Innovation and Interdependence: The Case of Gene-Editing Technology*

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Abstract

Technological breakthroughs often reshape patterns of international exchange and interdependence, posing unique challenges for governments. We argue that innovation reduces policy autonomy among national governments in two ways. First, lower barriers to entry create opportunities for forum-shopping by researchers, firms, and other actors. This facilitates regulatory arbitrage as actors evade national rules by relocating to more permissive jurisdictions. Second, public unease about new technologies creates the potential for backlash against controversial applications. This backlash can spill across borders: accidents or misuse in one jurisdiction undermine support for research and commercial development elsewhere. Together, these processes can generate inefficient cycles of accelerated progress disrupted by damaging controversies. We test these mechanisms in the case of gene editing, finding support for the theory in data on scientific employment patterns and a survey experiment examining public backlash. Our results demonstrate that technological disruption increases interdependence and undermines states' ability to regulate in isolation.

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1 Introduction

Technological innovation is a defining feature of contemporary social and economic life. Recent advances in fields like robotics, biotechnology, digital finance, and artificial intelligence promise improved welfare through enhanced health, productivity, and economic growth. They also pose significant risks: many emerging technologies can be misused to cause harm or violate ethical norms. The recent revolution in gene-editing technology, for example, has been celebrated for facilitating new medical therapies and also criticized for enabling controversial modifications of human DNA.

Governments navigate this tradeoff by regulating the development and application of emerging technologies. Regulations seek to guide the path and speed of technological progress, balancing the economic and social potential of technological change against the risk of disruption and harm. Countries often make different choices in this environment — imposing more restrictive or permissive rules on the use of a particular technology — as governments align regulations with public preferences and social norms.

In addition to provoking a regulatory choice within countries, technological disruption can also shape patterns of interdependence between them. Many technological breakthroughs lower barriers to entry, erode incumbent advantages, and expand access to more actors in more jurisdictions. For example, improvements in rocketry and control system technology have allowed new governments and private companies to participate in space exploration. Advances in 3-D printing are similarly lowering costs and expanding participation in the development of medical devices and prosthetics. Innovation in blockchain technology has enabled novel financial transactions and new opportunities to mine and trade cryptocurrencies (Reinsberg) [2021). As technologies like these develop, countries' regulatory efforts become more closely linked via two mechanisms.

First, the reduction in entry barriers creates opportunities for forum shopping by firms

and individuals. As innovation reduces the material costs and expertise needed to participate in an industry, actors gain more freedom of movement. They may engage in regulatory arbitrage, exploiting differences in national regulations by relocating scientific and commercial development to more permissive jurisdictions. In some cases, governments will face pressure to weaken standards to lure researchers, firms, and capital from elsewhere. While regulatory arbitrage and competition are well-established features of national governance (Genschel & Plumper) [1997), we argue that technological shocks often exacerbate the problem by lowering costs and increasing cross-border mobility.

The second mechanism is rooted in public attitudes regarding emerging technologies. Because these technologies involve risks of harm or misuse, they generate apprehension among citizens and potential consumers (Zhang & Dafoe, 2019). When controversies occur, they may spur public backlash and undermine support for related research and commercial development. We argue that backlash frequently spills across national boundaries, such that controversies in one state affect public attitudes in another. As a result, one government's decision to weaken regulation can damage confidence in the technology around the world. Unlike forum shopping, we are not aware of existing scholarship that examines international spillovers in public backlash. Nonetheless, we argue that it is an important challenge in the governance of new technologies.

These mechanisms reshape the flow of people, information, and production across borders, complicating the role of national governments as they regulate emerging technologies. The most direct effect is an increase in interdependence among countries. Interdependence refers to situations in which "the ability of one participant to gain his ends is dependent to an important degree on the choices or decisions that the other participant will make" (Schelling, 1960a) p.5). In other words, it entails a reduction in a state's ability to realize its goals autonomously. Both mechanisms described above weaken the power of national governments to regulate technology in isolation. Increased opportunities for arbitrage make it easier for

targets of regulation to evade national rules. The potential for spillovers in public backlash mean that effective national governance cannot insulate a country from poor regulation in another jurisdiction. These dynamics increase the need for international policy coordination to manage interdependence.

In combination, the two mechanisms create incentives for states to mismanage risk, generating inefficient cycles of accelerated progress disrupted by damaging controversies. The freedom to forum shop diminishes governments' ability to constrain the use of new technologies; it may also encourage states to compete by weakening regulatory standards. This may temporarily speed technological progress, but also increases the systemic risk of controversial accidents or misuse. These controversies, in turn, can arouse public anxiety, undermine support, and stall continued progress.

We examine these processes in the case of gene editing, a field in the midst of a technological revolution. Gene editing refers to the targeted manipulation of an organism's genetic material. The emergence of CRISPR^T and associated techniques in the last decade provides a dramatically more accurate, efficient, and economical method for editing genes. In recognition of the technology's revolutionary capacity for "rewriting the code of life," CRISPR architects Emmanuelle Charpentier and Jennifer Doudna received the Nobel Prize in Chemistry in 2020 The development of CRISPR dramatically expanded the application of gene-editing technology while also triggering concerns about unethical or harmful misuse. We argue these conditions will increase the two forms of interdependence described above.

We probe these mechanisms with two sets of empirical tests. We first analyze a novel dataset on gene scientist employment to examine patterns of regulatory arbitrage. Specifically, we leverage the 2012 introduction of CRISPR as a temporal shock to examine how

¹The acronym CRISPR stands for clustered regularly interspaced short palindromic repeats. The term was coined by Ishino *et al.* (1987), who first noted the appearance of repeating DNA sequences in bacteria.

²Royal Swedish Academy of Sciences, "Press release: the Nobel Prize in Chemistry 2020," October 2020.

national regulation shapes the cross-border movement of gene-editing researchers. Our results are consistent with theoretical expectations: researchers are more likely to relocate to countries with weaker gene-editing regulations after 2012.

To test for spillovers in public backlash, we implement a survey experiment in which American respondents react to a hypothetical controversy involving the birth of genetically-altered infants. In addition to this basic treatment, we vary the country in which the inappropriate gene-editing activity occurred. We find that both foreign and domestic gene-editing controversies negatively affect domestic public support for gene-editing research.

Our paper adds to a growing literature on international competition, cooperation, and technological change (Canfil 2021) Drezner 2019; Jia et al. 2022 Milner & Solstad, 2020; Perlman, 2020b). We develop a theory of technology and interdependence in a domain, biotechnology, that has been largely neglected by scholarship in international relations and political science. While international relations scholars have paid close attention to the security implications of technological advancements (Ayoub & Payne, 2016) Buchanan & Keohane 2015), we know less about governance of scientific issues in non-security sectors. We expect biotechnology to increase in salience as governments and their citizens grapple with the unprecedented technological progress in this field.

More broadly, we identify two theoretical mechanisms — forum shopping and spillovers in public backlash — that link countries' fates as they govern emerging technologies. In doing so, we demonstrate how technological shocks interact with patterns of economic and political exchange to induce interdependence among countries (Keohane & Nye, 1977). These mechanisms have clear implications for the design of international institutions, which are likely to be charged with managing these spillovers (Koremenos et al., 2001).

The following section develops our theory of technological innovation and international interdependence. Section 3 provides background on the case of gene editing, summarizing

³For exceptions, see Oye & Wellhausen (2009) and Perlman (2020b).

the emergence and governance of this rapidly advancing technology and applies our theory to the technology's international spread. Section 4 describes our empirical strategy and presents our findings, and section 5 concludes.

2 Technological Innovation & Interdependence

A substantial body of research examines the political consequences of technological change. In international relations, scholars have long been interested in the effects of technology on interstate bargaining and conflict. Military technology is a key determinant of the offense-defense balance, affecting the severity of the security dilemma and the stability of cooperation between states (Jervis, 1978). Medical innovations such as vaccines and field medicine have drastically altered battlefield tactics and combat effectiveness (Fazal, 2014). More broadly, technological innovation and adoption shape the economic and military power of states. This provides strong incentives for governments to invest in technology, especially when they confront external threats (Taylor, 2012) and when the international system is more competitive (Milner & Solstad, 2020).

At the domestic level, technological developments shift the relationship between governments, opposition groups, and citizens. Surveillance technology, for example, has increased the coercive capacity of the state while also altering the balance of power between firms that supply these technologies and the states that use them (Soss & Weaver 2017; Xu, 2021). Access to mobile technology affects rebel and government tactics in civil wars (Bailard, 2015; Pierskalla & Hollenbach, 2013; Shapiro & Weidmann, 2015). Social media changes how citizens and advocacy groups organize for political change and alters government control tactics (Gilardi et al., 2022; King et al., 2017).

Technological innovation has also allowed citizens, firms, and political groups to forge new transnational links. Improvements in transportation, financial, and digital technologies lowered the costs of transnational exchange, spurred waves of globalization, and deepened economic interdependence (O'Rourke & Williamson, 2001) Keohane & Nye Jr. [1973]. Modern communications technology facilitates cooperation among advocacy groups (Hall et al., 2020) Keck & Sikkink [1999] as well as transnational extremist movements (Gohdes, 2018; Mitts, 2021).

As technology induces new patterns of exchange among substate actors, it often creates challenges for governments as they navigate domestic and international politics. Communication networks, financial flows, and complex supply chains generate new opportunities for coercion and cooperation among states (Keohane & Nye Jr, 1973; Farrell & Newman 2019). Shifts in production and employment generate domestic winners and losers and shape the political preferences of citizens (Drezner, 2019). As governments respond to these events, they must simultaneously grapple with the increasing cross-border mobility of information and economic assets that generally accompanies technological innovation. This enhanced mobility and the associated threat of exit can constrain governments' ability to craft domestic policy (Genschel & Schwarz, 2011) [Mansfield & Rudra, 2021).

2.1 Technology, Arbitrage and Backlash

We build on this growing body of work examining how technology and the transnational flows it facilitates undermine the policy autonomy of states. Specifically, we focus on two general mechanisms that arise in the wake of rapid technological innovation: enhanced opportunities for regulatory arbitrage and backlash from public controversies. Both mechanisms constrain the ability of states to govern in isolation and increase the need for international policy coordination.

We develop the logic of each mechanism below and then describe how they interact. While these effects of technological change are far from exhaustive, we argue they are important sources of international interdependence that shape the domestic and international political response to disruptive innovation.

2.1.1 Regulatory Arbitrage

Technological innovation creates new economic opportunities by lowering production costs and allowing new markets to emerge (Sheal 1998). After a new technology is introduced, researchers, entrepreneurs, and firms race to develop applications to exploit these opportunities. The stakes of this competition are substantial. The global market for artificial intelligence, for example, was approximately \$60 billion in 2021 and will exceed \$400 billion by 2028 The quantum computing and advanced robotics industries are similarly expected to grow drastically over the next decade. Successful pioneer firms capitalize on this growth, often securing a first-mover advantage that endures even as competitors subsequently enter the market (Lieberman & Montgomery, 1988 Agarwal & Gort, 2001).

As actors jockey for advantage, regulatory constraints can limit their ability to develop and commercialize new technologies. One strategy for overcoming these hurdles is to seek out a more permissive legal environment. In other words, firms or researchers may circumvent regulatory constraints by relocating to jurisdictions with more favorable rules.

Regulatory arbitrage is a strategic response to inconsistencies in national regulation. Novel technologies may increase the potential returns of arbitrage by increasing profit opportunities. More importantly, technological innovation can increase actors' ability to relocate research, development, and production across regulatory jurisdictions. Innovation reduces the capital and expertise needed to participate in an industry. These reduced costs increase freedom of movement, increasing opportunities for forum shopping.

Accelerated regulatory arbitrage creates several problems for governments. First, it

⁴ "\$422.37+ Billion Global Artificial Intelligence (AI) Market Size Likely to Grow at 39.4% CAGR During 2022-2028," *Bloomberg*, 27 June 2022.

⁵ "Quantum Computing Market to Grow Exponentially; Increasing Product Applications to Generate Remunerative Market Opportunities: Fortune Business Insights," *Global News Wire*, 4 March 2022.

reduces a government's ability to regulate the application of novel technologies. When cross-border mobility is limited, actors are largely forced to accept the regulations that governments impose. Technological shocks increase the exit options available to firms and researchers. These options allow them to evade unfavorable rules and weaken the hand of national governments.

A second problem is the potential for regulatory competition among governments. The economic returns associated with new technologies creates pressures to lower regulatory barriers. Lax regulations can lure capital and production from more stringent regulations, giving governments an incentive to undercut each other's rules. This can generate a deregulatory spiral in which governments collectively weaken standards.

In international relations scholarship, regulatory arbitrage and competition are most closely associated with issues like tax policy, financial regulation, and environmental standards (Trachtman [1993]; Angelini & Cetorelli, [2003]; Konisky, [2007]; Genschel & Schwarz, [2011]). These are policy domains in which nations differ in their regulatory approach, economic output is responsive to rules, and assets have high cross-border mobility. We argue that these conditions increasingly characterize fields undergoing rapid technological change. Technological advancement often lowers entry barriers and encourages diffusion to a wider set of regulatory jurisdictions. It frequently occurs in industries with enormous growth potential and where initial economic advantages can yield substantial returns. As a result, we should expect emerging technologies to exhibit higher levels of regulatory arbitrage and the potential for more intense regulatory competition. [6]

An implication of this argument is that technological applications will advance more rapidly and with more limited oversight than a counterfactual world with fewer arbitrage

⁶Notably, other processes can sometimes counteract competitive pressures to generate a "race to the top" (Genschel & Plumper) [1997] Prakash & Potoski, [2006]). In particular, if jurisdictions with sufficient market size adopt stringent rules, these can encourage higher standards elsewhere. Our argument is not that competitive pressures will dominate these countervailing forces, but that the pressure to engage in regulatory competition increases in the face of rapid technological advancement.

opportunities. If innovation enables actors to relocate research and development activity to more permissive jurisdictions — a hypotheses we test in Section 4— we should expect an increase in the systemic risk of accidents or misuse. These events can spark public controversies that spill across borders, which we examine below.

2.1.2 Spillovers in Public Controversies

The second governance challenge arising from technological change stems from public attitudes about emerging technologies. Public opinion affects the trajectory of the technological development in at least three ways. First, beliefs about the safety and morality of new technologies shape consumer demand for associated products. Second, public opinion affects the ability of firms to attract investors. Finally, public attitudes influence regulation, which determines the permissible ends to which the technology may be applied.

We argue that public attitudes about new technologies are often fragile. Rapid technological advances challenge existing systems of practice and thought. The ramifications of disruptive technologies frequently do not nest neatly into existing ideological or political cleavages; instead, they create unexpected coalitions and give rise to a mix of emerging public narratives. As a result, we view emerging technologies as particularly vulnerable to backlash when controversies arise. If the technology intersects traditional political divides, elites may lack incentives to provide a narrative for individuals to anchor their own beliefs (Druckman et al.), 2013). With no pre-existing reference frame to anchor individuals' views and moderate extreme reactions, high profile events can create quick and profound shifts in public opinion. Controversies spark public backlash, lead to reductions in public funding, and engender knee-jerk regulatory responses that constrain even responsible scientific activity.

The recent history of gene therapy provides an example of such backlash. In 1999, 18-year old Jesse Gelsinger joined a clinical trial at the University of Pennsylvania for a developmental

gene therapy treatment. Unlike the other trial participants, Gelsinger suffered an unexpected immune response that ultimately lead to his death. The tragic loss led to an immediate and precipitous drop in public support and consumer demand for gene therapies. As Jennifer Doudna recalls, the incident "made the whole field of gene therapy go away, mostly, for at least a decade. Even the term gene therapy became kind of a black label" (Rinde, 2019).

This example illustrates how an uncertain environment with few consistent cues engenders instability in public attitudes. As a result, emerging technologies that rely on public support often progress in fits and starts, with periods of promising technological advancements interrupted by crises of public confidence. There is evidence for this dynamic in the related field of genetically-modified organisms, where media exposure to controversies has been found to meaningfully affect public opinion (Prakash & Kollman, 2003) Drezner, 2008; Vigani et al., 2012). Ciocca et al. (2021) similarly note the potential for "hype-induced backsliding" in the field of artificial intelligence.

Notably, public backlash is not limited to the jurisdiction in which a controversy occurs. Citizens and elites may respond to accidents or misuse abroad by reducing their support for the technology at home. We argue backlash is usefully conceptualized as a negative spillover that spans national borders. As discussed above, the choice to reduce regulatory barriers can bring economic rewards and enhance the prestige of a country's scientific establishment. There are also costs associated with lax regulation, however, including the potential for domestic public backlash. Because backlash spreads across jurisdictions, these costs are not fully internalized by the home country. Controversies may damage support for new technologies even in jurisdictions that are comparatively well-regulated. While regulatory arbitrage has been documented in other contexts, we are not aware of existing scholarship that examines the potential for spillovers in public backlash. Nonetheless, we expect that it is an important source of interdependence that occurs whenever a technology is associated with safety risks or ethical concerns.

Two scope conditions are necessary for innovation to increase the sources of interdependence described above. First, the new technology must lower barriers to entry or otherwise expand access in the industries to which it is applied. Many technological breakthroughs do this by lowering the material costs of an activity or production process; others may reduce the human capital required to participate. Innovation in computing systems, telecommunications, and information technology facilitates broader participation in a range of economic pursuits. This scope condition is necessary to trigger increased arbitrage behavior, which requires the ability to apply and develop the technology in multiple jurisdictions. It also suggests technologies that concentrate rather than expand participation will not be plagued by arbitrage concerns. For example, nuclear weapons technology has diffused to relatively few countries and only to government actors, largely because states intentionally constructed high barriers to entry in this field.

Second, a technology must be susceptible to misuse, unethical applications, or harmful accidents in order to arouse public anxiety. We argue that this is a relatively common trait of emerging technologies – from nuclear energy in the 1950s to artificial intelligence today – and expect the potential for public backlash to pose a recurring challenge for technology governance.

Both regulatory arbitrage and backlash spillovers increase the need for international policy coordination. Each process represents a form of interdependence that undermines the efficiency of decentralized, national-level governance. This is precisely the class of governance problems appropriate for international coordination, which is designed to manage spillovers and resolve market failures among states (Keohane & Nyel 1977; Keohane 1984) Farrell & Newman 2015; Koremenos 2016). The most straightforward way that international coordination can mitigate these spillovers is by harmonizing national rules via international standards. Even in the presence of enforcement problems, international rules can serve as a focal point to draw national regulations toward a single regulatory standard (Schelling).

1960b; Perlman, 2020a). International institutions may also boost public confidence by monitoring for potential violations and certifying the responsible conduct of research. We discuss both mechanisms of global governance in Section 4.3. We now turn to the case of gene-editing technology, the subject of our empirical tests.

3 Gene Editing: Technological and Political Landscape

While the ability to modify genetic material is not new, scientific advances have transformed the field over the past decade. The goal of gene editing is typically to suppress or alter naturally-occurring biological traits of an organism. Historically, the field evolved from splicing together naturally-occurring genetic material (producing "recombinant" DNA) in the 1970s to using cells' own DNA-repair technology to selectively edit specific genes (using "programmable nucleases") in the early 2000s (Gupta et al., 2014).

The emergence of the CRISPR method in 2012 represents a particularly significant breakthrough in gene-editing technology. The name CRISPR — an acronym for "clustered regularly interspaced short palindromic repeats" — refers to a series of repeating DNA sequences originally found in bacteria. These sequences provide bacteria with adaptive immunity by allowing them to recognize and destroy the DNA of harmful viruses. Scientists adapted this technique for programmable gene editing (Jinek et al., 2012). The CRISPR system is significantly more accurate, efficient, and economical than previous methods for editing.

In the years since its development, CRISPR has become the dominant gene-editing technology (Carroll, 2018) 7 A report in Stanford Medicine notes that while "other gene-editing tools have emerged in recent years... none seems to match the precision, low cost and usability of CRISPR" (Shwartz 2019). Like other breakthrough technologies, CRISPR dramatically reduces the costs associated with editing genes: by 2019, a gene-editing template that cost

⁷Figure A1 in the appendix shows the frequency of CRISPR patent applications compared to rival methods like TALENs and ZFNs.

\$1000 to design using rival technologies could be produced with CRISPR for \$65 (Shwartz, 2019).

Lower costs have expanded the use of gene-editing technology to laboratories around the world. Diffusion is also facilitated by conditions imposed by scientific journals, which require authors to make their data and materials available to other researchers. Much of the biological material – including the plasmids used to edit genes – is handled by third-party distributors. Appendix Figure A2 displays the number of researchers registered with a popular genetic material repository by country of origin. American researchers are the largest group, followed by China, France, Japan, India, and Germany An employee of this repository estimated that 25% of requests are for transfers of CRISPR-related materials.

Many have cheered the spread of gene-editing technology, which has stimulated a "biotechnological revolution" in basic research, clinical care, agriculture, and other fields (Knott & Doudna, 2018). Researchers routinely "knock out" genes in mice or other animals to study gene function and expression. New gene therapies are being developed to treat cancer and correct harmful genetic mutations (Khan et al., 2016). Agricultural producers are applying CRISPR to both plants and livestock. Research teams have successfully altered the DNA of mosquitos to prevent the transmission of malaria (Gantz et al., 2015). More recently, geneediting technology has been used to develop diagnostic tests and treatments for COVID-19 (Straiton, 2020).

3.1 Governance of Gene-Editing Technology

Gene editing is governed by a fragmented patchwork of norms, national laws, and international guidelines. When targeted gene editing first became feasible in the 1970s, scientists

⁸US researchers are overwhelmingly the most frequent depositors of CRISPR plasmids (see Appendix Figure A3).

⁹Interview by authors, 11.25.2019.

attempted to construct self-governing arrangements for gene-editing research. In 1973, leading geneticists announced a voluntary moratorium on gene-editing experiments involving certain viruses and toxins (Berg et al.) 1974). The moratorium was maintained for two years until it was replaced by formal guidelines adopted by the National Institutes of Health. Scientists involved in drafting the original guidelines argue that this decentralized approach was successful in constraining potentially inappropriate applications (Berg & Mertz, 2010).

In recent years, similar efforts have sought to establish new norms for the research community. A 2019 conference of geneticists called for a global five-year ban on editing DNA in human eggs, sperm, or embyros that are brought to term (Lander et al., 2019). However, there is dissent about this approach even among the most prominent gene researchers (Cohen 2019). The lack of consensus creates uncertainty about appropriate applications of gene-editing technology, potentially contributing to misuse. In addition, it is unclear whether voluntary, decentralized rules can succeed in an era when gene-editing technology is more accessible and diffusely distributed than in the 1970s.

As gene-editing technology progressed, national regulations began to supplement scientific norms. Early U.S. guidelines built upon the partial gene-editing moratorium of 1973-4 (Baskin et al., 2016). Other states followed suit as the technology became more widespread. Currently, there is significant variation in the structure and rigor of national rules. Some countries, for example, maintain a legal ban on the alteration of human germline cells. Some have less formal "guidelines" prohibiting germline editing, while others are more permissive in the constraints they place on the technology (Araki & Ishii) 2014; Ishii, 2017). Figure 1 displays a composite measure of national gene-editing regulations combining information from three recent surveys of regulatory policies. Countries are shaded according to

¹⁰According to Ishii (2017), this group includes Canada, Brazil, Australia, and much of Western Europe.

¹¹Source data are from Araki & Ishii (2014), Isasi et al. (2016), and Baylis et al. (2020). For details on these measures and the construction of the composite measure, see Section 4.1.

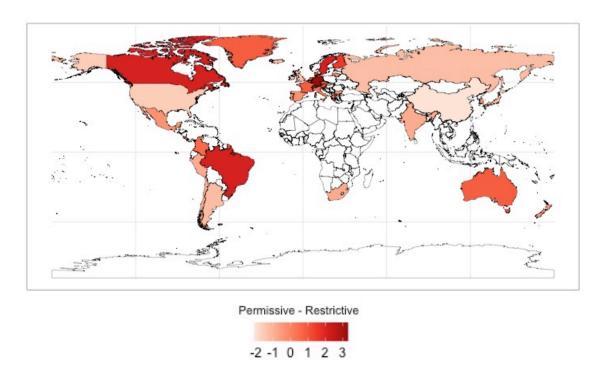


Figure 1: National Regulation of Gene-editing Technology. Thirty-nine countries are rated by the permissiveness of national gene-editing technology regulations. Ratings combine data from Isasi et al. (2016), Araki & Ishii (2014), and Baylis et al. (2020). See Section 4.1 for details on the coding and source data.

regulatory rigor, with darker shades indicating more restrictive national rules. [12]

Inconsistent rules across countries stem, in part, from different historical experiences and cultural expectations regarding the appropriate use of gene-editing technology. For example, Germany's experience with unethical experiments during the Nazi regime has conditioned the state's regulation of human subjects research. South Korea developed relatively strict biological research guidelines in response to a high-profile controversy regarding the falsification of data in a cloning experiment (Resnik et al.) [2006). The United States maintains comparatively weaker regulations for gene editing, consistent with a policy process that is more receptive to industry influence. Permissive regulations in China are driven in part by pressure to outpace Western countries in technological innovation as well as resistance to

¹²Countries with no identifiable gene-editing regulations are not colored.

 $^{^{13}}$ One biotechnology expert referred to US regulation of gene-editing technology as "the Wild West" (Interview by authors, 9/21/2020).

international biotechnology standards (Kleiderman & Ogbogu, 2019). In many cases, however, outdated national regulations have simply not kept up with rapid advances in the field (Baylis, 2019).

At the global level, there is growing interest in international coordination. Notably, the prospects for multilateral cooperation have not been not plagued by the political cleavages common to other issue areas (e.g., geopolitical rivalries or North-South divisions). A set of legacy international agreements, negotiated in the 1990s in reaction to concerns about cloning, provide a precedent for global governance of genetic research. In recent years, however, formal international institutions have been slow to develop rules despite calls for new global standards. The World Health Organization is among the few international organizations explicitly addressing the issue, releasing a set of non-binding recommendations in 2021 for appropriate oversight of human genome editing (WHO) [2021].

3.2 Arbitrage and Backlash in Gene-Editing Technology

We argue that CRISPR is an ideal-typical case of a disruptive technology that reduces the ability of states to independently regulate gene-editing research and applications due to its low cost and high public salience. While there are longstanding concerns about regulatory arbitrage in genetic research. The recent innovations have increased both the demand and ability to circumvent regulatory constraints.

Gene-editing technologies are inputs to an array of commercial applications that are expected to grow substantially over the next decade. In 2019, the gene-editing market was

¹⁴The 1997 Oviedo Convention prohibits human cloning, genetic screening for non-health purposes, and the misuse of innovations in biomedicine and bans. The 1997 Universal Declaration on the Human Genome and Human Rights and subsequent UNESCO declarations address genetic data and trade in genetic resources.

¹⁵In 2016, environmental activists unsuccessfully pushed for the UN Convention on Biological Diversity to expand its mandate to regulate synthetic biology and gene drive organisms.

¹⁶The potential for regulatory arbitrage was raised in the 1970s as several clinical trials moved to Europe and South America to sidestep burdensome rules in the United States (Baskin *et al.*) 2016).

worth approximately \$3.8 billion and is projected to exceed \$10 billion in the next five years (Ugalmugle & Swain) [2020]. The most direct applications are in the healthcare industry, where firms are developing gene therapies to address a range of disorders and chronic illnesses. Among these are CRISPR Therapeutics, co-founded by Nobel laureate Emmanuelle Charpentier to develop gene-based medicines, which went public in 2016 and has since increased more than fivefold in market value. Other sectors like agriculture, veterinary medicine, and industrial production processes also increasingly draw on gene-editing technology (Brinegar et al.) [2017). Given the array of market applications, the competition for these economic returns is fierce. Patent applications associated with gene-editing technologies grew from less than 1,500 in 2000 to over 12,000 in 2019.

Given these economic benefits, firms have lobbied governments to lower regulations on the technology. European plant breeders have pressed the EU to relax gene-editing restrictions, arguing that existing rules put them at a competitive disadvantage [18] South Korea recently instituted a review of rules on gene therapy research in order to maintain its competitiveness in medical technology (Ji-young, 2017). In the United States, the government bowed to agricultural producers' demands to weaken restrictions on gene-edited crops and livestock (Cancryn & Crampton, 2021). Other countries have announced similar regulatory reviews or new public initiatives to capitalize on gene-editing technologies. [19]

When regulatory barriers remain, actors have moved research and development to more permissive rules. Baylis (2019) describes several incidents in which US-based scientists relocated embryonic genetic research on embryonic DNA to clinics in Mexico and Ukraine "so as

¹⁷Data are from lens.org and reflect searches for "gene editing."

¹⁸Max Planck Institute, "Regulating genome edited organisms as GMOs has negative consequences for agriculture, society and economy", https://www.mpg.de/13748566/position-paper-crispr.pdf]

¹⁹Policymakers in New Zealand are reviewing the country's gene-editing regulations (Morton, 2019), and the Russian government recently announced a collaboration with Rosneft to develop gene-editing technology (Morton, 2020).

to not violate US federal law" (46). Isaacson (2021) similarly reports that entrepreneurs seeking to develop gene-editing applications in reproductive care openly acknowledge the ability to evade strict regulations.

As with other technologies, however, rapid progress has been accompanied by public anxiety and fears of misuse. Actors in both academia and industry are keenly aware that continued research depends on managing public anxiety about gene editing. Participants at a 2015 conference on gene editing, for example, called for slowing down the more controversial germline gene-editing research "in order to create a safe political space" (Isaacson, 2021, 288). Historically, controversies regarding one application of gene-editing technology have diminished investor interest more broadly (Gardner, 2020), as reflected in the Jesse Gelsinger tragedy. Advocates for scientific and national regulation of gene editing frequently cite "increasing legitimacy and trust" as a primary goal (Kuzma et al., 2018, 23). Aiyegbusi et al. (2020), for example, identify public perceptions of gene therapies as "central to their uptake and use."

Concern about inappropriate genetic modification escalated in 2018, when the Chinese scientist He Jiankui announced the birth of the world's first gene-edited infants. He used CRISPR to genetically alter several embryos in order to render them immune to HIV (Cyranoski) 2019). Unlike the Gelsinger tragedy, however, it unleashed a response that spilled across national borders. The revelation sparked international outcry, raising concerns about safety, consent of the participants, and the risks of modifying traits that will pass to subsequent generations. Calls for a global moratorium on some avenues of gene-editing research swiftly followed the revelation of He's experiment (Lander et al.), [2019). Recognizing the po-

²⁰The Ukraine clinic now boasts patients from Ukraine, the US, Israel, and Spain in further evidence of forum-shopping for gene-editing products (Baylis) 2019 47).

²¹A Silicon Valley entrepreneur responded to questions about FDA regulations of CRISPR for gene-editing babies by saying that "clinics did not have to be in the US. There would likely be other countries where the procedure would be allowed, and people who could afford gene-edited babies would be willing to travel" [Isaacson] [2021] 286).

tential for public backlash, leading scientists were quick to condemn the research. A senior colleague accused He of "jeopardizing the entire field of genetic engineering" (Isaacson, 2021, 306). Chinese scientists working in the field of gene editing expressed concern after the sentencing "that the international condemnation that followed He's explosive announcement in 2018 might have a wider chilling effect on CRISPR work in China" (Cyranoski, 2020).

The economic incentives and public anxiety associated with the gene-editing revolution align with our theory of technological disruption and interdependence. In the following section, we look for evidence of regulatory arbitrage and public backlash spillovers in this field.

4 Empirical Tests

We present two empirical tests of the theory outlined above. First, we leverage data on scientific employment to examine patterns of regulatory arbitrage. Because we expect the CRISPR revolution to exacerbate forum shopping, we use the year in which CRISPR was introduced as a cutpoint in the analysis. We test whether gene-editing researchers are systematically more likely to move to countries with weaker regulations after 2012. We also examine whether countries with permissive regulations benefit from increased patent applications and clinical trial development in this period.

For the second test, we identify the presence of public backlash using an original online survey experiment on American respondents. We randomly assign information about a hypothetical gene-editing controversy and examine its effect on public support for gene-editing research and policy. The experiment varies whether the controversy occurs domestically or in a foreign country, allowing us to test whether foreign misuse of gene-editing technology affects public attitudes towards gene-editing use and policy in the United States.

²²After initially heralding the achievement, China sentenced He and two colleagues to three years in prison for "illegal medical practice" (Cyranoski 2020).

4.1 Regulatory Arbitrage

We argued above that actors can evade strict regulations by relocating to jurisdictions with weaker rules, and that this behavior should increase in the wake of significant technological breakthroughs. To test this claim, we analyze employment patterns of over 100,000 gene researchers. We also look for evidence of forum shopping in the commercial development of gene-editing technology using data on clinical trials and patent applications.

Our independent variable for these tests is the rigor of national regulations governing gene-editing technology. We develop a composite national regulatory score drawn from three sources. First, Isasi et al. (2016) classify national regulations on a range of gene editing-related issues, including gene therapy, human germline editing, and genetic diagnosis. Countries are rated as "permissive," "intermediate," or "restrictive" on each issue; we transform these into a 1-3 scale of increasing regulatory rigor and average across the fields to generate a single national regulatory score. Second, Araki & Ishii (2014) provide a separate classification of countries based on the regulation of heritable genetic editing [23] Finally, Baylis et al. (2020) examine national rules regarding the use of genetically modified in vitro embryos in laboratory research.

These three measures are positively correlated but prioritize different applications of gene-editing technology. We combine them into a broad measure of each country's regulatory environment via principal components analysis. This provides a continuous, cross-national measure of gene-editing regulation for 39 countries that engage in gene research and clinical development. Cross-national variation in these regulations is visualized in Figure 1 The

 $^{^{23}}$ The categories include "ban based on legislation," "ban based on guidelines," "restrictive," and "ambiguous," which we transform into a 1-4 scale.

²⁴Baylis et al. (2020) categorize countries' regulatory approach as prohibitive, prohibitive with exceptions, indeterminate, or permissive based on a review of national legislation, guidelines, and codes of conduct which we transform into a 1-4 scale.

²⁵The three data sources vary widely in geographic coverage. Thirty-nine countries are classified by at least two sources. For these countries, we impute the missing scores before estimating the principal components.

most restrictive regulatory environments include Germany, Sweden, Switzerland, and Brazil. The most permissive are China, Ireland, and the United States. The regulatory scores are centered at zero and range from -2.1 to 3.3.

We first examine whether gene scientists relocate to countries with more permissive regulatory standards at higher rates following the introduction of CRISPR in 2012. If our theory is correct, institutions located in countries with more permissive regulatory standards will be comparatively more attractive destinations for researchers in the post-CRISPR era. To test this claim, we examine employment patterns of researchers who have published scientific papers in the field of gene editing. We collect these data from *PubMed*, a large database of biomedical publications. To obtain a sample of gene researchers, we extract all published articles on the topic of "genetic engineering" from 2002-2021 [26] We match these articles with a separate database, *Web of Science*, to identify the institutional affiliations of the authors in the *PubMed* sample [27] The search yields approximately 120,000 papers and over 100,000 unique gene researchers.

Using this record of scholarly publications, we construct a dataset of researcher movement at the level of the directed country-dyad-year. An observation reflects the number of gene researchers who relocate from country i to country j in year t^{28} In the year 2005, for example, twelve scientists who were most recently employed in Japan published papers while employed in the United Kingdom. Another ten moved in the opposite direction, relocating from the United Kingdom to Japan. We also include observations representing scientists who remain

 $^{^{26}}$ We begin in 2002 because this is the first year PubMed records full author names.

²⁷We exclude two sets of researchers: 1) those without a listed institutional affiliation, and 2) those with very common names (appear 100 times or more in our sample). We classify each researcher's country of employment using the geographic information included in their institutional affiliation.

²⁸A constraint of the source data is that we only observe a scientists' country of employment in the years that they publish. Because the ability of scientists in a given country to publish research may be related to the state of gene-editing technology, we restrict the sample to researchers who published at least one paper before the introduction of CRISPR in 2012. The data therefore reflect employment relocations among gene scientists who were active researchers before the technological shock.

in their "home country" (e.g., 737 researchers who were most recently employed in the UK remained there and published papers in 2005). These counts of gene scientist relocations serve as the dependent variable in the tests below.

Of the 262,377 employment records we observe in our sample, 27% represent relocation across international borders while 73% remain in their country of prior employment. International relocations occur for several reasons. Scientists move abroad in search of institutional prestige, more generous funding, familial ties, or other reasons. We argue, however, that the regulatory environment of each country shapes decisions on the margin, and that the effect of regulatory differences will be larger in the CRISPR era.

To test these patterns systematically, we construct a variable (Regulatory Difference) that subtracts the origin country regulatory score from the researcher's country of current employment. Positive values mean that the destination country has stricter regulations than the origin country. In the statistical models below, we interact this variable with an indicator for the time period (2012-present) when we theorize that CRISPR technology enhanced forum-shopping opportunities. These models exploit the technological shock of CRISPR to estimate how employment patterns respond to regulation in the wake of technological breakthroughs.

Control variables reflect each country's national economic output, human capital, and commitment to research funding. We include GDP to account for each country's economic capacity and GDP per capita for its level of development. We proxy scientific capital with a count of the number of patent applications in each country (Patents). Finally, we include a measure of R&D expenditure for each country. Data for all control variables are drawn from the World Bank's World Development Indicators (WDI). Separately, we add an indicator for same-country pairings to account for the high propensity of researchers to remain employed in the same country over time. We further include dyad fixed effects in some specifications to control for features specific to each country pair.

	DV: Scientific Relocation		
	(1)	(2)	(3)
Regulatory Difference	-0.010 (0.010)	$0.038 \\ (0.028)$	$0.038 \\ (0.028)$
CRISPR	9.355*** (3.409)	9.423** (4.370)	1.971** (2.597)
Regulatory Difference \times CRISPR	-0.025^* (0.014)	-0.034^{**} (0.017)	-0.034^{**} (0.017)
Controls		✓	✓
Dyad FE			✓
Observations	22,730	22,730	22,730

Table 1: Employment Relocation of Gene Researchers: Linear model estimates for the volume of gene-editing researchers who relocate to institutions in another country. Column 2 includes the following controls (not shown): GDP origin country, GDP destination country, GDP per capita origin country, GDP per capita destination country, Patent Applications origin country, Patent Applications destination country, R&D origin country, and R&D destination country. Standard errors are clustered by country dyad. Statistical significance is denoted by: *p<0.1; **p<0.05; ***p<0.01.

We report the results of linear models in Table 1 Standard errors are clustered by country dyad. Column 1 presents a baseline model with the Regulatory Difference measure, an indicator for 2012-present (CRISPR), and the interaction term. The positive and statistically significant coefficient for (CRISPR) indicates a higher propensity for employment relocation after 2012. This aligns with our theory, which suggests technological shocks generally increase cross-border mobility. More importantly, the interaction term Regulatory Difference × CRISPR is negative and significant, suggesting that permissive jurisdictions become substantially more attractive destinations starting in 2012. This result is consistent with enhanced opportunities for arbitrage in the CRISPR era. Columns 2 and 3 find broadly similar results after adding the full set of covariates as well as dyad fixed effects. Substantively, a one-unit shift in Regulatory Difference reduces scientific relocation by approximately

0.03 researchers per year. This is a modest effect, but can be meaningful once aggregated over the time period. For example, the results suggest that a high-regulation country like Germany has lost approximately 31 gene-editing researchers to regulatory arbitrage from 2012-2019.

One possible challenge to inference is that permissive jurisdictions (e.g., China, Ireland, United States, and Russia) may be generally more attractive destinations for scientists for reasons other than their regulatory environment. To address this, we conduct a placebo test that examines whether scientists in unrelated fields exhibit relocation patterns that correlate with national gene-editing regulations. The results, reported in Appendix Table A1, reveal no significant change in relocation patterns after 2012.

To gauge whether incentives for arbitrage after 2012 extend beyond employment relocation to other outcomes, we analyze patterns of scientific coauthorships, gene therapy clinical trials, and patent application in Table 2. These tests are structured at the country-year level and include the same control variables listed above. In Column 1, we examine whether gene-editing researchers in higher-regulation countries are incentivized to seek more international coauthors as an alternative means to evade national rules. We find that researchers in strict regulatory environments generally have more international coauthors, and this tendency is strengthened after the introduction of CRISPR in 2012 (p=0.09). We see no significant evidence that weaker regulations are associated with more clinical trial development or patent applications after 2012.

4.2 Public Backlash

We next examine cross-national spillovers arising from public controversies via a survey experiment. The survey examines backlash among the general public in response to a hypothetical, norm-violating application of gene-editing technology. To gauge the spillover effect, we examine both the effect of controversial activity in one's own country as well as activity

	$Dependent\ variable:$		
	Int'l Coauthors	Clinical Trials	Patents
	(1)	(2)	(3)
Regulation	2.743***	-3.184***	-664.499***
	(0.530)	(0.552)	(141.227)
CRISPR	11.089***	1.415	-42.111
	(1.133)	(1.053)	(209.687)
Regulation×CRISPR	1.203*	-0.714	-56.107
	(0.716)	(1.054)	(213.015)
Controls	✓	✓	✓
Observations	647	647	592

Table 2: Effect of National Regulations on Coauthorship, Clinical Trials, Patents. The table displays coefficient estimates and dyad-clustered standard errors from a linear model. Statistical significance is denoted by: *p<0.1; **p<0.05; ***p<0.01.

in a foreign country.

The online survey was conducted in July 2020 on a sample of 1,075 Americans quota-sampled to US census margins. We embed an experiment in the survey designed to address two questions. First, do controversies over the use of gene editing reduce public support for the technology and its potential applications? Second, does public backlash spill across national jurisdictions?

In the experiment, all respondents receive a summary of gene-editing technology. It reads:

All organisms, from bacteria to lizards to humans, have molecules called DNA, or deoxyribonucleic acid. These DNA molecules contain the genetic code for each

²⁹See Appendix A.3 for full survey text and Table A2 for sample summary statistics. While the survey was fielded on a sample of 1,200 respondents, we restrict our sample to the 1,075 individuals who passed attention checks. Our survey was conducted on the platform Lucid, and our pre-registration plan can be found under EGAP 20200505AA.

organism. DNA provides the instructions that determine an organism's physical characteristics and control how it develops, functions, and reproduces.

In recent years, scientists have developed new gene-editing technologies that can permanently alter an organism's DNA. These technologies allow scientists to make targeted changes to DNA molecules in plants and animals, modifying their biological traits. For example, scientists have edited the genes of wheat plants to make them easier to grow.

Some respondents are then randomly assigned to a treatment condition where they read about a gene-editing controversy. Among treated respondents, we randomize whether the controversy occurs in the US, UK, or China. The treatments are presented as a hypothetical news article set in near future. To increase external validity, we model the experimental intervention on the real-world controversy surrounding He Jiankui. We present the text for the UK treatment condition here:

Birth of Genetically Altered Babies in the UK Provokes Outcry January 25, 2021

[LONDON]—A British research team announced that they have used a new geneediting technology to alter the DNA of a group of infants. In an unprecedented intervention, scientists on the research team deleted a set of genes believed to be linked to breast and prostate cancer. The deleted genes are not considered essential to basic biological functions in humans, but the long-term effects of their removal are unclear. The research team plans to periodically examine the infants throughout their lives to assess any side effects of the genetic alteration.

The disclosure this week of the research — carried out in the UK — has sparked urgent debate about the ethics of genetic alteration. The infants' birth represents a significant and controversial leap in the use of gene-editing technology. The

British study has also increased concerns about a future in which parents produce "designer babies" with selectively improved traits, such as height or intelligence.

After treatment assignment, respondents rate their agreement with the following four statements on a scale of 0 (no agreement) to 10 (complete agreement).

- Research in the US involving gene editing should be more strictly regulated.
- US patients should have access to medical treatments that involve gene editing.
- The US government should provide funding for gene editing research.
- Most US scientists conduct their research in a safe and responsible manner.

The statements estimate the extent of public confidence in the safety of gene-editing technology and support for continued development. Respondents' answers constitute our dependent variables in the analyses below.

Our theory of public backlash against emerging technologies implies two patterns of response. First, we expect that respondents who read about a controversy in their own country will be less supportive of gene-editing research. This "domestic public backlash" should heighten demand for strict regulation, depress calls for patient access to gene therapy, decrease support for funding gene-editing research, and decrease confidence in the safety of scientific research. Second, we expect foreign controversies to similarly reduce public support for gene editing among US respondents. A "public backlash spillover" occurs if the controversial use of gene editing generates a domestic backlash even when the scandal occurs in another country.

We report treatment effects for each outcome of interest in Figure 2. Coefficients in the figure represent the treatment effect of exposure to a gene-editing controversy, compared to the control (no controversy) condition. Within each panel, we display the effect of a

³⁰See Table A3 in the appendix for point estimates and standard errors. Table A4 reports similar results among respondents who passed an alternative attention check.

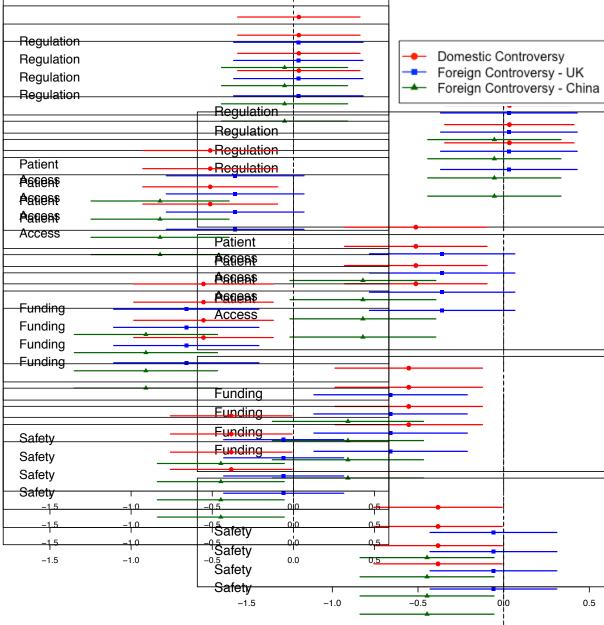


Figure 2: Public Response to Gene-Editing Controversy. The figure shows the treatment effect of a hypothetical gene-editing controversy in the US, UK, and China, with 95% confidence intervals. The panels report effects on the four dimensions of public support listed above.

domestic controversy, a foreign controversy in the United Kingdom, and a foreign controversy in China.

We find evidence of a domestic public backlash in three of four outcomes. When American respondents read about a hypothetical misuse of gene-editing technology by American

researchers, they significantly reduce support for patient access to gene therapies (p = 0.02) and public funding for gene-editing research (p < 0.01). They also have diminished perceptions of the safety and responsibility of scientific research (p = 0.09) in the United States. On average, the domestic controversy treatment shifts opinion on each of these outcomes by approximately 0.5 points. Contrary to expectations, respondents do not increase demand for strict regulations in reaction to domestic gene-editing scandals (p = 0.61). This null finding may reflect a ceiling effect, as even respondents in the control condition call for strict regulations in high numbers (see Appendix Figure A4 for the distribution of responses).

There is clear evidence that backlash is not limited by national jurisdiction. As in the domestic scenario, neither of the foreign scandals significantly affects attitudes about gene-editing regulation. However, support for public funding of gene-editing research significantly decreases in response to foreign controversies in the UK and China (p < 0.01). Respondents also reduce confidence in the responsibility of US scientists (p = 0.07) and support for gene therapies (p < 0.01) in the China condition. The UK controversy does not affect perceptions of safety but does decrease support for gene therapies (p = 0.07). Notably, the effects of domestic and foreign controversies are statistically indistinguishable across all four outcomes.

Together, these results suggest that the public does not discriminate between domestic and foreign research controversies. We find clear evidence that, for some public policy outcomes, the spillover effect of controversial research in one national jurisdiction negatively impacts domestic support for gene editing in another jurisdiction.

In Appendix Section A.4, we demonstrate that spillovers in public backlash occur empirically using data from Twitter posts surrounding the He Jiankui controversy. In a sample of over 50,000 tweets, we find that posts increase in volume, negative sentiment, and moral outrage following the scandal. Consistent with the spillover effect, negative sentiment and moral outrage occur at higher levels outside of the jurisdiction (China) in which the scandal occurred.

4.3 Discussion

Our findings provide evidence for two sources of interdependence that afflict national governance of gene-editing technology. We show that weaker national regulations lure scientific talent and boost gene therapy development, creating pressures for governments to engage in regulatory competition. We also demonstrate that the risks of weak national rules are not fully internalized by the home country. If a government's permissive regulation increases the risk of inappropriate behavior, the resulting backlash spills across national boundaries.

Taken together, our results suggest that governments have compelling incentives to lower regulatory barriers beyond the level they would otherwise prefer. Each country can obtain individual economic benefits from weakening rules, while the risks of doing so are diffusely spread across multiple jurisdictions. If governments respond rationally to these incentives, effective regulation will be under-produced and the systemic risk of misuse will rise. However, the resulting public backlash in domestic constituencies could reduce demand for the technology or lead to knee-jerk regulatory reactions, halting continued progress. While these inefficient boom-and-bust cycles can occur in the absence of interdependence, they are more likely to occur in the presence of regulatory arbitrage and public opinion spillovers.

By illuminating patterns of interdependence among states, our findings make an implicit case for international policy coordination. In doing so, we reinforce calls for instruments of global governance to manage "emerging technologies that affect the global commons" (Oye et al., 2014). International institutions are designed to manage interdependence and reduce transaction costs (Keohane, 1984; Haggard & Simmons, 1987). The recent guidelines adopted by the World Health Organization (WHO, 2021) are consistent with this function. The recommendations establish a floor of basic ethical and safety protections and encourage harmonization of disparate rules governing the technology. If successful, this would limit the

³¹While we do not directly test the effect of regulation on scientific scandals, we observe a positive correlation between weak regulatory environments and retractions in gene editing studies (see Appendix A.6).

scope for regulatory competition and reduce the risk of scandalous applications.

5 Conclusion

In this paper, we put forward a theory of and provide evidence for international interdependence created by technological advancement. We demonstrate first that states are subject to regulatory arbitrage by scientists and practitioners in the field of gene-editing. The accessibility of cheap, powerful technology creates opportunities and incentives for regulatory arbitrage. To measure regulation, we construct a new index of gene-editing restrictions across three gene-editing issues: gene therapy, germline editing, and in-vitro embryonic modification. Using novel sources of data on scientific employment, gene therapy trials, and patents, we find evidence of arbitrage behavior all three domains.

We also argue that controversial applications of gene-editing technology trigger public backlash that can spill across national boundaries. As far as we are aware, we are the first to identify this theoretical mechanism that links public attitudes in one country to policy decisions in another. The effect of these controversies can be dramatic: in the history of gene editing, high-profile scandals led to collapsed public support, the abandonment of commercial applications, and harsh regulatory responses. We demonstrate the mechanism in an original survey experiment on American respondents and confirm its external validity using social media data from a real-life controversy. Our findings support the existence of a public backlash spillover that can undermine confidence in gene-editing technology.

While our tests focus on the field of gene editing, we argue these dynamics recur in the governance of disruptive technological innovations more generally. Additional work can assess the generalizability of our theory by expanding empirical tests to other fields. In addition to contemporary emerging technologies like artificial intelligence, historical disruptions such as the nuclear energy and information technology revolutions may have similarly enhanced

international interdependence via these mechanisms.

More broadly, our paper helps outline a new agenda for understanding how technologies affect interstate cooperation and the demand for global governance. Emerging technologies shape a range of transnational spillovers in addition to the two we emphasize here. Genetic manipulation of the natural environment, including vegetation or insect populations, can easily traverse national jurisdictions. Similarly, the use of digital currencies may disrupt international financial systems or exacerbate collective action problems like carbon emissions. Advancements in artificial intelligence and robotics could reshape labor demand in ways that interact with the politics of trade, human rights, or military competition. Future work should test and expand upon these effects of technological innovation.

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1 Appendix

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A.1 Tables

	DV: Mental Health Scientist Relocation		
	(1)	(2)	
Regulatory Difference	0.048 (0.047)	-42.89 (33.28)	
CRISPR	0.118 (0.147)	14.09 (9.63)	
Regulatory Difference \times CRISPR	0.023 (0.078)	-0.006 (0.0146)	
Controls		✓	
Dyad FE		✓	
Year FE		✓	
Observations	20,592	14,435	

Table A1: Placebo Test: Employment Relocation of Mental Health and Eating Disorder Researchers: Linear model estimates for the volume of mental health and eating disorder researchers who relocate to institutions in another country. Column 2 includes the following controls (not shown): GDP origin country, GDP destination country, GDP per capita origin country, GDP per capita destination country, Patent Applications origin country, Patent Applications destination country, R&D origin country, and R&D destination country. Standard errors are clustered by country dyad. Statistical significance is denoted by: *p<0.1; **p<0.05; ***p<0.01.

Variable	Sample Proportion		
Party ID			
Democrat	0.36		
Republican	0.36		
Independent	0.28		
Age			
18-30	0.15		
31-45	0.36		
46-60	0.30		
over 60	0.19		
Education			
High School or Less	0.30		
Some College	0.32		
Bachelor's Degree	0.22		
Post-Graduate	0.17		
Gender			
Female	0.51		
Male	0.49		
Ethnicity			
White	0.83		
Black or African American	0.14		
Asian	0.01		
Other	0.02		
Hispanic			
Yes	0.13		
No	0.87		
Household Income			
< \$25,000	0.55		
\$25-50,000	0.21		
\$50-75,000	0.14		
> \$75,000	0.10		
Region			
Northeast	0.21		
Midwest	0.20		
South	0.38		
West	0.22		

Table A2: Survey sample statistics. For each category, we report the proportion of respondents who fit into the category among those that answered the relevant question.

	Dependent variable:			
	Regulations	Access	Funding	Safety
	(1)	(2)	(3)	(4)
US Controversy	0.030	-0.515**	-0.551**	-0.380**
	(0.193)	(0.213)	(0.220)	(0.192)
UK Controversy	0.037	-0.365*	-0.641***	-0.055
·	(0.203)	(0.216)	(0.228)	(0.188)
China Controversy	-0.048	-0.820***	-0.910***	-0.450**
v	(0.198)	(0.217)	(0.225)	(0.200)
Observations	1,197	1,193	1,198	1,199
Adjusted R ²	-0.002	0.010	0.012	0.004

Table A3: Survey Results. Estimated treatment effects and robust standard errors for the survey experiment. Effects are relative to the control condition (no additional informattiton). Statistical significance is denoted by: *p<0.1; **p<0.05; ***p<0.01.

	Dependent variable:			
	Regulations	Access	Funding	Safety
	(1)	(2)	(3)	(4)
US Controversy	-0.109 (0.197)	-0.644^{***} (0.223)	-0.572^{**} (0.231)	-0.416^{**} (0.199)
UK Controversy	-0.531^{**} (0.237)	-0.742^{***} (0.269)	-1.165^{***} (0.286)	0.080 (0.219)
China Controversy	-0.104 (0.223)	-1.170*** (0.240)	-1.347^{***} (0.254)	-0.447^{**} (0.224)
Observations Adjusted R ²	955 0.002	$952 \\ 0.023$	954 0.032	955 0.006

Table A4: Survey Experiment Results on Respondents who pass Manipulation Check. Results of the survey experiment on the sample of respondents who successfully pass a manipulation check. We check for attention by asking treated individuals in which country gene-editing occurred. Statistical significance is denoted by: *p<0.1; **p<0.05; ***p<0.01

A.2 Figures

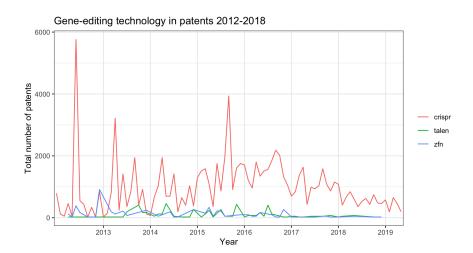


Figure A1: Gene-editing Patent Applications, 2012-2018: The figure displays annual patent applications related to CRISPR (red), TALENS (green), and ZFN (blue) technologies. Data from Oribit Intelligence.

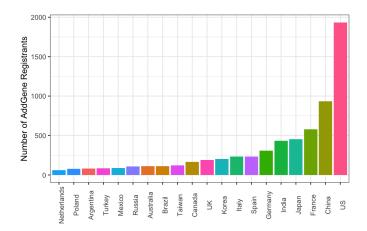


Figure A2: AddGene Registered Researchers by Country of Origin: Number of researchers registered on the AddGene website by country of origin. Data collected by authors. AddGene is an organization that stores and disseminates genetic material used in published studies. AddGene has served as a popular repository for CRISPR plasmids since Jinek et al. (2012) used it to store materials from their landmark paper. Researchers register on the AddGene website and pay a fee for the plasmid transfer. They are then free to replicate the parent study or alter the plasmids for their own research purposes. Although CRISPR-related materials are a minority of AddGene's repository, they are among the most commonly requested plasmids from researchers.

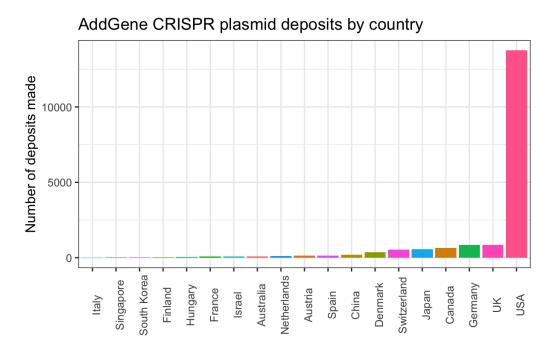


Figure A3: AddGene depositors by Country of Origin. Data collected by authors.

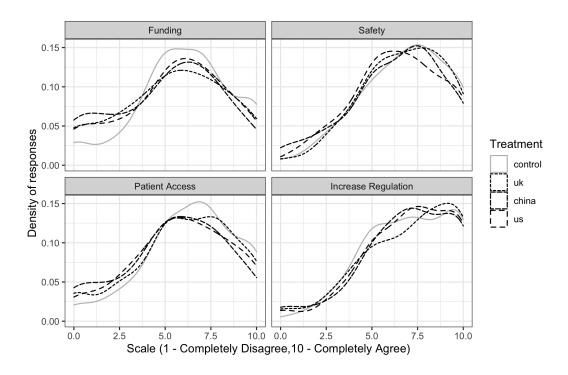


Figure A4: Distribution of Responses for Outcome Variables. The figure displays the distribution of responses by treatment condition for each of four outcomes.

A.3 Survey Experiment Consent and Text

In line with the APSA Principles and Guidance for Human Subjects Research, the author who provided the funding for this experiment submitted the survey protocol to the relevant Institutional Review Board (IRB) Human Subjects Committee prior to launching the survey experiment. The IRB reviewed this survey experiment and granted an exemption under federal regulation 45 CFR 46.104 (2)(ii) (IRB Protocol ID 2000027424). The survey does not contain deceptive material, intervene in political processes, or collect sensitive and/or personally identifiable information.

Respondents were recruited through Lucid, an automated marketplace that connects researchers with online research participants. The authors compensated Lucid \$1 per completed interview. Lucid contracts with suppliers who provide financial incentives to survey respondents in the form of cash, gift cards, or loyalty reward points. All respondents are voluntary participants based in the United States. For further details, see https://luc.id/wp-content/uploads/2019/10/Lucid-IRB-Methodology.pdf.

Before beginning, potential respondents are informed that the study is voluntary and assured that their responses will be kept confidential. We then ask for their informed consent:

You are invited to participate in a research study that will take approximately 15 minutes to complete. You will be asked to answer some questions about yourself and your preferences.

There are no known or anticipated risks to you for participating.

Participation in this study is completely voluntary. You are free to decline to participate, to end participation at any time for any reason, or to refuse to answer any individual question without penalty or loss of compensation. The researcher will not know your name, and no identifying information will be connected to your survey answers in any way. The survey is therefore anonymous.

If at any time you have questions or concerns about the survey or your rights or welfare as a research subject, contact [Author name] at [Author email].

If you would like to talk with someone other than the researchers to discuss problems or concerns, to discuss situations in the event that a member of the research team is not available, or to discuss your rights as a research participant,

you may contact the [Author's university] Human Subjects Committee, [phone number], [email]. Additional information is available at [Link to statement of research participant's rights at Author's university].

If you would like to participate, simply click the 'I agree to participate' box below, then click the >> button to start the survey.

After a set of demographic questions, all respondents are provided the following information:

Now you will read some information related to recent advances in biotechnology. All organisms, from bacteria to lizards to humans, have molecules called DNA, or deoxyribonucleic acid. These DNA molecules contain the genetic code for each organism. DNA provides the instructions that determine an organism's physical

In recent years, scientists have developed new gene-editing technologies that can permanently alter an organism's DNA. These technologies allow scientists to make targeted changes to DNA molecules in plants and animals, modifying their biological traits. For example, scientists have edited the genes of wheat plants to

characteristics and control how it develops, functions, and reproduces.

Respondents are then randomly assigned to one of four conditions:

1. Control - no additional information

make them easier to grow.

- 2. Domestic Controversy
- 3. Foreign Controversy (UK)
- 4. Foreign Controversy (China)

Those assigned to conditions 2-4 additionally read a hypothetical news article regarding a gene-editing controversy. We show the text for the Foreign Controversy (UK) here.

Below you will read a hypothetical news article about the use of gene-editing technology. The article describes events that could take place in the future. After you have read about the situation, we will ask for your opinions.

Birth of Genetically Altered Babies in the UK Provokes Outcry January 25, 2021

[LONDON]—A British research team announced that they have used a new geneediting technology to alter the DNA of a group of infants. In an unprecedented intervention, scientists on the research team deleted a set of genes believed to be linked to breast and prostate cancer. The deleted genes are not considered essential to basic biological functions in humans, but the long-term effects of their removal are unclear. The research team plans to periodically examine the infants throughout their lives to assess any side effects of the genetic alteration.

The disclosure this week of the research — carried out in the UK — has sparked urgent debate about the ethics of genetic alteration. The infants' birth represents a significant and controversial leap in the use of gene-editing technology. The British study has also increased concerns about a future in which parents produce "designer babies" with selectively improved traits, such as height or intelligence.

Finally, we ask respondents to rate their agreement with four statements on a scale from zero to ten.

Please indicate your level of agreement with the following statements, with "0" representing complete disagreement and "10" representing complete agreement.

- Research involving gene editing should be more strictly regulated in the US
- US patients should have access to medical treatments that involve gene editing
- The US government should provide funding for gene editing research
- Most US scientists conduct their research in a safe and responsible manner

A.4 Public Backlash: social media data

We follow (Müller et al., 2020) in scraping and analyzing tweets with the keyword CRISPR. Using Twitter's API through Barrie & Ho (2021)'s R package, academictwitter, we pulled approximately 50,000 tweets that contain the word "CRISPR" in the 50 days prior to and after the He Jiankui controversy. The bottom panel of Figure A.5.1 shows a histogram of the appearance of CRISPR in tweets over this time period. Using a bag-of-words procedure, we take the average sentiment of each tweet by identifying the proportion of positive words

¹We exclude replies and retweets in our analysis.

in the tweet. Higher proportions indicate higher levels of positive sentiment. Only English-language tweets are included in the sentiment analysis. Only tweets with at least one word with a positive or negative valence are included in this sample. The top panel of Figure A.5.1 displays change in average sentiment over time.

While sentiment in tweets does capture overall public opinion of Twitter users towards CRISPR technology, new research suggests that certain forms of expression on social media are more likely to drive conversations (Brady et al., 2021). In particular, tweets expressing moral outrage receive higher levels of positive feedback online and are therefore more likely to be seen and to influence online sentiment (Brady et al., 2021). We use Brady et al. (2021)'s measure of moral outrage to understand whether controversial events in CRISPR technology influence not only sentiment, but the type of language that drives greater engagement with the overall conversation. We note that moral outrage is a form of negative sentiment. We measure the number of words stems associated with moral outrage in each tweet. The middle panel of Figure A.5.1 displays change in moral outrage over time. Table A.5.1 reports the pre-post change in sentiment and outrage expressed in tweets.

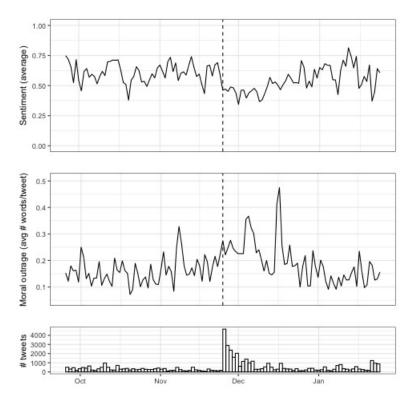


Figure A.5.1: CRISPR tweet sentiment: Top panel: sentiment analysis of tweets including the word "CRISPR" from September 2018 - January 2019. Higher values indicate more positive sentiment. Middle panel: moral outrage in same sample of tweets. Higher values indicate more moral outrage. Bottom panel: histogram of number of tweets per day. Dashed black line on November 26, 2018, the day the He Jiankui controversy became public. Data collected by authors.

Subsetting the data to only tweets with identifiable geolocation data for the associated Twitter user, we replicate the main analysis by user country. We limit our analysis to English-language tweets, which likely affects the composition of countries in this sample. Figure A.5.2 displays the results for countries with more than 10,000 unique tweets that mention CRISPR. As Figure A.5.2 shows, the revelation of the gene-editing controversy produced negative sentiment in the days afterwards for every country in the sample. Importantly, the scandal did not occur in any of these countries. (China is not included in the sample as the country blocks access to Twitter for regular users.²)

²https://help.twitter.com/en/rules-and-policies/state-affiliated-china

	Dependent variable:		
	Sentiment	Moral Outrage	
	(1)	(2)	
Post-He Jiankui	-0.110***	0.062***	
	(0.006)	(0.004)	
Constant	0.609***	0.158***	
	(0.005)	(0.003)	
Observations	21,856	50,839	
\mathbb{R}^2	0.013	0.004	
Adjusted R ²	0.013	0.004	
Note:	*p<0.1; **p<0.05; ***p<0.01		

Table A.5.1: Change in Tweet Sentiment. Average tweet sentiment and number of expressions of moral outrage before and after the news of the He Jiankui controversy broke on November 26, 2018. Sentiment refers to expressions of positive sentiment and is only measured for tweets with at least one word that expresses positive or negative sentiment; positive values indicate more positive sentiment. Moral outrage refers to words that are categorized as expressing moral outrage and indexes the number of words per tweet that reflect this sentiment; positive values indicate increased outrage. Robust standard errors in parentheses. Sample is all English-language tweets mentioning "CRISPR" from September 2018 - January 2019.

In addition to expressing a particular form of negative sentiment, posts that use the language of moral outrage also generate greater engagement in the form of likes and retweets (Brady et al., 2021). Using a dictionary of moral outrage terms (e.g., "abhor", "hate", "shame"), we measure the extent of moral outrage expressed in tweets. The top panel of Table A.4.2 confirms that tweets that contain higher levels of moral outrage are liked and retweeted more often. Tweets with greater positive sentiment are liked more (but not retweeted more). The bottom panel shows that moral outrage tweets are not more liked or retweeted post-scandal. In contrast, negative sentiment tweets are retweeted and liked more often post-scandal. These results 1) confirm that moral outrage tweets have greater engagement in the realm of gene-editing and 2) show that negative sentiment tweets are engaged with at

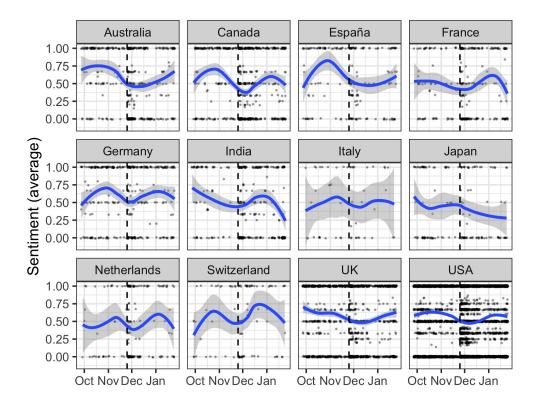


Figure A.5.2: CRISPR tweets by location: Sentiment analysis of tweets including the word "CRISPR" from September 2018 to January 2019 by Twitter user location. Only countries with over 10,000 CRISPR-related tweets included. Higher values indicate more positive sentiment. Blue line is a fitted loess model; grey bar indicates 95% confidence interval. Dashed black line on November 26, 2018, the day the He Jiankui controversy became public. Data collected by authors.

higher rates post-scandal. Combined with prior findings about higher numbers of negative sentiment and moral outrage tweets in response to scandal, this suggests even greater levels of public opinion shifts to anti-gene-editing sentiment after scientific scandals. The finding is particularly salient when one considers that public policy attitudes are driven by social cues from peers in addition to elites (Kertzer & Zeitzoff, 2017).

These results confirm the external validity of our experimental findings. The He Jiankui scandal was salient to the public as evidenced by the steep increase in tweets about gene-editing technology after the scandal was made public. These tweets were also more negative and contained higher levels of moral outrage. Finally, geo-located tweets confirm that the

	Dependent variable:			
	Retweets	Likes	Retweets	Likes
	(1)	(2)	(3)	(4)
Moral Outrage	0.715*** (0.185)	1.373*** (0.349)		
Sentiment			-0.022 (0.205)	0.877** (0.402)
Observations	50,839	50,839	21,856	21,856
	(5)	(6)	(7)	(8)
Moral Outrage	1.066** (0.444)	2.332** (1.023)		
Sentiment			0.556 (0.385)	2.035** (0.944)
Post-He Jiankui	-0.022 (0.107)	-0.421^* (0.236)	0.352 (0.364)	0.058 (0.746)
Moral Outrage*Post-He Jiankui	-0.455 (0.487)	-1.214 (1.075)		
Sentiment*Post-He Jiankui			-0.846^* (0.452)	-1.816^* (1.023)
Observations	50,839	50,839	21,856	21,856

Table A.5.2: Tweet Virality. Correlation between average tweet sentiment and number of expressions of moral outrage with tweet virality (likes and retweets). Sentiment refers to expressions of positive sentiment and is only measured for tweets with at least one word that expresses positive or negative sentiment; positive values indicate more positive sentiment. Moral outrage refers to words that are categorized as expressing moral outrage and indexes the number of words per tweet that reflect this sentiment; positive values indicate increased outrage. Robust standard errors in parentheses. Sample is all tweets mentioning "CRISPR" from 9-25-2018 to 1-25-2019. Bottom panel displays pre-post results on virality.

scandal, which occurred in China, had effects on international public opinion about geneediting technology.

A.5 Regulations and Scientific Retractions

We directly test for a correlation between regulatory stringency and scientific scandals through data on scientific retractions in the field of genetic editing. Retractions may occur for many reasons including ethical malfeasance, data manipulation, and data errors. Retractions are characterized as scandals in the scientific community (Azoulay et al., 2017) and have significant negative effects on the field of study in which they occur (Azoulay et al., 2015). Within our sample of gene-editing papers, 674 unique scientists were involved in 121 redacted papers as indicated in the PubMed database. We identify a clear negative correlation between the level of regulation in a country and the number of redactions ($\rho = -0.39$, p = 0.02) as well as the proportion of retracted papers ($\rho = -0.28$, p = 0.12) in a given country. Figure A.6.1 visualizes these relationships.

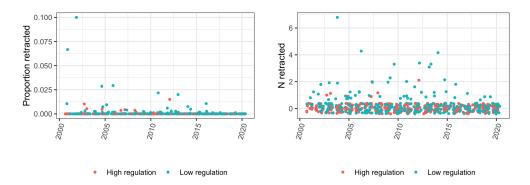


Figure A.6.1: Regulations and Scientific Retractions. Left panel depicts proportion of papers published by scientists located in a given country in a given year that were retracted. Right panel depicts raw numbers of retracted papers by country-year.

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