

# IDENTIFYING PROVIDER PREJUDICE IN HEALTHCARE\*

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## Abstract

We use simple economic insights to assess the case for prejudicial behavior in healthcare where treatment disparities by race and gender are not explained by access, preferences, or severity. We demonstrate that treatment disparities, by themselves, cannot distinguish between two competing views of provider behavior: physicians may consciously or unconsciously underprovide treatment to minority groups despite similar benefits (prejudice), or they may use race and gender to determine treatment because they are associated with lower benefit because of physiology, behaviors, or pre-hospital factors. We demonstrate that these two views can only be distinguished using data on patient outcomes: for patients with the same propensity to be treated, prejudice implies a higher return from treatment for treated minorities, while non-prejudicial behavior implies that returns are equalized. Using data on heart attack treatments, we do not find empirical support for prejudice-based explanations. Despite receiving less treatment, women and blacks receive slightly lower benefits from treatment, perhaps due to higher stroke risk, delays in seeking care, and providers over-treating minorities due to equity and liability concerns.

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## **I. Introduction**

A large literature in social-science and medicine finds evidence of race and gender disparities in wages, hiring decisions, mortgage approvals, and medical treatments. Heckman (1998) and List (2004) have noted that the presence of disparities cannot distinguish between two competing explanations for why they exist: are employers, lenders, or physicians, consciously or unconsciously withholding jobs, loans and medical treatment from minority groups despite similar benefit (prejudice), or is membership in a minority group associated with lower productivity or benefit (non prejudicial behavior)? In this paper, we use simple economic insights to develop a general framework for using observational data to distinguish between these two explanations.

We focus our inquiry on the enormous literature on racial and gender disparities in healthcare where treatment differences are not explained by access, preferences, or severity. Disparities in medical treatment are often ascribed to prejudicial providers [Healy (1991), Bogart et al. (1994), Schulman et al. (1999), van Ryn and Fu (2003), Fincher et al. (2004) and Green et. al. (2007)]. This view is shared by the Institute of Medicine's (IOM) *Unequal Treatment* report, which concluded that provider bias was among the leading determinants of disparities (Smedley, Stith and Nelson, 2003).

We define prejudice to occur when blacks and women (henceforth referred to as minorities) receive less care than other patients who have the same expected

being used to determine benefit at the time of treatment.<sup>1</sup>

While both prejudicial and non-prejudicial behavior can result in minority groups receiving less treatment, they imply vastly different policies. The prejudicial use of race and gender in determining patient benefit requires educating physicians about patient benefits, and encouraging physicians to treat minority patients more aggressively to overcome underuse. In contrast, non-prejudicial use of a patient's race and gender would require making minority patients more appropriate for treatment and that would require reducing pre-hospital delays (and pre-hospital prejudice), improving the skills of providers who treat them, or developing race and gender specific treatments. In sharp contrast to prejudice, policies designed to eliminate disparities caused by non-prejudicial behavior would not encourage providers to close the treatment gap.

It is intuitive to distinguish between the two possibilities by comparing outcomes for the marginal patient being treated for minority and majority groups, but in the absence of observing exactly what the physician observes about the patient, it is impossible to identify the marginal patient and measure the benefit from treatment for this patient. We address this complication by proposing a new test for provider prejudice. The intuition for our test is simple: we compare outcomes two patients who have the same propensity to receive treatment. This propensity to receive treatment is allowed to depend on minority status—both because minority patients may benefit less and because there is the incorrect belief that they benefit less (prejudice). If there is prejudice, then a minority patient with the same propensity to receive treatment as a majority patient, should receive a *higher* benefit from treatment (because this patient was in effect more appropriate for the treatment than his propensity would indicate). Our framework offers a test based on observational data that is complementary to that in List (2004) who used data from field experiments to make the distinction between prejudice and statistical discrimination.

Our approach to identifying provider prejudice provides several advantages over ostensibly simpler methods. Regression adjustment, audit-studies involving patient-actors, and Implicit Association Tests (IAT) are often used to assess the case for prejudice by comparing loan approval or treatment rates by race and gender (Munnell et al. 1996, Schulman et al., 1999,

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<sup>1</sup> The literature often uses the terms prejudice, stereotyping, bias and discrimination interchangeably. We use the terms prejudice and bias to refer to provider behavior that results in allocative inefficiency. We avoid using the term stereotyping because both prejudice and statistical discrimination involve stereotyping—the key is whether physician behavior leads to greater efficiency or inefficiency in allocating treatment. We have chosen not to use the term discrimination to avoid confusion with the alternative hypothesis of statistical discrimination.

Smedley, Stith and Nelson, 2003 and Green et al., 2007). But the fact that providers or bankers may offer less to minorities is not by itself evidence of prejudice, if the benefit to these groups is in fact lower on average because of lower appropriateness stemming from biological differences in how the treatment is absorbed, or behavioral factors such as lower compliance or ambulance delays in being taken to the hospital (although it could be evidence of more macro prejudice against minorities). Yet because it is impossible to measure outcomes for simulated patients, it is not possible to distinguish between the two explanations. Similarly, it may be tempting to see how physicians of one race treat patients of their demographic versus others, as has been done in recent studies of racial bias in motor vehicle searches (Antonovics and Knight, 2005; Anwar and Fang, 2006). However, this approach can only identify whether there is differential bias across physicians of different race, and previous evidence has found no difference in the how white and black physicians treat heart attack patients (Chen et al., 2001). Moreover, since the care of minority patients is highly concentrated within a small set of physicians, studies (including audit studies) that focus on whether these physicians treat white and black patients differently exclude the substantial number of physicians who only treat patients of one race. In theory, such providers may be the most or the least prejudicial, and patients may sort into providers in way that makes this evidence very unrepresentative of the discrimination being faced by a typical patient (Heckman, 1998).

In Section II we develop the theoretical model underlying our analysis. Section III discusses the etiology of heart attacks and their principal treatment in the modern era-- reperfusion, and introduces data from the Cooperative Cardiovascular Project (CCP). In Section IV we detail our estimation strategy, paying particular attention to how the theoretical model developed in Section II will be evaluated using the CCP data. Section V presents results. We do not find empirical support for alternative explanations based on prejudice or bias. If anything, despite receiving less treatment, women and blacks receive lower benefits. Differences in follow-up care or provider skill do not appear to explain our results, suggesting that the behavior that we observe is fairly pervasive and not isolated to a handful of providers. Our favored interpretation of these results is that since few medical trials separately estimate the benefits of treatments by race or gender, medical care providers may be unaware of the extent to which the benefits of treatment are lower among women and minorities, or may be unwilling to make treatment decisions without a documented medical reason based solely on a statistical association between benefits and gender or race. A sense of equity in treatment that may be heightened by the

malpractice system may explain this behavior. Thus, the differences in treatment that we observe are not the result of prejudice by providers in the current treatment decision, even though they may be the result of discrimination more broadly (for example, in ‘pre-hospital’ factors such as ambulance response times or the fact that the treatment was primarily designed for males). These are very different conclusions from what simpler tests of prejudice have concluded.

## II. Theory

Our goal is to evaluate whether disparities in treatment rates represent prejudicial behavior by providers (implying underuse in blacks and women). A simple model of patient treatment choice will guide our empirical work. We assume that treatment is provided to each patient whenever his or her benefit from the treatment exceeds a minimal threshold. Within this framework, there are two ways in which a patient’s race or gender could affect treatment choice: race or gender could be related to the expected benefit of treatment, or could incorrectly alter physicians perception of benefit. Both effects could be at work, and separating their effects is the key empirical challenge. Of course, a physicians perception of benefit also depends on the diligence with which a patients appropriateness for treatment is assessed, and we develop a simple economic framework to measure the role of diligence in treatment decisions.

### *Treatment without Prejudice*

We first discuss treatment disparities by gender in a world without prejudice; the case for black/white disparities is analogous, with *female* being replaced by *black*. We can estimate the gain from treatment for men and women by estimating the following model of survival:

$$(1a) \quad Survival_i = \alpha_i^{male} Treatment_i + X\Pi^{male} + u_i$$

$$(1b) \quad Survival_i = \alpha_i^{female} Treatment_i + X\Pi^{female} + u_i$$

We have allowed for heterogeneous treatment effects ( $\alpha_i$ ) which allow the improvement in survival from being treated to be specific to a patient.<sup>2</sup> A patients benefit from treatment is:

$$(2) \quad \alpha_i = Benefit_i = X\Phi + Female_i\beta + \varepsilon_i$$

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<sup>2</sup> Estimation of (1) by OLS yields  $\alpha^{male} = E(\alpha_i | Treatment=1, Male)$  and  $\alpha^{female} = E(\alpha_i | Treatment=1, Female)$ , the average benefit from the treatment for patients receiving the treatment. This is true if receipt of treatment is uncorrelated with the unobservables affecting survival ( $u_i$ ), and if the unobservables affecting survival are uncorrelated with the idiosyncratic benefit from treatment ( $\varepsilon_i$ ). Since this is Roy selection, the receipt of treatment can be correlated with a patients idiosyncratic gain, which we do not observe, but is observed and used by providers (Heckman and Vytlacil, 2007).

We allow the benefit from treatment to depend on the patient's gender, observable patient characteristics ( $X$ ) such as age, medical history, and lab results (all of which can also be interacted with gender), and other factors that are known to the medical care provider when making the treatment decision but unobserved by the econometrician ( $\varepsilon$ ). At present, we assume that the variance of the  $\varepsilon$ 's conditional on the  $X$ 's do not vary by gender. But if providers spend less time ascertaining a female patient's appropriateness for treatment (i.e. are less diligent with female patients), then the variance of the  $\varepsilon$ 's will be smaller for women. We develop tests for this important assumption in the next section.

Note that gender could be statistically related to the benefit of treatment in Equation (1) because of differences in biology, pre-existing medical conditions, follow-up care, presence of family member, or quality of the provider. Similarly, factors that are correlated with race and observed by providers but unobserved to the econometrician, will be absorbed into the coefficient on gender or race. All such differences by gender and race should be taken into account by the medical provider if the goal is to maximize the benefit to the patient. In our framework, differences in treatment that arise from differences in expected benefit are not the result of prejudice in the current treatment decision, even though they may be the result of discrimination more broadly (for example, in ambulance response times or the fact that the treatment was primarily designed for males).

Each patient receives treatment if their benefit from treatment exceeds zero:

$$(3) \quad Pr(Treatment_i=1) = Pr(Benefit_i > 0) = Pr(\alpha_i > 0) \\ = Pr(X\Phi + Female\beta + \varepsilon_i > 0)$$

Thus, in the terminology of Heckman, Urzua and Vytlacil (2006), our model allows for *essential* heterogeneity where the decision to provide treatment to each patient is made with knowledge of their idiosyncratic gain from treatment. This framework follows directly from a Roy model of treatment allocation, where a patient receives treatment if the gain from the treatment exceeds zero.

In this framework, the average benefit of treatment on patients receiving the treatment is:

$$(4) \quad E(\alpha_i | Treatment=1) = E(X\Phi + Female\beta + \varepsilon | X\Phi + Female\beta + \varepsilon > 0) \\ = E(I + \varepsilon | I + \varepsilon > 0), \text{ where } I = X\Phi + Female\beta$$

The conditional mean  $E(I + \varepsilon | I + \varepsilon > 0)$  is an unknown function of  $I$  (where  $I$  is estimated from equation 3), and depends only on  $I$  if the distribution of  $\varepsilon$  does not vary by gender (which is our

maintained assumption, that we will relax in the next section). We do not assume a particular parametric structure for this truncated mean. Equation (4) states that in the absence of prejudice, a man and a woman receiving treatment who have the same propensity to get the treatment (same  $I$ ) will have the same expected benefit from the treatment.

It is important to note that Equation (4) does not imply that the treatment-on-the-treated parameter is the same for all men and women in the absence of prejudice. Rather, it implies that the treatment-on-the-treated effect will tend to be larger among men if men have a higher propensity to be treated, since  $E(I + \varepsilon | I + \varepsilon > 0)$  is increasing in  $I$ . One can see this by noting that the average benefit to men and women, across all values of  $I$  may be expressed as:

$$(5a) \quad E(\text{Benefit} | \text{Treatment}=1, \text{Male}) = \int E(\alpha_i^{\text{male}} | \text{Treatment}=1, \text{Male}, I) dF(I|\text{Male})$$

$$(5b) \quad E(\text{Benefit} | \text{Treatment}=1, \text{Female}) = \int E(\alpha_i^{\text{female}} | \text{Treatment}=1, \text{Female}, I) dF(I|\text{Female})$$

Equation (5) makes it clear that even if the treatment on the treated parameter is identical for men and women of the same  $I$ , if women have a lower distribution of  $I$  (e.g., their characteristics make them less appropriate for the treatment), this will result in finding that women have lower average benefits; that estimation of equation (1) produces  $\alpha^{\text{female}} < \alpha^{\text{male}}$ . However, if the distributions of  $I$  are equalized through reweighting for men and women (by calculating benefits for men as  $\int E(\text{Benefit} | \text{Treatment}=1, \text{Male}, I) dF(I|\text{Female})$ ), the comparison of reweighted treatment benefits will be identical for men and women; that is,  $\alpha^{\text{female}} = \alpha^{\text{male}}$  implies that physician behavior is non-prejudicial.

### *Treatment with Prejudice*

We define prejudice to exist when the treatment of two patients with the same true benefit from treatment differs because of their race or gender. This may occur because the provider incorrectly believes treatment is less effective for the minority group, or because the provider withholds treatment from the minority group despite positive benefits. In either situation, there is underuse of treatment for the minority group because treatment is being withheld from patients who would benefit from treatment.

The presence of prejudice in assessing benefit from treatment changes equation (2) to

$$(2') \quad \text{Benefit}^{\text{Perceived}} = X\Phi + \text{Female}(\beta - \tau) + \varepsilon$$

Women experience prejudice if  $\tau > 0$  in Equation (2'). We have modeled prejudice as occurring when providers incorrectly believe that women benefit less from treatment than they actually do,

but we could equivalently have modeled prejudice as providers using a higher benefit hurdle to determine whether women should be treated.

The probability of a patient receiving treatment is just the probability that a patient's perceived benefits exceed the minimum threshold:

$$\begin{aligned}
 (3') \quad Pr(Treatment=1) &= Pr(Benefit^{Perceived} > 0) \\
 &= Pr(X\Phi + Female(\beta-\tau) + \varepsilon > 0) \\
 &= F(X\Phi + Female(\beta-\tau)) = F(I^*), \text{ where } I^* = X\Phi + Female(\beta-\tau)
 \end{aligned}$$

Equation (3') highlights why differences in treatment rates by gender (or race) that are estimated by regression adjustment are not by themselves evidence of prejudice. Women may be less likely to receive treatment because of lower expected benefits of treatment ( $\beta < 0$ ) or because of prejudice ( $\tau > 0$ ), and a negative coefficient on gender combines both effects. Even if the coefficient on *Female* is zero in a regression model of treatment receipt, it could mask the fact that there is prejudice against minority patients *and* that their benefit from treatment is higher. Thus, contrary to the empirical literature, it is not possible to identify the effect of taste-based prejudice from the coefficient on *Female* in Equation (3) alone.

However, the effect of prejudice can be separately identified from the effect of women having lower benefits if information on the treatment effect among the treated population is available. Under prejudice, the treatment-on-the-treated parameter is defined as:

$$\begin{aligned}
 (4') \quad E(Benefit^{True} | Treatment=1) &= E(X\Phi + Female\beta + \varepsilon | Treatment = 1) \\
 &= E(X\Phi + Female\beta + \varepsilon | X\Phi + Female(\beta-\tau) + \varepsilon > 0) \\
 &= E(I^* + Female\tau + \varepsilon | I^* + \varepsilon > 0) \\
 &= Female\tau + E(I^* + \varepsilon | I^* + \varepsilon > 0)
 \end{aligned}$$

Equation (4') offers two insights. First, it states that we can conclude that there is prejudice against women ( $\tau > 0$ ) if the treatment-on-the-treated effect is larger for women than it is for men with the same propensity to get treatment (where the propensity to receive treatment,  $I^*$ , is allowed to be affected by prejudice). The intuition here is simple—if prejudiced providers lower a minority patients propensity to receive care, then a comparison of a minority and majority patient of the same propensity should result in the minority patient receiving higher benefits (because their true propensity was higher). Once again, to ensure that we're picking up the effect of prejudice and not gender differences in the distribution of  $I$ , the distributions of  $I$  must be equalized (by calculating benefits for men as  $\int E(Benefit | Treatment=1, Male, I)$



$dF(I|Female)$ ). Thus, the comparison of reweighted treatment benefits provides a test for prejudicial providers. This is the key empirical implication that we will test in our empirical work.

Second, this equation notes that we can identify the effect of prejudice when  $I$  is very small. When  $I$  is very small—as it will be in patients with the lowest propensity to receive treatment,  $E(I^* + \varepsilon | I^* + \varepsilon > 0)$  is zero, and any difference in benefits is the consequence of prejudice. Hence, ‘identification at infinity’ provides another way to isolate the potential for prejudice without invoking any assumptions about the distributions of  $\varepsilon$ .

Our model shares the same notion of taste-based prejudice as Knowles, Persico and Todd (2001), and Balsa and McGuire (2003). In all three papers, the decision to take an action (search a motorist for contraband or treat a patient) is determined by whether the expected benefit exceeds a threshold, and prejudice exists when the threshold (or equivalently, the difference between perceived and actual benefits) differs by race or gender. The model of racial bias in motor vehicle searches developed in Knowles et al. is conceptually similar to our model of prejudice, but differs in one important respect, which makes our empirical test of prejudice more complicated. In the Knowles et al. model, the returns to searching motorists in the absence of prejudice are equalized across all subgroups in equilibrium – otherwise, police would always search motorists with higher returns and motorists would react by carrying less contraband. The key difference in our setting is that patients are not choosing any action analogous to carrying contraband, so there will be some subgroups of patients (those with high propensity to be treated) who continue to have higher returns in equilibrium even in the absence of prejudice. By contrast, in Knowles et al. because of an equilibrium condition that relies on motorists responding to the threat of being searched, there is no distinction between the marginal and treatment on treated effects in the absence of prejudice. Thus, in the absence of prejudice, the Knowles et al. model implies that the returns to search are identical across subgroups unconditionally, while our model implies that the returns to treatment are identical across subgroups conditional only on the propensity to receive treatment ( $I$ ).

Our model differs from Balsa and McGuire (2003) in how we define non-prejudicial behavior. In Balsa and McGuire, non-prejudicial behavior occurs when providers use different decision rules to determine a patient’s appropriateness for care; there is no outcomes based test. In our framework, finding evidence of these triage-rule differences is not informative about prejudice because it may be correct for physicians to use different triage rules, including using

information on race and gender to determine appropriateness. Our approach is analogous to how statistical discrimination is modeled by labor economists (Altonji and Pierret, 2001).

### Graphical Intuition

The graphical intuition for our model can be seen in Figure 1a for the case of no prejudice, and in Figure 1b for the case of prejudice against women. The expected benefit from treatment is given on the vertical axis, while the index  $I$  (which determines the propensity of being treated) is given on the horizontal axis. There is a distribution of unobservable characteristics that is centered on the expected benefits curve, but patients receive treatment only if their perceived benefit exceeds zero (the minimum threshold). The thick curve in Figure 1a represents the treatment-on-the-treated effect for a patient with index  $I$ , that is it gives  $E(\textit{Benefit} \mid \textit{Benefit} > 0)$ . Treatment-on-the-treated approaches the minimum threshold of zero benefit for a patient with a low propensity of being treated (a very negative  $I$ ), since no patient is ever treated with a benefit below this threshold. For a patient with a high propensity of being treated (a very positive  $I$ ), truncation becomes irrelevant and the treatment-on-the-treated effect asymptotes to the unconditional benefit of treatment.

When there is no prejudice, as in Figure 1a, if men have a higher general propensity to receive treatment (the distribution of  $I_{Male}$  is to the right of the distribution of  $I_{Female}$ ), then they will also have greater benefits from treatment – a movement along the curve in Figure 1a. But in the absence of prejudice, any two people with the same propensity will have identical treatment effects if treated. This underscores the need to equate the distributions of  $I_{Male}$  and  $I_{Female}$ , before comparing the average treatment on the treated for men and women.

Figure 1b shows how treatment-on-the-treated differs for men and women when there is prejudice against women ( $\tau > 0$ ). Because physicians underestimate the benefits from treatment for women, a woman with the same perceived benefit as a man will have a higher true benefit from treatment. Consequently, when two people of the same  $I$  are compared, the treatment effect among women is higher. Moreover, the true benefit of treatment among those being treated will always be greater than  $\tau$  (since  $\textit{Benefit} - \tau > 0$ ), even among individuals with low propensity of treatment. Thus, even after reweighting to equate the distributions of  $I_{Male}$  and  $I_{Female}$ , the treatment-on-treated effect will be higher for women than for men whenever there is prejudice against women.

## Evaluating the Single Index Assumption

Our test relies on the assumption that in equation (4') the term  $E(I^* + \varepsilon | I^* + \varepsilon > 0)$  depends only on the index, i.e. we must assume a single-index selectivity model, so that the truncated mean of the error in equation (4') depends only on  $I^*$ . This would not be the case if providers spend less time ascertaining a female patient's appropriateness for treatment. If providers are less diligent with female patients, then the variance of the unobservable factors determining treatment ( $\sigma_\varepsilon$ ) will be smaller for women. A smaller error variance for women would imply a lower truncated mean – leading to lower expected benefits from treatment for women. This would look like prejudice against men (even though it is the opposite!) so it is important to rule out.

To allow for different variance in unobservable factors determining treatment by groups, let  $\varepsilon_g = \varepsilon \sigma_g$ , where  $\sigma_g$  is a scale factor determining the variance in unobservable factors for each group (e.g.  $\sigma_{female} < \sigma_{male}$  would correspond to less variance for women in unobservable factors determining treatment). With this modification, Equations 2-4 would be rewritten as:

$$(2'') \quad Benefit^{Perceived} = X\Phi + Female(\beta - \tau) + \varepsilon \sigma_g$$

$$(3'') \quad Pr(Treatment=1) = Pr(Benefit^{Perceived} > 0) \\ = Pr(I^{**} + \varepsilon > 0), \text{ where } I^{**} = I^*/\sigma_g = X\Phi/\sigma_g + Female(\beta - \tau)/\sigma_g$$

$$(4'') \quad E(Benefit^{True} | Treatment=1) = Female \tau + \sigma_g E(I^{**} + \varepsilon | I^{**} + \varepsilon > 0)$$

Even if females differ from males in having a different variance of errors, estimation of equations (3'') and (4'') still preserve all the necessary information for our test, and further provide a number of tests for whether the variance of the errors differ across groups. The first test for differences in the variance of unobservables follows naturally from the fact that the probability of treatment depends on  $I^{**} = I^*/\sigma_g$ , implying that we would expect to observe a stronger relationship (larger coefficients) between observable patient characteristics and the propensity to be treated for any group for which unobservable factors played a smaller role in treatment ( $\sigma_g$  smaller). However, this is at best an indirect test since it relies on the assumption that the true  $\Phi$  is the same across groups. For example, one might believe that inattention lowers the effect of both observables and unobservables on treatment for minorities, so that  $\Phi/\sigma_g$  remains unchanged.

A more powerful test for differences in the variance of the unobservables comes from the relationship between the treatment propensity ( $I^{**}$ ) and the treatment on the treated effect in

Equation 4'''. As was the case when we assumed that all groups had the same variance of unobservables (Equation 4'), we can still identify the effect of prejudice when  $I$  is very small – since  $E(I^{**} + \varepsilon | I^{**} + \varepsilon > 0)$  is zero for patients with the lowest propensity to receive treatment, and any difference in benefits is the consequence of prejudice ( $\tau$ ). However, Equation 4''' states that the treatment-on-treated effect will rise less quickly with propensity to receive treatment when the variance in unobservables ( $\sigma_g$ ) is smaller (the derivative  $\delta E(\text{Benefit}^{True} | \text{Treatment}=1) / \delta I^{**}$  is increasing in  $\sigma_g$ ). The intuition is that the lower variance results in a lower mean for the truncated distribution of benefits, and for any given proportion of the distribution lying above the treatment threshold, the benefit for the patients being treated will lie closer to the threshold.

In summary, if there is less diligence in the treatment of minority patients, we would see smaller variance in the unobservable factors used to determine treatment for minority groups. This implies that we would expect to see (1) larger coefficients on the observable characteristics determining treatment (2) smaller benefits from treatment and (3) less of a relationship between benefits and treatment propensity among minority groups. While we will maintain the equal diligence assumption (statistically equivalent to assuming single index) for our primary analyses, we will also test its validity empirically in Section V.

### **III. Heart-Attacks: Biology, Treatments, and Data**

#### ***Heart-Attack Biology and Treatments***

Heart attacks (more precisely, acute myocardial infarction (AMI)) occur when the heart-muscle (the myocardium) does not receive sufficient oxygen, because of a blockage in one of the coronary arteries which supply blood to the heart. The blockage is typically caused by a blood clot that occurs because of coagulation induced by the rupture of atherosclerotic plaque inside the coronary arteries. Angioplasty (where a balloon on a catheter is inflated inside the blocked coronary artery to restore blood flow) and timely thrombolytics (are administered intravenously and break down blood clots by pharmacological means) are two treatments that are used for immediate reperfusion (opening up the coronary artery). Following the clinical literature, we define a patient to have received reperfusion if any of these therapies was provided within 12 hours of the heart attack (in our data from the mid-1990s, over 90 percent of patients receiving reperfusion received thrombolytics). The decision to reperfuse a patient reflects decision making by a team of people: the triage nurse, emergency room physician, and the cardiologist. We view

this team as the key decision makers in our model; their skill and level of coordination will vary greatly from one hospital to another.

We focus our empirical work on the treatment of AMI for a number of reasons. First, cardiovascular disease, of which heart attacks are the primary manifestation, is the leading cause of death in the US. A perusal of the leading medical journals would indicate that heart attack treatments are constantly being refined, and a large body of trial evidence points to significant therapeutic gains from many of these treatments. In this context, racial and gender disparities in treatments may directly translate into lost lives, and there is a rich tradition of studying racial and gender disparities in treatments and outcomes after heart attacks (Allison et al. (1996), Canto et al. (2000), Barnato et al. (2005), Vaccarino et al. (2005), Rathore et al. (2000), Skinner et al. (2005), Jha et al. (2007)). One advantage of studying heart attacks in the Medicare population is that black/white differences in insurance (at least for in-hospital care) aren't a source of confounding.

Second, as a consequence of what is known about heart attack treatments from randomized controlled trials, and more specifically for our setting, the benefits from reperfusion, we are able to assess whether our regression estimates of the benefits from reperfusion are comparable to those found in the medical literature, or whether they are confounded by selection-bias. We focus on reperfusion, where our use of chart data allows us to replicate the RCT evidence that is summarized by the Fibrinolytic Therapy Trialists' Collaborative Group (1994).

Third, because mortality post-AMI is high (survival rates at one year are less than 70 percent), a well-defined endpoint is available to test the efficacy of heart attack treatments. This would not be true if we focused on treatment disparities for more chronic conditions such as diabetes, chronic obstructive pulmonary disease, or arthritis.

Our fourth reason for focusing on heart attacks is that it is an acute condition for which virtually all patients are hospitalized and receive some medical care, thereby allowing us to evaluate the case for provider prejudice comprehensively. "911" operators and emergency medical teams are especially trained to recognize the symptoms of heart attacks. This would be less true of chronic diseases that progress gradually. Nor do we believe that patient preferences matter as much for treating heart attacks—at least during the acute phase of the heart attack the therapeutic emphasis is on maximizing survival, which is achieved by timely reperfusion, and hospital staff (not patients and their families) make treatment decisions. While providers may

specialize in the use of surgical or medical management of heart attacks, as in Chandra and Staiger (2007), the fact that patients are generally taken to the nearest hospital for treatment, renders the nature of treatment received as exogenous to the patient preferences. This feature of heart attack treatments would not be true of cancer therapies where two clinically identical patients may chose different therapies based on their idiosyncratic valuation of side-effects and treatment duration.

### ***Data***

Because acute myocardial infarction is both common and serious, it has been the topic of intense scientific and clinical interest. One effort to incorporate evidence-based practice guidelines into the care of heart attack patients, begun in 1992, is the Health Care Financing Administration's Health Care Quality Improvement Initiative Cooperative Cardiovascular Project (CCP). Information about more than 200,000 patients admitted to hospitals for treatment of heart attacks in 1994/1995 was obtained from clinical records. The CCP is considerably superior to administrative/claims data (of the type used by McClellan et al. (1994)) as it collects chart data on the patients—detailed information is provided on laboratory tests, enzyme levels, the location of the myocardial infarction, and the condition of the patient at the time of admission. Detailed clinical data were abstracted from each patient's chart using a standard protocol. Further details about the CCP data are available in Marciniak et al. (1998), O'Connor et al. (1999), and in the appendix to this paper. The choice of sample and variables is identical to what we used and described in Barnato et al. (2005) and Chandra and Staiger (2007).

## **IV. Estimation**

### **A. Basic Specifications**

We use data on heart attack treatments to estimate the key components of our model, using receipt of reperfusion within 12 hours of the initial heart attack as our treatment. The propensity to receive treatment ( $I$  in the theoretical model) is estimated by obtaining the index from a probit model that regresses whether a patient received reperfusion within 12 hours of the heart attack on gender, race, and all the CCP risk-adjusters ( $X$ ):

$$(7) \ Pr(\text{Reperfusion}) = F(\text{Female} * \text{Black} * X\beta)$$

In equation (7), the effect of race and gender on the probability of receiving reperfusion is allowed to vary across every comorbidity (direct effects for female and black are also included).

Differences in the coefficients by race or gender may reflect ignorance, prejudice, or knowledge about the benefits of treatment (for example, the effect of diabetes or age may operate differently in blacks).<sup>3</sup>

Our test for prejudice is to determine, among patients with the same propensity to be treated, whether the survival benefit to reperfusion is greater for women relative to men, or blacks relative to whites. Survival is measured as a binary variable that measures survival to a certain date (e.g. survival to 7 days, or survival to 1 year). This suggests estimating models of the following type for women (there will be an identical equation for the black-white disparity, where female is replaced with black).

$$(8) \quad Survival_i = \alpha_0 + \alpha_1 Reperfusion + \alpha_2 (Reperfusion * Female) + X_i + e$$

In equation 8,  $\alpha_1$  is the survival gain from reperfusion for men,  $\alpha_2$  is the differential benefit for women ( $\alpha_1 = E(Benefit_{men} | Treatment=1)$ ;  $\alpha_2 = E(Benefit_{men} - Benefit_{females} | Treatment=1)$ ), and X includes full gender-race interactions (including the main effect of race and gender) with the CCP risk-adjusters. We interpret  $\alpha_2$  as the difference in the average treatment on the treated effects for men and women. However, as we noted in the theory section,  $\alpha_2$  is estimated over different distributions of the propensity to receive reperfusion for men and women, and is therefore not a precise test of our model. In other words, evidence that  $\alpha_2 < 0$  is not sufficient evidence to conclude that there is no prejudice against women, if women are generally less appropriate for treatment and consequently, have lower treatment propensities. In the absence of prejudice, our model states that the treatment on treated (TT) effect is the same only for men and women with the *same propensity* to be treated, or equivalently for any population of men and women that have the same propensity distributions (e.g. are matched on propensity to be treated).

Our estimation strategy relies on using the richness of the CCP data to invoke a 'selection on observables' assumption to estimate treatment on the treated for reperfusion therapy. To ascertain the credibility of this assumption we can compare the estimates in Table 2 to those obtained from clinical trials to evaluate the plausibility this assumption. A summary of nine trials was published in the journal *Lancet* by the Fibrinolytic Therapy Trialists' Collaborative Group (FTTCG, 1994). This was the same time-period as the CCP data and each trial evaluated

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<sup>3</sup> As we note in Section V, we find little support for these concerns in the empirical work: the predicted propensity to receive reperfusion from the fully interacted model was correlated 0.997 with the propensity from the model with no race or gender interactions.

fibrinolytic therapy in heart-attack patients. Across these nine trials, reperfusion within 12 hours reduces 35-day mortality by 1.9 percentage points (an 18 percent reduction in mortality). In the second column of Table 3 we estimate the effect of reperfusion (pooling across men and women) on 30-day survival as 1.9 percentage points (s.e.=0.2 percentage points). We focus on the second column because randomization balances the distribution of characteristics between the treatment and control arms of the trial.<sup>4</sup> We take this evidence as supporting the case for simpler regression models that invoke a ‘selection on observables’ assumption.

To test whether the treatment-on-treated effect is the same for men and women using the framework of equation (8) it is key that this parameter be estimated for men and women with similar propensities to treated women. Therefore, we reweight the above equations at the time of estimation two different strategies. First, we use the method of Barsky et al. (2002) to make the propensity distribution of both treated and untreated males (or whites) look like the propensity distribution of treated females (or blacks), and also reweight untreated female (or black) propensities to look like the treated female (or black) distribution of propensities. In other words, the treatment effect for men and women (or whites and blacks) is estimated for a population with the propensity distribution observed among the treated in the female (or black) population. We compute the weights by calculating the value of the 100 percentiles of the propensity distribution among treated females. By construction 1 percent of treated women are in each of these percentiles. Suppose that  $m_p$  percent of one of the other three groups (treated or untreated men, or untreated women) are in the  $p$ th percentile of the treated female distribution of propensity. For such patients we will assign them a weight of  $1/m_p$  at the time of estimation. We provide evidence of the success of this strategy in the results section.

An alternative to the reweighting strategy, which examines whether the treatment-on-treated effect is the same for men and women (or whites and black) with the same propensity to be treated, is to allow the treatment effect in equation 8 to depend directly each individual’s propensity to be treated. We estimate  $E(I^* + \varepsilon | I^* + \varepsilon > 0)$  non-parametrically since it’s a function of the propensity to receive treatment ( $I$ ) with 100 indicator variables for the percentiles of  $I$  interacted with reperfusion (and included directly in  $X$ ), thus allowing the return to

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<sup>4</sup> Standard propensity score weighting to make the distribution of propensities for the untreated resemble that of the treated (that is, weighting the untreated by  $\text{Pr}(\text{Reperfusion})/1-\text{Pr}(\text{Reperfusion})$ ), estimates the effect of reperfusion as 2.1 percentage points which is very similar to the trial estimate of 1.9 points.



reperfusion to vary flexibly with a patient's propensity to receive reperfusion. This requires us to estimate:

$$(9) \text{ Survival}_i = \alpha_0 + \alpha_1(\text{Reperfusion}_i * \text{Percentile of I}) + \alpha_2(\text{Reperfusion}_i * \text{Percentile of I} * \text{Female}) + \text{XII} + e$$

In equation 9,  $\alpha_1$  measures the survival gain from reperfusion for men at different percentiles of I, and  $\alpha_2$  is the differential benefit for women. We can also recover the *Reperfusion\*Percentile* effects to examine whether the benefits of reperfusion are increasing in the index (as they should if the Roy-model framework underlying our theoretical model is a correct assumption.) Thus, this specification allows the return to reperfusion to depend non-parametrically on the index

CCP risk-adjusters closes this gap for women but not for blacks. However, as noted in the theory section, the probit coefficients on race and sex capture both differences in the benefit from treatment, as well as potential differences in prejudice for these groups. As such, these negative coefficients do not provide us with a test of prejudice. Finally, in the last column of Table 2 we add a Female\*Black interaction. This interaction, which is insignificant, demonstrates that the racial disparity in treatment does not vary by gender.

In Table 3 we present our key results from the estimation of equation 8 for differences by gender (Panel A) and race (Panel B). The coefficient on Reperfusion reports the survival benefit of Reperfusion for men (or whites in Panel B), and the interaction effect with sex or race reports the differential benefit for women (and blacks in Panel B). Focusing on the unweighted results in Table 3, we see that Reperfusion improves the probability of surviving 30 days for men by 3.4 percentage points. But for women the effect is 1.4 percentage points less. If there were underuse in women, we would have expected the opposite – higher returns to reperfusion for women. The unweighted results by race are similar. The effect of reperfusion is 2.6 percentage points less for blacks. Thus, the unweighted results find significantly lower effects of reperfusion for both blacks and women, suggesting that if anything there is overuse among blacks and women relative to whites and men – that is, the lower returns for blacks and women suggest that they face a lower hurdle in the decision to treat.

However, as noted by our theory, this is not a sufficient test for prejudice because the treatment-on-the-treated effect will tend to be larger among men because men have a higher propensity to be treated, since  $E(I^* + \varepsilon | I^* + \varepsilon > 0)$  is increasing in  $I$ . In column 2, we use the method of Barsky et al. (2002) to reweight the distributions of propensities in untreated females, treated males, and untreated females to resemble that of treated females. Panel B performs the analogous weighting for blacks. As expected, the estimated treatment effect for males and whites (the direct effect of reperfusion) declines after reweighting, because we are now estimating the treatment effect in a (reweighted) population with a lower propensity to receive treatment. Correspondingly, the estimated differences in the treatment effect between men and women or whites and blacks (the interaction of reperfusion with female or black) is smaller, and we are no longer able to reject the null hypothesis that the effect of reperfusion for patients receiving reperfusion is the same by gender and race. These weighted results are the direct test of the theoretical model, and find no evidence of prejudice. Prejudice and underuse would imply higher returns to treatment for females and blacks, but our point estimates suggest the opposite. Even

the upper bound of the 95% confidence interval on these estimates would suggest very little prejudice against females or blacks.

In Figure 2, we examine the success of our weighting strategy to equalize the distribution of propensities by gender (Panel A) and by race (Panel B). The unweighted and reweighted propensity distributions are plotted by gender and race for the untreated (left hand side) and for the treated (right hand side). For reference, we have also included the propensity distribution for treated females (or blacks), since this is the target distribution that is being matched with reweighting. In the unweighted distributions, it is apparent that whites and men have higher propensities to receive treatment, particularly among patients receiving treatment. Therefore, our model would predict that unweighted estimates of the treatment effect will tend to be larger for whites and men, even in the absence of any prejudice. In contrast, the re-weighted distributions (where greater weight is put on the observations that are similar to those of the treated minority group) look identical to those for treated women and blacks.

An alternative to rebalancing the distribution of propensities is to control for  $E(I^* + \varepsilon | I^* + \varepsilon > \theta)$  directly. In column (3) of Table 3 we include 100 indicator variables for the percentiles of  $I$  as a non-parametric way to control for appropriateness. We also included interactions of these indicators with Reperfusion to test whether the benefits of reperfusion increase with the index (this is why we cannot report the direct effect of reperfusion), as they should if providers are ranking patients correctly and then working down the distribution of appropriateness. The results of non-parametric specification column (3) is identical to those obtained from propensity score reweighting suggesting that concerns about imperfect reweighting aren't a source of bias. Moreover, in Figure 3, we plot the estimated benefit from reperfusion at every percentile of  $I$ , using local linear regression to smooth the effect across percentiles. The benefit from reperfusion is increasing with  $I$ , as is required by our model and demonstrates that physicians are able to rank patients so that patients who benefit most from reperfusion are most likely to receive the treatment. There is evidence of 'harm' for patients with very low propensities of being treated (suggesting that the minimum hurdle to determine receipt of treatment is being set at a level where survival is reduced by the treatment), but there are very few patients getting treatment in this range of propensities.

## **VI. Understanding Differences in Benefits**

The above results may be summarized as finding no evidence of higher returns from treatment for women and blacks of similar appropriateness. If anything, we have consistent but statistically insignificant evidence that the benefits for these groups is lower than that for men and blacks. Before concluding that this evidence represents evidence of no-prejudice, we have to rule out a number of other explanations. Some of these reasons may actually reflect prejudice and it is important to rule them out prior to concluding that we have found support of non-prejudicial behavior.

#### *A. Differences in Triage Rule and Diligence*

First we consider the idea that physicians are incorrectly using different triage rules to rank patients (by sex or race) for treatment. If this were true, the reason for the slightly lower benefits for women and blacks could be that less appropriate patients from these populations were getting the treatment because physicians spend less time and effort in doing the triage for these groups. These pressures may be exacerbated by the demands of working in a high-pressure, time-sensitive environment where the potential for subconscious biases such as ‘implicit discrimination’ is high (Bertrand, Chugh and Mullainathan, 2005). In such an environment providers may be biased in how patients are ranked as appropriate for care – leading to both fewer women and blacks receiving reperfusion and less benefit.<sup>5</sup> We can evaluate this concern directly by estimating separate probit models of receiving reperfusion by race and gender (e.g. allowing risk factors to have different impacts on propensity to treat by gender and race), and then examining whether the predicted values from this more flexible model is similar to a model in which race and gender enter only directly (as would be the case if they only effected the threshold). The predicted values are very similar, suggesting that physicians are not using different triage rules for women and blacks: in the full sample, the correlation between the fitted values is 0.996. For women it is 0.993 and for males it is 0.9958. Similarly, the correlations are 0.949 for blacks and 0.998 for whites.<sup>6</sup>

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<sup>5</sup> More malevolent versions of this story would involve providers offering treatment to the most appropriate white patients and randomly offering it to blacks, or, in the extreme, withholding it to the most appropriate black patients.

<sup>6</sup> In theory, it may be the case that on average, the common-effects model produces estimates of receiving reperfusion that are similar to those from the race-specific model, but there are significant departures in the tails of the appropriateness distribution. In Jha et al. (2007), we explore this possibility for a number of heart attack treatments, and note that the two models give very similar results across the entire distribution of appropriateness.

The similarity of predictions where the effects of comorbidities are allowed to vary by race and gender to those where they are not, supports the view that provider diligence is similar across minority and majority groups (diligence is measured by higher variance of the  $\epsilon$ 's). As we noted in Section IV, probit coefficients identify  $\beta/\sigma$  so models estimated separately by gender and race will produce larger coefficients for the group with the smaller variances, i.e. the group for whom providers spend less time ascertaining appropriateness. Under the assumption that men and women and blacks and whites have the same weights on the  $X$ 's, we can therefore conclude that diligence is the same between these groups. However, this would be a premature assessment if providers are more likely to underweight all information, both the parts that are observed to us and the parts that are not, at the time of triaging minority patients. In this case, *both* the betas and the sigmas would scale down and the predicted effects across models with and without race and gender specific effects would be similar. But, as discussed in Section IV, if the variance of unobservables were smaller in minority groups, then the slope of curve relating treatment benefits to the propensity of receiving treatment would also be smaller in minority groups. Figure 4 demonstrates that this is not the case— it explores this possibility graphically by reporting the estimated benefit from reperfusion at every propensity to receive reperfusion (it is estimated as in Figure 3 but done separately by gender and race). For blacks and whites, the slope of the line relating benefits to propensities is identical. Women have a steeper slope than men, suggesting that there is no evidence of lower diligence for women.

The similarity of rankings from the two models also helps us rule out other forms of triage-bias. If providers were maximizing different dimensions of benefit in different populations (e.g. maximizing survival in blacks but survival and quality of life in whites) then the coefficients from the separate probits would be different and result in weakly correlated predictions.

### *B. Differences in Knowing Who Benefits From Treatment*

An alternative explanation for the fact that minorities and women receive lower returns is that providers are unaware of how to rank patients: in other words, even though the previous section demonstrates that providers use the same decision rule to rank patients, they should not. This form of bias would occur if medical textbooks and clinical trials are biased towards studying the etiology of disease in whites and men, and physicians assume that this knowledge applies equally to women and blacks. In this case, the return to reperfusion for women or blacks

would be lower on average, and again would depend less on their propensity to receive care (and could even cross, e.g., with low propensity women benefiting more from treatment than low propensity men). However, as was just discussed in the previous section, the benefit for blacks and women does not depend less on their propensity – and for women appears to depend more on the propensity to receive reperfusion. The evidence in Figure 4 demonstrates that if anything the benefits to treatment are lower for female and blacks at all propensities (except the very highest for females). Moreover, while estimates for blacks and whites are roughly parallel, the relationship between propensity and the effect of reperfusion is steeper for females than for males – the opposite of what we would expect if physicians were using the wrong decision rule to treat women.

### *C. Differences in Follow-up Care*

Next we investigate the concern that reperfusion provides similar short-run benefits to blacks and women, but that a lack of follow-up care leads to worse outcomes after the patient is discharged from the hospital (usually within a week of the AMI). If this were the case, we would expect the effect of treatment on short-term mortality to be similar, while the effect of treatment on long-term mortality would diverge. We explore the time-path of survival in Table 4 and Figure 5, where we estimate separate models for survival at horizons of 3 days through 1 year. We find that for women, receiving reperfusion is actually associated with harm during the acute phase of treatment (the first 3 days). The clinical literature notes that strokes are a possible side-effect of reperfusion, and while we cannot directly verify that women are having strokes, the early mortality at 3 days is consistent with this explanation. After this initial phase, there is no further deterioration of benefits in women; lower benefits for women are largely the consequence of higher acute mortality associated with reperfusion. Thus, women appear to face greater acute risk from the procedure, but then experience similar survival benefits in the post-acute phase.

Blacks also experience somewhat greater mortality risk from the treatment in the acute phase, although the difference from whites at 3 days is not significant. In contrast to the result for women, however, blacks in our sample appear to receive lower (but not harmful) benefit in the post-acute phase—their estimated benefits of treatment remain small even at one year, and the gap in the treatment effect between whites and blacks continues to grow through 90 days. Thus, it appears that (like women) reperfusion may harm blacks more during the acute phase of treatment, but also (unlike women) blacks may benefit less from the treatment after the acute

phase. One possible reason for the lack of post-acute benefit of reperfusion among blacks could be a lack of follow-up care. But another possible explanation, which we discuss further in the conclusion, is that blacks are known to take longer to arrive at the hospital because of ambulance delays and other factors, and trial evidence suggests that the effectiveness of reperfusion declines sharply with time since the heart attack.

#### *D. Differences in Hospital Skill*

One potential explanation for the lower benefits for blacks is that blacks and whites go to different hospitals, and that hospitals which treat blacks are not good at the management of heart attacks [Chandra and Skinner (2004) and Skinner et al. (2005)]. To explore this theory, we modified equation 8 to include hospital fixed effects as well as hospital fixed effects interacted with whether a patient received reperfusion. With these fixed effects we allow hospitals to vary in quality and also in their reperfusion specific expertise. The estimates are reported in Table 4. Including these fixed-effects resulted in nearly identical estimates to the results without fixed-effects reported in Table 4, and demonstrates that the fact that women and minorities receive slightly lower returns is a within hospital phenomena. This, in turn, means that it is a fairly pervasive phenomena and is not isolated to a few hospitals.

#### *E. Differences in Cost of Treatment*

Finally, we explored the possibility that the benefit from treatment, and therefore the decision to treat, may also depend on the costs of treatment. Thinking about the costs as well as the health benefits of treatment is useful for two reasons. First, differences in the cost of treatment by race or gender may offset the survival differences, e.g. the larger impact of reperfusion on survival for men may be offset by higher costs of doing the procedure in men. Second, prejudice could appear in a more subtle form if medical care providers placed a smaller weight on costs in the decision to treat whites and men – implicitly placing a higher value on their life. Empirically, we found that the story about costs did not affect the results – because the effect of reperfusion on cost was similar for men and women and for whites and blacks.<sup>7</sup>

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<sup>7</sup> To perform this test, we redefined benefit as the benefit net of costs from treatment:  $B - \lambda C$ , with  $S$  representing the survival benefit from treatment, and  $C$  the cost for a patient.  $\lambda$  is a measure of survival per dollar tradeoff (when  $\lambda=0$ , the medical care provider focuses solely on survival benefits). A minimum value for a life year commonly used in cost-effectiveness studies would be about \$20k per life year, which implies a value of  $\lambda=0.01$ .

## VII. Conclusions

We used a simple economic intuition to develop a framework that provides a way to ascertain the role of prejudice, including less diligence, in determining treatments for minority patients. Our framework relies on the Beckerian intuition that prejudice can be ascertained by comparing outcomes for the marginal patient (a test that cannot be implemented without observing everything that providers observe), but we recast this intuition by combining information on the propensity to receive treatment with estimates of the benefits from treatment for those patients who receive the treatment. This outcome-based test for prejudice is in contrast to much of the empirical literature on prejudice.

In contrast to what models of prejudice would predict, we find that providers appear to know of the lower benefits that accrue to women and blacks and consequently use gender and race to correctly determine patient appropriateness for care. Even the upper bound of our estimates does not support the case for prejudice against women and blacks. In fact, if anything, we find that women and blacks have slightly lower benefits relative to a model of non-prejudicial behavior. There are a number of reasons that we might observe such a pattern. Medical care providers may be aware of the lower benefits of treatment in minorities, but choose to increase treatment rates for these groups (or lower treatment for other groups) because of perceived equity concerns. This behavior may be reinforced by malpractice concerns if providers worry that their decision not to treat a patient based on their race or gender could expose them to litigation. Alternatively, since few medical trials separately estimate the benefits of treatments by race or gender, medical care providers may be unaware of the extent to which the benefits of treatment are lower among women and minorities, or may be unwilling to make treatment decisions without a documented medical reason based solely on a statistical association between benefits and gender or race. In fact, the absence of such trials or medical knowledge may be a form of macro-prejudice against minorities (Healy, 1991), but this is a different explanation than bias in the clinical encounter.

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For reasonable values of  $\lambda$ , between 0 and .1, we were unable to reject the hypothesis that the returns to men and whites were similar to those for women and blacks. These results are available upon request.



Other mechanisms for the slightly lower benefits for women and blacks are less benign. For example, the returns to any treatment might be low unless other treatments are also being provided (for example, a provider may be less likely to provide reperfusion to black patients knowing that they are less likely to receive other complimentary therapies such as high-quality diabetes care or follow-up care after discharge from the hospital), leading to an equilibrium in which no treatments are ever provided to minorities. While this remains a possibility, in earlier work we noted that racial disparities in care within hospitals and within areas are not correlated (see Baicker et al., 2004). For example, the racial disparity in the receipt of high-quality care was correlated less than 0.10 with the disparity in colorectal cancer screening or cardiac catheterization. The lack of correlation in disparities would suggest that explanations grounded in self-fulfilling equilibria are not first-order. It is also possible that lower levels of provider effort cause lower benefits in minorities. But we did not find evidence that providers triage patients into care differently, as would be the case if they used all the information at the time of admission for men and whites, but not for women and blacks.

In future work it will be important to document the precise biological mechanism that leads to the slightly lower returns to treatment for female and black heart attack patients. For women, we interpret the risk of early mortality from reperfusion as evidence that they are at higher risk of having a stroke—and while the presence of stroke-risk is consistent with the trial results (see FTTCG, 1994) this would need to be established more rigorously. Such a mechanism suggests that alternative treatments must be developed for women that reduce the probability of early strokes. For blacks, lower long-term benefits from reperfusion may reflect delays in obtaining treatment. Sheifer et al. (2000) and Ting et al. (2008) demonstrate that it takes blacks longer to arrive at the hospital because of ambulance delays or because of delays in recognizing the symptoms of a heart attack. In our data 54 percent of whites arrived within 6 hours of the start of reported chest pain relative to 43 percent of blacks, and blacks were more likely to be coded as having no chest pain (22 percent versus 15 percent) or having unknown time since chest-pain at the time of arrival (9 percent versus 13 percent). Because providers do not know the exact status of the delay confronting minority patients they may treat black heart attacks identically to white AMIs, but doing so could result in lower benefits for black patients if they are less appropriate for reperfusion by virtue of their delayed arrival. Efforts to reduce late arrivals for minority patients are very different from those that emphasize the prejudice channel.

While we reject the role of prejudice in explaining treatment disparities for heart attacks, we are cautious about generalizing these conclusions to treatment disparities across all medical conditions. It is certainly possible that prejudice may be more pertinent in an ambulatory setting where patient-doctor communication is more relevant for treatment decisions. Alternatively, prejudice may be less relevant in ambulatory settings where the physician has more time to make a decision, which helps to reduce the scope for implicit bias, which is most prevalent in a time-sensitive environment (Bertrand, Chugh and Mullainathan, 2005). Regardless of the specific context and explanation, our paper provides a general economic framework to evaluate the role of a variety of bias-based explanations for disparities in care, versus less pernicious alternatives. In theory, this framework can be used to assess the role of prejudice in other settings such as hiring, loan applications, and admissions decisions. In all of these settings, disparities in treatment rates alone do not provide evidence of prejudice, but can when combined with estimates of the benefits of the intervention.

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Figure 1: Illustration of Unprejudiced Provider Behavior (Panel A)

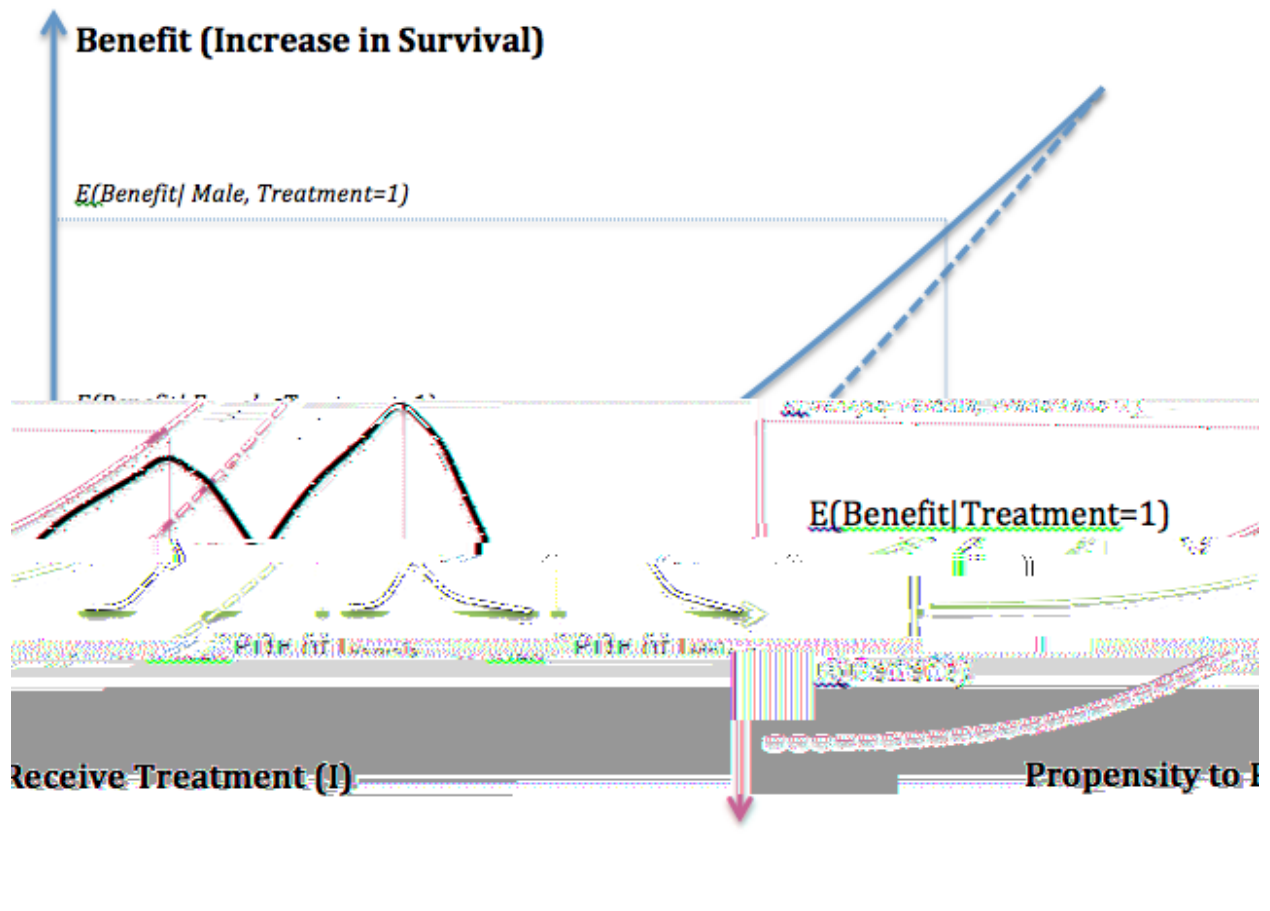


Figure illustrates the relationship between the expected benefit from treatment for men and women on the vertical axis, and the index  $I$  (which determines the propensity of being treated) on the horizontal axis. The thick curve represents the treatment-on-the-treated effect for a patient with index  $I$ . It approaches a minimum benefit of zero for a patient with a low propensity of being treated. The dashed line is the average treatment effect for patients with an index  $I$ . The two distributions  $f(I_{\text{male}})$  and  $f(I_{\text{female}})$  illustrate the difference in propensities to be treated between men and women.

Figure 1b: Illustration of Prejudice

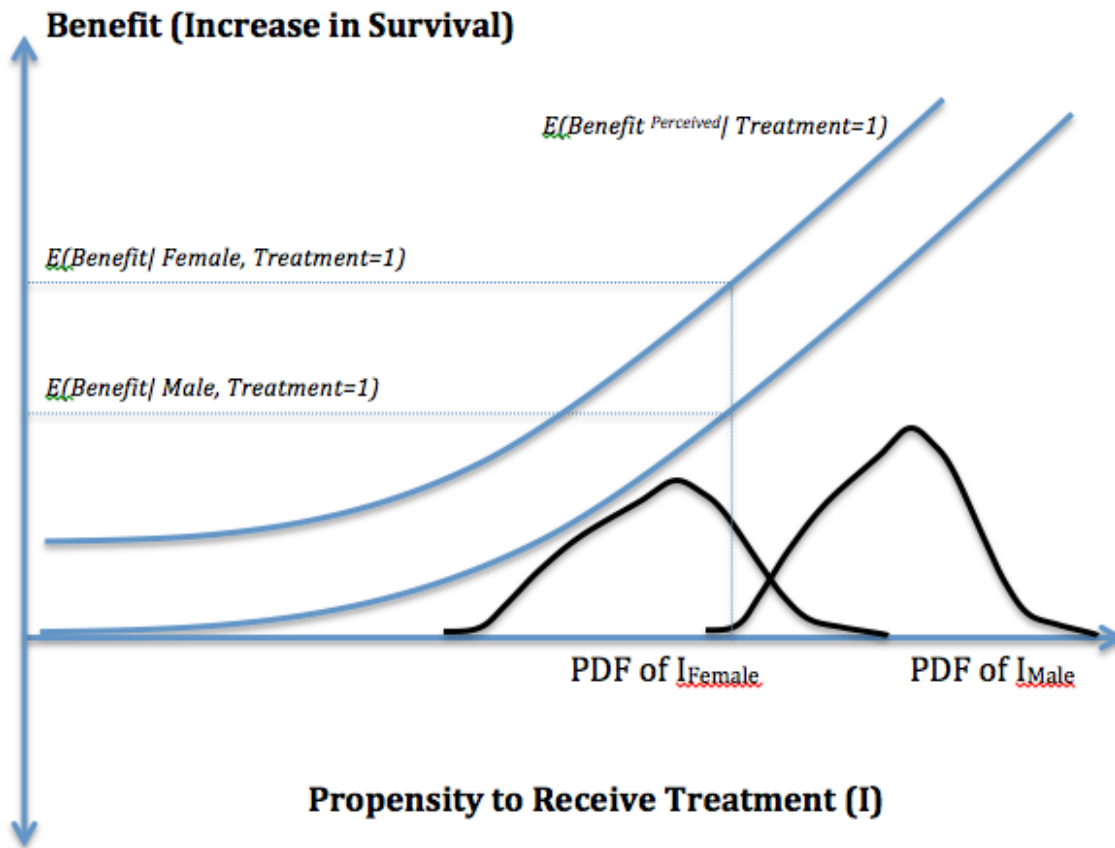
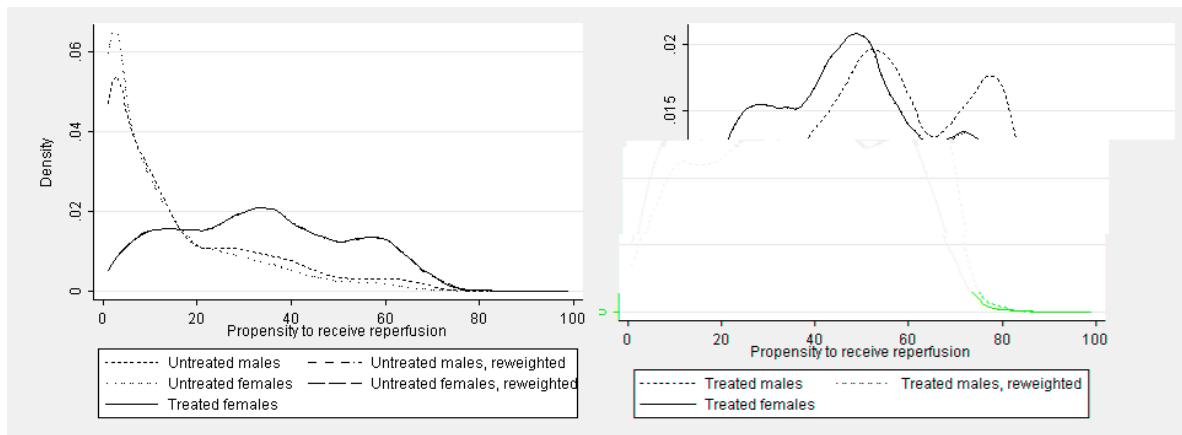


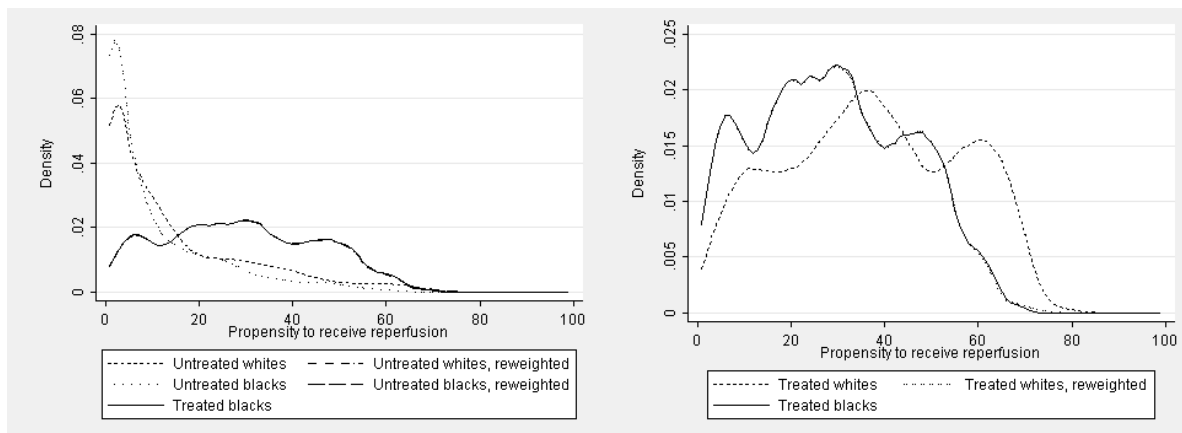
Figure illustrates how treatment-on-the-treated differs for men and women when there is prejudice against women. The treatment effect among women is higher at every point, reflecting the fact that the benefit of treatment must exceed a higher minimum threshold for women because of the perception that it is less effective in women.

Figure 2: Reweighting the Distributions of Propensities by Sex and Race

Panel A: Reweighting by Sex



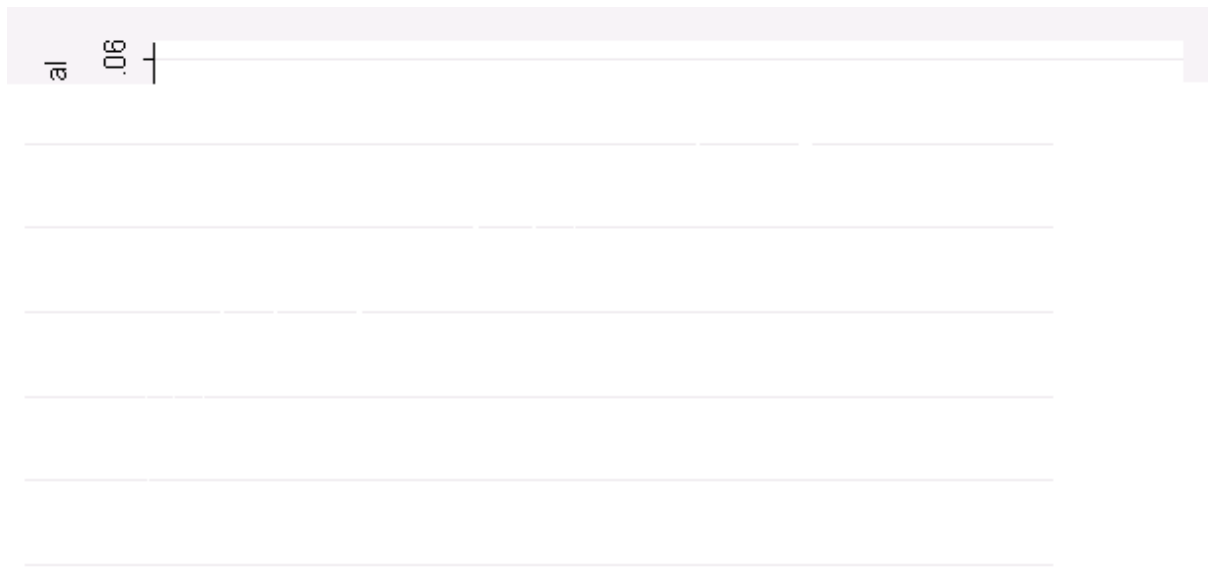
Panel B: Reweighting by Race



Note: the unweighted and reweighted kernel densities of the probability of receiving reperfusion are plotted by gender (panel A) and race (panel B) for the untreated (left hand side) and for the treated (right hand side). For reference, we have also included the propensity distribution for treated females (or blacks), since this is the target distribution that is being matched with reweighting. All reweighting is done using the method of Barsky et al. (2002).



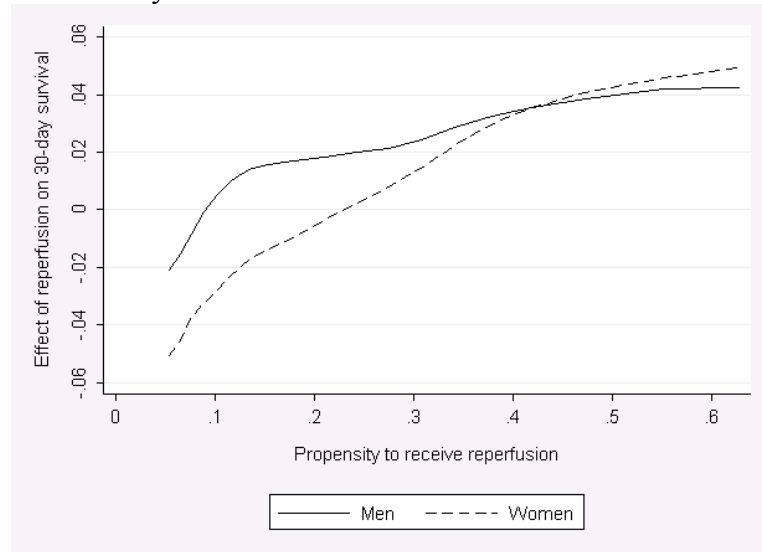
Figure 3: Effect of Reperfusion on 30 Day Survival, by propensity to receive reperfusion



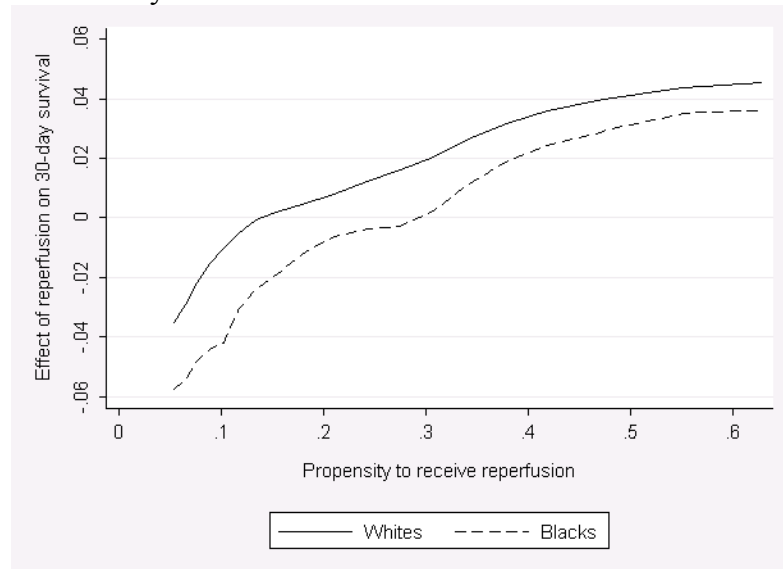
Note: Propensity to receive reperfusion is a patient's probability of receiving reperfusion within 12 hours of the heart-attack. It is calculated as fitted values from a probit model that includes sex, race and all covariates, and all interactions with sex and race. After ranking patients on this index, we calculated the effect of reperfusion on survival at each percentile of the index (see Table 3, column 3 and text for details).

Figure 4: Effect of Reperfusion on 30 Day Survival, by propensity to receive Reperfusion, and Sex and Race

Panel A: By Sex



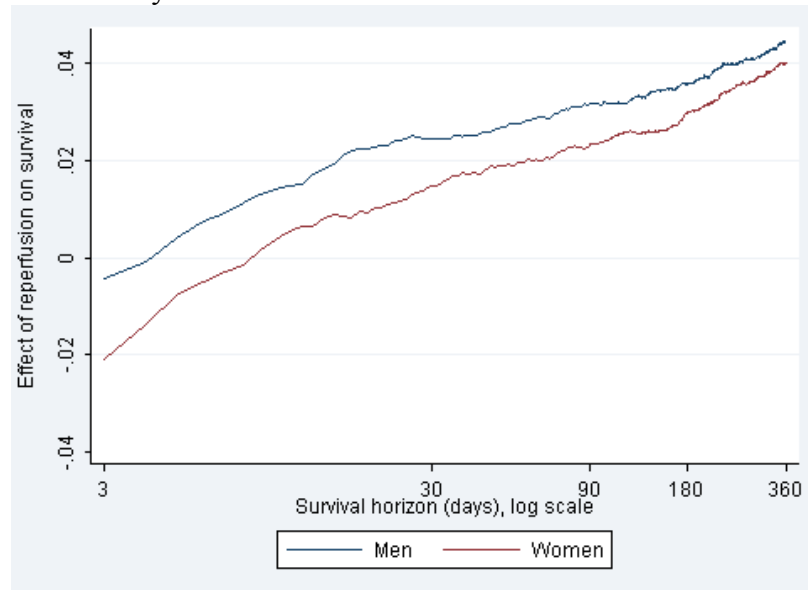
Panel B: By Race



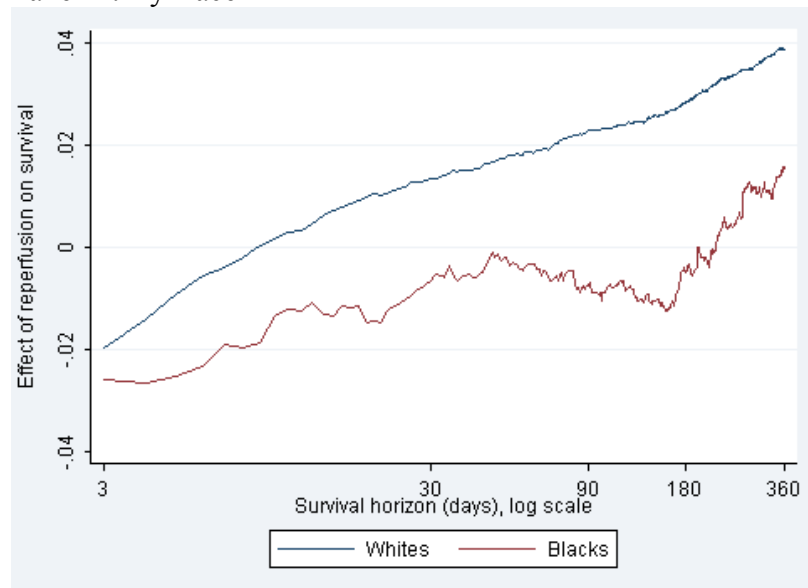
Note: Propensity to receive reperfusion is a patient's probability of receiving reperfusion within 12 hours of the heart-attack. It is calculated as fitted values from a probit model that includes sex, race and all covariates, and all interactions with sex and race. After ranking patients on this index, we calculated the effect of reperfusion on survival using local linear regression using a bandwidth of 10% of the sample and a linear kernel.

Figure 5: Effect of Reperfusion on Survival over Time

Panel A: By Sex



Panel B: By Race



Note: Figure reports the effect of reperfusion on the probability of survival at 3-365 days after the heart-attack. For each day of survival, we estimated a separate regression. The propensity of receive reperfusion is reweighted to resemble the distribution of treated minority patients (treated females in Panel A, and treated blacks in Panel B).

Table 1: Means by Sex and Race of Selected Variables

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	Men	Women	Whites	Blacks
Sample Size	70,154	68,803	130,672	8,285
Age	75.25	78.08	76.72	75.60
Pr (CHF)	0.19	0.25	0.21	0.27
Pr (Diabetes)	0.28	0.33	0.30	0.42
Pr (Hypertension)	0.56	0.68	0.60	0.80
Predicted (30 day Survival)	0.82	0.80	0.81	0.81
Pr (3 day Survival)	0.92	0.90	0.91	0.92
Pr (30 day Survival)	0.83	0.80	0.81	0.83
Pr (360 day Survival)	0.70	0.65	0.68	0.67
Pr (Reperfusion <= 12 hours)	0.21	0.16	0.19	0.12

Predicted 30-day survival is a summary measure of patient sickness at the time of the heart-attack. The prediction is made using all the CCP data, but not using race or gender.

Table 2: Probit estimates of the Association between Gender and Race on the Probability of Receiving Reperfusion with 12 Hours

	(1) No Controls	(2) Full Controls	(3) Full Controls
Female	-0.170 (0.008) [-0.045]	-0.043 (0.009) [-0.008]	-0.044 (0.010) [-0.008]
Black	-0.255 (0.018) [-0.061]	-0.251 (0.021) [-0.042]	-0.260 (0.031) [-0.044]
Female*Black			0.017 (0.042) [0.003]
N	138,957	138,957	138,957

Note: Dependent variable is the whether patient received reperfusion within 12 hours. Table reports probit coefficients with marginal effects evaluated at the point of full sample means in square-brackets. Probit model includes all CCP risk-adjusters, but does not allow for further interactions between sex, race and these covariates.

Table 3: Gender and Race Differences in 30 day Survival from Reperfusion within 12 Hours, OLS Estimates using reweighting

	Unweighted	Reweighted	Non-Parametric
	(1)	(2)	
<u>A: Gender Difference</u>			
Reperfusion	0.034 (0.003)	0.024 (0.003)	
Reperfusion*Female	-0.014 (0.005)	-0.010 (0.006)	-0.009 (0.005)
N	138,957	138,957	138,957
<u>B: Race Difference</u>			
Reperfusion	0.029 (0.003)	0.013 (0.003)	
Reperfusion*Black	-0.026 (0.012)	-0.020 (0.012)	-0.019 (0.012)
N	138,957	138,957	138,957

Note: Results for each column and in each panel are from a linear regression in which the dependent variable is a dummy for 30-day survival, and that controls for sex, race and all covariates, and all interactions with sex and race. Regression in column (2) reweights everyone using the method of Barsky et al. (2002) to look like treated females (Panel A) or treated blacks (Panel B). Column 3 includes 100 percentiles of I interacted with the receipt of Reperfusion. Robust standard errors reported in parentheses.

Table 4: Gender and Race Differences in the Effect of Reperfusion within 12 hours on Survival at Different Time-points, OLS Estimates

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Survival at:	3-day	7-day	30-day	90-day	360-day
<u>A: Gender Difference</u>					
Reperfusion	-0.004 (0.003)	0.009 (0.003)	0.024 (0.003)	0.032 (0.004)	0.045 (0.004)
Reperfusion*Female	-0.016 (0.004)	-0.012 (0.005)	-0.010 (0.006)	-0.009 (0.006)	-0.005 (0.006)
N	138,957	138,957	138,957	138,957	138,957
<u>B: Race Differences</u>					
Reperfusion	-0.020 (0.002)	-0.004 (0.003)	0.013 (0.003)	0.023 (0.003)	0.039 (0.004)
Reperfusion * Black	-0.006 (0.009)	-0.015 (0.010)	-0.020 (0.012)	-0.030 (0.013)	-0.023 (0.015)
N	138,957	138,957	138,957	138,957	138,957

Note: Results for each column and in each panel are from a linear regression in which the dependent variable is a dummy for survival as indicated, and that controls for sex, race and all covariates, and all interactions with sex and race. Regression reweight everyone using the method of Barsky et al. (2002) to look like treated females (Panel A) or treated blacks (Panel B). Robust standard errors reported in parentheses.

Table 5: Gender and Race Differences in the Effect of Reperfusion within 12 hours on Survival at Different Time-points, OLS Estimates with Hospital and Hospital Expertise Fixed-Effects

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Survival at:	3-days	7-days	30-days	90-days	360-days
<u>A: Gender Gap in Benefit</u>					
Reperfusion*Female	-0.017 (0.004)	-0.013 (0.005)	-0.010 (0.006)	-0.009 (0.006)	-0.005 (0.006)
Hospital FE	Yes	Yes	Yes	Yes	Yes
Hospital FE * Reperfusion	Yes	Yes	Yes	Yes	Yes
N	138,957	138,957	138,957	138,957	138,957
<u>B: Race Gap in Benefit</u>					
Reperfusion * Black	-0.010 (0.009)	-0.019 (0.010)	-0.020 (0.012)	-0.027 (0.012)	-0.039 (0.014)
Hospital FE	Yes	Yes	Yes	Yes	Yes
Hospital FE * Reperfusion	Yes	Yes	Yes	Yes	Yes
N	138,957	138,957	138,957	138,957	138,957

Note: Results for each column and in each panel are from a linear regression in which the dependent variable is a dummy for survival as indicated. Controls are included for sex, race and all covariates, and all interactions with sex and race, along with hospital and hospital\*reperfusion fixed effects. Regression reweight everyone using the method of Barsky et al. (2002) to look like treated females (Panel A) or treated blacks (Panel B). Robust standard errors reported in parentheses.