Tetty Havinga

Conceptualizing Regulatory Arrangements: complex networks of actors and regulatory roles
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CONCEPTUALIZING REGULATORY ARRANGEMENTS: COMPLEX NETWORKS OF ACTORS AND REGULATORY ROLES

Tetty Havinga*

Abstract
Food regulation nowadays involves a broad variety of actors, including government agencies as well as other stakeholders. In this chapter I explore which actors are involved in private and mixed forms of food regulation. This question is important when considering such issues as the effectiveness, legitimacy and accountability of regulatory regimes. I argue that a dichotomous distinction between public (governmental) and private (non-governmental) regulation is not an adequate conceptualisation for analysing the reconfiguration of relationships between the actors involved in food regulation. We need more sophisticated distinctions linking the type of actor to the role they play in the regulatory process. I disentangle the regulatory process into multiple regulatory roles/actions in order to analyse complex patterns of actor involvement in a regulatory regime.

Key words

Introduction
The regulation and governance of food has changed dramatically in recent decades. Traditional food regulation consisted of national governments enacting food legislation and enforcing compliance with the laws. Several developments in society have contributed to the emergence of national and transnational private or mixed forms of food regulation.

Food supply chains are increasingly becoming international. Fresh products are sourced all over the world to assure Western consumers a continuous stream of fruits and vegetables throughout the year. Many processed food products are made up of a variety of ingredients, drawn from all over the world. Regulating complex global food supply chains is difficult for national governments since their jurisdiction is limited to their own national territory.

Food safety issues has been transformed from a predominantly neutral, technical issue that is the preserve of food experts into a contested political

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matter that is debated in parliament, on television and in the newspapers. Food
scares and incidents such as mad cow disease (BSE), contaminated milk, and
EHEC have received wide media attention. The risks connected with food have
become increasingly visible.

Traditional government food regulation was perceived as no longer ade-
quate, similarly to government regulation in other domains. Traditional regula-
tion is believed to be ineffective and inflexible, while it disregards the respon-
sibilities of citizens and organizations. The way both the UK government and
the EU handled the BSE crisis is often cited as a turning point, which demon-
strated an urgent need to transform food regulation.

These social developments resulted in a shift towards private and mixed
forms of regulation and a shift from national government regulation towards
European Union regulation. In an attempt to restore consumer confidence and
to retain export markets, governments in Brussels and the EU member states
enacted stricter food safety regulations and reorganized the government food
safety system. Food producers and suppliers expressly bore primary responsi-

bility for food safety while national governments became responsible for con-
trolling the adequacy of risk control mechanisms of companies in a food chain.
As a result of their legal responsibility and from a fear of potential damage to
their reputation in case unsafe food products should find their way to market,
the food industry and food retailers developed initiatives to decrease food
safety risks and increase consumer confidence in safe food.

These reforms involved the emergence of new forms of food regulation, in-
cluding transnational private food standards, corporate social responsibility
initiatives, and codes of conduct. These transformations involve a new relation-
ship and a changed distribution of responsibilities between government bodies
on the one side and private actors on the other. New forms of food regulation
are characterised by a less dominant role for the government and more res-
ponsibilities for private actors (Havinga 2006, Henson and Humphrey 2011,
Marsden et al. 2010, Oosterveer 2005). These new forms of food regulation
include both public actors and also private actors, such as firms, NGOs and
other organisations both inside and outside the food production chain. Govern-
ment regulation is moving from a prescriptive, command-and-control system
towards an enforced self-regulatory approach (Braithwaite 1982, Hutter
2011, Martinez et al. 2007) in which the government lays down broad stan-
dards and leaves it to companies to develop, implement and monitor risk man-
agement systems. An example is the European Food Law, which requires all
food businesses to have controls that demonstrate that they manage food
safety risks.

Recent trends are the emergence of retailer-led food governance and
global coalitions for setting food safety standards, an increased use of global
business-to-business standards and third party certification (Fulponi 2006, Harrison 1997, Hatanaki and Busch 2008, Havinga 2006, Marsden et al. 2000, 2010). Another trend is the increase of consumer hallmarks for ‘new’ issues, such as animal welfare, fair trade, sustainability, healthy food, and halal food.

Marsden et al. (2010) describe the new situation as the Public-Private Model of Food Regulation.

**Various forms of food regulation**

Regulation is a means for controlling harm in the public interest in order to protect consumers, citizens and the environment (Hutter 2011: 10). Henson and Caswell (1999) distinguish between direct and indirect regulation of food safety (see figure 1).

**Figure 1: Types of regulation of food safety**

<table>
<thead>
<tr>
<th>Source of the rules</th>
<th>Character of the rules</th>
</tr>
</thead>
<tbody>
<tr>
<td>Direct</td>
<td>Indirect</td>
</tr>
<tr>
<td>Public</td>
<td></td>
</tr>
<tr>
<td>Dutch Commodities Act (Warenwet)¹</td>
<td>Product liability law²</td>
</tr>
<tr>
<td>UK 1990 Food Safety Act</td>
<td>TBT-GATT agreement³</td>
</tr>
<tr>
<td>EU General Food Law</td>
<td></td>
</tr>
<tr>
<td>Hybrid Public-private</td>
<td></td>
</tr>
<tr>
<td>Industrial code of hygienic practice</td>
<td>Liability insurance policy</td>
</tr>
<tr>
<td>Private</td>
<td></td>
</tr>
<tr>
<td>Private food safety certification scheme (GlobalGap, MSC)</td>
<td>Food company complaints procedure</td>
</tr>
</tbody>
</table>

Direct regulation entails obligatory prescriptions and requirements for the production and handling of food to assure the production of safe food. Even though indirect regulation does not provide prescriptions for the production and the product, it is nevertheless expected to act as an incentive to implement food safety controls indirectly by shifting the cost-benefit balance or imposing process requirements. For example, liability law is assumed to promote food safety by encouraging producers to do their best to produce safe products (Havinga

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³ Art. 185-193 bk 6 BW.
⁴ GATT Agreement on Technical Barriers to Trade (TBT).
A food company’s consumer complaint procedure is an example of private, indirect regulation because the expectation is that learning from complaints will increase the product’s quality. Direct regulation is intended to provide food safety whereas indirect regulation may aim at increasing food safety; however, a contribution to food safety may also be an unintended de facto effect.

Until recently, most research on food safety has focused on direct forms of food safety regulation. Recently, however, not only public regulatory arrangements but also private forms of direct food safety regulation have started to attract attention (Fuchs et al 2011, Fulponi 2006, Havinga 2006, Henson and Reardon 2005, Marsden et al. 2000, 2010, Martinez et al. 2007, Rothstein 2005, Van Waarden 2006).

This paper deals with two questions in the context of the reconfiguration of the relationships between the various actors: which actors are involved in the new food regulatory arrangements? And what roles do these actors play?

Public and private actors in regulatory arrangements

Some authors only distinguish between public and private actors (e.g. Spruyt 2001; Josling, Roberts and Orden 2004: 156), state and non-state actors or government and non-government actors. Daniel Stewart (2010) implicitly distinguishes state actors from non-state actors in his analysis of the availability of judicial review to actions of non-state actors. Bingen and Busch (2006: 246) observe ‘a constantly evolving and changing global network of relationships among public and private, mandatory, voluntary, and national and international standards bodies instead of neatly delineated public and private responsibilities’.

Several authors on environmental regulation and sustainability have replaced this public-private dichotomy by the following threefold classification: state, market and civil society. In their analysis of non-state, market-driven governance systems, Bernstein and Cashore (2007: 356) and Abbott and Snidal (2009a, 2009b) distinguish between two types of non-state actors: firms and NGOs. A similar threefold classification is made by Levi-Faur (2010), who distinguishes between civil, market, and state actors. For Levi-Faur a non-governmental organization can be controlled by civil society actors (CiNGO), market actors (MaNGO), or state actors (GoNGO). ‘This distinction between different types of NGOs which act as regulators will allow us to develop a clearer understanding of hybrid designs of regulatory institutions.’ Because of the ambivalent definition of NGO I prefer to speak of civil actors (alongside government or state actors and firms or market actors).
Figure 2: Threefold classifications of actors in regulation

<table>
<thead>
<tr>
<th>Abbott &amp; Snidal 2009</th>
<th>Levi-Faur 2010</th>
<th>Marsden et al 2010</th>
</tr>
</thead>
<tbody>
<tr>
<td>States</td>
<td>State actors</td>
<td>Policy &amp; regulatory interests</td>
</tr>
<tr>
<td>Firms</td>
<td>Market actors</td>
<td>Private interests</td>
</tr>
<tr>
<td>NGOs</td>
<td>Civil actors</td>
<td>Consumer &amp; social interests</td>
</tr>
</tbody>
</table>

The Governance Triangle

Abbott and Snidal (2009a) visualize the variety of regulatory standard-setting (RSS) in their Governance Triangle (see figure 3). They distinguish three groups of actors directly involved in RSS: states, firms and NGOs. Abbott and Snidal consider RSS rules to be a form of regulation, even though they are voluntary (that is not legally mandatory).

Figure 3: The governance triangle (Abbott and Snidal 2009a, 50)

The seven zones in the triangle indicate the three forms of single-actor standards (1-3), three forms of dual-actor standards (4-6) and one form involving direct participation of all three groups of actors (7). Abbott and Snidal included only four regulatory food standards in their governance triangle5; in

5 Marine Stewardship Council (MSC) 1997, Max Havelaar (MH) 1988 (fair trade coffee) and International Federation of Organic Agriculture Movements (IFOAM) 1972 (organic →
another publication they include another two food standards\textsuperscript{6}. Further food standards could easily be included. Most food regulatory arrangements are situated in zone 1 (state regulation) and zone 2 (industrial self-regulation).

Abbott and Snidal show the evolution of the governance triangle (see figure 4). In the pre-1985 period there are only a few RSS schemes, mostly in Zone 1 (state). The decade between 1985 and 1994 shows the emergence of RSS schemes, especially firm-schemes, as well as the first multi-stakeholder and NGO schemes. The post-1994 period shows a continued proliferation of firm schemes and an increasing number of NGO schemes as well as the emergence of collaborative schemes (NGOs and firms or tripartite, zone 6 and 7).

\textit{Figure 4: Evolution of the governance triangle over time}

\begin{figure}[h]
\centering
\includegraphics[width=0.8\textwidth]{governance_triangle.png}
\caption{Evolution of the governance triangle over time}
\end{figure}

The governance triangle is useful for analyzing changes in regulatory standards over time or to compare regulatory arrangements between different sectors or countries. Food regulatory schemes are dominated by state regulation (zone 1) and by firm schemes (zone 2). This applies particularly to food safety standards. The food standards that involve civil actors are mainly concerned with other issues such as sustainability or fair trade, but it is quite possible that civil actors will become more important in food regulation in the future. We can already witness an increase in the numbers of these actors in the regulatory

\footnote{food), all three in zone 6 (dual-actor standards with involvement of firms and NGOs) and WHO Code of Marketing for Breast-milk Substitutes (BM) 1981 in zone 1 (state standard).}

\footnote{GlobalGap 1997 and SQF 1994 (SafeQualityFood), Abbott and Snidal 2009b: 513, both in zone 2.}
domain, devoted to social interests (e.g. fair trade), environmental interests (e.g. organic food), or animal welfare (e.g. free-range meat). It remains to be seen whether future regulation in this field will develop in the way Abbott and Snidal sketch out.

However, an analysis focused on actor types also has its drawbacks. Let us look at the example of GlobalGAP (Global Partnership for Good Agricultural Practices). GlobalGAP is a private sector body originally set up by European supermarkets chain to set standards for the certification of agricultural products around the globe. GlobalGap is classified as a single-actor standard in the Governance triangle (Zone 2). In fact, however, GlobalGap is not a single-actor standard, but a standard set by actors from one single category of actors (firms). Multiple actors are involved in GlobalGap, all with very different positions and interests. GlobalGap is a membership organization with three different types of members: retail and food service members, producer/supplier members and associate members such as certification bodies, consultancies and the crop protection industry. Conflicts of interest exist within these member categories, for example between multinational supermarket corporations and small local retailers or between aquaculture industries in Norway and Thailand. Moreover, other stakeholders such as environmental NGOs, scientific experts or government officials are also involved in working groups or meetings of GlobalGap.

The focus on actor types in the regulatory triangle neglects that a ‘single-actor standard’ might involve multiple actors with conflicting interests. Furthermore, retail food standards (e.g. the British Retail Consortium Global Standards), halal certification (e.g. the quality system of the Halal Correct Certification Foundation) and traditional industrial self-regulation (e.g. the South Africa Olive Commitment to Compliance scheme or the Unilever Supplier Qualification System) are Zone 2 standards in the governance triangle, just like GlobalGap, although they differ significantly in terms of the actors involved. We should therefore differentiate between various actors.

Authors on food regulation tend to focus less on NGOs or civil society than do authors on environmental regulation, probably because civil society actors do not play a significant role in many food regulatory arrangements. Because of this, I neglected to include civil society organisations in an analysis of shifts in food safety regulation (Havinga 2006). However, in the field of food safety it is particularly important to distinguish between different types of private actors, so I distinguished three important institutional actors in this field: state (government agencies), industry (food industry and farmers) and third parties.8

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7 See footnote 6 for some examples of zone 6 standards.
8 This concept of ‘third party’ differs from the concept of Levi-Faur.
Havinga: Conceptualizing Regulatory Arrangements

(private auditing and certification organisations, retailers and consumer organisations) (Havinga 2006). Fuchs et al. (2011, 360) stress the importance of retailers in food regulation when they distinguish four types of stakeholders in private retail food standards: 1) retailers, 2) producers: food industry, growers, fishers, 3) certification bodies and 4) civil society and NGOs.

Dilling et al. (2008: 3-4) distinguish three actor constellations from which the para-legal systems of the business world emerge: single enterprises (corporate self-responsibility), transnational corporate networks, and non-governmental organisation-business partnerships.

**Analysing networks and production chains**

In another body of literature, authors on food regulation do analyse the networks or production chains connected to a particularly regulatory regime.

Marsden et al. (2010: 120) distinguish between three types of interest in the new hybrid food policy-formation network:

1. policy and regulatory interests,
2. private interests, and
3. consumer and social interests.

These types of interest correspond to the three actor categories in the regulatory triangle proposed by Abbott and Snidal (state, firms, NGOs, see figure 2). Marsden et al. (2010: 283, italics in original, 290) argue that ‘it is the interactions between public and private actors that are essential to a fuller understanding of the food system’ and the boundaries between public, private and civil society interests become less clear with time. Other authors also observe the blurring of the public-private distinction and the rise of hybrid organisations and networks combining government and non-governmental actors (e.g. Bingen and Busch 2006, Black 2002, Havinga 2006, Hutter 2011, Levi-Faur 2010).

In their analysis of the new regulation and governance of food, Marsden et al. pay attention to a variety of actors such as UK and EU government bodies, corporate retailers, farmers, producers, manufacturers, global standard-setting organisations, private certification bodies, logistic firms, environmental and social NGOs. They paint a picture of various complex networks involving many organisations and regulations (see figure 5, for example).
Figure 5 Regulation in the supply chain and the role of different global organizations


Bush and Oosterveer (2007: 391) describe export firms as key actors in the shrimp trade, linking local producers in Thailand and Vietnam with global trade networks. They conclude that certification bodies are unable to audit the chain below the level of exporters, where capital and information flow through informal, diffuse trade networks. This governance arrangement involves a complex mix of state and non-state actors: bureaucrats, retailers, wholesalers, im-

9 BRC (British Retail Consortium), EurepGap (Euro-Retailer Produce Working Group Good Agricultural Practice), FAO (Food and Agriculture Organisation), FSA (Food Standards Agency), HACCP (Hazard Analysis and Critical Point), IFS (International Food Standards), IPPC (International Plant Protection Convention), OIE (Organisation Mondiale de la Santé Animale/ World Organisation for Animal Health), SPS (Sanitary and Phytosanitary Measures), SQF (Safe Quality Food), WHO (World Health Organisation), WTO (World Trade Organisation).
porters and exporters, local trade network, producers, NGOs, processing companies and consumers.

In a study of public-private protection of forests and fish (Forest Stewardship Council and Marine Stewardship Council), Van Waarden (2010) points to the involvement of a wide range of actors from all phases in the value chain, with different interests, as one of the factors contributing to the success of these schemes: small scale and large fisheries, fishery associations and cooperatives, suppliers, manufacturers, distribution, retailers, scientists, indigenous people's organisations, fishery communities, and environmental groups.

To conclude, the broad threefold categorisation of actors involved in regulation (state actors, firms and civil society) is useful in analysing the balance between these three actor types in comparative perspective, whether historical, as Abbot and Snidal do, or when comparing countries, industries or fields of regulation. However, a more sophisticated categorisation of firms is needed for analysing the involvement of actors in non-governmental and hybrid systems of food regulation. Consider three private food regulatory regimes: GlobalGap, BRC, and a Dutch Halal certification scheme. All three would be classified as firm schemes in zone 2 according to Abbott and Snidal. Yet the three are only similar in some ways. When analysing the effectiveness and legitimacy of a regulatory regime it is important to distinguish between non-governmental actors which are regulated (regulatees), non-governmental actors which are part of the production chain but are not regulated themselves by the regulation under consideration (such as suppliers and retailers), and non-governmental actors providing services to the regulated industry (such as certification and auditing organisations). The increasing role of transnational and international government bodies in food regulation means it is inevitable also to distinguish between different levels of state actors. Following Marsden et al. (2010), civil society interests can be broken down into social interests (e.g. child labour, fair trade, occupational health, public health), environmental interests (e.g. biodiversity, sustainability, insecticides) and consumer interests (e.g. food safety, food security, choice and price). Figure 6 provides a survey of these distinctions.

10 Havinga 2008.
11 Animal rights groups do not fit into this categorization very well.
**Figure 6 Types of actors involved in regulation and their position in the supply chain**

<table>
<thead>
<tr>
<th>Type of actor</th>
<th>Position in regulation/supply chain</th>
<th>Actor role</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>State</strong></td>
<td>- transnational level</td>
<td></td>
</tr>
<tr>
<td></td>
<td>- (federal) level</td>
<td></td>
</tr>
<tr>
<td></td>
<td>- state level</td>
<td></td>
</tr>
<tr>
<td></td>
<td>- local level</td>
<td></td>
</tr>
<tr>
<td><strong>Non-state/ private</strong></td>
<td>Mar-</td>
<td>Inside regulated supply chain</td>
</tr>
<tr>
<td></td>
<td>ket/firms</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>- Associations of regulated industry</td>
</tr>
<tr>
<td></td>
<td></td>
<td>- Firms up- or downstream in supply chain</td>
</tr>
<tr>
<td></td>
<td></td>
<td>not target of regulation at hand</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>Outside regulated supply chain</td>
<td>- Providing services (e.g. standard setting,</td>
</tr>
<tr>
<td></td>
<td></td>
<td>auditing, certification, disinfectants,</td>
</tr>
<tr>
<td></td>
<td></td>
<td>education, consultancy)</td>
</tr>
<tr>
<td><strong>Civil society (NGOs)</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>- Environmental interest groups</td>
</tr>
<tr>
<td></td>
<td></td>
<td>- Consumer interest groups</td>
</tr>
</tbody>
</table>

**Regulatory roles**

Levi-Faur (2010) distinguishes further between three roles: regulator, regulatee, and third party\textsuperscript{12}. Combining these three roles with three actor types, he distinguishes 27 types of third party regulatory designs (see figure 7).

**Figure 7** Types of Third party regulatory designs (Levi-Faur)

<table>
<thead>
<tr>
<th>Type of Third Party Enlisted</th>
<th>State Actor as Regulatores (e.g. regulatory agencies; GoNGOs)</th>
<th>Market Actor as Regulator (e.g. Mangos)</th>
<th>Civil Actor as Regulators (e.g., CINGOs)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>S      M      C</td>
<td>S      M      C</td>
<td>S      M      C</td>
</tr>
<tr>
<td>State Actor as regulatees</td>
<td>SSS    SMS    SCS</td>
<td>MS      MMS      MCS</td>
<td>CSS    CMS    CCS</td>
</tr>
<tr>
<td>Market Actor as Regulates</td>
<td>SSM    SMM    SCM</td>
<td>MSM      MMM      MCM</td>
<td>CSM    CMM    CCM</td>
</tr>
<tr>
<td>Civil Actors as Regulates</td>
<td>SSC    SMC    SCC</td>
<td>MSC      MMC      MCC</td>
<td>CSC    CMC    CCC</td>
</tr>
</tbody>
</table>

**Triplets Key:** First letter means the Regulator; Second represents the enlisted third party; Third letter the regulatee; S=State; M=Market; C=Civil


Mattli and Woods (2009) include implementation in their definition of regulation: ‘the organisation and control of economic, political, and social activities by means of making, implementing, monitoring, and enforcing of rules.’ Adding implementation implies that we should consider the regulatees as part of the regulatory regime.

\textsuperscript{12} Levi-Faur 2010 distinguishes three main strategies in regulation: 1. first party regulation: Regulator=regulatee (self-regulation), 2. second party regulation: regulator independent of regulatee (e.g. state regulation of business, or retailer imposing regulation on suppliers), 3. third party regulation – relationship between regulator and regulatee is mediated by a third party as independent auditor (e.g. EurepGAP).

The meaning of the concepts ‘first party’, ‘second party’ and ‘third party’ is not consistently used in the literature. E.g. Van Waarden 2012: first party = self-regulation by actor, second party = actor involved in transaction imposing standards, third party = independent actor not involved in transaction; Bingen and Busch 2006: first party = supplier, second party = buyer, third party = independent third assessing compliance, fourth party = state/regulatory agency. Havinga (2006) categorized retailers setting food safety standards for their suppliers as third party regulation.
For Abbott and Snidal (2009a: 63) the regulatory process comprises five main stages (or five tasks):

1. Agenda-setting (placing an issue on the regulatory agenda)
2. Negotiation (negotiating, drafting and promulgating standards)
3. Implementation (implementing standards within the operations of targets of regulation such as firms)
4. Monitoring compliance
5. Enforcement (promoting compliance and responding to non-compliance)

Agenda-setting is added to rule-making. Abbott and Snidal use the abbreviation ANIME to refer to these five stages.

In my examination of private forms of food regulation such as GlobalGap and BRC, there seems to be one important phase missing in ANIME. In traditional command-and-control regulation, after making the rules, the next phase is implementing the rules. However, in private regulation drafting and promulgating the rules is not enough. Rules from GlobalGap or BRC, for instance, are not just being implemented in farms or the food industry. It seems to be crucial that these rules were first promoted and eventually made mandatory by dominant market parties such as big supermarket chains.

Regulatees have to accept or adopt a regulation. Adopting a regulation means deciding to accept the rules and to aim at compliance. By and large, an organization may adopt a regulation because it is legally mandatory, because compliance with the regulation is an obligation imposed by a dominant actor in the market (such as the combined retailers) or because the regulatee considers adoption beneficial for some reason (e.g. improving reputation, market share, price). Henson and Humphrey meet this objection by including adoption in the five functions involved in making a food standard operational: standard-setting, adoption, implementation, conformity assessment and enforcement (2011: 155-156). They are right to argue that the distinction between standard setting and adoption clarifies the issue of compulsion and obligation.¹³

Conclusions

When I started working on this paper, my purpose was just to add some distinctions of actors to the three main actor types (state, firm, NGO) and to include some regulatory activities among the three elements of regulation (rule enforcement).

¹³ Both Meidinger and Van Waarden distinguish different functions for certification schemes. Van Waarden (2010) distinguishes three main functions for a certification standard organization such as FSC and MSC: standard setting, accreditation, and trademark assurance. Meidinger arrives at four functions: standard setting, certifying compliance, accrediting certifiers and labelling products (2008: 265).
making, monitoring and enforcement). However, my work on the paper, which involved reading and rereading the literature, revealed the high level of complexity of regulation. Many distinctions are only important in the context of particular regulatory arrangements or a particular inquiry. Besides, it often is more important to know whether the different tasks or functions (e.g. drafting the rules, adopting the rules, implementing the rules, assessing conformity and enforcing compliance) are carried out by the same organisation or by three or more different organisations than to know that all tasks are carried out by market actors (making this a single-actor regulation in the governance triangle).

Black (2008) sees accountability problems because regulatory roles and responsibilities are distributed among several actors (deciding on goals, drafting standard, monitoring, enforcement). It is impossible to call a standard-setter to account for enforcement of the rules or to call the enforcer to account for rules he did not make. Black (2008) asks, ‘How to call to account a constellation of regulators?’

To make it even more complex: the involvement of actors and the roles they perform develop over time. Bernstein and Cashore (2007) showed for non-state, market-driven governance systems that political legitimacy is constructed in a three-phase process with different relationships between the actors and participation of different actors. Bernstein and Cashore distinguish between the initiation phase, the phase of widespread support and the phase of political legitimacy. A picture of the actors and roles involved in Eurepgap in 1997 differs significantly from the picture of GlobalGap in 2010 (Van der Kloet and Havinga 2008, Van der Kloet 2011). The picture will be different again at some other moment in time.

Five main functions seem to apply to every regulatory arrangement. These functions are similar to the functions Henson and Humphrey (2011) distinguish for food standards, but are worded slightly differently to make them universally applicable. For a regulatory arrangement to be effective, rules have to be laid down and subsequently adopted and implemented, and compliance with the rules has to be monitored and enforced. All regulatory activities can be thought of as part of one of these functions.

1. Standard setting (rule making)
2. Adoption
3. Implementation
4. Monitoring compliance (assessing conformity)
5. Enforcement
### Figure 8 Catalogue of regulatory activities connected to five main functions

<table>
<thead>
<tr>
<th>Actions</th>
<th>Sideline actions</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>I. Rule making</strong></td>
<td></td>
</tr>
<tr>
<td>1. First step (initiator) agenda setting</td>
<td>- Lobbying</td>
</tr>
<tr>
<td>2. Determine goals of regulation</td>
<td>- Risk assessment</td>
</tr>
<tr>
<td>3. Negotiation about the rules</td>
<td>- Mobilizing resistance or support</td>
</tr>
<tr>
<td>4. Drafting the rules</td>
<td>- Adapting the rules (to new situations and acquired experience)</td>
</tr>
<tr>
<td>5. Lay down the rules (decide on and promulgating the rules)</td>
<td></td>
</tr>
<tr>
<td>6. Further regulations and implementing rules</td>
<td></td>
</tr>
<tr>
<td><strong>II. Adoption</strong></td>
<td></td>
</tr>
<tr>
<td>7. Adopting the rules</td>
<td>- Promoting and supporting the rules</td>
</tr>
<tr>
<td>8. Imposing the rules on suppliers or other actors in the supply chain (make the rules compulsory for other actors)</td>
<td>- Education &amp; coaching</td>
</tr>
<tr>
<td></td>
<td>- Facilitating</td>
</tr>
<tr>
<td><strong>III. Implementation</strong></td>
<td></td>
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<tr>
<td>9. Implementing rules within firms (or other targets of regulation/regulatees)</td>
<td>- Promoting and supporting the rules</td>
</tr>
<tr>
<td></td>
<td>- Education &amp; coaching</td>
</tr>
<tr>
<td></td>
<td>- Facilitating</td>
</tr>
<tr>
<td><strong>IV. Monitoring</strong></td>
<td></td>
</tr>
<tr>
<td>10. Testing</td>
<td>- Accreditation certification bodies</td>
</tr>
<tr>
<td>11. Inspection</td>
<td>- Trademark assurance</td>
</tr>
<tr>
<td>12. Audit, verification</td>
<td>- Evaluation and adaptation of monitoring strategy</td>
</tr>
<tr>
<td>13. Certification</td>
<td></td>
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<td>14. Documentation</td>
<td></td>
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<tr>
<td>15. Tracing non-compliant and undesirable behaviour</td>
<td></td>
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<tr>
<td><strong>V. Enforcement</strong></td>
<td></td>
</tr>
<tr>
<td>16. Sanctioning non-compliance (warning, fine, withdrawal certificate)</td>
<td>- Accreditation of certification bodies</td>
</tr>
<tr>
<td>17. Legal enforcement (prosecution, civil law claim, appeal)</td>
<td>- Trademark assurance</td>
</tr>
<tr>
<td></td>
<td>- Developing instruments and strategy in response to non compliance</td>
</tr>
<tr>
<td></td>
<td>- Evaluation and adaptation of enforcement strategy</td>
</tr>
</tbody>
</table>

14 These activities may repeat on another level, e.g. EU, Member states, agency and so on and after some time in the future.

15 These activities may repeat up- or downstream in the food chain when requirements are passed on: e.g. supermarket – manufacturer – import firm – export firm – farmer.

16 Monitoring is a continuous process.
We need to analyse particular regulatory regimes and establish which actors are involved, how they are interrelated and what role they play. A crude distinction between three parties -- state, market and civil society -- masks interdependence, conflicts of interest and power. Knowledge of which particular actors are involved and what their role is in the regulatory social field reveals power relations that may have an effect on the reliability of a certificate, the level of compliance with prescriptions, and the openness of decision making. These insights might trigger government agencies to interfere or to monitor more actively.
References


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