

**ADVERTISING REGULATIONS
IN PHARMACEUTICAL MARKETS:
PRODUCT
VERSUS
ENLIGHTENMENT**

Junichiro Ishida
Tsuyoshi Takahara

June 2019

The Institute of Social and Economic Research
Osaka University
6-1 Mihogaoka, Ibaraki, Osaka 567-0047, Japan

Advertising Regulations in Pharmaceutical Markets: Product Versus Enlightenment*

Junichiro Ishida[†]

Tsuyoshi Takahara[‡]

June 17, 2019

Abstract

This paper analyzes the optimal content regulation of direct-to-consumer advertisement (DTCA) in a pharmaceutical market, with particular focus on the distinction between product and enlightenment advertisement. Firms are allowed to freely promote their own specific products under product DTCA, whereas they can only advertise the presence of a disease and its typical subjective symptoms under enlightenment DTCA. The content regulation changes the nature of market competition and the incentive to invest in advertisement, thereby yielding substantial welfare and policy implications. The overall welfare impact of the content regulation is ambiguous and depends, among other things, on the cost effectiveness of advertisement and the market-size distortion induced by product DTCA. We also analyze the effect of free market pricing and argue that a less stringent advertisement regulation, i.e., product DTCA, is often complementary to a less stringent price regulation.

JEL Classification Code: M37, I11, I18

Keywords: DTCA, Advertisement content regulation, Prescription decision

*The authors acknowledge the financial support of the Japan Society for Promotion of Science (JSPS) via Grants-in-Aid for Scientific Research (15H05728).

[†]Institute of Social and Economic Research, Osaka University. E-Mail: jishida@iser.osaka-u.ac.jp

[‡]School of Business Administration, Kwansei Gakuin University. E-Mail: takahara@a0.rusk.to

1 Introduction

Marketing activities, such as TV advertisement, play a very important role in the pharmaceutical industry. According to Sufrin and Ross (2008), Pfizer, Merck, and Bristol-Meyer-Squibb – the three biggest pharmaceutical firms in the US – spend a greater share of expenditure on marketing than R&D activities in 2005. Although a large fraction of advertisement is directed at physicians (prescribers), pharmaceutical firms are increasingly more involved in direct-to-consumer advertisement (DTCA), especially after the deregulation of DTCA of prescription drugs by the US Food and Drug Administration (FDA) in 1997. Owing to some unique features of the industry, however, many experts and practitioners suspect the effectiveness of DTCA in pharmaceutical markets by noting the possibility that it may distort prescription decisions. There is now a heated debate worldwide about whether DTCA should be regulated, and if so, in what ways.

A body of evidence suggests that DTCA in the pharmaceutical industry can in fact be a double-edged sword. As proponents of DTCA insist, it could serve an educational purpose by informing patients of a particular medical condition and a potential treatment, thereby encouraging them to contact a health care provider at an opportune time. Opponents of DTCA argue, however, that it often misinforms patients, by selectively omitting risk factors or causes of the condition and overemphasizing drug benefits (see Section 2.6 for more discussion on this).¹ Among those, Ventola (2011) notes:

If a patient's request for an advertised drug is clinically inappropriate and the health care provider is unable or unwilling to correct the patients perception that it is a good choice, this situation may lead to unnecessary or harmful prescribing. An additional problem mentioned by critics is that patients may withhold information to fit a particular profile that they saw in DTC ads in an attempt to get the doctor to prescribe a drug they want but that might not be appropriate for them.

To sum up, giving information to patients is beneficial to the extent that it brings them to clinics in search of medical treatment, but anything more than that could just disturb the physician-patient relationship and cause unnecessary complications. If this is the case, there may be a scope

¹The most notable example may be the case of Rofecoxib which is a nonsteroidal anti-inflammatory drug marketed by Merck. While approved in 1999, it was later withdrawn over safety concerns after disclosures that Merck withheld information about the drug's risks from physicians and patients for over five years.

for welfare improvement by regulating the content of DTCA. One way to do this is to require that DTCA only describe the presence of a disease and its typical symptoms and treatments but without any information about particular brand names or products – the type of advertisement often called enlightenment or help-seeking advertisement. This is precisely what has been done in many countries outside of the US where DTCA is permitted only if it is enlightenment in nature.²

At a glance, enlightenment DTCA can be an effective remedy for the distortion caused by DTCA in pharmaceutical markets because it only provides minimal information to patients while still serving the educational purpose. However, the welfare impact of such a regulation is not necessarily so straightforward, especially in pharmaceutical markets that are typically oligopolistic and dominated by a few major firms, as it alters the strategic nature of market competition. In particular, under enlightenment DTCA, advertisement effectively becomes a public good, which diminishes each firm's incentive to engage in marketing activities. This can be welfare-reducing as it may leave a large fraction of potential patients uninformed and receiving no treatment at all in the end.

In this paper, we investigate how the content regulation of DTCA (hereafter, simply the content regulation) affects pharmaceutical firms' expenditures on advertisement and welfare to contribute to the current debate surrounding the pharmaceutical industry. To this end, we analyze a market of prescription-only drugs for a particular disease with two horizontally differentiated firms (pharmaceutical suppliers). The two firms decide how much to advertise to reach out potential patients who are initially unaware of the disease and need to be informed via advertisement to receive any medical treatment before it becomes too serious. Within this framework, we consider two DTCA regulation policies: in one case, the firms are allowed to promote their own specific brand names (product DTCA); in the other, they can only provide general information about a disease and its typical symptoms without specifying their own brand names (enlightenment DTCA).

The difference in the nature of DTCA gives different incentives to the pharmaceutical firms and induces different responses from potential patients. Under enlightenment DCTA, no specific information about a particular product is available, and all informed patients are largely neutral with respect to the two available products. Under product DTCA, on the other hand, informed patients

²Outside of the US, New Zealand is the only OECD country which allows DTCA that includes product names. In other countries, there are some regulations on the extent that a firm can promote its own brands or products, virtually prohibiting product DTCA.

may develop a biased preference for a particular product, depending on the extent of exposure to DTCA. Borrowing from Brekke and Kuhn (2006), we call them selective and captive throughout the analysis: patients are called selective when they are exposed to both firms' products and captive when exposed only to one of them. In either case, patients are informed and hence visit a (representative) physician in search of medical treatment, which captures the demand-expansion effect of DTCA Grossman and Shapiro (1984). We assume, however, that only selective patients can ask the physician to compare the two products and be prescribed a suitable drug while captive patients, with a biased attachment, insist on the product they have in mind and needs to be persuaded if an alternative measure is to be taken.

The key assumption we make here is that this persuasion process is, broadly speaking, costly for the physician. The presence of captive patients may then create a welfare loss by leading to inappropriate prescribing if the physician deems the persuasion cost too much to bear. A subtle point to note is that the presence of the persuasion cost, which imposes an extra burden on the physician, creates two forms of market distortion under product DTCA: on one hand, a fraction of captive patients may receive no treatment; on the other hand, they may also be prescribed the suboptimal product. To make clear distinction between them, we refer to the former, which limits the equilibrium industry demand below its efficient level, the "market-size distortion," and to the latter as the "prescription distortion." These two forms of distortion depend on various parameters in a somewhat complicated way and play a major role in our welfare analysis.

Main results: We obtain several results regarding the costs and benefits of product DTCA vis-a-vis enlightenment DTCA. Our argument centers around the interactions between the two forms of market distortion mentioned above and the strategic investment incentives that are crucially shaped by the context regulation in effect.

First, it is intuitively clear that the firms invest more in advertisement under product DTCA than under enlightenment DTCA. In fact, the advertisement competition is excessively strong under product DTCA because of the potential of business stealing, as a consequence of which the firms overinvest in advertisement compared to the first-best level. Under enlightenment DTCA, on the other hand, the incentives to inform patients via advertisement are substantially weaker due to its public-good nature, and the firms indeed underinvest compared to the first-best level. Despite the apparent distortion in prescription decisions induced by product DTCA, the welfare effect of

the content regulation is thus ambiguous because the stronger, though excessive, incentives to invest in advertisement reduce the number of uninformed patients, ultimately helping them receive treatment in time.

With this result as a building block, we proceed to characterize the welfare impact of the content regulation. When the cost of advertisement is relatively high, the interests of the firms and potential patients are better aligned, and there often exists a Pareto-improving policy choice. The optimal policy in this case is largely determined by the extent of the market-size distortion induced by product DTCA: enlightenment DTCA is preferred when the market-size distortion is more severe while product DTCA is preferred when it is less so. As the cost of advertisement decreases, however, the overinvestment problem becomes more of a factor, and a serious conflict arises between the firms and potential patients as a consequence: while the overinvestment problem is purely detrimental to the firms, it is beneficial from the viewpoint of patients as it reduces the likelihood of being left out and helps them seek treatment in time. In general, patients are made better off product DTCA as the cost gets smaller while the firms are made worse off.

It is also worth emphasizing that although the presence of the persuasion cost is the source of market distortion in our setup, the welfare impact of the persuasion cost is not entirely negative. This is because an increase in the persuasion cost affects the two forms of distortion in different ways: while it aggravates the prescription distortion, it reduces the market-size distortion by allowing each firm to capture patients who would otherwise be prescribed the rival firm's product. The latter effect provides the firms an extra incentive to invest in advertisement, which can potentially be welfare-enhancing by reducing the number of uninformed patients. We find that this “distortions as incentives” mechanism can indeed be strong enough to compensate for the loss from the prescription distortion and benefit not only the firms but also patients under some conditions.

Finally, although most of our key insight can be illustrated under the assumption of regulated prices, we also extend the baseline model to incorporate the firms' price setting behavior to evaluate the impact of free market pricing in this context. We find that when the regulated price is set low enough to begin with, free market pricing not only benefits the firms but also patients despite the higher equilibrium price. This suggests that the two forms of regulation are often complementary to each other in that a less stringent advertisement regulation, i.e., product DTCA, tends to be compatible with a less stringent price regulation – the type of policy mix adopted in the US – and

hence should be evaluated jointly.

Literature: The existing literature on DTCA in the pharmaceutical industry is mostly empirical (Rosenthal et al., 2003; Donohue et al., 2004; Iizuka, 2004; Iizuka and Jin, 2007; Bradford et al., 2010; Sinkinson and Starc, 2018), and there are only a handful of theoretical analyses, despite the ongoing debate over the desirability of DTCA.³ Among them, our model is most closely related to and draws on Brekke and Kuhn (2006) who investigate the strategic relationship between DTCA and detailing (marketing to physicians). A crucial difference is that their focus is on how much information to disseminate to physicians via marketing while their DTCA is enlightenment in nature as its only role is to bring them to clinics.⁴ Here, we attempt to shed light on a different aspect of the problem by making a clear distinction between product and enlightenment DTCA. To the best of our knowledge, this is the first work which emphasizes this distinction and analyzes the impact of the content regulation of DTCA.

Our work also has an obvious connection to analyses of advertisement in industrial organization. In conventional analyses of advertisement, consumers precisely know their preferences (or payoff functions). Here, the situation we deal with is somewhat special in that physicians, with their expertise, possess far more knowledge about what patients need than patients themselves: specifically, we assume that patients are lined up on a Hotelling line without knowing their exact location which only physicians can tell. However, even though physicians are informationally superior, it is still patients who must make an initial move and contact a physician on their own. This unique industry structure gives rise to the possibility of “misinformation” and a scope for welfare improvement by policy interventions. Although we cast our analysis in a specific context of advertisement competition, we believe that this problem can provide a broader perspective as it is a realistic and pervasive issue in the medical industry to what extent patients should be informed.

³A recent exception is Bar and Lillard (2014) who analyze a situation where there are two alternative treatments, and one of them is advertised directly to patients by a profit-maximizing pharmaceutical firm.

⁴In their model, there are captive and selective physicians, but on such distinction is made for patients.

2 The Model

2.1 Environment

We consider a market of prescription-only drugs for a particular disease that comprises of two pharmaceutical firms, potential patients (male) and a representative physician (female). In this market, potential patients with unit mass are horizontally and uniformly distributed on the Hotelling line extended on $[0, 1]$. Each patient is identified with his location x which captures his “type” (e.g., disease type and/or physical constitution). The pharmaceutical products are supplied by the two firms, each indexed by $i = 0, 1$, where firm 0 is located at the left end of the Hotelling line while firm 1 is located at the right end. We suppose that the two firms supply pharmaceutical products for the same disease although their products are horizontally differentiated by their chemical compounds and consequent treatment effects.

2.2 Pharmaceutical firms

In the first stage of the game, the two firms simultaneously choose the levels of DTCA, which directly affect each potential patient’s decision of whether or not to seek medical treatment by visiting a physician. Let θ_i denote the fraction of patients who are exposed to firm i ’s advertisement. The demand for firm i ’s product (product i) is assumed to be a function of (θ_i, θ_{-i}) , $i \neq -i$ and denoted as $Q_i(\theta_i, \theta_{-i})$. Firm i ’s profit is then given by

$$\pi_i = p_i Q_i(\theta_i, \theta_{-i}) - \frac{c}{2} \theta_i^2,$$

where p_i denotes the price of product i , and c the cost effectiveness of advertisement.

As most of our key insight can be illustrated under regulated prices, we assume for now that the pharmaceutical prices are fixed exogenously at $p = p_0 = p_1$ based on factors that are not directly related to the issue at hand. The assumption of regulated prices is a relevant one in most countries outside the US while we will extend our analysis to allow the firms to set prices endogenously in section 5. To ensure the existence of interior solutions, we also assume that c is large relative to p .

Assumption 1. $2c > p$.

2.3 Patients

A patient receives medical treatment when he visits a physician. The utility derived by patient x from consuming a unit of product i is given by

$$u(x, i) = v - \tau|x - i| - \lambda p,$$

where $v \in (0, \infty)$ denotes the gross effectiveness of a pharmaceutical product, $\lambda \in [0, 1]$ the copayment rate, and $\tau \in (0, \infty)$ the “transport cost” which measures the quality/characteristic diversion between the two products.

The key assumption is that a patient’s type x can be observed only by the physician but not by the patient himself – a defining characteristic of the pharmaceutical industry – as assumed in Brekke and Kuhn (2006). The two products have the same effectiveness but differ in their treatment effects across patients, depending on the specific product-patient type match. We assume that each product is effective enough to ensure that it is beneficial even for the most “distant” patient (at $x = 0.5$).

Assumption 2. $2(v - \lambda p) > \tau$.

The utility of patients who receive no treatment (either by their own or the physician’s choice) is normalized to 0.

2.4 Advertising regulation

Throughout the analysis, we consider two regulatory regimes on the content of DTCA: product and enlightenment. Under product DTCA, a pharmaceutical firm is allowed to provide information about a specific product, including its brand name or the names of chemical components.⁵ Then,

⁵An important premise of our analysis is that product DTCA is effective in raising the demand for its own product (via business stealing) and indeed generates different effects from enlightenment DTCA. Of course, if product and enlightenment DTCA make no difference, the whole debate over DTCA becomes groundless; the current debate over DTCA reflects physicians’ perception that product DTCA does make a difference at some level. Although empirical evidence on this issue may not be not conclusive (see, for instance, Iizuka and Jin (2007) who find no significant effect of DTCA on the market share), there is in fact some evidence in support of this claim (Kalyanaram, 2008, 2009; Liu and Gupta, 2011; Dave and Saffer, 2012; Sinkinson and Starc, 2018). Most notably, Sinkinson and Starc (2018) measure the impact of DTCA by pharmaceutical firms and find substantial evidence of business stealing.

when the firms are allowed to engage in product DTCA, patients are classified into three (publicly observable) informational types, depending on the extent of exposure to the firms' DTCA:

- A patient is *selective* if he is exposed to advertisement from both firms, which occurs with probability $\theta_i\theta_{-i}$;
- A patient is *captive of product i* if he is exposed to advertisement from only one of the firms, which occurs with probability $\theta_i(1 - \theta_{-i})$;⁶
- A patient is *uninformed* if he is exposed to advertisement from neither, which occurs with probability $(1 - \theta_i)(1 - \theta_{-i})$.

In contrast, when the firms are restricted to enlightenment DTCA, they can only provide general information about the presence of a disease and its symptoms without specifying particular brand names or products. Under enlightenment DTCA, therefore, there are no captive patients. As such, a patient becomes selective with probability $\theta_i + \theta_{-i} - \theta_i\theta_{-i}$ and uninformed with $(1 - \theta_i)(1 - \theta_{-i})$.

2.5 Physician

We assume that there is a representative physician to whom all potential patients visit when they become informed. The physician can prescribe either product 0 or product 1, or else can choose to provide no treatment (or an outside treatment). The physician is partly altruistic in that she cares about the welfare of attending patients net of the “persuasion cost.” The utility of the physician depends on the attending patient's informational type.

First, if a patient is selective, he is neutral and simply accepts the physician's advice as it is. In this case, the physician incurs no persuasion cost and her utility from prescribing product i is given by $u(x, i)$. The physician thus simply prescribes the more suitable product for the patient.⁷

Second, if a patient is captive, he has developed a biased preference in favor of a particular product and makes a specific request for it. We assume in this case that the physician either

⁶We assume that a patient becomes captive when he is exposed only to one of the products. Although this may appear a strong assumption to impose, our conclusions hold in a qualitative as long as there is a positive fraction of patients who become captive.

⁷Under Assumption 2, it is never optimal to provide no treatment in this case.

prescribes the product as requested or provides no treatment.⁸ If the physician takes the option of no treatment, she must incur a persuasion cost d which includes time, effort, and psychological discomfort of persuading a patient, as well as the risk of disrupting the future relationship. As such, the physician prescribes the requested product i if and only if $u(x, i) \geq -d$.

2.6 Discussion on distorted prescription decisions

The key premise of our analysis is that product DTCA may disturb the physician-patient relationship and distort prescription decisions to some extent. We believe, at some intuitive level, that this is not only possible but also almost inevitable. On the one hand, it is hardly surprising that patients sometimes develop a biased preference for a particular product: first, it is very difficult, if not impossible, for patients to understand all the possible consequences of prescription drugs; at the same time, health-related problems are always emotionally arousing, and it is also very difficult to stay perfectly rational by taking emotion entirely out of the decision process. On the other hand, physicians are also humans with limited resources and persuasion is never free. A combination of these two factors necessarily gives rise to some distortion in prescription decisions.

To motivate our analysis further, there is also ample evidence which suggests that giving detailed information to patients via product DTCA creates seemingly unnecessary complications and psychological pressures to physicians. Murray et al. (2003) report that many physicians feel that DTCA worsens the time efficiency of the visit and have a negative response to the recent increase in DTCA.⁹ Moreover, it is well documented that many physicians have prescribed inappropriate drugs which are requested by patients.¹⁰ Kravitz et al. (2005) also report that patients' requests

⁸We assume that it is more costly to persuade a captive patient into taking the other product and hence do not consider this possibility. More generally, we may assume that the physician can prescribe the unrequested product by incurring a cost D where $D > d$. Our setup can then be interpreted as the one in which D is prohibitively large. We focus on this extreme case because our results do not change in a qualitative sense even when D is lower and finite because any market distortion that can arise in this process will be captured thoroughly by two distortion measures, M and ϕ , which we will define later.

⁹Robinson et al. (2004) also report that "few (physicians) believe that these advertisements are a positive trend in health care."

¹⁰Frosch et al. (2010) and Ventola (2011) survey literature on the effect of DTCA on the physicians' prescription decision. In Frosch et al. (2010), they conclude that the request for the advertised product by the patients increases both appropriate and inappropriate prescribing, though it is unclear which effect is greater. Weissman et al. (2004) show that the physicians who consider DTCA positively prescribe more DTCA drug than the physicians who consider DTCA negatively. Dai et al. (2005) insists that the marketing activities by Merck may grow the use of Vioxx, which is discontinued due to fatal side effects.

have a profound effect on prescription decisions in major depression and adjustment disorder. These results lend support for the view that physicians do indeed face some persuasion costs when patients are exposed to product DTCA. Additionally, the persuasion cost may seem to differ across symptoms. For instance, Kravitz et al. (2005) find that the influence of patients, especially brand-specific requests, on the prescription of antidepressants is higher for adjustment disorder than for major depression and argue that DTCA may stimulate prescribing more for questionable than for clear indications. This line of argument is again consistent with the existence of persuasion costs because it is natural to think that persuasion is more costly in the absence of clear and verifiable scientific evidence.

3 Analysis

3.1 Prescription decisions

We solve the model backward, starting with the final stage where the representative physician decides whether to prescribe a drug and, if so, which one. Given that a patient visits the physician, he must be either selective or captive. The physician's prescription decision clearly depends on the patient's informational type as we describe below.

Suppose first that a patient is selective (exposed to both products). Under Assumption 2, the physician recommends product 0 (1) if $0.5 \geq x$ ($x > 0.5$) and the patient simply accepts the advice. As such, the two firms split the market evenly in this contingency. Since the two firms compete for selective patients, this is the competitive segment of the market.

If a patient is captive (exposed only to one of the products), on the other hand, the physician chooses either to prescribe the product he requests or to provide no treatment. The market is now monopolized by one firm, i.e., this is the monopolistic segment of the market. The physician prescribes a requested product if and only if the benefit exceeds the persuasion cost, i.e.,

$$v - \lambda p - \tau|x - i| \geq -d.$$

The demand for each product in the monopolistic segment is then given by

$$M := \min \left\{ \frac{v - \lambda p + d}{\tau}, 1 \right\}.$$

By Assumption 2, $M > 0.5$, so that each firm can capture a larger market share, partly at the expense of the physician who is forced to incur the persuasion cost.

Note that the presence of the persuasion cost generates two forms of distortion in prescription decisions. First, there is a fraction of captive patients who end up with receiving no treatment. For the sake of exposition, we call this the “market-size distortion,” the size of which is captured by M , as it limits the industry demand below its efficient level. We say that the market-size distortion is more severe when M is small and close to 0.5, while it is less so when M is large and close to 1. Second, there is also a fraction of captive patients who are prescribed the suboptimal product. This form of distortion is called the “prescription distortion,” which we will discuss in depth when we analyze the welfare of patients in section 4.3.

3.2 Advertisement expenditures

Each firm decides how much to invest in advertisement, taking the other firm’s choice as given. First, under product DTCA, a patient is selective with probability $\theta_i \theta_{-i}$ and captive of i with probability $\theta_i(1 - \theta_{-i})$. The demand for product i under product DTCA is then obtained as

$$Q_i^P = \frac{\theta_i \theta_{-i}}{2} + \theta_i(1 - \theta_{-i})M,$$

where the superscript P stands for product DTCA. It follows from this that firm i ’s problem under product DTCA can be written as

$$\max_{\theta_i \in [0,1]} p Q_i^P - \frac{c}{2} \theta_i^2 = p \left[\frac{\theta_i \theta_{-i}}{2} + \theta_i(1 - \theta_{-i})M \right] - \frac{c}{2} \theta_i^2.$$

The best-response function is given by

$$\theta_i = \frac{p}{2c} [2M - \theta_{-i}(2M - 1)].$$

Solving this, the equilibrium expenditure, denoted by θ^P , can be obtained as

$$\theta^P = \frac{2Mp}{2c + p(2M - 1)}.$$

Note that $1 > \theta^P$ by Assumption 1.

Second, there are no captive patients under enlightenment DTCA. The demand under enlightenment DTCA is hence given by

$$Q_i^E = \frac{\theta_i + \theta_{-i} - \theta_i \theta_{-i}}{2},$$

where, as above, the superscript E stands for enlightenment DTCA. As such, firm i 's problem under enlightenment DTCA can be written as

$$\max_{\theta_i \in [0,1]} pQ_i^E - \frac{c}{2}\theta_i^2 = p \frac{\theta_i + \theta_{-i} - \theta_i \theta_{-i}}{2} - \frac{c}{2}\theta_i^2.$$

We can then obtain the best-response function as

$$\theta_i = \frac{p(1 - \theta_{-i})}{2c}.$$

It follows from this that the equilibrium expenditure, denoted by θ_i^E , can be obtained as

$$\theta^E = \frac{p}{2c + p}.$$

It is intuitively clear that the firms have more incentive to invest in advertisement when they are allowed to engage in product DTCA. To see this, observe that

$$\theta^P > \theta^E \Leftrightarrow \frac{2Mp}{2c + p(2M - 1)} > \frac{p}{2c + p}.$$

This is further reduced to

$$2M(2c + p) > 2c + p(2M - 1),$$

which holds for any $M \in (0.5, 1]$. In words, there is a private gain from DTCA when the firms are allowed to promote their own products, giving them more incentive to invest in advertisement. Moreover, we can show that the difference in the advertisement expenditures widens as c gets smaller. We summarize these findings below.

Proposition 1. *The advertisement expenditure is higher under product DTCA than under enlightenment DTCA. Moreover, the difference between θ^P and θ^E is strictly decreasing in c .*

Proof. *We have already seen that $\theta^P > \theta^E$ always holds. Note also that*

$$\frac{\partial \theta^P}{\partial c} - \frac{\partial \theta^E}{\partial c} = -\frac{4Mp}{[2c + p(2M - 1)]^2} + \frac{2p}{(2c + p)^2}.$$

This is negative if

$$\frac{4Mp}{[2c + p(2M - 1)]^2} > \frac{2p}{(2c + p)^2},$$

which can be written as

$$2M(2c + p)^2 > [2c + p(2M - 1)]^2.$$

With some computation, this is reduced to

$$4c^2(2M - 1) + 4cp + p^2[2M - (2M - 1)^2] > 0,$$

which holds for any $M \in (0.5, 1]$.

3.3 Equilibrium market demand

The effect of the content regulation on the demand for each product is ambiguous with two opposing forces at work: although product DTCA expands the potential market size due to higher advertisement expenditures, prescription decisions are necessarily distorted. To see this, observe

that

$$\Delta Q := Q^P - Q^E = \frac{\theta^P[\theta^P + 2(1 - \theta^P)M]}{2} - \frac{\theta^E(2 - \theta^E)}{2}.$$

This shows that for any $M \in (0.5, 1]$ and $\theta^P = \theta^E$, $\Delta Q < 0$, i.e., if the equilibrium advertisement expenditures are the same between the two regimes, the demand for each product is higher under enlightenment DTCA. A necessary condition for $\Delta Q > 0$ is hence that θ^P is sufficiently larger than θ^E , although this happens for a wide range of parameter values as we will see below.

To conduct comparative statics on ΔQ , we first obtain

$$\Delta Q = \frac{4M^2cp}{[2c + p(2M - 1)]^2} - \frac{p(4c + p)}{2(2c + p)^2}.$$

From this, $\Delta Q > 0$ if and only if

$$\frac{8M^2c}{4c + p} > \left[\frac{2c + p(2M - 1)}{2c + p} \right]^2. \quad (1)$$

Two facts are worth noting here. First, ΔQ naturally converges to 0 as c tends to infinity, because the advertisement expenditures converge to zero under both regimes. Second, observe also that (1) always holds at $c = 0.5p$, regardless of M , since the difference in the advertisement expenditures widens as c gets smaller. Aside from these properties, the exact shape of ΔQ depends crucially on the value of M , as the next proposition fully characterizes.

Proposition 2. $\Delta Q > 0$ for all $c \in [0.5p, \infty)$ if $M \geq \sqrt{0.5}$. For $M \in [0.5, \sqrt{0.5})$, there exists an interior threshold $c_Q(v, d) \in (0.5p, \infty)$ such that $\Delta Q > 0$ if and only if $c_Q(v, d) > c$. Moreover, $c_Q(v, d)$ is increasing in both v and d .

Proof. See Appendix.

[Figure 1 about here]

3.4 First best

The content regulation influences the equilibrium advertisement expenditure by changing the nature of competition between the two firms. To illustrate this point, it is instructive to derive the

first-best level of advertisement expenditures, defined as the one that maximizes the joint profit of the two firms, and compare it to the equilibrium level.

The first-best level under product DTCA can be obtained by solving the following problem:

$$\max_{\theta \in [0,1]} p \left[\frac{\theta^2}{2} + \theta(1 - \theta)M \right] - \frac{c}{2}\theta^2$$

Let θ^{P*} denote the first-best level under product DTCA, which is given by

$$\theta^{P*} = \frac{Mp}{c + p(2M - 1)}.$$

It is clear that $\theta^P > \theta^{P*}$, i.e., the firms overinvest in advertisement under product DCTA. This is due to the business-stealing effect of advertisement: a firm can “steal” a fraction of captive patients by reaching out to them and transforming them into selective.

Similarly, the problem under enlightenment DTCA is defined as

$$\max_{\theta \in [0,1]} \frac{p}{2}(2\theta - \theta^2) - \frac{c}{2}\theta^2$$

As above, let θ^{E*} denote the first-best level under enlightenment DTCA, which is given by

$$\theta^{E*} = \frac{p}{c + p}.$$

It follows from this that $\theta^{E*} > \theta^E$, i.e., the firms underinvest in advertisement under enlightenment DTCA. Aside from the fact that there is no business-stealing effect, this is due to the fact that advertisement under enlightenment DTCA is a public good which also increases the demand for the rival product as well. We summarize these findings in the following proposition.

Proposition 3. *The firms overinvest in advertisement under product DTCA and underinvest under enlightenment DTCA.*

4 Optimal content regulation of DTCA

4.1 Welfare measures

Denote by π^k the equilibrium profit under regime $k = P, E$. Similarly, we denote by PS^k what we call the patient surplus (to be defined more precisely below). We consider a regulatory authority with the objective function given by

$$W^k = \omega\pi^k + (1 - \omega)PS^k, \quad k = P, E,$$

where the parameter $\omega \in [0, 1]$ reflects the authority's policy stance where it is more “pro-business” when ω is close to one while it is “pro-patient” when ω is close to zero.¹¹

4.2 Firm profit

We first examine how the content regulation affects the firms' equilibrium profit. Letting $\Delta\pi := \pi^P - \pi^E$, $\Delta\theta := \theta^P - \theta^E$ and $\Delta C := (\theta^{P^2} - \theta^{E^2}) = \Delta\theta(\theta^P + \theta^E)$, we obtain

$$\begin{aligned} \Delta\pi &= p\Delta Q - \frac{c}{2}\Delta C \\ &= \frac{p\theta^P[2M - (2M - 1)\theta^P]}{2} - \frac{p\theta^E(2 - \theta^E)}{2} - \frac{c}{2}\Delta\theta(\theta^P + \theta^E) \\ &= -p(1 - M)\theta^E(1 - \theta^E) - \frac{\Delta\theta}{2} \left[\frac{(2M - 1)p + c}{2}(\theta^P + \theta^E) - 2Mp \right]. \end{aligned} \quad (2)$$

The impact of the content regulation on the equilibrium profit can be decomposed into two factors. Evaluated at $\theta^P = \theta^E$, any difference in the profits can be attributed to the distortion in prescription decisions. The first term of the right-hand side of (2) captures this effect which is negative for any $1 > M$. The second term captures the difference in the gains from demand expansion. Note that the sign of this effect is ambiguous, because the firms tend to overinvest under product DTCA, diminishing their profit margins with excessive competition in advertisement.

We have already characterized ΔQ in Proposition 2. Also, from Proposition 1, we know $\Delta C \geq 0$, meaning that $\Delta Q < 0$ is a sufficient condition for $\Delta\pi < 0$. This implies that the firms are more

¹¹We do not take into account the persuasion cost incurred by the physician.

likely to prefer enlightenment DTCA to product DTCA. In fact, this is always the case when M is relatively small, i.e., the market-size distortion is relatively large, as the next proposition suggests.

Proposition 4. $\Delta\pi < 0$ for all $c \in [0.5p, \infty)$ if $\sqrt{0.75} \geq M$. For $M \in (\sqrt{0.75}, 1]$, there exists an interior threshold $c_\pi(v, d) \in (0.5p, \infty)$ such that $\Delta\pi < 0$ if and only if $c_\pi(v, d) > c$. Moreover, $c_\pi(v, d)$ is decreasing in both v and d .

Proof. See appendix.

[Figure 2 about here]

4.3 Patient surplus

From the viewpoint of policymaking, the welfare of patients often carries more weight than that of firms. Since the payoff of uninformed patients is invariably zero, we only need to consider selective and captive patients. Let PS_S and PS_C denote the patient surplus for selective and captive patients, respectively. Also, define PS^k as the total patient surplus under regime $k = P, E$, where

$$\begin{aligned} PS^P &= \theta^{P^2} PS_S + 2\theta^P(1 - \theta^P) PS_C \\ PS^E &= \theta^E(2 - \theta^E) PS_S. \end{aligned}$$

First, selective patients always end up with the most suitable product. The patient surplus for this group of patients is hence given by

$$\begin{aligned} PS_S &= 2 \int_0^{\frac{1}{2}} (v - \lambda p - \tau x) dx \\ &= v - \lambda p - \frac{\tau}{4}. \end{aligned}$$

On the other hand, captive patients are prescribed the requested product as far as the patient benefit exceeds the persuasion cost. The patient surplus for this group of patients is given by

$$\begin{aligned} PS_C &= \int_0^M (v - \lambda p - \tau x) dx \\ &= (v - \lambda p)M - \tau \frac{M^2}{2}. \end{aligned}$$

Define

$$\phi := \frac{4(v - \lambda p) - 2\tau M}{4(v - \lambda p) - \tau},$$

such that $M\phi := \frac{PS_C}{PS_S}$ which we use as a measure of the overall distortion in prescription decisions (more on this point later in section 4.4).

The welfare ranking among the three types of patient is clear: selective patients earn the highest payoff while uninformed ones earn the lowest (which is zero). There are hence two ways to improve the patient surplus, either to increase the number of selective patients or to decrease the number of uninformed patients. More precisely, similarly as above, define $\Delta PS := PS^P - PS^E$, which is given by

$$\begin{aligned} \Delta PS &= \theta^{P^2} PS_S + 2\theta^P(1 - \theta^P)PS_C - \theta^E(2 - \theta^E)PS_S \\ &= -2\theta^P(1 - \theta^P)(PS_S - PS_C) + \Delta\theta(2 - \theta^P - \theta^E)PS_S \end{aligned} \quad (3)$$

Again, the impact of the content regulation can be decomposed into two factors. The first term of the right-hand side of (3) captures the loss due to the distortion in prescription decisions under product DTCA: this fraction of patients would have been selective under enlightenment DTCA but become captive under product DTCA. Since $PS_S > PS_C$, this term is always negative. The second term captures the demand-expansion effect under product DTCA, which is always positive.

To conduct comparative statics on ΔPS , we write (3) as

$$\Delta PS = \frac{4Mp[MpPS_S + (2c - p)PS_C]}{[2c + p(2M - 1)]^2} - \frac{p(4c + p)PS_S}{(2c + p)^2}.$$

From this, $\Delta PS > 0$ if and only if

$$\frac{4M^2[p + (2c - p)\phi]}{4c + p} > \left[\frac{2c + p(2M - 1)}{2c + p} \right]^2. \quad (4)$$

Clearly, there is a close relationship between ΔPS and ΔQ ; we in fact obtain (1) if ϕ in (4) is replaced with 1. First, for the same reason as for ΔQ , ΔPS converges to 0 as c tends to infinity.

Second, as the demand-expansion effect dominates, this condition holds at $c = 0.5p$. Following the same procedure, we can establish the following claim which runs parallel to Proposition 2, except for the effect of d which we will discuss in more depth in the next subsection.

Proposition 5. $\Delta PS > 0$ for all $c \in [0.5p, \infty)$ if $M\sqrt{\phi} \geq \sqrt{0.5}$. For $M \in [0.5, \sqrt{0.5\phi^{-1}})$, there exists an interior threshold $c_{PS}(v, d)$ such that $\Delta PS > 0$ if and only if $c_{PS}(v, d) > c$. Moreover, $c_{PS}(v, d)$ is increasing in v .

Proof. See appendix.

The overall welfare effects of the content regulation are summarized in Figure 4. There are two crucial factors: the cost of advertisement c and the market-size distortion (which in turn depends on various parameters such as v , p and d). When the cost of advertisement is relatively high, the interests of the firms and potential patients are better aligned, and there often exists a Pareto-improving regime, depending on the extent of the market-size distortion: enlightenment DTCA is preferred when the market-size distortion is more severe (M close to 0.5) while product DTCA is preferred when it is less so (M close to 1). As c decreases, however, a conflict arises between them. In general, product DTCA becomes the preferred choice for patients when c is relatively small, because the benefit of product DTCA is maximized while the distortion in prescription decisions becomes more attenuated.¹² However, the firms are generally worse off under product DTCA because the overinvestment problem becomes more serious as the cost of advertisement gets smaller.

[Figures 3 and 4 about here]

4.4 Market distortions as incentives

As mentioned earlier, the distortion in prescription decisions is ultimately detrimental to patients for two reasons. First, the size of the market for captive patients is restricted as only a fraction M of them receive any treatment (the market-size distortion). Second, among those who do receive treatment, some of them are prescribed the suboptimal product (the prescription distortion). The

¹²To see this, consider an extreme case where $c = 0.5p$ and hence $\theta^P = 1$. In this case, the patient surplus is maximized because patients are all selective. Patients are clearly made better off under product DTCA as it causes no distortion while serving the educational purpose to the full extent.

distortion in prescription decisions can be decomposed into two parts, where the ratio ϕ measures the size of the prescription distortion (corrected for the market size). Obviously, there arises no prescription distortion ($\phi = 1$) when $\tau = 0$, i.e., when the two products are perfect substitutes.

One crucial determinant of the overall distortion $M\phi$ is the persuasion cost d . An interesting point to note is that the effect of the persuasion cost is not entirely negative because a change in d affects M and ϕ in different ways: an increase in d raises M (less market-size distortion) but lowers ϕ (more prescription distortion). The former effect is conducive to more investment and can be welfare-improving under product DTCA (as evidenced by the facts that Q^P and π^P are both increasing in d). The effect on the patient surplus is, however, more ambiguous as it also involves the prescription distortion. Below, we dissect how a change in the persuasion cost d affects the patient surplus.

Since the distortion due to the persuasion cost only arises under product DTCA, we restrict our attention only to this case. Recall that the patient surplus under product DTCA is given by

$$PS^P = \theta^P[\theta^P PS_S + 2(1 - \theta^P)PS_C].$$

Taking partial derivative with respect to d yields

$$\frac{\partial PS^P}{\partial d} = 2[\theta^P PS_S + (1 - 2\theta^P)PS_C] \frac{\partial \theta^P}{\partial d} + 2\theta^P(1 - \theta^P) \frac{\partial PS_C}{\partial d}.$$

The first term of the right-hand side reflects the indirect effect of d , which works through its impact on θ^P . Since θ^P is increasing in d (see the proof of Proposition 4), the indirect effect is unambiguously positive. The second term captures the direct effect, where we have

$$\frac{\partial PS_C}{\partial d} = -\frac{d}{\tau} < 0.$$

As such, PS^P (and hence ΔPS) increases with d if the positive indirect effect dominates the negative direct effect. The following result confirms that there is a range of situations where this holds true.

Proposition 6. *If $v - \lambda p > 3d$, PS^P increases with d for a sufficiently large c .*

Proof. Observe that PS^P is increasing in d if

$$[\theta^P PS_S + (1 - 2\theta^P)PS_C] \frac{\partial \theta^P}{\partial d} > \theta^P (1 - \theta^P) \frac{d}{\tau}.$$

With some computation, this condition can be written as

$$\theta^P PS_S + (1 - 2\theta^P)PS_C = \frac{2Mp(PS_S - PS_C) + (2c - p)PS_C}{2c + p(2M - 1)} > \frac{Md}{\tau},$$

which converges to

$$PS_C = \frac{M(v - \lambda p - d)}{2\tau} > \frac{Md}{\tau} \Leftrightarrow v - \lambda p > 3d,$$

as c tends to infinity.

Q.E.D.

The proposition suggests that market distortions, in the form of a higher d , can be a blessing in this environment because they induce the firms to invest more in advertisement. Of course, this welfare improvement is at the expense of the physician who needs to incur more cost, and it will ultimately come down to how we evaluate this welfare loss (which we ignore in the current analysis).

5 A model with free market pricing

We have thus far assumed that the pharmaceutical prices are regulated and fixed exogenously at some predetermined level. Although this is a reasonable assumption in many countries, there is an important exception, namely the US, where pharmaceutical firms can set the prices relatively freely. Here, we extend the baseline analysis to incorporate price setting and illustrate how it would affect the equilibrium allocation.

5.1 Optimal pricing

We now consider a setting where each firm simultaneously chooses both the investment level θ_i and the price p_i . Given the price pair (p_0, p_1) , patient x in the competitive segment ends up with product 0 if

$$v - \tau x - \lambda p_0 > v - \tau(1 - x) - \lambda p_1 \Leftrightarrow \frac{\tau - \lambda(p_0 - p_1)}{\tau} > x.$$

Similarly, patient x in the monopolistic segment ends up with product i if

$$v - |x - i| - \lambda p_i > -d \Leftrightarrow v - \lambda p_i + d > |x - i|.$$

The market demands for firm i are hence given by

$$Q_i^P = \theta_i \theta_{-i} \frac{\tau - \lambda(p_i - p_{-i})}{2\tau} + \theta_i(1 - \theta_{-i})(v - \lambda p_i + d),$$

under product DTCA and

$$Q_i^E = (\theta_i + \theta_{-i} - \theta_i \theta_{-i}) \frac{\tau - \lambda(p_i - p_{-i})}{2\tau},$$

under enlightenment DTCA.

To gain some intuition for the effect of the price competition, it is helpful to look at each segment separately. First, if all patients are selective, each firm maximizes

$$p_i \frac{\tau - \lambda(p_i - p_{-i})}{2\tau},$$

and the (symmetric) optimal price is given by $p_S := \frac{\tau}{\lambda}$. Similarly, if all patients are captive, each firm maximizes

$$p_i(v - \lambda p_i + d),$$

and the optimal price is given by $p_C := \frac{v+d}{2\lambda}$. Naturally, the equilibrium price is bounded between p_S and p_C (and is in fact a weighted average of the two). This implies that free market pricing

has little bite when p_C and p_S are sufficiently close to each other, in which case the equilibrium allocation can be approximated by that under price regulation.

In what follows, we assume that the price competition is intense enough so that the optimal price is lower in the competitive segment than in the monopolistic segment.

Assumption 3. $p_C > p_S \Leftrightarrow v + d > 2\tau$.

5.2 Advertisement expenditures

To analyze the firms' investment problems, we make additional assumptions that are analogous to Assumptions 1 and 2 to simplify the analysis and focus our attention on more relevant cases. First, c is assumed to be sufficiently large, so that the optimal investment is bounded below 1. Second, the value of each product is high enough to ensure that it is beneficial even for the most distant patient. The following assumptions provide sufficient conditions for these two properties.

Assumption 4. $2c > p_C$.

Assumption 5. $2(v - p_C) > 1$.

Under these assumptions, the problem faced by firm i under product DTCA is formulated as

$$\max_{\theta_i \in [0,1], p_i} p_i Q_i^P - \frac{c}{2} \theta_i^2 = p_i \left[\theta_i \theta_{-i} \frac{\tau - \lambda(p_i - p_{-i})}{2\tau} + \theta_i(1 - \theta_{-i})(v - \lambda p_i + d) \right] - \frac{c}{2} \theta_i^2.$$

This yields a pair of first-order conditions:

$$\begin{aligned} \frac{p_i}{c} \left[\theta_{-i} \frac{\tau - \lambda(p_i - p_{-i})}{2\tau} + (1 - \theta_{-i})(v - \lambda p_i + d) \right] &= \theta_i, \\ \theta_i \theta_{-i} \frac{\tau + \lambda p_{-i}}{2\tau} + \theta_i(1 - \theta_{-i})(v + d) &= \theta_i \theta_{-i} \frac{\lambda p_i}{\tau} + 2\lambda \theta_i(1 - \theta_{-i}) p_i. \end{aligned}$$

With abuse of notation, denote by θ^k and p^k the equilibrium investment and price levels, respectively, under regime $k = P, E$. Imposing symmetry, the equilibrium values must solve

$$\theta^P = \frac{2M(p^P)p^P}{2c + p^P[2M(p^P) - 1]}, \quad p^P = \frac{\theta^P \tau + 2\tau(1 - \theta^P)(v + d)}{\lambda[\theta^P + 4\tau(1 - \theta^P)]} = \frac{\theta^P p^C + 4\tau(1 - \theta^P)p^M}{\theta^P + 4\tau(1 - \theta^P)}, \quad (5)$$

where $M(p) := \min\{v - \lambda p + d, 1\}$ to denote its dependence on p . Although the problem is now substantially complicated, we can still show that it always yields a well-defined solution.

Proposition 7. *There exists a unique pair (θ^P, p^P) that satisfies (5). Moreover, θ^P is increasing in c while p^P is increasing, with $\lim_{c \rightarrow 0.5p_S c} \theta^P = 1$, $\lim_{c \rightarrow 0.5p_C} p^P = p_S$, $\lim_{c \rightarrow \infty} \theta^P = 0$, $\lim_{c \rightarrow \infty} p^P = p_C$.*

Proof. *See appendix.*

In contrast, the problem under enlightenment DTCA is quite straightforward and defined as

$$\max_{\theta \in [0,1], p_i} p_i Q_i^E - \frac{c}{2} \theta_i^2 = p_i (\theta_i + \theta_{-i} - \theta_i \theta_{-i}) \frac{\tau - \lambda(p_i - p_{-i})}{2\tau} - \frac{c}{2} \theta_i^2.$$

The first-order conditions are given by

$$\begin{aligned} \frac{p_i(1 - \theta_{-i})[\tau - \lambda(p_i - p_{-i})]}{2\tau c} &= \theta_i, \\ \frac{\tau + \lambda p_{-i}}{2} &= \lambda p_i. \end{aligned}$$

The optimal price is simple and obtained independently of (θ_i, θ_{-i}) , meaning that there is no strategic relationship between pricing and advertisement. The equilibrium values, denoted by θ^E and p^E , are given by

$$\theta^E = \frac{p^E}{2c + p^E} = \frac{\tau}{2\lambda c + \tau}, \quad p^E = p_S = \frac{\tau}{\lambda}.$$

5.3 Welfare implications of free market pricing

Consider a regulatory authority which contemplates to deregulate pharmaceutical pricing by granting firms a higher degree of discretion over pricing. The welfare impact of such a reform depends obviously on the regulated price that has been imposed prior to the deregulation. For the sake of argument, suppose that the regulated price was initially set at $p = p_S$ (the lowest possible equilibrium price), as it is perhaps natural to assume that pharmaceutical prices would rise after the deregulation. Under this circumstance, the equilibrium price jumps up to $p^P > p_S$ under product DTCA while it remains the same under enlightenment DTCA. Welfare implications of free market pricing

ing can then be illustrated by examining how the two welfare measures respond to an exogenous variation in p .

As in the case with price controls, it is first instructive to observe the effect of a change in p on the advertisement expenditures. With some computation, we obtain

$$\begin{aligned}\frac{\partial \theta^P}{\partial p} &= \frac{4Mc + 2pM'(2c - p)}{[2c + p(2M - 1)]^2} = -\frac{\partial \theta^P}{\partial c} \frac{c}{p} + \frac{2pM'(2c - p)}{[2c + p(2M - 1)]^2}, \\ \frac{\partial \theta^E}{\partial p} &= \frac{2c}{(2c + p)^2} = -\frac{\partial \theta^E}{\partial c} \frac{c}{p},\end{aligned}$$

which indicates that the effect of an increase in p mirrors that of a decrease in c , the only difference being that it may also change the market-size distortion under product DTCA. In particular, when $M = 1$ and hence $M' = 0$ (no market-size distortion), an increase in p is equivalent to a decrease in c , thereby allowing us to directly apply the results obtained in the previous section. The reason for this is clear if we look at the firms' objective function: if $M = 1$, the problem under product DTCA can be normalized by p , so that only the price-cost ratio matters for the equilibrium allocation.

When $M < 1$, on the other hand, there is an additional effect on the market-size distortion, and the problem can no longer be normalized by the price. Since $M' = -\lambda < 0$, this makes the effect of an increase in p somewhat weaker. We can still show, however, that the overall effect is positive under some mild conditions. To this end, it suffices to show that $4Mc > 2\lambda p(2c - p)$ for all $c \in [0.5p, \infty)$. This condition clearly holds if c is sufficiently small and close to $0.5p$. As $c \rightarrow \infty$, this becomes

$$M \geq \lambda p \Leftrightarrow \frac{v+d}{2\lambda} \geq p,$$

which generally holds because $\frac{v+d}{2\lambda}$ is the upperbound of the equilibrium price. This argument suggests that the equilibrium advertisement expenditure is generally increasing in p under either regime.

Given this result, we now turn to the equilibrium firm profit. Taking partial derivative with respect to p , we obtain

$$\frac{\partial \pi^P}{\partial p} = \frac{\theta^P}{2} [(2M'p + 2M)(1 - \theta^P) + \theta^P] + [Mp(1 - 2\theta^P) + (p - c)\theta^P] \frac{\partial \theta^P}{\partial p}. \quad (6)$$

The first term of the right-hand side is the direct effect of a price increase which is always positive. An increase in p also induces an indirect effect via an increase in θ^P . Since the firms generally overinvest, this indirect effect is always negative as any further increase in the investment level from its equilibrium level can only lower the firm profit, given the price. Despite this apparent tradeoff, we can show that the direct effect dominates, so that an increase in the price level generally benefits the firms (see Proposition 8 below).

The situation is more complicated for the patient surplus. Taking partial derivative with respect to p , we obtain

$$\frac{\partial PS^P}{\partial p} = \theta^P \left[\theta^P \frac{\partial PS_S}{\partial p} + 2(1 - \theta^P) \frac{\partial PS_C}{\partial p} \right] + [2\theta^P(PS_S - PS_C) + 2(1 - \theta^P)PS_C] \frac{\partial \theta^P}{\partial p}, \quad (7)$$

where

$$\frac{\partial PS_S}{\partial p} = -\lambda p, \quad \frac{\partial PS_C}{\partial p} = -\lambda(M + M'p) - \tau MM'.$$

Again, the first term captures the direct effect which is now negative. The second term is the indirect effect which stems from an increase in θ^P . In contrast to the equilibrium profit, the indirect effect is positive because more investment is always beneficial for patients. The effect of a price increase on the patient surplus is more ambiguous due to this tradeoff but is still positive when the price is relatively low to begin with.

Proposition 8. (i) π^P is increasing in p for all $p \in (0, p_C]$. (ii) PS^P is increasing in p if p is sufficiently small.

Proof. See appendix.

The first part of the proposition implies that the two forms of regulation are complementary in that the benefit of a less stringent advertisement regulation, i.e., product DTCA, is amplified when it is combined with a less stringent price regulation. As such, in the region where $\Delta PS > 0$, product DTCA and concurrent price liberalization are the right policy mix for a pro-business authority. This is, however, not necessarily true for the welfare of patients as they suffer directly from a higher equilibrium price. Still, when the price level is sufficiently low to begin with, this negative effect

is more than offset by an increase in the investment level.¹³ The intuition behind this result is that at a price sufficiently close to zero, the firms make almost no investment, and there are hence no informed patients who directly suffer from a price increase. Although more patients become informed and are forced to incur more cost as the price increases, this effect is of second order and is dominated by the indirect effect which is of first order.

6 Conclusion

This paper investigates the effect of regulating the content of DTCA in a pharmaceutical market with emphasis on the distinction between product and enlightenment DTCA. Owing to a unique information structure of the market, the two different forms of DTCA generate different incentives to the pharmaceutical firms and induce different responses from potential patient. We argue that although product DTCA may distort prescription decisions, this welfare loss is partly offset by stronger incentives to invest in advertisement, which subsequently reduce the fraction of uninformed patients.

The overall welfare impact is ambiguous and depends, among other things, on the cost effectiveness of advertisement and the market-size distortion. When the cost of advertisement is relatively high, the interests of the pharmaceutical firms and potential patients are better aligned, and there often exists a Pareto-improving policy choice: enlightenment DTCA is preferred when the market-size distortion is more severe while product DTCA is preferred when it is less so. As the cost of advertisement decreases, however, a conflict emerges between the firms and patients. In general, product DTCA emerges as the preferred choice for patients as the prescription distortion is more attenuated while serving the educational purpose, but the firms are made worse off due to the overinvestment problem.

References

BAR, T. AND D. R. LILLARD (2014): “Direct to Consumer Advertising of Pharmaceutical Drugs: Information and Persuasion,” Working Paper 19794, National Bureau of Economic Research,

¹³The same reasoning suggests that the patient surplus is increasing in p if c is sufficiently large.

- BRADFORD, W. D., A. N. KLEIT, P. J. NIETERT, AND S. ORNSTEIN (2010): “The Effect of Direct to Consumer Television Advertising on the Timing of Treatment,” *Economic Inquiry*, 48, 306–322.
- BREKKE, K. R. AND M. KUHN (2006): “Direct to Consumer Advertising in Pharmaceutical Markets,” *Journal of Health Economics*, 25, 102–130.
- DAI, C., R. S. STAFFORD, AND G. C. ALEXANDER (2005): “National Trends in Cyclooxygenase-2 Inhibitor Use Since Market Release: Nonselective Diffusion of a Selectively Cost-Effective Innovation,” *Archives of Internal Medicine*, 165, 171–177.
- DAVE, D. AND H. SAFFER (2012): “Impact of Direct-to-Consumer Advertising on Pharmaceutical Prices and Demand,” *Southern Economic Journal*, 79, 97–126.
- DONOHUE, J. M., E. R. BERNDT, M. ROSENTHAL, A. M. EPSTEIN, AND R. G. FRANK (2004): “Effects of Pharmaceutical Promotion on Adherence to the Treatment Guidelines for Depression,” *Medical Care*, 42, 1176–1185.
- FROSCH, D. L., D. GRANDE, D. M. TARN, AND R. L. KRAVITZ (2010): “A Decade of Controversy: Balancing Policy With Evidence in the Regulation of Prescription Drug Advertising,” *American Journal of Public Health*, 100, 24–32.
- GROSSMAN, G. M. AND C. SHAPIRO (1984): “Informative Advertising with Differentiated Products,” *The Review of Economic Studies*, 51, 63–81.
- IZUKA, T. (2004): “What Explains the Use of Direct-to-Consumer Advertising of Prescription Drugs?” *Journal of Industrial Economics*, 52, 349–379.
- IZUKA, T. AND G. Z. JIN (2007): “Direct to Consumer Advertising and Prescription Choice,” *Journal of Industrial Economics*, 55, 771–771.
- KALYANARAM, G. (2008): “The Order of Entry Effect in Prescription (RX) and over-the-Counter (OTC) Pharmaceutical Drugs,” *International Journal of Pharmaceutical and Healthcare Marketing*, 2, 35–46.

- (2009): “The Endogenous Modeling of the Effect of Direct-to-Consumer Advertising in Prescription Drugs,” *International Journal of Pharmaceutical and Healthcare Marketing*, 3, 137–148.
- KRAVITZ, R. L., R. M. EPSTEIN, M. D. FELDMAN, C. E. FRANZ, R. AZARI, M. S. WILKES, L. HINTON, AND P. FRANKS (2005): “Influence of Patients Requests for Direct-to-Consumer Advertised Antidepressants: A Randomized Controlled Trial,” *Journal of American Medical Association*, 293, 1995–2002.
- LIU, Q. AND S. GUPTA (2011): “The Impact of Direct-to-Consumer Advertising of Prescription Drugs on Physician Visits and Drug Requests: Empirical Findings and Public Policy Implications,” *International Journal of Research in Marketing*, 28, 205–217.
- MURRAY, E., B. LO, L. POLLACK, K. DONELAN, AND K. LEE (2003): “Direct-to-Consumer Advertising: Physicians Views of Its Effects on Quality of Care and the Doctor-Patient Relationship,” *Journal of American Board of Family Medicine*, 16, 513–524.
- ROBINSON, A. R., K. B. HOHMANN, J. I. RIFKIN, D. TOPP, C. M. GILROY, J. A. PICKARD, AND R. J. ANDERSON (2004): “Direct-to-Consumer Pharmaceutical Advertising: Physician and Public Opinion and Potential Effects on the Physician-Patient Relationship,” *Archives of Internal Medicine*, 164, 427–432.
- ROSENTHAL, M. B., E. R. BERNDT, J. M. DONOHUE, A. M. EPSTEIN, AND R. G. FRANK (2003): *Demand Effects of Recent Changes in Prescription Drug Promotion*, MIT Press, vol. 6 of *Frontiers in Health Policy Research*, 1–26.
- SINKINSON, M. AND A. STARC (2018): “Ask Your Doctor? Direct-to-Consumer Advertising of Pharmaceuticals,” *The Review of Economic Studies*, 86, 836–881.
- SUFRIN, C. B. AND J. S. ROSS (2008): “Pharmaceutical Industry Marketing: Understanding Its Impact on Womens Health,” *Obstetrical and Gynecological Survey*, 63, 585–596.
- VENTOLA, C. L. (2011): “Direct-to-Consumer Pharmaceutical Advertisement: Therapeutic or Toxic?” *Pharmacy and Therapeutics*, 36, 669–684.

WEISSMAN, J. S., D. BLUMENTHAL, A. J. SILK, M. NEWMAN, K. ZAPERT, R. LEITMAN, AND S. FEIBELMANN
(2004): “Physicians Report On Patient Encounters Involving Direct-To-Consumer Advertising,”
Health Affairs, 23, 219–233.

Appendix. Proofs of Propositions

Proof of Proposition 2

Proof. Define θ^k and Q^k , $k = P, E$, as functions of v , p , d and c . The effects of v and d are identical. Note that Q^P is increasing in v if

$$2c + p(2M - 1) > 2Mp,$$

which for any $c > 0.5p$. This suggests that ΔQ is increasing in v and decreasing in d .

With respect to a change in c , observe that $\Delta Q = \frac{1}{8} > 0$ at $c = 0.5p$ and $\lim_{c \rightarrow \infty} \Delta Q = 0$. It is also straightforward to obtain

$$\begin{aligned} \frac{\partial Q^P}{\partial c} &= [\theta^P + (1 - 2\theta^P)M] \frac{\partial \theta^P}{\partial c} = -\frac{4M^2 p[2c - p(2M - 1)]}{[2c + p(2M - 1)]^3}, \\ \frac{\partial Q^E}{\partial c} &= (1 - \theta^E) \frac{\partial \theta^E}{\partial c} = -\frac{4cp}{(2c + p)^3}. \end{aligned}$$

It follows from this that ΔQ is decreasing in c if

$$\frac{M^2[2c - p(2M - 1)]}{[2c + p(2M - 1)]^3} > \frac{c}{(2c + p)^3}.$$

Alternatively, the condition can be written as

$$F(c)G(c)^3 > \frac{1}{M^2},$$

where

$$F(c) := \frac{2c - p(2M - 1)}{c}, \quad G(c) := \frac{2c + p}{2c + p(2M - 1)}.$$

Evaluated at $c = 0.5p$, the condition is reduced to

$$F(0.5p)G(0.5p)^3 = \frac{4(1 - M)}{M^3} > \frac{1}{M^2} \Leftrightarrow \frac{4}{5} > M.$$

As $c \rightarrow \infty$, on the other hand, we have

$$\lim_{c \rightarrow \infty} F(c)G(c)^3 = 2 > \frac{1}{M^2} \Leftrightarrow M > \sqrt{0.5}.$$

Note also that FG^3 is increasing in c if

$$F'G + 3FG' > 0, \tag{8}$$

where

$$F'(c) = \frac{p(2M-1)}{c^2}, \quad G'(c) = -\frac{4p(1-M)}{[2c + p(2M-1)]^2}.$$

It then follows that (10) can be written as

$$\frac{p(2M-1)(2c+p)}{c^2[2c + p(2M-1)]} > \frac{12p(1-M)[2c - p(2M-1)]}{c[2c + p(2M-1)]^2},$$

which is simplified to

$$\frac{p(2M-1)(2c+p)}{c} > \frac{12p(1-M)[2c - p(2M-1)]}{2c + p(2M-1)}.$$

Note that the left-hand side is strictly decreasing while the right-hand side is strictly increasing. This means that FG^3 may increase at the beginning but must eventually decrease as c gets sufficiently large. Combined with the fact that FG^3 must converge to 2 regardless of M , this fact allows us to pin down the shape of ΔQ . There are three possible cases we need to investigate.

Case 1 ($\sqrt{0.5} > M$): In this case, $FG^3 > M^{-2}$ when c is relatively small. At some point, FG^3 starts decreasing and converges to 2 which is lower than M^{-2} . This means that ΔQ first decreases and then increases, ultimately converging to 0. As such, there exists a unique threshold $c_Q(v, d)$ such that $\Delta Q > 0$ if and only if $c_Q > c$. Since ΔQ is increasing in v and d , the threshold c_Q must also be increasing in v and d .

Case 2 ($0.8 \geq M \geq \sqrt{0.5}$): In this case, FG^3 must invariably be smaller larger than M^{-2} , which implies that ΔQ is monotonically decreasing with $\Delta Q > 0$ for all $c > 0.5p$.

Case 3 ($1 > M > 0.8$): In this case, FG^3 is lower than M^{-2} when c is relatively small. As c gets larger, it increases but then decreases and converges to 2 which is larger than M^{-2} . This means that ΔQ first increases and then decreases with $\Delta Q > 0$ for all $c > 0.5p$.

Q.E.D.

Proof of Proposition 4

Proof. Since

$$\frac{\partial \Delta C}{\partial c} = 2\theta^P \frac{\partial \theta^P}{\partial c} - 2\theta^E \frac{\partial \theta^E}{\partial c},$$

we have

$$\begin{aligned} \frac{\partial \Delta \pi}{\partial c} &= p[M - (2M - 1)\theta^P] \frac{\partial \theta^P}{\partial c} - p(1 - \theta^E) \frac{\partial \theta^E}{\partial c} - c\theta^P \frac{\partial \theta^P}{\partial c} + c\theta^E \frac{\partial \theta^E}{\partial c} - \frac{\Delta C}{2} \\ &= (c\theta^P - Mp) \frac{\partial \theta^P}{\partial c} - c\theta^E \frac{\partial \theta^E}{\partial c} - \frac{\theta^{P^2} - \theta^{E^2}}{2}. \end{aligned}$$

Observe that

$$\frac{\theta^{P^2}}{2} = -\frac{\partial \theta^P}{\partial c} \frac{Mp}{2}, \quad \frac{\theta^{E^2}}{2} = -\frac{\partial \theta^E}{\partial c} \frac{p}{4},$$

which gives us

$$\begin{aligned} \frac{\partial \Delta \pi}{\partial c} &= \left(c\theta^P - \frac{Mp}{2}\right) \frac{\partial \theta^P}{\partial c} - \left(c\theta^E + \frac{p}{4}\right) \frac{\partial \theta^E}{\partial c} \\ &= \frac{Mp[2c - p(2M - 1)]}{2[2c + p(2M - 1)]} \frac{\partial \theta^P}{\partial c} - \frac{p(6c + p)}{4(2c + p)} \frac{\partial \theta^E}{\partial c} \\ &= -\frac{2(Mp)^2[2c - p(2M - 1)]}{[2c + p(2M - 1)]^3} + \frac{p^2(6c + p)}{2(2c + p)^3}. \end{aligned}$$

From this, $\Delta \pi$ is increasing in c if

$$\frac{1}{M^2} > \frac{4[2c - p(2M - 1)]}{6c + p} \left(\frac{2c + p}{2c + p(2M - 1)} \right)^3 = H(c)G(c)^3, \quad (9)$$

where

$$H(c) := \frac{4[2c - p(2M - 1)]}{6c + p}.$$

As above, HG^3 is increasing in c if $H'G + 3HG' > 0$. With some computation, we obtain

$$H'(c) = \frac{16p(3M - 1)}{(6c + p)^2},$$

so that this condition becomes

$$\frac{16p(3M - 1)(2c + p)}{(6c + p)^2[2c + p(2M - 1)]} > \frac{48p(1 - M)[2c - p(2M - 1)]}{[2c + p(2M - 1)]^2(6c + p)},$$

which is further simplified to

$$\frac{(3M - 1)(2c + p)}{(6c + p)} > \frac{3(1 - M)[2c - p(2M - 1)]}{[2c + p(2M - 1)]}. \quad (10)$$

Note that the right-hand side is decreasing in c while the left-hand side is increasing.

Evaluated at $c = 0.5p$, (10) becomes

$$\frac{3M - 1}{3(1 - M)} > \frac{2(1 - M)}{M}.$$

This holds if and only if $M > \frac{2}{3}$. In the limit as $c \rightarrow \infty$, on the other hand, (10) becomes

$$\frac{3M - 1}{3(1 - M)} > 3,$$

which holds if and only if $M > \frac{5}{6}$. This means that: if $M > \frac{5}{6}$, HG^3 is increasing for all $c \in [0.5p, \infty)$; if $\frac{5}{6} > M > \frac{2}{3}$, HG^3 first increases and then decreases.

To complete the proof, we now check the boundary conditions for (9). It is easy to verify that (9) holds at $c = 0.5p$ if $M > \frac{2}{3}$, so that $\Delta\pi$ increases at the beginning. At the other end, as $c \rightarrow \infty$, this holds if $\sqrt{0.75} > M$. Note also that $\sqrt{0.75} \approx 0.866 > \frac{5}{6}$. This means that for $M > \sqrt{0.75}$, $\Delta\pi$ increases at first, then decreases and converges to 0, proving that there exists a threshold $c_\pi(v, d)$ such that $\Delta\pi > 0$ if and only if $c > c_\pi(v, d)$. For $\sqrt{0.75} \geq M \geq \frac{5}{6}$, on the other hand, $\Delta\pi$ is

increasing for all c , and hence $\Delta\pi < 0$ for all c . We can also show that $\Delta\pi$ is increasing in v . Since a decrease in v is equivalent to a decrease in M with p and d fixed, this means that $\Delta\pi < 0$ for all c if $\sqrt{0.75} > M$.

Finally, since $\Delta\pi$ is increasing in v , it is clear that c_π is decreasing in v . To show that $\Delta\pi$ is increasing in d , recall that the profit under product DTCA is given by

$$\pi^P = \frac{p\theta^P[2M - (2M - 1)\theta^P]}{2} - \frac{c}{2}\theta^{P^2},$$

from which we obtain

$$\begin{aligned} \frac{\partial\pi^P}{\partial d} &= 2[Mp^P(1 - 2\theta^P) + (p^P - c)\theta^P]\frac{\partial\theta^P}{\partial d} + p^P\theta^P(1 - \theta^P) \\ &= -(2M - 1)p^P\theta^P\frac{\partial\theta^P}{\partial d} + p^P\theta^P(1 - \theta^P). \end{aligned}$$

Note also that

$$\frac{\partial\theta^P}{\partial d} = \frac{2p(2c - p)}{[2c + p(2M - 1)]^2} > 0.$$

Given this, π^P is increasing in d if

$$1 - \theta^P > \frac{2p(2M - 1)(2c - p)}{[2c + p(2M - 1)]^2} \Leftrightarrow 2c > p(2M - 1),$$

which holds for any $c > 0.5p$.

Q.E.D.

Proof of Proposition 5

Proof. Observe that $\Delta PS = \frac{PS_S}{4} > 0$ at $c = 0.5p$ and $\lim_{c \rightarrow \infty} \Delta PS = 0$. It is also straightforward to obtain

$$\begin{aligned} \frac{\partial PS^P}{\partial c} &= 2[\theta^P PS_S + (1 - 2\theta^P)PS_C]\frac{\partial\theta^P}{\partial c} = -\frac{8Mp[2Mp(PS_S - PS_C) + (2c - p)PS_C]}{[2c + p(2M - 1)]^3}, \\ \frac{\partial PS^E}{\partial c} &= 2(1 - \theta^E)PS_S\frac{\partial\theta^E}{\partial c} = -\frac{8pc}{(2c + p)^3}PS_S. \end{aligned}$$

It follows from this that ΔPS is decreasing in c if

$$\frac{M^2 [2c\phi - p[(2M+1)\phi - 2]]}{[2c + p(2M-1)]^3} > \frac{c}{(2c+p)^3}. \quad (11)$$

Here, we extend the definition of F in the proof of Proposition 2 and, with slight abuse of notation, define

$$F(c; \phi) = \frac{2c\phi - p[(2M+1)\phi - 2]}{c},$$

so that (11) is given by

$$F(c; \phi)G(c)^3 > \frac{1}{M^2}.$$

Note that $F(c; 1) = F(c)$.

Evaluated at $c = 0.5p$, (11) is reduced to

$$F(0.5p; \phi)G(0.5p) = \frac{4(1 - M\phi)}{M^3} > \frac{1}{M^2} \Leftrightarrow \frac{4}{1 + 4\phi} > M.$$

As $c \rightarrow \infty$, on the other hand, it converges to

$$\lim_{c \rightarrow \infty} F(c; \phi)G(c) = 2\phi > \frac{1}{M^2} \Leftrightarrow M > \sqrt{0.5\phi^{-1}}.$$

Moreover, as in Proposition 2, we can show that FG^3 is increasing in c if

$$\frac{p[(2M+1)\phi - 2](2c+p)}{c^2[2c + p(2M-1)]} > \frac{12p(1-M)[2c - p[(2M+1)\phi - 2]]}{c[2c + p(2M-1)]^2},$$

which is simplified to

$$\frac{p[(2M+1)\phi - 2](2c+p)}{c} > \frac{12p(1-M)[2c - p[(2M+1)\phi - 2]]}{2c + p(2M-1)},$$

If $(2M+1)\phi \leq 2$, FG^3 is weakly decreasing in c . If $(2M+1)\phi > 2$, we can essentially follow the same argument as in Proposition 2. In either case, we can conclude that: (i) if $M \geq \sqrt{0.5\phi^{-1}}$,

$\Delta PS > 0$ for all $c \in (0.5p, \infty)$; (ii) if $\sqrt{0.5\phi^{-1}} > M$, there exists an interior threshold c_{PS} such that $\Delta PS > 0$ if and only if $c_{PS} > c$.

Finally, it is easy to verify that PS^P is increasing in v while PS^E is independent of it. That c_{PS} is increasing in v immediately follows from this fact.

Q.E.D.

Proof of Proposition 7

Proof. Define

$$S(p) := \frac{2M(p)p}{2c + p[2M(p) - 1]} = \frac{2(v+d)p - 2\lambda p^2}{2c - p[2(v+d) - 1] - 2\lambda p^2}, \quad P(\theta) := \frac{\theta p_S + 4\tau(1-\theta)p_C}{\theta + 4\tau(1-\theta)}.$$

It is clear that S is upward sloping in p if $M(p) = 1$. If $M(p) < 1$, we have

$$S(p) = \frac{2(v+d)p - 2\lambda p^2}{2c - p[2(v+d) - 1] - 2\lambda p^2}.$$

Given this, S is upward sloping in p if

$$S'(p) = \frac{2(v+d-2\lambda p)(2c-p) + 2(v+d-\lambda p)p}{[2c - p[2(v+d) - 1] - 2\lambda p^2]^2} > 0.$$

This condition can be written as

$$(v+d-2\lambda p)(2c-p) + (v+d-\lambda p)p > 0,$$

which holds for any $p \in [0, p_C]$. Similarly, P is downward sloping in θ if

$$\begin{aligned} P'(\theta) &= \frac{(p_S - 4\tau p_C)[\theta + 4\tau(1-\theta)] - (1-4\tau)[\theta p_S + 4\tau(1-\theta)p_C]}{[\theta + 4\tau(1-\theta)]^2} \\ &= -\frac{4\tau(p_C - p_S)}{[\theta + 4\tau(1-\theta)]^2} < 0, \end{aligned}$$

which clearly holds. This shows that there is always a unique pair that satisfies (5).

It is also clear that S shifts downward as c increases while P is not affected. This means that θ^P is decreasing in c while p^P is increasing. Moreover, it is easy to verify that $\theta^P \rightarrow 1$ and $p^P = p_S$ as $c \rightarrow 0.5p_S$ and $\theta^P \rightarrow 0$ and $p^P = p_C$ as $c \rightarrow \infty$.

Q.E.D.

Proof of Proposition 8

Proof. By rearranging (6), we obtain

$$\frac{\partial \pi^P}{\partial p} = \frac{\theta^P}{2} \left[(2M'p + 2M)(1 - \theta^P) + \theta^P - 2Mp \frac{\partial \theta^P}{\partial p} \right].$$

Since

$$\frac{\partial \theta^P}{\partial p} = \frac{4Mc + 2M'p(2c - p)}{[2c + p(2M - 1)]^2},$$

π^P is increasing in p if

$$2M'p \left[1 - \theta^P - \frac{2Mp(2c - p)}{[2c + p(2M - 1)]^2} \right] + 2M(1 - \theta^P) + \theta^P - \frac{8M^2pc}{[2c + p(2M - 1)]^2} > 0.$$

Substituting θ^P yields

$$2M'p \left[\frac{2c - p}{2c + p(2M - 1)} - \frac{2Mp(2c - p)}{[2c + p(2M - 1)]^2} \right] + \frac{4Mc}{2c + p(2M - 1)} - \frac{8M^2pc}{[2c + p(2M - 1)]^2} > 0,$$

which is simplified to

$$M'p(2c - p) + 2Mc > 0.$$

When $M = 1$, $M' = 0$ and this condition clearly holds. When $M < 1$, we have

$$2Mc > \lambda p(2c - p). \tag{12}$$

This condition holds at $p = 0$. Note also that the right-hand side is decreasing in p at first and then becomes increasing while the left-hand side is strictly decreasing. This means that (12) holds for all $p \in (0, p_C]$ if it holds at $p = p_C$. This is the case if

$$2 \left(v - \lambda \frac{v + d}{2\lambda} + d \right) c = (v + d)c > \frac{v + d}{2} \left(2c - \frac{v + d}{2\lambda} \right),$$

which holds for any c .

Similarly, it follows from (7) that PS^P is increasing in p if

$$[2\theta^P(PS_S - PS_C) + 2(1 - \theta^P)PS_C] \frac{\partial \theta^P}{\partial p} > \theta^P [\lambda p \theta^P + 2(1 - \theta^P)[\lambda(M + M'p) + \tau MM']],$$

where

$$PS_S - PS_C = (v - \lambda p)(1 - M) + \frac{\tau}{4}(2M^2 - 1).$$

With some computation, we obtain

$$\begin{aligned} & [2Mp(PS_S - PS_C) + (2c - p)PS_C] \frac{2Mc + M'p(2c - p)}{2c + p(2M - 1)} \\ & > Mp [\lambda Mp^2 + (2c - p)[\lambda(M + M'p) + \tau MM']], \end{aligned}$$

As $p \rightarrow 0$, this condition is reduced to

$$\frac{4Mc^2PS_C}{2c + p(2M - 1)} > 0,$$

which always holds.

Q.E.D.

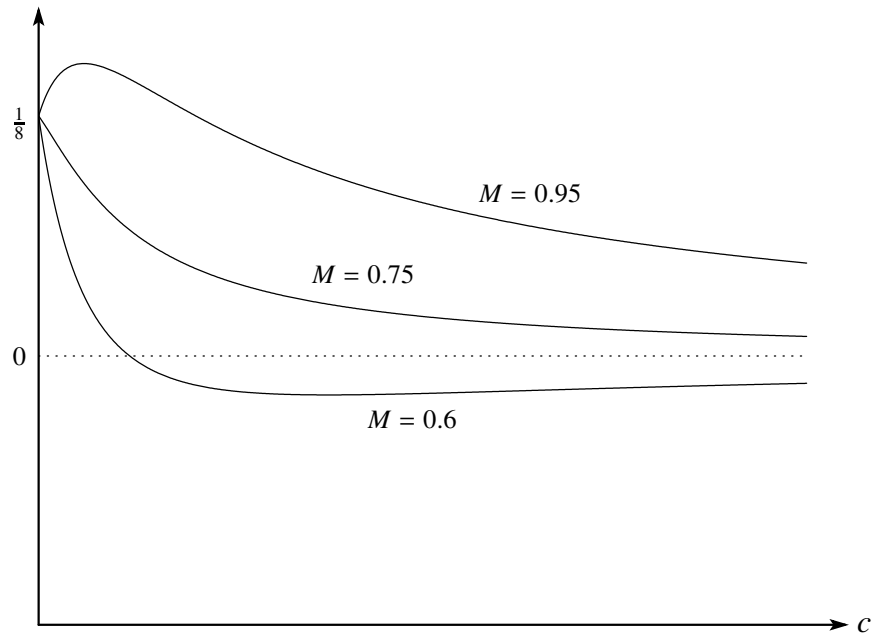


Figure 1: The difference in market demand ΔQ ($p = 0.6$)

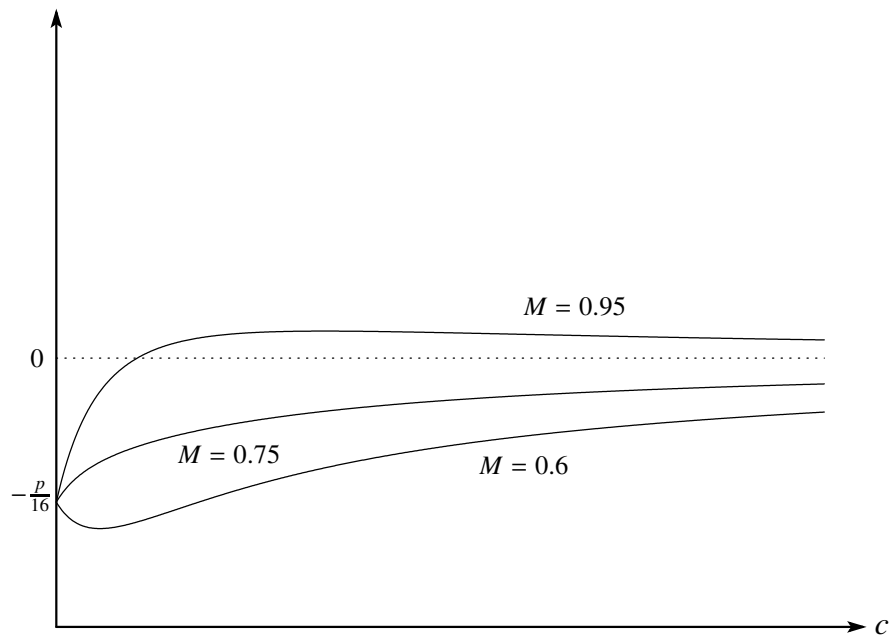


Figure 2: The difference in firm profit $\Delta \pi$ ($p = 0.6$)

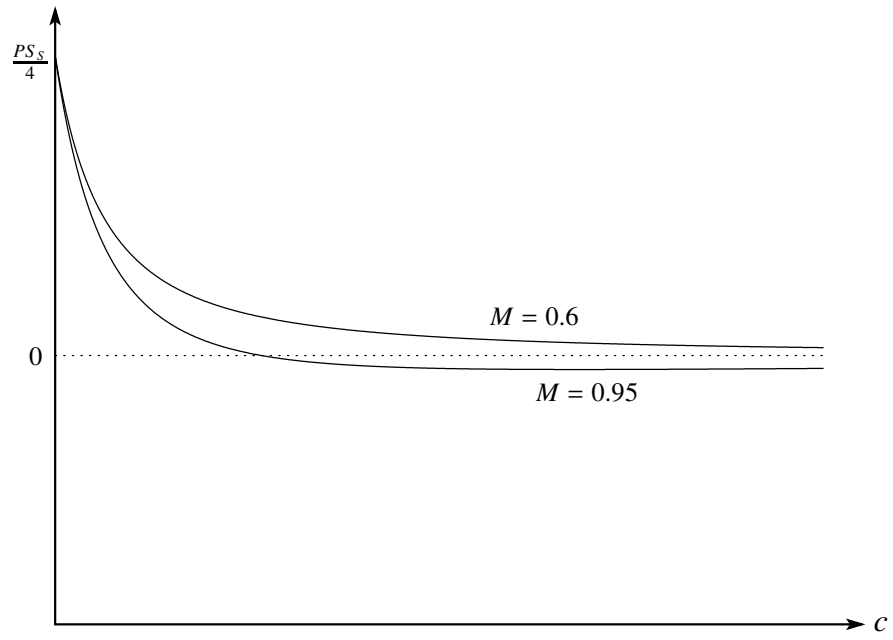


Figure 3: The difference in patient surplus ΔPS ($v = 1.1, p = 0.6, \tau = \lambda = 1$)

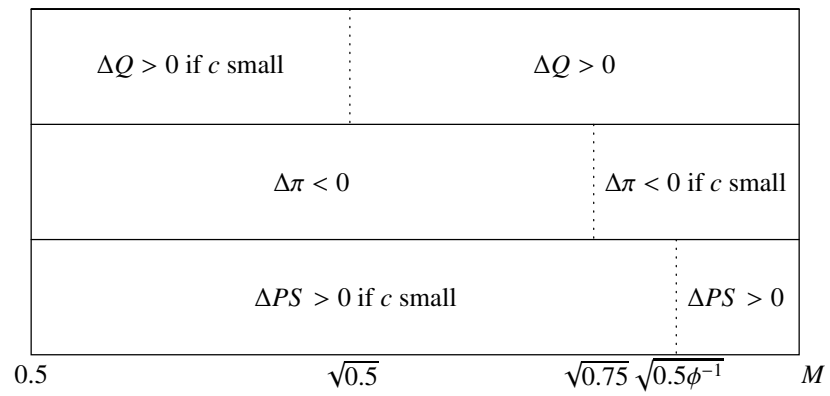


Figure 4: The welfare effects of the content regulation: summary