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Genetic fallout in bio-cultural landscapes: Molecular imperialism and the cultural politics of (not) seeing transgenes in Mexico

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Abstract

This article explores the trajectory of the global controversy over the introgression (or not) of transgenes from genetically modified maize into Mexican indigenous maize landraces. While a plurality of knowledge-making processes were deployed to render transgenes visible or invisible, we analyze how a particular *in vitro* based DNA-centered knowledge came to marginalize other forms of knowledge, thus obscuring other bio-cultural dimensions key to the understanding of gene flow and maize diversity. We show that dominant molecular norms of proof and standards of detection, which co-developed with the world of industrial monocropping and gene patenting, discarded and externalized non-compliant actors (i.e. complex maize genomes, human dimensions of gene flow). Operating in the name of high science, they hence obscured the complex biological and cultural processes that maintain crop diversity and enacted a cultural–political domination over the world of Mexican landraces and indigenous communities.

Keywords

co-production, epistemic cultures, genetically modified organisms, laboratory–field relations, legibility–illegibility, maize biocultures, Mexico’s indigenous landraces, normalization

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Introduction

In 2001, an article in the journal *Nature* by David Quist and Ignacio Chapela reported scientific findings of transgenes that had escaped from genetically modified (GM) maize and introgressed into Mexican indigenous maize landraces (Quist and Chapela, 2001). This news came a few months after reports that thousands of truckloads or railcars of unauthorized StarLink GM maize for human food in the United States had been contaminated. In just 4 days, the StarLink contamination case was confirmed and provoked thoroughgoing action, including a huge compensation bill and the recall of hundreds of product lines (Lezaun, 2003). The *Nature* findings were highly unwelcome to the Mexican government, which had established a moratorium on transgenic maize planting in Mexico in 1998, as well as to GM commercial interests, which claimed GM crops were both safe and containable. The findings suggested that no remote place could escape uncontrolled transgene fallout and highlighted the new, uncharted transnational artificial genetic landscapes created by the global agricultural biotechnology enterprise.

The Mexican GM maize issue was a resoundingly global controversy. Mexico is a global center of origin and ongoing diversification of maize, and its maize landraces constitute a genetic pool seen as critical for the future of international crop-breeding and food security. The controversy connected small farming communities with international arenas and biotech laboratories, and raised pressing issues over indigenous peoples' rights, farmers' livelihoods, neoliberal globalization, biological and cultural diversity, and commoditization of life (Foyer, 2010; Kinchy, 2012; Wainwright and Mercer, 2011). However, although socio-environmental groups have striven to foreground these political, cultural, and economic issues, the GM maize controversy has been 'scientized' and channeled into narrowly technical 'risk' and 'detection' framings (Kinchy, 2012; Wynne, 2001). In this article, we show how the technoscientific culture around molecular biology and DNA-based GM detection, given sovereignty in this controversy, enacted much more than a specific, solely technoscientific analytical cut. This technoscientific culture embodied and projected an extensive and powerful framing which performed a particular political-economic and cultural normativity by denying recognition to other key biological and human dimensions of Mexican maize.

The first publication reporting the presence of transgenes in Mexican landraces (Quist and Chapela, 2001) was immediately strongly attacked (see Delborne, 2008). The next peer-reviewed publication (Ortiz-García et al., 2005) reported no transgenes in samples from the same region, providing reassurances that were widely publicized that no transgenes existed in Mexican maize. It was not until 2007–2009 that further *positive* findings, which since 2001 had multiplied and circulated in non-peer-reviewed reports and unpublished documents, were published in peer-reviewed scientific journals (Dyer et al., 2009; Piñeyro-Nelson et al., 2009b; Serratos et al., 2007). In an age of high-throughput post-genomics, 8 years to confirm the presence of transgenes in analytical samples is a surprisingly long time.

Our inquiry starts from this basic question: Why did it take 4 days to confirm the presence of unauthorized GM maize and undertake costly remedial action in the case of StarLink in the United States, yet many years to do the same in Mexico? This situation warrants a closer look at sociotechnical processes that turn, or fail to turn, invisible

transgene flows into evidence of 'contamination'. We argue that a key element in the Mexican controversy is that a particular scientific subculture, molecular biology, with its taken-for-granted epistemic norms stretching far beyond the laboratory, gained global authority over what constituted good evidence. The apparently purely technical issues of transgene detection embody much larger issues of political and cultural asymmetries of power, abetted systematically through science.

The point that knowledge often involves making invisible phenomena visible, bringing them to produce recognizable and repeatable signs, traces, and tracks, has become normal currency of Science and Technology Studies (STS) work. However, its corollary, that such knowledge production and its renderings of visibility also perform a corresponding rendering of other forms of *invisibility*, or *non-knowledge*, has only come to the fore more recently to further elaborate on sociology of scientific knowledge in public arenas. A key dimension of both these forms of epistemic ordering to produce forms of invisibility as well as visibility is that these ordering processes occur through mutual ordering, or co-production, with corresponding orders of social practices and relations (Jasanoff, 2004). How can invisible transgenes be turned into visible signals that unmask their presence within local maize landraces? In Mexico, there were multiple knowledge-making processes available for rendering transgenes visible or invisible. But only the DNA-based approach that used state-of-the-art polymerase chain reaction (PCR) was deemed a worthy basis for articles published in top international scientific journals. This reflected (and reinforced) the hegemony of molecular biology, which focused on laboratory practices of molecular control and reduced its intellectual frame to only those factors that were amenable to such control, while non-compliant actors, non-human or human, were discarded and externalized.

Our article analyzes key beliefs, norms, and commitments embodied in the technical standards granted privilege to adjudicate the controversy. Those technical standards also embody commitments that impose normative structure far beyond the laboratory alone, as we see later, for example, with their neglect of important differences of genome structure between modern hybrid and indigenous maize varieties. In shaping what is counted as normal, these commitments also implicitly perform a particular worldview, reinforcing or co-producing a dominant political and social order. The technoscientific culture of molecular biology thus enacted a domination not only over Mexican indigenous farming cultures but also over other salient epistemic cultures, such as population genetics, agroecology, and anthropology. This article shows how such dominations obscured the complex bio-cultural processes that maintain crop diversity, delegitimized farmers' agroecological knowledge and forms of life, and threatened the very processes that maintain biodiversity.

The first section of this article introduces our theoretical perspective. The second sketches the trajectory of the controversy. In the third, we explore the plurality of knowledge-making processes deployed to render transgenes visible or invisible and show how a particular *in vitro* based DNA-centered knowledge came to marginalize other forms of knowledge and to obscure bio-cultural dimensions of gene flow and crop genetic diversity. The fourth section examines the tensions between the standardizing gaze of (commercial) PCR detection, and the fluid and heterogeneous Mexican landrace maize genomes. We conclude by discussing how dominant molecular norms of proof and

standards of detection embody a cultural–political domination over the bio-cultural world of Mexican landraces and the indigenous communities that cultivate them.

The article is based on interviews (some repeat visits) with about 30 key scientific and policy protagonists, as well as related peer-review reports and correspondence. We also make use of unpublished material, obtained from our informants, that allowed us to document all the sampling surveys and almost all the manuscripts on this issue submitted to scientific journals between 2000 and 2009.

The STS of not seeing

Given the inevitably selective processes that order scientific knowledge cultures, each ‘epistemic culture’ (Knorr-Cetina, 1999) has its own distinct practices of producing and dealing with non-knowledge. Several discussions from STS illustrate the point and can be taken to suggest that the very productivity and power of technoscientific standards may be quintessentially what also produces invisibility of non-standard actors, both human and non-human. Kleinman and Suryanarayanan (2013) describe the normative knock-on effects of particular epistemic cultures as multi-layered: the effects range and graduate over time and through routinization, from explicit choices to the naturalized practices of encultured technoscientific paradigms. They come to embody normative ‘choices’ constitutionally (Jasanoff, 2011). Such normative standards – which structure, extend, and empower each technoscientific paradigm – can exclude and effectively delete salient variables and conditions that happen to be non-compliant with the standard’s normative gaze. This occurs both within and beyond the domain itself, depending on perceived reach into applied worlds. Busch (2011) has also delineated this ambivalent property of the normativity of technoscientific paradigms in his analysis of standards and certification schemes. MacKenzie et al. (2013) have extended the discussion into the broad domain of life sciences and related forms. Here, especially, aims to standardize encounter a teeming variety of exuberantly emergent forms of non-compliance.

With respect to agricultural genetically modified organisms (GMOs) and their potential impacts, Böschen et al. (2006), Bonneuil (2006) and Böschen (2009) have identified contrasting and competing interactive epistemic cultures, such as molecular biology, ecology, and systems agronomy. These authors have shown that molecular biology is distinguished from related scientific specialties by its characteristic function, including its research objects (the gene as a commanding molecule in the programmable cell factory, as distinct from being in flux in meta-populations and ecosystems), epistemic objects, scales and units of analysis, methods of producing data, ways of constructing explanations, framing of risks, and attitudes to unknowns. Each function determines what is relevant or irrelevant to the epistemic project and also contributes to the bringing into being and consolidation of a corresponding human cultural order.

The epistemic culture of molecular biology is notable with respect to its culture of non-knowledge. Through her ethnographic work with the contrasting epistemic cultures of high-energy physics and molecular biology, Knorr-Cetina (1999) demonstrated that the latter is far less engaged with the limits and boundaries of its own knowledge. High-energy physics is attentive to limits such as disturbances, errors, unexpected events, and uncertainties. However, molecular biologists seldom explore possible unknowns when

an experiment fails or delivers unforeseen results. Instead, they tend to vary the conditions of the experiment until the *expected* outcome emerges, a heuristic strategy that Knorr-Cetina (1999) calls ‘semi-blind variation’ (p. 110).

In the case of the Mexican GM maize controversy, *epistemic culture* is an analytic that helps to illustrate the relationships between molecular biology’s particular understanding of (trans)gene flow and understandings of other, marginalized, epistemic cultures, such as (agro)ecology, population biology, or anthropology of farmers’ seed practices. In this case, molecular biology’s claim to sound science involved invocation of stringent norms of proof for detecting transgenes, while strategies such as semi-blind variation provided normative force to exclusion of uncertainties, anomalies, non-standard forms, and complexities. Attention to this culture of non-knowledge sheds light on how the controversy involved blindness to the dynamics and natural–cultural hybridities of maize genomes that have evolved over centuries in Mexican landscapes.

There is a useful complement to the epistemic cultures perspective in Proctor and Schiebinger’s (2008) analysis of the production of scientific ‘ignorance’ as social practice. Proctor and Schiebinger identify multiple kinds of ignorance. One kind, that which is actively and strategically fabricated, has been well documented in the Mexican GM maize controversy (Delborne, 2008; Foyer, 2010; Kinchy, 2012). Indeed, in his analysis of the first year of the controversy, Delborne (2008) speaks of the intensive deployment of strategically fabricated ignorance in terms of ‘epistemological tyranny of the intellectual majority’ and the ‘institutional majority’. However, Proctor and Schiebinger outline another highly relevant yet less discussed kind of ignorance, the effect of *framing* (Wynne, 1989, 1992) – to focus on *this* as salient is inevitably to ignore *that*, possibly without even knowing that one has made that ‘choice’. This selective ignorance is embodied in paradigms, or epistemic cultures, that always articulate theoretical, material, and social options in ways that neglect phenomena outside of their scope. Ignorance as an effect of framing is an issue in the controversy over Mexican GM maize because the framings privileged by the scientific journals were orchestrated by a set of technoscientific mechanisms that selected who and what were, or were not, relevant and meaningful. These included a selective set of questions, a selective set of data, a selective set of technical skills to create and interpret correspondent data, specific sources of knowledge and information, and specific norms of proof.

These selective framings also embodied particular epistemic–cultural beliefs and commitments: that the knowledge produced by molecular biology is universally superior to knowledge produced by farmers, agronomy, ecology, population genetics, and anthropology; that the laboratory and the molecular scale, not the field, is the relevant locus for sound science; and that a few genetically homogeneous cultivars designed for global markets and their infrastructures can adequately represent, genomically and politically, Mexico’s complex maize bio-cultural landscape. These epistemic norms resonate with a political economy in which agriculture oriented to global markets competes on unequal terms with indigenous maize biocultures.

In the Mexican GM maize controversy, the beliefs and norms of the epistemic culture of molecular biology gained force from two related processes. First, there was a chain of extraction, reduction, and translation that allowed signs observed on gels in laboratories to speak for the distant presence or absence of transgenes in fields. And second, there

was a purification and standardization in the laboratory micro-world, which were materially and symbolically projected onto the diverse, fluid, and extensive macro-worlds of indigenous maize. Actor–Network Theory (ANT) has contributed much to the understanding of these types of processes. For example, Latour (1995) analyzes the production of stable networks across innumerable transfer situations. Much scientific effort aims to produce conceptual and material objects that remain stable from field to laboratory and back (Callon et al., 2009: 48–68). These *immutable mobiles* differ from fuzzy objects that would lose their salient properties when displaced. Once purified and disciplined in the laboratory, immutable mobiles are projected back into field situations, reordering them according to laboratory standards. In this controversy however, there was also the key mediating role of international scientific journals in giving sole authority to a molecular framing of the issue.

20th century genetics and molecular biology have constituted the cultivar and the gene as immutable mobiles – objects of exact science, modernization policies, and global trade (Bonneuil, in press; Fox-Keller, 2000). Van der Ploeg (1993) offers a powerful anthropological study of such constitutive processes in another center of origin of agriculture, indigenous Andean potato-farming. The study illustrates how modernizing agricultural experts and institutions promoted a singular laboratory-optimized genotype, effectively requiring diverse potato-farming environments, conditions, and practices to be standardized in reflection of the ‘sovereign’ laboratory (Scott, 1998). While such optimization was scientifically justified in its own terms, those terms themselves demanded questioning. They also performed an epistemic culture of non-knowledge or ignorance through their mutually corresponding epistemic and institutional framing.

ANT has illuminated how technological standardization works to universalize scientific knowledge through laboratory practices that reshape the world through the circulation of commensurable objects (O’Connell, 1993). However, it fails to address the power asymmetries generated through these processes. STS work by scholars like Star (1991), has shed important light on how technological standardization black-boxes heterogeneity and difference and disables certain non-aligned forms of life in reflection – and reinforcement – of asymmetric power (see also MacKenzie et al., 2013). James C. Scott’s extensive anthropological work also highlights the issue of power asymmetries and how these are obscured in natural knowledge cultures. For example, drawing on Foucault, he shows how states and markets have organized societies and ecosystems so as to render them ‘legible’ (Scott, 1998) for purposes of control. As noted by Scott and Star, as well as Busch (2011), such strategies of extraction, simplification, standardization, and legibility-making attempt deletion of objects, phenomena, and forms of life that do not comply with centers of surveillance, calculation, and power. In return, non-compliant actors may resist such standardization and legibility through manifesting resistance or through cultivating autonomy through invisibility (Scott, 1990, 2009).

Critical STS resources – the analytic of epistemic cultures, complemented by the model of ignorance and exclusion as an effect of framing, and by co-production, the ANT account of immutable mobiles, and analyses of relevant power–knowledge asymmetries – provide a perspective from which to explore connections between the most technical and social aspects of non-knowledge in the Mexican GM maize controversy. In the following sections, we discuss how DNA detection methodologies enacted

selective reduction and standardization of what are in situ fluid, as well as connected in-the-field genomes, so as to produce an in vitro knowledge of them that may have co-produced a corresponding cognitive, political, commercial, and cultural order (Jasanoff, 2004, 2006). The aim of this case history is to elucidate the multi-layered ways in which scientific knowledge – including as a form of non-knowledge – can act in the name of nature to reflect and embody, and thus perform, particular cultural values and power relations.

Transgene escape: the trajectory of a research and policy problem

In the 1970s, research had found that 20 percent of samples from local landraces of maize in Chiapas showed traits introgressed from Green Revolution hybrid varieties (Ortega-Paczka, 1973). Two decades later, in 1990, the US geneticist John Doebley published the first molecular evidence for gene flow between maize and wild *Zea* relatives. He warned that ‘an engineered gene in maize could spread ... throughout the teosinte population’ that grow in some regions of Mexico and proposed ‘not to grow transgenic maize in those regions’ (Doebley, 1990: 443). However, policymakers and mainstream science did not pick up this unwelcome early evidence of gene flow. In 1988, GM maize field trials were authorized nationwide in Mexico when President Salinas de Gortari initiated negotiation of the North American Free Trade Agreement (NAFTA). Enacted in 1994, NAFTA led to an import boom of US maize in Mexico. US commercial cultivation of transgenic maize in Mexico started in 1995.

Before 2001, in both the United States and Mexico, minimal attention was paid to agroecological implications of maize transgene flow as a research or policy issue. In the United States, the issue was not addressed by the National Academy of Sciences’ report on the impacts of GMOs (National Research Council, 2002), and none of the hundreds of biosafety research projects submitted to the Biotech Risk Assessment Program between 1992 and 2000 addressed it.¹ In Mexico, the issue did emerge within the small group of experts on the Mexican National Agricultural Biosafety Committee (CNBA). Among these, the agronomist and biotechnologist Antonio Serratos co-organized two meetings to address transgene escape in 1995 and 1997 (Serratos et al., 1997, 2000). However, although a few articles on gene flow were published in the meeting proceedings, these did not find their way into international journals. And while a few members of the CNBA persuaded the Ministry of Agriculture to issue a de facto moratorium on Mexican GM corn field trials and commercial growing in 1998, pro-GM interests counter-lobbied. These efforts included a letter to *Science* arguing that there was ‘no need for concern’ about gene-escape from transgenic maize (Martinez-Soriano and Leal-Klevezas, 2000).

The issue of maize transgene flow mushroomed dramatically in November 2001 when Quist and Chapela’s *Nature* article reported ‘the presence of introgressed transgenic DNA constructs in native maize landraces grown in 2000, in remote mountains in Oaxaca, Mexico’ (Quist and Chapela, 2001: 541). One year after the United Nations (UN) Cartagena protocol on global biosafety, and three years after the Mexican moratorium on GM corn field trials, the discovery of transgenes in a remote southern area

suggested a radical failure of Mexico's national biosafety policy, as well as the need for research 'to trace the flow of genetic material over bio-geographical regions' (p. 542).

However, these findings – and the challenges they posed to industry and government reassurances of control and containment – provoked immediate and intense contestation (Delborne, 2008: 2011). In April 2002, the editor of *Nature* took the unprecedented step of withdrawing the journal's support for the original article, stating that 'the evidence available is not sufficient to justify the publication of the original paper' (Kaplinsky et al., 2002: 600). Criticisms of Quist and Chapela's article actually focused on a point that was secondary to the central issue, the finding of transgenes in samples of indigenous landraces. The critics instead alleged that the study over-interpreted inverse polymerase chain reaction (iPCR) results that indicated fragmentation of transgene inserts. However, they also mostly succeeded in undermining the credibility of the central finding of transgenes in indigenous maize.

Nonetheless, especially in Mexico, the *Nature* article prompted a hot political struggle around GM crops, including research and policy activism and widespread anti-genetic engineering campaigns from peasant, indigenous, and environmental organizations (Foyer, 2010; Kinchy, 2012). A Ministry of Environment agency, the Instituto Nacional de Ecología (INE), and the Department of Agriculture sampled maize throughout Mexico and confirmed Quist and Chapela's findings (Alvarez-Morales, 2002; Ezcurra et al., 2002). Much positive data were released by the authorities and by non-governmental organizations (NGOs) in international conferences and press releases (Herrera et al., 2002). However, none of these were published in peer-reviewed scientific journals (CECCAM et al., 2003; Foyer, 2010) until 2007, and, as we describe below, the only intervening journal article was instead one (Ortiz-García et al., 2005) that claimed to show no such GM contamination of indigenous maize fields.

In 2002, a group of organizations and rural communities petitioned the Commission for Environmental Cooperation (CEC), a body created under the NAFTA, to attend to a wider set of issues than environmental risk assessment alone, such as the cultural value of maize and peasants' livelihoods. In their report on maize and biodiversity, CEC experts echoed some of these dimensions, resulting in rejection of the experts' recommendations by the governments of the United States and Canada. However, the CEC report still framed the issues mainly around risks and narrow technical questions. It affirmed the presence of transgenes in this way:

Transgenes have entered some landraces of maize in Mexico. This finding was confirmed by scientific studies sponsored by the Mexican government. However, no peer-reviewed summaries of this work have been published and information released to the public has been vague. (Secretariat of the CEC, 2004: 16)

While it acknowledged transgene escape, this framing of the issues required that data be published in peer-reviewed journals in order to be deemed trustworthy and relevant to policy. This requirement became a strategic resource for those corporate and policy actors who were concerned to cool the controversy and lift the ban on GM maize until further reports of transgenes in Mexican maize were published in scientific journals in 2007.

In a second phase of the controversy, from 2004 to 2008, the pro-GM policy line of the Department of Agriculture won its battle against the more cautious Mexican government environmental authorities. The latter had achieved Mexico's ratification of the UN 2000 Cartagena Protocol on Biosafety. However, the agricultural authorities retaliated with Mexico's 2003 signing of the North American Biotechnology Initiative, and a 2005 biosafety law that accommodated lifting the GM maize moratorium (Foyer, 2010). This policy shift was supported by a 2005 article in the prominent US journal *Proceedings of the National Academy of Sciences (PNAS)*. The article, published through a fast-track procedure, reported no findings of transgenes in the Oaxaca mountains from 2003 to 2004 (Ortiz-García et al., 2005). Negating the many positive results found and reported by public agencies and researchers, including in the year 2003 (Comisión Intersecretarial de Bioseguridad de los Organismos Genéticamente Modificados (CIBIOGEM), 2004; Landavazo Gamboa et al., 2006), this one negative article was the most intensely publicized and globally recognized report on the topic. Until 2007, the positive studies could not get published in international scientific journals, despite submissions from 2002 onward – all rejected – to *Nature*, *PNAS*, and others.

Curiously, Ortiz-García, the lead author of the 2005 *PNAS* article, had been one of the co-authors of an article that reported positive findings, which had then recently been rejected by *Nature* (interview with S. Ortiz-García, November 21, 2008). Her other co-authors of the *PNAS* article included the directors of INE, a senior scientist of the US GMO detection company Genetic ID (GID), and US Ohio State University professor Alison Snow. The article was the main, indeed only, reference used to discredit Quist and Chapela over the issue of whether transgenes had ever really escaped into Mexican landraces. At the very least, the *PNAS* article suggested that transgenes had disappeared from the Oaxaca region. Reports of the *PNAS* article in *Nature* and *Science* with titles like, 'Four years on, no transgenes found in Mexican maize' and 'Calming fears, no foreign genes found in Mexican maize', exuded blanket scientific reassurance and invited extrapolation of the negative findings to all Mexican territory (Kaiser, 2005; Marris, 2005). In Mexico, this high-profile absence/reversibility narrative helped government and industry actors to claim that 'it is proven that the transgenes do not remain in the environment' (Biotech industry representative in Mexico, quoted by Fitting (2011: 59); see also Prakash (2005)).

The publication, between 2007 and 2009, of three studies showing positive results in several states of Mexico opened a new phase of the controversy. The first, coordinated by former CNBA member Antonio Serratos and published in *Frontiers in Ecology and the Environment*, reported the presence of 'transgenic proteins in maize' in the Federal District (i.e. around Mexico City) (Serratos et al., 2007). A second article, authored by Piñeyro-Nelson et al. (2009b), a research team from the Universidad Nacional Autónoma México (UNAM) in Mexico City, was published in *Molecular Ecology*. This study reported positive results (at overall frequencies of around 1%) for samples collected in 2001 and 2004 in Oaxaca. These results confirmed Quist and Chapela's findings. A third article, co-authored by a group of US and Mexican scientists, was published in *PLoS One* (Dyer et al., 2009). The article reported findings of the GM protein in more than 3 percent of the 419 seed-lots sampled in 2002 across 14 Mexican states.

These three positive sets of findings, however, attracted little attention in scientific and media arenas compared with Ortiz-García et al.'s (2005) article. Only Piñeyro-Nelson et al.'s findings were covered in another major publication, a 'news' section in *Nature* (Dalton, 2008). Neither did the findings influence the Mexican policy arena. This was striking, particularly in light of the 2009 public volte-face by one of the two US co-authors of the 2005 Ortiz-García 'denial' findings (Snow, 2009). The author declared herself fully convinced by Piñeyro-Nelson et al.'s work. Given the structure of institutional power in Mexico, neither farmers', environmental, and indigenous activist movements nor independent scientists could use these unquestioned positive scientific findings to alter the pro-GM policy line that had solidified around 2005. Once political closure had been achieved, the new scientific findings seemed unable to reopen the debate over transgenes. GM maize field trials were officially authorized in 2009 (a decision suspended recently by judicial interventions).

An issue relevant to their lack of policy influence is that these three publications were in journals of lesser standing than *Nature* and *PNAS*. The world of scientific publications is highly hierarchical, and the importance of articles both within and beyond science itself is determined by their host journals' 'impact factors'. In 2009, *Nature*'s impact factor was 34.48 and *PNAS*'s was 9.43, while *Frontiers in Ecology and the Environment*, *Molecular Ecology*, and *PLoS One*'s were between 4.35 and 6.92. Correspondingly, different detection methodologies had different epistemic authority. The *Frontiers in Ecology and the Environment* and *PLoS One* articles, which reported studies that used agronomic or immunological protein assays rather than higher profile DNA-based methodologies, failed to attract weight. Piñeyro-Nelson et al.'s 2009 article did use DNA-based assays, but these were attacked by GID scientists as misinterpretations of the PCR results or even 'indicative of contamination in the laboratory' (Schoel and Fagan, 2009: 4143). We return to this later.

Why was a single publication of negative findings enough to cool down the controversy by 2005, while several other simultaneous surveys that confirmed transgenic findings in Mexican landraces did not enjoy international scientific journal publication? Why, while most biologists acknowledge that maize gene flow occurs at significant rates, was further positive DNA-based evidence repeatedly required? Why was DNA evidence deemed the only legitimate proof, while protein detection from simple immunoassays was deemed illegitimate, despite having been accepted in the US StarLink case? The next section analyzes how such a particular standard of evidence – whose influence on the overall trajectory of the controversy has been shown – became hegemonic and how it also embodied and performed a much larger set of non-scientific commitments.

Making transgenes visible: constructing the authority of in vitro knowledge

We have tracked all the GM sampling surveys conducted in Mexico after Quist and Chapela's 2001 article and documented their subsequent fates in scientific and public arenas. On the one hand, as Table 1 shows, of the more than 18 surveys, only 1 – that published in the high-profile international journal *PNAS* (Ortiz-García et al., 2005) – reported not finding transgenes in landraces. On the other hand, of the more than 17 surveys that reported positive findings, only three achieved international journal publication.

Table 1. An overview of GM Maize sampling surveys in Mexico (2001–2009) and their fate.

Surveys post Quist and Chapela (2001)	Detecting transgenes	Not detecting transgenes
Data published in a peer-reviewed international journal	3: Serratos et al. (2007); Piñeyro-Nelson et al. (2009); Dyer et al. (2009)	1: Ortiz-García et al. (2005), <i>PNAS</i> , fast track
Data submitted in a manuscript but rejected by peer-reviewed international journals	6+: Alvarez-Buylla et al. to <i>Nature</i> , 2002; Piñeyro-Nelson et al. to <i>PNAS</i> , 2007; Piñeyro-Nelson et al. to <i>Plos One</i> , 2008; Dyer et al. to <i>PNAS</i> (Mexico Rural Household Survey (ENHRUM), 2002); Dyer et al. to <i>Applied Ecology</i> (idem.); Dyer et al. to <i>Agriculture & Human Values</i> (idem.)	0
Data not submitted to any international journal (with year of collecting)	9+: INE – 2001–2009 (2003 and 2004 excluded); Grupo ad-hoc (CIBIOGEM-Sagarpa) – 2001, 2002; INIFAP – 2003 (Landavazo Gamboa et al., 2006); ETC group-CECCAM-CASIFOP (Press release in 2003)	–

GM: genetically modified; *PNAS*: *Proceedings of the US National Academy of Science*; INE: Instituto Nacional de Ecología; CIBIOGEM: Comisión Intersecretarial de Bioseguridad de los Organismos Genéticamente Modificados.

When the same institution conducted a survey with several collecting years, we counted it as several different surveys.

These three published articles are the survivors of obstacle-course-like peer-review processes: the work by Piñeyro-Nelson et al. (2009b) was published only after rejected submissions to *Nature* in 2002, *PNAS* in 2007, and *PLoS One* in 2008; similarly the work by Dyer et al. (2009) had been previously rejected by *PNAS*, *Applied Ecology*, and *Agriculture & Human Values*. This suggests that after 2001, positive results faced more difficulties than negative results in attempts to achieve publication in authoritative journals. *Nature* and *PNAS* operated as gate-keepers, channeling and shaping what knowledge was ultimately produced and which kinds of evidence constitute reliable proof. In this section, we examine how these high-profile journals both privileged a particular way of detecting transgene flow and also discarded others' understandings, thereby disempowering subaltern voices on more than gene flow alone.

How can invisible transgenes be turned into visible signals and meanings? We have observed four ways of making transgenes visible during the controversy: (1) phenotypic judgments derived from consideration of the whole plant, (2) the search for agronomic traits specifically expressed in the GM plants, (3) the search for distinct protein products of the transgenes in plant cells, and (4) the search for transgenic DNA sequences in the plant genome.

The first approach was implemented by some indigenous community organizations, members of the *Red en Defensa del Maíz* activist network. They claimed that shamans and some experienced farmers could visually detect 'contaminated' plants because they look monstrous (interview with Aldo Gonzales, April 2, 2009). Monster hunting, resonating with cosmologies that associate monstrosities with a moral hurt, constituted a tool for raising community awareness against GMOs.

The second avenue tested the expression of agronomic traits specific to the GM plants. As most transgenic constructs confer tolerance to a specific herbicide, the finding of a landrace that is unaffected by application of a particular herbicide constitutes evidence of transgene introgression. This type of test was easy to conduct, in either an experimental plot or the field, by applying the herbicide to a leaf with a pencil. The zone would turn yellow if non-transgenic and remain green if transgenic.

The third option tested the expression of transgene protein products through immunological methods (enzyme-linked immunosorbent assay (ELISA) tests, lateral flow, strip band, etc.). These techniques are cheap, reliable, and simple. They were the most used methods to identify preservation routines along the food chain by industry and regulatory agencies in the United States until the mid-2000s. In 2000, Friends of the Earth sampled food products and engaged a then-young detection company, GID, which ran immunological tests in taco shells, finding a protein (Cry9C) belonging to an unauthorized GM maize (StarLink) commercialized by Aventis for animal feed. Within a few days, Kraft Foods, having double-checked and confirmed this contamination, recalled 300 different product lines. This caused large drops in US corn shipments, and Aventis was forced to pay almost US\$1 billion to affected farmers, consumers, and companies (Lezaun, 2003). So, immunological detection methods were compelling enough to authorize strong policy measures.

The fourth approach was based on the molecular identification of DNA sequences specific to the genetic construct transferred to the plant genome in the transformation event (e.g. the Southern blot hybridization technique and qualitative and quantitative PCR techniques). The crucial process of PCR amplification of DNA from samples is notoriously sensitive to variable factors (including contamination, but also genuine differences in genome complexity between hybrid and landrace maize genomes), especially when quantification (quantitative polymerase chain reaction (qPCR)) is being attempted. PCR methods, although precise and technologically sophisticated, are hard to handle properly and prone to both false-positives and false-negatives. They also require knowledge of the precise DNA sequences of the GM constructs, which are often restricted from scientific access as commercially confidential.

During the controversy, these DNA methodologies became the gold standard of evidence to the detriment of the three other ways of rendering transgenes visible. By late 2000, Quist and Chapela could have reported transgene findings based on rapid

immunoassay methods, as was done in the StarLink case or in the 1999 case in which Greenpeace Mexico reported detection of GM *Bt* maize on maize shipments from the United States. But to optimize the academic and policy impact of their discovery, Quist and Chapela preferred to wait and carefully construct both their PCR methodology and their audience (Delborne, 2011). Their report, however, was criticized for using *only* PCR methods, which were deemed prone to false-positives (Christou, 2002; Kaplinsky et al., 2002). Quist and Chapela (2002), therefore, conducted DNA hybridization (i.e. Southern blot) tests to confirm their findings (p. 602). This episode set the standard: when new manuscripts were submitted to *Nature* to provide evidence for the presence of transgenes in Mexican landraces, they had to use both PCR and Southern blot to have a chance of acceptance. While phenotypic, agronomic, or immunological evidence was regarded as sufficient in other cases (e.g. the StarLink case), *Nature* and *PNAS* only published articles based on DNA methods. This hegemony of DNA-based hard evidence had deep political impacts in the Mexican controversy.

Unsurprisingly, indigenous groups' claims to ascertain 'contaminated' maize plants visually (as monsters) had no authority in the international scientific arena or in the national policy arena. Most maize scientists consider the existence of monstrous forms in maize a well-known and longstanding phenomenon. Furthermore, the DNA-based gold standard of evidence embodied the idea that the laboratory, not the field, was the legitimate place to produce knowledge. This meant that, a priori, community organizations could not produce legitimate data. For instance, the regional indigenous organization in Oaxaca where Quist had first detected transgenes in 2000, Union de Comunidades Forestales Zapoteco-Chinanteca (UZACHI), wanted to conduct detection and monitoring itself. It had a fairly well-equipped laboratory, and its director had learned molecular techniques. But the INE refused to defer to UZACHI for the testing of the samples collected in UZACHI communities (interview with Lilia Perez, April 4, 2009). Seeking hard evidence with international credibility in mind, INE preferred to subcontract public and private biotechnology laboratories for the analytical work (interview with S. Ortiz-García, November 21, 2008). A coalition of NGOs and community organizations decided to use immunological tests to conduct their own survey, independent from the government. They reported the presence of transgenes in landraces in several Mexican states (CECCAM et al., 2003). This community science strategy aimed both at producing evidence more rapidly than studies stuck in long peer-review processes and at 'empowering local farming communities' (interview with Ana de Ita, January 26, 2009). But these NGO findings and their subsequent 'open letter' to the government failed to influence policy or to reach the international press. Even the fact that they found Cry9C, the allergen protein present in StarLink, in many samples did not make a difference. A Mexican scientist noted that 'the problem in Mexico ... is that nobody cares about what is not published in an international journal', while reflecting on the strong policy gate-keeping influence of academic journals (interview with Francisca Acevedo, November 21, 2008). Government officials dismissed these results as anecdotal and unreliable because they were not published in a peer-reviewed journal.

Not only were farmers' and indigenous organizations' knowledge and claims disempowered by the DNA-based proof requirement enforced by top scientific journals, but also Mexican agronomists' in situ expertise was discarded. Heirs of a research school

initiated by Hernández Xolocotzi (1913–1991) who had surveyed maize landraces with Green Revolution scientists in the 1950s, maize genetic resource specialists possess an in-depth knowledge of maize landraces at the crossroads of botanical, genetic, agronomic, and ethnoecological perspectives. These maize scientists (calling themselves ‘maiceros’) can morphologically identify any maize plant as one of the approximately 60 races identified in Mexico. They can also phenotypically distinguish any modern commercial hybrid from local landraces or from intermediary forms. This expertise provides evidence about whether a plant that turned positive in a detection test is a GMO commercial hybrid (i.e. for some reason ‘lost’ in a farmer’s field) or a landrace that has introgressed a transgene into its own genome. But this phenotypic expertise was rejected by the molecular biologists who served as reviewers for the top scientific journals (Referee 1 on Alvarez-Buylla et al.’s rejected manuscript, *Nature* to Dr Ezcurra, 22 September 2002, personal communication from S. Ortiz-García). Maiceros’ knowledge was deemed inadequate.

The requirement for DNA-based methodologies as the gold standard of evidence not only marginalized farmers and Mexican maize scientists but also made it harder for Mexican molecular biologists to contribute effectively to the debate. In the early 2000s, these DNA-based techniques were indeed mastered only by a few Mexican laboratories, for research rather than for routine testing. However, it proved hard for these laboratories to meet the standards imposed by high-profile journals. This is why two of the three 2007–2009 publications that reported positive findings (Dyer et al., 2009; Serratos et al., 2007) relied only on immunoassays. A third Mexican group – composed of Elena Alvarez-Buylla’s unit at UNAM, Rafael Rivera’s group at Cinvestav, and scientists from the INE – worked hard in 2002 to meet *Nature*’s standards of proof, saying, ‘We had to repeat PCR assays in large numbers ... Develop a satisfying Southern blot methodology took us a hard work ... We met many kinds of hurdles’ (interview with Elena Alvarez-Buylla, December 10, 2008). Finally, the manuscript that reported positive results was rejected by *Nature* for lack of ‘standard procedure and controls’ for PCR and Southern Blot (Referee 2 comment, in *Nature* to E. Ezcurra, 23 September 2002, personal communication from E. Alvarez-Buylla).

It was exactly this obstacle of producing publishable data that led to a split within the group. After *Nature*’s refusal, INE sent their samples to the US-based GID. From these, GID obtained inconsistent and ‘strange’ (interview with S. Ortiz-García, November 21, 2008) quantitative PCR results and, rather than consider the possibility that the genomic heterogeneity and complexity of Mexican landraces (see below) could explain such results, concluded that ‘contamination’ had occurred in the two Mexican laboratories. Persuaded by GID, INE, which had no expertise in molecular methods, stopped delegating the analytical work to Alvarez-Buylla and Rivera’s laboratories and began collaboration with GID instead. This resulted in the switch to the negative 2005 *PNAS* publication (interviews with Ortiz-García, Alvarez-Buylla, and Piñeyro-Nelson, March 18, 2009). Having to provide both DNA-based and immunological evidence, the two public Mexican laboratories took some further years to verify and publish their positive data (Piñeyro-Nelson et al., 2009b) – which also by then included a direct ‘blind’ test of GID’s own claims to exclusive methodological purity.

Thus, in the 2000s, the growing authority of DNA-based detection techniques, and the corresponding stringent norms of proof of leading scientific journals, proved instrumental in silencing other voices and their forms of maize expertise. This growing authority was fueled by rapid advances in molecular biotechnologies (namely, new generations of PCR techniques) and strengthened by the view of DNA as the universal basic molecule of life, the demand that DNA technologies be the most precise and accurate (Fox-Keller, 2000; Rabinow, 1996).

The long march of a bio-cultural understanding of gene flow

Spearheaded in Mexico after World War II by the International Maize and Wheat Improvement Centre (CIMMYT), the Green Revolution paradigm included a division of labor between breeders as innovators and farmers as mere users, a strong analytical separation between the genotype and the environment, and a standardization of the latter around the laboratory-manipulated ‘optimal’ genotype (Van der Ploeg, 1993). This genetic agricultural modernization project succeeded in increasing production in the large farms of the northern states. Nevertheless, it failed to meet the needs of most Mexican farmers since indigenous maize landraces rather than modern hybrids are still grown on an estimated 80 percent of maize acreage (Aquino et al., 2001).

By 1970, some Mexican agricultural scientists had already reported that farmers could not afford Green Revolution technological packages and that hybrids were not adapted to farmers’ environments (Cotter, 2003: 290–309). Led by Hernández Xolocotzi (1977), these scientists pioneered agroecological approaches – which combined ethnobotany, agronomy, genetics, and ecology – and called for a re-Mexicanization of research. Although initially marginalized, this agroecological and ethnoecological research school was developed again in the mid-1990s by the anthropologist Steve Brush at the University of California (UC), Davis, through a new worldwide program to implement participatory management of crop diversity (Bonneuil and Fenzi, 2011; Brush, 2000).

The new program funded *in situ* maize diversity conservation projects in Mexico and fueled new research on farmers’ seed systems. This bio-cultural approach provided a deeper understanding of the farmers’ very diverse knowledge practices – cosmologies, seed sharing, mixing of seed, farmer selection, cultural transmission of practical farming savvy, cooking knowledge, and complex multiple use criteria in continual play. Farmers’ practices had been operating as a distributed evolutionary process acting on the genetic structure and dynamics of maize populations at a landscape level over decades and centuries in a wide range of ecological conditions. The farmers’ system, which manages and selects from numerous ‘meta-populations’ in a wide range of environments, accounts both for the existence of some genetic continuity and for the maintenance of diversity among numerous maize landraces; the system also maintains some stability at the phenotypic level (Bellon, 1996; Hugo Perales et al., 2003; Pressoir and Berthaud, 2004).

However, anti-reductionist and field-sensitive approaches remained marginal. As a population geneticist at CIMMYT put it, ‘The then head of the CIMMYT viewed us as “tree huggers” ... We preached in the desert’ (interview with J. Berthaud, June 17, 2009). Molecular biology and its promised molecular intensification of the Green Revolution

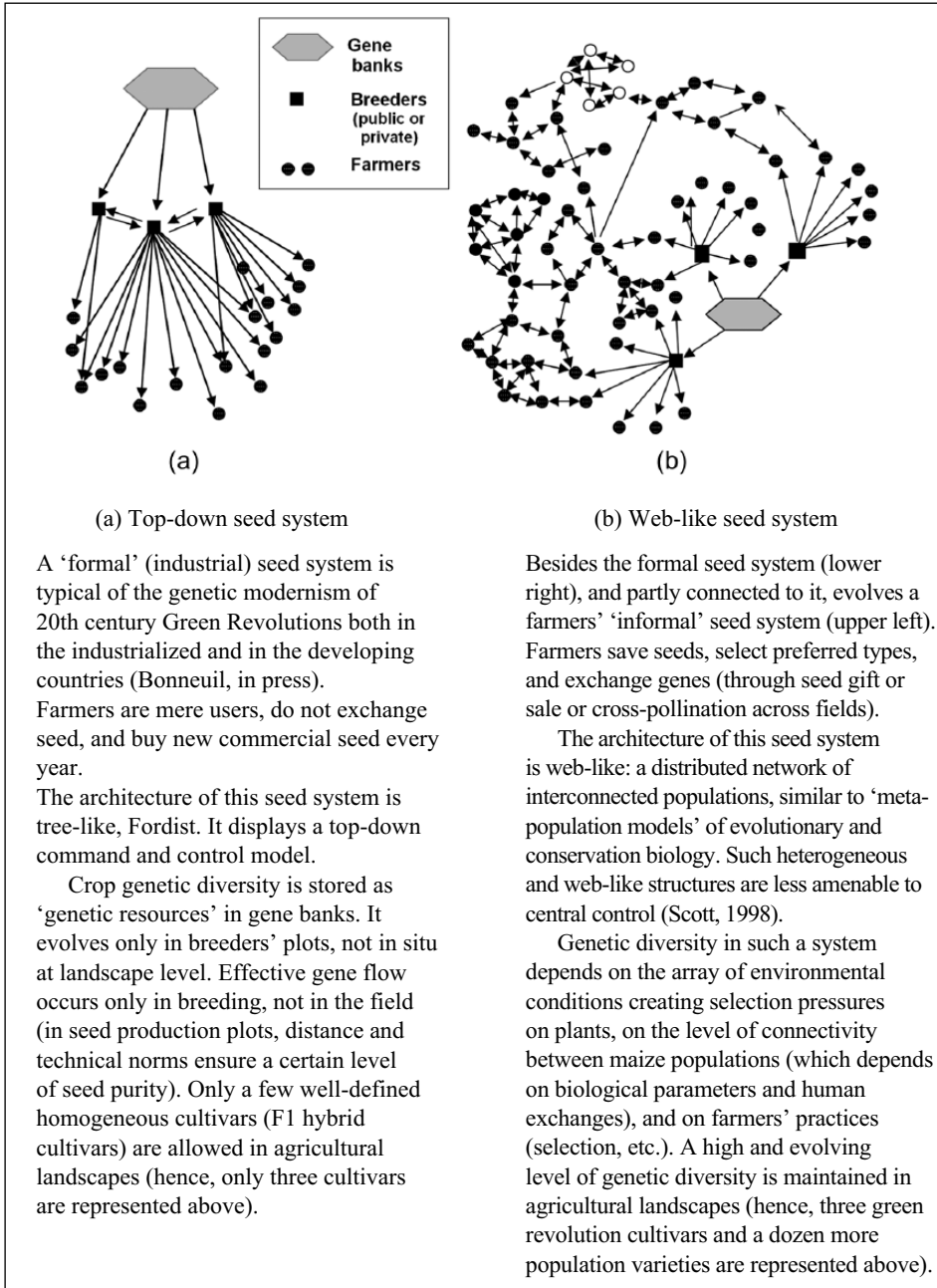


Figure 1. Contrasting seed systems, contrasting knowledge cultures (a) green revolution seed system and (b) farmers' web-like seed system.

paradigm, on the one hand, and bio-cultural approaches to gene flow, on the other, constituted two different epistemic cultures. Each developed in integration with a different seed

system, as outlined in Figure 1: a top-down industrialized maize world, in one case, and a web-like bio-cultural indigenous maize seed system, in the other. The two cultures differ deeply in their views of what genes can do, how genes flow, and what maize diversity is.

The epistemic culture of molecular biology gives preference to static and homogeneous forms of life – which are deemed better for generating robust commercial technoscience – reliable mass production, global markets, and intellectual property (Bonneuil, in press). It views crop diversity as a standing reserve, which local farmers hardly know how to harness, and sets a modernist boundary between ‘nature’ and ‘culture’. Crop biodiversity is abstracted from its ecological and social contexts to be used as a resource for crafting elite commodity cultivars (Bonneuil and Fenzi, 2011). These abstracted ‘genetic resources’ are centralized in ex situ seed-banks and processed through molecular control laboratories, which constitute the centers of calculation of a tree-like seed system. Co-produced with industrial monocropping of commercial hybrids for global markets (see Figure 1(a)), the molecular biology epistemic culture tends to see diversity, genome variation, and in situ gene flow only as problems to be controlled.

In contrast, the bio-cultural epistemic culture views maize diversity as a web of dynamic ecological and human connections, ‘not just as “genetic resource,” but as diversity in the field’ (interview with M Bellon, 30 April 2009). Rather than basing sound science on purified systems amenable to property rights, the bio-cultural approach views environmental and genetic heterogeneity, web-like in-situ gene flows (see Figure 1(b)), and autonomous diverse traffic across the nature/culture border as its very object of investigation (Bellon, 1996; Brush, 2000; Dyer and Taylor, 2008; Van Heerwaarden et al., 2012).

Making farmers invisible

The tension between molecular biological and bio-cultural understandings of gene flow was far more than epistemic. Strikingly, farmers’ roles in shaping gene flows at landscape levels were erased from the two articles published in the higher standing journals. Within the constraints of a short letter to *Nature*, Quist and Chapela (UC, Berkeley researchers) were led to obscure their longstanding collaboration with Oaxacan indigenous communities and to elide the agency of farmers in the shaping of maize genetic landscapes (Quist and Chapela, 2001; interview with Chapela, 2 November 2006). This spiriting away of farmers to meet *Nature*’s narrative genre repeats the 17th-century scientific revolution’s ‘invisible technician’ accounts (Shapin, 1989). In the *PNAS* article (Ortiz-García et al., 2005), there are 21 references to ‘farmers’, but farmers are always treated as mere *receptors* of scientific knowledge. Indigenous farmers, who may have ‘unknowingly’ planted GM grain, are depicted as an epistemically vacuous public with its own (non-scientific, ‘local’, and ‘traditional’) ‘cultural perceptions’ and irrational fears of GMOs. Hence, there was need for a patronizing scientific expertise (Wynne, 1996) for the state to inform and reassure those ‘traditional farmers’ (Ortiz-García et al., 2005). Only the lower ranking later articles acknowledged farmers’ agency in the dynamic shaping of maize genetic diversity at landscape levels and reported having interviewed dozens of farmers to derive knowledge of the ‘seed systems’ that crucially informed their sampling or modeling strategies (Dyer et al., 2009; Piñeyro-Nelson et al., 2009b; Serratos et al., 2007). The lower ranking articles reflected an alternative, bio-cultural, way of making scientific knowledge, one rejected by *Nature* and *PNAS*.

Such high-profile journals also contributed to the production of ignorance about the human dimensions of (trans)gene flow through the letters that they selected (and rejected) for publication after Quist and Chapela's article. Although *Nature* only published technical molecular biology critiques, we found additional submitted comments from the bio-cultural perspective that were rejected by *Nature* (Bellon and Berthaud, 2004; Soleri, personal communication, 2009). In these discarded comments, gene flow was not discussed as a surprise to be proven with hard molecular data, but as an already well-known reality, which might be key to maize diversity, depending on both biological parameters and farmers' practices.

Although marginalized by the molecular biology epistemic culture, the interdisciplinary (agronomic, anthropological, genetic, and ecological) study of farmers' seed systems turned out to be essential knowledge. Because they collected maize in farming communities on a regular basis, the bio-cultural epistemic community practitioners were better prepared to design robust sampling schemes than were laboratory-based molecular biologists. In light of farmers' practices and maize pollination biology, bio-cultural practitioners knew that ears from the same plot and from the same household are closely genetically related and on this basis noted a key limitation of Ortiz-García et al.'s (2005) *PNAS* article. Molecular biologists took the seeds deriving from a single cob as the unit of analysis. As a result, Ortiz-García et al. calculated that the probability of transgene presence in Oaxaca mountains was below 0.01 percent. However, more robust sampling schemes determined the 'effective population size' – a population genetics concept, alien to molecular biology's epistemic culture – by taking the farm-household as the unit of analysis. This more refined bio-culturally informed sampling and probabilistic analysis, based on population genetics and statistical and anthropological knowledge of farmers' seed systems, corrected this assessment to 1–4 percent (using a sampling scheme encompassing only 124 different households), even before their differences in interpretation of PCR and gel-band readings (Cleveland et al., 2005).

It is astonishing that such a crucial issue as sampling did not enter the controversy before 2005, and it only did so through journals of lesser standing than *Nature*, *Science*, and *PNAS*. This reflects the fact that peer-reviewers mobilized by high-profile journals were molecular biologists who had poor knowledge of and little interest in farmers' practices, population genetics, and statistics. Typical here is the comment made by a reviewer for *Nature* in 2002:

If such gene flow is as frequent as is claimed by the authors ... it is necessary that the experimentation and analysis is absolutely watertight [...] My advice to the authors in the first instance is that they focus on fewer plants than in the present study and carry out the DNA and other analyses in much more detail than at present. (*Nature* Reviewer 1 of Alvarez-Buylla et al., 's 2002 submitted manuscript)

Such understanding of what constitutes a 'watertight' proof illustrates molecular biology's particular approach to making sense of and ordering the world: not using large data sets from carefully sampled materials in statistical analyses, nor modeling ecological or social complexity, but rather using sophisticated laboratory DNA tools to track, isolate, and follow one (or a few) completely abstracted genes or gene-products at a time, in controlled environments, while working with artificially standardized living forms (Bonneuil, 2006; Knorr-Cetina, 1999). While the epistemic objects of anthropologically informed population geneticists are genes as collectively circulating entities in socio-agro-ecosystems

across large time and space ranges, molecular biology's epistemic objects are genes as manipulatable entities in an experimental system nested in the micro-worlds of the controlled laboratory. The production of robust knowledge in molecular biology's epistemic culture thus requires that genes be abstracted/extracted from their cultural/ecological entanglements.

The extraction/purification approach of the epistemic culture of molecular biology produces ignorance (non-knowledge) simultaneously with knowledge. Moreover, ignorance coincides with and co-produces a denial of the value or relevance of indigenous maize culture, including its human and non-human participants. This example illustrates Proctor's model of ignorance-making through framing – the active erasure of phenomena treated as out of scope, or as disorderly noise threatening the cognitive order and security of the dominant epistemic culture (Proctor and Schiebinger, 2008). This entrenched epistemic culture transforms gene flow from a complex, fluid, and distributed bio-cultural process, that can be acted upon and spoken for by a variety of actors, into a laboratory object that only some particular scientists can speak for in selected authoritative journals. It reduces socio-natures to *what biotechnology* (and its molecular-commercial culture) *can see*, to follow Scott's (1998) analysis. We return to this visibility question in the conclusions.

Making genomes legible: imperial molecular metrologies

The reductionist erasure of the cultural and ecological dimensions of maize genetic landscapes – together with corresponding types of identity and power distribution – cannot deliver on its promise to produce the type of knowledge considered most relevant without an additional implicit selective commitment. This particular assumption, of the homogeneity of maize landraces, in the image of commercial hybrid maize seeds, becomes normalized and performed in practice as if natural. The norm enacted is that only a few genetically homogeneous, standard, and tractable maize varieties ought to be used as research or calibration objects, even if a wide variety of genetically heterogeneous and complex varieties constitute Mexico's actual maize-genomic landscape. This section analyzes this standardizing molecular metrology, which carries with it a politics that values homogeneity – a product of the co-evolution of industrial agriculture and modern genetics (Bonneuil, in press) – and which is epistemically and politically blind to the heterogeneity of the bio-cultural world of Mexican landraces.

Legible varieties

When Quist and Chapela sent their first manuscript to *Nature*, one of the reviewers was not 'convinced that this work is strong enough for *Nature*' (Referee 2, May 2001, personal communication from David Quist) and questioned its methodology, especially the choice of a negative control as yardstick for comparison with positive samples:

My main problem with the science is in the choice of control. Surely the appropriate control would be samples of seed from the native plant that were stored before GM maize was grown in Mexico. (Referee 2, May 2001, personal communication from Quist)

This requirement assumed that before the putative arrival of transgenes in Oaxaca communities, there was one ‘native plant’, which could act as the control – a single cultivar, or genotype, homogeneously spread in the whole area. Familiar with the sheer diversity of Oaxacan maize culture, the authors replied that

It is difficult, however, to know what [the referee] suggests by ‘native plant’... The level of genetic heterogeneity in maize landraces... make such resolve equivocal at best, and usually impossible. (Quist and Chapela to *Nature*, 14 May 2001, personal communication)

Regretting that ‘the authors have made no attempts to address my fundamental criticism [concerning] the appropriate control’, the referee advised rejection of the revised manuscript: ‘I just don’t think the paper is anywhere near interesting or careful enough to merit publication’ (*Nature* to Quist, 14 July 2001, personal communication from Quist)

Derived from the comparison of near-isogenic lines – a standard norm of proof among plant biotechnologists – this requirement from the reviewer implies the existence of a single homogeneous cultivar as a necessary control. In this particular epistemic culture, the production of exact knowledge implies ontological closure around a single ‘native plant’ that can stand for the ‘before’. This unrealistic laboratory-molecular imaginary has its roots in the 20th century’s genetic modernism (Bonneuil, in press; Fox-Keller, 2000), which valued genetically uniform life forms (e.g. clones, pure lines, and F1 hybrids) and conceptualized biodiversity as stocks of genes rather than as ongoing flows (cf. the static notion of ‘genetic resources’). As a result, the possibility of ambiguous intermediary forms or sur-reptitious circulations between the ‘modern’ and the ‘traditional’ is discarded.

The reviewer’s methodological requirement reflects an apparently unwitting normalization of industrialized and genetically homogeneous agricultural crops. The assumption of homogeneity is performed, not tested. It is imposed on Mexican landscapes and cultures and on competing knowledge systems that are more respectful of those cultures’ non-compliant complexities. This reflects an imaginary of a legible order of selective homogeneous genomes, one that had become the dominant material reality of agricultural research stations, biotech laboratories, industrial agriculture, and Green Revolution landscapes. This high modernist way of constituting cultivars and their genomes also reflected a political project to discipline agriculturally and biologically diverse cultures and to govern rural societies. It is much harder for a state or for a global seed company to govern an informal, distributed, heterogeneously complex bio-social seed nexus than it is to govern a formal, streamlined, standardized, literally reduced – in terms of genetic, agronomic, and cultural complexity – top-down system. Our analysis of molecular biology’s modernist standardizing gaze extends Scott’s (1998) concept of legibility. Scott showed that in order to ‘improve’ and maintain control over rural socio-environmental complexes, modernist states, agribusiness companies, and their scientific agents had to render them legible, homogenized so as to become commensurable with a centralized mode of knowing and an industrial mode of production. As we demonstrate next, this control strategy has deep molecular epistemic dimensions.

Calibrating genomes

Together with the ‘choice’ of controls, the issue of calibration of PCR detection assays played a key role in the controversy, while also embodying far-reaching cultural assumptions, material conditions, and power relations. GID was founded by US molecular biologists in 1996. It soon became the world’s biggest commercial GMO testing company. From 2001, GID led a campaign against immunological tests, which was launched with a study that undermined strip tests commercialized by a competitor (Fagan et al., 2001). Although this work was much criticized, once again for lack of an appropriate sampling method (Lezaun, 2003), GID succeeded in imposing PCR as the method of choice. GID promoted qPCR or real-time PCR (rtPCR) (developed in 1999) as the best ways to detect GMOs. rtPCR is run in a closed system, less prone to contamination, and hence potentially less prone to false-positives. Another advantage for commercial development is rtPCR’s amenability to automation and routinized technician labor. But it also has some limits: to get proper data and to avoid false-negatives it requires demanding preliminary calibration work (namely, choice of primer, reference gene, etc.) and hence relies upon particular assumptions about homogeneity and regularity of seed sample DNA. To allow quantification, it also has to make assumptions about DNA amplification rates with different materials and conditions.

In 2002, after *Nature* rejected the manuscript that confirmed Quist and Chapela’s findings and reported many positives in two states of Mexico, one of the junior co-authors, Sol Ortiz-García from INE, sent samples to GID. As mentioned above, GID’s report emphasized many inconsistencies in the results, obtained with qualitative and quantitative PCR (GID Report, December 2002, personal communication by S. Ortiz-García). While discreetly working hard in-house to tame these inconsistencies through further calibration and optimization of their rtPCR technique, rather than asking about what possible biological unknowns might have generated them (affirming Knorr-Cetina’s (1999: 110) account of molecular biology epistemic culture’s habit of ‘semi-blind variation’, GID scientists told Ortiz-García that these strange results could only come from contamination in the public research laboratories that had prepared the samples (i.e. UNAM and Cinvestav’s laboratories). Ortiz-García and INE officials gradually became convinced that only GID would provide reliable data for publication. They therefore adopted GID’s assumption that Mexican maize genomes should behave like those to which GID was accustomed – typically big, homogeneous, easily sampled commercial shipments of modern hybrid seeds – akin to those assumed by the *Nature* referee dismissing Quist and Chapela, described in the preceding section. INE stopped collaboration with UNAM and Cinvestav, thus dropping Ortiz-García’s co-authorship of the positive transgene findings manuscript. Unannounced to those colleagues, the Ortiz-García INE group instead worked with GID to prepare the 2005 *PNAS* publication, which, on the basis of GID’s PCR analysis, reported no transgenes in all samples collected in Oaxaca in 2003 and 2004. This publication omitted mention of the many surveys that reported positives in several Mexican states in 2001, 2002, and 2003 samplings, including INE surveys. All these positives were retrospectively recast as false-positives, in light of the apparent DNA contamination in public laboratories not specialized in the business of molecular detection. This was although some of those positive findings resulted from

non-DNA methods! The year following the publication of the *PNAS* article, in 2006, the Mexican Government bought the license for GID's proprietary testing technology, which is now used throughout the world by its Global Laboratory Alliance® members, including a dozen governments. So, the issue of which kind of methodology is relevant for producing evidence in detection controversies is not only about the sensitivity, stability, or robustness of particular tools but, ultimately, about *who* gets the business to monitor the global circulation of standardized commodities.

Piñeyro-Nelson et al.'s (2009b) publication, authored by, among others, scientists from the two laboratories that GID accused of 'contamination' a few years earlier, challenged the kind of legibility at work in GID's detection methodologies. In particular, they underlined that the molecular culture's blindness to research objects other than globally traded modern hybrid genomes exposed GID's PCR methodology (and hence Ortiz-García et al.'s (2005) *PNAS* article) as prone to false-negatives:

while Genetic ID's method may be well suited for commercial use on commercial crop varieties in the U.S., they ... [are] working with the limited genetics of commercial hybrid maize... built upon a very homogeneous genetic background, a cross of Northern Flint x Southern Dent. Both inbreds ... are quite different from Mexican maize landraces. (Piñeyro-Nelson et al., 2009a: 4146)

At stake here is the tension between current commercial quantitative PCR technologies and maize diversity. Maize-genomics research has shown that Mexican maize landrace genomes are about 10 percent bigger and more variable in DNA sequence terms than laboratory pure lines and commercial hybrids, and they show a surprisingly limited level of collinearity with US cultivars. 'Collinearity' refers to the way genes are ordered along the chromosomes (Walbot, 2008). Due to more active transposons, gypsies, and mutational locus variability, this higher genetic heterogeneity and fluidity in maize landraces point to many plausible inhibition mechanisms of PCR amplification by various compounds present in landraces (e.g. secondary metabolites, oils, and polysaccharides). These compounds need to be understood and controlled, at least analytically, if not practically, for their variable effects, rather than discarded as 'strange', as 'false-positives', or as (incompetent) 'contamination'.

Still a contentious issue in 2001, when Quist and Chapela suggested it, it is now established that GM crop genomes undergo uncontrolled post-release 'event-specific variations' – changes in the location and sequence of the transferred sequence and its flanking sequences themselves (e.g. recombination, mutation, rearrangements due to transposons) (Matsuoka et al., 2002). For instance, Piñeyro-Nelson et al. (2009a) reported a case in which GID produced a false-negative finding by failing to detect transgenes in a well-known NK603 cultivar, which they had submitted to GID as a 'blind' test. This could be attributed to inefficient primer binding, due to post-release, event-specific variation in that cultivar (p. 4148). In qPCR, the amplification of the target sequence (a sub-sequence of the transgenic construct) is compared to the amplification of a control sequence, or 'reference gene', which is assumed to behave similarly in all maize genomes. Commercial rtPCR utilizes only one or a few reference gene(s), such as the zein gene. But variability in these control sequences has been reported, and these differences lead to differential amplification. This messiness related to internal-standard variability may produce the kind of

inconsistent results that are then scored negative in commercial testing methodologies, thus leaving additional room for false-negatives (Piñeyro-Nelson et al., 2009a: Figure 1).

In summary, current DNA-based transgene detection methodologies, which are sold and defined as the gold standard worldwide, carry an implicit assumption of universality and homogeneity of maize genomes, as if modern industrial agriculture hybrid maize was the universal norm. This is simply incompatible with Mexican maize-genomic complexity and diversity. In this way, hard DNA-based proofs and commercial PCR methodologies enact a cultural domination of the world of industrial hybrids over the bio-cultural world of indigenous maize. But maize genomes are not made of discrete bounded genes, with similar sequences that sit in an orderly manner like beads on the DNA strand. Typical of Latour's recalcitrant objects, they resist such metrological alignment by the biomolecular genetic reductionism that co-evolved with commercial global industrial monocrop agriculture. These remaining bio-social recalcitrants, ambiguities, and variability, which exist at the periphery of contemporary bio-empire (Jasanoff, 2006), suggest a path to alternative modernities – made of what could be reclaimed as 'other' bio-cultural entanglements, knowledge, and identities (Foyer, 2010).

Conclusion

We can now address the question raised in the 'Introduction': why did it take a decade to ascertain that transgenes have escaped from US GM corn industrial hybrids into local maize landraces in a global center of maize diversity? First, in Mexico and worldwide, many scientific and policy actors were happy to delegate to high-profile scientific journals the function of gate-keeping what was to be defined as scientifically sound, and hence policy relevant and authoritative, among the wide range of detection data, assumptions, and claims advanced by a diversity of actors and knowledge cultures. Requiring peer-reviewed DNA and PCR-based proof proved a politically powerful way to contain the larger controversy sparked in 2001 within an esoteric, controlled, and apparently purely technical question of scientific methodology.

In the elite arenas and journals that were given authority to channel and arbitrate the controversy, only one out of four knowledge practices (namely, PCR) was considered reliable to make transgenes visible. This exclusion of other forms of expertise marginalized the voices of related social groups, including not only indigenous and socio-environmental organizations but also disciplinary agronomists, population biologists, and anthropologists who had joined to investigate the entangled biological and cultural processes that produce or harm crop genetic diversity. As such, the controversy over Mexican GM maize appears as a classic instance of imperialist subjugation of subaltern knowledge cultures through the restrictive discourse of 'science' in general and the monopoly form of molecular biology DNA detection in particular.

Rural communities, and the scientists who took them seriously, were not given a voice in high-profile scientific institutions and journals or in policy circles. Instead, the framings and norms of proof of only one particular epistemic culture, molecular biology, were made to stand hegemonically for sound science. Moreover, in the later exchanges of this controversy, a particular, globally influential molecular biology perspective, seen clearly in GID's attacks on opposing analyses, incorporated normative assumptions.

Alternative approaches recognized other significantly different ‘normalities’, such as factors highly relevant for Mexican maize agriculture, but also implied different normative standards for molecular biological practices. PCR-based molecular biology has a blindness to sampling issues; to the dynamic, bio-cultural, and extensive web-like functioning of genetic landscapes; and to the fluidity and diversity of maize genomes. Thus, a singular reliance on PCR-based approaches failed to order the messiness of Mexican maize gene landscapes into unequivocally robust tests. Yet, at the same time, this hegemonic culture – which linked laboratory norms to journal and peer-review normative standards and to media and political networks – erased from recognition and visibility some non-compliant forms of life and knowledge that may have helped understand gene flows, in all their exuberantly emergent, lively bio-social complexity.

As the inventor of PCR himself noted, ‘the remarkable part is that you will pull out a little piece of DNA from its context’ (Mullis quoted in Rabinow, 1996: 6). The narrowing of the understanding of gene flow in Mexican agricultural landscapes and in the GM debate to a focus on ascertaining the presence/absence of transgenes as ‘matter out of place’ reduced the scientific and policy gaze around a particular vision of the gene and of the world. It imposed a view of cultivars and (trans)genes as self-sufficient, non-relational, stable, and standardized units, as if their mode of existence and impact could be abstracted from their ecological and social bonds. Such a framing embodies and reinforces a cultural domination by the standardized (first) world of industrial maize – with its standardized genomes enabling patent claims, commodity circulation, and centralized governing – over the complex, heterogeneous, and web-like bio-cultural world of indigenous landraces and their peoples (see Figure 1).

The controversy over the detection of transgenes in Mexican maize landraces is not merely technical or even sociotechnical. This case is a political and cultural struggle, a new episode of a contentious cultural encounter between worldviews, values, forms of knowledge, and forms of life that started with the Green Revolution. The molecular techniques and norms of proof that we have followed in Mexico, and from there around the globe, are key instruments in this cultural confrontation. As Sheila Jasanoff (2006) has shown, biotechnology holds much of its imperial power from particular metrologies, at work in risk assessment framings, standards and guidelines, databases, gene banks, patent offices, detection methods, and so on. These metrologies include (1) an ontological dimension, seen in the ‘genetic resources’ modernist paradigm and the denial of the bio-cultural nature and existence of gene flow; (2) an epistemic dimension, seen in a hegemonic domination of molecular biology’s epistemic culture over a bio-cultural epistemic culture; and (3) a socio-ecological dimension, such as the need to partially align the conditions of the real world and the prevailing epistemic culture so that the products from the biotech firms and detection laboratories may be said to work. These three imperial metrological standardizations commit to the making of a world less livable for some forms of life than for others.

In the Green Revolution era, the cultivar as a whole was the unit that polarized most efforts to produce exact science (controlled experiment with homogeneous ‘material’), productive inputs (elite cultivars), and ownable and tradable commodities (‘Distinct Uniform and Stable’ cultivars). With the ‘biotech revolution’, the gene and its practical manipulation into specific transgenic constructs have become the unit of scientific intervention, ownership, trade, and endless ‘improvement’. Here, the standardization of GMO

detection techniques has become a major, contested stake for the control of the global flow of patented transgenes in world markets and agroecosystems. The global biotech enterprise keeps grinding away in the laboratories of power, externalizing and denying recalcitrant, non-compliant Mexican actors and processes and reducing global historical and political struggles to 'respect for sound science'. Meanwhile, the more modest agents of a non-imperialist world struggle to maintain their very existence within a more accommodating ontology, a more plural world that could allow multiple nature cultures to thrive.

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