

PERSPECTIVE

Control use of data to protect privacy

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Massive data collection by businesses and governments calls into question traditional methods for protecting privacy, underpinned by two core principles: (i) notice, that there should be no data collection system whose existence is secret, and (ii) consent, that data collected for one purpose not be used for another without user permission. But notice, designated as a fundamental privacy principle in a different era, makes little sense in situations where collection consists of lots and lots of small amounts of information, whereas consent is no longer realistic, given the complexity and number of decisions that must be made. Thus, efforts to protect privacy by controlling use of data are gaining more attention. I discuss relevant technology, policy, and law, as well as some examples that can illuminate the way.

We live in an era of an explosion of data. For a variety of reasons, including massive collection by both the private sector and governments, as well as the ease of computing correlations—from which information can be derived even about people whose data are not in the set—the old methods for protecting privacy no longer work. An old protection made new, managing use, now seems the most appropriate way to secure privacy. Controlling use is complex, but combining technology, policy, and law is the best way to control incursions from businesses and governments.

The principles governing data protection are 40 years old. The Fair Information Practices (FIPs) were developed in response to the rise in the 1960s of computerized data systems. Coming originally from a report from the U.S. Department of Health, Education, and Welfare (1), the FIPs were revised by the Organization of Economic Cooperation and Development (OECD) (2). The more expansive OECD privacy principles have been the basis for many national and international privacy regulations.

Notice, consent, context

User control sits at the heart of the FIPs. Transparency and/or notice says that there should be no data collection system whose existence is secret; access, that there should be a way for the data subject to find out what information is in her record and how it is used; consent—sometimes called choice—that data collected for one purpose not be used for another without user permission; redress, that the data subject must have the ability to correct inaccuracies; and integrity and security, that the data collector keeps reliable records and protects them. In 1998, the U.S. Federal Trade Commission (FTC) identified these as the “five core principles of privacy protection” and noted that notice was fundamental, calling choice or consent the “second widely accepted core principle” (3).

Whereas the U.S. and Europe have taken different routes to protecting privacy—the U.S. using

sector-specific protections (financial data, banking information, health records), Europe pursuing broader data-protection schemes—both emphasized notice and consent. But, although the FIPs made sense when an individual could discern and react to a data-collection event, this is no longer true.

Consider data collection from a smart phone. The combination of information from the user

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and aggregated data from others can improve her experience. For companies, such data promotes faster, more-targeted services (and advertising), ties the consumer more strongly to the business, and boosts profits. For researchers, massive data illuminates connections that might not have been apparent and may uncover correlations that are actually causations.

Because data collection involves compilation of massive amounts of small bits of data, notice and consent are difficult for users to manage. Should collection of phone location data increase when a traffic accident blocks a popular route? What if the user is on a private assignment that day? That a service that provides up-to-date route information also collects up-to-date location data is not something all users realize (although they should). Frequent queries about permission for collection create a situation in which the user inattentively clicks “Yes”—not exactly a win for privacy.

Notice simply doesn't make much sense in a situation where collection consists of lots and lots of small amounts of information (Fig. 1). Written to cover all contingencies, privacy notices are not designed for human use. A 2008 study

showed that the average reader would need 244 hours simply to read the privacy policies for all websites she accessed in a year (4).

Consent is often not an option. Almost a decade ago, Fred Cate noted, “If consent is required as a condition for opening an account or obtaining a service, a high response rate can always be obtained” (5), whereas a 2014 President's Advisory Committee on Science and Technology (PCAST) report on big data and privacy observed, “Only in some fantasy world do users actually read these notices and understand their implications before clicking their consent.” (6).

Sometimes the user is not even given a choice about consent. Because of overwhelming complexity, Google, whose Android platform dominates the consumer smart phone market (7), decided to put permissions for information access into groups. Thus, a user lacks the ability to conduct fine-grained decisions on which information to permit apps to access (8). The user moves on, rarely examining—or withdrawing—consent afterward.

A fundamental problem is that seemingly innocuous data may trigger a privacy incident. Using the history of buying patterns of other customers, Target predicted a teenager's pregnancy from her vitamin purchases (9), and the ride-share firm Uber claimed to be able to discern one-night stands from the usage patterns of rider pick-up and drop-off data (10). Solon Barocas and Helen Nissenbaum noted, “The willingness of a few individuals to disclose information about themselves may implicate others who happen to share the more easily observable traits that correlate with the traits disclosed.” (11).

Context matters in privacy. That idea first espoused by Nissenbaum a decade ago (12) is gaining support in policy circles, including in the White House Consumer Bill of Rights (13) and a recent FTC report (14). Massive amounts of data create such personal and societal benefits that collection is unlikely to stop.

Controlling use

The FIPs protected privacy through notice and consent, but for reasons of complexity (too many tiny collections, too many repurposings), those are no longer effective. Nonetheless, notice and consent provide benefits: notice, for transparency, and consent, for certain types of data or use, as well as for controlling context (15). But the value of big data means we must directly control use rather than using notice and consent as proxies (6). That is true no matter who the collector is.

This is easier said than done. Big data provides the patterns that allow us to use resources efficiently. Determining how to continue to collect and use big data, but control its use, is complex. The tools are technology, policy, and law, and there are some examples that can illuminate the way.

Once the most solitary of activities, reading is losing the privacy between the reader and the page. Amazon and other purveyors of e-books have discovered multiple ways of tracking activity: where readers start, what they reread, whether they mark a passage, if they finish the text (16).

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There are other approaches that make tracking user reading more difficult, rather than less so.

One such is Shibboleth, software that enables a user at one participating institution, say, the University of Michigan, to access electronic resources at another, say, the University of Illinois. A user authenticates on the University of Michigan. The user, however, is identified to the University of Illinois not by personal identifier such as name or e-mail address but by her right to the resource. This could be because she is a member of the University of Michigan community (student or staff), a participant in a particular course, or one of a set of users authorized to access particular resources. Unless the information is specifically needed, the University of Illinois does not learn the user's actual online identity. The Family Educational Rights and Privacy Act, which protects the privacy of student educational records, and the fact that librarians view reader privacy as fundamental motivated this privacy-protective architecture.

A potentially powerful approach to controlling data usage is “accountable http,” a variant of the

http protocol. Proposed by MIT researchers Oshani Seneviratne and Lalana Kagal, [httpa creates a system to track information usage \(17\)](#). The system consists of a user who wishes to access data that have usage restrictions (e.g., no sharing, no sharing without informing the data owner, etc.); a data provider using an httpa server; and a Provenance Tracking Network (PTN). The PTN is a network of servers that log each data access and usage, either from the original data provider or any user downstream.

The magic behind the system is httpa, a protocol that conveys usage restrictions between the data providers and data users, creating a log in the PTN for each time a protected resource is accessed. These logs do not enforce compliance but can be used to determine it. This general approach to controlling data usage has only been tested in a small-scale effort; whether it can scale to the Internet is unclear. But it might be valuable in limited settings, such as patient health data, where a motivator might be the Health Insurance Affordability and Accountability Act (HIPAA), the U.S. law that restricts the sharing of patient medical data.

Online identities are used ubiquitously across the Internet to access restricted resources (e.g., pay-for-use subscriptions or library memberships confined to a university community), to post comments in restricted settings such as YouTube, and to conduct business at a bank or online broker. Although the need for secure, interoperable, and easy-to-use credentials for online identities was clear, development and adoption of such tools was slow.

The U.S. federal government stepped in, creating the National Strategy for Trusted Identities in Cyberspace (NSTIC) to provide funding for pilot programs and standards efforts that would provide both privacy and security. Using access to federal government sites as a lever, NSTIC requires that private-sector identity providers protect the privacy of information regarding user activities on federal sites (18).

Tracking when a user goes on a .gov website can reveal their private information, e.g., interest in HIV/AIDs or in penalties for unpaid taxes. Federal rules prevent identity providers from using tracking information from federal sites

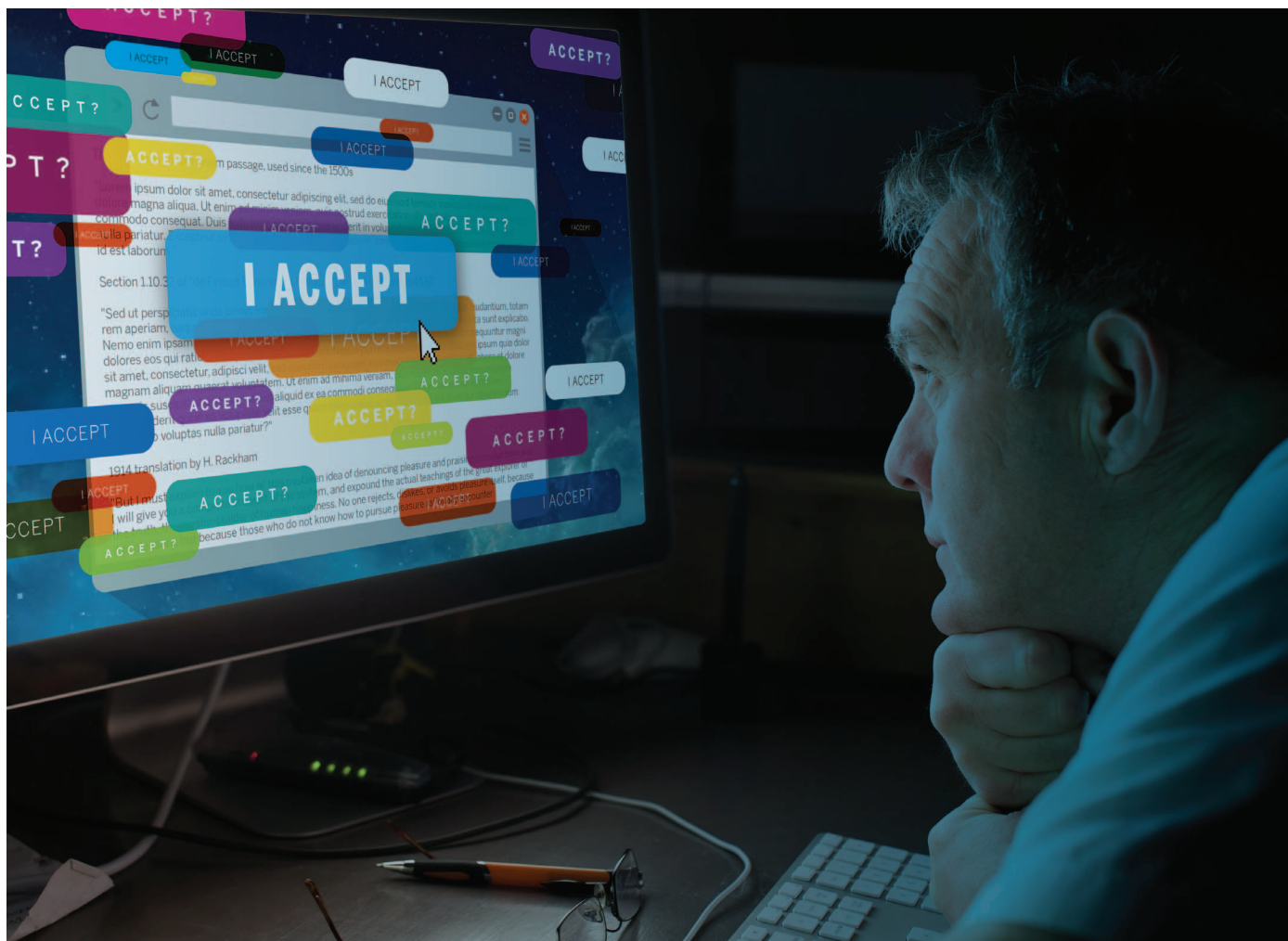


Fig. 1. Permission to run software has become complicated. In signing “I accept”—typically necessary to use an application—the user agrees to collection and use of information present on their device. Such data may not only be revelatory, it may also have been collected without the user’s knowledge or understanding of what can be discovered from this information.

for anything but authentication, audit, or complying with the law (19). In other words, no ads, no sharing the information with a third party, and no using the information to promote the identity provider's products. A signed-on user has greater privacy protections when visiting the National Cancer Institute website than when visiting the American Cancer Society site. Policy controls data usage and is then manifested in technical design.

Laws can provide shields against inappropriate data usage. The 1970 U.S. Fair Credit Reporting Act (FCRA), which predates the FIPs, does not control collection. Instead the FCRA strictly limits the circumstances under which a person's credit information can be accessed (essentially for credit, employment, and in response to court orders) (20).

A different example of control is the Genome Information Nondiscrimination Act (GINA) of 2008, which protects against discrimination in health insurance and employment based on genetic data. But GINA, too, has its limits. If a woman discovers through genetic testing that she is BRAC1- or BRAC2-positive, with a 55 to 65% or 45% chance, respectively, of developing breast cancer by age 70, GINA protects her ability to obtain health insurance and employability but says nothing about her ability to obtain disability, long-term care, or life insurance in the face of the BRAC1 and BRAC2 data.

There are other examples of how technology, policy, and law combine to control use. A well-known one is in medical research. The HIPAA privacy rule governs how researchers within health care organizations handle the health information of individuals; it also governs researchers who interact with such data (21). There are a number of ways this is done: through the law itself; its implementation in regulations (21); and Institutional Review Boards, which examine researchers' access and use of patient data, as well as by social controls. A researcher who is careless about the privacy of health records will find future access to such records very difficult.

Privacy and national security

An example that doesn't tend to appear when discussing privacy and big data is national-security collection. Yet the Snowden leaks disclosed massive collection both domestically and abroad. These disclosures started a national discussion on collection and use, although not, for obvious reasons, on notice and consent, which have little role in national-security collection.

I recently participated in a National Academies study on software alternatives to bulk signals intelligence collection (22). Bulk collection, specifically of telephony metadata—NSA receives daily downloads of telephony metadata (to, from, time, data, and length of call data) from major service providers—has generated much consternation. Metadata are data about the call, not its content, but mobile phones and the fact that cell phones are usually associated with a single individual mean that metadata themselves are remarkably revelatory (23, 24). Both a presiden-

tially appointed review group on intelligence and communications technologies and the Privacy and Civil Liberties Oversight Board, an executive-branch oversight board, recommended ending the government telephony metadata program (25, 26).

Our charge was somewhat different—technical alternatives to the collection—and our conclusion was also somewhat different. Because the program provides information that cannot be found in other ways, we believe there are no alternatives providing the same information (22). In particular, if past events become interesting in the present—a non-nuclear nation is discovered to be pursuing nuclear weapons or a new target is identified as a terrorist—past history may present new leads (22). Such past history will be available, in general, only if there were bulk collection in the past.

We made no judgment on whether the bulk collection program should continue; that is a policy decision, not a technical analysis, and out of scope for the study. We observed that the only way to protect privacy in the face of bulk collection is to control use—the same solution as the one to the private-sector big data collection issue.

We had no evidence that NSA was inappropriately using bulk data that were being collected. Nonetheless, we found that there were possible improvements for controlling use. We recommended increased utilization of automated controls and auditing, noting that manual controls and auditing will also always be necessary (22). Automating controls on data usage means data usage rules must be stated with great precision. That has its own advantages, including preventing inconsistencies (one such, on the meaning of archive, resulted in inappropriate access to the database) (22). Automated controls on data usage will also provide transparency. We also proposed research into privacy-protective methods of auditing by outsiders (22).

Controls on use

Our point was that "Controls on use ... offer an alternative to controls on collection as a way of protecting privacy." The same is true outside the national-security domain. Privacy intrusions are everywhere. The technologies—smart phones and their apps; the ubiquity of Google, which performs 68% of searches in the United States (27) and over 90% in Europe (28); and Internet-connected sensors in automobiles, bridges, cargo trucks, and so forth—are novel, but the fact that technologies and changing social mores create privacy intrusions is not. In 1890, a similar situation—yellow journalism and hand-held cameras—induced Samuel Warren and Louis Brandeis to write "The right to privacy," which laid a foundation for U.S. privacy protections. Warren and Brandeis observed that, "it has been found necessary from time to time to define anew the exact nature and extent of such protection" (29). Today is such a time. The nature and extent of redefinition will be of control on use, and determining the right controls, and the right ways to exercise them, will be challenging—but that is what we must do.

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