

The Impact of Academic Patenting on the Rate, Quality, and Direction of (Public) Research Output

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Abstract

We examine the influence of faculty patenting activity on the rate, quality, and content of public research outputs in a panel dataset spanning the careers of 3,862 academic life scientists. Using inverse probability of treatment weights (IPTW) to account for the dynamics of self-selection into patenting, we find that patenting has a positive effect on the rate of publication of journal articles, but no effect on the quality of these publications. Using several measures of the “patentability” of the content of research papers, we also find that patenters may be shifting their research focus to questions of commercial interest. We conclude that the often-voiced concern that patenting in academe has a nefarious effect on public research output is, at least in its simplest form, misplaced.

Keywords: patenting, economics of science, technology transfer, self-selection.

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1 Introduction

In the past few decades, universities and other public-sector research organizations have become much more proactive in their efforts to commercialize scientific discoveries (see Henderson et al., 1998; Jaffe and Lerner, 2001; Jensen and Thursby, 2001; Mowery et al., 2001; Thursby and Thursby, 2002). Underlying this well documented upswing in university patenting has been a sharp increase in the number of individual academic scientists who are listed as inventors on patents. As its incidence has increased, however, academic patenting has generated intense controversy, much of which has centered on the long-term effect of patenting on the development of future scientific knowledge.

At this juncture, every available indicator suggests that a growing number of university faculty will become involved in the commercialization of scientific research. As the literature shifts to evaluating the consequences of faculty patenting for the traditional research process, a number of questions will require investigation. In this paper, we focus on two issues that have come to the fore in debates about academic patenting. First, in what direction and to what degree does faculty patenting affect the rate of production of public scientific outputs? Second, does patenting directly influence either the quality or the content of the subsequent-to-the-patent research performed by the scientist?

These questions are important and, we believe, largely unresolved. On one hand, surveys of academic scientists have found that patenting skews scientists' research agendas toward commercial priorities, causes delay in the public dissemination of research findings, and crowds out effort devoted to producing public research (Blumenthal et al., 1996; Campbell et al. 2002; Krinsky, 2003). In stark terms, this work has portrayed a tradeoff between patenting and the progress of academic science. On the other hand, the few studies that have econometrically assessed the scientist-level relationship between patenting and publishing have come to a very different conclusion. Agrawal and Henderson (2002) estimated fixed-effect regressions of the effect of patenting in a 15-year panel of 236 scientists in two MIT departments. They found that patenting did not affect publishing rates. Markiewicz and DiMinin (2005) constructed a sample of 166 academic patenters that were matched to an

equivalent number of non-patenting scientists. In a fixed effects specification, they found a statistically positive effect of researchers' patent stocks on their publication counts. In a third study, Stephan et al. (forthcoming) exploited a survey of doctorate recipients and used instrumental variables to estimate the cross-sectional relationship between patenting and publishing; they found that patenting and publishing relate positively.

Our findings concur with — and significantly extend — this latter set of results. With careful adjustment for selection into patenting, we find that both the flow and the stock of scientists' patents are positively related to subsequent publication rates. Moreover, this increase in output does not come at the expense of the quality of the published research: we find that the average quality of publications is unaffected by patenting. However, we present three distinct pieces of evidence which indicate that patenting induces a moderate shift in the content of scientists' research. First, faculty holding patents are more likely to coauthor papers with researchers in firms. Second, patenters' publications more frequently appear in journals that have a higher proportion of company-affiliated authors. Finally, we develop a measure of the latent “patentability” of research based on the title keywords of articles and find it to be significantly higher in the subsequent-to-the-patent papers of patenting scientists.

At a minimum, we interpret our results as refuting the simple form of the claim that academic patenting has a deleterious effect on the production of public science. However, our findings do confirm that patenting has had real effects on the direction of scientific progress. Although it is legitimate to ask whether the continued migration of commercial interests into universities will further induce scientists to select research projects on the basis of their perceived value in the private sector, assessing the welfare implications of this change will require a more refined understanding of the relationship between research outputs that are “applied” (*i.e.*, less likely to become an important foundation for subsequent scientific research) versus those that are “patentable” (*i.e.*, focused on questions of industrial usefulness). In the context of the life sciences, for example, it is not a priori clear that there is a trade-off between the academic influence and the patentability of a research project (see Stokes, 1997).

In addition to presenting findings pertinent to an ongoing policy debate, our study makes two other contributions. First, we have assembled a comprehensive, longitudinal dataset: it is a prospective, 3,862-person random sample drawn from the population of life scientists in academia between 1967 and 1999. For the individuals in the sample, we have reconstituted entire career histories, including patent and publication information, as well as many employer-level variables. We believe that this is the most inclusive dataset available for assessing the relationship between patenting and public research productivity among academic scientists.

Second, we attempt to disentangle correlation from causality in the assessment of the effect of patenting. As we will show, patent holders differ from other researchers on many observable characteristics (see also Stephan et al., forthcoming). More accomplished researchers are much more likely to patent, and controlling for the stock of past publications, scientists with a recent good run are also more likely to patent. This evidence calls into question the ability of traditional fixed-effect specifications to consistently estimate causal effects, since patenters and non-patenters do not appear to follow similar trends in publication rates *before* the initiation of patenting. We use Inverse Probability of Treatment Weighted (IPTW) estimation (Robins et al., 2000; Hernán et al., 2001) to account for the dynamics of self-selection of researchers into patenting. This methodology, which generalizes the propensity score to settings in which treatment is staggered over time, accounts for selection into patenting on the basis of observable characteristics, including (in our case) lagged productivity and the latent patentability of a scientist’s research trajectory. While this approach naturally cannot rule out selection based on unobservable factors, we were able to generate an extensive list of covariates to model the probability of selection into patenting.

In addition to these two primary contributions, the paper indirectly relates to the literature on the tension between applied and basic research (Cohen and Levinthal, 1989; Rosenberg, 1990; Henderson and Cockburn, 1994; Ding, 2005). This group of studies has sought to understand why for-profit firms fund basic research. It has generally concluded that basic and applied research are complements, although several distinct mechanisms are

at play, and not all of them correspond to true spillovers of knowledge across activities (Stern, 2004). This work bears an obvious similarity to our effort to assess the nature of the relationship between basic and commercial scientific projects conducted by individual scientists.

The rest of the paper proceeds as follows. In the next section, we provide an overview of the controversies surrounding academic patenting. Section 3 presents our econometric methodology. Section 4 describes the construction of the sample and data sources, presents descriptive statistics, and reports our econometric results. Section 5 concludes.

2 Basic, Applied, and Commercializable Research: Where Do We Stand?

Both the current level and the trend line for academic patenting leave little doubt that the contemporary research university has become a locus of commercially-oriented innovation. However, this development is not without controversy; many observers have decried the emergence of academic patenting and other forms of commercial science for its potentially adverse effects on the further advancement of science (Krimsky, 2003). Among critics' concerns, the most fundamental revolves around the potential effect of academic patenting on the traditional incentives in science. It is commonly acknowledged that the reward system in science is rooted in peers' acknowledgment of important research advances (Merton, 1973; Dasgupta and David, 1994). Scientists' incentives to create and quickly publish research findings are clear when promotions, salary increases, and professional accolades are awarded on the basis of contributions to the corpus of public scientific findings. Seen in this light, the relevant question about university patenting becomes, to what degree does the availability of the option to patent alter the incentive or ability of scientists to contribute public (i.e., non-excludable) advances to the scientific literature?

We are not persuaded that the presence of the option to patent meaningfully decreases scientists' incentives to invest in the production of public science. In fact, for two reasons we foresee the possibility that patenting and publishing are complementary activities. First, an

academic researcher's scientific reputation is his/her most important currency in the effort to capitalize on intellectual property in the market for university-originated technology. Second, with respect to the production of new scientific knowledge, there are likely to be (intra-person) scope economies that emerge when a scientist is involved in the development of both academic and commercial science. As we describe below, there are, though, more compelling reasons to believe that the emergence of academic patenting has and will continue to cause a shift in the *content* of scientists' research.

Scientist reputation, patents, and the market for university inventions. Scientists are thought to be strongly motivated by the intrinsic satisfaction of solving vexing problems (Stern, 2004), the search for fame and status (Merton, 1973 [1942]), and the up-or-out promotion rule inherent to the tenure system (Carmichael, 1988). How does patenting influence these traditional incentives to produce academic research? While the direct effect of intellectual property rights is probably small, academic patenting could still influence incentives in subtle ways. In particular, the market for university inventions is rife with asymmetric information. As the literature frequently notes, academic discoveries often require years of additional development to yield marketable products; there is likely to be a great deal of uncertainty surrounding the commercial and scientific merit of discoveries at this primitive stage; and exhaustive due diligence regarding the value of a discovery is costly.

Because of these information problems, we argue that scientists' reputations are essential in the market for university technology. By acting as a signal of invention quality, the prominence of a patenting faculty in the community of science diminishes the search and screening costs that potential licensees must incur in the process of identifying promising university technology. Furthermore, university technology transfer officers are aware of the certification role of scientific eminence. Other things equal, because the discoveries of prominent scientists are more marketable in industry, TTOs are more likely to choose to file for patents on the discoveries of high-status scientists.¹ Therefore, the ex post search, screening,

¹Along these lines, Shane and Khurana (2003) showed that startup firms are more likely to be founded to capitalize on university technology if the intellectual property was created by full professors. Elfenbein (2004) found that discoveries made by scientists with extensive publication records were more likely to find a licensing partner.

and contracting problems in the market for ideas increase faculty's ex ante incentives to maintain their reputation on the scientific labor market, as doing so enhances both the odds of finding an industrial match for their inventions, and the value of their patents conditional on a match.²

Non-pecuniary and pecuniary spillovers. A likely consequence of applying for a patent is that academic scientists become acquainted with researchers in companies. As these acquaintances develop into relationships, we expect that industry contacts might become sources of ideas for new research projects. The notion that connections with researchers in industry serve as fruitful sources for unearthing interesting research questions emerges in Agrawal and Henderson's (2002) interviews with MIT scientists. In addition, there is a natural analogy to the complementarities observed between applied and basic research in industrial firms. Rosenberg (1998), for example, documented that innovations born out of contact with commercial enterprises in the applied field of chemical engineering ushered a new era of basic discoveries in chemistry. The possibility of within-scientist economies of scope is also consistent with evolutionary theories of technological and scientific progress in which major advances are understood to represent insightful combinations of disparate pieces of knowledge (e.g., Hull, 1988; Weitzman, 1998). Insofar as access to diverse information facilitates the development of new and fruitful lines of scientific inquiry, patenting may facilitate the creation of the professional ties that productively broaden researchers' information networks.

Knowledge is not the only input to the research process that may transcend the university-industry divide; it is also possible to envisage pecuniary spillovers between patenting and

²Some critics would counter that there is an automatic tradeoff between patenting and publishing because it is time consuming to disclose inventions and flesh out patent applications. In addition, crowding out would occur if faculty members devote a substantial block of time to conduct the research that leads to patentable discoveries. However, two facts mitigate these concerns. First, scientists are assisted in the patent application process by their university's technology transfer office, whose existence enables a division of labor between invention and commercialization activities (Hellman, 2005). Second, qualitative evidence suggests that patent applications are often byproducts of traditional scientific efforts, and that patents and scientific articles routinely encode related pieces of knowledge. For example, in her study of tissue engineering, Murray (2002) shows that many scientists choose the path of dual-knowledge disclosure, a practice whose output she labels "paper-patent pairs." We therefore doubt the claim that patenting necessarily requires substantial amounts of time and always crowds out traditional scientific research (also see Thursby et al., 2005).

publishing. Useful commercial discoveries often lead to industrial sources of funding for the laboratory of the patenting scientist. Even without access to new pools of knowledge, the ability to hire additional post-doctoral scientists or graduate students might result in higher scientific output for a scientist's lab. A related point is that many seminal scientific achievements have been made possible only by technological advances in instrumentation. In the biomedical fields and other areas of science, technological and scientific advances are therefore interdependent: new understandings are often beholden to progress in instrumentation. If patenting scientists are more likely to be in a position to negotiate access to state-of-the-art equipment in corporate laboratories (Owen-Smith and Powell, 2001a), or if they are more likely to have developed the technical expertise to understand and modify research equipment, complementarities between the capital stock of their laboratory and that of their industrial partners might also increase publication output.³

Patenting and the *direction* of scientific advance. While we expect to find that patenting has at worst a neutral effect on the rate and quality of scientists' publication output, there is also a case to be made that it will influence the content of the output.

In formulating this argument, it is useful to begin with an over-simplified description of the controversy surrounding the commercialization of university science. Suppose that there are two types of academic scientists: purists, who disapprove of commercial encroachments into the university and select research topics solely on the basis of scientific merit, and commercialists, who participate in university patenting and frequently associate with firms in industry. Scientists in this latter camp investigate two kinds of research questions: like purists, they explore issues of basic scientific relevance. In addition, they allocate some fraction of their time to investigating discoveries with patentable, commercial application. Although this characterization may exaggerate the actual level of difference between purists and commercialists in some institutions, Owen-Smith and Powell (2001b) present qualitative

³Note that whether the relevant spillovers are technological or pecuniary, it is not the act of seeking intellectual property rights that, in itself, changes the nature and quantity of output produced by a scientist. Rather, patenting, by making the scientist's research visible to new constituencies, will lead to collaborations (intellectual or financial) that would not have occurred in the absence of the patent application, and between individuals with potentially complementary scientific backgrounds or access to non-overlapping social networks. It should be clear that any spillovers of this type will arise over time, not contemporaneously.

evidence that there is in fact a division along these lines in many academic departments: traditional scientists who, like Nobel Prize winner John Sulston, oppose the convergence of academe and commerce represent the purist pole, and serial patenters and entrepreneurs constitute the other (Sulston, 2003).

If this characterization is approximately accurate, scientists that choose to patent and thereby shift into the commercialist camp will begin to allocate their research time across a different, wider set of research questions than they had done when they were purists. Once a scientist accepts the label of commercialist, we can expect a within-person change such that a scientist will be more likely to pursue projects for which part of the pay-off for conducting the research will be a patent or some other form of commercial recognition. We do not anticipate that all or even a majority of a scientist's work will shift, but rather that some share of it will be focused on new (to the scientist) research questions. Thus, we expect to discover that patenting is associated with a shift in scientists' focus toward exploring scientific questions with commercial application.

A second and possibly more meaningful mechanism for why patenting may result in a shift in scientists' research foci relates to our previous assertion that patents are a form of translational publication that facilitates the formation of relationships between academic scientists and members of the industrial research community. Through the university's efforts to commercialize their technologies, patenting scientists gain visibility in industry circles. As this visibility leads to associations with researchers in corporate laboratories, academic scientists become interested in and familiar with scientific questions of central importance to industry. These contacts presumably expose university scientists to new (relative to their previous work) areas of commercially-useful scientific inquiry. As we have argued above, *ceteris paribus*, exposure to new and diverse information may enhance scientists' productivity. In addition, one should expect the nature of the work that arises after a scientist bridges the university-industry divide to be different from what preceded it, if only

because academic scientists are likely to become intrigued by questions of interest to industry researchers.⁴

We will remain agnostic regarding the welfare implications of this potential change in research agenda. Among critics of the increasing dependence of universities on private-sector funding, distortion in the choice of topics is a frequently assumed and vigorously lamented consequence. An implicit assumption of these policy discussions is that engagement with the world of commerce will necessarily produce applied research, in the sense that future generations of researchers will be less likely to build on it because of its narrow focus. Increasingly, however, scholars of technological change recognize that ideas might simultaneously have high scientific value and important commercial potential (Stokes, 1997). In the conclusion, we briefly discuss how the present study could be expanded to adjudicate between the optimistic and pessimistic interpretations of the results we present below.

3 Econometric Considerations

Estimating the causal effect of academic patenting on research output must confront a basic selectivity problem: researchers choose whether, when, and how much to patent. As a result, traditional econometric techniques, which assume that exposure to “treatment” occurs randomly, cannot recover causal effects. Past researchers have used two types of methodological approaches to deal with this selection problem: instrumental variables techniques and fixed effects estimation. Using a random cross-section of academic scientists, Stephan et al. (forthcoming) implement the first of these approaches, but the instruments they use — characteristics of the scientist’s university and characteristics of a peer group of scientists — are unlikely to be legitimately excluded from their second stage. Markiewicz and DiMinin (2005) use a fixed effects specification in a panel dataset of matched patenting and

⁴Reliable evidence of a shift in research priorities is still scant. The most systematic data come from Blumenthal et al. (1986). They surveyed academic life scientists, asking whether respondents had considered commercial potential when choosing research projects. 30% of life science faculty with industry funding replied affirmatively, compared to just 7% of faculty without private sector funding. This correlation suggests that industry funding (often associated with patenting) skews scientists’ research agenda, but the causality could just as easily flow in reverse, from researchers’ interests to funding sources.

non-patenting researchers. In so doing, they purge their estimates from any influence of unobserved heterogeneity that is constant over time. However, it is well-known that for difference-in-differences estimation to be valid, it must be the case that the average outcome for the treated and control groups would have followed parallel paths over time in the absence of treatment. This assumption is implausible if pretreatment characteristics that are thought to be associated with the dynamics of the outcome variable are unbalanced between treatment and control units. Below, we provide strong evidence that selection into patenting is influenced by transitory shocks to scientific opportunities. In this respect, estimating the causal effect of academic patenting on research output presents similar challenges to that of estimating the effect of a job training program on wages. In the job training example, treated individuals have lower earnings on average (relative to their pre-treatment average) in the year immediately preceding enrollment into the program; therefore, the fixed effects estimator is likely to overestimate the treatment effect. Conversely, we will show that patenting scientists have higher output (relative to their average in the pre-patenting regime) in the year immediately preceding their first patent application; as a result, the fixed effect estimator is likely to *underestimate* the effect of patenting on publishing rates.

To overcome these challenges, we make use of a novel approach that has recently gained acceptance in biostatistics: Inverse Probability of Treatment Weighted (IPTW) estimation (Robins et al., 2000; Hernán et al., 2001). These estimators are akin to propensity-score matching techniques (Rosenbaum and Rubin, 1983; Dehejia and Wahba, 2002) in that they make the (untestable) assumption that selection into treatment is based on variables that are observable to the econometrician, but extend it to the case of time-varying treatments. In particular, IPTW estimation allows one to recover average treatment effects even in the presence of time-varying confounders, i.e., time-varying variables that (1) are correlated with future values of the dependent variable; (2) predict selection into treatment; and (3) are themselves predicted by past treatment history. As we will show below, this applies to the case of academic patenting, since publication rates are strongly auto-correlated, the probability of patenting increases after a recent flurry of publications, and past patenting history influences future publication rates.

Consider a study in which treatment decisions are made in $T + 1$ distinct periods $0, 1, \dots, T$. At each time t , for each individual i , “prognostic factors” Z_{it} and an outcome of interest y_{it} are measured, and a discrete treatment $TREAT_{it} \in \{0, 1\}$ is chosen. For any variable W , denote \widetilde{W}_{it} its history up to time t .

Let $y_{it}^{\tilde{a}}$ be the value of y that would have been observed at time t had i chosen treatment sequence $\tilde{a}_{it} = (a_{i0}, a_{i1}, \dots, a_{it})$ rather than his observed treatment history \widetilde{TREAT}_{it} . Note that, even if a_{ik} is dichotomous in each year k , there will be 2^k treatment histories and thus 2^k possible counterfactuals, only one of which is observed for each individual.

By definition, the average treatment effect of treatment history \tilde{a} on the outcome y is the difference $E[y^{\tilde{a}}] - E[y^{\tilde{0}}]$, the average difference between outcomes when following \tilde{a} and outcomes when never treated. We model the mean of $y^{\tilde{a}}$ conditional on treatment and exogenous covariates X as:

$$E[y_{it}^{\tilde{a}} | \widetilde{TREAT}_{it}, X_{it}] = \beta_0 + \beta_1' X_{it} + \beta_2 \Psi(\widetilde{TREAT}_{it}) \quad (1)$$

where $\Psi(\cdot)$ is a “dose-response function.” For example, if $\Psi(\cdot)$ puts a weight of 1 on $TREAT_{it}$ in each time-period, then it is the *stock* of patents that influences publishing rates. Conversely, if $\Psi(\cdot)$ puts a weight of 1 on $TREAT_{it}$ and a weight of 0 on $TREAT_{ik}$, $k = 0, \dots, t-1$, then only the instantaneous *flow* of patents has a causal effect on outcomes. In the empirical work, we will experiment with various specifications for $\Psi(\cdot)$.

Following Robins (1999b), we introduce the Sequential Conditional Independence Assumption (SCIA), which provides a formal way to extend the assumption of selection on observables to the case of dynamic treatments:

$$y_{it}^{\tilde{a}} \perp\!\!\!\perp TREAT_{it} | TREAT_{i,t-1}, Z_{i,t-1}, X_{it}$$

for all i and t , where the $\perp\!\!\!\perp$ sign denotes statistical independence. Robins (1999b) shows that under SCIA, the average treatment effect β_2 is identified and can be recovered by estimating

$$y_{it} = \beta_0 + \beta_1' X_{it} + \beta_2 \Psi(\widetilde{TREAT}_{it}) + \varepsilon_{it} \quad (2)$$

by weighted least squares, where the weights correspond to the inverse probability of following actual treatment history \widetilde{TREAT}_{it} up to time t for individual i . Note that (2) differs from (1) in that the observed outcomes y have been substituted for the counterfactual outcomes $y^{\tilde{a}}$.

Implementing IPTW estimation is relatively straightforward. Under SCIA, the selection bias can be removed by weighting the regression by:

$$w_{it} = \frac{1}{\prod_{k=0}^t Prob(TREAT_{ik} | \widetilde{TREAT}_{i,k-1}, \widetilde{Z}_{i,k-1}, \widetilde{X}_{ik})}$$

Each factor in the denominator is the probability that the researcher received her own observed treatment at time k , conditional on past treatment history and her past history of “prognosis factors” for treatment, whether time-varying or fixed over time. Therefore, the denominator of w_{it} represents the conditional probability that an individual followed his or her own history of treatment up to time t . Suppose that all relevant time-varying confounders are observed and included in Z_{it} . Then, weighting by w_{it} effectively creates a pseudo-population in which Z_{it} no longer predicts selection into treatment and the causal association between treatment and outcome is the same as in the original population. We refer to $\widehat{\beta}_2$ when eqn. (1) is weighted by w_{it} as the Inverse Probability of Treatment Weighted (IPTW) estimator of β_2 .

At this juncture, it is useful to pause and ask, why, if selection is assumed to depend only on observables, would it be invalid to just include all determinants of selection on the right-hand side of the outcome equation and to proceed with estimation by ordinary least squares? The answer is twofold. First, weighting the outcome equation by the inverse probability of treatment controls for these factors without making strong functional form assumptions; it can be thought of as regressing outcomes on treatment and a very flexible function of the variables in the selection equation. In the presence of staggered treatments and time-varying confounders, there is another important consideration. Under the usual assumption regarding orthogonality of the regressors to the error term, β_2 can be estimated consistently. However, such an estimate will not correspond to any causal parameter of interest, because the time-varying confounders are themselves affected by past treatment

history. In this situation, controlling directly for intermediate outcomes (for instance by including a lagged dependent variable as a regressor) would lead to an underestimate of the magnitude of the treatment effect.

The probabilities $Prob(TREAT_{ik}|\widetilde{TREAT}_{i,k-1}, \widetilde{Z}_{i,k-1}, \widetilde{X}_{ik})$ may vary greatly between subjects when time-varying confounders are strongly associated with treatment. This variability can result in extremely large outlying values for w_{it} . These outliers will contribute heavily to the pseudo-population, and the resulting IPTW estimator will have a very large variance. This problem can be alleviated by replacing w_{it} by a “stabilized” weight sw_{it} :

$$sw_{it} = \prod_{k=0}^t \frac{Prob(TREAT_{ik}|\widetilde{TREAT}_{i,k-1}, \widetilde{X}_{ik})}{Prob(TREAT_{ik}|\widetilde{TREAT}_{i,k-1}, \widetilde{Z}_{i,k-1}, \widetilde{X}_{ik})}$$

Although this modification does not influence the consistency of IPTW estimators, it does increase their efficiency (Hernán et al., 2000). Despite its simplicity and intuitiveness, IPTW estimation also presents some significant drawbacks. First and foremost, the assumption of no unobserved confounding is a strong one. Past research in the program evaluation literature has shown that techniques assume selection on observables perform well (in the sense of replicating an experimental benchmark) when (1) researchers use a rich list of covariates to model the probability of treatment; (2) units are drawn from similar labor markets, and (3) outcomes are measured in the same way for both treatment and control groups (Dehejia and Waba, 2002; Smith and Todd, 2005). All of these conditions would appear to be met in our setting and data, but this should not lead researchers to believe that IPTW estimation represents a universal solution for endogeneity problems. A second limitation is that IPTW estimates are just identified: the assumption of no unobserved determinants of selection into treatment cannot be tested; neither can misspecification of the selection equation used to estimate the weights. Third, the causal effect estimated by IPTW models is the population average treatment effect (ATE). In social science applications, however, the average treatment effect on the treated (ATET) might also be of policy interest. However,

Lechner and Miquel (2005) show that ATET is not identified without further assumptions on the joint distribution of the counterfactual outcomes.⁵

Informative censoring. Although we focused the first part of the discussion on the problem of non-random selection into patenting, a second problem arises because some subjects might exit the sample for endogenous reasons. For instance, scientists might leave academia because of low scientific productivity, or because they receive attractive offers to join commercial firms. Even if treatment was randomly allocated across units, this type of informative censoring could jeopardize the validity of the statistical estimates. We deal with this problem by treating censoring as just another time-varying treatment. As Robins et al. (2000) note, from this point of view, adjusting for censoring is only to say that our interest lies in estimating the causal effect of $TREAT$ on y if, contrary to the fact, all subjects had remained in the sample rather than having followed their censoring history. We model the exit decision as a function of constant and time-varying observable factors, and compute weights corresponding to the probability of exit given these observables:

$$sw_{it}^* = \prod_{k=0}^t \frac{Prob(EXIT_{ik} | \widetilde{TREAT}_{i,k-1}, X_{ik})}{Prob(EXIT_{ik} | \widetilde{TREAT}_{i,k-1}, \widetilde{Z}_{i,k-1}, X_{ik})}$$

sw_{it}^* is the inverse of the ratio of a scientist’s probability of exiting academia up to year t divided by that probability calculated as if there had been no time-dependent determinants of censoring except past treatment history and X . Hernán et al. (2001) shows that consistent estimates for β_2 can be obtained by combining the weight corresponding to the inverse probability of treatment sw_{it} and the weight corresponding to the inverse probability of censoring sw_{it}^* . The denominator of the final weight, $sw_{it}^* \times sw_{it}$, is the probability that a subject would have followed his own treatment and censoring history up to year t , conditional on observables. As a result, we label this methodology *Inverse Probability of Treatment and Censoring Weighted* (IPTCW) estimation in the rest of the paper.

⁵One might worry about performing statistical inference using “second stage” IPTW estimates, since the weights that are used as input in the outcome equation are themselves estimated. In contrast to two-step selection correction methods, Wooldridge (2002) has shown that the standard errors obtained in this case are conservative.

Estimation of the weights. The procedure followed to compute the weights depends on the way in which treatment is defined. According to a first definition, treatment is a *flow*: $TREAT_{it} = 1$ whenever researcher i applies for at least one patent in year t , and 0 otherwise. This formulation implies that treatment does not necessarily have a lasting impact on the individual. In contrast, the *regime* formulation defines $TREAT_{it} = 1$ for all years subsequent to the first patent application. Defining treatment this way implies a one-time shift on the outcome of interest, with subsequent treatment episodes having no effect on the dependent variable.

In the flow formulation, the weights are computed by estimating pooled cross-sectional logit specifications on the whole dataset. To compute the denominator of sw_{it} , one estimates a logit model for:

$$Prob(TREAT_{it} = 1) = \alpha_0 + \alpha_1 TREAT_{i,t-1} + \Phi(\tilde{Z}_{i,t-1}, \alpha_2) + \alpha_3 X_{it} + \delta_t \quad (3)$$

where $\Phi(\tilde{Z}_{i,t-1}, \alpha_2)$ corresponds to a parametric function of past values for time-varying confounders, X_{it} includes both time-varying and fixed-over-time characteristics of individuals in the sample (such as years of experience, gender, characteristics of the Ph.D-granting institution, etc.), and δ_t represents calendar year effects. In practice, we specify Φ as a linear function of publication flow in year $t-1$ and stock of publications up to year $t-2$, the number of past collaborations with industrial firms, patentability of the scientist's flow of publication in year $t-1$ and its stock up to year $t-2$, and employer characteristics. Let T_1 denote the set of years in which scientist i gets at least a patent and T_2 the set of years during which i gets no patents. The estimate of the denominator of sw_{it} is $\prod_{t \in T_1} \hat{p}_{it} \prod_{t \in T_2} (1 - \hat{p}_{it})$, where \hat{p}_{it} refers to the predicted probability obtained after estimating eqn. (3). The numerator of sw_{it} stems from an almost identical specification, except that one omits the term $\Phi(\tilde{Z}_{i,t-1}, \alpha_2)$ from the list of covariates.

This approach needs to be slightly modified when treatment is modeled as a regime shift rather than as a flow, because the probability of getting treatment remains constant and equal to one once a scientist enters the patenting regime. As a result, it is only necessary to fit the model on a subset of the data, that of scientist-year observations up to the year when

the scientist applies for his/her first patent. In this risk set, $TREAT_{i,t-1}$ is uniformly 0. To compute the denominator of sw_{it} we estimate a logit model for

$$Prob(TREAT_{it} = 1) = \alpha_0 + \Phi(\tilde{Z}_{i,t-1}, \alpha_2) + \alpha_3 X_{it} + \delta_t \quad (4)$$

and to compute the numerator of sw_{it} we estimate a logit model for

$$Prob(TREAT_{it} = 1) = \alpha_0 + \alpha_3 X_{it} + \delta_t \quad (5)$$

Our estimate of the denominator of sw_{it} for scientist i in year t is $\prod_{k=0}^t (1 - \hat{p}_{ik})$ if scientist i did not apply for at least one patent by year t , and $\prod_{k=0}^{t-1} (1 - \hat{p}_{ik}) \times \hat{p}_{it}$ if scientist i applied for his first patent in year t . Estimation of sw_{it}^* proceeds in the same fashion.

Relationship of IPTCW estimation with propensity-score matching methods.

Rosenbaum and Rubin (1983) refer to $Prob(TREAT_i = 1|X, Z)$ as the propensity score, and show how to use this probability to estimate treatment effects when selection into treatment depends only on observables. Recently, Heckman et al. (1997) have combined the propensity score with difference-in-differences to estimate the causal effect of undergoing a job training program. Abadie (2005) proposes a semiparametric difference-in-differences estimate that weights observations by the inverse probability of (own) treatment. Although the goals of these earlier papers resemble ours, we follow a different approach because the structure of our data differs significantly from the typical program evaluation setup. Labor econometricians generally study programs for which a “before” and “after” period can be unambiguously delineated for both treated and untreated units. In contrast, in our setting and many others, selection into treatment can occur at different times and/or in several disjoint episodes. Matching on the propensity score is difficult in these cases. Intuitively, an untreated individual might be a good control for a treated subject in one period (in the sense that the difference in their propensity scores is close to 0) and a bad control for the same treated subject in another period. In contrast, IPTCW estimation readily generalizes to the case of treatments that are staggered over time.

Robustness to unobserved heterogeneity. If our selection model does not capture some relevant determinants of the patenting decision, and these omitted factors influence research

output directly, IPTCW estimates will be biased. Since there is overwhelming evidence of positive selection in the cross-sectional dimension of the data (“better” scientists are both more likely to patent and publish heavily), such unobserved heterogeneity likely leads us to *overestimate* the treatment effect. Therefore, neither fixed effects nor the IPTCW approach provides a fully satisfactory solution to the problem of estimating the causal effect of patenting on publishing rates. In combination, however, these estimators implicitly define a confidence interval, with the fixed effects estimate providing a lower bound, and the IPTCW estimate providing an upper bound. The evidence presented below will show that, in practice, these bounds are sufficiently tight to inform the policy debate surrounding academic patenting.

4 Data and Sample Characteristics

We examine the association between patenting and publishing in a panel dataset of academic life scientists employed at universities and non-profit research institutes. This area was chosen because the biomedical fields have accounted for the preponderance of university patenting and licensing activity (Mowery et al., 2001). While we have not selected scientists because they have patented, we have sampled from scientific disciplines that we know to have significantly contributed to a vibrant area of technological development. We began by drawing 12,000 doctoral degree recipients from UMI Proquest Dissertations, which lists Ph.D. recipients from more than one thousand universities. In forming the sample, we randomly selected individuals, but only those with Ph.D.s in scientific disciplines that have informed commercial biotechnology.⁶ This assures a random sample of Ph.D.s in areas in which academic research may have significant, short-term commercial value.

⁶To identify the scientific disciplines that have been most important to biotechnology, we coded the educational backgrounds of the Ph.D.-holding, university-employed scientific advisory board members of all publicly traded biotechnology firms. The source of information on scientific advisors’ degrees was the IPO prospectuses of the 533 U.S.-based biotechnology firms that were filed with the U.S. Securities and Exchange Committee. We then stratified the random draw from UMI to correspond to the disciplines and Ph.D. years of firms’ scientific advisors. For example, 22 percent of biotechnology company scientific advisors hold biochemistry Ph.D.s; we drew a corresponding proportion of biochemists into our sample. Table 1 lists the Top 15 disciplines from which scientists in our sample are selected.

Next, we obtained scientists' publication records from the ISI's *Web of Science* database. Because the *Web of Science* includes authors' affiliations, we were able to identify Ph.D. graduates who pursued careers outside of academe. After removing individuals that (i) had no publications in any post-graduate year, (ii) published exclusively under corporate affiliations, or (iii) exited academe early in their careers,⁷ we were left with 3,862 scientists, all of whom we know to have been employed at U.S. universities or public research institutions. Each scientist is observed from the year after he or she earned a Ph.D. until 1999, unless the individual exited academia.⁸ The final panel contains 58,601 person-year observations between 1967 and 1999.

4.1 Variables

A brief description of the patenting process in academia is useful to interpret the results we will present. The process begins when a faculty member discloses an invention to the university's Technology Transfer Office (TTO).⁹ The commercial potential of this invention is then evaluated by the TTO, which may decide to seek patent rights on the invention. Concurrently, the TTO will market the innovation to potential licensing partners in industry. A typical licensing agreement specifies a 40% royalty rate for the individual faculty inventor, to be assessed on the gross licensing revenues the invention accrues.

Research outputs. From the *Web of Science* we computed annual paper publication counts for each scientist. We count all papers on which a scientist is listed as an author (in other words, we treat sole authored and coauthored papers as equivalents). Second, we used the affiliation data available in the *Web of Science* to identify all instances in which

⁷Ph.D.s with academic affiliations lasting less than five years were dropped from the dataset to exclude post-doctoral fellows that later moved to jobs in industry.

⁸We assume a researcher has exited academia when he or she fails to publish for five consecutive years, or in fewer instances, when the scientist begins to publish almost exclusively under a corporate affiliation. In either case, we censor observation in the year in which a scientist last publishes under a university affiliation.

⁹Faculty members are contractually obligated to disclose potentially commercializable discoveries developed on university premises to the TTO; they do not have the option to patent university-originated discoveries without going through the official channels. On average, TTO received 78 invention disclosures in 2003, but filed only 40 new patent applications (AUTM, 2003). Of course, these numbers vary widely across institutions depending on whether involvement with the world of commerce corresponds to a well-established culture within the institution.

a scientist wrote a paper that was coauthored with one or more individuals in a corporate research and development lab. We consider the rate of publication of papers with coauthors in industry as an indicator of the degree to which scientists are engaging in commercially-oriented research. We also keep track, for each journal in which our scientists published, of the relative prevalence of authors with corporate affiliations.¹⁰ In particular, for each scientist and in each year, we compute, following Lim (2004), an average Journal Commercial Score (JCS) by weighting each publication by the proportion of corporate authors who publish in the corresponding journal, summing the weights corresponding to all the articles published by the scientist in a given year, and dividing this sum by the (unweighted) number of articles he/she published during the year.

We use a two-pronged approach to measure the quality of the articles published. First, we make use of the order of authorship, computing the proportion of articles in which the scientist appears in first or last position. This choice is motivated by a robust social norm in the life sciences which systematically assigns last authorship to the principal investigator (generally the head of the laboratory), first authorship to the junior author who was responsible for the actual conduct of the investigation, and apportions the remaining credit to authors in the middle of the authorship list, generally as a decreasing function of the distance from the extremities of the list. In the second approach, we make use of the Journal Citation Reports, published yearly by the Institute for Scientific Information. ISI ranks journals by impact factor (JIF) in different scientific fields. The impact factor is a measure of the frequency with which the “average article” in a journal has been cited in a particular year. We weight each article published by the scientists in our sample by the corresponding journal’s JIF, sum these weights for all the published output in a given year, and divide by the yearly publication count. The resulting variable can be thought of as a measure of quality for the average article published by one of our scientists in a given year.¹¹

¹⁰For example, 35.7% of the affiliations for the authors publishing articles in the *Journal of Medicinal Chemistry* correspond to corporations. In contrast, the number is only 1.60% for the *Journal of General Physiology*.

¹¹Basically a ratio between citations and recent citable items published, JIFs suffer from built-in biases: they tend to discount the advantage of large journals over small ones, of frequently-issued journals over less frequently-issued ones, and of older journals over newer ones. Nonetheless, they convey quite effectively the idea that the *New England Journal of Medicine* (Impact Factor = 23.223 in 1991) is a much more influential

Patents. The patents of the academic scientists in our data were assembled from the NBER patent database (Hall et al., 2001). To identify academic patenters, we matched the scientists in our dataset to the list of inventors in the NBER patent database. Matches were done on the basis of first and last names, and we used information on assignee (university) and geographic region to eliminate false matches. For each scientist in our data, we generated flow and stock measures of patent applications, as well as corresponding dummy variables.

Control variables. Following a number of studies of the determinants of scientists' productivity, we were also able to construct a rich set of control variables to account for individual and institutional attributes that may influence rates of publication and patenting. To account for life-cycle effects (Levin and Stephan, 1991), we include the number of years since a scientist earned his or her Ph.D. An extensive literature in the sociology of science has documented gender differences in productivity (e.g., Long and Fox, 1995), so we include a "scientist is female" dummy variable. Because the time involved in publishing scientific research varies across fields, the regressions include a set of dummies for researchers' dissertation subject areas. Some of the regressions control for quality differences among researchers through the inclusion of scientist fixed effects. In specifications without fixed effects, we enter a dichotomous measure of the quality of a scientists' Ph.D.-degree granting institution — a dummy variable indicating whether or not a scientists' doctoral program was ranked in the Top 20. Specifically, we collected Gourman Report rankings for all institutions in our dataset. Gourman rankings for graduate schools were issued for the first time in 1980. We assigned universities their original rating for all years prior to 1980 and updated them every other year for the subsequent period. We also included in the models the stock of patents issued to the Ph.D-granting institution in the five years preceding the doctorate, to further control for the "imprinting" of norms regarding commercial activities during graduate training.

publication than the *Journal of General Internal Medicine* (Impact Factor = 1.056 in 1991). In an ideal world, rather than assigning an identical weight to all publications appearing in a given journal, we would instead weight each publication by the number of citations it garnered from other scientists. At the present time, querying the *Web of Science* to collect this information is prohibitively time-consuming since this database does not provide time-varying citation data.

From previous research, we know that institutional context exerts a strong influence on the propensity to commercialize research, either in the form of a well-funded technology licensing office, or through the presence of prominent peers who themselves are engaged in this activity (Di Gregorio and Shane 2003; Lach and Schankerman 2004; Stuart and Ding, 2006). As a result, we also include in our models a number of employer-level variables. These covariates are updated each year and when scientists change employers. First, we include a quality rank dummy variable analogous to the one constructed for Ph.D.-granting institutions. There are a variety of reasons why scientists at prominent universities are likely to be more productive, including the availability of more resources and easy access to high quality colleagues. Second, we used the AUTM surveys to create a technology transfer office (TTO) dummy variable, which is set to one in all years in which a scientist’s employing university has an active TTO. Third, a university’s stock of patents is entered in the model, among other things to further control for institutional differences in support for patenting.

Patentability. In the regressions for selection into patenting used to construct the inverse probability of treatment weights, it would obviously be desirable to account for differences among scientists in the inherent “patentability” of their research. In past studies, latent patentability was thought to be unobservable, and researchers used field fixed effects as controls in order to hold constant individual scientists’ research agenda. In contrast, we attempt to measure patentability directly. To construct such a measure, we use the title words in scientists’ publications to identify the areas in which they have conducted research, and then apply weights to these areas based on an (endogenous-to-the-sample) measure of the extent to which other scientists working in these areas have patented their discoveries. Intuitively, we use the publications of scientists that have already applied for patent rights as the benchmark for patentable research, and then compare the research of each scientist in our dataset to this benchmark to generate a research patentability score for each scientist-year. Specifically, the research patentability score for scientist i in year t is defined as:

$$PATENTABILITY_{it} = \sum_{j=1}^J w_{j,t-1}^i \frac{n_{ijt}}{\sum_k n_{ikt}}$$

where $j = 1, \dots, J$ indexes each of the scientific keywords appearing in the titles of the journal articles published by scientist i in year t ,¹² n_{ijt} is the number of times each of the keywords j has appeared in scientist i 's articles published in year t , and w_{jt}^i is a weight for each keyword that measures the frequency with which word j is used in the titles of articles published by scientists who have entered the patenting regime in year t or earlier, relative to those who have not entered the patenting regime as of year t (the calculation of w_{jt}^i is detailed in Appendix I). Intuitively, the patentability of a scientist's research can change because of a change in the direction of the research of that scientist, or because other patenters' research increasingly comes to resemble that of the scientist. The former effect is captured by the ratio $\frac{n_{ijt}}{\sum_k n_{ikt}}$, the latter by the weights $w_{j,t-1}^i$. Because the benchmark in year $t - 1$ is used to weight title words in year t , year-to-year changes in the research patentability score will only reflect actions of the scientist (through their choices of title keywords), rather than contemporaneous changes in the benchmark.¹³

Finally, to capture the idea that the inherent patentability of past research might still influence the current propensity to patent, we compute a depreciated stock of the research patentability score using a perpetual inventory model. Through the impact of the depreciation rate δ , this formulation captures the fact that the recent substantive research orientation of a scientist's research should influence current behavior more strongly than scientific trajectories that unfolded in the more distant past.¹⁴

$$STOCK_RP_{it} = (1 - \delta)STOCK_RP_{i,t-1} + FLOW_RP_{it} = \sum_{\tau=0}^t (1 - \delta)^{t-\tau} \cdot FLOW_RP_{i\tau}$$

¹²We relied on title words in journal articles instead of journal- or author-assigned keywords because the *Web of Science* database did not begin to include keyword descriptors until 1992. However, the titles of biomedical research papers typically indicate the research area and the methodology used in the paper. We find high overlap between title words and keywords in the papers for which both are available.

¹³Previous researchers have developed other measures of proximity in technological space. For instance, Jaffe (1986) used a cosine-based measure to assess the proximity between the R&D portfolio of any given pair of firms. While this approach works well for measuring technological distance between dyads, it is not well suited to our setting, since we need to measure the distance between the scientific trajectory of any given scientist relative to that of a benchmark *group* of (patenting) scientists.

¹⁴We set δ equal to 0.15 — the Griliches constant — which has been used by many innovation researchers on whose work this paper builds. We verified that our core results are not sensitive to this arbitrary choice.

4.2 Descriptive Statistics

Among the 3,862 researchers in our sample, 717 (18.60%) hold one or more patents. In Figure 1, we plot the distribution of patents for the patenting researchers in our sample. The histogram illustrates a rapid drop off after one — most patenters are listed on 1 or 2 patents throughout their career, and very few scientists in our data receive more than 10 patents. Figure 2 displays the distribution of scientists' total publication counts by the end of our observation period, broken out by their patenting status. Consistent with the conventional wisdom that patenting is concentrated among the group of academically productive scientists, the distribution for the patenter subsample is much less skewed than that for the non-patenter subsample.

Table 2 presents the summary descriptive statistics for variables used in our analysis. Table 3 reports, by scientists' patenting status, the mean research and employer characteristics measured at five career stages. Researchers who have sought and received patent rights for their discoveries appear more productive at each of these five stages: they publish approximately one and a half times as many research papers as those who have never patented. The average JIF of these publications, however, does not seem to differ between the two groups. At all career stages, scientists who have applied for patent rights appear closer to commercial research than their non-patenting counterparts, as indicated by the fact that they have collaborated more often with researchers in the private sector. Finally, patenters are more likely to work in settings where a technology transfer office exists and patenting activity is intensive.

4.3 Results

We present four sets of results. Table 4 focuses on the antecedents of selection into patenting, and on the determinants of exit from academia. It provides evidence on the importance of time-varying confounding, and displays the specifications from which our probability of treatment and censoring weights are derived. Using these weights as inputs, the following

tables present results pertaining to the effect of patenting on the rate (Table 5), quality (Table 6), and direction (Tables 7) of scientific output.

Determinants of patenting activity. We begin by presenting results pertaining to the probability of applying for a patent in a given year (flow formulation) or for the first time (regime formulation). The results are displayed in Table 4. It is important to note that the list of independent variables and the risk set differ significantly across the flow and regime models. In the former, all scientist-year observations are included, and the list of independent variables includes a lag structure for patenting in order to address the possibility of structural state dependence. In the latter, the observations corresponding to years subsequent to the year of the first patent application are not part of the risk set; consequently, no lag structure for the dependent variable can be part of the set of right-hand side variables.

The econometric analysis confirms that time-varying confounders are important determinants of patenting activity for these scientists. First, controlling for the stock of publications up to year $t - 3$ and the flow of publications in year $t - 2$, the probability of patenting in year t is significantly increasing in the flow of publications in year $t - 1$: at the mean of the data, a standard deviation increase in the flow of lagged publications increases the probability of patenting by 7.30% for the flow specification (column 1a) and by 13.3% for the regime specification (column 2a).¹⁵

This conditional correlation strikes us as being an important finding, for it can help distinguish between competing interpretations of the association between scientific productivity and involvement with the world of commerce. In the first interpretation, commercialization activities correspond to attempts by academics to monetize established reputations and professional status. In the second interpretation, publications and patents are co-occurring outputs that encode the same set of scientific insights; patents, just like publications, reflect genuine shocks to scientific opportunities. We see the correlation between the onset of patenting and the lagged flow, but not the stock, of publications as much more consistent

¹⁵In a companion paper (Azoulay et al., 2005), we confirm that this result is robust to much more flexible specifications of the lag structure.

with the latter interpretation.¹⁶ The plausibility of this interpretation is reinforced by a peculiar aspect of US patent law, which grants inventors a one-year grace period from the date of publication for the filing of a patent application (Merges, 1997, p. 226). In other words, an academic inventor wishing to maximize the effective life of a patent would apply to the USPTO exactly 364 days after the date of publication, provided that he/she is willing to forego patent protection in foreign jurisdictions.¹⁷

We also find that previous ties to industry in the form of coauthorships, and the stock of patents for the university where the scientist obtained his/her doctorate increases the likelihood of patenting activities. Similarly, scientists working in areas of science that are inherently more amenable to patenting are, unsurprisingly, more likely to patent. At the mean of the data, a high (in the top quartile) research patentability score increases the probability of patenting by 14.48% (column 1a) and by 21.7% (column 2a).¹⁸ Just as in the case of publications, the onset of patenting appears simultaneous with a change in the content of a scientist's research in a direction that makes it more similar to that of scientists who have already applied for patent rights. But because it is the flow, and not the stock of this measure that seems to matter, the evidence is consistent with the idea that a patent application does not constitute merely a response to changes in the formal and informal incentives faced by academic scientists over their careers, but also reflects the seizing of opportunities along a novel research trajectory.

In light of the results above, the shortcomings of fixed-effects estimation strategies become clearer. Selection into patenting is influenced by transitory shocks to outcome variables of interest, such as publications and their commercial content. While scientist fixed effects

¹⁶This interpretation is also consistent with Murray and Stern's (2005) analysis of paper-patent pairs, but it suggests that this phenomenon is not confined to the single journal whose articles they analyze. Of course, since we do not examine the actual content of patents and papers, we can only provide circumstantial evidence in favor of a substantive linkage between these two forms of output. In practice, it seems likely that patentable claims will be spread over a number of papers revolving around a common theme, some published before, some after the filing of the patent application.

¹⁷This result also provides strong evidence against the crowding-out hypothesis, at least in its simplest form. If the patent application process carried a high opportunity cost of time, one would expect this to be reflected in the output of patenting scientists before their first patent application. The opposite is true.

¹⁸This conclusion is not altered when using a more flexible functional form to model the distributed lag of the latent patentability score (Azoulay et al., 2005).

purge econometric estimates from selection bias stemming from immutable characteristics, they will fail to account for the transitory dynamics highlighted above.

Determinants of exit from academia. Models 3a and 3b display the results corresponding to specifications modeling the probability of exit from academia. A priori, one might imagine that academic scientists leave academia because they do not achieve success in the publication game. One might also conjecture that very productive academics are more likely to be poached by the private sector, leading to a premature exit from the academic ranks. We find support for both stories. Even controlling for the stock of past publications, a dry spell in academic productivity significantly increases the likelihood of exit. The stock of patents up to year $t - 2$ and research patentability are found to have no meaningful effect, but a patent application in year $t - 1$ is associated with a 36% increase in the probability of exit (column 3a).

Effect of patenting on the rate of publication output. Table 5 is divided into three sets of results, corresponding to three definitions of the patenting effect: flow (Models 1a, 1b, and 1c), regime (Models 2a, 2b, and 2c), and stock (Models 3a, 3b, and 3c). Within each set, the first column reports on the determinants the rate of publication using the conditional fixed effect poisson model of Hausman et al. (1984). As noted earlier, these estimates are likely to understate the causal effect of patenting. The second column is a “naïve” specification for the count of research publications, using Poisson Quasi-Maximum Likelihood Estimation (PQMLE).¹⁹ The corresponding estimates are likely to be biased upwards by unobserved heterogeneity. The third column is identical to the second except that it also incorporates our inverse probability of treatment and censoring weights. Under the sequential conditional independence assumption, these estimates correspond to the average treatment effect of patenting. Table 5 yields three robust results: (a) regardless of the method employed, the

¹⁹Because the Poisson model is in the linear exponential family, the coefficient estimates remain consistent as long as the mean of the dependent variable is correctly specified (Gouriéroux et al., 1984). Further, “robust” standard errors are consistent even if the underlying data generating process is not Poisson. In fact the PQMLE can be used for any non-negative dependent variables, whether integer or continuous (see Santos Silva and Tenreiro, Forthcoming). In all specifications, we trim the sample by dropping observations with extreme values of the selection and censoring weights — those below the 0.5 percentile and above the 99.5th percentile.

estimated effect of patenting is positive and statistically significant; (b) the IPTCW estimates are always higher than the conditional fixed effect estimates; and (c) in the cross-section, the magnitude of the effect is much lower once we account for self-selection into patenting. The formula $(e^\beta - 1) \times 100\%$ (where β denotes an estimated coefficient) provides a number directly interpretable in terms of elasticity. For example, the estimates in columns 2a, 2b, and 2c imply elasticities of publishing with respect to patenting equal to .138, .519 and .288, respectively.

Effect of patenting on the quality of publication output. Table 6 uses two distinct measures of publication quality. The first is the proportion of publications in which the researcher appears in first or last position in the authorship list (Models 1a and 1b). The second is the average journal impact factor for the articles published in a given year (Models 2a and 2b).²⁰ For the proportion, we estimate the model using the quasi-maximum likelihood fractional logit estimator of Papke and Wooldridge (1996). For the average JIF, estimation is performed using the Poisson QML approach as in Table 5. The estimated effects are small in magnitude and statistically insignificant, suggesting that patenting has very little impact on the quality of scientific output.

Effect of patenting on the content of publication output. Measuring the direction of scientific progress is always more challenging than merely measuring scientific output. In Table 7, we propose three distinct ways of measuring the commercial content of scientists' publications, and we show that our conclusions are not sensitive to the choice of measure. We begin by using the research patentability score described in section 4 as the dependent variable, and we perform estimation using the PQML estimator in columns 1a and 1b (since RP is a non-negative, albeit continuous, dependent variable). Patenting increases modestly the latent patentability of the research published in the post-patenting regime, even when we adjust for confounding (our weights take into account the fact that a shock to patentability in period $t - 1$ is associated with an increased likelihood of patenting at time t). For example,

²⁰Note that these two measures are not defined whenever a scientist has no output in a given year. As a result, the estimation sample shrinks by about a third.

the estimates in Model 2b imply that entering the patenting regime increases RP by a statistically significant 5.3%.

Models 2a and 2b provide a different angle on the same question by focusing on the institutional affiliations of our scientists' coauthors. In the years of positive output, we compute the fraction of total publications accounted for by articles in which at least one coauthor has an industry affiliation. At the mean of the data, the IPTCW estimates imply that entering the patenting regime increases this proportion by a statistically significant 36%. The naïve cross-sectional estimate is of a similar magnitude.

Finally, Models 3a and 3b use the average Journal Commercial Score (JCS) as the dependent variable. Starting from a journal-specific index that measures the proportion of authors publishing in the journal that have an industry affiliation, we compute the scientist-specific score by averaging these weights over all articles published in a given year.²¹ Patenting appears to increase the average JCS in a statistically significant fashion, but the magnitude of the effect is modest: at the mean of the data, the IPTCW estimates correspond to 4.2% increase in average JCS for patenting scientists.

Taken together, however, these results paint a consistent picture whereby patenting increases the rate of scientific output while maintaining its quality, but also changes the content of these publications by connecting them more tightly to the world of commerce.

Sensitivity analysis. If the fixed effect specifications understate the causal effect of patenting, but the IPTCW specifications overstate it, Models 2a and 2c in Table 5 imply that the average treatment effect of patenting on publication rates, expressed as an elasticity, lies within the interval [0.138; 0.288]. In order to gauge the robustness of inverse probability of treatment-weighted estimation to confounding by unobserved variables, we conduct a sensitivity analysis. Following Robins (1999a) and Brumback et al. (2004), we ask how much unmeasured confounding would there need to be for the confidence interval around our treatment effect to include 0? This approach compels us to parameterize the bias from unobserved

²¹Note that this measure has the advantage of not conflating the effect of patenting on the content of publications with its effect on the quantity of publication. As in the case of the average JIF, however, it suffers from the shortcoming that it is not defined whenever a scientist does not publish in a given year.

confounding — a functional form choice that is guided by little else than intuition regarding the cause and direction of bias. As such, the exercise does not provide a formal specification test for our results. Yet, its results are reassuring in the sense that our estimates appear robust to substantial amounts of unmeasured confounding. Estimation details and results are provided in Appendix II.

5 Discussion and Conclusion

While past research had established that commercialists are disproportionately recruited from the ranks of elite scientists and institutions (Zucker et al., 1998), our results build on this prior literature by showing that patenting is often accompanied by a flurry of publication activity in the year preceding the patent application, even after accounting for the lagged stock of publications. This result highlights the fact that academic patenting, rather than merely reflecting the influence of time-invariant demographic factors, also responds to variation in scientific opportunities (Azoulay et al., 2005). We also find that academic scientists who patent are more productive than otherwise equivalent scientists that are not listed as inventors on patents, but that publication quality appears similar in the two groups. Thus, the evidence appears to reject the assertion that the increase in patenting in academe has come at the cost of diverting researchers' time, interest, and attention from their traditional focus on standard scientific research. However, we also find that scientists alter the content of their research after they patent in ways that make their output more relevant to questions of commercial interest.

These results depend on the strong assumption that the outcomes we examine be independent of patenting conditional on the history of observables. As in all observational studies, this assumption cannot be tested from the data. It is obviously better to include a large set of potential confounders to model the probability of selection, but we recognize that in practice, the assumption of selection on observables may still not be precisely or even approximately tenable. We take solace in the results of a sensitivity analysis showing that our core result is in fact robust.

There are two other avenues, all outside the scope of this analysis, through which patenting in academic science could yet have a significant — and possibly deleterious — effect on the advancement of scientific knowledge. As a result, beyond the first-order effect of a scientist’s decision to patent on his or her individual productivity, our conclusions must remain tempered.

First, as patenting within a department or research area continues to grow, is there a point at which a negative effect on the collective output sets in, either because researchers are deterred or blocked by intellectual property rights held by others, or because concerns about intellectual property rights diminish open communications among scientists? This “tragedy of the anti-commons” has recently been investigated by Murray and Stern (2005), who provide evidence that scientific papers paired with patent applications are less likely to be cited after the patent is granted by the USPTO (though the effect they uncover is modest in magnitude). In the context of this paper, we present evidence that patenting changes the content of individual scientists’ research trajectory, but there is a strong leap from our results to normative statements regarding the welfare implications of this change in research agenda.

Academic patenting might also alter the career trajectories of the graduate students and post-doctoral fellows that work in patenters’ laboratories. For instance, patenters may have much thicker and more diverse relationships with researchers in firms than non-patenting scientists, which may in turn facilitate apprentice scientists’ job searches in the private sector. Therefore, patenters may (perhaps unintentionally) encourage their students to select private-sector careers above academic posts. Conversely, if patenters enlist the help of scientists-in-training in the research streams that lead to patents, and if these projects are different from the research topics that intrigue non-patenters, apprentices training under patenters may be less appealing to academic departments searching for new faculty. In short, the most significant impact of patenting on public research output may well lie in the consequence of the behavior for non-patenting and soon-to-be scientists. We plan to investigate this topic in future research.

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Table 1
Top 15 Scientific Disciplines in the Sample

UMI Subject			
Code	UMI Subject Description	Frequency	
487	Biochemistry	855	(22.1%)
306	Biology, General	563	(14.6%)
410	Biology, Microbiology	466	(12.1%)
419	Health Sciences, Pharmacology	239	(6.2%)
490	Chemistry, Organic	212	(5.5%)
786	Biophysics, General	210	(5.4%)
369	Biology, Genetics	191	(4.9%)
433	Biology, Animal Physiology	170	(4.4%)
982	Health Sciences, Immunology	167	(4.3%)
307	Biology, Molecular	102	(2.6%)
301	Bacteriology	61	(1.6%)
287	Biology, Anatomy	54	(1.4%)
571	Health Sciences, Pathology	52	(1.3%)
349	Psychology, psychobiology	37	(1.0%)
572	Health Sciences, Pharmacy	33	(0.9%)

Legend: Table 1 reports the Top 15 disciplines from which our sample was drawn. These disciplines have spawned the greatest number of biotechnology firm founders, scientific advisors and executives. The table also reports the frequency and the proportion of scientists in our sample for each of these 15 scientific disciplines.

Table 2
Descriptive Statistics

	Mean	Std. Dev.	Min.	Max.	<i>N</i>
Patent Flow (=1 if one or more patent app. in year)	0.030	0.170	0	1	58,601
Patent Regime (=1 after first patent app.)	0.111	0.314	0	1	58,601
Patent Stock	0.331	1.612	0	57	58,601
Research Publication Flow	1.729	2.378	0	35	58,601
Fraction of First or Last Authored Publications (Flow)	0.619	0.397	0	1	38,043
Average JIF of Publications (Flow)	3.956	3.103	0.005	30.334	38,043
Average Journal Commercial Score of Pubs. (Flow)	0.076	0.055	0.001	1	38,043
Fraction of Pubs. with Industry Coauthors (Flow)	0.075	0.223	0	1	38,043
Research Patentability Score (Flow)	0.084	0.108	0	5.185	58,601
Employer Graduate School in Top 20	0.231	0.422	0	1	58,601
Employer has TTO	0.488	0.500	0	1	58,601
Employer Patent Stock ($\times 0.01$)	0.718	1.452	0	21.890	58,601
Experience (Career Age)	10.203	7.121	1	32	58,601
Calendar Year	1986.1	7.741	1968	1999	58,601
Female	0.211	0.408	0	1	3,862
Ph.D. Univ. Grad. School in Top 20	0.308	0.462	0	1	3,862
Ph.D. Univ. 5-year Patent Stock ($\times 0.01$)	0.190	0.409	0	5.660	3,862

Table 3
Mean Scientist and Employer Characteristics at Five Career Stages,
by Patent Application Status

	Experience = 5		Experience = 10		Experience = 15		Experience = 20		Experience = 25	
	<i>Yes</i>	<i>No</i>	<i>Yes</i>	<i>No</i>	<i>Yes</i>	<i>No</i>	<i>Yes</i>	<i>No</i>	<i>Yes</i>	<i>No</i>
<i>Scientist ever applied for patent rights</i>										
(1) Count of Research Publications (Flow)	1.627 (1.892)	1.281 (1.791)	2.482 (2.848)	1.791 (2.197)	3.227 (3.554)	1.951 (2.477)	3.408 (3.851)	2.126 (2.821)	3.053 (3.660)	2.159 (2.990)
(2) Fraction of First or Last Authored Pubs.	0.609 (0.400)	0.605 (0.417)	0.618 (0.386)	0.628 (0.395)	0.582 (0.370)	0.624 (0.384)	0.600 (0.366)	0.580 (0.394)	0.601 (0.357)	0.576 (0.392)
(3) Average JIF of Research Publications	4.560 (3.656)	4.117 (3.382)	4.040 (3.133)	3.949 (3.017)	4.062 (2.730)	3.797 (2.778)	3.879 (2.841)	3.595 (2.646)	3.859 (2.597)	3.467 (2.730)
(4) Average JCS of Publications	0.074 (0.049)	0.077 (0.056)	0.077 (0.069)	0.077 (0.058)	0.083 (0.070)	0.075 (0.051)	0.069 (0.038)	0.073 (0.051)	0.069 (0.052)	0.074 (0.053)
(5) Fraction of Pubs. with Industry Coauthors	0.128 (0.295)	0.051 (0.194)	0.098 (0.236)	0.076 (0.231)	0.102 (0.237)	0.082 (0.231)	0.111 (0.251)	0.113 (0.261)	0.112 (0.228)	0.096 (0.239)
(5) Research Patentability (RP)	0.085 (0.093)	0.073 (0.121)	0.107 (0.097)	0.091 (0.105)	0.116 (0.090)	0.096 (0.097)	0.142 (0.118)	0.100 (0.088)	0.117 (0.091)	0.101 (0.086)
(6) Employer Grad. School in Top 20	0.290 (0.455)	0.265 (0.441)	0.299 (0.458)	0.216 (0.412)	0.246 (0.431)	0.197 (0.398)	0.196 (0.398)	0.180 (0.384)	0.172 (0.378)	0.170 (0.376)
(7) Employer has TTO	0.438 (0.498)	0.385 (0.487)	0.576 (0.495)	0.485 (0.500)	0.685 (0.465)	0.590 (0.492)	0.729 (0.445)	0.682 (0.466)	0.822 (0.383)	0.728 (0.445)
(8) Employer Patent Stock ($\times 100$)	0.815 (2.011)	0.538 (1.355)	1.247 (2.774)	0.642 (1.283)	1.252 (1.987)	0.750 (1.157)	1.306 (1.828)	1.091 (1.528)	1.538 (2.203)	1.215 (1.629)
<i>Number of scientists</i>	169	3,640	278	2,319	321	1,516	321	980	169	464

Legend: Table 3 reports the mean and standard deviation (in parentheses) of scientist research and employer characteristics measured at five career stages: the 5th, 10th, 15th, 20th and 25th year after a scientist was granted a Ph.D. At each professional age, the table is further broken out by whether a scientist has ever applied for a patent right.

Table 4
Probability of Patenting and Exiting Academia

	Model 1a	Model 1b	Model 2a	Model 2b	Model 3a	Model 3b
<i>Dependent Variable</i>	<i>Patent Flow</i>		<i>Patent Regime</i>		<i>Exit Academia</i>	
	<i>Denominator</i>	<i>Numerator</i>	<i>Denominator</i>	<i>Numerator</i>	<i>Denominator</i>	<i>Numerator</i>
Experience = [5, 8]	0.578 (0.115)**	0.663 (0.114)**	0.509 (0.128)**	0.589 (0.125)**		
Experience = [9, 15]	0.680 (0.133)**	0.849 (0.126)**	0.412 (0.142)**	0.563 (0.132)**	0.201 (0.062)**	-0.005 (0.057)
Experience = [16, 22]	0.481 (0.155)**	0.704 (0.145)**	0.236 (0.174)	0.442 (0.152)**	0.072 (0.088)	-0.266 (0.077)**
Experience = [23, 35]	0.017 (0.203)	0.259 (0.186)	-0.321 (0.253)	-0.112 (0.232)	0.298 (0.119)*	-0.130 (0.101)
Female	-0.934 (0.143)**	-0.972 (0.144)**	-0.899 (0.134)**	-0.929 (0.133)**	0.151 (0.054)**	0.242 (0.054)**
Patent Flow _{t-1}	1.197 (0.077)**	1.273 (0.078)**			0.322 (0.063)**	
Patent Stock _{t-2}	0.219 (0.029)**	0.243 (0.029)**			-0.013 (0.018)	
Publication Flow _{t-1}	0.030 (0.014)*		0.056 (0.017)**		-0.214 (0.024)**	
Publication Flow _{t-2}	0.025 (0.014)†		0.017 (0.020)		-0.095 (0.020)**	
Publications Stock _{t-3}	0.001 (0.002)		0.001 (0.003)		-0.010 (0.003)**	
High Res. Patentability _{t-1}	0.166 (0.061)**		0.195 (0.092)*		0.036 (0.064)	
Research Patentability Stock _{t-2}	-0.028 (0.112)		0.014 (0.127)		-0.001 (0.092)	
Has Industry Coauthors _{t-1}	0.443 (0.084)**		0.314 (0.093)**		0.116 (0.059)†	
Employer Grad. School in Top 20	0.079 (0.096)		-0.043 (0.103)		0.037 (0.060)	
Employer has TTO	0.128 (0.085)		0.098 (0.095)		0.006 (0.052)	
Employer Patent Stock _{t-1} in Top Quartile	0.035 (0.026)		0.089 (0.028)*		0.026 (0.017)	
Ph.D. Univ Grad. School in Top 20	0.054 (0.083)	0.082 (0.082)	0.014 (0.087)	0.044 (0.086)	-0.147 (0.053)**	-0.179 (0.053)**
Ph.D. Univ. 5-yr Patent Stock (×100)	0.116 (0.089)	0.149 (0.090)†	0.197 (0.090)*	0.256 (0.087)**	-0.096 (0.073)	-0.075 (0.075)
Constant	-5.989 (0.314)**	-5.932 (0.314)**	-5.984 (0.315)**	-5.936 (0.313)**	-5.368 (0.265)**	-5.523 (0.265)**
Observations	58,601	58,601	52,799	52,799	58,476	58,476
Number of researchers	3,862	3,862	3,862	3,862	3,862	3,862
Log pseudo-likelihood	-6302.1	-6389.2	-3567.9	-3607.6	-8835.4	-9070.4
Wald χ^2	1708.2	1632.3	425.9	353.4	569.6	311.3
Number of Variables	51	42	49	40	50	39

Notes:

- (1) Models 2a-2b exclude observations after a researcher has filed for his or her first patent application.
- (2) Models 3a-3b exclude observations after a researcher has accumulated 30 years' professional experience (at which point he or she is no longer considered at risk of exiting academia).
- (3) All models control for Ph.D. subject and calendar year dummies.
- (4) Robust standard errors in parentheses, clustered around individual researchers.
- (5) † significant at 10%; * significant at 5%; ** significant at 1%.

Table 5
Effect of Patenting on the Rate of Publications: Poisson Models

	Model 1a	Model 1b	Model 1c	Model 2a	Model 2b	Model 2c	Model 3a	Model 3b	Model 3c
<i>Scientist Fixed Effects</i>	<i>Yes</i>	<i>No</i>	<i>No</i>	<i>Yes</i>	<i>No</i>	<i>No</i>	<i>Yes</i>	<i>No</i>	<i>No</i>
<i>IPTC Weights</i>	<i>No</i>	<i>No</i>	<i>Yes</i>	<i>No</i>	<i>No</i>	<i>Yes</i>	<i>No</i>	<i>No</i>	<i>Yes</i>
Experience = [5, 8]	0.148 (0.013)**	0.192 (0.018)**	0.177 (0.018)**	0.149 (0.013)**	0.181 (0.018)**	0.171 (0.018)**	0.152 (0.013)**	0.195 (0.018)**	0.178 (0.018)**
Experience = [9, 15]	0.243 (0.020)**	0.429 (0.030)**	0.385 (0.030)**	0.239 (0.020)**	0.405 (0.030)**	0.363 (0.030)**	0.249 (0.020)**	0.428 (0.030)**	0.381 (0.030)**
Experience = [16, 22]	0.199 (0.032)**	0.538 (0.048)**	0.457 (0.047)**	0.183 (0.031)**	0.485 (0.048)**	0.414 (0.047)**	0.203 (0.032)**	0.518 (0.049)**	0.440 (0.048)**
Experience = [23, 32]	0.052 (0.044)	0.499 (0.072)**	0.361 (0.072)**	0.046 (0.044)	0.455 (0.072)**	0.328 (0.070)**	0.052 (0.044)	0.448 (0.073)**	0.332 (0.072)**
Female		-0.211 (0.051)**	-0.242 (0.047)**		-0.186 (0.051)**	-0.224 (0.046)**		-0.206 (0.051)**	-0.235 (0.047)**
PhD Univ. Grad School in Top 20		0.075 (0.041) [†]	0.093 (0.042) [*]		0.069 (0.041) [†]	0.089 (0.040) [*]		0.080 (0.041) [†]	0.094 (0.042) [*]
PhD Univ. 5-Year Patent Stock (×100)		0.036 (0.047)	0.045 (0.047)		0.039 (0.046)	0.047 (0.045)		0.036 (0.047)	0.045 (0.047)
Patent Flow	0.099 (0.017)**	0.475 (0.046)**	0.252 (0.054)**						
Patent Regime				0.129 (0.015)**	0.418 (0.045)**	0.253 (0.043)**			
Patent Stock							0.016 (0.002)**	0.054 (0.008)**	0.044 (0.008)**
Constant		0.217 (0.049)**	0.204 (0.048)**		0.199 (0.049)**	0.186 (0.048)**		0.224 (0.049)**	0.206 (0.048)**
Log pseudo-likelihood	-77531.9	-118905.6	-114174.6	-77492.4	118332.9	-113230.0	-77526.8	-118678.4	-114001.7
Wald χ^2	2923.6	1200.1	803.7	2941.1	1340.1	896.2	2931.5	1228.5	818.2
Number of covariates	31	41	41	31	41	41	31	41	41

Notes:

- (1) Number of observations = 58,298; number of researchers = 3,862.
- (2) 303 observations with extremely large or small IPTC weights have been excluded from the estimation sample.
- (3) All models control for calendar year dummies; all cross-sectional models also control for Ph.D. subject dummies.
- (4) All cross-sectional models display robust standard errors in parentheses, clustered around researchers.
- (5) [†] significant at 10%; ^{*} significant at 5%; ^{**} significant at 1%.

Table 6
Effect of Patenting on the Quality of Publications

	Model 1a	Model 1b	Model 2a	Model 2b
	Fractional Logit QML Estimates Proportion of First or Last- Authored Publications		Poisson Model QML Estimates Average JIF of Publications	
	<i>Unweighted</i>	<i>IPTCW</i>	<i>Unweighted</i>	<i>IPTCW</i>
Experience = [5, 8]	-0.093 (0.029)**	-0.094 (0.029)**	-0.086 (0.013)**	-0.084 (0.013)**
Experience = [9, 15]	0.041 (0.034)	0.038 (0.035)	-0.190 (0.018)**	-0.187 (0.018)**
Experience = [16, 22]	0.146 (0.047)**	0.146 (0.047)**	-0.272 (0.027)**	-0.278 (0.027)**
Experience = [23, 32]	0.165 (0.069)*	0.131 (0.071) [†]	-0.351 (0.040)**	-0.366 (0.039)**
Female	-0.007 (0.038)	-0.001 (0.038)	0.030 (0.022)	0.031 (0.022)
PhD Univ. Grad School in Top 20	0.053 (0.033)	0.049 (0.033)	0.137 (0.021)**	0.136 (0.021)**
PhD Univ. 5-Year Patent Stock (×100)	0.049 (0.042)	0.055 (0.042)	0.086 (0.030)**	0.093 (0.029)**
Patent Regime	-0.034 (0.041)	-0.051 (0.044)	0.043 (0.025) [†]	0.034 (0.025)
Constant	0.840 (0.056)**	0.838 (0.056)**	1.374 (0.026)**	1.564 (0.029)**
Log pseudo-likelihood	-22150.0	-21484.3	-91565.7	-88813.9
Wald χ^2	275.8	280.7	646.2	698.9

Notes:

- (1) Number of observations = 37,859; number of researchers = 3,862; number of variables = 41.
- (2) 184 observations with extremely large or small IPTC weights have been excluded from the estimated sample.
- (3) All models control for PhD subject and calendar year dummies.
- (4) Robust standard errors are reported in parenthesis, clustered around researchers.
- (5) [†] significant at 10%; * significant at 5%; ** significant at 1%.

Table 7
Effect of Patenting on the Commercial Content of Publications

	Model 1a	Model 1b	Model 2a	Model 2b	Model 3a	Model 3b
	Poisson Models QML Estimates Research Patentability		Fractional Logit QML Estimates Proportion of Pub. with Industry Coauthors		Fractional Logit QML Estimates Average Journal Commercial Score	
	<i>Unweighted</i>	<i>IPTCW</i>	<i>Unweighted</i>	<i>IPTCW</i>	<i>Unweighted</i>	<i>IPTCW</i>
Experience = [5, 8]	0.031 (0.017) [†]	0.031 (0.017) [†]	0.073 (0.068)	0.085 (0.069)	0.013 (0.015)	0.013 (0.015)
Experience = [9, 15]	0.018 (0.017)	0.019 (0.017)	0.091 (0.087)	0.113 (0.088)	0.004 (0.019)	0.006 (0.019)
Experience = [16, 22]	0.029 (0.019)	0.029 (0.019)	0.070 (0.113)	0.096 (0.115)	0.012 (0.025)	0.013 (0.026)
Experience = [23, 32]	0.012 (0.022)	0.011 (0.023)	0.027 (0.157)	0.049 (0.161)	0.049 (0.035)	0.061 (0.036) [†]
Female	-0.027 (0.013) [*]	-0.028 (0.013) [*]	-0.046 (0.092)	-0.040 (0.094)	-0.007 (0.017)	-0.006 (0.017)
PhD Univ. Grad School in Top 20	-0.033 (0.012) ^{**}	-0.033 (0.012) ^{**}	-0.304 (0.084) ^{**}	-0.312 (0.088) ^{**}	-0.067 (0.018) ^{**}	-0.066 (0.018) ^{**}
PhD Univ. 5-year Patent Stock (×100)	0.012 (0.018)	0.011 (0.018)	0.121 (0.098)	0.115 (0.096)	-0.020 (0.025)	-0.029 (0.024)
Patent Regime	0.047 (0.014) ^{**}	0.052 (0.016) ^{**}	0.350 (0.080) ^{**}	0.333 (0.087) ^{**}	0.040 (0.021) [†]	0.045 (0.022) [*]
Constant	-4.668 (0.121) ^{**}	-4.668 (0.121) ^{**}	-4.387 (0.217) ^{**}	-2.304 (0.120) ^{**}	-2.498 (0.026) ^{**}	-2.500 (0.026) ^{**}
Log pseudo-likelihood	-12566.8	-12096.2	-9037.2	-8775.7	7639.2	7391.2
Wald χ^2	1265.6	1262.6	322.7	316.3	423.3	383.5

Notes:

- (1) Number of observations = 37,859; number of researchers = 3,862; number of variables = 41.
- (2) 184 observations with extremely large or small IPTC weights have been excluded from the estimated sample.
- (3) All models control for PhD subject and calendar year dummies.
- (4) Robust standard errors are reported in parenthesis, clustered around researchers.
- (5) [†] significant at 10%; ^{*} significant at 5%; ^{**} significant at 1%.

Figure 1
Distribution of Patent Count for Patenting Scientists

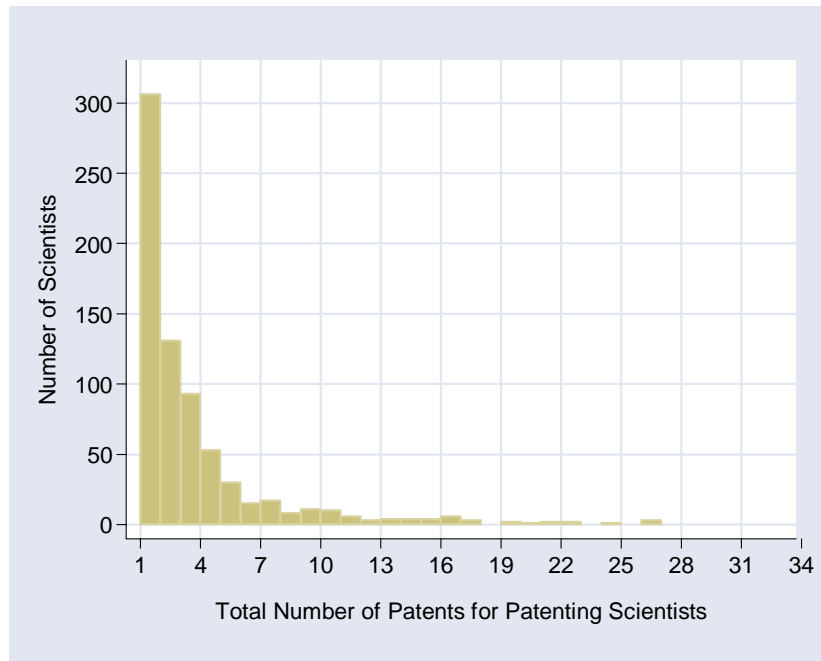
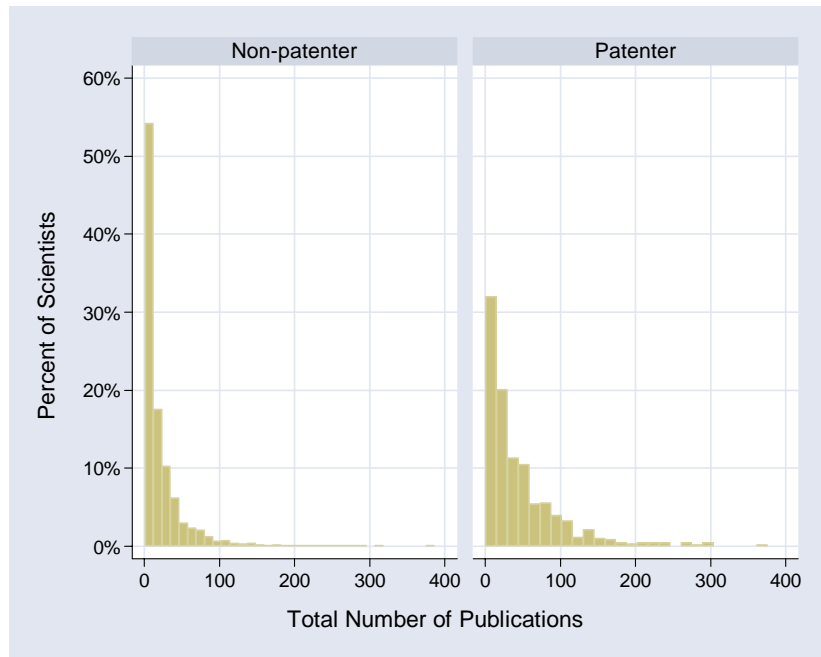


Figure 2
Distribution of Publication Count for Patenting and Non-patenting Scientists



Appendix I: Keyword Weights

w_{jt}^i , the patentability weight for each keyword j in year t is defined as:

$$w_{jt}^i = \frac{\sum_{s \in I_t^p - \{i\}} \frac{m_{sjt}}{\sum_k m_{skt}}}{\sum_{s \in I_t^{np} - \{i\}} m_{sjt}}$$

where m_{sjt} denotes the number of times keyword j has appeared in articles published up to year t by scientist s , I_t^p is the subset of scientists in our sample that have already applied for one or more patents as of year t , and I_t^{np} is the subset of scientists in our sample that have not yet applied for any patent as of year t . The weight is also indexed by scientist i , because i 's publications are taken out of the set of articles used to compute the formula above.

To create the numerator of w_{jt}^i , we first create a row-normalized matrix with each scientist in the patenting regime listed in a row and each of the keywords used to describe their papers up to year t listed in a column. The sj^{th} cell in the matrix, $[m_{sjt}/\sum_k m_{skt}]$, refers to the proportion of title keywords for scientist s that corresponds to keyword j . We then take the column sums from this matrix, i.e., we sum the contributions of individual patenting scientists for keyword j . Turning next to the denominator, we proceed in a similar manner, except that the articles considered only belong to the set of scientists who have not applied for patents as of year t . The numerator is then deflated by keyword j 's frequency in the titles of non-patenters' publications (in the rare case of keywords exclusively used by patenters, we substitute the number 1 for the frequency).

The weights w_{jt}^i are large for keywords that have appeared with disproportionate frequency as descriptors of papers written by scientists already in the patenting regime, relative to scientists not yet in the patenting regime.

Two things should be noted about the construction of these weights. First, $w_{jt}^i = 0$ for all keywords that have never appeared in the titles of papers written by scientists that have patented before t . Second, the articles written by scientist i him/herself do not contribute at all to the weights w_{jt}^i . Therefore, no scientist can directly influence year-to-year changes in these weights.

The final step for each scientist i in the dataset is to produce a list of the keywords in the individual's papers published in year t , calculate the proportion of the total represented by each keyword j , apply the appropriate keyword weight $w_{j,t-1}^i$, and sum over keywords to produce a composite score. The resulting variable increases in the degree to which keywords in the titles of a focal scientist's papers have appeared relatively more frequently in the titles of other academics who have applied for patents. This score is entered in the regressions to control for the research patentability of scientists' areas of specialization and to assess changes in the direction of their research.

To illustrate the construction of the research patentability measure, Table A1 lists some representative keywords, along with their patentability weights in the year 2000. Consider the keyword "Endoplasmic-Reticulum" (italicized in the table) in group 1. In 2000, it had previously appeared 22 times as a keyword in one or more articles of scientists who had patented prior to 2000. Among them is Carlos B. Hirschberg, professor and chair of molecular and cell biology at Boston University, who is listed as an inventor on a patent filed in 1996. To compute the numerator of the patentability weight for this keyword, we begin with the fraction of Hirschberg's research using "Endoplasmic-Reticulum" in the title. In his 29 ISI-listed research papers published between 1970, when he was granted a Ph.D. and 2000, 234 unique keywords have been used a total of 445 times. The word "Endoplasmic-Reticulum" was used 5 times, hence the fraction of Hirschberg's research stock devoted to "Endoplasmic-Reticulum" is 0.011. This procedure is repeated for the other 21 patenting scientists who have used the word. The sum of these fractions taken over all 21 patenting scientists is reported in the second column of the table. Next, to compute the denominator in the above equation, we examine the keywords of all scientists who had not yet received a patent by 2000 for the appearance of Endoplasmic-Reticulum. In the research publications of 3,841 such scientists, this keyword

has appeared on 46 occasions. The patentability weight for Endoplasmic-Reticulum is obtained by dividing the sum of proportions of keyword use among patenting scientists (column 2) by the frequency of use for this same keyword among non-patenting scientists (column 3).

Appendix II: Sensitivity Analysis

We present a sensitivity analysis for unmeasured confounding, following Robins (1999a) and Brumback et al. (2004). The basic approach is similar to that adopted by other researchers who have evaluated the robustness of non-experimental estimators in the presence of selectivity (see for example Altonji et al. (2005) for the case of Heckman two-step estimators and Rosenbaum (2002, chap. 4) for the propensity score). Namely, we attempt to quantify the amount of unmeasured confounding that would be necessary for our results to lose statistical significance. The exercise assumes a particular functional form for the bias from unmeasured confounding, generates new outcomes adjusted for the bias, and then performs IPTCW estimation by substituting these adjusted outcomes for the actual outcomes. For clarity, we begin by presenting the method in the case of a binary treatment with only two periods: baseline and end-of-study. The results are then generalized to the more relevant case of repeated treatments in a panel context.

Cross-sectional case. Let y be the outcome of interest (in our case, publications) at the end of the study, A be a binary treatment administered at baseline (*e.g.*, patenting), and V be a set of observables measured at baseline. Let y_i^a denote the outcome that person i would have experienced if he/she had, possibly contrary to the fact, received treatment a ($a = 0$ or $a = 1$). The method proposed by Robins (1999a) quantifies unmeasured confounding through the function Δ defined by:

$$\Delta(a, v) = E[y_i^a | A = a, V = v] - E[y_i^a | A = 1 - a, V = v]$$

Δ represents, among the set of individuals with baseline characteristics $V = v$, the average difference in potential outcomes between those treated with $A = a$ and those treated with $A = 1 - a$. For example, $\Delta(1, v)$ is the difference between the average outcome experienced by the treated when they are treated and the average outcome experienced by the untreated had they been treated instead. Note that the assumption of no unmeasured confounding is synonymous with $\Delta(a, v) = 0$ for all a and v . The effect of interest can be written $\theta = E[y_i^1 - y_i^0 | V = v]$. A naïve estimator for the treatment effect is:

$$\theta_{naive} = E[y | A = 1, V = v] - E[y | A = 0, V = v] \quad (\text{AII}_1)$$

The function Δ links θ and θ_{naive} through the identity

$$\theta = \theta_{naive} + \Delta(1, v) \cdot Prob[A = 0 | V = v] - \Delta(0, v) \cdot Prob[A = 1 | V = v] \quad (\text{AII}_2)$$

If $\Delta(1, v) > 0$ but $\Delta(0, v) < 0$, then on average, treated individuals will have higher potential outcomes to both treatment and no treatment than untreated individuals (*i.e.*, more talented scientists are more likely to patent). This seems to correspond to the type of bias that threatens the validity of the estimates presented in Section 4. Therefore we choose the following functional form for Δ :

$$\Delta(a, y_{i0}) = \alpha \cdot (2a - 1) \cdot y_{i0}$$

where α is a sensitivity parameter and y_{i0} denotes baseline outcome. With $\alpha > 0$, θ_{naive} is biased upward by $\alpha \cdot y_{i0}$. Robins (1999a) shows that for any user-specified α , one can obtain unbiased estimates of the treatment effect by replacing each observed outcome y_i with the adjusted outcome $y_i^\alpha = y_i - \Delta(A, V) \cdot Prob[1 - A | V]$ and recalculating θ_{naive} according to:

$$\bar{\theta}_{naive}^\alpha = E[y_i^\alpha | A = 1, V] - E[y_i^\alpha | A = 0, V] \quad (\text{AII}_3)$$

Intuitively, $\bar{\theta}_{naive}^\alpha$ subtracts the bias term in eqn. [AII₂] from θ_{naive} . To counteract bias from unmeasured confounding, we adjust the outcome of a treated individual downward by subtracting $\alpha \cdot y_{i0} \cdot Prob[A = 0|V]$, and that of an untreated individual by adding $\alpha \cdot y_{i0} \cdot Prob[A = 1|V]$.

Generalization to the panel case. The method of adjustment described above generalizes as follows for time-varying confounding. First, we compute the adjusted outcome

$$y_{it}^\alpha = y_{it} - \sum_{k=0}^{t-1} \Delta_{ik} \cdot Prob[1 - TREAT_{ik} | TREAT_{i,k-1}, \tilde{Z}_{i,k-1}, \tilde{X}_{i,k-1}] \quad (\text{AII}_4)$$

where $\Delta_{ik} = \alpha y_{ik}(2 \times TREAT_{ik} - 1)$. Second, we re-estimate the causal model with y_{it}^α in place of y_{it} . Subject to weak regularity conditions, the resulting augmented IPTCW estimator is consistent, assuming of course that the confounding function Δ is correctly specified. In what follows, we will focus on the case where treatment is specified as a “regime,” *i.e.*, once the scientist has patented for the first time, the treatment variable never again reverts to 0. For such monotonic treatments, the sum above simplifies to

$$\alpha y_{im} Prob[TREAT_{im} = 0 | \tilde{Z}_{i,m-1}, \tilde{X}_{i,m-1}] - \alpha \sum_{k=0}^{m-1} y_{ik} Prob[TREAT_{ik} = 1 | \tilde{Z}_{i,k-1}, \tilde{X}_{i,k-1}]$$

for scientists who initiate patenting in period $m \leq t - 1$, whereas for those initiating patenting at $m \geq t$, it remains

$$-\alpha \sum_{k=0}^{t-1} y_{ik} Prob[TREAT_{ik} = 1 | \tilde{Z}_{i,k-1}, \tilde{X}_{i,k-1}]$$

Choice of α and estimation details. From eqn. [AII₄], α is measured on the same scale as the dependent variable. We vary delta from 0.00 to 1.00 publications, in increments of 0.10. A priori, we regard negative values for α as implausible, since $\alpha < 0$ implies that non-patenters have higher potential outcomes regardless of whether they patent. One can think of increasing α as progressively stacking the deck against the finding of a positive effect of patenting on the rate of publication. $\alpha = 0$ corresponds to the estimate found in Table 5, Model 2c. The corresponding 95% confidence interval is [0.168; 0.337].

IPTCW estimation proceeds in two steps, where estimates of the first step (from which the weights derive) are used as inputs in the second step. As Newey and McFadden (1994) have shown, the standard errors that result from the second step when the first-step estimates are assumed known instead of estimated will be biased. In the specific case of IPTCW estimation, Robins (1999b) has shown that the naïve asymptotic standard errors are upward-biased. However, this does not necessarily hold true for the augmented model. Since computing the standard errors analytically can be very cumbersome in these circumstances, we have used the block bootstrap (based on 1,000 replications) where we treat each scientist as a sampling unit, thus allowing for arbitrary serial correlation and heteroskedasticity. We account for the two-stage nature of IPTCW estimators by bootstrapping the entire estimation procedure.

Results. As can be seen on Table A2, the threshold value of α such that the 95% confidence interval around the treatment effect falls just short of containing 0 is about 0.53. If one is willing to use a one-tailed test, the threshold value of α is slightly larger, about 0.58. This might be warranted if one holds a strong prior regarding the direction of bias from unmeasured confounding.

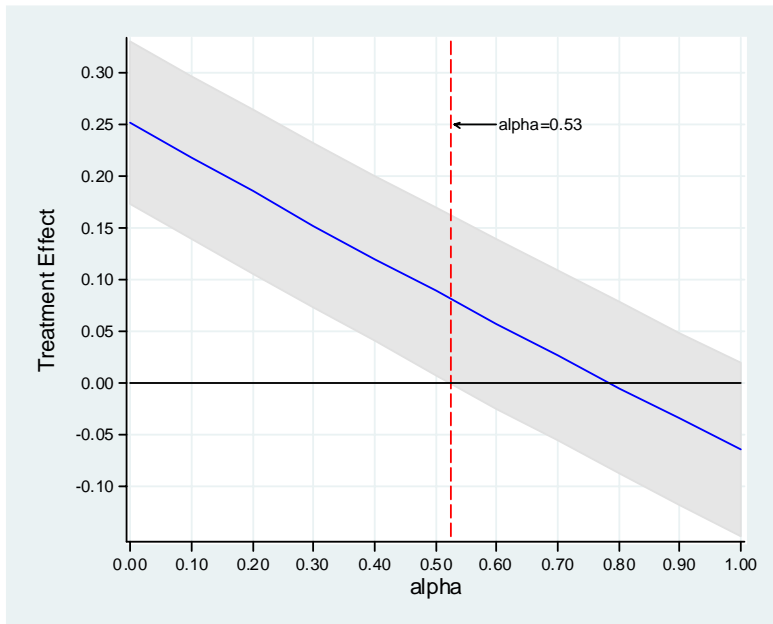
Table A1: Sample Title Keywords in the year 2000

	(1)	(2)	(3)	(4)
	Number of times keyword used by patenting scientists	Sum over all patenting scientists of keyword's proportion of total keywords used	Number of times keyword used by non- patenting scientists	Keyword weight: Column (2) / Column (3)
	$\sum_{s \in I_t^p - \{i\}} m_{sjt}$	$\sum_{s \in I_t^p - \{i\}} \frac{m_{sjt}}{\sum_k m_{skt}}$	$\sum_{s \in I_t^{np} - \{i\}} m_{sjt}$	$\frac{\sum_{s \in I_t^p - \{i\}} m_{sjt}}{\sum_{s \in I_t^{np} - \{i\}} m_{sjt}}$
<i>Group 1</i>				
Polyhedrosis-Virus	31	0.144	14	1.030
Trypanosoma-Cruzi	37	0.123	14	0.877
Copolymerization	29	0.067	11	0.612
Schistosoma-Mansoni	44	0.083	17	0.491
Phosphoribosyltransferase	36	0.092	27	0.342
Autocrine	33	0.073	22	0.330
Follicle-Stimulating-Hormone	45	0.061	19	0.323
Etoposide	34	0.052	17	0.306
Atherosclerosis	43	0.101	35	0.289
Methotrexate	73	0.170	63	0.271
<i>Endoplasmic-Reticulum</i>	54	0.114	46	0.247
Antitumor	106	0.220	92	0.240
Integrin	113	0.229	98	0.234
Leukotriene	43	0.078	34	0.231
Monooxygenase	38	0.051	33	0.156
<i>Group 2</i>				
Enzyme	498	1.069	1148	0.097
Escherichia-Coli	552	3.251	1500	0.093
RNA	393	1.383	1097	0.120
Transcription	385	1.795	974	0.120
Receptor	1,328	0.941	3513	0.097
<i>Group 3</i>				
Tropomyosin	7	0.012	62	0.015
Peroxisomal	6	0.023	56	0.023
Aplysia	4	0.015	102	0.026
Photosystem-Ii	4	0.006	80	0.007
Dynein	3	0.038	89	0.062

Legend: To illustrate the construction of keyword weights, we have chosen representative words in three categories. Group 1 keywords are typical of those that appear frequently in the work of patenting scientists, and infrequently in the work of non-patenting scientists. These words receive high patentability weights. Group 2 comprises keywords that occur frequently in the journal articles of both patenting and non-patenting scientists. Words in this group garner intermediate weights. Group 3 contains keywords that are very common in the research of non-patenting scientists but uncommon in the work of patenters. In consequence, these keywords receive low weight.

Table and Figure A2: Sensitivity Analysis

α	<i>Estimate of the Treatment Effect</i>	<i>naïve SE</i>	<i>bootstrap SE</i>	<i>95% Confidence Interval (based on bootstrap SE)</i>	
0.00	0.253	0.043	0.040	0.174	0.331
0.10	0.219	0.043	0.040	0.140	0.298
0.20	0.186	0.044	0.040	0.106	0.265
0.30	0.153	0.045	0.041	0.073	0.233
0.40	0.121	0.045	0.041	0.041	0.201
0.50	0.089	0.046	0.041	0.008	0.170
0.60	0.058	0.046	0.042	-0.024	0.139
0.70	0.027	0.047	0.042	-0.056	0.109
0.80	-0.004	0.047	0.042	-0.087	0.079
0.90	-0.035	0.048	0.043	-0.118	0.049
1.00	-0.065	0.049	0.043	-0.149	0.020



Effect estimates and 95% pointwise confidence bands are graphed versus the scale parameter α .