Hepatitis delta virus (HDV) is a cause of acute and chronic liver disease for which no effective therapy is currently available. Previous research has demonstrated that prenylation of the large HDV antigen is essential for viral assembly. A new report (see the related article beginning on page 407) describes a novel small animal model for HDV replication and demonstrates that prenylation inhibitors are highly effective at clearing viremia and thus have potential relevance for the treatment of chronic delta hepatitis.


Delta hepatitis is the least frequent but most severe form of viral hepatitis. Acute infection with hepatitis delta virus (HDV) is often severe and has a fatality rate of at least 5%, even in children. Chronic delta hepatitis leads to cirrhosis in at least two thirds of persons and can present as end-stage liver disease in early adulthood (1). Hepatitis D also has the distinction of being the form of viral hepatitis that is most resistant to current antiviral therapies (2).

Virology of HDV

The unique clinical features of delta hepatitis are matched by the unique qualities of its viral agent. HDV is the smallest known human pathogen and uses a unique replicative strategy (3). The HDV genome is merely 1.7 kb in length and consists of a single-stranded circular molecule of RNA of negative polarity that is held in a rigid rod-like form by extensive intramolecular base pairing (4). The antigenome of HDV RNA has a single open reading frame that encodes hepatitis delta antigen. Importantly, HDV requires the presence of another (helper) virus for productive replication and spread (5). The helper virus is the hepatitis B virus (HBV), a very significant human pathogen in its own right. Being linked to HBV and only occurring in persons who have hepatitis B surface antigen (HBsAg) in serum, HDV is spread in the same ways as hepatitis B, through parenteral or sexual exposure to blood or body fluids. Up to 5% of the world’s population is infected with HBV, and probably 5% of these have HDV superinfection (6). Delta hepatitis has a worldwide distribution, but areas of dramatically high endemicity associated with severe clinical outcomes have been identified in the Amazon basin, Central Africa, and parts of Eastern Europe. Prevention of hepatitis B will also prevent hepatitis D. In a triumph of public health, increasing vaccination rates for hepatitis B have led to a dramatic decline in HDV incidence. Nevertheless, there are still millions of persons with chronic HDV infection, and until hepatitis B is fully eradicated, new cases of delta hepatitis will occur.

HDV circulates in serum as a 35- to 40-nm particle containing viral RNA and HDV antigen, encapsidated by HBsAg (7, 8). The envelope protein of HBV allows for the specific binding and uptake of virions into hepatocytes. Inside the cell, the viral genome is released and replicated by a double-rolling circle mechanism (4). Replication of HDV, like that of other viruses, requires multiple enzymatic activities. Although many viruses encode the majority of their replicative and processing enzymes, HDV apparently does not. The sole enzymatic activity HDV possesses is a ribozyme (similar to that in plant satellite viruses) that autocleaves the circular RNA, producing a linear molecule (9). Other enzyme activities are apparently provided by the host cell.

Address correspondence to: Jay H. Hoofnagle, Liver Diseases Research Branch, National Institute of Diabetes and Digestive and Kidney Diseases, National Institutes of Health, Bethesda, Maryland, USA.

Phone: (301) 496-1333; Fax: (301) 480-7926; E-mail: hoofnaglej@extra.niddk.nih.gov.

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Nonstandard abbreviations used: hepatitis delta virus (HDV); hepatitis B virus (HBV); hepatitis B surface antigen (HBsAg); small HDV antigen (S); large HDV antigen (L).
The prenylation of the L HDAg and show how this knowledge can be used to develop therapies for delta hepatitis. Since prenylation is critical for virion formation, prenylation inhibitors might be effective in blocking viral replication. The hurdle for demonstrating activity of prenylation inhibitors (many of which were developed as anticancer agents) (13) was the lack of availability of a practical model for studying the viral life cycle. To this end, the authors developed a murine model using hydrodynamic transfection of HDV cDNA into an established HBV transgenic mouse. They reasoned that if HBsAg was provided, introduction of HDV RNA might result in HDV viremia. This was indeed the case. The authors then showed that viremia was blocked by prenylation inhibition.

At present, therapy of delta hepatitis is difficult and unsatisfactory. The nucleoside analogues that block HBV replication have little or no effect against HDV replication or the disease that it causes. IFN-α inhibits HDV replication, but therapy usually requires high doses given for a prolonged period, which are often poorly tolerated (2). The work by Bordier et al. (12) can be viewed as a new phase of therapeutics for HDV. In the typical first stage, known and established antiviral drugs are taken off the shelf and tried. Such was the case with studies of IFN-α and the nucleoside analogue lamivudine for HDV (2, 14). In the second stage, fundamental knowledge about the virus is applied, and drugs are used on the basis of such knowledge. With prenylation inhibitors, hepatitis D may enter the second stage of drug discovery, an excellent example of translational research. For the third phase, drugs are specifically designed and synthesized on the basis of the structure and life cycle of the virus. Delta hepatitis may be the least common form of viral hepatitis, but in many ways it is highly deserving of focused drug discovery.


Figure 1
Replication of HDV. Viral particles consist of HDV RNA (black lines) and HDV antigen (blue) encapsidated by HBsAg (red). The HBsAg allows for binding and uptake of the virion into hepatocytes. Inside the cell, the RNA genome (negative polarity) is released and undergoes replication by the double-rolling circle pattern (double arrows). The antigeneomic RNA (positive polarity) serves as the template for production of more genomic RNA but can also be edited by intracellular host dsRNA adenosine deaminase, changing A to I so that with production of genomic RNA, C rather than U will be transcribed (right side of the figure). The editing changes the UAG stop codon on the positive strand to a UGG, which is read as tryptophan, resulting in a protein with an additional 19 amino acids, the large delta antigen (large δ) that inhibits replication. Small delta antigen (small δ), the product of the unedited transcript (left side of the figure), binds to HDV RNA and promotes replication. Large delta antigen has a prenylation site that creates a hydrophobic molecule that binds both to HDV RNA and to HBsAg, leading to viral assembly and release. Thus, inhibition of prenylation will interfere with viral assembly and release but should affect replication minimally, if at all.

depends on cellular enzymes that the virus redirects toward production of viral antigen and virions. HDV also clearly uses other cellular enzymes, not all of which are well defined but perhaps include a helicase to unwind the intramolecular base pairing of the HDV RNA circular genome and a polymerase for production of viral RNA. These findings are important because knowledge of the viral life cycle can point to targets for antiviral therapy. The difficulty (and unique feature) of delta hepatitis is that most of the revealed targets are normal cellular proteins. Thus, HDV hides inside the envelope of HBV as well as behind normal cellular enzymes: the wolf in sheep’s clothing.

Inhibition of HDV by inhibition of prenylation
In this issue of the JCI, Bordier et al. (12) extend the previous studies of
Alzheimer disease (AD) is characterized by the progressive accumulation of amyloid β protein (Aβ) in areas of the brain serving cognitive functions such as memory and language. The first of two separate reports (see the related articles beginning on pages 415 and 440) reveals that intrinsic T cell reactivity to the self-antigen Aβ exists in many humans and increases with age. This finding has implications for the design of Aβ vaccines. The second report demonstrates that a number of FDA-approved nonsteroidal anti-inflammatory drugs are capable of lowering Aβ levels in mice. The work suggests that further testing of the therapeutic utility of these types of compounds for the potential treatment of AD is warranted.


Alzheimer disease (AD) has received a lot of recent attention, particularly in areas related to novel treatments. Recently, the potential therapeutic usefulness of the immune system has become apparent, leading to the question of whether it can be used to directly or indirectly influence AD-related pathology in beneficial ways.

Active immunization with amyloid β (Aβ) peptides takes advantage of the immune system to generate antibodies that can somehow decrease Aβ-related pathology in mouse models of AD (1). Similarly, passive immunization involves direct administration of anti-Aβ antibodies, bypassing the need for an active immune response (2, 3). Since genetic, pathologic, and animal studies suggest that the buildup of Aβ in the brain leads directly or indirectly to cell dysfunction, cell death, and cognitive impairment, increased generation of anti-Aβ antibodies has the potential to prevent or treat AD by decreasing amyloid burden and its consequences in the brain. Though the first clinical trials for Aβ vaccination were halted due to CNS inflammation in a small subset of subjects, active and passive immunization strategies remain a viable potential therapy worth continued exploration. If positive effects can be seen in future trials, it will be important to minimize unwanted toxicity. In this issue of the JCI, Monsonego and colleagues (4) further characterize the innate immune response to Aβ in humans, thus revealing important details about how the elderly body reacts to Aβ, and opening new avenues to modify existing vaccination protocols. Also in this issue, Eriksen and colleagues (5) studied traditional NSAIDs that appear to have a nontraditional, COX-independent effect on decreasing Aβ42 production. While these drugs are often used to treat inflammation, they appear to have a novel effect on amyloid precursor protein (APP) cleavage, which is only now becoming apparent and which may be useful in the future as a therapeutic.

Aβ-reactive T cells increase with age

Monsonego et al. (4) found that some healthy, elderly individuals, as well as individuals with AD, contain elevated baseline levels of Aβ-reactive T cells. While the general trend is toward a diminished immune response with aging, this demonstrates a selective increase in Aβ-reactive T cells in older individuals with and without dementia. The reason for this selective expansion of Aβ-reactive T cells in elderly individuals remains unclear. It is often presumed that cognitively normal middle-aged and elderly individuals are similar in that they lack AD pathology; however, Aβ deposition in plaques appears to begin about 10–20 years prior to the onset of even the earliest symptoms sugges-