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The Limits of Government Regulation of Science

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The court ruled that federally funded scientific research, especially at universities, should be free from prior restraint—calling into question the validity of CUI conditions on research grants.

advocates believe the greater risk is that the mutated virus could escape or that knowledge about these mutations could get into the wrong hands. They suggest that research of this kind should not be funded or undertaken in the first place. Where, as here, the research has already been conducted, they urge scientific journals not to publish any sensitive methods or results (1).

The HHS request reveals a troubled relationship between security and science. This is not the first time a government has requested that a journal not publish information. In 1979, the U.S. Department of Energy secured an injunction against the magazine The Progressive to prevent the publication of an article about building a hydrogen bomb, even though the information was in the public domain; the injunction was later vacated when the article was published elsewhere (2). In 2005, the Proceedings of the National Academy of Sciences refused to comply with an HHS request to decline publishing a mathematical model of botulism in the milk supply (3). The H5N1 case, however, is the first time government has sought to redact information after an institutionalized HHS review process.

Constitutional Limits on Government Restrictions of Scientific Publications

The First Amendment to the U.S. Constitution affords considerable protection to political, artistic, and scientific expression, that could trigger "strict scrutiny" by the Supreme Court (4). The court is most vigorous in reviewing government restraints from human-to-human, with a >50% case-fatality rate. The National Science Advisory Board for Biosecurity (NSABB), which advises the U.S. Department of Health and Human Services (HHS), recommended that two journals, Science and Nature, redact key information before publication. The NSABB and HHS expressed concerns that published details about the papers' methodology and results could become a blueprint for bioterrorism (1).

The U.S. government's request not to publish key scientific findings sparked considerable controversy. To many researchers, knowledge about what mutations enable respiratory transmission is essential to surveillance of and early action against variants of H5N1. They worry that government intrusion into scientific innovation would discourage vital research. However, security about building a hydrogen bomb, even though the information was in the public domain; the injunction was later vacated when the article was published elsewhere (2). In 2005, the Proceedings of the National Academy of Sciences refused to comply with an HHS request to decline publishing a mathematical model of botulism in the milk supply (3). The H5N1 case, however, is the first time government has sought to redact information after an institutionalized HHS review process.

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tion whether or not to comply (6). Given the absence of legal force or undue inducements or penalties, the government’s request to withhold information does not violate the First Amendment.

There are situations in which a government has the authority to block scientific communications. The clearest case is when research has been properly classified under federal law and the person seeking to communicate findings obtained it under the terms of a security clearance—whether they are still working for the government or not, so long as procedural requirements are met (7). Although a researcher is obliged to keep classified information confidential, publishers who obtain that information lawfully have a right to publish. In the Pentagon Papers case, the Supreme Court held that President Nixon did not overcome the “heavy presum-

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on research grants. The wider the scope of CUI conditions, the more likely that courts will invalidate them (4).

The Supreme Court’s “unconstitutional conditions” doctrine holds that government may not place conditions on public funding that require the recipient to surrender First Amendment rights. Thus, government has no obligation to provide research funding, but if it chooses to, it cannot restrain the free expression of researchers without a compelling state interest. For example, a federal appellate court recently struck down HHS guidelines requiring recipients of AIDS prevention funding to pledge their opposition to prostitution, reasoning that it was an unconstitutional condition (10).

The unconstitutional conditions doctrine, however, is hard to decipher. For example, the Supreme Court upheld HHS prohibitions on

Access to Sensitive Data Under the Freedom of Information Act

A functioning democracy requires that citizens be able to access information in the government’s possession, but not if access poses an unacceptable security risk. The Freedom of Information Act (FOIA) balances these concerns by affording access to federal agency records unless the records fall within a statutory exemption. Federal agencies support much of the research in the United States, including both of the recent H5N1 studies. Could the public obtain sensitive data that have been redacted from publications through a FOIA request? If so, governmental requests to redact sensitive information would be fruitless.

FOIA applies only to “agency records,” so a threshold issue is whether university research data acquired under a grant constitute an agency record. In 1980, the Supreme Court ruled that research data produced under an NIH grant and used in regulatory proceedings by the U.S. Food and Drug Administration did not constitute an agency record subject to FOIA because it was retained by the non-governmental grantee. The court found that FOIA required the agency to either produce or obtain permanent custody of the data (16).

The “ Shelby amendment,” enacted in 1999, expanded public access to data produced at universities and other nonprofit research entities under federal grants. The public can request the data if they were produced under a federal grant and “cited publicly and officially by the Federal Government in support of an agency action that has the force and effect of law” (17). Federal agencies could take care not to officially cite highly sensitive data, thereby avoiding a successful FOIA request. However, it is not always simple or easy to refrain from referencing sensitive research. The NIH, for example, might reasonably refer to the H5N1 research as justification for revising biosecurity policies.

Even if sensitive data do become part of an agency record, FOIA provides the federal government with ample authority to refuse a request on security grounds. FOIA provides nine exemptions under which records that would otherwise have to be disclosed
may be withheld, one of which is for “matters that are specifically authorized under criteria established by an executive order to be kept secret in the interest of national defense or foreign policy and are in fact properly classified pursuant to such an executive order” (18). Through this exception, Congress has acknowledged broad executive authority to classify records so long as it is done lawfully pursuant to an executive order.

President Obama’s 2009 Executive Order 13526 revises existing classification standards (19). Although it was designed to reduce the amount of classified materials, the executive order affords agencies considerable discretion to classify on security grounds. Consistent with prior policy, the executive order mandates that “basic scientific research information not clearly related to the national interest shall not be classified.” However, the order permits the classification of “scientific, technical, or economic matters relating to the national security,” provided that disclosure is reasonably expected “to cause identifiable or describable damage to the national security.” Furthermore, agencies may classify data that meet the executive order’s standards even if the data were not classified at the time of the FOIA request (19). Thus, federal agencies have wide authority to prevent the release of research information through a FOIA request simply by classifying it, provided that there are legitimate national security justifications.

In 2010, President Obama issued a further executive order stating that CUI is not automatically exempt from FOIA (20). Thus, to ensure that sensitive biological research information is not disclosed, agencies would have to classify it. [Certain nonbiological research, such as nuclear energy, is automatically exempt from FOIA, as are the locations where select biological agents are held (21).] Some research data also might be protected under FOIA exemptions for trade secrets or predecisional deliberative memoranda within the government, but these options are limited (22).

The law, then, draws a distinction between classified and controlled unclassified information. However, from a constitutional perspective, it would be troubling if the result turned solely on the label the government placed on the data. If the result did turn on the label, the government could simply relabel research from CUI to classified and thus prohibit its dissemination. Although decisions to classify can be challenged, prevailing is difficult, and unnecessary classification is common (23). This appears to place too much discretion in the hands of public officials.

The problem of government discretion is compounded by highly inconsistent practices among federal agencies in the classification systems they use. There is inconsistency of structure (the labels attached, such as classified, CUI, SBU, or other terminology), as well as in the application of that structure to individual documents (no clear standard exists for deciding whether to classify particular information). In short, the line between classified and CUI remains unclear, as agencies struggle to apply President Obama’s executive orders (24).

Balancing Scientific Freedom, Constitutional Values, and Biosecurity

The federal government has the power to prevent the dissemination of sensitive life sciences research, but there are good reasons to exercise that power sparingly. The current system of deliberation by a federal expert advisory board and HHS-issued voluntary recommendations is preferable to formal government mandates. Although we do not have all the data, the NSABB process in the H5N1 cases appears reasonable, given that unredacted publication could enable bad actors with scientific skill to replicate the studies, with profoundly harmful effects. The federal government has promised to share the researchers’ methods and conclusions with scientists with a need to know, which substantially advances scientific objectives.

Can the review process for high-risk biologic research be improved further? The NSABB’s origins can be traced to the so-called Fink report issued in 2004 by the National Research Council (21). However, vital aspects of the Fink report have not been implemented. In particular, the Fink report proposed an institutional review process for biological “experiments of concern”—those falling into seven research classes, making the pathogen considerably more attractive as a bioterrorism agent (e.g., by enhancing virulence or transmissibility or by rendering vaccines ineffective). This approach was patterned on the Institutional Biosafety Committees (IBCs) required by NIH for recombinant DNA research at institutions receiving federal funding, which generally have been considered to be successful (21).

HHS, in partnership with institutions, will have to ensure that the IBC model works effectively: (i) institutions must develop the requisite expertise to review dual-use research of concern; (ii) HHS must specify the categories of research requiring institutional review—minimally including the seven types of high-risk experiments; and (iii) HHS must set clear and consistent standards for institutional review. If IBCs are formally designated to conduct the institutional review function, HHS will have to clarify whether NSABB will guide and oversee the process (21). In addition, because IBCs may recommend that researchers voluntarily restrict access to methods or results in some instances, it will be important for HHS to develop a system for managing access to sensitive data and for disseminating it to those with a need to know in a fair manner.

If HHS improves its functioning, the institutional review process can ensure a sound balance between scientific freedom and national security. A fair, transparent process undertaken by research institutions, with a balanced approach to scientific benefits and public safety, together with HHS guidance and oversight of high-risk research, is preferable to government constraints on scientific information by force of law.

References and Notes
10. Alliance for Open Society International v. USAID (2nd Cir. 2011).