The potential threat of the smallpox virus as a bioterror weapon has long been recognized, and the need for developing suitable countermeasures has become especially acute following the events of September 2001. Traditional antiviral agents interfere with viral proteins or functions. In a new study, Yang et al. focus instead on host cellular pathways used by the virus. A drug that interferes with the cellular ErbB-1 signal transduction pathway, activated by smallpox growth factor, sheds new light on how the virus replicates in the cell. Drugs that target the ErbB-signaling pathways represent a promising new class of antiviral agents.
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Developing medical interventions against the threat of bioterrorism has been a part of the national biomedical research agenda at least since the days of the Cold War. However, the immediacy of such a threat was not fully appreciated until the events of the fall of 2001. The attacks on the World Trade Center and the Pentagon and the subsequent release of anthrax spores that infected 22 people and killed 5 transformed a remote possibility into somber reality (1).

Of special concern are category A agents and the need to develop countermeasures against these agents, which are known to cause fatal diseases such as anthrax, smallpox, plague, botulism, tularemia, and viral hemorrhagic fevers (2). The need for countermeasures against the Variola major virus, which causes smallpox, is especially acute because the virus is highly lethal and easily transmitted from person to person and because no effective treatments have yet been approved for use in humans. As a result of the successful global eradication of smallpox in 1977, routine vaccinations have been discontinued, and most people today have little or no immunity against the virus. Concerns about unaccounted-for stocks of smallpox virus surfaced after the fall of the Soviet Union, when it was revealed that massive quantities of the virus had been produced, further heightening the sense of urgency for the development of new therapies (2–4). An effective smallpox vaccine exists, and several new vaccine candidates are now being tested. Should a deliberate release of smallpox occur, however, it is important to have drug treatments readily accessible to protect against disease and to reduce any adverse effects of a live or attenuated smallpox vaccine. Currently, no such treatments are available, although cidofovir, a nucleoside analog that inhibits viral replication, is being tested clinically for treatment of some poxvirus infections (5, 6).

CI-1033 prevents viral replication in vitro

Traditional antiviral drugs are generally directed against the proteins and functional pathways of the virus itself. For example, numerous therapies have been developed against HIV that interfere with viral reverse transcriptase, protease, and integrase as well as viral components involved in the binding and fusion of the virus to the target cell (7). However, many viruses evolve rapidly, particularly under selective pressures, and drug resistance almost invariably develops.

Because viruses typically rely upon cellular pathways to self-propagate, another antiviral approach would be to develop drugs that interfere with viral functions that are dependent on the functional machinery of the cell. Such an approach has been adopted by Yang et al., who report in this issue of the JCI (8) on a class of drugs originally developed as antitumor agents that show promise against orthopoxviruses and perhaps other viruses as well.

It has previously been shown that a growth factor (GF) encoded by the genomes of all orthopoxviruses (smallpox growth factor [SPGF] by variola virus, vaccinia growth factor [VGF] by vaccinia virus) binds to and activates the ErbB-1 kinase, a member of the epidermal GF receptor family of tyrosine kinases (9, 10). Because the poxvirus-encoded GFs are important for viral pathogenesis (11, 12), it seemed likely that inhibiting the cellular GF receptor might be a useful approach to controlling poxvirus infection. Yang et al. now demonstrate that inhibitors of cellular ErbB-1 do in fact disrupt important processes of the viral replication cycle and may represent an important new approach to antiviral chemotherapy (8).

This paper also provides new insight into the role of the poxvirus-encoded GFs in viral pathogenesis, suggesting that poxvirus GFs may play a direct role in virus replication. Previous studies have suggested that VGF acts on cells to stimulate metabolism, thereby increasing the number of cells capable of supporting efficient viral replication (11). Yang et al. (8) examined the effect of the ErbB inhibitor CI-1033 on the growth of variola and vaccinia virus—a smallpox-like virus—in infected monkey kidney cells in vitro. The drug had no effect on the overall yield of newly made virus in cell culture experiments in which all the cells in the culture were infected simultaneously, but it did have an effect on the appearance of plaques, which arise from the initial infection of a single cell and require local spread of the virus from the infected cell to surrounding uninfected cells. Two distinct forms of infectious virions are produced in poxvirus-infected cells: intracellular mature virus (IMV), which is released only following death and lysis of infected cells, and extracellular-enveloped virus (EEV), which is actively extruded from cells by interaction with actin tails (13) (Figure 1). The release of EEV from infected cells is thought to be the principal mechanism for rapid spread of the virus in the infected host. Yang et al. (8) show that the ErbB inhibitor CI-1033 greatly reduces the release of EEV from cells infected with either vaccinia or variola. This reduction in EEV release is likely due, at least in part, to inhibition of the viral GF activation of ErbB-1 because, even in the absence of CI-1033, deletion of the GF gene from vaccinia virus has an effect...
on the release of EEV similar to that of CI-1033 on wild-type virus. However, the drug appears to have additional antiviral effects when added to cells infected with mutant virus as shown by the reduction in the size of plaques. It has recently been shown that actin tail formation, which appears to be involved in EEV release, is regulated by phosphorylation of the viral morphogenesis protein A36 by cellular Src (c-Src) (14, 15).

Since activated ErbB-1 is known to activate c-Src (16), this provides one possible explanation for the role of GF in EEV release. Other possible targets for GF-activated ErbB-1 are indicated in Figure 1.

**CI-1033 helps control viral infection in vivo**

In mice infected with vaccinia, treatment with CI-1033 clearly altered the course of disease (8). Mice were first infected with virus and treated with CI-1033 in the presence and absence of a monoclonal antibody that neutralizes the IMV form of infectious virions. At moderate challenge doses of virus (close to the LD$_{50}$), CI-1033 treatment greatly increased animal survival and had a moderate effect on viral load in the lung. A much greater effect of the drug was seen when it was administered in conjunction with monoclonal antibody. This result is consistent with what is known about immunotherapies for poxvirus disease: maximum protection is seen when both EEV and IMV are neutralized (17, 18). Interestingly, the combination of CI-1033 and anti-IMV antibody also significantly stimulated T cell immunity in the infected mice (8). The mechanism of this effect remains to be explored.

**A new class of antiviral drugs**

The concept of interfering with the dependence of virus replication on cellular machinery is not new. For years, HIV/AIDS researchers have attempted to blunt viral replication by interfering with the cascade of aberrant activation signals that make the cell permissive for viral replication (19, 20). The benefit of using drugs such as cyclosporin and mycophenolate, which suppress immune cell activation, to treat HIV infection remains questionable because of the chronic nature of the disease; however, such an approach may be an effective strategy for treating an acute infection, such as smallpox.

In essence, the approach suggested by Yang et al. (8) may serve to turn the tables on the virus by interfering with the very pathways that are required for viral replication and extrusion. Because it is less likely that such a strategy would allow the virus to develop drug resistance and because different members of a virus family will likely use the same host pathways to propagate, this strategy approaches the concept of a “universal” antiviral therapy, at least within the virus family in question. Indeed, replication of many viruses, including poxviruses, is known to be dependent upon the ErbB class of tyrosine kinases, and the screening for and development of drugs that block this and other classes of receptors should be vigorously pursued. This strategy would be especially useful in developing countermeasures against newly emerging infectious diseases and those that are introduced deliberately, as in a bioterror attack.

Over the past several decades, thousands of promising anticancer drugs, including CI-1033, have been developed by pharmaceutical companies to interfere with GF-mediated...
ed signal transduction cascades such as those coordinated by the ErbB class of receptor tyrosine kinases (21, 22). Yang et al.’s study (8) shows that variola and related viruses are dependent upon some of the same pathways that the host cell uses for growth and development. Inhibitors of the ErbB-1 pathway as well as other cell-signal transduction pathways required for viral replication represent a largely untapped source of potential antiviral drugs and merit further exploration.

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Cytokines regulate an array of biological processes by activating cell surface receptor complexes, a process that initially involves oligomerization and activation of the JAK family of tyrosine kinases. In turn, JAKs phosphorylate the cell surface receptor, and signaling proteins such as STATs are recruited to these phosphotyrosine sites on the receptor; the proximity of the STATs allows them to be phosphorylated by the JAKs (Figure 1). Dimerization of the phosphorylated STATs leads to nuclear migration and regulation of gene expression (1). To control excessive cytokine effects, the cytokine signal is negatively regulated by a number of proteins, including protein tyrosine phosphatases such as Src homology 1–containing tyrosine phosphatase (SHP1), protein inhibitor of activated STAT (PIAS), and SOCS (2). The latter family is comprised of cytokine-inducible SH2-containing protein and SOCS1–SOCS7. SOCSs are furthermore induced by cytokine signaling and therefore form a closed-loop, negative-feedback control mechanism (Figure 1).

SOCS2, a new player in growth hormone receptor signaling
While cytokines and their receptors have traditionally been the domain of immunol-