Corporate Ownership and Firm Performance: Evidence from Fertility Clinics*

Ambar La Forgia
Haas School of Business
University of California, Berkeley
ambar@berkeley.edu

Julia Bodner
Copenhagen Business School
jbo.si@cbs.dk

ABSTRACT

Corporate investors are often credited with boosting the performance of acquired firms but have also been criticized for generating profits at the expense of consumer well-being. This tension between shareholder and stakeholder interests is particularly evident in the healthcare sector, where information frictions can contribute to underinvestment in quality. This paper finds that corporate ownership can improve healthcare outcomes in a setting where patients have access to service pricing and quality information – the market for In Vitro Fertilization (IVF). We construct a novel dataset on US fertility clinic chains between 2004 and 2018, during which time chain ownership grew from 5% to 20%. Estimates from a difference-in-differences model show that post-transaction clinic volume increases by 25.8%, and IVF success rates increase by 7.0%. Direct ownership of clinics drives this increase in IVF success rates: acquired clinics exhibit a 13.6% increase, while affiliated clinics experience no change. We also provide evidence that fertility chains implement changes to clinic processes and procedures that enhance quality, lead to larger improvements among underperforming clinics, and expand the IVF market instead of capturing market share from independent clinics.

*We are grateful to Morten Bennedsen, Zack Cooper, Mathijs De Vaan, Paul Gertler, Matt Grennan, Atul Gupta, Hans Christian Kongsted, Adam Leive, Adam Sacarny, Toby Stuart, Ashley Swanson, and Steve Tadelis for their comments. We also thank several private equity fund managers and fertility service providers (who have asked to remain anonymous) for their business and clinical insights.

1. INTRODUCTION

Corporate investors can play a crucial role in generating firm growth and value. For example, by improving managerial processes and easing financial constraints, corporate ownership has been shown to increase performance in the manufacturing, retail, and service sectors (Bloom, Sadun, and Van Reenen 2015; Bernstein and Sheen 2016; Fracassi, Previtero, and Sheen 2022). However, corporate takeovers often generate private benefits for shareholders at the expense of societal benefits or value for stakeholders (Shleifer and Summers 1988). The potential misalignment between shareholder financial interests and consumer well-being is a growing concern in the healthcare sector, with most recent studies documenting a decline in quality after corporate acquisitions (Eliason et al. 2020; Gupta et al. 2021).[1] In contrast, this paper provides evidence that corporate ownership can lead to better healthcare outcomes using data on US fertility clinics.

The market for fertility services provides distinct advantages to studying the impact of corporate ownership. Most healthcare markets are characterized by information frictions that can contribute to underinvestment in quality and dull the market's ability to discipline the behavior of corporate entities (Arrow 1963; Dranove and Satterthwaite 1992; Gaynor 2006). The business model of fertility clinics, by contrast, better resembles that of the retail and service sector, where both the price and quality of goods and services are more readily observable. One of the main services used to treat infertility is In Vitro Fertilization (IVF), which has the singular goal of producing a live baby. IVF can cost up to $20,000 per cycle, and patients typically pay for IVF upfront and out-of-pocket because of limited insurance coverage. Despite the high cost, the probability of success is low: For women under 35, the live birth rate per IVF cycle is 40.5%, dropping to 4.5% for women over 42 (ASRM 2012). Out of concern that patients were not informed about low IVF success rates, congress passed a quality transparency law in 1992 that made fertility clinic report cards publicly available.[2] The transparency of IVF prices and quality may help align shareholder and patient interests and explain why the effect of corporatization differs across industries.

To study the impact of corporate ownership, we combine hand-collected data on fertility clinic transactions from business intelligence databases with clinic-level data from the Centers for Disease Control (CDC) Fertility Clinic Success Rates Reports between 2004 and 2018. We identify 11 fertility chains, which are groups of clinics that are owned by or affiliated with a single for-profit corporation. Out of 528 fertility clinics,

[1] Corporate ownership refers to "corporate investors (e.g., public companies, venture capital/private equity firms, insurance companies, and health systems) acquiring a majority and/or controlling interest" in previously independent organizations (American Medical Association 2019).

[2] The Fertility Clinic Success Rate and Certification Act (FCSRCA) of 1992 requires all US clinics that perform IVF to report their outcomes to the CDC. Yearly reports are published with standardized information on IVF success rates and are widely used by patients (Bundorf et al. 2009; Kowitt 2020).

the share of clinics in a chain increased from 5% to over 20% between 2004 and 2018.[3] This growth has generated concerns that investors will put less emphasis on clinical or operational improvements and instead treat "fertility medicine as a cash cow" (Robbins 2017). However, fertility chains argue they can help clinics "deliver high-quality, convenient care to patients while implementing cost savings, improving processes, and driving growth" (Krause 2019). We estimate changes in clinic growth and quality using difference-in-difference methods, which compare clinics before and after corporate ownership to a control group of independent clinics. We focus on two key outcomes: 1) clinic volume, measured as the overall number of IVF cycles and transfers, and 2) the success of IVF treatment, measured as the live birth rate per transfer.[4]

Our results show that after a clinic joins a fertility chain, IVF cycles increase by 25.8%, IVF transfers increase by 21.6%, and the live birth rate increases by 7.0%. We investigate several mechanisms that could explain these performance improvements. Rather than improving outcomes, clinics could instead select younger or healthier patients. We do not find evidence of patient selection: results are quantitatively similar whether or not we include controls for patient characteristics and infertility diagnosis, and we do not find evidence that clinics that become part of chains systematically treat patients that would be more likely to experience IVF success. Fertility chains could also be better at selecting clinics that would generate performance improvements. While clinic selection is an inherent feature of this setting, we conduct analyses that mitigate concerns that clinic selection explains our results. First, event studies adjusted for staggered treatment timing show no observable pre-trends before a clinic transaction. Second, results are quantitatively similar in specifications using *state* × *year* or *market* × *year* fixed effects, which would account for state or market-level changes that could impact the demand or provision of fertility services. Results are also quantitatively similar when excluding markets that became more concentrated because of corporate ownership. Third, we find quantitatively similar results using a matched sample based on pre-transaction clinic characteristics.

Corporate owners often claim to facilitate fertility clinic growth and quality improvements by providing clinics with access to resources and knowledge (The Economist 2019; Kowitt 2020). Through text analysis of press releases, we confirm that fertility chains advertise providing financial resources such as capital and managerial resources such as revenue cycle management and marketing and patient engagement services. Press releases also suggest that clinics receive access to best practices, protocols, and training and that fertility chains facilitate knowledge sharing between clinics through research consortiums and complex case review meetings. While we cannot

[3] By 2018, 9 of the 11 fertility chains were acquired by PE firms, and 2 chains were acquired by larger international healthcare chains. We define treatment time as the year a clinic became part of a chain.

[4] The live birth rate is calculated as the number of live births divided by the number of transfers. Any birth with at least one live-born infant is counted as a single live birth (i.e., twin births are counted as one live birth). Since age is the most important predictor of IVF success, we present the live birth rate by patient age category (under 35, 35-37, 38-40 and, 41-42) and as a weighted average.

precisely measure units of resources or knowledge being transferred, we use the data at our disposal to show patterns consistent with resource and knowledge transfers leading to improved outcomes.

Differences in ownership and control may influence the corporate parent's ability to transfer resources and knowledge. In this setting, some clinic transactions are structured as acquisitions: the parent company of the fertility chain owns and manages the clinic's assets. Other transactions are structured as affiliations: a clinic contracts with a fertility chain for management services and capital or financing options, which may involve no or partial ownership of clinic assets. In an acquisition, the corporate parent has greater control over clinical processes needed to improve performance, whereas, in affiliations, clinic owners retain greater decision-making authority (Grossman and Hart 1986). Greater corporate control is likely more important for facilitating knowledge sharing needed to increase IVF success than it is for resource sharing needed to fund clinic growth. [5] Consistent with this argument, acquired and affiliated clinics both increase volume, but only acquired clinics increase live birth rates (13.6% increase).

We also show that clinics change processes and procedures in ways that enhance quality and are consistent with new knowledge leading to improved outcomes. For example, *acquired* clinics achieve the IVF "gold standard": they reduce multiple births, which pose significantly higher risks for patients, and increase singleton births by enough to have a net positive increase in the live birth rate. We find that this quality-enhancing result coincides with acquired clinics reducing the number of embryos placed in the uterus per transfer, suggesting that clinics improve techniques and processes when conducting single embryo transfers (Reimundo et al. 2021; Mizrachi and McQueen 2022).[6] We also find that acquired clinics achieve higher IVF success rates among older patients, whose cases are often more complex. One reason for these improvements is that acquired clinics significantly increase the use of preimplantation genetic testing, which can help physicians choose higher-quality embryos, especially in older patients (Maxwell and Grifo 2018; ACOG 2020). Lastly, we find that the initially lowest-performing clinics experience the largest improvements relative to the highest-performing clinics, and that clinics acquired by higher-quality chains experience larger increases in live birth rates.

In support of chains providing resources needed to expand clinic operations and attract new patients, we find that fertility clinics lead to market expansion rather than business stealing from independent clinics. For every IVF cycle performed by a chain clinic in a market, there is one additional IVF cycle in that market and no reductions in IVF cycles for independent clinics. As another strategy to understand the role of resource transfers, we study differences in outcomes based on PE investment into fertility chains.

[5] Depending on the terms of the contract, chains may also be reluctant to transfer knowledge to affiliated clinics if they are "free to walk away at any time with the acquired knowledge" (Garicano and Rayo 2017).
[6] For each transfer, a physician can transfer one embryo or multiple embryos. A transfer of multiple embryos has a higher initial success rate but has a greater chance of multiple birth. A single embryo transfer has lower initial success rates but less than 1% probability of a multiple birth. Therefore, it is more difficult to increase the live birth rate via single embryo transfers.

Consistent with PE firms easing financial constraints and facilitating clinic growth, we find that PE investment into fertility chains primarily drives increases in clinic volume.

Overall, these results are consistent with fertility chains providing access to new resources and knowledge needed to increase clinic volume and IVF success rates. The positive impact of corporate ownership on fertility clinic quality is considerably different from the predominantly negative or null effects on quality found in the recent healthcare literature.[7] For example, Eliason et al. (2020) find that patients of dialysis clinics experience increases in mortality and hospitalization following acquisitions by a large corporate chain. Similarly, Gupta et al. (2020) find that PE acquisitions of nursing homes increase short-term mortality and decrease the reported well-being of patients. One explanation for these negative outcomes is that information frictions prevent healthcare markets from functioning effectively.[8] Specifically, information asymmetries between medical experts and patients make it difficult for patients to assess the quality of care. Government subsidies and insurance coverage also make it difficult for patients to observe and internalize the cost of care.

In contrast, the price and quality of services are readily observable in fertility markets, resulting in fewer information frictions. Fertility clinics often list their prices online and are extensively discussed with patients before treatment, and patients assume the majority of these costs. The main goal of IVF – to have a live baby – is also straightforward, and outcome data is readily available to prospective patients. In fact, Bundorf et al. (2009) found that the introduction of fertility clinics report cards led consumers to alter their choice of clinics. The salience and transparency of the live birth rate may motivate corporate owners to invest in quality to attract patients (Jin and Leslie 2003; Dranove and Jin 2010). The limited role of third-party payers may also lead corporate owners to engage in more price competition to increase IVF affordability and gain market share (Brown 2019; Sinaiko 2019).[9]

The features of the fertility setting help to reconcile the different impacts of corporate ownership observed in healthcare relative to other settings. For example, our findings are consistent with the positive effects of corporate ownership on health ratings in restaurants (Bernstein and Sheen 2016), sales of consumer goods in retail stores (Fracassi, Previtero, and Sheen 2022), workplace safety in publicly traded firms (Cohn, Nestoriak, and Wardlaw 2021), and on managerial practices in manufacturing firms (Bloom, Sadun, and Van Reenen 2015). Additionally, our results complement Gandhi et al. (2020), who find that only after CMS introduced a 5-star rating system did PE-

[7] Focusing on corporate acquisitions of hospitals, Andreyeva et al. (2022) document negative effects on hospital readmissions, Cerullo et al. (2022) find no changes in patient outcomes except for a slight improvement in mortality among Medicare beneficiaries hospitalized with AMI, and Gao, Sevilir, and Kim (2021) find no changes in patient experience or mortality. La Forgia (2022) finds that physician practice acquisitions by management companies that provide financial services lead to reductions in quality.

[8] An emphasis on financial outcomes at the expense of quality has also been reported in higher education, a setting with similar features to healthcare (Eaton, Howell, and Yannelis 2020).

[9] For example, fertility chains argue that efficiencies driven by economies of scale enable lower costs to be passed on to patients and advertise payment programs to help patients finance fertility (see Section 2.4).

backed nursing homes divert resources toward the measures evaluated by the system. Altogether, these results provide evidence that increasing price and quality transparency in healthcare can help align shareholder profit motives with key patient interests.

This paper also provides insights into the increasingly complex ownership structures accompanying mergers and acquisitions in healthcare. There is substantial evidence that healthcare consolidation leads to higher prices (see Gaynor, Ho, and Town (2015) for a review) with mixed effects on quality (Koch, Wendling, and Wilson 2017; Dunn and Shapiro 2018; Beaulieu et al. 2020). This paper extends the literature by studying the direct effect of corporate ownership and, in particular, sheds light on the chain business model common to other industries. By comparing acquired clinics to affiliated clinics, our findings suggest that affiliations can facilitate resource transfers necessary to improve volume but may not generate sufficient incentives or alignment to invest in quality improvement. The influence of ownership and control complements differences found in acquisitions involving directly owned vs. franchised restaurants (Bernstein and Sheen 2016), majority vs. minority-owned power plants (Demirer and Karaduman 2022), and in-network vs. out-of-network managers (Braguinsky et al. 2015).

Lastly, this paper contributes to the research on the economics of the fertility industry. Much of the literature has focused on the impact of state fertility coverage mandates on the utilization of fertility services and treatment choices (Schmidt 2007; Henne and Bundorf 2008; Bitler and Schmidt 2012; Hamilton et al. 2018). Despite expansions in state-mandated IVF coverage, the cost of IVF in the US still constrains usage to a homogenous patient population: patients utilizing IVF are predominantly white, privately insured, high-income, and highly educated (Chandra, Copen, and Stephen 2014; Galic et al. 2021). Though we are unable to measure price changes, we find that corporate ownership appears to expand access to IVF and improve IVF success rates. In the long term, these improvements may enable more women to delay motherhood, invest in education and establish their careers, contributing to greater gender equality (Gershoni and Low 2021).

2. THE EMPIRICAL SETTING

2.1 The Fertility Industry

The main providers of infertility services are fertility clinics, which assist couples or individuals who wish to conceive but are unable to naturally. The most effective way to treat infertility is through Assisted Reproductive Technology (ART), where In Vitro Fertilization (IVF) represents over 99% of ART procedures. In the U.S., fertility clinics are predominantly for-profit businesses, and treatment costs are remarkably high.[10] The cumulative cost of IVF is estimated to be between $40,000 and $60,000 because the

[10] Fewer than 15% of fertility clinics are part of an academic medical center (AMC), a majority of which are nonprofit (Patrizio et al. 2022). While the AMC may be nonprofit, the fertility clinic itself may still be organized as a for-profit subsidiary, which can be owned by physicians (Appelbaum and Batt 2020).

average patient undergoes multiple IVF cycles (Fertility IQ 2022). These costs are either financed privately by patients or subsidized by insurance companies. However, even with some coverage, patients pay for the majority of services out of pocket, which can amount to 50% of disposable household income (Chambers et al. 2009; McLaughlin et al. 2019).[11]

In 2021, the global fertility industry was valued at $26 billion, and analysts forecast continued growth at 7% annually (Zion Market Research 2022). By the end of the century, it is estimated that nearly 400 million babies will have been born via IVF (Faddy, Gosden, and Gosden 2018). The growing demand for infertility services is driven by heterosexual couples delaying parenthood, as well as more single individuals and same-sex couples choosing to have biological children (Kaufman 2020; Witte 2020). This projected demand and the high margins of fertility clinics have attracted considerable investment from PE firms, venture capitalists, and management companies (Landi 2022; Pringle 2022). This investment has transformed the industry from stand-alone clinics to large fertility chains. For example, a recent study found that nearly 15% of clinics in 2018 were owned by PE firms (Borsa and Bruch 2022), with dozens of deals announced in the subsequent years (Patrizio et al. 2022).

The distinct characteristics of the fertility sector potentially makes it more amenable to performance improvements following corporate ownership. The salience of IVF success rates and transparency in prices and outcomes across clinics may motivate corporate owners to invest in the resources and knowledge needed to generate improvements and attract patients. Additionally, within the US, there is minimal government regulation over IVF processes and practices, and federal funding of embryo research is explicitly banned (See Calandrillo and Deliganis (2015) for a review). This contrasts with countries like the UK, which regulate the storage of embryos, and requires that all clinics comply with a code of practice before licensure. France and Sweden have even imposed bans on the number of embryos that can be transferred during IVF to minimize multiple births. In the US, fertility clinics have considerably more autonomy over clinical and operational decisions. There are also few areas of healthcare with such a homogenous patient population. The fragmented organization of clinics, the minimal regulations, and more homogenous patients may give corporate owners a greater ability to standardize care.

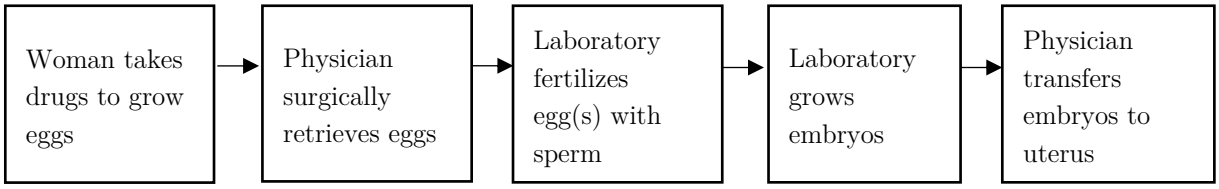
2.2 The IVF Process

This paper focuses on fertility clinics that provide IVF. A fertility clinic is typically comprised of a medical office for patient visits and procedures and a laboratory for creating, testing, and storing eggs and embryos. Clinics employ reproductive

[11] Insurance coverage varies widely by state and employer. State coverage mandates do not apply to self-insured companies or smaller companies and often only cover some costs of care, such as fertility-enhancing drugs but not necessarily IVF. Still, some coverage can considerably reduce the total out-of-pocket costs associated with IVF and influence treatment decisions (Hamilton et al. 2018). For this reason, our primary specification includes *state* × *year* fixed effects to account for any state-level policy changes.

endocrinologists, embryologists, nurses, surgical technicians, and administrators who work collaboratively to provide fertility services. While the desired outcome of IVF is straightforward, the process of achieving a live birth is complex. The IVF process consists of several stages that involve over 350 steps performed over 4 to 6 weeks per cycle (McCaffrey, Forman, and Copperman 2022). At a high level, a patient undergoes the five phases of treatment shown in Figure 1. Patients often need several cycles to achieve a live birth, with many patients undergoing at least 3 IVF cycles.

Figure 1. The Basic IVF Process of a Single Cycle



Note: Author's illustration adapted from Fertility IQ "What is IVF?"

Each step of the IVF process depicted in Figure 1 involves subjective decisions that contribute to variation in fertility outcomes across clinics and physicians (Lintsen et al. 2010; Mizrachi and McQueen 2022; Morin 2022). For example, identifying and grading embryo quality and, therefore, which embryos to transfer to the uterus is considered a subjective assessment (Schoolcraft and Meseguer 2017).[12] Similarly, Mizrachi and McQueen (2022) conclude that differences in physician embryo transfer techniques, but not experience, drive much of the variation in the success of an embryo transfer. This variation is highlighted by an internal study conducted by a fertility chain which found that for comparable patients treated in the same clinic, the probability of a live birth varied from 50.6% for the lowest-performing physician and 66.4% for the highest-performing physician (Morin 2022).

Differences also exist in the core decision made between a reproductive endocrinologist and a patient on whether to transfer a single embryo or multiple embryos: Transferring multiple embryos increases the success of pregnancy but also results in multiple births in 30% of pregnancies relative to single embryo transfers (Kissin, Boulet, and Jamieson 2016). Multiple births are associated with significant fetal and maternal risks, such as pre-term delivery, low birth weight, and pre-eclampsia. These worse outcomes led the American Society for Reproductive Medicine (ASRM) to issue changes in recommended IVF guidelines to lower multiple birth rates by encouraging single embryo transfers (ASRM 2013, 2017). Therefore, the embryo transfer decision

[12] Other factors that a physician can influence include the choice of ovarian stimulation protocol, the choice of hormone medications to achieve follicle growth, and the timing of the ovulation trigger (USCFertility 2022). There is also variation in technical skill and expertise when determining sperm and egg quality, as well as during the fertilization process and embryo transfer (Morin 2022).

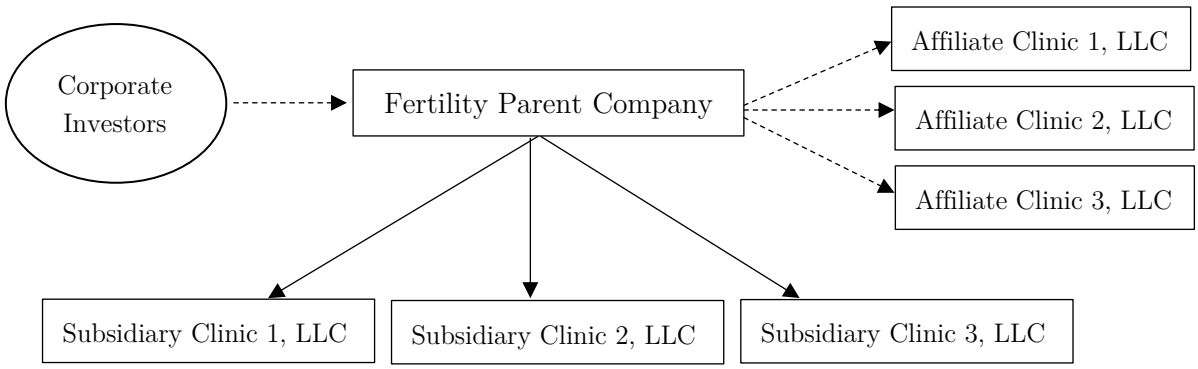
generates a tension between increasing a clinic's live birth rate through multiple embryo transfers and complying with large-scale efforts to reduce multiple births from IVF.

2.3 Organization of Fertility Chains

This paper focuses on fertility clinics that become part of a fertility chain. A fertility chain represents a group of clinics that are owned by or affiliated with a single for-profit corporation that is either publicly traded or receives investment from PE firms.[13] In these chains, fertility clinics are connected to a fertility parent company, as depicted in Figure 2. In *acquisitions*, the fertility parent company acquires, owns, and directly manages the assets of the fertility clinic.[14] In *affiliations*, the clinic signs a contract with the fertility chain for select management services and capital or financing options. For example, a clinic may affiliate with a chain to receive access to marketing and patient engagement services. The contracts may resemble outsourcing agreements in which the parent company has no ownership stake or may be structured as joint ventures in which the parent company has a partial ownership stake. The commonality of these affiliations is that the clinic owners maintain greater control of clinic operations than in an acquisition. Some corporate entities pursue a mix of ownership structures, while others focus more exclusively on acquisitions or affiliations.

Fertility chains often receive external investment from PE firms. PE firms invest in chains in two ways. They can either acquire a pre-existing fertility chain to help finance further growth or invest in a renowned clinic to establish that clinic as the new fertility parent company of a chain (Zoeller, Muller, and Janiga 2020). In Section 3.1 we will provide more details on the characteristics of fertility chains.

Figure 2. Organization of a Fertility Chain



[13] Healthcare providers are often reluctant to use business terminology such as "chain" and, instead, use terms such as "network." In the fertility setting, chain clinics share in central management and standardized business practices, but rather than big box-style rebranding, clinics signal chain membership on their websites and marketing materials.

[14] Corporate practice of medicine (CPOM) laws often prohibits non-physician-owned business entities from exerting undue influence on clinical practice. On paper, corporations comply with these laws using complex corporate structures involving a subsidiary management service organization purchasing the non-clinical assets of a clinic. In practice, the corporations still exert significant control over day-to-day clinic operations, generating controversy over the efficacy of CPOM. See Appendix A for details.

2.4 Value Proposition of Fertility Chains

Press releases, company websites, news articles, and industry reports provide insights into the goals, strategies, and service offerings of fertility chains. These materials suggest that chains seek to increase clinic performance by facilitating knowledge sharing among clinics and providing them with financial resources and managerial capabilities. For example, one report claims, "private equity firms are taking stakes in clinics and pushing for consolidation in hopes of seeing cost savings, process improvement, and growth" (Cantrell 2019). Beyond just private equity, "clinics united into national chains have been sharing best practices, introducing newer technologies, and offering more flexible payment plans for customers" (Robbins 2017). Below we explore this anecdotal evidence surrounding resource and knowledge transfers within fertility chains. All sources and documentation of quotes are reported in Appendix Table B1.

Financial Resources and Managerial Capabilities. One reason fertility clinics join fertility chains is to gain access to resources. For example, several chains emphasize providing clinics with long-term "financial stability and growth opportunities" and "strong financial support" from PE firms. These resources can help clinics hire new physicians and build new locations: "[Fertility chain] plans for continued growth through the addition of physicians and satellite offices." Similarly, another chain advertises that they apply "business and operations strategies that expand [clinic] markets and their market share. This may involve the development of new practice locations, embryology laboratories or ambulatory surgery centers, in order to […] achieve strategic growth objectives." Ultimately, these investments can help increase clinic profitability. One chain advertises that "clinics practices' patient revenues increased 21% from 2007 to 2009" because of access "to capital and best-in-class business and clinical services."

Marketing materials also highlight that fertility chains provide managerial resources to streamline back-office administration. Since fertility clinics are typically run by physicians focused on clinical medicine and not trained in business practices, clinics may benefit from better management practices. For example, one chain advertises providing "operational and financial management, revenue cycle management, patient marketing and sales, information systems support, and various other services, including patient support." One managerial capability that is particularly highlighted by various chains is marketing, where one chain suggests its clinics should expect "increased patient volume as a result of [the Fertility Chain's] marketing efforts."

Chains also attract patients by offering and heavily marketing new pricing models that help patients finance treatment. For example, one fertility chain has its own subsidiary fertility financing company where "the company's Fertility Loan Specialists will work closely with [the Fertility Chain] to ensure the funds are secured prior to the commencement of [patient] treatment." Similarly, one chain launched "IVF Refund and Multi-Cycle Programs [that] offer patients the assurance that if multiple IVF cycles are necessary, they will not need to expend additional financial resources to receive them."

Clinical Knowledge. A second cited benefit of joining a chain is the ability to share and generate clinical knowledge with other clinics. For instance, a fertility chain advertises that it was "created to break down barriers to idea-sharing and collaborative care." This sentiment is echoed by a fertility clinic citing "access to […] the most advanced on-going research in the field of reproductive medicine" as a reason for joining a chain. Similarly, a clinic owner suggests their patients will benefit from "improved access to the best treatment protocols and unique programs for specific conditions, […] increased access to clinical trials and research initiatives […] access to an expanded network of […] experts who will come together to review and assist in complex cases." Multiple physicians echo that chains help standardize clinics' practices via treatment protocols. For example, one physician expects "to further revolutionize patient care with new access to proven treatment protocols."

Fertility chains also create internal processes to facilitate knowledge sharing and establish best practices within the chain. Many chains create committees with physicians across clinics who meet regularly to discuss clinical research, patient cases, and in some instances, conduct their own research and clinical trials. For example, one chain states that when "research proves that techniques improve conception rates, [Fertility Chain] incorporates those techniques into their standard care wherever possible." Another chain says that "treatment breakthroughs are quickly applied to multiple centers, thereby furthering the positive impact for patients." Chains also advertise using "proprietary platforms, applications, and data and analytics" to track clinic performance and help clinics improve their clinical processes. Lastly, some chains implement continued medical education and training programs to improve IVF success rates. For example, the CEO of one chain shared: "We'll look at pregnancy per transfer by physician with a blinded letter for each physician. And we'll be able to see how everybody stacks up. And if people fall below a standard deviation, we have that doctor go work with somebody who is above a standard deviation to get retrained."

Fertility chains also advertise strategic goals for the organization that are in line with the latest medical research. For example, many chains advertise increasing single embryo transfers: "Striving for One Embryo-One Baby. [Fertility Chain's] founding philosophy to achieve successful pregnancy one healthy baby at a time." This goal is likely motivated by the ongoing efforts from professional associations encouraging the reduction of multiple births, leading fertility chains to advertise achieving lower multiple birth rates as both a marketing and reputation tool. Additionally, the chains want to attract employers who offer subsidized fertility benefits to their employees but are sensitive to the much higher costs associated with multiple births.[15] Altogether, the available documentation suggests that when clinics transition from independent,

[15] A multiple birth can cost up to 20 times for than a singleton birth. Therefore, self-insured employers have an incentive to reduce multiple births to reduce the birth costs they would incur from offering fertility benefits (Lemos et al. 2013). In fact, several fertility benefits management companies advertise only partnering with clinics with low multiple birth rates to attract employers (Winfertility 2022).

standalone clinics into fertility chains, they receive access to resources and knowledge meant to improve financial and clinical performance.

3. DATA AND DESCRIPTIVE STATISTICS

3.1 Data Description

We construct a panel dataset of fertility clinic transactions to estimate the impact of corporate ownership on clinic performance. We combine data from several sources to create a novel dataset of corporate-backed fertility chains and outcomes between 2004 to 2018. See Appendix A for additional details on data construction.

Clinic Characteristics. All clinics that perform Assisted Reproductive Technology (ART) must submit data to the CDC annually under the Fertility Clinic Success Rate and Certification Act (FCSRCA) of 1992.[16] ART includes all fertility treatments in which either eggs or embryos are handled; over 99% of ART is IVF. The CDC then compiles and publishes Fertility Clinic Success Reports (download here: https://www.cdc.gov/art/artdata/index.html), which are meant to inform prospective patients of their probability of achieving a live birth. We will refer to these data as the CDC ART data. While the data are consistent within a year, the variable names and data collected have undergone considerable changes over time, limiting which variables can be studied in a panel framework. After extensive data cleaning, we create a consistent clinic identification number and identify each clinic's service offerings, patient infertility diagnoses, number of IVF cycles and transfers, and IVF success rates. We also use PDF versions of the CDC ART data to extract additional information, such as clinic addresses and laboratories used.

Market Characteristics. We define a market as a clinic's Core Based Statistical Area (CBSA), which consists of one or more counties with an urban center of at least 10,000 people plus adjacent counties that are socioeconomically tied to the urban center by commuting.[17] Fertility clinics in the sample are present in 145 out of 927 CBSAs (See Harris et al. (2017) for details on the geographic distribution of fertility clinics).[18] Past research on fertility clinic competition has defined the market for fertility services as a

[16] To ensure data quality, the medical director of a clinic must verify by signature that the success rates are accurate. Additionally, a random sample of clinics is audited each year, and a validation team examines the clinics' medical records and compares them to the reported data.

[17] The choice of fertility clinic is not typically related to distance in areas with multiple clinics. Patients are often "willing and able to travel long distances to use the provider of their choice regardless of distance, time, or expense" (Harris et al. 2017). Around half of all clinics also have at least one satellite clinic, with a mean driving time of 66 minutes between the satellite and main clinic, which can greatly expand a clinic's catchment area (McGarity et al. 2022).

[18] One fertility clinic in Alaska is not located in a CBSA and so was assigned its county-level market characteristics. Clinics in Puerto Rico are excluded from analyses with market-level controls because several variables are unavailable during the sample period.

metropolitan statistical area (Bundorf et al. 2009; Hamilton and McManus 2012). However, CBSAs, which include both metropolitan and micropolitan statistical areas, help capture additional clinics located in smaller urban areas. We use data from the US Census Bureau and the Bureau of Labor Statistics to obtain market-level population estimates, the median household income, and the unemployment rate. We also calculate the Herfindahl-Hirschman Index (HHI) based on a clinic's total IVF cycles to measure the competitiveness of the market.

Patient Characteristics. The CDC ART data are aggregated to the clinic level and do not contain patient information beyond infertility diagnoses. For secondary analyses, we use National Center for Health Statistics (NCHS) Vital Statistics Data to account for market-level patient demographics for women who gave birth after receiving any infertility treatment. The NCHS data include a mother's race/ethnicity, educational attainment, insurance, and clinical conditions such as hypertension and diabetes, but are only available from 2009-2018 with an indicator for receiving infertility treatment.

Corporate Ownership. We identify clinic transactions through press releases, archived versions of clinic and fertility chain websites, the CDC ART data, and the following business intelligence databases: Irvin Levin, SDC Platinum, and Pitchbook data. These business sources often provide the announced date and descriptions of the terms of the transaction.[19] Based on their ownership structure, we classify fertility clinics into three main categories: 1) Acquisition: the clinic's assets (office, lab, or both) are acquired and owned by the corporate parent; 2) Affiliation: a clinic contracts with a fertility chain for selected management services and capital or financing options, and the corporate parent either has no ownership rights or partial ownership, and 3) Independent, a clinic that is never part of a fertility chain during the sample period. To our knowledge, independent clinics are never corporate-owned or part of a fertility chain during the sample period.[20] Additionally, fertility chains often build new clinics as part of their growth strategy, which we classify as a "de novo" clinic.

We identify 11 fertility chains between 2004 and 2018 (see Appendix Table A1 for details). These chains match those identified and discussed in industry reports and articles studying fertility clinic business models (Dresner Partners 2018; Borsa and Bruch 2022; Patrizio et al. 2022). These chains can either form as the result of PE investment or exist as for-profit corporations before being acquired by a PE firm or larger corporate entity. In our sample, 5 chains were created by PE firms, 1 chain was publicly traded

[19] Databases such as Pitchbook often provide high-level detail on whether the transaction involved an acquisition, a joint venture, or other types of equity arrangements. When these data are unavailable, we analyze company press releases and websites, which often signal the type of transaction.

[20] An independent clinic may be a hospital-based clinic. Hospital-based clinics may also be part of a fertility chain. Some independent fertility specialists may be part of a multi-specialty women's health practice or a military health provider. Absent an acquisition or affiliation with a chain, these organizational forms are static from 2004 to 2018 and therefore, accounted for using clinic fixed effects.

before being acquired by a PE firm, 1 chain was formed as the subsidiary of a physician practice management company a PE firm acquired, and the remaining chains were pre-existing physician-founded corporations acquired by either a PE firm (2 chains) or an international health care chain (2 chains). We define the clinics in this set of 11 chains as the treatment group and treatment time as the year a clinic first became part of the chain. In additional analyses, we will also examine the timing of PE investment separate from the effect of joining the fertility chain.

3.2 Outcome Variables

We focus on two outcomes that measure clinic performance: 1) clinic volume and 2) success of IVF treatment. Clinic volume is measured as the total number of IVF cycles and the total number of IVF transfers performed in a year (including donor and non-donor cycles and transfers). A cycle starts with the intent of retrieving an egg for immediate fertilization or to be frozen for future use. A transfer represents the part of an IVF cycle when one or more embryos (the result of the fertilized egg) are transferred into the uterus of a woman with the intent to establish a pregnancy (SART 2021a). Not all cycles become transfers because eggs may not develop, the patient may become ill, the patient may choose to stop treatment, or the fertilization of the egg may not be successful, among other reasons (Bedrick et al. 2019). In regression analyses, we log the volume variables to better fit the distribution of the data (Appendix Figure C1).

The success of IVF is measured by the live birth rate. The live birth rate represents the number of live births divided by the number of transfers using fresh or frozen non-donor eggs (i.e., the patient's own eggs).[21] The CDC ART data reports the live birth rate by patient age bins (under 35, 35-37, 38-40 and 41-42). While we present live birth rates separately by age bins, in most analyses, we present the live birth rate as an age-bin weighted average. We also show results decomposing the live birth rate into the singleton birth rate and the multiple birth rate (i.e., twin births are counted as one live birth). All live birth measures focus on cases where at least one embryo was transferred within 12 months of the start of the cycle; therefore, fertility preservation cycles where patients freeze their eggs or embryos for future use with no intent to become pregnant within 12 months are excluded.

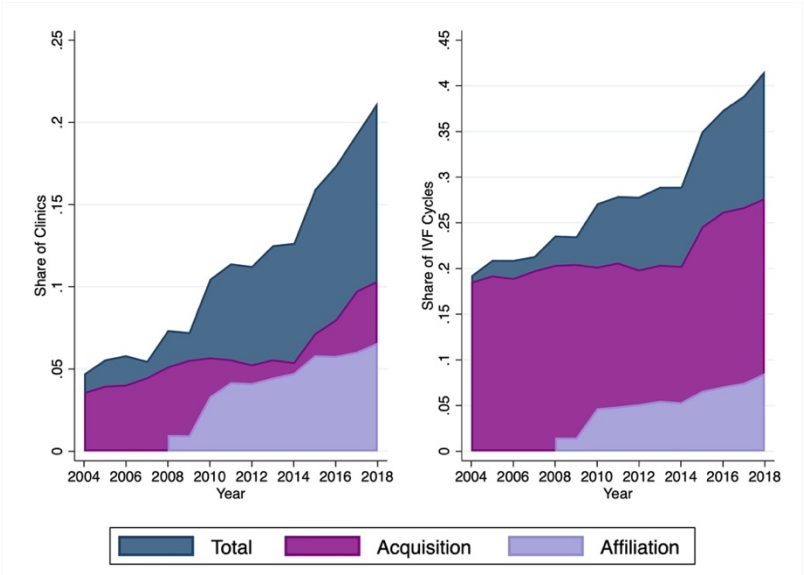
3.3 Descriptive Statistics

The final analytic sample includes 528 clinics and 6,274 clinic years. To construct this sample, we 1) exclude data reported for patients over the age of 42 because of changes in data availability over the sample period, 2) exclude clinics that perform fewer than 20 cycles a year on average, and 3) exclude clinics that are in the sample for less

[21] An IVF cycle may result in only one viable embryo or multiple viable embryos. With multiple viable embryos, one or more embryos may be transferred within a few days of creation (a fresh embryo transfer), with the remainder frozen for future use. If the fresh embryo transfer is unsuccessful, the frozen embryos can be thawed and transferred subsequently.

than 3 years (except for de novo clinics).[22] Figure 3 shows that by 2018, over 20% of clinics are in a fertility chain and perform over 40% of IVF cycles in the country. In total, there are 62 corporate transactions: 33 clinics are structured as acquisitions, and 29 clinics are structured as affiliations. Additionally, 23 clinics are newly built by a chain, 15 clinics are always in a chain, and 428 are never part of a chain (i.e., independent clinics). However, in most analysis, we exclude 14 independent clinics with multi-year gaps in reporting to the CDC, resulting in 414 independent clinics.[23]

Figure 3. Share of Clinics and IVF Cycles in a Fertility Chain



Note: The left-hand figure shows the share of clinics in a fertility chain each year, and the right-hand figure shows the share of IVF cycles performed by corporate clinics. "Total" includes all clinics ever part of a fertility chain, including those always in the chain or that were newly built by the chain.

Table 1 provides additional fertility clinic and market-level characteristics. Prior to a corporate transaction, clinics appear to perform more cycles and transfers and have higher live birth rates than independent clinics, suggesting that chains may target better-performing clinics. De novo clinics also provide evidence of the role of corporate ownership as they also experience greater volume and IVF success compared to independent clinics. Despite differences in these outcomes, patients do not appear to be inherently different across clinics: there are similar distributions of patients under 35 and of patient diagnoses for infertility. Similarly, a mother's reported education, race/ethnicity, insurance type, and health factors are comparable across clinic categories (Appendix Table C1). These statistics confirm that most patients who use infertility treatment are white, privately insured, and highly educated.

[22] We also exclude 17 clinic-years when a clinic has fewer than 10 IVF cycles in their first or last year of data, as this signals a clinic opening or closure and may not accurately reflect a clinic's fertility program.
[23] Multi-year gaps typically occur if a clinic paused IVF cycles to restructure or substantially update its clinic. Clinics could also have been failing to adhere to FCSRCA reporting requirements. Including these clinics in the control group yields slightly larger effect sizes than those in Table 2.

Table 1. Fertility Clinic Summary Statistics, 2004-2018

	Fertility Chain			Independent
	Acquisition	**Affiliation**	**De Novo**	
	Pre-transaction mean	Pre-transaction mean	Mean of all years	Mean of all years
Clinic Volume				
IVF Cycles	547.36	493.61	491.42	284.17
IVF Transfers	449.40	392.30	363.52	223.19
Log(IVF Cycles)	6.04	5.88	5.82	5.16
Log(IVF Transfers)	5.83	5.66	5.47	4.94
Birth Rates (%)				
Live Birth Rate	41.67	41.80	40.89	36.78
Singleton Birth Rate	30.40	30.86	29.99	26.87
Multiple Birth Rate	11.26	10.97	10.87	9.84
Patient Characteristics (%)				
Share of Patients < 35 (transfers)	51.64	51.88	50.94	51.30
Share of Patients 35-37 (transfers)	24.21	23.23	22.63	23.54
Share of Patients ≥ 38 (transfers)	24.16	24.89	26.43	25.16
Diagnosis, Tubal Factor	11.16	11.17	7.79	12.93
Diagnosis, Ovulatory Dysfunction	11.30	11.67	9.13	12.45
Diagnosis, Diminished Ovarian Reserve	24.10	23.34	20.75	21.43
Diagnosis, Endometriosis	7.96	7.74	6.50	8.03
Diagnosis, Uterine Factor	3.79	4.12	1.65	4.04
Diagnosis, Male Factor	25.32	23.70	21.58	26.96
Diagnosis, Other	12.97	12.55	19.11	11.06
Diagnosis, Unknown	10.41	11.36	11.69	9.72
Market Characteristics (CBSA)				
Total Population (Age 20-49)	1,498,221	1,777,190	2,721,922	1,910,081
Population Female (%)	50.23	50.46	50.17	50.29
Unemployment Rate (%)	6.24	5.97	5.65	6.08
Median Household Income ($)	55,362	54,168	62,843	58,475
Market Concentration (HHI)	4,373	3,666	4,023	4,410
Observations				
Number of Clinics	33	29	23	414
Clinic-Years	283	193	138	4948

Notes: All summary statistics are at the clinic-year level. Clinic volume, birth rates, and patient characteristics include adjustment for year effects to account for changes in reporting in the CDC ART data. Market concentration is calculated using a clinic's total IVF cycles. 15 clinics are excluded because they are in a chain before the sample period and 14 independent clinics are excluded because of multi-year reporting gaps. In total, there are 528 clinics and 6,274 clinic-years.

In Appendix Table C2, we present the results of a targeting regression to better understand the probability of a clinic acquisition or affiliation based on pre-transaction characteristics. The targeting results suggest chains target clinics that perform more IVF cycles, have a higher live birth rate, and, though marginally significant, target clinics in

more competitive markets with larger populations aged 20-49.[24] Additionally, clinics are less likely to be targeted in markets with a greater share of Medicaid patients.

Overall, these descriptive statistics help inform potential identification challenges. While there are differences in the types of clinics selected by fertility chains, there do not appear to be observable differences in the types of patients treated by clinics before the transaction. This pattern is consistent with the homogenous nature of patients treated by fertility clinics. Still, in the empirical analyses that follow, we use several strategies to account for differences between corporate and independent clinics.

4. The Effect of Corporate Ownership on Fertility Clinic Outcomes

Our empirical strategy aims to identify the causal effects of corporate ownership on fertility clinic volume and clinical performance and study potential mechanisms. Our primary strategy utilizes a difference-in-differences (DD) specification to compare changes in outcomes for fertility clinics before and after a corporate transaction (treated clinics) with concurrent changes for independent clinics that were never part of a fertility chain (control clinics). De novo clinics and clinics always observed in a chain are excluded from analyses unless otherwise specified. We estimate extensions of this DD model using an event study framework and a matching estimator, among other analyses, that together provide compelling evidence that corporate ownership positively impacts clinic volume and IVF success.

The preferred specification includes clinic fixed effects (θ_c) to adjust for time-invariant clinic characteristics and calendar *state* × *year* fixed effects (θ_{st}) to flexibly allow for time-varying factors that are common to all clinics in a state. We also include a vector of controls ($\boldsymbol{X}_{ct}$) that include an indicator for whether two clinics combined their data reporting to the CDC (i.e., had reported as two separate clinics but then began reporting as a single clinic) and an indicator for the first year a clinic was in the sample to account for partial year reporting when a clinic first enters the data.[25] Each estimation uses cluster-robust standard errors at the clinic level.

Eq. 1: $$Y_{ct} = \beta Post_{ct} + \gamma \boldsymbol{X}_{ct} + \theta_c + \theta_{st} + \epsilon_{ct}$$

Equation 1 is a within-clinic regression, where $Post_{ct}$ is a binary variable equal to one if clinic c is acquired by or affiliated with a fertility chain in year t. The coefficient of interest, β, captures the relationship between becoming part of the chain and Y_{ct}.

[24] Fertility markets are highly concentrated on average, in part, because 28% of clinics are monopoly or duopoly providers. However, 54% of clinics are in CBSAs with 5 or more clinics, and an HHI of 2134, suggesting that most clinics are in moderately concentrated CBSAs. See Appendix D for more details.

[25] We identify 10 clinics that combined reporting – this could be the result of a true merger, or the clinics remain separate entities that report under a single clinic. In either case, we take the weighted average of their outcome variables in each year and create an indicator variable equal to 1 post-merger. This is the most conservative approach; magnitudes are slightly larger when not accounting for these mergers.

Since there may be differences in clinic outcomes by ownership structure, we use interactions between $Post_{ct}$ and whether a clinic transaction was structured as an acquisition or affiliation, as seen in Equation 2.

$$\text{Eq. 2:} \qquad Y_{ct} = \beta_1(Post \times Acquisition)_{ct} + \beta_2(Post \times Affiliation)_{ct} + \gamma \boldsymbol{X}_{ct} + \\ + \theta_c + \theta_{st} + \epsilon_{ct}$$

The identifying variation is primarily based on the staggered timing of clinic transactions and the comparison of treatment and control clinics in their overlapping periods. To interpret the β coefficients as the causal effect of the transaction, we must assume that the trends in outcomes of these corporate clinics would have been similar to the trends in outcomes of independent clinics in the absence of the transaction. The concern with this identification strategy is that the timing of the transaction may be correlated with other contemporaneous factors that impact the outcomes, such as changes in the patient population and the non-random selection of clinics. Additionally, recent literature in econometrics has shown the issue of negative weights that arise from DD with staggered treatment timing because of comparisons not only between treated and control units but between already treated and eventually treated units (see Baker et al. (2022) and Roth et al. (2022) for reviews). We conduct a series of diagnostic and robustness checks and additional analyses that mitigate concerns of treatment effect heterogeneity and the role of patient and clinic selection.

4.1 Main Effect on Clinic Volume and IVF Success Rates

Table 2 shows the estimates of the pooled regression from Equation 1 in Panel A using both *state* × *year* (the preferred specification) and year fixed effects. After a clinic becomes part of a fertility chain, IVF cycles increase by 25.8%, and IVF transfers increase by 21.6%. There are also significant changes to IVF treatment success: The live birth rate increases by 2.6 percentage points (7.0% of the mean). However, pooling together clinics masks the heterogeneity in outcomes by ownership type. Table 2 Panel B reveals that while both acquired and affiliated clinics significantly increase clinic volume, only acquired clinics significantly increase the live birth rate. After an acquisition, the live birth rate increases by 5.1 percentage points (13.6%), and we fail to find evidence of changes in the live birth rate of affiliated clinics.[26]

Table 2 shows that effect sizes are similar when accounting for time-invariant differences across states rather than within states over time (Columns 2, 4, and 6). Additionally, we find quantitatively similar effects using *CBSA* × *year* fixed effects, which would account for market-level changes that could impact the demand or provision

[26] To assess whether the lack of statistical significance is due to low statistical power, Appendix E provides power curves based on simulation analysis. For affiliated clinics, we have 60% and 80% power to detect an effect size of 1.8 pp and 2.4 pp in the live birth rate and 9.5% and 12% in log IVF cycles, respectively (Appendix Figure E1). Power curves are similar for acquired clinics.

of fertility services (Appendix Table D1). Accordingly, results are not statistically different when including the logged values of the CBSA-level population aged 20-49 and median household income as controls (Appendix Table D2). We also show that outcomes are quantitively similar when excluding markets that became more concentrated because of corporation ownership (Appendix Table D3).[27]

Table 2. Effect of Corporate Ownership on Fertility Clinic Outcomes

	(1)	(2)	(3)	(4)	(5)	(6)
	Log(Cycles)		**Log(Transfers)**		**Live Birth Rate**	
Panel A: Pooled						
Post	0.258***	0.282***	0.216***	0.237***	0.026**	0.018*
	(0.071)	(0.063)	(0.071)	(0.063)	(0.011)	(0.010)
Panel B: Ownership Structure						
Post × Acquisition	0.282***	0.297***	0.214**	0.217**	0.051***	0.043***
	(0.098)	(0.087)	(0.099)	(0.088)	(0.013)	(0.012)
Post × Affiliation	0.238**	0.268***	0.217**	0.253***	0.004	-0.003
	(0.100)	(0.088)	(0.099)	(0.087)	(0.016)	(0.014)
Clinic FE	X	X	X	X	X	X
State × Year FE	X		X		X	
Year FE		X		X		X
Dep. Var. Mean	5.252	5.256	5.035	5.040	0.374	0.375
Clinic-Years	5666	5809	5666	5809	5666	5809
R^2	0.899	0.887	0.898	0.886	0.625	0.579

Notes: Panel A shows the β estimates of Equation 1, and Panel B shows the β_1 and β_2 estimates of Equation 2. The live birth rate is calculated as the number of live births divided by number of transfers. The dependent variable mean captures the predicted mean for control clinics and treatment clinics before the transaction. Standard errors are clustered at the clinic level. Statistical interpretations remain unchanged when using wild bootstrap standard errors to adjust for small sample sizes (Appendix Table D4). Significance levels: *p<0.1, **p< .05, ***p<0.01

4.2 Treatment Effect Timing

Using a DD research design with multiple periods and treatment times could result in "bad" comparisons between clinics treated earlier and clinics treated later in the sample. The diagnostic test developed by Goodman-Bacon (2021) decomposes treatment effects into multiple, weighted, two-by-two DD estimators.[28] In Appendix Table D5, we show that approximately 90% of the weight is attributable to "good" comparisons of treated to never-treated clinics, and less than 10% is attributable to comparisons between treated clinics at different times (i.e., early vs. later treated clinics)

[27] Most acquisitions or affiliations occur across markets. Only 3 CBSAs became more concentrated because of a corporate transaction (as measured by an increase in HHI).

[28] The Goodman-Bacon decomposition requires a balanced panel, which limits this analysis to clinics with 15 years of data (52% of clinics and 64% of observations). In Appendix Table F3 we show results of our primary specification in Table 2 are quantitatively similar in the balanced panel.

or to within clinic variation. Furthermore, the small amount of weight placed on within-clinic variation suggests the inclusion of controls does not drive our results.

Next, we use an event study framework to evaluate whether the treatment and control clinics had differential trends before acquisition or affiliation. The event study is an extension of Equation 2, where instead of aggregating years before and after a transaction, indicators are included for each year relative to the transaction year. To further address potential concerns of treatment effect heterogeneity, we utilize the two-stage DD method developed by Gardner (2021): the first stage regresses unit and time effects on the outcomes variable using only untreated and not-yet-treated observations, and the second stage regresses these adjusted outcomes on relative treatment time indicators in the full sample. The average treatment effect is "identified from a comparison of mean outcomes between treated and untreated groups after removing group and period fixed effects" (Gardner, 2021).[29] In Appendix Figures D1 and D2, we show robustness to using estimates from a two-way fixed effects (TWFE) model and the weighted group-time estimator developed by Sun and Abraham (2021).

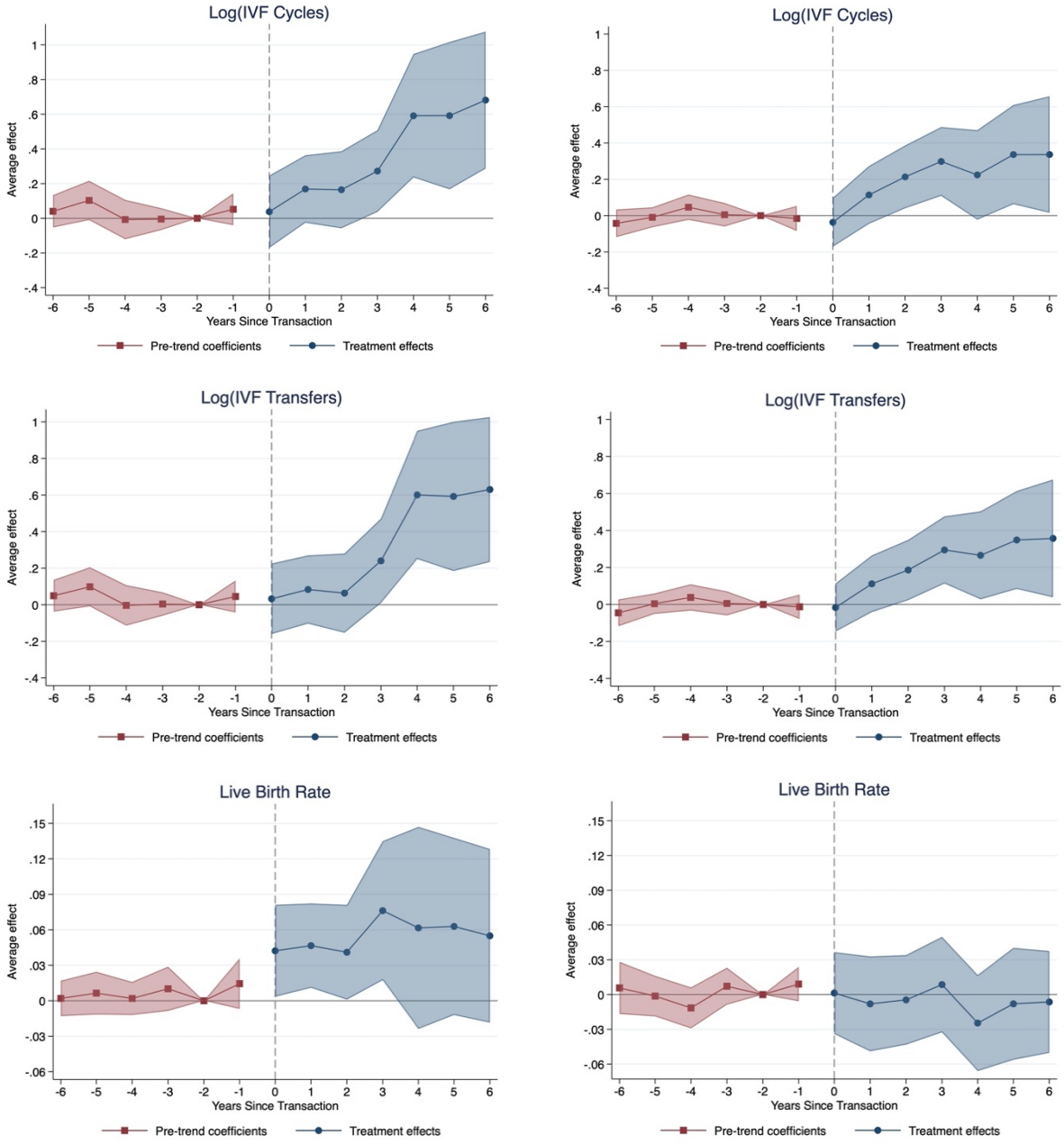
Figure 4 shows event study estimates for the 6 years before and after acquisition or affiliation for IVF cycles, IVF transfers, and the live birth rate (see Appendix Figure D3 for pooled results). We set the reference period to two years before the transaction to rule out possible anticipatory effects. For both ownership structures, there are no significant pre-trends before the transaction for IVF cycles and transfers: F-tests of joint significance show that the pre-transaction years are not statistically different from zero. After transaction, the cycles and transfers increase steadily before stabilizing after year 4. This pattern is consistent with the time it would take to expand operations. For example, volume-enhancing changes, such as investments in new office space, hiring and training new staff, and marketing to attract patients, will likely take time.

With respect to the live birth rate, the event study results are less precise. Before acquisition, pre-trends are relatively flat and not statistically different from zero. After acquisition, there is evidence that the live birth rate increases in the year of acquisition and remains above 4 percentage points. Pre-trends are flatter, and post-transaction estimates are more precisely estimated and show similar increases using TWFE and the Sun and Abraham (2021) method (Appendix Figure D1 and D2), lending further credibility to the increase in the live birth rate. These event study results may be suggestive of both immediate and longer-term effects of knowledge sharing in chains for acquired clinics. The immediate change in the live birth rate could result from accessing chain-wide protocols, while the additional increases in the live birth rate observed after year two could result from continued efforts to standardize care and learning from peers. For affiliated clinics, the live birth rate remains close to zero before and after affiliation.

[29] We use Gardner (2021) as our primary event study specification because of the simple, intuitive steps and assumptions, minimal programming, and flexibility when implementing DD with an unbalanced panel, multiple treatment groups and interacted fixed effects (Cunningham 2021). For the event studies, we use the *did2s* package developed by Butts (2021) and *eventstudyinteract* package developed by Sun (2021).

Figure 4. Event Study Results by Ownership Structure

(a) Acquisition (b) Affiliation



Notes: This figure shows the β_1 and β_2 estimates of Equation 2 interacted with indicators for the year relative to the transaction year. The reference period is two years before the transaction. Bands indicate 95% confidence intervals constructed from clinic-level clustered standard errors. For acquisitions and affiliations, respectively, the p-value from an F-test of joint significance are as follows: 0.281 and 0.696 for log(cycles), 0.275 and 0.757 for log(transfers) and 0.680 and 0.469 for the live birth rate.

4.3 The Role of Patient Selection

One identification concern is whether patient characteristics changed after a clinic joined a fertility chain in ways that would influence IVF success rates. For example, if observable patient characteristics changed after acquisition or affiliation, this suggests

patient selection on unobservables could bias the estimates. This selection could result from patients of higher or lower risk selecting certain clinics or from clinics potentially "cherry-picking" patients that would have more successful IVF outcomes, such as younger patients. Below we provide evidence that changes in patient characteristics do not appear to drive changes in the live birth rate.

First, the homogenous patient population mitigates concerns about patient differences across clinics. Patients that receive IVF treatment are predominantly white, privately insured, high-income, and highly educated (Chandra, Copen, and Stephen 2014; Galic et al. 2021). Furthermore, the single largest predictor of IVF success is a patient's age (SART 2021b). Predictive models based on pre-treatment patient characteristics find that patient age explains 85% of the total variation in the live birth rate and that patient infertility diagnosis, race/ethnicity, and body mass index are not strongly predictive of IVF success (Xu et al. 2022).

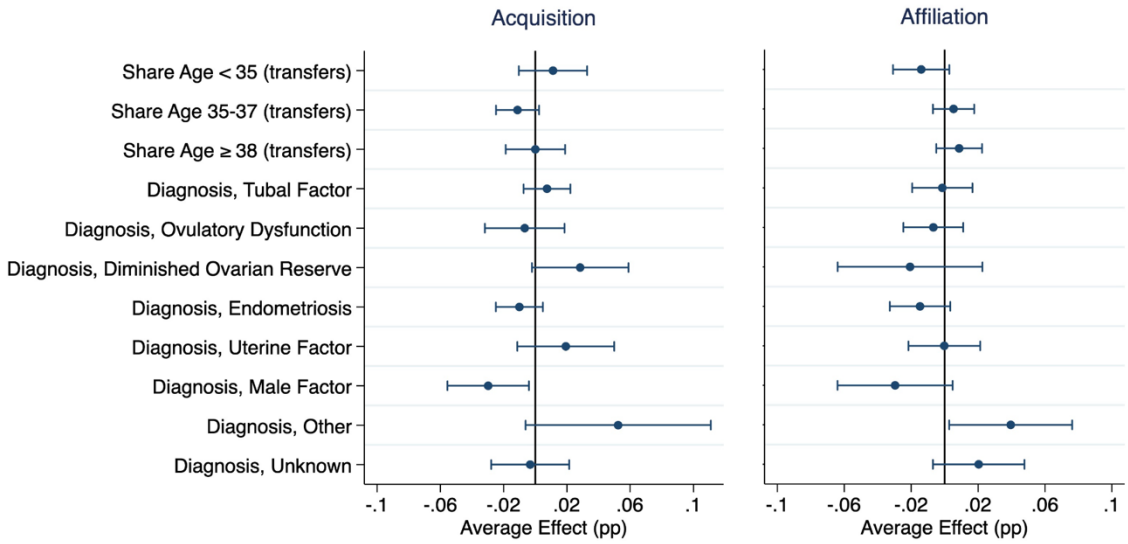
Second, we do not find evidence that clinics systematically treat patients that could be more or less likely to experience IVF success. Figure 5 presents changes in the share of patients in different age groups and patient infertility diagnosis as the outcome variable of Equation 2. For acquired clinics, there is a small reduction in the share of patients aged 35-37 but no change in patients under 35 or 38 and over, which suggests no clear pattern of patient selection based on age. For affiliated clinics, there is a small reduction in the share of patients under 35. However, in Section 5, we show that 1) increases in clinic volume are similar across all age categories in both acquired and affiliated clinics, 2) acquired clinics increase the live birth rate across all age categories, especially among patients over 38, and 3) affiliated clinics do not increase the live birth rate, even among patients under 35 (Figure 6 and Appendix Table G1). These results further minimize concerns that outcomes are driven by selection on patient age.

Patient diagnosis patterns are largely similar in acquired and affiliated clinics, and the changes are unlikely to influence IVF success. For example, even though there are significantly lower rates of patients with male factor infertility, there is limited evidence that male factor infertility impacts IVF outcomes (Shamonki et al. 2004; Vaegter et al. 2017). One study found that among the diagnosis categories, a tubal factor diagnosis is associated with the lowest live birth rate, and ovulatory dysfunction is associated with the highest (Vaegter et al. 2017). Figure 5 shows limited evidence of post-transaction changes for these diagnosis categories.

Third, we find quantitatively similar effects to our primary estimates when including patient diagnoses in Figure 5 as controls, confirming that patient infertility diagnoses have minimal influence on live birth rates (Appendix Table F1 Panel A). We also find quantitatively similar results when including controls for market-level maternal characteristics for patients who delivered a baby and reported using infertility treatment (Appendix Table F1 Panel B). These characteristics include a mother's race, mother's level of education, mother's insurance status, and maternal health factors. Lastly, we

find qualitatively similar increases in the live birth rate when weighting by yearly clinic volume to account for differences in clinic size (Appendix Table F1 Panel C).

Figure 5. Effect of Corporate Ownership on Clinic-Level Patient Characteristics



Notes: This figure displays β_1 and β_2 estimates of Equation 2 using patient characteristics as the outcome variables. Bars are 95% confidence bands. Standard errors are clustered at the clinic level.

Overall, these results suggest that changes in patient characteristics do not drive changes in IVF success across clinic types. A limitation of this study is that we can only observe IVF success rates per transfer rather than per patient. This means that the same patient may undergo multiple transfers during the year, and we cannot identify whether the success occurs on the first transfer or subsequent transfer.[30] Still, this is a commonly used success rate in IVF research and provides the most granular level to estimate success (Awadalla et al. 2021; Cozzolino et al. 2022; Mizrachi and McQueen 2022). Specifically, the occurrence of a transfer requires the creation of a viable embryo and, therefore, precludes patients with failed cycles. This allows for a more "apples to apples" comparison of patients across clinics.

4.4 The Role of Clinic Selection

Selection is an inherent feature of this setting: Fertility chains select the clinics they want in their chain, and clinics select the chain they want to join. Since being part of a fertility chain is not randomly assigned, we cannot unambiguously conclude that corporate ownership causes changes in clinic volume and IVF success. For example, chains may acquire clinics that they believe will achieve the best outcomes in the future.

[30] In cross-sectional analysis using new variables reported in 2017 and 2018, we can observe success rates per intended egg retrieval for new patients with no prior ART treatment between their first retrieval and their cumulative retrievals. As seen in Appendix Table F6, the cumulative rates are slightly larger than the first retrieval rates, but both are both positive and not statistically different from each other.

Below we provide additional discussion and analysis that helps mitigate but does not eliminate the role of clinic selection in explaining our results.

First, the primary specification helps account for potential differences between the treatment and control groups by including clinic fixed effects, which adjust for time-invariant clinic characteristics such as location and reputation. Clinic fixed effects also account for whether a clinic was part of an academic medical center. Including *state* × *year* fixed effects account for time-varying factors common to all clinics in a state. For example, these fixed effects would account for increases in demand for IVF if changes in state insurance coverage laws increase the affordability of care. Second, as seen in Figure 4, the event studies reveal clear changes in the outcomes before and after the transaction year. There are also no observable pre-trends in the years before the transaction. This suggests that the selection mechanism is unlikely to fully explain the changes observed after an acquisition or affiliation.

Third, a standard approach to address the endogeneity due to selection is to match treated units with similar characteristics in the pre-transaction period to untreated units. In this setting, clinic volume is the most salient difference between treatment and control clinics: on average, acquired and affiliated clinics appear to perform more IVF cycles and transfers than independent clinics before transaction (Table 1). While differences are slightly mitigated by logging the volume outcomes, higher volume clinics may still have different capabilities that more readily manifest in increased volume and IVF success rates in the future.

In Appendix Table F2, we show that the effects of the DD specification in Equation 2 are quantitatively similar using a matched control group. Specifically, we use 1-1 coarsened exact matching on a clinic's IVF cycles in the year before the transaction (see Appendix Table F3 for summary statistics on the matched sample). We also show robustness to matching on clinic cycles, live birth rate and the share of patients under 35. However, matching on outcome variables within a DD framework is susceptible to regression to the mean bias (Daw and Hatfield 2018). As an alternative strategy, we limit the control group to independent clinics that perform at least 150 cycles a year. This restriction increases the average number of IVF cycles of independent clinics from 284 to 426 cycles, which is closer to the pre-transaction volume of acquired and affiliated clinics. The results in Appendix Table F4 Panel A show qualitatively similar results for affiliated clinics and quantitatively similar results for acquired clinics. Together, the matched sample and sample restricted to high-volume clinics suggest that effects are not driven by differential selection of clinics based on size.

Lastly, we conduct robustness checks for the years a clinic was in the sample. Clinics open and close during the sample period and so may not be present in the data for all years between 2004 and 2018. For both corporate clinics and independent clinics, it could take several years for their fertility program to stabilize as they build their reputation and expertise after opening. We find that effects are quantitatively similar when limiting the sample to clinics present in all 15 years of data (Appendix Table F4

Panel B). The CDC ART data also includes a flag for whether a clinic self-reported restructuring, defined as a change in at least two of the three key staff positions (practice director, medical director, or laboratory director), or if the clinic was on the verge of closing. Independent clinics may experience a restructuring unrelated to joining a chain because of retirement, expansion, general changes in leadership, or impending closure. To ensure these changes are not inadvertently leading to a pseudo-treatment effect, we show that results are quantitatively similar when excluding independent clinics that are ever restructured or closed (Appendix Table F4 Panel C).

4.5 Additional Robustness

Unique features of the data also warrant additional robustness checks. In our preferred specification, we assume that once a clinic is treated, it remains treated. However, in some cases, a fertility chain experiences a second acquisition event (i.e., a larger chain acquires a smaller chain). Our results are robust to controlling for the second acquisition event (Appendix Table F5 Panel A). Another challenge included a change to data reporting by the CDC in 2018. While data cleaning efforts were made to homogenize data across years, we show results are robust to dropping the year 2018 (Appendix Table F5 Panel B). Lastly, there were two cases where it was less clear whether the clinic was an acquisition or an affiliation. There were also two chains that did not receive outside funding until 2018 and, therefore, may not resemble the other chains in the sample. We show that results are robust to excluding all these cases (Appendix Table F5 Panel C).

5. MECHANISMS

The previous analyses find that corporate ownership significantly increases clinic volume and IVF success rates and provide evidence that patient and clinic selection do not drive results. In this section, we provide suggestive evidence that the transfer of resources and knowledge following corporate ownership most likely explain the changes in clinic volume and IVF success rates. Resource transfers include any transfer of financial resources (i.e., capital) or managerial capabilities (i.e., marketing) from the corporate parent to the target clinic. Knowledge transfers include the sharing of new or superior clinical information. Corporate owners can transfer knowledge through top-down clinical directives (i.e., protocols, monitoring, and mandatory trainings) and by facilitating knowledge sharing among clinics through the creation of research consortiums and complex case review meetings. While resource and knowledge transfers can work in tandem to improve clinic volume and IVF success rates, resource transfers likely have a greater impact on clinic volume, and knowledge transfers likely have a greater impact on IVF success rates.

The following analyses are collectively intended to show patterns consistent with resource and knowledge transfers leading to improvement in outcomes. First, we conduct text analysis of press releases to supplement the quotes from other marketing materials

provided in Section 2.4. This text analysis confirms that fertility chains' stated goals and strategies are to help clinics grow and improve quality. Second, we discuss and conduct analyses most supportive of knowledge transfers: 1) We argue that increases in live birth rates among acquired clinics, but not affiliated clinics, are consistent with acquisitions better facilitating knowledge transfers because of greater incentive alignment and corporate control, 2) We show that clinics change processes and procedures in ways that enhance quality, consistent with new knowledge leading to improvements in IVF outcomes, and 3) We find that the lowest performing clinics pre-transaction experience the largest performance improvements, as do clinics acquired by higher-quality chains, consistent with clinics learning from other clinics in their chain. Third, we discuss and conduct analyses most supportive of resource transfers: 1) We find that fertility clinics lead to market expansion rather than business stealing from independent clinics, suggesting fertility chains provide resources needed to expand clinic operations and attract new patients, and 2) We show that PE investment into fertility chains largely drives increases in clinic volume, consistent with PE firms easing financial constraints and facilitating clinic growth.

5.1 Press Releases

There are often press releases associated with the announcement of a clinic acquisition or affiliation with a fertility chain. These press releases typically contain the stated goals and strategies of the chain and may shed light on how outcomes would change (Fracassi, Previtero, and Sheen 2022). While announcements are used as marketing tools that present transactions favorably, collectively, they suggest fertility chains emphasize clinic growth.

Table 3. Press Release Text Analysis

Concept	Identification	Number of Deals	Percentage
Expansion/Growth	(mentioning "growth" or "expand")	27	75%
Knowledge	(mentioning any of the below)	29	81%
	“knowledge”	7	19%
	"research"	20	56%
	"standard"	14	39%
	"systems"	3	8%
	"protocols"	5	14%
	“process”	4	11%
Resources	(mentioning any of the below)	29	81%
	"resources"	14	39%
	"financial"	19	53%
	"capital"	8	22%
	"management"	11	31%
	"marketing"	10	28%
	"technology"	17	47%

We find informative press releases on 36 transactions (Table 3). None of the texts mention layoffs or restructuring. However, 75% mention growth opportunities or expansion. To achieve growth, the press releases mention adding resources (81%) such as managerial capabilities and capital. Most press releases (81%) also emphasize standardization and knowledge sharing. For example, the chains make references to developing protocols, updating clinical processes, and conducting research. Overall, the stated strategies are consistent with resource and knowledge transfers that could help clinics increase volume and improve IVF success rates.

5.2 Ownership Structure

Differences in ownership structure can shed light on a corporate parent's ability to improve firm performance (Grossman and Hart 1986). Specifically, greater ownership typically confers greater control over the operations of the target firm. For example, Bernstein and Sheen (2016) study the effect of PE investment on restaurants and find effects are stronger among directly owned restaurants compared to franchised restaurants. Demirer and Karaduman (2022) use minority acquisitions as a placebo test under the assumption that firm efficiency would only change in majority but not minority acquisitions in their study of power plants.

In our setting, marketing materials and press releases suggest that both acquired and affiliated clinics receive access to resources and knowledge in a chain. However, in conversations with corporate investors, we learned that they prefer to acquire clinics because of their ability to better control operations. Affiliations typically occur because clinics' owners do not want to relinquish autonomy but still want access to financial resources and management services. Since the clinic owners retain greater decision-making authority and the transactions are more focused on resource transfers, we may expect that affiliations would only impact clinic volume but not IVF success rates.

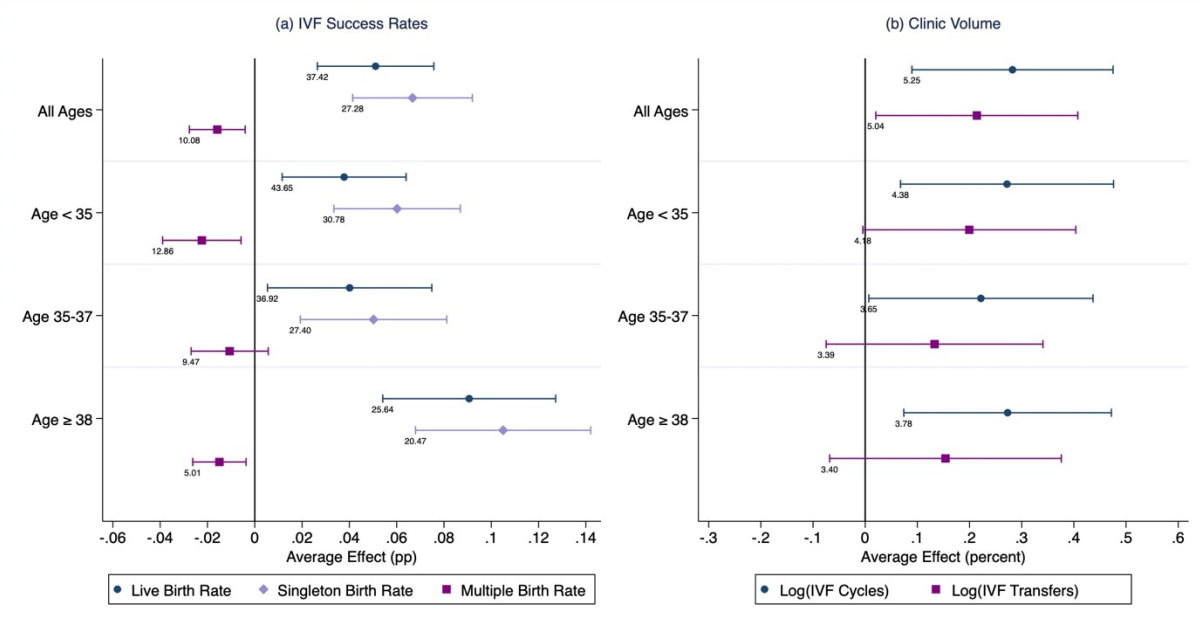
Furthermore, the interests of affiliated clinics may not be congruent with those of the chain because they are seeking to maximize their own profits rather than that of the organization. As seen in Table 1, affiliated clinics may believe their IVF success rates are already superior and have less incentive to invest in quality improvement efforts. Affiliated clinics may also seek to benefit from the chain's reputation, which may further reduce incentives to share knowledge or learn from other clinics. Overall, differences in control and interests of affiliated clinics compared to acquired clinics provide a placebo test for the role of knowledge transfer. Consistent with this argument, we find that both acquisitions and affiliations lead to increases in clinic volume, but only acquisitions lead to increases in the live birth rate (Table 2).

5.3 Procedure and Technology Changes and Quality Improvement

Changes in Singleton vs. Multiple Birth Rates. Improving IVF success rates is a major challenge for fertility clinics. The "gold standard" is to simultaneously decrease

multiple births and increase singleton births by enough to have a net positive effect on the live birth rate. Reducing multiple births is considered quality-enhancing because multiple births have a greater incidence of obstetric and neonatal complications (Kissin, Boulet, and Jamieson 2016). However, transferring multiple embryos (approximately 30% probability of a multiple birth) is associated with a greater probability of success than a single embryo transfer (less than 1% probability of a multiple birth). Since the live birth rate is the key metric clinics use to attract patients, this may create an incentive to transfer multiple embryos at once to increase IVF success rates.[31] Additionally, transferring multiple embryos allows more room for error in the embryos chosen for transfer, whereas a single embryo transfer requires more precision and expertise to identify the highest quality embryo to transfer (Reimundo et al. 2021).

Figure 6. The Effect of Corporate Ownership on Fertility Clinic Outcomes by Patient Age and Birth Type, Acquired Clinics



Notes: This figure displays β_1 estimates of Equation 2 by patient age category (i.e., only displays results for acquired clinics). The dependent variable mean based on the predicted mean for control clinics and treatment clinics before the transaction is displayed under each 95% confidence bar. Standard errors are clustered at the clinic level. See Appendix Table G1 for the full regression results.

As a result, if the chain were sharing knowledge to improve IVF processes and quality, we would expect increases in the live birth rate to be driven by increases in singleton births large enough to compensate for reductions in multiple births. In Figure 6(a), we graphically present results from Equation 1 by the overall live birth rate, multiple birth rate, and singleton birth rate for acquired clinics. For overall IVF success

[31] There is much debate on patient preferences for twins since patients perceive twins as avoiding future IVF attempts but may not comprehend the associated risks (Fiddelers et al. 2011; Shenoy et al. 2017; Mendoza et al. 2018). However, a metanalysis suggests patient education is an effective strategy in reducing the desire for twins and increasing the use of single embryo transfers (Sunderam et al. 2018).

rates, singleton births increase by 6.7 percentage points, and multiple births decrease by 1.6 percentage points. A similar pattern is observed within each patient age group.

The increase in the live birth rate driven by singleton births is consistent with new knowledge enabling greater success of single embryo transfers. For example, chains emphasize adopting new procedures that improve embryo selection, such as preimplantation genetic testing (PGT) and intracytoplasmic sperm injection (ICSI): "cutting edge technology enables embryologists and fertility specialists to assess the genetic and chromosomal makeup of an embryo prior to its transfer into a woman's uterus."[32] As shown in Table 4, we find strong evidence that acquired clinics increase the use of PGT but find no changes in ICSI use (potentially because ICSI was introduced in 1991 and experienced rapid adoption). One chain also describes implementing the use of day 5 blastocyst embryos because "this advanced IVF lab technique allows the embryo to mature as far as it can outside the human body, again allowing embryologists and physicians an enhanced ability to select the best single embryo for transfer." While we are unable to measure this outcome, we find that, per transfer, acquired clinics reduce the average number of embryos transferred to the uterus (Table 4). In contrast, no such improvements are seen in affiliated clinics.

Table 4. The Effect of Corporate Ownership on Procedure and Lab Changes

	(1)	(2)	(3)	(4)	(5)
	PGT Rate	**ICSI Rate**	**Avg. # of Embryos Transferred**	**Prob. Lab Name Change (Any)**	**Prob. Lab Name Change (Single)**
Post × Acquisition	0.070***	-0.004	-0.286***	0.443***	0.357***
	(0.027)	(0.026)	(0.053)	(0.078)	(0.103)
Post × Affiliation	0.016	0.000	0.075*	0.225***	0.169*
	(0.010)	(0.033)	(0.038)	(0.085)	(0.088)
Dep. Var. Mean	0.058	0.676	2.232	0.124	0.084
Clinic-Years	3863	4888	4890	5666	5210
R^2	0.698	0.766	0.845	0.745	0.700

Notes: This table shows the β_1 and β_2 estimates of Equation 2. The dependent variable mean captures the predicted mean for control clinics and treatment clinics before the transaction. The CDC ART data does not report PGT until 2007 and changes how the data on PGT, ICSI and the number of embryos transferred are collected in 2017 and 2018. Therefore, PGT rate uses data from 2007-2016, and ICSI rate and number of embryos transferred from 2004-2016. Column 4 includes all clinics and column 5 limits the sample to clinics that only had a single change in the lab name during the sample period. Standard errors are clustered at the clinic level. Significance levels: *p<0.1, **p< .05, ***p<0.01

Fertility chains can also provide resources to help modernize a clinic's laboratory and implement protocols to standardize laboratory processes. For example, one chain

[32] See Appendix Table B1 for quote sources in this section. Note that new procedures or techniques are often referred to as technological advancements by fertility specialists. Different technology (i.e., devices or platforms) can be used to conduct the procedures.

emphasizes that "continuous improvement in laboratory processes and patient care protocols have to lead to increased success rates." While we cannot directly measure whether a clinic changes or makes updates to their laboratory, the CDC ART data publishes the name of the laboratory used by each clinic each year. The name change may indicate a significant overhaul or signal a rebranding with no meaningful changes to the laboratory. In Table 4 Column 4, the outcome variable is zero for all clinics before a name change and 1 after the first time a clinic changes its laboratory name. In Table 4 Column 5, we limit the sample to clinics that only experience 1 name change, as this may best capture a real change. We find that post-transaction, the probability a laboratory changes for acquired clinics increases between 35.7 and 44.3 percentage points and between 16.9 and 22.5 percentage points for affiliated clinics (the estimates for acquired and affiliated clinics are statistically different from each other). These results provide evidence that fertility chains may update clinic laboratories and facilitate the use of new technology and techniques that enhance the quality of IVF.

Changes by Patient Age. As an additional strategy to explore the role of knowledge transfer, we consider whether IVF success rates change for patients of different complexity (Stan and Vermeulen 2013). The qualitative materials collected from chain websites suggest physicians within a chain regularly meet to discuss complex patients (Appendix Table B1). One of the most important predictors of IVF success is a patient's age, with patients of older age representing more complex cases. If accessing new or superior knowledge contributes to improved techniques and processes, we would expect the largest improvements for older patients, as they have the most to benefit.

Figure 6 provides empirical support for this argument, where patients aged 38 and older experience increases in the live birth rate at almost double the rates of patients under 35 or patients ages 35-37. The volume of IVF cycles and transfers also increases by the same amount for older patients as other age groups and increases in singleton births drive the increase in the live birth rate. One potential reason for these improvements is the increased use of PGT (Table 4): studies have found that usage particularly increases the success of single embryo transfers among older women (Maxwell and Grifo 2018; ACOG 2020).

5.4 Clinic Heterogeneity and Learning

Changes by Baseline Clinic Rates. The existence of knowledge transfer assumes physicians receive access to new or superior knowledge or generate knowledge through collaboration after joining a fertility chain. The event studies in Figure 4 suggest both immediate and longer-term effects of acquisitions on the live birth rate, though the wide confidence intervals may indicate differences in effect sizes across clinics. For example, clinics with high birth rates pre-transaction may be less likely to experience positive effects, and those with lower birth rates may have the most to benefit from accessing

the knowledge of the chain. Similarly, clinics that already had higher clinic volume may benefit less from resources meant to expand clinic operations.

In Table 5, we divide acquired and affiliated clinics into three terciles based on their pre-transaction IVF cycles, IVF transfers, and the live birth rate, and interact an indicator for each category with the post-transaction indicators in Equation 2. Specifically, all acquired clinics are divided into terciles, and separately, all affiliated clinics are divided into terciles based on their pre-transaction means for each outcome. This strategy allows a clinic to potentially be initially high performing on live birth rate but low performing on clinic volume. Table 5 provides evidence that all acquired clinics experience improvements, but that the largest increases in clinic volume and live birth rates occur among initially lower-performing clinics relative to those that were higher-performing. These results suggest joining a fertility chain creates a "rising tide lifts all boats" effect, in which all clinics improve, but especially lower-performing clinics.[33]

Table 5. The Effect of Corporate Ownership on Fertility Clinic Outcomes by Pre-Transaction Clinic Performance

	(1)	(2)	(3)
	Log(Cycles)	**Log(Transfers)**	**Live Birth Rate**
Post × Acquisition(Low=1)	0.366***	0.287*	0.059***
	(0.140)	(0.146)	(0.022)
Post × Acquisition(Med=1)	0.302	0.209	0.058**
	(0.205)	(0.212)	(0.024)
Post × Acquisition(High=1)	0.133	0.107	0.037***
	(0.105)	(0.099)	(0.009)
Post × Affiliation(Low=1)	0.521***	0.490**	-0.010
	(0.201)	(0.206)	(0.016)
Post × Affiliation(Med=1)	0.232*	0.220*	0.044
	(0.131)	(0.127)	(0.029)
Post × Affiliation(High=1)	-0.035	-0.055	-0.034
	(0.069)	(0.074)	(0.026)
Dep. Var. Mean	5.251	5.034	0.374
Clinic-Years	5666	5666	5666
R^2	0.900	0.899	0.626

Notes: This table displays β_1 and β_2 estimates of Equation 2 with clinics divided into terciles based on their pre-transaction averages for each outcome variable (these categories are mutually exclusive). For example, *Acquisition(Low=1)* is an indicator equal to 1 if an acquired clinic was in the bottom tercile of acquired clinics based on its pre-transaction mean. The dependent variable mean captures the predicted mean for control clinics and treatment clinics before the transaction. Standard errors are clustered at the clinic level. Significance levels: *p<0.1, **p< .05, ***p<0.01

[33] To further support this claim, in Appendix Table G2 we use the difference between an acquired clinic's own live birth rate and the average live birth rate of the chain as the outcome variable of Equation 2. The results confirm that clinics that are below *and* above their chain's average before acquisition experience significant increases in their relative live birth rates after acquisition, but that initially below-average clinics experience the largest increases (8.3 percentage points vs. 3.9 percentage points).

To better understand the fertility chain's role in facilitating improvements in the live birth rate, we conduct additional subsample analyses. If clinics are indeed learning from their chain, we would expect clinics acquired by high-performing chains to experience larger increases in the live birth rate than those acquired by lower performing chains. To test this, we create an indicator for whether a *fertility chain* is relatively high performing (above median), or low performing (below median) based on the average live birth rate of the clinics in the chain before the first chain transaction occurs during our sample period (there are not enough chains to create terciles). We then interact these high and low-performing chain indicators with the post-transaction indicators in Equation 2. The estimates on these interaction terms are statistically significant and reveal that live birth rates increase by 7.3 percentage points in clinics acquired by high-performing chains, and by 2.8 percentage points in clinics acquired by low-performing chains (Appendix Table G3). These findings suggest that fertility chains with pre-existing superior knowledge can facilitate larger increases in live birth rates.

Volume-Outcome Relationship. Rather than improving from accessing new knowledge, physicians may improve their outcomes by performing more IVF cycles. A recent study by Wilkinson et al. (2022) did not find a significant association between a clinic's volume and its live birth rates. Similarly, a review by Mizrachi and McQueen (2022) finds no evidence of differences in IVF success rates based on physician experience: Even among fellows, outcomes were stable throughout their training. The authors conclude that because embryo transfer is "performed by a single operator on their own, and thus, after initial training, there is limited opportunity for physicians to compare their technique to other colleagues and improve" (Mizrachi and McQueen 2022, p. 816). Therefore, rather than within-physician learning from increased volume, physicians may be more likely to improve from knowledge sharing within the chain.[34]

The event study results provide insights into the volume-outcome relationship (Figure 4). If improvements among acquired clinics only happened via learning-by-doing from increased volume, we would expect the improvement in the live birth rate to follow a similar trajectory as the increase in clinic volume. Instead, in the first couple of years post-acquisition, there are immediate increases in the live birth rate without commensurate increases in volume. Additionally, affiliated clinics see large increases in volume but no changes to the live birth rate. We also re-estimate the terciles in Table 4 Column 1 based on pre-transaction IVF cycles but use live birth rate as the outcome (Appendix Table G4). The estimates of this regression show that acquired clinics in the top tercile of volume (which saw no significant increases in volume) still significantly increase the live birth rate by 4.2 percentage points. Together, these results suggest the volume-outcome relationship does not appear to drive the increase in the live birth rate.

[34] Volume could still be an important mechanism to reinforce a newly learned or adopted procedure. We aim to rule out that volume alone drives changes in the live birth rate.

5.5 Market Expansion

The marketing materials of fertility chains place a large emphasis on growth. For example, chains advertise providing clinics with financial resources to fund add-on locations and hire new clinical and administrative staff. Additionally, chains advertise providing clinics with managerial capabilities such as marketing services and patient engagement programs to attract and retain patients throughout their IVF journey. Clinic growth could help increase access to IVF given the unmet demand for fertility services driven by the scarcity of clinics and cost of IVF (Chambers et al. 2009; Greil et al. 2016).

However, rather than expand the market and increase access to care, corporate clinics may instead be engaging in business stealing. That is, capturing market share of independent clinics by treating patients that would have been treated by independent clinics in the absence of an acquisition or affiliation. To study market expansion vs. business stealing, we implement the following instrumental variables approach:[35]

Eq. 3: $$Total\_Corporate_{jy} = \gamma Corporate\_Clinics_{jy} + \zeta \boldsymbol{X}_{jy} + \theta_j + \theta_y + \nu_{jy}$$

Eq. 4: $$M_{jy} = \delta Total\_Corporate_{jy} + \phi \boldsymbol{X}_{jy} + \theta_j + \theta_y + \epsilon_{jy}$$

The first stage in Equation 3 instruments for the total number of IVF cycles ($Total\_Corporate_{jy}$) performed by corporate clinics in market j in year y using the number of corporate clinics ($Corporate\_Clinics_{jy}$) in market j in year y. We control for market fixed effects (θ_m), year fixed effects (θ_y), and market-level controls ($\boldsymbol{X}_{jy}$) including the log of median household income and log of total population aged 20-49.

To test whether corporate ownership leads to business stealing, we estimate the second stage (Equation 4) using the total number of IVF cycles performed by independent clinics in a market as outcome M_{jy}. If an increase in IVF cycles by corporate clinics is the result of business stealing, then we would expect $\delta = -1$. To test whether corporate ownership leads to market expansion, we instead use the total number of IVF cycles performed by *all* clinics in a market as outcome M_{jy}. If fertility chain growth is market expanding, then we would expect $\delta = 1$. In other words, for every 1 IVF cycle performed by a corporate clinic, there is 1 additional IVF cycle in that market.

In Table 6, the market is defined as a clinic's CBSA, where Columns 1 and 2 show the δ estimates from Equation 4 using the total number of IVF cycles.[36] In Columns 3 and 4, we present estimates using the total number of live births (not the rate). By construction, the instrument is strongly predictive of total IVF cycles and live births by corporate clinics because as the number of corporate clinics in a market increases, so will the number of corporate cycles and live births. The number of corporate clinics is

[35] We implement this approach to account for multiple transactions by different clinic chains in the same market and because it provides an intuitive test of market expansion.

[36] We find quantitatively similar results when defining the market as commuting zones (developed by the Economic Research Service in 2000), which do not depend on population size (Appendix Table G5).

assumed to only impact total *market* cycles and live births through the increase in total *corporate* cycles and live births.

As seen in Table 6, we find no support for business stealing and strong evidence in support of market expansion. We observe no reduction in cycles for independent clinics (Column 1), and for every additional cycle performed by a corporate clinic, there is one additional cycle at the market level (Column 2). These results are consistent with chains providing resources needed to ease clinic capacity constraints and expand the set of patients utilizing IVF. Given the large unmet demand for fertility services, corporate clinics likely increase access to IVF.[37] There are also similar patterns observed for the number of live births (Columns 3 and 4), providing evidence that the entry of corporate clinics does not significantly impact the IVF outcomes of independent clinics.

Table 6. Market Expansion Analysis, IV Estimates

	(1)	(2)	(3)	(4)
	Total Market Cycles		**Total Market Live Births**	
	Independent Clinics	All Clinics	Independent Clinics	All Clinics
Total Corporate Cycles	-0.002	0.998***		
	(0.148)	(0.148)		
Total Corporate Live Births			-0.201	0.799***
			(0.131)	(0.131)
First Stage: F-Stat	80.607	80.607	64.675	64.675
Market-Years	1930	1930	1930	1930

Notes: This table displays the δ estimates of Equation 4. The market is defined as the CBSA of the clinic. *Total Corporate Cycles* and *Total Corporate Live Births* represent the total number of IVF cycles and total number of live births performed by corporate clinics each year in a CBSA, instrumented using the number of corporate clinics each year in a CBSA. The first stage F-stat shows the Kleibergen-Paap Wald rk F statistics. We cannot reject that the estimates in Columns 2 and 4 are statistically different from 1 (the p-value from F-tests are 0.989 and 0.134, respectively). The sample includes all clinics (including clinics always in a chain and newly opened by a chain) in a CBSA that ever had an independent clinic. Standard errors are clustered at the market level. Significance levels: *p<0.1, **p< .05, ***p<0.01

5.6 The Role of Private Equity

Private equity (PE) firms can enable significant growth among acquired firms by alleviating financial constraints relative to other types of ownership (Eaton, Howell, and Yannelis 2020; Fracassi, Previtero, and Sheen 2022). Therefore, we may expect that the

[37] An increase in cycles may also be the result of supplier-induced demand: physicians pressure patients into IVF instead of alternative treatments or into conducting multiple cycles. However, this doesn't necessarily make patients worse-off. More cycles per patient could be the direct effect of increasing the use of single embryo transfers to reduce multiple births, which can often require multiple rounds of IVF to achieve a live birth. Similarly, alternative treatments such as intrauterine insemination have a higher incidence of multiple birth and often take longer to achieve pregnancy than IVF.

financial resources provided by PE firms would have more salient effects on clinic growth and volume (Braun et al. 2021; Singh et al. 2022). As described in Section 2, 5 fertility chains were created following PE investment, 4 chains already existed and later received PE investment, and 2 chains were acquired by larger international healthcare chains without PE investment. The variation in ownership and timing allows us to decompose the effect of PE investment on outcomes.

Appendix Table G6 Panel A shows the results of Equation 2 decomposing the effect between post-transaction years when a clinic was part of a chain with and without PE funding. Specifically, we create a post-transaction indicator equal to 1 when a clinic is part of a chain without PE funding and 0 when the chain has PE funding (*Post_NoPE*), and another post-transaction indicator that is the inverse (*Post_YesPE*). We find large and significant increases in the live birth rate *both* when a clinic is acquired by a chain without PE (5.1 percentage points) and with PE funding (5.2 percentage points). In contrast, almost all the volume effect for acquired clinics occurs because of PE funding (10.6% increase in cycles without PE and 31.9% increase with PE). This result is consistent with clinics accessing knowledge upon first joining the chain but not experiencing significant growth until after PE investment. In other words, PE investment may help facilitate clinic growth but appears to play less of a role in increasing the live birth rate.[38]

6. CONCLUSION

This paper studies how corporate ownership impacts firm performance in the fertility industry. By 2018, over 20% of fertility clinics (performing over 40% of IVF cycles) were part of a fertility chain backed by private equity or a large global corporation. Our results show that affiliated and acquired clinics increase the volume of IVF cycles and transfers by over 22%, whereas only acquired clinics significantly increase the live birth rate. The 5.1 percentage points increase observed in acquired clinics represents a statistically and economically meaningful increase of 13.6% in the live birth rate. We provide compelling qualitative and quantitative evidence that resource and knowledge transfers driven by corporate ownership are the most likely explanation for the improvement in clinic performance.

Acquired clinics increase the quality of care by simultaneously reducing multiple births and increasing singleton births and achieve the greatest increase in live births among older patients. These improvements coincide with decreases in the number of embryos transferred and a significant increase in preimplantation genetic testing, which has been found to improve IVF success rates among older patients. These results are consistent with the marketing materials and press releases of fertility chains that argue

[38] In Appendix Table G6 Panel B, we use the year the network received PE funding as the year of treatment. These results confirm the role of PE: after investment, IVF cycles increase by 18.9 percent, IVF transfers increase by 14.2 percent, and the live birth rate increases by 2.0 percentage points.

that by facilitating knowledge sharing, they can improve IVF success rates. We also find that corporate clinics increase volume mainly through market expansion rather than business stealing and that PE investment into fertility chains largely drives increases in clinic volume. These results are consistent with access to new resources facilitating clinic growth. Lastly, we do not find evidence that results are driven by changes in patient characteristics that could influence IVF success or by differences in the types of clinics selected for acquisition or affiliation.

By studying fertility clinics, this paper provides a case study of corporate ownership in healthcare with relatively minimal market frictions and information asymmetries common to other healthcare settings. For these reasons, positive outcomes may be possible in other healthcare settings with similar characteristics, such as dermatology and cosmetic surgery and ophthalmology, and some urgent care or retail clinics. More broadly, our findings suggest that increased price and quality transparency may be an effective tool to better align shareholders with stakeholder interests, and sheds light on the potential impact of the Transparency in Coverage Act that went into effect in July 2022.

However, the DOJ and FTC have recently announced a shift in their antitrust enforcement toward PE-backed healthcare organizations (Cumming 2022). A key concern, as stated by Deputy Assistant Attorney General Andrew Foreman, is that rather than "function as a maverick or a disruptor in health care markets," investors will "cause the target company to focus solely on short-term financial gain and not on advancing innovation or quality" (Foreman 2022). We find evidence of the former, whereby creating fertility clinic chains, corporate investors may help facilitate knowledge sharing among physicians. Previously independent privately-owned clinics had limited means and incentives to collaborate and learn from each other. Joining a fertility chain can help reduce barriers to collaboration, and a centralized management system can allow for distribution of information and standardization of care. We also find limited evidence that fertility clinic acquisitions or affiliations lead to changes in market concentration. Still, given the recency of the phenomenon, there may eventually be anti-competitive effects as clinic acquisitions continue.

Ultimately, the findings of this paper are societally important and shed light on the future of the fertility industry. Current changes in abortion laws could affect patient access to IVF (Halleman et al. 2022; Jokisch Polo 2022). Given the potential income loss, abortion bans could ultimately lead corporate owners to close or move fertility clinics, laboratories, and egg-freezing facilities to less restrictive states, further exacerbating inequalities in access to fertility care (Pringle 2022). Abortion bans also raise concerns over maternal and infant health because women pregnant via IVF are more likely to develop life-threatening pregnancy complications, and babies are more likely to suffer congenital disabilities (Hansen et al. 2013; Reddy et al. 2007).

Projected demand for IVF has also generated increased scrutiny of fertility clinic performance (Faddy, Gosden, and Gosden 2018; Walsh 2021). While the live birth rate

has increased considerably in the past two decades, most patients still have less than a 40% chance of delivering a baby, and rates vary considerably across clinics. New technologies are currently being developed that utilize artificial intelligence to standardize care and improve success rates, raising questions about which clinics and patients will receive access to these technologies (Kesari 2022). More broadly, the striking improvement in IVF success rates within fertility chains highlights the tension between clinical knowledge as a competitive advantage versus a public good that could collectively improve fertility outcomes. Future research must consider implications for equity in access and outcomes.

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APPENDIX

Appendix A. Data Collection and Sample

A.1 CDC ART Data

We downloaded both excel files of the data and PDF files from the CDC directly: https://www.cdc.gov/art/artdata/index.html. The PDF files contain additional information about whether a clinic restructured or failed to report data, as well as address information and laboratory information. For almost each year of data there are changes in how the CDC reports the data. For this reason, not all variables can be consistently identified over the sample period and it is necessary to include year fixed effects to account for potential measurement error from year-to-year changes in reporting. Below we describe two additional data issues and our data cleaning approach.

1) Changes in 2018: In the year 2018, when fewer than 4 observations were reported for an outcome, the number was obscured by a "star" value. We replace "star" equal to 1 in one dataset (the main dataset used in the analysis) and equal to 4 in another dataset (as a robustness check) and find results do not differ between these two extremes. Additionally, since our analytic sample excludes clinics with fewer than 20 cycles a year over the sample, this removes most of these missing data cases regardless of the 1 or 4 replacement strategy. Lastly, we show results are quantitatively similar when excluding the year 2018 altogether (Appendix Table F4 Panel A).
2) Non-reporting clinic: Every year, the CDC ART data lists clinics that conducted IVF cycles but failed to report their outcomes (approximately 8% of fertility clinics each year). In most cases, this is because a clinic is about to close or is restructuring, which are both indicators provided in the CDC data (these are not mutually exclusive – clinics can close or restructure and still report data to the CDC in the year they are closing or restructuring). In our empirical analyses we exclude all clinics with less than 3 years of data and with gaps of 2 years or more in reporting to the CDC, since these clinics are likely undergoing substantial changes to their IVF program (effect sizes are slightly larger but not statistically different if we include these clinics). This restriction removes most of the clinics with non-reporting years. However, in robustness checks, we also show that results are robust to excluding clinics that ever closed or ever restructured, as well as to limiting the sample to clinics present in all 15 years of data (Appendix Table F3). Altogether, the consistency of results whether or not we exclude clinics based on reporting criteria mitigates concerns that reporting bias is affecting our results.

The main variables in the data are constructed as follows:

- IVF Cycles. Total number of cycles using fresh or frozen eggs from both donors or non-donors.
    - From 2004-2016 fresh and frozen cycles were reported separately, so we added them together to create a single consistent variable. In 2017 and 2018, the total cycles variable included fertility preservation cycles but also reported which

percent of cycles were for fertility preservation, so we removed those from the total cycles estimate.
    - In any analysis reported or weighted by age category, we only include non-donor cycles, because donor cycles are not reported by age.
- IVF Transfers. Total number of transfers using fresh or frozen eggs from both donors or non-donors.
    - From 2004-2016 fresh and frozen transfers were reported separately, so we added them together to create a single consistent variable.
    - In any analysis reported or weighted by age, we only include non-donor transfers, because donor transfers are not reported by age.
- Live Birth Rate. This is the percentage of fresh or frozen embryo transfers from non-donor eggs that resulted in a live birth.
    - In most analysis the live birth rate is a weighted average of four patient age categories: under 35, 35-37, 38-40 and 41-42. In analysis by age group, we take the weighted average of patients aged 38-40 and 41-42 to have a more proportional number of patients in each group (under 35, 35-37 and over 38).

A.2 Clinic Classification

Clinic Transaction Year. We define the transaction year as the year the clinic became part of a fertility clinic chain and is therefore, no longer an independent clinic. Some chains do not exist until they are created by private equity firms during the sample period, in which case, the transaction year is the first year the chain was formed.

Corporate Ownership. Clinics can become part of a chain in the following three ways:
1) Acquisition: An acquisition refers to an event where assets of the clinic (office, lab, or both) are acquired, owned, and managed by the parent company of the fertility chain. To comply with corporate practice of medicine laws, which prohibit non-physician-owned business entities from practicing medicine, these acquisitions typically follow what is referred to as a "friendly PC" model.[1] In this model, the chain has a subsidiary management company that acquires the seller's assets, and the selling physician serves as the owner of a separate professional corporation (PC). A long-term management service agreement is then signed between the friendly PC and management company for the company to manage the operations of the clinic. These agreements may include restrictive covenants and non-compete agreements (for example, physicians often enter 5-year employment agreements, and are restricted from opening a geographically proximate

[1] A more comprehensive legal description can be found here: https://www.chapman.com/publication-Health-Care-Management-Service-Organizations. CPOM has been heavily criticized for not being effective given the legal workarounds and lack of state enforcement. There have also been several high-profile lawsuits by physicians again PE-backed firms for "profound and pervasive direct and indirect control over the physicians' practice of medicine" (See https://www.lifesciencesperspectives.com/2022/01/26/california-physicians-allege-pe-backed-provider-violates-corporate-practice-law/). Note that CPOM would only apply to cases where a clinic is acquired by a publicly traded or PE-backed firm and not when a clinic is acquired by another clinic in a chain (before the chain received external investment). Clinic to clinic acquisitions are sometimes referred to as mergers in press releases, but usually the clinics remain physically separate entities that are part of the same chain.

competitor clinic), productivity thresholds, and capital commitments to fund clinic growth. The parent company, therefore, exerts control over clinic operations.
2) Affiliation: An affiliation refers to any interorganizational partnership, alliance, venture or collaboration between a clinic and a fertility chain where the clinic contracts with the chain for selected management services and capital or financing options. For example, a clinic may contract with a chain to receive access to marketing and patient engagement services. These affiliations may resemble outsourcing agreements in which the parent company has no ownership stake or may be structured as joint ventures in which the parent company has a partial ownership stake. While affiliations cover a diverse set of arrangements, the common feature is that clinic owners maintain greater control of clinic operations.
3) De novo: De novo growth refers to a new location built and opened as part of the chain and is therefore, owned and managed by the chain.

Data Collection Process. Below we explain how we determine which clinics are part of a corporate chain and their ownership structure.

Create list of fertility chains: We used documents published by consultants and legal firms outlining fertility clinic mergers and acquisitions (for example: www.dresnerpartners.com%2Face-files%2FFertility_June_2018.pdf) and used business intelligence databases such as Irvin Levin, SDC Platinum and Pitchbook to identify fertility chains that existed during the sample period (some chains no longer exist by name because they were acquired by other chains). In addition, since many times the name of a clinic includes the name of the chain, we were able to use CDC Fertility Clinic Success Reports (referred to as the CDC ART data) to identify chains. A recent publication by reproductive endocrinologists also helped confirm the identification of chains (Patrizio et al. 2022). Details on the specific chains are provided in Section A.3.

Identify clinics in fertility chains: We focused on one chain at a time to identify clinics in that chain. In addition to the business intelligence databases, we used archived and current versions of chain websites and clinic websites, EDGAR database for SEC filings for those that are part of a publicly traded company, searched for press releases using the name of the chain, and searched for whether the name of the clinic or the clinic's laboratory included the name of the fertility chain in the CDC ART data. We then manually searched for each clinic in the data to ensure we did not miss any clinics through the process above and ensure that none of our independent clinics were under corporate ownership. Specifically, we used www.google.com to search the name of the clinic in combination with any of the following terms: "management company", "private equity", "acquired", "acquisition", "merger", "partnership", "alliance" and "affiliation."

1) **Clinic transaction year.** The year of transaction was recorded as the year of the announcement date via a press release or PitchBook or date provided in a SEC filing. If this was not available, we used the date the clinic appeared on a chain website using the WayBack Machine. However, if the CDC reports showed a clinic change names in a year different than the sources above, we used the ART data year and noted this choice. The CDC reports signal a change through a change in the clinic's name to include the chain name and provides an indicator for whether a clinic restructured, which often coincides

with an acquisition or affiliation. In all cases, if there was a discrepancy, the changes occur in the CDC ART data in the year before the announced date in a press release, especially if the press release is from early in the year. For example, if a press release is from January 1, 2017, but the clinic changes name to a chain name in 2016, then we record the year of transaction as 2016.

2) **Clinic ownership structure.** In addition to collecting data on the transaction year, we also classify clinics into the ownership structures previously described.
    - **a)** When available, we use the description of the financial terms of the transaction as provided in the business intelligence databases or the SEC filings to determine whether a clinic was an acquisition or affiliation.
        - **i.** Note that contracts for acquisitions and affiliations are complex. Acquisitions are easier to identify because of legal filing requirements and recording in business intelligence databases. The exact type of affiliation, and particularly the terms of an affiliation, are not possible to determine unless the information has been explicitly reported. For this reason, affiliations include a variety of different arrangements from contracts for services to joint venture agreements. Our assumption is that a fertility chain exerts greater control over clinical operations in an acquisition than in an affiliation.
    - **b)** If this information was not available, we used press releases and information provided on chain websites in combination with state corporate filing data to deduce the nature of the transaction. For example, if a press release used the term "partnership" this could reflect either and acquisition or affiliation. We therefore search for the clinic's corporate filing records (using OpenCorporates.com or state corporate filing websites) to see if a clinic filed to become a subsidiary of a chain (likely signals an acquisition) or remained an independent legal entity (likely signals an affiliation). However, this is not always the case and depending on the state, corporate filing details may not be available.
        - i. From the chain websites we were also able to deduce their typical organizational model. For example, one chain's internal marketing materials emphasize that all their member clinics follow a joint venture model where physicians retain majority control. Another chain's website lists the clinics it owns separately from affiliate clinics. In combination with the steps above, we were able to make an informed decision of a clinic classification based on the chain's strategy.
        - ii. Additionally, if there is evidence of a merger – the clinic's operations or reporting were merged with another clinic in the chain – then we used this as evidence of an acquisition (i.e., these are not mergers of equal – one clinic acquires another clinic and then "merges" their data reporting). In the ART data, clinics that are merged typically stop reporting outcomes under their own name and start reporting under the clinic that acquired them. We record, combine and control for these mergers in the analysis.
    - **c)** Each clinic's classification is recorded/checked by 4 different individuals (two Research Assistants with experience working in private equity and finance, and

the two authors of the paper) and cases that are not clear are discussed. If consensus cannot be reached, we provide the most likely classification given the nature of the chain and the classification of other clinics in the chain. These clinics are also flagged to be removed or reclassified during robustness checks. Any measurement error in the classification is not likely to be systematic and instead would introduce noise rather than bias into the estimates.

We also follow the additional rules:

a) When a clinic is a "founding" clinic of a chain, it is always recorded as an acquisition since this is the central clinic owned by the parent company.

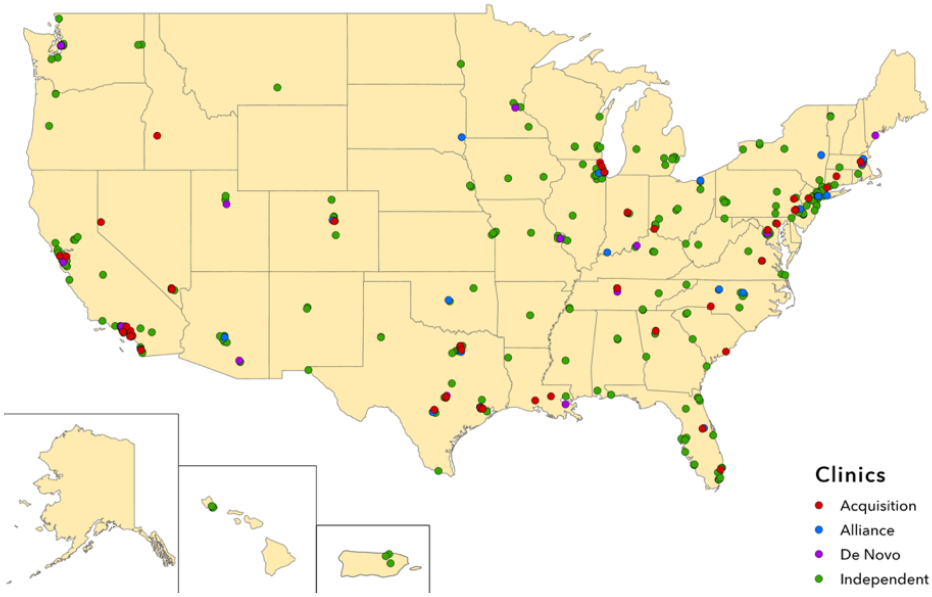
- **i.** For chains that did not previously exist and were newly formed following external investment, the date of corporate transaction for a founding clinic is the date the chain formed.
- **ii.** For founding clinics that started as a single independent practice that then began acquiring or affiliating with other clinics before external investment, their date of corporate transaction is their date of founding. In other words, these clinics are considered always treated in the sample because there is no official chain creation date. For any clinics acquired or affiliated with that clinic in the future, we use the date they joined the founding clinic's chain.

b) If a clinic had multiple transactions (i.e., the chain was acquired by another chain or company), we only record the date of first transaction under the assumption that the clinic has already received treatment. However, in robustness checks we will control for the second transaction. If a clinic leaves a chain, the post-transaction indicator turns to zero. Results are robust to dropping the years after a clinic left a chain.

A.3 Fertility Clinic Chains

We identified 11 fertility chains active from 2004 to 2018. Information about each chain is provided in Table A1 below and a map of the distribution of chain clinics in provided in Figure A1.

Figure A1. Location of US Fertility Clinics, 2018



Note: Location based on coordinates of the address provided by the clinic to the CDC in the Fertility Clinic Success Reports. "De Novo" refers to a newly built clinic that is part of a chain.

Table A1. Characteristics of Fertility Chains

Chain Name	Year Founded	Acquisition Year	Acquiring Firm	Additional Details
Boston IVF	1986	2018	NMC Health (Publicly traded international health care chain)	Growth initially focused on New England but now has a national presence.
Reproductive Medicine Associates of New Jersey (RMANJ)	1999	2017	The Valencian Infertility Institute (Privately held international fertility chain)	The merger with IVI led to rebranding to the "RMA Chain" to capture its national presence. IVI owns 70% of the company.
Huntington Reproductive Center (HRC) Fertility	1988	2017	Jinxin Fertility (backed by PE firm Warburg Pincus and Sequoia Capital)	Started as two found partners that expanded through Southern California. Jinxin was taken public in 2019.
InVitro Sciences (IVS)	1998	2017	Sverica Capital Management (PE firm)	Subsidiary of the practice management company Women's Health USA – was spun out by Sverica in 2019 and rebranded as First Fertility.

Prelude Chain	2016	2016	Lee Equity Partners (PE firm)	Prelude went on to acquire Vivere Health in 2017 and Inception Fertility in 2019. Also owns an egg storage company.
Inception Fertility	2015	2016	Lee Equity Partners (PE firm)	Founded in 2015 as company, but no clinics in the chain until PE funding in 2016. Clinics managed under the "Aspire" fertility brand.
Colorado Center for Reproductive Medicine (CCRM)	2015*	2015	TA Associates (PE firm)	*CCRM's first clinic was founded in 1987. Did not expand until acquisition in 2015. In 2021 was acquired by Unified Women's Healthcare.
Ovation Fertility	2015	2015	MTS Health Investors (PE firm, name has changed to WindRose)	Focuses more on lab management than clinics, though provides both services. Sold to Morgan Stanley Capital Partners in 2019.
Sher Institute for Reproductive Medicine (SIRM)	1982	2014	Integramed (backed by PE firm Sagard Capital Partners)	First private IVF practice. Bi-coastal growth focus. Many locations closed or rebranded post buyout.
Integramed	1985	2012	Sagard Capital Partners (PE firm)	Integramed owned Shady Grove Fertility until Shady Grove was acquired by Amulet Partners to form a new chain called US fertility. Integramed declared bankruptcy in 2020.
Vivere Health	2010	2010	LLR Partners (PE firm)	Founded as a fertility management company and equity partner. Acquired by Prelude Chain in 2017.

Appendix B. Qualitative Data

This appendix provides quotes compiled from fertility clinic and chain press releases, websites, and marketing materials to understand the stated purpose of the fertility chains and the reasons clinics would join a chain. When possible, we used archived websites to reflect the materials during the sample period and used quotes from all the chains in the sample. See the endnotes of the appendix to links to the sources.

Table B1. Selected Quotes from Archival Material Reflecting Fertility Clinic Motives for Chain Affiliation or Acquisition

	Examples	
Resource transfer	Financial Resources and Growth Strategies	"We […] develop new practices or strengthen existing ones by applying business and operations strategies that expand their markets and their market share. This may involve the development of new practice locations, embryology laboratories or ambulatory surgery centers, in order to strengthen the performance of a practice and achieve strategic growth objectives."[1] "Our partner practices' patient revenues increased 21% from 2007 to 2009. Customers that chose partner relationships with us gain access to capital and best-in-class business and clinical services. The combination of expertise and economies of scale offers them a unique formula for profitable growth."[2] "Joining us allows you to continue improving people's lives by helping them make the family of their dreams while enjoying the financial stability and growth opportunities you're looking for in the long term."[3] "Strong financial support from a leading New York–based private equity firm. For REs, [Fertility Chain] provides higher success rates, access to the first multi-center network of fertility centers offering cryogenic egg vitrification, world-class marketing and lead generation, more patients, new revenue streams, and strong financial support."[4] "[Fertility Chain] partners to offer strategic opportunities for independent practices, including: implementing creative growth strategies supporting streamlined operational costs, payer alignment, merger and acquisition plans, marketing, and risk management services."[5] "Internationally recognized for its extensive clinical experience, advanced technologies, and groundbreaking research within the field of reproductive endocrinology [Fertility Chain] will reinvest in and enhance every aspect of clinical care."[6] "Joining […] [Fertility Chain] will enable our patients and patients throughout Dallas-Fort Worth access to its leading fertility services, innovative technology and cutting-edge labs"[7] "[Fertility Chain] plans for continued growth through the addition of physicians and satellite offices"[8]

	Managerial Capabilities	"We feel very strongly about the benefits that the full complement of management support, patient sales and marketing, electronic patient information systems, and one-of-a-kind [the clinic's] IVF programs will bring to our faculty, students and patients."[9] "[The clinic] will receive [Fertility Chain's] full complement of support services, including operational and financial management, revenue cycle management, patient marketing and sales, information systems support, and various other services, including patient support […]."[10] "[Fertility Chain's] primary focus is to improve quality outcomes by enabling the strategic expansion of growing fertility practices by providing capital and operational support including marketing services, pharmacy services and back office services."[11] "Under the agreement, the newly formed LLC will receive [the Fertility Chain's] full complement of support services, including operational and financial management, revenue cycle management, patient marketing and sales, information systems support, and various other services, including patient support"[12] "Increased patient volume as a result of [Fertility Chain's] marketing efforts to educate men and women in their 20s and 30s about the benefits of preserving their fertility at its peak, as well as to target potential gamete donors.[13] "At [Fertility Chain], we make the following commitments to our patients: […] • Offer transparent pricing with no out-of-network fees or hidden costs • Use the most advanced technology to increase the odds of a successful pregnancy • Simplify the treatment process, meeting critical timelines, avoiding missed medications, and reducing travel time • Streamline communications between you and your clinicians, eliminating the frustration of missed calls • Use advanced digital technologies to help you easily manage all aspects of your fertility journey • Clearly explain the fertility treatment process, empowering you to make decisions that are right for you"[14] "Each loan program is designed to fit your individual circumstances and, once approved, the company's Fertility Loan Specialists will work closely with [Fertility Chain] to ensure the funds are secured prior to the commencement of your treatment."[15] "[Fertility Chain] announced the launch of two new programs designed to help ease the financial burdens for fertility patients that may need multiple in vitro fertilization (IVF) cycles. The [Fertility Chain's] IVF Refund and Multi-Cycle Programs offer patients the assurance that if multiple IVF cycles are necessary, they will not need to expend additional financial resources to receive them."[16]

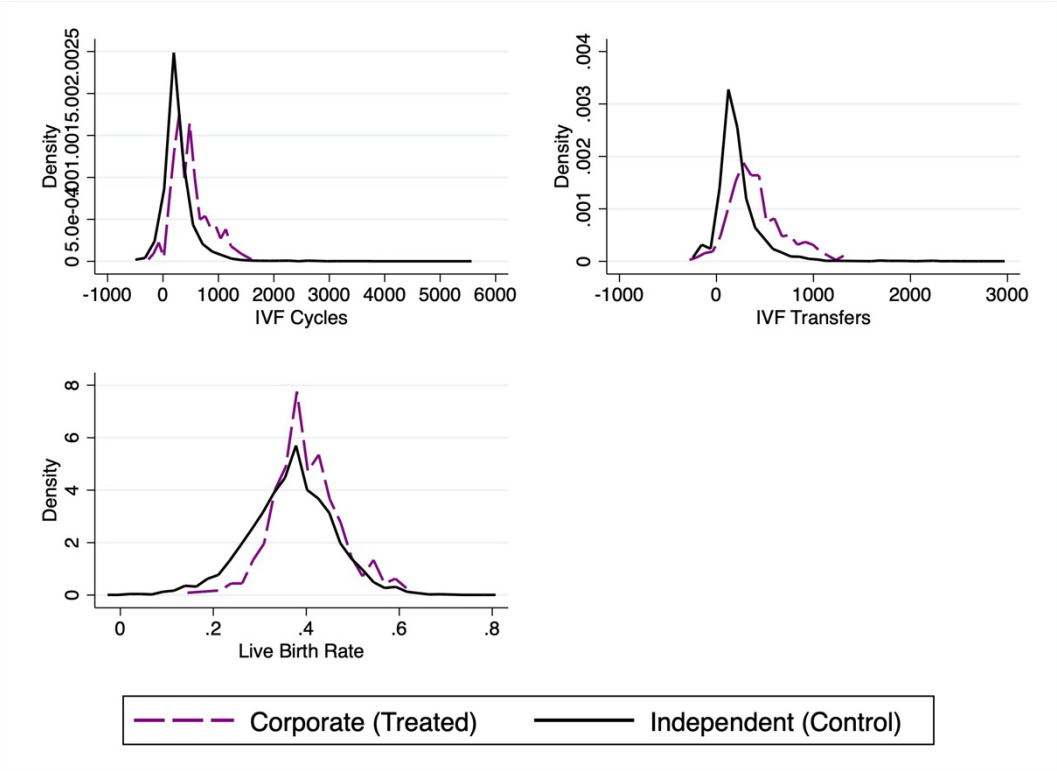
Knowledge transfer	Access to Protocols, Best Practices and New Knowledge	"Patients seeking treatment at their fertility center […] will receive improved access to the best treatment protocols and unique programs for specific conditions, new technologies that improve clinical success rates, and increased access to clinical trials and research initiatives, which often offer significant discounts on treatments and medications."[17] "Partnering with [Fertility Chain] allows us to greatly expand our work with other top-tier centers, and to leverage the strengths of this national network to further revolutionize patient care with new access to proven treatment protocols and an expanded focus on fertility preservation."[18] "Our collaboration with [Fertility Chain] will give us access to the most advanced research in the field of reproductive medicine, and will further enable us to deliver leading care to our patients"[19] "[Fertility Chain's] mission is to shift the paradigm of the IVF market by raising the standard of care, streamlining fragmented components into an integrated system, and enhancing the overall patient experience."[20] "Our collaboration with [Fertility Chain] will give us access to the most advanced research in the field of reproductive medicine, and will further enable us to deliver leading care to our patients."[21] "Best Practices Are Standard Care. [Fertility Chain] believes that patients should not have to pay more for best practices. Therefore, when research proves that techniques improve conception rates, [Fertility Chain] incorporates those techniques into their standard care wherever possible."[22] "[Fertility Chain] will give our embryologists access to a broader base of knowledge drawn from all […] labs […] and because of [Fertility Chain's] size, we will be able to take advantage of the latest techniques and equipment for services offerings we don't currently provide in house, including long-term storage, egg donation and genetic testing. By bringing these services in house, we can better control our patients' experience."[23]
	Information Sharing Process	"[Fertility Chain] was created to break down barriers to idea-sharing and collaborative care. "At its best, reproductive medicine is a tightly woven community working side by side on research, clinical trials and educational efforts. The [Fertility Chain] umbrella gives us the freedom to explore clinical and laboratory breakthroughs together, and that is exciting news for the future of infertility care."[24] "Patients will also have access to an expanded network of [Fertility Chain] experts who will come together to review and assist in complex cases."[25] "The successes at [Fertility Chain] led to the desire to share these techniques with additional clinics, thereby providing patients increased geographical access to top-quality infertility care. [Fertility Chain] is in a unique position where treatment breakthroughs are quickly applied to multiple centers, thereby furthering the positive impact for patients."[26] "The [Fertility Chain] is also a tech-enabled organization leading the industry in proprietary platforms, applications, and data and analytics. By using these purpose-built applications with other flagship technologies, we are improving patient experience and outcomes. Our proprietary […] tool keeps our network seamlessly

		connected when it comes to data and analytics, financial services, staff workflow automation, and hybrid workforce productivity."[27] "We'll look at pregnancy per transfer by physician with a blinded letter for each physician. And we'll be able to see how everybody stacks up. And if people fall below a standard deviation, we have that doctor go work with somebody who is above a standard deviation to get retrained."[28] "Under the terms of the agreement, [Fertility Chain] purchased the assets of [the clinic] and will provide a variety of services, including marketing, treatment programs for women who wish to get pregnant and a sophisticated electronic medical records system."[29] "Collaborative innovation – [Fertility Chain] scientists and physician partners share data and best practices, improving outcomes for patients and partner physician practices. Research – [Fertility Chain] is one of America's most prolific producers of IVF research, with collaborative studies continuously under way to advance the state of the art in IVF."[30]
	Single Embryo Transfer Strategy	"Striving for One Embryo-One Baby. [Fertility Chain's] founding philosophy to achieve successful pregnancy one healthy baby at a time. Although advanced embryo culturing has led to favorable pregnancy results using fresh embryo transfers, they often were the result of multiple embryo transfers, resulting in high-risk pregnancies with twins and triplets. Through elective vitrification and the adoption of blastocyst biopsy/PGS, [Fertility Chain] has improved the quality of patient care by transferring fewer embryos, reducing miscarriages and increasing healthy singleton live births."[31] "[Fertility Chain] reported a significant increase in the number of IVF cycles employing pre-implantation genetic testing. This cutting edge technology enables embryologists and fertility specialists to assess the genetic and chromosomal makeup of an embryo prior to its transfer into a woman's uterus. [Fertility Chain] now also performs almost exclusively Day 5 embryo transfers (at the blastocyst stage of development) for those patients who request or need a fresh embryo transfer. This advanced IVF lab technique allows the embryo to mature as far as it can outside the human body, again allowing embryologists and physicians an enhanced ability to select the best single embryo for transfer into the patient's uterus."[32] "The One Healthy Baby at a Time Promise. Reducing Risk for Mom and Baby. By routinely practicing Single Embryo Transfer (SET), [Fertility Chain] has drastically reduced the risk for the mother and child." "`Our rate of single embryo transfer [and subsequently our low multiple pregnancy rate] is higher than the national average, with no difference in the number of embryos transferred in fee for service versus Shared Risk patients,' comments [physician]. With advances in technology, eSET has allowed patients to have a healthy singleton pregnancy while significantly lowering the risks associated with multiple pregnancies without comprising chances of success."[33]

Appendix C. Additional Descriptive Statistics

This appendix provides additional summary statistics of the data and sample. Figure C1 shows the distribution of clinic volume and IVF success rates between corporate clinics and independent clinics before the transaction, Table C1 provides descriptive statistics based on CBSA-level patient characteristics, and Table C2 provides a targeting regression of the probability a clinic becomes acquired or affiliated with a chain using characteristics at the CBSA-level (estimation details shown before the table).

Figure C1. Distribution of Clinic Volume and IVF Success Rates for Clinics Before Corporate Transaction and Independent Clinics



Note: This histogram shows the distribution of clinic volume (IVF cycles and IVF transfers) and of IVF success rates (Live Birth Rate) in the pre-transaction period for corporate clinics and over the full sample period for independent clinics. The regression includes *state* × *year* fixed effects, with robust standard errors clustered at the clinic-level.

Table C1. CBSA-Level Characteristics of Patients Reporting Using Infertility Treatment, 2009-2018

	Fertility Chain			Independent
	Acquisition	**Affiliation**	**De Novo**	
	Pre-transaction mean	Pre-transaction mean	Mean of all years	Mean of all years
Education (%)				
Less than High School	1.06	1.55	1.35	1.37
High School	5.57	6.60	5.85	6.72
Some College	11.39	10.12	10.58	11.06
Associate or Bachelor's Degree	47.57	46.02	45.70	44.61
Graduate Degree	34.41	35.71	36.53	36.24
Race/Ethnicity (%)				
White	71.71	75.27	71.41	71.71
Black	6.02	6.22	4.89	5.69
Hispanic/Latina	9.68	7.83	9.62	9.76
Other Race	12.58	10.67	14.09	12.83
Insurance (%)				
Private	89.97	91.80	90.37	88.99
Medicaid	3.61	4.59	5.65	5.30
Self Pay	3.48	1.89	1.56	2.34
Other	2.94	1.73	2.42	3.37
Clinical Factors				
Body Mass Index	25.27	25.82	25.43	25.74
Pre-Pregnancy Diabetes	0.79	0.81	0.90	1.20
Pre-Pregnancy Hypertension	2.34	2.54	2.09	2.76
Previous Birth	28.72	28.58	30.61	30.10
Observations				
Number of Clinics	23	25	21	400
Clinic-Years	147	79	112	3249

Note: This data is from the NCHS natality data for patients reporting using any infertility treatment to deliver a baby. This reporting flag was only available starting in 2009, therefore, the data represent years 2009-2018. Since the NCHS data is reported at the county-level, we created a CBSA-level average using the county data weighted by the population in the county that was female. Clinics that that experienced a corporate transaction prior to 2009 are removed because they are now always considered part of a chain.

Targeting Regression (Table C2, below). To explore what types of fertility clinics are targeted, we estimate the probability a clinic is acquired or affiliated with a chain for the years before the transaction:

Appendix Eq. 1: $$Prob(Chain)_{ct} = \alpha_s + \alpha_t + \beta \boldsymbol{X}_{ct} + \epsilon_{ct}$$

Here, *Prob(Chain)* is set to 100 in the year before acquisition or affiliation (and zero otherwise) and all years post-transaction are dropped. We include state and year fixed effects, and $\boldsymbol{X}_{ct}$ is a vector of different clinic and market characteristics. The results are presented in Table 2, where each column groups different combinations of characteristics. Because the listed patient characteristics are only available between 2009-2018, we do not present them in combination with the market-level characteristics. The characteristics most predictive of acquisition or affiliation are total IVF cycles and the live birth rate, while clinics are much less likely to be targeted in markets with higher proportions of patients on Medicaid or other government insurance. Larger population aged 20-49 and more competitive markets are also somewhat predictive of being targeted, though become insignificant once we simultaneously control for clinic volume and IVF success rates.

Table C2. Probability Clinic Becomes Part of a Fertility Chain, CBSA Level Characteristics

	Mean	(1)	(2)	(3)	(4)	(5)
Clinic Outcomes						
Log(IVF Cycles)	5.220	0.645***			0.627***	
		(0.161)			(0.161)	
Live Birth Rate	37.051	5.500***			5.529***	
		(1.543)			(1.536)	
Market Characteristics (CBSA)						
Log(Total Population)	13.615		0.279*		0.099	
			(0.156)		(0.205)	
Log(Median Household Income)	10.949		0.610		-0.065	
			(1.194)		(1.201)	
Market Concentration						
HHI (tercile = 1)				0.933*	0.553	
				(0.479)	(0.626)	
HHI (tercile = 2)				0.605*	0.303	
				(0.339)	(0.420)	
Patient Characteristics (CBSA)						
Education (reference=Graduate School)						
Less than High School	1.359					8.608
						(13.105)
High School	6.671					-3.772
						(2.434)
Some College	11.056					-4.519*
						(2.651)
Associate or Bachelor's Degree	44.764					-0.539
						(2.037)
Race/Ethnicity (reference=White)						
Black	5.720					5.789*
						(2.958)
Hispanic/Latina	9.717					-2.939
						(2.175)
Other Race	12.772					-4.624*
						(2.621)
Insurance (reference=Private)						
Medicaid	5.214					-5.278**
						(2.474)
Self Pay	2.375					1.456
						(4.153)
Other	3.316					-6.316***
						(1.991)
Clinic-Years		5424	5385	5385	5385	3475
R^2		0.030	0.024	0.024	0.031	0.040
Ymean		1.143	1.151	1.151	1.151	1.324

Note: This table shows the estimates of Appendix Equation 1, which estimates the probability a clinic becomes part of a fertility chain. The dependent variable is an indicator for a clinic transaction in the following year (100 if yes, 0 otherwise). The sample is restricted to independent clinics and acquired or affiliated clinics for the years before transaction.

Appendix D. Robustness and Diagnostic Checks of Main Effects on Clinic Volume and Performance

This appendix provides robustness and diagnostic checks to the primary results presented in Table 2 of the manuscript. Table D1 replicates Table 2 using *CBSA* × *Year* fixed effects, Table D2 replicates Table 2 including market-level controls, Table D3 replicates Table 2 excluding markets that became more concentrated post-transaction (more details provided before the table), Table D4 replicates Table 2 showing p-values derived from wild bootstrap standard errors, and Table D5 shows results of the Goodman-Bacon (2021) decomposition. Figure D1 provides event study graphs from the canonical two-way fixed effects model by ownership structure, Figure D2 provides event study graphs using the weighted group-time estimator developed by Sun and Abraham (2021) by ownership structure and Figure D3 provides event study graphs pooling ownership types using the two-stage method of Gardner (2021). By showing event studies using these three different methods, we provide robustness to different types of estimators and approaches recently developed in the difference-in-differences literature.

Table D1. Effect of Corporate Ownership on Fertility Clinic Outcomes, Including *CBSA × Year* Fixed Effects

	(1)	(2)	(3)
	Log(Cycles)	**Log(Transfers)**	**Live Birth Rate**
Panel A: Pooled			
Post	0.309***	0.283***	0.020
	(0.086)	(0.083)	(0.013)
Panel B: Ownership Structure			
Post × Acquisition	0.395***	0.356***	0.043***
	(0.099)	(0.096)	(0.014)
Post × Affiliation	0.236*	0.220*	0.001
	(0.123)	(0.120)	(0.019)
Dep. Var. Mean	5.287	5.064	0.373
Clinic-Years	4766	4766	4766
R^2	0.908	0.907	0.682

Notes: Panel A shows the estimates of Equation 1, and Panel B shows the estimates of Equation 2, except we include *CBSA × Year* fixed effects. Therefore, observations where there is only one clinic in a CBSA-year are dropped from the regression. The live birth rate is weighted by the number of transfers within each patient age category: under age 35, 35-37, 38-40 and 41-42. The dependent variable mean captures the predicted mean for control clinics and treatment clinics before the transaction. Standard errors are clustered at the clinic level. Significance levels: *p<0.1, **p< .05, ***p<0.01

Table D2. Effect of Corporate Ownership on Fertility Clinic Outcomes, Including Market-Level Controls

	(1)	(2)	(3)	(4)	(5)	(6)
	Log(Cycles)		**Log(Transfers)**		**Live Birth Rate**	
Panel A: Pooled						
Post	0.257***	0.275***	0.215***	0.233***	0.025**	0.017*
	(0.071)	(0.063)	(0.071)	(0.063)	(0.011)	(0.010)
Panel B: Ownership Structure						
Post × Acquisition	0.278***	0.289***	0.212**	0.213**	0.050***	0.042***
	(0.099)	(0.086)	(0.099)	(0.088)	(0.013)	(0.012)
Post × Affiliation	0.239**	0.264***	0.217**	0.251***	0.005	-0.003
	(0.100)	(0.089)	(0.099)	(0.088)	(0.016)	(0.014)
Clinic FE	X	X	X	X	X	X
State × Year FE	X		X		X	
Year FE		X		X		X
Dep. Var. Mean	5.258	5.262	5.039	5.046	0.375	0.375
Clinic-Years	5627	5770	5627	5770	5627	5770
R^2	0.898	0.887	0.898	0.885	0.622	0.578

Notes: Panel A shows the estimates of Equation 1, and Panel B shows the estimates of Equation 2. All regressions include the following CBSA-level controls: the log of the total population aged 20-49 and the log of the median household income (the unemployment rate and household income are highly correlated, so we only included household income). These variables are not available for Puerto Rico, therefore, clinics in Puerto Rico are not included in this sample. The live birth rate is weighted by the number of transfers within each patient age category: under age 35, 35-37, 38-40 and 41-42. The dependent variable mean captures the predicted mean for control clinics and treatment clinics before the transaction. Standard errors are clustered at the clinic level. Significance levels: *p<0.1, **p< .05, ***p<0.01

Market Concentration Analysis (Table D3, below). Like most healthcare markets, US fertility markets are highly concentrated because there are not many providers. One reason is that reproductive endocrinology is a relatively new field of medicine having started in the 1970s, with the first IVF baby born in the US in 1981. While there has been much progress and growth, in 2014, for example, there were only 442 fertility clinics providing IVF services in the US and the average HHI was 4343. However, as seen in the table below (data from 2014), 28% of clinics are in CBSAs with 1-2 clinics, whereas 51% are in CBSAs with 6 or more clinics. The HHI is calculated using the share of total IVF cycles by year for each parent organization (the parent is the chain for corporate clinics and the clinic itself for independent clinics) in a CBSA. By comparison, the mean HHI for hospitals at the MSA-level was around 5500 for hospitals and 3300 for specialist physicians in 2014 (Fulton 2017).

Number of Clinics	% of Clinics in CBSA	HHI
1-2	28.02	8578
3-5	20.96	4156
6-9	17.31	2720
10+	33.71	1770

As mentioned in the manuscript, fertility chains are slightly more likely to target clinics in more competitive markets and most chain transactions occur across markets rather than within markets. To confirm this, we calculate that number of markets where an acquisition or an affiliation increased the number of clinics belonging to a single chain. For example, if there were three independent clinics in a CBSA before an acquisition, and one clinic was acquired, then there would be no consolidation at the chain level in that market. However, if a second clinic in that CBSA was acquired by the same chain, then that CBSA would become more consolidated as the result of the acquisition. We identify 5 CBSAs (out of 147) where an acquisition or affiliation led to chain consolidation (the 5 CBSAs include Boston-Cambridge-Newton, Chicago-Naperville-Elgin, Dallas-Fort Worth-Arlington, Los Angeles-Long Beach-Anaheim and New York-Newark-Jersey City).

We then calculate whether a market became more concentrated because of an affiliation or an acquisition. Defining a market as a CBSA, we locate market-years where an acquisition or affiliation occurs in the following year. Next, we calculate a counterfactual HHI based on a chain's pre-transaction shares but post-transaction ownership within a market. This counterfactual HHI represents the post-transaction change in HHI only driven by the clinic acquisition or affiliation in that market. Using this methodology, we identify 3 counties and 3 CBSAs as having transactions that induce increases in HHI. In Appendix Table D3 we provide evidence that results are robust to excluding these markets, suggesting that the impact of corporate ownership on fertility clinic outcomes is not driven by changes in market concentration. Furthermore, as seen in Appendix Table D1, results are also robust to the inclusion of $CBSA \times year$ fixed effects, which would help account for market level changes.

Table D3. Effect of Corporate Ownership on Fertility Clinic Outcomes, Robustness to Changes in Market (CBSA) Concentration

	(1)	(2)	(3)
	Log(Cycles)	**Log(Transfers)**	**Live Birth Rate**
Panel A: Exclude Markets Where Corporate Transaction Increased HHI			
Post × Acquisition	0.261***	0.186*	0.051***
	(0.095)	(0.096)	(0.012)
Post × Affiliation	0.239**	0.211**	0.001
	(0.099)	(0.098)	(0.015)
Dep. Var. Mean	5.267	5.044	0.376
Clinic-Years	5271	5271	5271
R^2	0.905	0.905	0.635
Panel B: Exclude Markets With Any Corporate Consolidation			
Post × Acquisition	0.272**	0.191	0.040***
	(0.119)	(0.117)	(0.013)
Post × Affiliation	0.118	0.106	0.003
	(0.082)	(0.084)	(0.021)
Dep. Var. Mean	5.233	5.015	0.379
Clinic-Years	4091	4091	4091
R^2	0.907	0.907	0.627

Notes: This table shows the estimates of Equation 2 run on sample excluding different markets. Panel A excludes data from 3 CBSAs where a clinic transaction increased the HHI in that market. Panel B excludes data from 5 CBSAs where a clinic transaction increased the number of clinics under the same ownership in the same market. The dependent variable mean captures the predicted mean for control clinics and treatment clinics before the transaction. Standard errors are clustered at the clinic level. Significance levels: *p<0.1, **p< .05, ***p<0.01

Table D4. Effect of Corporate Ownership on Fertility Clinic Outcomes, Wild Bootstrap Standard Errors

	(1)	(2)	(3)	(4)	(5)	(6)
	Log(Cycles)		**Log(Transfers)**		**Live Birth Rate**	
	b/wild p-val		b/wild p-val		b/wild p-val	
Panel A: Pooled						
Post	0.258***	0.282***	0.216***	0.237***	0.026**	0.018*
	0.000	0.000	0.004	0.000	0.021	0.072
Panel B: Ownership Structure						
Post × Acquisition	0.282**	0.297***	0.214**	0.217**	0.051***	0.043***
	0.011	0.000	0.046	0.017	0.001	0.001
Post × Affiliation	0.238**	0.268***	0.217**	0.253***	0.004	-0.003
	0.015	0.004	0.031	0.004	0.794	0.843
Clinic FE	X	X	X	X	X	X
State × Year FE	X		X		X	
Year FE		X		X		X
Dep. Var. Mean	5.252	5.256	5.035	5.040	0.374	0.375
Clinic-Years	5666	5809	5666	5809	5666	5809
R^2	0.899	0.887	0.898	0.886	0.625	0.579

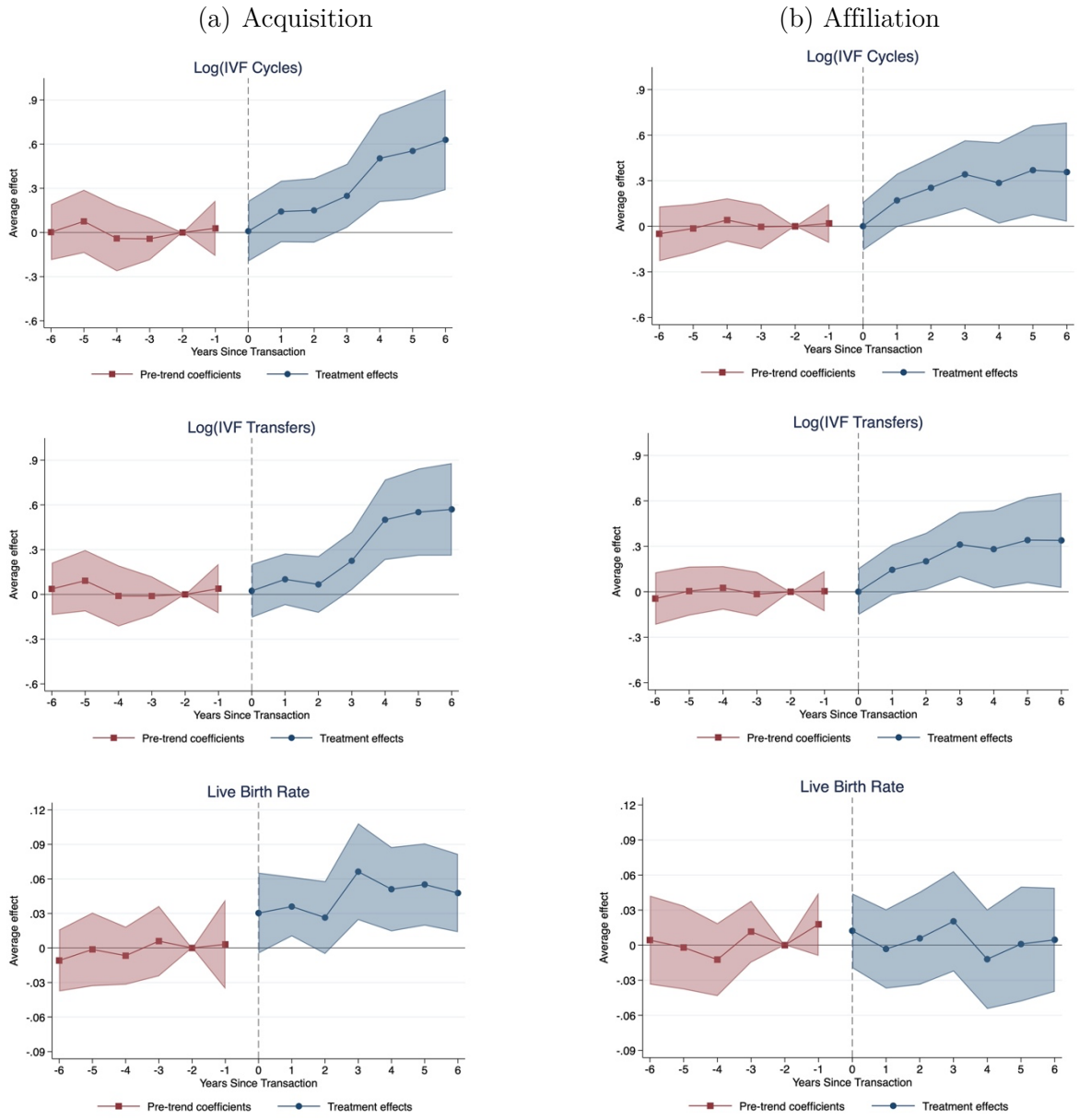
Notes: Panel A shows the estimates of Equation 1, and Panel B shows the estimates of Equation 2. The live birth rates are weighted by the number of transfers within each patient age category: under age 35, 35-37, 38-40 and 41-42. The dependent variable mean captures the predicted mean for control clinics and treatment clinics before the transaction. Standard errors are clustered at the clinic level. Significance levels: *p<0.1, **p< .05, ***p<0.01

Table D5. Goodman-Bacon Decomposition of Treatment Effects

		(1)	(2)	(3)
	Weight	**Log(Cycles)**	**Log(Transfers)**	**Live Birth Rate**
Panel A: Pooled				
Post		0.245***	0.205***	0.019*
		(0.072)	(0.072)	(0.011)
Timing groups	0.093	0.113	0.109	-0.006
Never_v_timing	0.881	0.278	0.229	0.021
Within	0.026	-0.399	-0.254	0.026
Panel B: Ownership Structure				
Post × Acquisition		0.267***	0.197**	0.041***
		(0.096)	(0.099)	(0.012)
Timing groups	0.052	0.129	0.125	0.009
Never_v_timing	0.907	0.294	0.212	0.042
Within	0.041	-0.146	-0.048	0.053
Post × Affiliation		0.223**	0.213**	-0.002
		(0.104)	(0.102)	(0.016)
Timing groups	0.029	0.115	0.111	-0.009
Never_v_timing	0.966	0.216	0.208	-0.004
Within	0.005	2.368	1.991	0.474

Notes: Panel A shows the estimates of the Goodman-Bacon decomposition when pooling together both ownership structures, and Panel B shows the decomposition by ownership structure. The decomposition requires a balanced panel, in this setting, that represents 246 clinics and 3690 clinic-years that had 15 years of data. Standard errors are clustered at the clinic level. Significance levels: *p<0.1, **p< .05, ***p<0.01

Figure D1. Event Study Results by Ownership Structure, TWFE

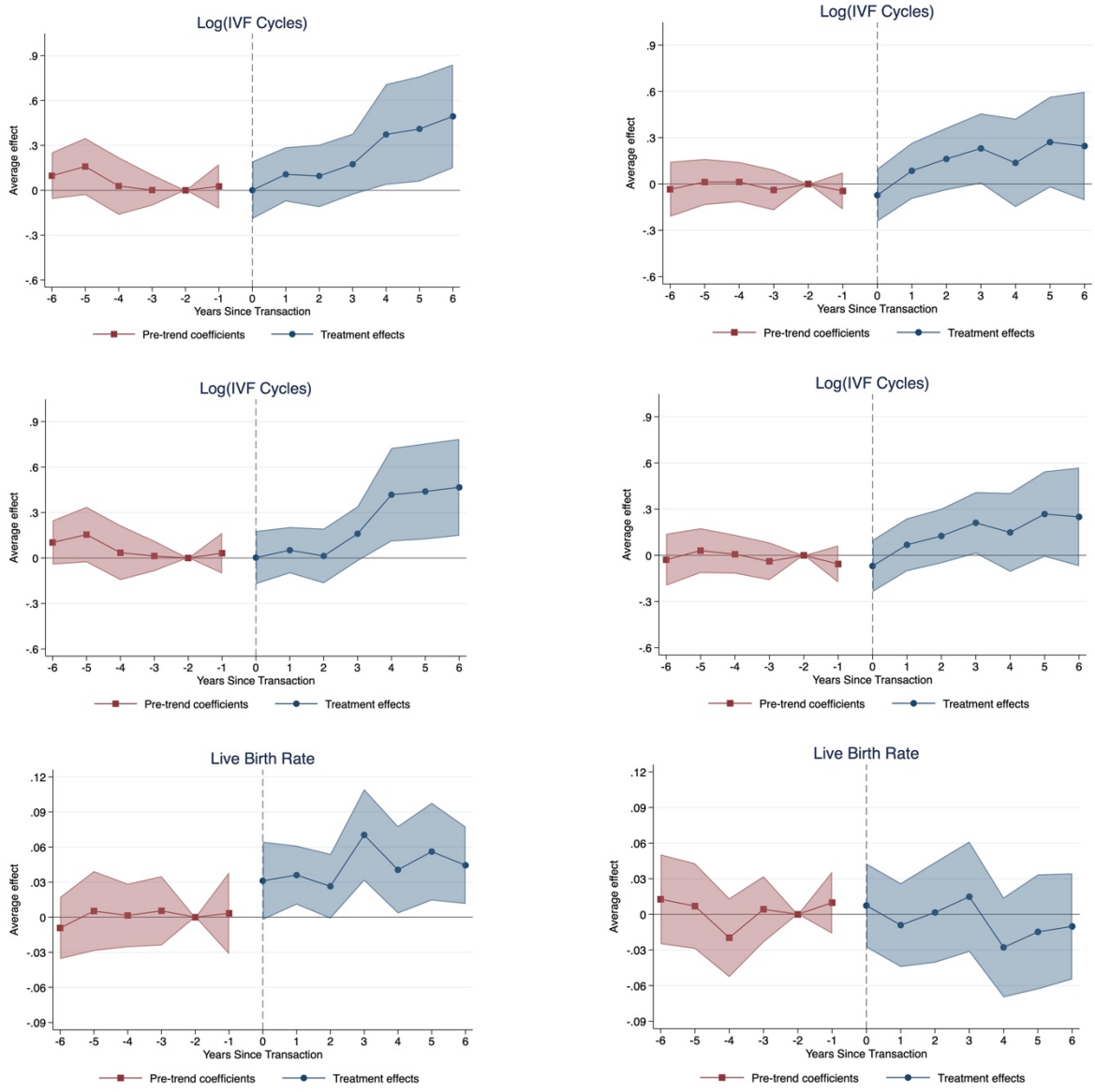


Notes: This figure shows the β_1 and β_2 estimates of Equation 2 interacted with indicators for the year relative to the transaction year. The reference period is two years before the transaction. Bands indicate 95% confidence intervals constructed from clinic-level clustered standard errors.

Figure D2. Event Study Results by Ownership Structure, Sun and Abraham (2021)

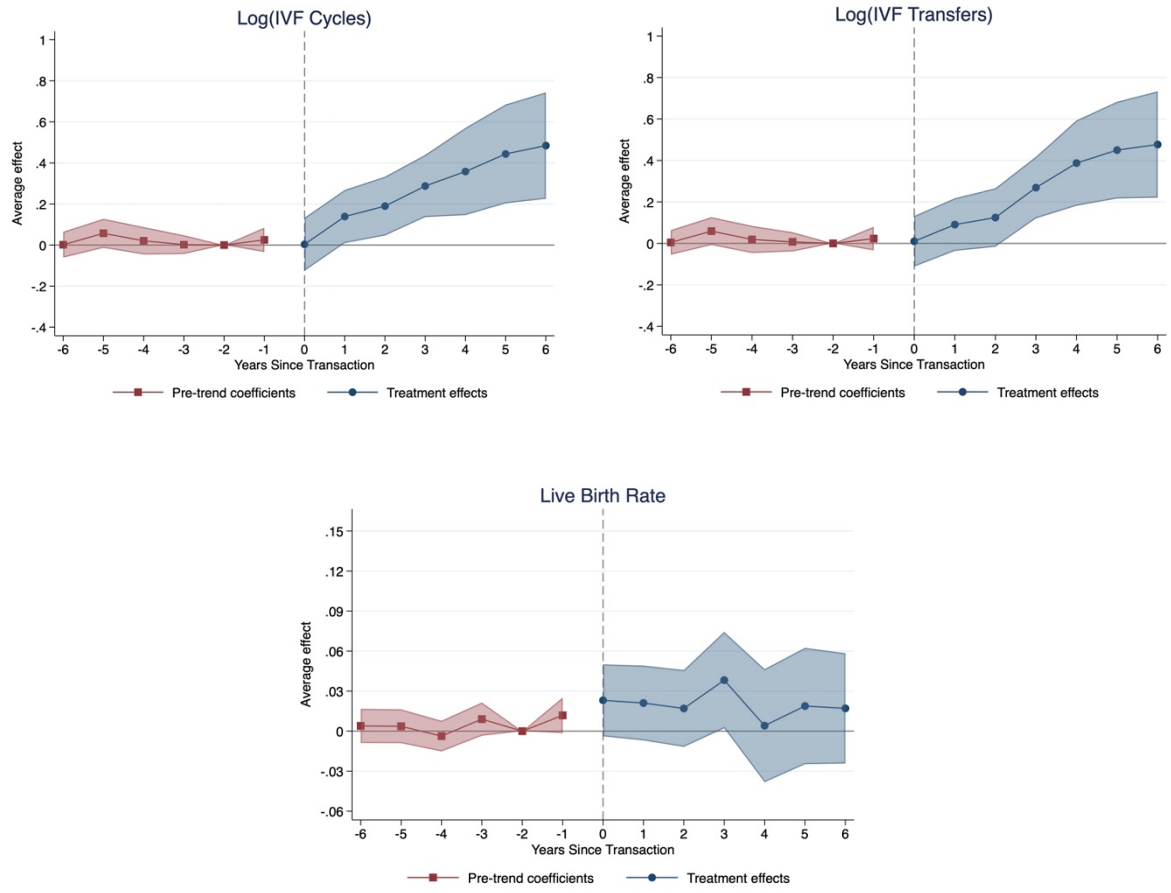
(a) Acquisition

(b) Affiliation



Notes: This figure shows the β_1 and β_2 estimates of Equation 2 interacted with indicators for the year relative to the transaction year. The reference period is two years before the transaction. Bands indicate 95% confidence intervals constructed from clinic-level clustered standard errors. These event studies were created using the user written stata command eventstudyinteract by Sun (2021).

Figure D3. Pooled Event Study Results, Two-Stage DD (Gardner 2021)



Notes: This figure shows the β estimates of Equation 1 interacted with indicators for the year relative to the transaction year. The reference period is two years before the transaction. Bands indicate 95% confidence intervals constructed from clinic-level clustered standard errors. These event studies were created using the user written stata command did2s by Butts (2021).

Appendix E. Power Calculations

This appendix provides power curves based on simulation analysis to assess to what extent we are powered to detect different effect sizes. For both clinic volume and IVF success rates we simulate power to assess our ability to statistically reject the null hypothesis that $\beta_1 = 0$ and $\beta_2 = 0$ in Equation 2. We calculate power for a range of values, allowing for a type I error rate of 5%. We implement this procedure in three steps:

1) Calculate the parameters of Equation 2 (shown below as a reminder) for the control variables and fixed effects using only data from independent clinics and corporate clinics before transaction.

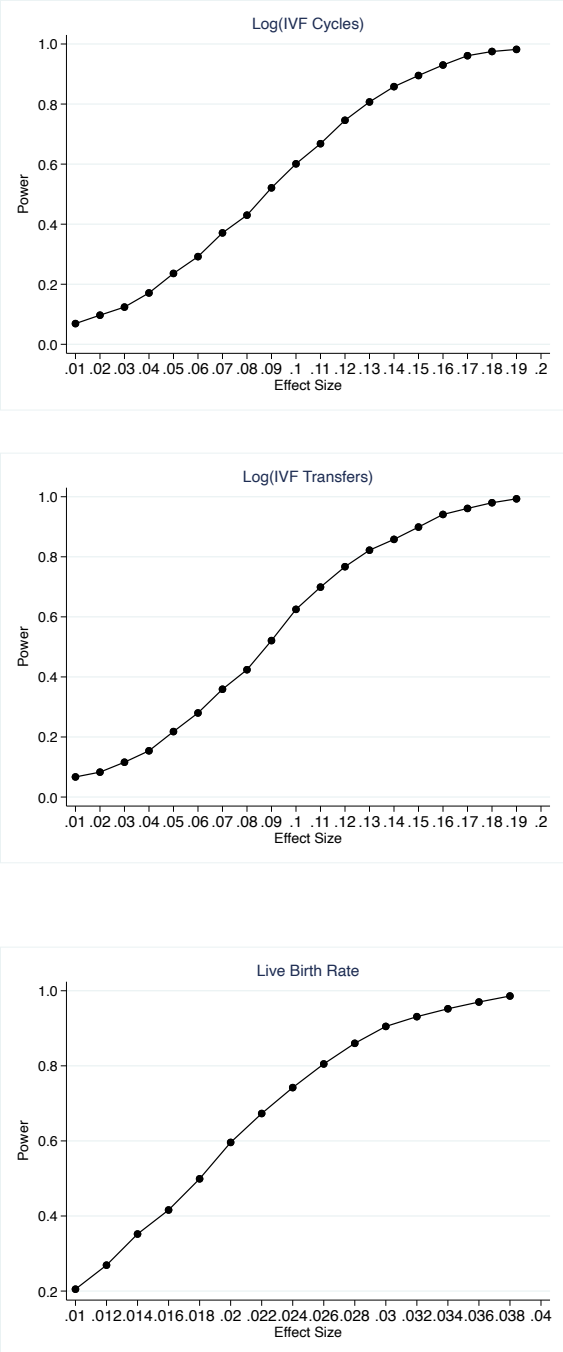
Eq. 2: $$Y_{ct} = \beta_1(Post \times Acquisition)_{ct} + \beta_2(Post \times Affiliation)_{ct} + \gamma \boldsymbol{X}_{ct} + \theta_c + \theta_{st} + \epsilon_{ct}$$

2) For each clinic, simulate outcomes based on the proposed true effect size of β_1 and β_2 and a random error, drawn from a normal distribution with mean zero and standard deviation calculated from the residuals from step 1. Repeat this process 1000 times to construct 1000 simulated samples.
3) Estimate Equation 2, clustering standard errors at the clinic level, in each of the 1000 simulated samples, and record the percentage of cases in which the *p-value* when testing $\beta_1 = 0$ and $\beta_2 = 0$ is below 0.05.

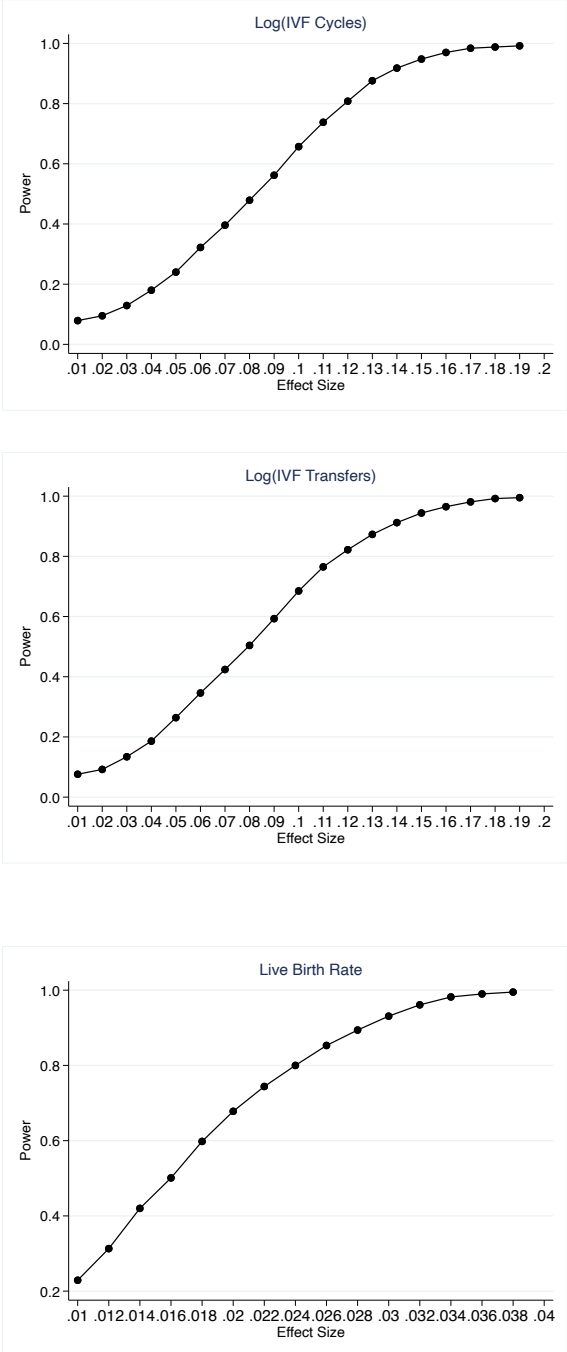
As seen in Figure E1, the simulations suggest the difference-in-differences regressions are well powered. For example, at 80% power, we can detect a 13% and 12% change in total cycles for acquired and affiliated clinics respectively, which is roughly half of the observed increase seen in most analysis. For the live birth rate, at 80% power we can detect a 2.6 pp and 2.4 pp change for acquired and affiliated clinics, respectively. While these are meaningful effect sizes, acquired clinic effect sizes are more than 1pp above this value in all regressions and affiliated clinic effect sizes are often close to zero.

Figure E1. Power Curve for Fertility Clinic Outcomes by Ownership Structure

(a) Acquisition



(b) Affiliation



Notes: Figures plot power against hypothesized effect sizes assuming a type I error rate of 0.05. For the singleton birth rate, at 80% power we can detect a 2.2 pp and 2.0 pp change for acquired and affiliated clinics, respectively. For the multiple birth rate, at 80% we can detect a 1.5 pp and 1.3 pp change for acquired and affiliated clinics, respectively. Output for these additional outcome variables available upon request.

Appendix F. Assessing the Role of Patient and Clinic Selection

This appendix provides robustness checks to the main DD that mitigate concerns of patient selection and the non-random selection of clinics into fertility chains. Table F1 shows results of the live birth rate, singleton birth rate and multiple birth rate using different patient controls and weights, Table F2 shows results using different matched control groups, Table F3 shows the summary statistics for the matched samples, Table F4 shows results using alternative samples, Table F5 shows additional robustness that account for unique features of the data and Table F6 shows cross-sectional analysis using new variables available in 2017 and 2018 for first time patients using IVF.

Table F1. Effect of Corporate Ownership on Fertility Clinic Outcomes, Patient Robustness

	(1) **Live Birth Rate**	(2) **Singleton Birth Rate**	(3) **Multiple Birth Rate**
Panel A: Including Clinic-Level Patient Diagnosis as Controls (2004-2018)			
Post × Acquisition	0.052***	0.067***	-0.015**
	(0.012)	(0.012)	(0.006)
Post × Affiliation	0.007	0.011	-0.005
	(0.016)	(0.014)	(0.006)
Dep. Var. Mean	0.374	0.273	0.101
Clinic-Years	5666	5666	5666
R^2	0.626	0.620	0.512
Panel B: Including CBSA-Level Patient Characteristics as Controls (2009-2018)			
Post × Acquisition	0.036***	0.064***	-0.028***
	(0.012)	(0.013)	(0.006)
Post × Affiliation	0.013	0.020	-0.007
	(0.014)	(0.014)	(0.008)
Dep. Var. Mean	0.389	0.293	0.096
Clinic-Years	3742	3742	3742
R^2	0.656	0.642	0.550
Panel C: Main DD Estimation Weighted by Clinic IVF Cycles (2004-2018)			
Post × Acquisition	0.030***	0.039***	-0.009*
	(0.011)	(0.013)	(0.005)
Post × Affiliation	0.003	0.007	-0.004
	(0.016)	(0.016)	(0.005)
Dep. Var. Mean	0.390	0.295	0.095
Clinic-Years	5666	5666	5666
R^2	0.752	0.776	0.685

Notes: This table shows the estimates of Equation 2 including different sets of patient controls. In Panel B, the sample excludes years before 2009 because the NCHS data did not report whether a patient used infertility treatment until 2009, and therefore, includes a smaller sample of corporate clinics. For the same sample period (2009-2018) without including vital statistics controls, the b/se for the live birth rate in acquired and affiliated clinics are .035 (.012) and .004 (.015), respectively. We also find nearly identical results when using county-level patients characteristics as controls (available upon request). The dependent variable mean captures the predicted mean for control clinics and treatment clinics before the transaction. Standard errors are clustered at the clinic level. Significance levels: *p<0.1, **p< .05, ***p<0.01

Table F2. Effect of Corporate Ownership on Fertility Clinic Outcomes, Matched Sample

	(1)	(2)	(3)
	Log(Cycles)	**Log(Transfers)**	**Live Birth Rate**
Panel A: Matched Sample (Cycles Only)			
Post × Acquisition	0.288**	0.264**	0.040***
	(0.119)	(0.114)	(0.015)
Post × Affiliation	0.202*	0.177*	-0.023
	(0.110)	(0.104)	(0.018)
Dep. Var. Mean	5.719	5.498	0.398
Clinic-Years	1514	1514	1514
R2	0.924	0.926	0.727
Panel B: Matched Sample (Cycles, Live Birth Rate, Share of Patients<35)			
Post × Acquisition	0.341***	0.300***	0.050***
	(0.120)	(0.106)	(0.016)
Post × Affiliation	0.166	0.152	-0.009
	(0.123)	(0.121)	(0.019)
Dep. Var. Mean	5.567	5.355	0.395
Clinic-Years	1314	1314	1314
R2	0.912	0.917	0.686

Notes: This table shows the estimates of Equation 2 using a matched control group. Panel A includes independent clinics matched based on 1-1 coarsened exact matching on a clinic's IVF cycles in the year before transaction (62 treated and 62 control clinics), and Panel B repeats the match process but matches on IVF cycles, the live birth rate, and the share of patients under 35 (55 treated and 55 control clinics). The dependent variable mean captures the predicted mean for control clinics and treatment clinics before the transaction. Standard errors are clustered at the clinic level. Significance levels: *p<0.1, **p< .05, ***p<0.01

Table F3. Fertility Clinic Summary Statistics: Matched Sample, 2004-2018

	Matched (IVF Cycles)			Matched (IVF Cycles, Live Birth Rate, Share Under 35)		
	Fertility Chain		**Independent**	**Fertility Chain**		**Independent**
	Acquisition	**Affiliation**		**Acquisition**	**Affiliation**	
	Pre-transaction mean	Pre-transaction mean	Mean of all years	Pre-transaction mean	Pre-transaction mean	Mean of all years
Clinic Volume						
IVF Cycles	563.81	515.43	440.56	487.01	482.00	333.79
IVF Transfers	453.57	401.08	347.91	388.09	376.40	260.75
Log(IVF Cycles)	6.05	5.90	5.56	5.91	5.84	5.38
Log(IVF Transfers)	5.83	5.67	5.34	5.69	5.62	5.17
Birth Rates (%)						
Live Birth Rate	41.80	41.98	37.74	41.13	41.95	38.17
Singleton Birth Rate	30.68	31.21	27.48	30.09	31.17	27.51
Multiple Birth Rate	11.10	10.79	10.19	11.01	10.79	10.60
Patient Characteristics (%)						
Share of Patients < 35 (transfers)	51.79	52.06	49.54	53.27	52.16	52.54
Share of Patients 35-37 (transfers)	24.17	23.17	23.87	23.78	23.04	23.70
Share of Patients ≥ 38 (transfers)	24.05	24.77	26.58	22.94	24.80	23.76
Diagnosis, Tubal Factor	10.94	10.72	11.75	11.56	10.74	13.24
Diagnosis, Ovulatory Dysfunction	11.14	11.28	10.94	11.58	11.56	11.88
Diagnosis, Diminished Ovarian	23.90	23.24	23.21	22.50	23.82	20.09
Diagnosis, Endometriosis	7.80	7.32	7.03	7.91	7.77	9.75
Diagnosis, Uterine Factor	3.74	4.06	3.88	3.71	4.15	3.86
Diagnosis, Male Factor	24.89	22.77	24.96	25.51	22.89	27.89
Diagnosis, Other	13.18	12.81	12.68	11.99	12.59	10.82
Diagnosis, Unknown	10.44	11.46	11.73	11.00	10.94	9.78
Observations						
Number of Clinics	33	29	62	29	26	55
Clinic-Years	283	193	849	237	180	744

Notes: All summary statistics are at the clinic-year level. Clinic volume, birth rates, and patient characteristics include adjustment for year effects to account for changes in reporting in the CDC ART data (therefore, there will be differences between these statistics and those reported in Table 1). The samples are constructed using 1-1 coarsened exact matching on a clinic's IVF cycles in the year before transaction or using a clinic's IVF cycles, live birth rates and share of patients under 35 years of age in the year before transaction. In the latter matched sample, not all treated clinics were able to be matched to an independent clinic.

Table F4. Effect of Corporate Ownership on Fertility Clinic Outcomes, Alternative Clinic Samples

	(1)	(2)	(3)
	Log(Cycles)	**Log(Transfers)**	**Live Birth Rate**
Panel A: Clinics with at Least 150 Cycles a Year			
Post × Acquisition	0.202**	0.174*	0.038***
	(0.094)	(0.093)	(0.013)
Post × Affiliation	0.158	0.151	-0.002
	(0.105)	(0.103)	(0.016)
Dep. Var. Mean	5.805	5.577	0.387
Clinic-Years	3541	3541	3541
R2	0.840	0.840	0.723
Panel B: Balanced Panel (15 Years)			
Post × Acquisition	0.309***	0.254**	0.051***
	(0.111)	(0.112)	(0.013)
Post × Affiliation	0.211*	0.190	0.013
	(0.122)	(0.120)	(0.019)
Dep. Var. Mean	5.550	5.340	0.375
Clinic-Years	3570	3570	3570
R2	0.905	0.906	0.634
Panel C: Excluding Independent Clinics that Restructured or Closed			
Post × Acquisition	0.263**	0.202*	0.047***
	(0.105)	(0.104)	(0.013)
Post × Affiliation	0.213**	0.194*	0.004
	(0.104)	(0.103)	(0.016)
Dep. Var. Mean	5.319	5.099	0.379
Clinic-Years	4238	4238	4238
R2	0.908	0.908	0.628

Notes: This table shows the estimates of Equation 2 for different samples. Panel A includes a sample of independent clinics that perform at least 150 cycles per year over the sample period, Panel B includes clinic present in all years of data from 2004 to 2015 and Panel C excludes independent clinics that ever restructured or closed. The dependent variable mean captures the predicted mean for control clinics and treatment clinics before the transaction. Standard errors are clustered at the clinic level. Significance levels: *p<0.1, **p< .05, ***p<0.01

Table F5. Effect of Corporate Ownership on Fertility Clinic Outcomes, Additional Robustness

	(1)	(2)	(3)
	Log(Cycles)	**Log(Transfers)**	**Live Birth Rate**
Panel A: Excluding the Year 2018			
Post × Acquisition	0.281***	0.237**	0.045***
	(0.105)	(0.105)	(0.014)
Post × Affiliation	0.211**	0.197**	-0.004
	(0.101)	(0.098)	(0.017)
Dep. Var. Mean	5.222	5.036	0.372
Clinic-Years	5287	5287	5287
R^2	0.903	0.904	0.645
Panel B: Controlling for a Second Acquisition			
Post × Acquisition	0.266***	0.200**	0.050***
	(0.095)	(0.094)	(0.013)
Post × Affiliation	0.233**	0.212**	0.004
	(0.100)	(0.099)	(0.016)
Dep. Var. Mean	5.252	5.035	0.374
Clinic-Years	5666	5666	5666
R^2	0.899	0.898	0.625
Panel C: Removing Clinics with Uncertain Classification			
Post × Acquisition	0.292***	0.221**	0.047***
	(0.103)	(0.104)	(0.012)
Post × Affiliation	0.254**	0.233**	0.011
	(0.115)	(0.114)	(0.018)
Dep. Var. Mean	5576	5576	5576
Clinic-Years	0.900	0.899	0.623
R^2	5.249	5.032	0.374

Notes: This table shows the estimates of Equation 2 for different samples and use of alternative control variables. Panel A includes an additional indicator for whether a clinic experienced a second acquisition event (1 for the second acquisition, 0 otherwise); Panel B removes the year 2018 to assess robustness to changes in data reporting in that year; and Panel C removes clinics for which there was potential uncertainty in clinic ownership and removes chains that did not receive external funding until 2018. The dependent variable mean captures the predicted mean for control clinics and treatment clinics before the transaction. Standard errors are clustered at the clinic level. Significance levels: *p<0.1, **p< .05, ***p<0.01

Table F6. Association between Corporate Ownership and Live Birth Rates for First Time IVF Patients, Years 2017 and 2018

	(1)	(2)
	Live Birth Rate (First Intended Retrieval)	**Live Birth Rate (All Intended Retrievals)**
Acquired=1	0.088***	0.093***
	(0.017)	(0.016)
Affiliation=1	0.057***	0.069***
	(0.015)	(0.017)
Dep. Var. Mean	0.370	0.429
Clinic-Years	759	759
R^2	0.263	0.238

Notes: This table shows an adaptation of Equation 2 without clinic fixed effects using data only available in 2017 and 2018. Rather than post-transaction indicators, "*Acquired==1*" is equal to 1 for acquired clinics and zero otherwise and "*Affiliation=1*" is equal to 1 for affiliated clinics and zero otherwise. The dependent variable mean captures the predicted mean for control clinics and treatment clinics in 2017 and 2018. Standard errors are clustered at the clinic level. Significance levels: *p<0.1, **p< .05, ***p<0.01

Appendix G. Mechanisms

This appendix uses alternative specifications and outcomes to explore underlying mechanisms driving the observed changes in fertility clinic volume and IVF success rates. Table G1 shows improvements in outcomes for patients of different ages as well as decomposes the live birth rate into multiple and singleton births, Table G2 shows clinic improvements in the live birth rate relative to the average of their fertility chain, Table G3 shows changes in the live birth rate by whether the fertility chain has below or above average live birth rates before the first chain transaction, Table G4 shows changes in the live birth rate by terciles of a clinic's pre-transaction clinic volume, Table G5 shows whether clinics engage in market expansion or business stealing using commuting zones as markets and Table G6 explores the role of private equity investment into fertility chains.

Table G1. Effect of Corporate Ownership on Fertility Clinic IVF Success Rates by Patient Age Category

	(1)	(2)	(3)	(4)	(5)
	Log(Cycles)	**Log(Transfers)**	**Live Birth Rate**	**Singleton Birth Rate**	**Multiple Birth Rate**
Panel A: Patients All Ages (Weighted)					
Post × Acquisition	0.282***	0.214**	0.051***	0.067***	-0.016***
	(0.098)	(0.099)	(0.013)	(0.013)	(0.006)
Post × Affiliation	0.238**	0.217**	0.004	0.011	-0.007
	(0.100)	(0.099)	(0.016)	(0.014)	(0.006)
Dep. Var. Mean	5.252	5.035	0.374	0.273	0.101
Clinic-Years	5666	5666	5666	5666	5666
R^2	0.899	0.898	0.625	0.618	0.508
Panel B: Patients Under 35					
Post × Acquisition	0.274***	0.202*	0.038***	0.060***	-0.022***
	(0.104)	(0.104)	(0.013)	(0.014)	(0.008)
Post × Affiliation	0.199**	0.176*	-0.003	0.007	-0.010
	(0.094)	(0.091)	(0.016)	(0.015)	(0.008)
Dep. Var. Mean.	4.378	4.180	0.437	0.308	0.129
Clinic-Years	5666	5666	5665	5665	5665
R^2	0.875	0.872	0.521	0.508	0.433
Panel C: Patients 35-37					
Post × Acquisition	0.222**	0.130	0.040**	0.050***	-0.011
	(0.109)	(0.106)	(0.018)	(0.016)	(0.008)
Post × Affiliation	0.265***	0.221**	0.014	0.023	-0.009
	(0.102)	(0.100)	(0.021)	(0.019)	(0.010)
Dep. Var. Mean	3.651	3.388	0.369	0.274	0.095
Clinic-Years	5666	5666	5656	5656	5656
R^2	0.824	0.802	0.405	0.414	0.265
Panel D: Patients 38 and Over					
Post × Acquisition	0.273***	0.153	0.091***	0.105***	-0.015***
	(0.101)	(0.113)	(0.019)	(0.019)	(0.006)
Post × Affiliation	0.263**	0.265**	0.024	0.012	0.010*
	(0.112)	(0.115)	(0.019)	(0.018)	(0.006)
Dep. Var. Mean	3.779	3.402	0.256	0.205	0.050
Clinic-Years	5666	5666	5655	5655	5655
R^2	0.866	0.816	0.417	0.406	0.233

Notes: This table shows the estimates of Equation 2 run for three separate age categories: all patients with results weighted by patient age group (Panel A), patients under the age of 35 (Panel B), patients between the ages of 35 and 37 (Panel C) and patients of age 38 or older (Panel D). Note that because of changes in data reporting the oldest patients included in the sample are 42 years old. The dependent variable mean captures the predicted mean for control clinics and treatment clinics before the transaction. Standard errors are clustered at the clinic level. Significance levels: *p<0.1, **p< .05, ***p<0.01

Table G2. Effect of Corporate Ownership on Fertility Clinic Live Birth Rate Relative to the Fertility Chain Average Live Birth Rate

	Difference in Live Birth Rate Between Clinic and Network Average
Post × Acquisition(BelowAvg=1)	0.082**
	(0.036)
Post × Acquisition(AboveAvg=1)	0.039***
	(0.015)
Post × Affiliation(BelowAvg=1)	0.034**
	(0.013)
Post × Affiliation(AboveAvg=1)	-0.009
	(0.021)
Dep. Var. Mean	0.050
Clinic-Years	5666
R^2	0.582

Notes: This table provides an extension of Equation 2 where the outcome variable is the difference between a clinic's own live birth rate and the average live birth rate of the clinic's chain (the average excludes a clinic's own live birth rate) for each year of the sample. We decompose the change between clinics above their fertility chain's average live birth rate pre-transaction (i.e., positive-valued difference) and clinics below their fertility chain's average live birth rate pre-transaction (i.e., negative-valued difference). The dependent variable mean captures the predicted mean for control clinics and treatment clinics before the transaction. Standard errors are clustered at the clinic level. Significance levels: *p<0.1, **p< .05, ***p<0.01

Table G3. The Effect of Corporate Ownership on the Live Birth Rate by Fertility Chain Pre-Transaction Live Birth Rates

	Live Birth Rate
Post × Acquisition(ChainBelow=1)	0.028**
	(0.012)
Post × Acquisition(ChainAbove=1)	0.073***
	(0.019)
Post × Affiliation(ChainBelow=1)	-0.006
	(0.021)
Post × Affiliation(ChainAbove=1)	0.029*
	(0.016)
Dep. Var. Mean	0.375
Clinic-Years	5666
R^2	0.625

Notes: This table provides an extension of Equation 2 where the post-transaction indicators are interacted with an indicator for whether the clinic is acquired by or affiliates with a fertility chain with below or above median live birth rates. To calculate each chain's live birth rate, we take the average live birth rate of the clinics that are already in the chain before the first transaction takes place in our sample. However, since some chains are newly created by PE firms during our sample period (i.e., had no clinics already in the chain), for these chains we use the average live birth rate of the flagship clinic(s) first acquired to create the chain in the years before the transaction occurs. Using these pre-transaction chain-level live birth rates, we then divide chains into those with below or above the median live birth rates. The dependent variable mean captures the predicted mean for control clinics and treatment clinics before the transaction. Standard errors are clustered at the clinic level. Significance levels: *p<0.1, **p< .05, ***p<0.01

Table G4. The Effect of Corporate Ownership on the Live Birth Rate by Terciles of Clinic Pre-Transaction IVF Cycles

	Live Birth Rate
Post × Acquisition(Low=1)	0.068***
	(0.021)
Post × Acquisition(Med=1)	0.043*
	(0.024)
Post × Acquisition(High=1)	0.042***
	(0.014)
Post × Affiliation(Low=1)	0.001
	(0.023)
Post × Affiliation(Med=1)	0.006
	(0.036)
Post × Affiliation(High=1)	0.006
	(0.016)
Dep. Var. Mean	0.374
Clinic-Years	5666
R^2	0.625

Notes: This table provides an extension of Equation 2 with clinics divided into terciles based on their pre-transaction average of IVF cycles. For example, *Acquisition(Low=1)* is an indicator equal to 1 if an acquired clinic was in the bottom tercile of acquired clinics based on its pre-transaction average of IVF cycles. The dependent variable mean captures the predicted mean for control clinics and treatment clinics before the transaction. Standard errors are clustered at the clinic level. Significance levels: *p<0.1, **p< .05, ***p<0.01

Table G5. Market Expansion Analysis, IV Estimates Based on Commuting Zones

	(1)	(2)	(3)	(4)
	Total Market Cycles		**Total Market Live Births**	
	Independent Clinics	All Clinics	Independent Clinics	All Clinics
Total Corporate Cycles	0.005	1.005***		
	(0.177)	(0.177)		
Total Corporate Live Births			-0.252*	0.748***
			(0.135)	(0.135)
First Stage: F-Stat	63.420	63.420	61.027	61.027
Market-Years	1876	1876	1876	1876

Notes: This table displays the δ estimates of Equation 4. The market is defined as the commuting zone of the clinic based on ERS 2000 delineations. *Total Corporate Cycles* and *Total Corporate Live Births* represent the total number of IVF cycles and total number of live births performed by corporate clinics each year in a commuting zone, instrumented using the number of corporate clinics each year in a commuting zone. The first stage F-stat shows the Kleibergen-Paap Wald rk F statistics. The sample includes all clinics (including clinics always in a chain and newly opened by a chain) in a commuting zone that ever had an independent clinic. Standard errors are clustered at the market level. Significance levels: *p<0.1, **p< .05, ***p<0.01

Table G6. Effect of Private Equity Investment on Fertility Clinic Outcomes

	(1)	(2)	(3)
	Log(Cycles)	**Log(Transfers)**	**Live Birth Rate**
Panel A: Private Equity Decomposition			
Post_NoPE × Acquisition	0.106	0.117	0.051***
	(0.151)	(0.133)	(0.018)
Post_YesPE × Acquisition	0.319***	0.234**	0.052***
	(0.102)	(0.103)	(0.012)
Post_NoPE × Affiliation	0.225**	0.219**	-0.009
	(0.102)	(0.098)	(0.019)
Post_YesPE × Affiliation	0.261**	0.219*	0.021
	(0.115)	(0.118)	(0.015)
Dep. Var. Mean	5.253	5.036	0.374
Clinic-Years	5666	5666	5666
R^2	0.899	0.898	0.625
Panel B: Effect of Private Equity Funding			
Post_PE	0.189***	0.142***	0.020**
	(0.052)	(0.051)	(0.008)
Dep. Var. Mean	5.323	5.102	0.375
Clinic-Years	6134	6134	6134
R^2	0.910	0.909	0.630

Notes: This table shows the estimates of Equation 2 with treatment times based on private equity investment into a chain. In Panel A, *Post_NoPE* is a post-transaction indicator equal to 1 for the years a clinic is in a chain without PE funding and 0 for the years the clinic is in the chain with PE funding. *Post_YesPE* is a post-transaction indicator equal to 1 for the years a clinic is in the chain with PE funding and 0 for the years the clinic is in the chain without PE funding (these are mutually exclusive). In Panel B, *Post_PE* is an indicator equal to 1 when a chain receives PE funding and zero prior to PE funding. Panel B includes that full sample of clinics. The dependent variable mean captures the predicted mean for control clinics and treatment clinics before the transaction (Panel A) or before PE investment (Panel B). Standard errors are clustered at the clinic level. Significance levels: *p<0.1, **p< .05, ***p<0.01

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