Letters

HIV Infection in the Laboratory

In their report "Risk of human immunodeficiency virus (HIV-1) infection among laboratory workers," Stanley H. Weiss et al. (1 Jan., p. 68) make repeated reference to "concentrated virus." We wish to remind readers that concentration of an infectious agent is far less important than its pathogenicity. The strain that has infected two laboratory workers to date is presumed to be a strain of HIV-1, designated HTLV III_B whose clinical origins are obscure. Another strain of HIV-1 called LAV, with a genome virtually identical to HTLV III_B (I), was isolated from an individual who is still alive and reported well at least 6 years after the initial isolation (2). It may be, therefore, that the pathogenicity of these particular strains is low. However, this does not mean that individuals will only become infected by large amounts of virus, but rather that only a few virus particles will succeed in being infectious. Any laboratory worker exposed to any amount of virus is at risk, whether the virus is "concentrated" or not.

We heartily concur that all laboratory workers should receive quarterly testing and suggest both a sensitive enzyme-linked immunosorbent assay (ELISA) and a Western blot in conjunction with appropriate counseling and confidentiality of test results. The beneficiaries of testing will be not only laboratory workers themselves but their loved ones.

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Response: Our focus on potential exposure to "concentrated" virus arose because of the fact that both the worker reported in the cohort study and the second worker shared laboratory procedures in different settings that involved the handling of large volumes of concentrated virus (1). It is well recognized from infectivity studies that potential human pathogens vary in their infectious dose. Thus some agents require fewer than ten infectious organisms to result in a human infection while others require significantly larger doses (for example, more than 10⁶ infectious organisms). For HIV the number of particles needed to cause infection in humans has not been quantified, but it would not appear to be one of the more infectious agents to which health care and laboratory workers are exposed. This is evidenced by the failure in numerous studies to document "casual" household transmission, the rarity of infections resulting from parenteral inoculation in the hospital setting, and the fact that, in our cohort, among the ten workers experiencing parenteral exposure to potentially infectious apparatus, none seroconverted, although the second worker who did seroconvert, who was not in our cohort, did experience parenteral exposure (1-3). It is recommended that biosafety level 3 practices and containment be followed for HIV, since it is a dangerous human pathogen. In our report we emphasized certain biosafety practices to prevent inapparent exposures or unnecessary risk resulting from the use of glass or sharp instruments, in agreement with the concerns of Fox and Fox for careful biosafety practice.

With regard to the speculation about laboratory strain variation and disease pathogenicity, HTLV-III_B differs from the LAV strain by 144 nucleotides (4), and HTLV- III_{B} has a polymorphic variant of the R gene different from that shared by LAV and ARV (5). Whether there are less pathogenic strains of HIV, as suggested from the follow-up of Clavel's LAV patient and in vitro correlates of the original 48 HIV isolates (3), or whether host or other factors explain different rates of disease progression, remains to be established. While the strain isolated from the laboratory worker was indistinguishable from a subclone of HTLV-III_B, it is noteworthy that this T lymphocyte-adapted laboratory strain could be isolated only from monocyte-macrophages of the individual (I). Careful molecular analysis of the isolates from the worker are under way to search for subtle changes in the nucleotide sequence of the virus that may explain this altered tropism. The difference between LAV and HTLV-III_B is significantly greater than that between HTLV-III_B and the isolate from the laboratory worker. Thus, if even subtle mutation can result in changes in tropism or pathogenicity, or both, then the apparent lack of pathogenicity noted for LAV may not be relevant for HTLV-III_B.

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Historiographic Distinctions

In his review (8 Jan., p. 198) of my book Darwin and the Emergence of Evolutionary Theories of Mind and Behavior (1), John Greene states: "Human nature has dimensions that escape, and must forever escape, the abstractions of science." I rather believe there is no other way to knowledge than the kind of thinking that drives science. On several points of fact and logic, though, I believe Greene has attempted an alternative.

Greene writes that "Richards fails to distinguish" between considerations that led Lamarck and Darwin to adopt an evolutionary theory and those that led them to advance certain mechanisms to explain species change. The distinction, a standard one, I most assuredly made, and precisely in those terms he suggests I neglected (1, pp. 47-48 and 79-81). I even referred to Greene's own theory about what led Lamarck to his initial formulation (1, p. 47). Greene's remarks about Herbert Spencer, a figure who quickly polarizes historians of biology, epitomize the difficulties I have with his review. Greene quotes me as praising Spencer's entire philosophical-scientific system-with the implication that anyone would be foolish to do so. But the truncated quotation he uses refers only to Spencer's ethical notions, especially of justice and altruism (I, p. 303). Greene writes that "Spencer himself eventually admitted that his ethical principles and social theory did not require evolutionary biology as a foundation" and uses this supposed admission to rebut my argument that Spencer's ethical ideas determined his evolutionary theory. I do not know on what grounds Greene bases this statement. In an earlier essay, he surmised: "The truth of the matter is that [Spencer's] social ideal had never really been grounded in biological science, much as he liked to pretend that it was" (2). A historian's surmise about Spencer is quite different from "Spencer himself eventually admitted. . . ." (Another assertion attributed to Lamarck is, I believe, a surmise.) The last phrase in the quotation from his essay indicates that at the time Greene himself believed Spencer never "eventually admitted...." Moreover, in Spencer's last major ethical work, Principles of Ethics (1893), which I discussed at length, he explicitly sought to derive his basic moral principles from evolutionary laws. In a way, Greene's series of counterclaims is beside the point. My primary thesis in the chapters on Spencer was not that his ethics depended on his evolutionary theory; it was that his evolutionary theory depended on his ethics. It is a simple error to render these two relations of dependency as logically equivalent.

Although Greene generously appraises