undetectable in serum using the insensitive assays at the time. In the anti-HBe-positive patients, the durable response of therapy was approximately 20%, again using the insensitive solution hybridization assay for HBV DNA (3-5).

Lamivudine is an oral agent with only minor side effects. The registration trials lasted one year, with a disappointing HBeAg seroconversion rate of approximately 18% to 20% (6). In the anti-HBe-positive patients, adequate viral suppression was achieved on therapy, but after withdrawal of lamivudine, more than 75% of patients relapsed (7,8,9). However, suppression of viral replication could be maintained beyond the first year by continued therapy (10). This was usually associated with improved ALT levels and improved histology (11). Until recently, there was no information that lamivudine had any effect on hard end points such as survival. Last year, however, data were published showing that long-term lamivudine use in cirrhotic patients with active viral replication decreased the incidence of liver-related complications such as hepatoma, liver failure and liver-related death (12).

Lamivudine use, however, has been severely restricted by the development of resistance. In a study by Hadziyannis et al (13), within the first year of use approximately 18% of infections became resistant to lamivudine, and over four years approximately 70% of patients had developed resistance. Lamivudine resistance is associated with a rebound in the concentration of virus in serum and, usually, with an increase in ALT and histological disease progression. Initially, it was thought that the development of resistance did not affect the rate of progression of disease, but recent evidence has disproved that hypothesis (12,14,15). In fact, in patients with cirrhosis who develop lamivudine-resistant infection, the rate of progression to death is rapid and this has become a feared outcome.

There are now a number of newer agents for use in HBV patients. These include adefovir (16,17) and entecavir (18,19), both licensed in many parts of the world but not in Canada. Other drugs in the pipeline include telbivudine, pradefovir, emtricitabine and clevudine. Although tenofovir is not licensed for HBV, it is also a potent anti-HBV drug and is available in Canada (20). It has been used to treat lamivudine-resistant HBV infection. Adefovir was the first drug after lamivudine to be commercialized. In the United States, adefovir was first licensed in 2002. It maintains full antiviral activity in lamivudine-resistant infection, so that in addition to being used as a first-line drug, it has been used to rescue lamivudine-resistant infection (21). Adefovir resistance does occur, although at a much lower rate than lamivudine. Hadziyannis et al (13) showed that in the first year of therapy, the resistance rate was in the range of 1% to 2% and after five years, the resistance rate was approximately 28% compared with 70% for lamivudine. Although adefovir is less potent than lamivudine, it has been recommended for first-line use because of the lower rate of resistance, particularly in anti-HBe-positive chronic HBV, where treatment may have lasted for many years (22).

Adefovir has been available in the United States for more than three years. It was submitted for licensing in Canada in June 2002 and received the notice of compliance in August 2003, a little over one year later. However, more than two years later, it is still not available in Canada. The reason is that the Patented Medicines Price Review Board (PMPRB) has not allowed Gilead Sciences Inc (USA), the manufacturer, to charge the price it wishes.

The PMPRB negotiates with pharmaceutical manufacturers to set drug prices in Canada. The drugs are reviewed using several principles; if the drug is used to treat a disease for which there are existing treatments of equivalent efficacy (eg, another statin or histamine 2 receptor blocker), the price should be in the same price range as other drugs used to treat the same disease. If the drug is a breakthrough drug (eg, for a previously untreatable disease or greatly improved efficacy over existing treatments), the price charged in Canada should be no more than the median of prices charged in seven industrialized countries (USA, Italy, France, Sweden, Switzerland, United Kingdom and Germany). Gilead Sciences Inc, the makers of adefovir, sought presale advisory assistance from the PMPRB, and on the basis of that advice, decided not to proceed with marketing in Canada (recently Gilead Sciences Inc has revised its approach and is undertaking premarketing activities). The substance of these discussions is not public, but it seems that the price that the PMPRB was prepared to allow was insufficient for the manufacturer. One can only presume that the PMPRB reviewed adefovir as a ‘me too’ drug so that the set price would be approximately the same as for lamivudine. This was not acceptable to Gilead Sciences Inc because the cost of one month’s therapy in, for example, the United States was at

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three times higher than that of lamivudine. Hence, the drug is not being sold in Canada.

Gilead Sciences Inc has, however, established an early access program that allows some physicians access to free drugs. As of November 16, 2005, 812 patients are receiving free drugs. Essentially, this is a gift on the part of Gilead Sciences Inc to Canada, the value of which (assuming on average one year of use) exceeds $5 million.

There are several reasons why the PMPRB decision must be revisited. Although we have no data, there must be thousands of patients with chronic HBV who have been treated with lamivudine since it was first licensed in approximately 1998. For 70% of those patients, long-term therapy is required and in approximately 70% of that proportion, lamivudine resistance must have developed. Thus, these patients are either without treatment or are continuing to take treatment that is ineffective. For those patients who were cirrhotic at the start of therapy (an unknown proportion), the risk of hepatic decompensation is high. These patients can be rescued with adefovir and if maintained on both adefovir and lamivudine, they may have a low likelihood of developing further resistance (at least over the first few years).

It is scandalous that the drug approval process in Canada is so slow. It took 13 months to review the data and approve the drug. Pegylated IFN-α-2a plus ribavirin for hepatitis C took three years to be approved. This is not acceptable. In the United States, the review process takes six months. Why it is necessary in Canada to repeat the review previously performed elsewhere is inexplicable. There is no need to reinvent the wheel. Both the Food and Drug Administration (FDA) and European agencies undertake extremely detailed and critical reviews. A look at the FDA website for any particular drug clearly shows the depth of the analysis. Health Canada does not provide details of the review process so it is not possible to assess the details of the review, but it is surely unnecessary to repeat what the FDA and the Europeans have done.

In the case of adefovir, the delay in access is particularly ironic because many of the patients who were studied in the registration trials came from Canada. Prominent Canadian hepatologists, Dr Jenny Heathcote in particular, played an important role in the development of this drug. Yet, it is still not available here.

In addition, the PMPRB review has been conducted in ignorance of the importance of adefovir. There are no hepatologists, infectious disease experts nor anyone who can provide informed advice on the board. No external consultations with such experts appear to have been undertaken. When questioned about the decision, the chairwoman of the board indicated that Gilead Sciences Inc was indeed free to sell adefovir and could charge whatever price they chose. However, Gilead Sciences Inc would have to refund the difference between the approved price and the actual price. This is a disingenuous answer designed to absolve the board of responsibility for the lack of availability of adefovir. After an initial protest against the decision by some hepatologists and by the Canadian Liver Foundation (Toronto, Ontario), the PMPRB contacted Gilead Sciences Inc for further discussions, the outcome of which is unknown.

Between them, Health Canada and the PMPRB have, through ignorance and inefficiency, created a situation in which a valuable drug is not available to Canadians who need it; patients for whom there is no other drug substitute. In my practice alone, there has been serious morbidity, and elsewhere, quite likely some deaths because adefovir was not available. It is particularly hard to swallow this decision, given that trastuzumab (Herceptin, Hoffmann-La Roche Ltd, Canada), at more than $100,000 for a one-year course, has been granted pricing approval. Other equally expensive drugs for HIV are also available (eg, tenofovir).

This is a situation that must be remedied. There are additional HBV drugs coming. They will likely also be expensive. However, as with HIV, we need to have an armamentarium of different drugs with different resistance profiles available to provide to our patients if and when resistance to current therapy emerges. In the near future, we will be using combination therapy rather than mono therapy and again will need to be able to select from a pool of available drugs, depending on pre-existing resistance profiles. Entecavir will be the next agent up for consideration. If the same principles are applied by the PMPRB, this drug will not receive pricing approval either. Yet, entecavir is the most powerful drug in its class (23). It has the best antiviral effect and the best resistance profile so far. This in itself will make it a valuable drug. We cannot wait another four to five years for entecavir to become available.

The PMPRB should be pressured to reconsider the pricing of adefovir and to do so with some urgency. Indeed, the whole drug review process in Canada needs to improve.

The views expressed in the article are the personal views of the author and do not necessarily reflect the views of either the Canadian Association for the Study of the Liver or Pulsus Group Inc.

Dr Sherman is on the advisory boards of Gilead Sciences Inc in Canada, Roche Canada and Bristol Myers Squibb, and has given sponsored talks for Roche and Bristol Myers Squibb.
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