COMPARISON FRICTION:

EXPERIMENTAL EVIDENCE FROM MEDICARE DRUG PLANS*

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<u>ABSTRACT</u>

Consumers need information to compare alternatives for markets to function efficiently. Recognizing this, public policies often pair competition with easy access to comparative information. The implicit assumption is that comparison friction—the wedge between the availability of comparative information and consumers' use of it—is inconsequential because information is readily available and consumers will access this information and make effective choices. We examine the extent of comparison friction in the market for Medicare Part D prescription drug plans in the United States. In a randomized field experiment, an intervention group received a letter with personalized cost information. That information was readily available for free and widely advertised. However, this additional step—providing the information rather than having consumers actively access it—had an impact. Plan switching was 28 percent in the intervention group, versus 17 percent in the comparison group, and the intervention caused an average decline in predicted consumer cost of about \$100 per year among letter recipients—roughly 5 percent of the cost in the comparison group. Our results suggest that comparison friction can be large even when the cost of acquiring information is small, and may be relevant for a wide range of public policies that incorporate consumer choice.

Keywords: field experiment; Medicare Part D; prescription drug insurance JEL classifications: D89, I11

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I. INTRODUCTION

Government services increasingly rely on consumer choice. For instance, the design of the largest new social program of the last decade, Medicare Part D prescription drug insurance, relies heavily on consumers making choices. Choice is a key feature of Social Security privatizations and proposed school voucher programs. The rationale for including choice and competition is straightforward. Individuals have heterogeneous preferences and choice allows them to opt for services that best match their preferences. Competition between providers then facilitates a menu of services being provided at the cost-efficient frontier. In the best case, consumers get services that fit their needs better and governments save money.

This best case requires that consumers be able to look at the menu of options and pick the one that most cost effectively matches their needs. Making such choices requires having informed consumers. Yet, service providers may not provide all relevant information voluntarily. When service providers have information relevant to choices that consumers do not, this information asymmetry can undercut the benefits of choice and competition. As a result, policymakers frequently pair choice mechanisms with transparency systems—such as public school report cards, nutritional labeling, toxic pollution reporting, auto safety and fuel economy ratings, and corporate financial reporting (Weil et al. 2006)—all intended to make comparative information readily available.

Simply making information available, however, does not ensure consumers will use it. We call *comparison friction* the wedge between the availability of comparative information and consumers' use of it. (It is analogous to search friction—the challenge for buyers and sellers in locating each other.) Traditionally, public policies assume that comparison friction is largely

inconsequential as long as comparative data is provided for free and the benefits from comparing are non-negligible.

This study estimates the effect of reducing comparison friction in the market for prescription drug insurance plans for senior citizens. Over a period of about two and a half years, we followed the choices made by seniors who participated in an experiment we designed that reduced comparison friction by delivering personalized cost information to seniors via a letter. That personalized information used aspects of the match between consumers and the available plans (specifically, the differences in out-of-pocket costs of the drugs an individual takes) that could be readily observed.

One group of seniors—the intervention group—was presented with personalized price information created by entering their drug data into Medicare's Plan Finder website. They saw the cost of all plans for their personal drug profile as well as how much they would save by switching to the lowest-cost plan. A comparison group was given only the address of this website. The distinction between the groups was that the comparison group had to actively visit a website (or call Medicare's toll-free number, or seek information from a third party), whereas the intervention group had information delivered to them. We concentrate specifically on people who enrolled in a plan, and examine the effects of comparison friction on the choice of plan during an open enrollment period at the end of a year and on the plan selected for the coming year.

The intervention was designed to reduce comparison friction related to expected cost of plans. Other forms of comparison friction related to differences in uncertainty about costs or about requirements insurers might have for obtaining medication were not addressed. The

transaction costs of taking action and deciding to switch plans (involving a phone call) were relatively low.

We found large effects of this simple intervention. The intervention group switched plans 28 percent of the time, whereas the comparison group only did so 17 percent of the time. The average cost savings of the intervention-across the entire intervention group including nonswitchers—was about \$100 per year, or about 5 percent of the average predicted cost for the comparison group. We did not find effects on potential variability of consumer costs or on plan quality, although our power to do so was limited. The effects on consumer cost appeared to persist over time, although those estimates are imprecise. The intervention encouraged some individuals to switch to the lowest-cost plan and some to switch to other lower-cost plans. Although our sample included a larger proportion of college graduates and of people dissatisfied with their plans than a nationally representative sample would have, estimates of the effects for non-college graduates and for people satisfied with their plans (\$129 and \$74, respectively) were substantial. We estimate that our sample had larger potential savings in dollars—and similar savings in percentage terms—from switching to the lowest-cost plan than a national sample would have. The effects of the intervention for individuals below and above \$400 in potential savings were similar in relative terms $(0.052 \text{ and } 0.075 \log \text{ points}, \text{ respectively})$. Overall, the effects in these subgroups suggest that the results have broader applicability in a range of settings beyond that of the experiment itself.

These results fit into a set of recent studies that focus on comparison friction, such as those that examine the effect of the Internet in reducing comparison friction in markets where government plays a relatively minor role. (For example, see Brynjolfsson and Smith, 2000; Scott Morton, Zettelmeyer and Silva-Risso, 2001; Brown and Goolsbee, 2002; and Ellison and Ellison,

2009.) Hastings and Weinstein (2008), in a study of school choice where government plays a major role, found parents were more likely to choose a school with higher average test scores after receiving difficult to gather publicly available information about school scores. Moreover, children improved their own test scores after attending a higher-scoring school. A key difference between that study and ours is that the information provided in that study was plausibly hard to gather. In our context, the comparative data was relatively easy to acquire. Our results as a whole suggest that comparison friction could be effectively large even when the cost of acquiring information is low. This is consistent with other findings in contexts ranging from finance to nutrition to health where people have often failed to make use of comparative information that is readily available (Fung, Graham, and Weil, 2007). The findings have implications for a wide range of public policies that incorporate consumer choice.

The rest of the paper proceeds as follows. Section II provides a very brief background on Medicare Part D. Section III presents a conceptual framework for our analysis of plan choices. Section IV describes our data sources. Section V uses several of these sources (a cross-sectional survey and several audits of information sources) to characterize demand for and supply of information and knowledge of Medicare drug plans, and provide context for the experimental analyses. Section VI describes the experiment and presents results. Section VII discusses their interpretation. Section VIII discusses directions for future research.

II. MEDICARE PART D

The Medicare Part D prescription drug benefit was established as part of the Medicare Modernization Act of 2003, with coverage first beginning in January 2006. The drug benefit was subsidized, with Medicare paying about three-quarters of the premium. At the outset and again at

the end of each year during an open enrollment period, individuals typically chose from among 40-60 plans, depending upon where they lived.

Costs of plans included a monthly premium common to all beneficiaries of that plan and a personalized component that depended upon use. Under a standard plan for 2007 for drugs on the plan's formulary of covered medications, individuals paid 100% of the first \$265 of total costs (determined by the quantity of prescriptions and their full prices negotiated by the plan), 25% of total costs \$266-\$2400, 100% of total costs above \$2400 until own out of pocket costs reached \$3850. Above the \$3850 threshold, individuals paid either \$2.15 for a 30 day supply of generics and \$5.35 for brand name drugs, or 5% of further total costs– whichever was greater.

Most insurers offered two or three plans, including one actuarially equivalent to the standard plan and one or two with higher premiums and more cost sharing. Some plans had cost-sharing over the initial range (that is, no deductible) and some had cost sharing over the middle range (that is, offered some coverage through the coverage gap known as the "doughnut hole"). Still other variants had cost sharing in the form of a fixed price per prescription (co-payments with amounts depending upon the specific tier into which the plan had classified a drug) rather than as a percentage of the cost.¹ Actuarially equivalent plans (61 percent of enrollees in standalone plans nationally in 2006) used one or more of these variants as an alternative benefit design that covered the same share of enrollees' drug costs, on average, as the standard benefit (18 percent

¹ For example, Humana Insurance Company offered three standalone prescription drug plans (PDPs) in 2007. Humana PDP Standard had a premium that differed by state, ranging from \$10.20 to \$18.20 per month, and used exactly the same cost sharing as the standard plan described above. Humana PDP Enhanced had a premium ranging from \$17.10 to \$27.50 per month. There was no deductible, and the costs were: \$5 for a 30 day supply of preferred generic drugs; \$30 for a 30 day supply of preferred brand drugs; \$60 for a 30 day supply of other non-preferred drugs; 25% coinsurance for a 30 day supply of specialty drugs; \$15 for a 90 day retail supply of preferred generic drugs (\$12.50 mail order); \$90 for a 90 day retail supply of preferred brand drugs (\$75 mail order); \$180 for a 90 day supply of other non-preferred drugs (\$150 mail order). After the total yearly drug costs (paid by the individual and the plan) reached \$2,400, the individual paid 100% of costs until own out of pocket costs reached \$3850, at which point there was the same catastrophic coverage cost sharing as with the standard plan. Humana Complete had a monthly premium ranging from \$69.50 to \$88.40 per month, and differed from the Enhanced plan only in that it provided cost sharing in the coverage gap: \$5 for a 30 day supply of preferred generic drugs, and \$15 for a 90 day supply of preferred generic drugs.

of enrollees). Enhanced plans (20 percent of enrollees) covered a greater share of the drug costs, often using some cost sharing in the doughnut hole.

Insurers had different coverage of drugs and dosage forms (known as the formulary), which sometimes differed among plans offered by insurers. The insurers differed along a variety of other dimensions (that generally did not vary among plans offered by insurers), such as utilization management tools (prior authorization, step therapy, quantity limitations), pharmacy accessibility, mail order discounts, customer service, and financial stability of insurer.²

Medicare beneficiaries were offered the opportunity to voluntarily enroll in drug coverage either through a standalone plan (complementing fee-for-service health insurance through Medicare) or through a Medicare Advantage plan (often a health maintenance organization). This study focuses on individuals who were enrolled in standalone prescription drug plans in 2006 (the first year the benefit was offered) and were not receiving low-income subsidies. Nationally, this group was about 8 million of 43 million seniors (MedPAC 2007).

During an open enrollment period from November 15 to December 31 in 2006 (and similarly in subsequent years), individuals could switch plans. Prior to this period, individuals received a *Medicare and You* handbook, which contained fourteen pages describing Part D plans and answering frequently asked questions. The handbook indicated that one could switch plans by calling the plan that one wanted to join, or by calling 800-MEDICARE. The handbook included one page with a list of plans offered in one's state, with information on the monthly premium, benefit type (basic, enhanced without gap, enhanced with gap), and the percentage of the 100 most common drugs that are covered by the plan.

 $^{^{2}}$ For example, all three Humana plans in 2007 used the same pharmacy network and mail order system, and covered the same drugs using the same formulary with the same utilization management.

There are several reasons to suspect substantial comparison friction in Part D plan selection. For example, Heiss, McFadden, and Winter (2006) found that about 70 percent of seniors agreed with the statement "There were too many alternative plans to choose from" and more than half had difficulty understanding how Medicare Part D worked and what savings to expect. Earlier research has found that seniors have difficulty navigating insurance choices within Medicare (Gold, Achman, and Brown 2003; Hibbard et al. 2001; McCormack et al. 2001).

Recently, a number of authors have examined the quality of seniors' choices and the effect of information in the context of Medicare Part D. Heiss, McFadden, and Winter (2010) used survey data and concluded that seniors' decision to enroll in 2006 responded to incentives provided by their health status and the environment but acknowledged that "enrollment is transparently optimal for most eligible seniors."

Abaluck and Gruber (2011) examined claims data from 2005 and 2006 and determined that most seniors did not choose plans on an efficient frontier, defined in terms of expected cost and its variance. In addition, relative to a rational model of choice, seniors placed more weight on plan premiums than on out-of-pocket costs, placed weight on plan's financial characteristics (e.g. the presence of a deductible) independent of the effect of those characteristics on their own costs, and showed unexpectedly low levels of risk aversion—all behaviors that contradict a rational, normative model of plan choice.

Based on an analysis of two years of panel data, Ketcham et al.(2010) found that seniors reduced their out-of-pocket costs for insurance and prescription drugs above the cost of their cheapest ex-post alternative between 2006 and 2007, although this improvement came both from active decisions to change plans and convergence among plans. Using a laboratory experiment, Bundorf and Szrek (2010) reported that both the benefits and costs of choice increase with the

number of options available. In another laboratory experiment, Hanoch et al. (2009) found that participants were less likely to correctly identify the plan that minimized total cost when presented with larger numbers of plans. Taken together, these results suggest some potential for choice errors, particularly for more difficult choices, in newer situations, and for less able individuals; they also suggest some potential for learning and adapting and a role for information.³

III. CONCEPTUAL FRAMEWORK

To visualize the choice problem facing the individual, we focus on three decisions. We define the following random variables representing the distribution of potential realizations for each individual:

- \tilde{b}_{ij} is the potential benefit to the individual *i* from plan *j* minus switching costs;
- \tilde{p}_{ij} is the component of potential consumer cost for plan *j* that can be predicted from comparative research (based on extrapolations of last year's drug use); and
- \tilde{c}_{ij} is the component of potential consumer cost for plan *j* that cannot be predicted from comparative research.

We assume a utility function for increments to the utility from current consumption from participating in a plan, such that $\tilde{u}_{ij} \equiv U(\tilde{b}_{ij} - \tilde{p}_{ij} - \tilde{c}_{ij})$ and marginal utility is decreasing in its arguments. Also, r_i is the comparison friction—specifically, each individual's known cost of undertaking comparative research about the costs of plans (expressed in the same units as \tilde{u}_{ij}).

³ Other research on Part D has examined the market structure and plan dimensions, such as the incentives that exist for adverse selection (Goldman, Joyce, and Vogt, 2011), prices for branded drugs (Duggan and Scott Morton, 2011), and welfare impacts of limiting the number of Part D plans (Lucarelli, Prince, and Simon, 2008). The cost management strategies do appear to have encouraged people to switch to cheaper medications (Neuman et al. 2007). Utilization has increased, while seniors' expenditures have decreased (Yin et al. 2008). About one-third of new public expenditure has crowded out previous private expenditure (Engelhardt and Gruber, forthcoming).

III.A. Plan Choice Without Research

Without research, the highest level of expected utility across all plans, taking the expectation over the joint distribution of all the random variables that determine \tilde{u}_{ij} is given in equation (1).

(1)
$$h_i^1 = \max_i E(\tilde{u}_{ii})$$

If research is not undertaken, then the plan j that maximizes expected utility in equation (1) will be selected—and if this is the current plan, the individual will not switch plans.

III.B. Plan Choice With Research

If research is undertaken, p_{ij} is a realization of \tilde{p}_{ij} . The highest level of expected utility across all plans is then given in equation (2), where r_i is additively separable from \tilde{u}_{ij} for simplicity of exposition. The individual selects the plan *j* that maximizes this expression.

(2)
$$h_i^2(p_{i1}, p_{i2}, ..., p_{ij}) = \max_j E(\tilde{u}_{ij} | \tilde{p}_{ij} = p_{ij}) - r_i$$

III.C. Deciding to Undertake Research

The decision to undertake research involves comparing h_i^1 to the expected value of h_i^2 when the predictable cost component is uncertain (because research has not yet been undertaken). This expected value is shown in equation (3), taken over the joint distribution of the predictable cost component of all plans and the cost of comparative research for an individual.

(3)
$$h_i^3 = E[h_i^2(\tilde{p}_{i1}, \tilde{p}_{i2}, ..., \tilde{p}_{ij})]$$

The individual undertakes research if the expected value of the maximum expected utility from undertaking research is greater than the maximum expected utility from the plan that would be chosen without research $(h_i^3 \ge h_i^1)$ and does not otherwise. The key conceptual distinction between h_i^3 and h_i^1 is that h_i^3 captures the option value of comparative research. That is, if the research could reveal substantial predicted savings from a plan, then paying a small cost of research tends to be worthwhile.⁴

III.D. Implications

Because research reduces uncertainty about costs, we would expect plan choices based on research to be more sensitive to cost differences than those made without research. Thus, we would expect an intervention that reduced the cost of comparative research to cause individuals to put more weight on cost (as a random variable with lower variance in their expected utility calculation) when evaluating plans—which would tend to reduce potential consumer cost of selected plans.

When r_i is reduced for some individuals, we would expect more people to undertake research and switching plans would be worthwhile for more people. However, the magnitude of the effect of the intervention on potential consumer cost is not necessarily determined by the magnitude of r_i because the cost savings must be sufficient to compensate for switching costs.

IV. DATA SOURCES

Several sources of data were used for this paper. A national phone survey of Part D beneficiaries was fielded to understand their knowledge of and experience with Medicare drug plans and their sources of information. An audit of potential sources of information was undertaken to understand the information available to them. A sample of pharmacy claims data was obtained to examine the potential savings available to beneficiaries who could change plans.

⁴ For example, assume there are only two plans—one is the current plan and a second is an alternative. Without research, assume the alternative has an equal chance of having large predicted savings on average (yielding X) or large predicted cost on average (yielding -Y) such that its expected utility (0.5X - 0.5Y) is negative. Under these conditions, the alternative will not be selected over the current plan. However, it will be worthwhile to do research that will reveal whether the plan will result in utility of X or -Y if the combination of utility from predicted savings X and the zero utility change from staying with the current plan is greater than the cost of the research—that is, $(h_i^3 \ge h_i^1)$ when $(0.5X + 0.5*0 - r \ge 0)$.

And a field experiment was conducted in which some beneficiaries received a personalized letter about drug plan choice, while others received general information.

IV.A. National Phone Survey

We commissioned a phone survey of Medicare beneficiaries over age 65 who were enrolled in stand-alone Medicare prescription drug plans. The survey was fielded in February and March 2007. A market research firm generated an initial sample of phone numbers; these numbers were intended to reach seniors with high probability and, ultimately, to generate a nationally representative sample of seniors. 26 percent of people reached by phone agreed to begin the survey. Of these, 49 percent did not meet screening criteria, 8 percent did not complete the survey, and 43 percent were both eligible and completed the survey. An additional 13 percent of respondents were removed from the sample due to incomplete data, leading to 348 responses.

IV.B. Audit of Information Sources

Actors, hired by the researchers, made 12 calls to Medicare, 5 calls to State Health Insurance Programs (SHIPs), 88 in-person visits to Boston area pharmacies (stratified by chain/independent/retail, urban/suburban, and community income), 8 in-person visits to Bostonarea senior centers, and 12 calls to other help-lines, identified via an internet search, during the open enrollment period in December 2006. For the in-person audits, an actor, aged approximately 65, posed as a Medicare beneficiary and asked for advice about choosing a drug plan using a set of questions developed by the research team. For the phone audit, a research assistant, posing as a relative of a Medicare beneficiary, asked these same questions.

IV.C. Sample of Pharmacy Claims

We derived drug profiles for 59 seniors with Medicare drug plans from the 2006 claims of a large pharmacy chain's stores in one state. For 41 of these seniors, we identified a sponsor but not a plan. For these individuals, we calculated costs for the lowest- and the highest-cost plan among those offered by the sponsor. Cost measures were then created using the Medicare Plan Finder website to compare costs of selected plans and the lowest-cost plans. The pharmacy data is likely missing some data on prescriptions that the individual filled at other pharmacies; a countervailing factor is that some individuals with insurance but without prescription use are omitted from the sample by construction.

IV.D. Experimental Intervention

The experiment and associated data collection consisted of three surveys and one mailing. A baseline phone survey was conducted in November 2006. A letter intervention was mailed in December 2006. Two follow-up phone surveys were fielded: one in April and May of 2007 and one in March and April of 2008.

Patients of the University of Wisconsin Hospital system age 65 and over made up the sample frame. Letters of invitation were mailed to 5,873 subjects who were then contacted by phone. Approximately half of those agreed to join the study. Of these, approximately 15 percent met screening criteria, and reported a plan name that could be later matched to the Medicare Plan Finder, leading to a baseline sample size of 451.

In the baseline survey, the participants answered detailed questions about their prescription drug use and basic questions about personal characteristics. Researchers constructed measures of beneficiary costs in each respondent's current drug plan and in all available drug plans by entering the respondent's drug utilization information into the Medicare Plan Finder website.

After baseline data were collected, participants were randomly assigned to intervention and comparison groups. Each participant received a personalized mailing. The materials participants received are shown in the Appendix.

The 2007 follow-up inquired about the plan chosen for 2007 and the choice process. 406 people completed the baseline survey, were randomly assigned in the experiment, and completed the 2007 follow-up survey—forming our main analytic sample—and 45 people completed the baseline survey and were assigned to the experiment but could not be reached for follow-up. Thus, although about half of those sent a letter of invitation did not agree to participate (which is relevant for external validity), the study does include 90 percent of people randomly assigned to a group in the experiment (which is particularly relevant for internal validity of the results from the experiment). 92 percent of intervention group and 88 percent of comparison group completed the 2007 follow-up survey.

The 2008 follow-up collected data on drugs used in 2007 and 2008, experiences in the 2007 plan, and the plan chosen for 2008.⁵ 305 of those 406 people completing the 2007 follow-up survey also completed the 2008 follow-up survey.

Data on actual plan enrollment in 2007 (from the 2007 follow-up survey) and 2008 (from the 2008 follow-up survey) was used to determine when individuals had switched plans. To assess the dispersion in costs across plans for the same individuals, we compiled data on the predicted and actual costs of every possible plan. Predicted cost for 2007 is the estimated annual cost measure computed by the Medicare Plan Finder for a given drug plan based on an individual's prescription drug use as reported at the time of random assignment in fall 2006. The

⁵ In order to construct cost measures, because the 2007 Medicare Plan Finder was no longer available in 2008, researchers entered respondents' reported 2007 drug utilization into a 2007 version of a private-sector counterpart of the Medicare Plan Finder, the Experion Plan Prescriber. The Plan Finder was used to construct cost measures for 2008, and tests based on the 2008 releases of both tools demonstrated a high-level of agreement (>90 percent).

Plan Finder computed the out-of-pocket cost for each plan, assuming that the drugs entered would be taken for the full year of 2007 and that chemically-equivalent generic drugs would be substituted for brand-name drugs. Actual realized cost for 2007 is the estimated annual cost measure computed by the Medicare Plan Finder for a given drug plan based on an individual's prescription drug use throughout 2007 (with dosages prorated to reflect that cumulative use) as reported at the time of the second follow-up survey in 2008. Predicted cost for 2008 and for 2009 is based on current drug use as reported in the 2008 follow-up survey.

V. CHOICE ENVIRONMENT

We used our phone survey to assess the context within which seniors made choices about whether or not to change prescription drug plans during open enrollment. A significant majority of respondents knew that different plans were better for different people (82 percent) and that they could only change plans during open enrollment (74 percent). (See Table I.) However, few had learned additional facts about the specific differences among plans. Only 37 percent knew that only some (rather than all) plans have a deductible. Only 55 percent knew that different plans have different co-payments for generic drugs, rather than all plans having the same co-payments.⁶

We found that over 80 percent of respondents were generally satisfied with their 2006 prescription drug plans. The percentage that switched plans between 2006 and 2007 was 10 percent, slightly above the reported national rate of seven percent.⁷ An additional 14 percent considered switching for 2007 but did not switch, which is consistent with the high levels of

⁶ In survey data collected in 2005, just prior to the beginning of the first open enrollment period, Winter et al. (2006) also found low knowledge about the structure of the benefit and the potential for differences among plans.

⁷ That national rate is for those not receiving the Low Income Subsidy (U.S. Department of Health and Human Services, 2007).

reported satisfaction.8

The leading sources of information that participants used to learn about drug plans were mailings from plans and mailings from Medicare. That material is not personalized and does not convey transparent information about out-of-pocket costs. The more interactive forms of information gathering, such as in-person, phone, or internet, were each used by less than 15 percent of respondents. Eighteen percent reviewed personalized plan comparisons.⁹

To better understand the information available in the existing choice environment and the costs of acquiring it, we audited five potential sources of advice on choosing a drug plan: the Medicare help-line (1-800-Medicare), state health insurance assistance programs (SHIPs), senior centers, other telephone help-lines, and retail pharmacies. In our calls to 1-800-Medicare, customer service representatives consistently entered personalized drug information, identified a low cost plan, and offered to enroll the caller—drawing upon Medicare's website tool, the Prescription Drug Plan Finder. Our calls to SHIPs generated either referrals to Medicare or offers of similar assistance. Our visits to senior centers sometimes resulted in general discussions about the drug benefit or partial demonstrations of the Medicare website but never in comparative information left in the hands of the auditor. A search for and audit of other sources of telephone advice indicated that few private-sector information sources had emerged.¹⁰ In general, these sources were either not helpful or referred the caller to Medicare or another public-sector

⁸ Our survey results are similar to Heiss, McFadden, and Winter (2010), who reported that 82 percent rated their 2006 plan good or better, 18 percent considered switching for 2007 but did not, and 11 percent switched plans from 2006 to 2007. Unpublished results from the same survey used in that research indicated that 60 percent did not consider switching because they were happy with their plan while 18 percent "wanted to avoid the trouble of going through the plan comparison and choice process again."

⁹ Our results are broadly consistent with the U.S. Department of Health and Human Services (2007), which reported results from a survey in January 2007 indicating that 85 percent of seniors were aware of the open enrollment period, 50 percent reviewed their current coverage, 34 percent compared plans, and 17 percent evaluated premiums, co-payments, and coverage.

¹⁰ A contributing factor may be Medicare policies, motivated by concerns about conflicts of interest that restrict the extent to which third parties can provide advice.

information source. In one noteworthy exception (a major pharmacy chain), the help-line offered personalized suggestions, using technology similar to Medicare's, and mailed a personalized report.¹¹

A small fraction of pharmacies offered personalized in-store assistance with plan choice to auditors who walked in. In four of the 88 pharmacies audited, staff people made personalized plan suggestions based on a Plan Finder. In five pharmacies (all in one chain), a staff person offered personalized plan information about the entire universe of available plans. Sixty-nine of the 88 pharmacies provided print materials. (Separately, tests we gave to recipients of print materials indicated that these materials alone were not sufficient for seniors to understand the cost implications of plan choice even in very simple cases.)

In sum, seniors could acquire personalized assistance from Medicare with minimal effort, but seniors who sought information through other channels were not consistently assisted or even consistently directed to Medicare. Personalized information was readily available but not widely diffused.

VI. INTERVENTION IN THE CHOICE ENVIRONMENT

To examine the extent to which a reduction in comparison friction would affect plan choices, we designed a randomized experiment in which the intervention lowered the cost of obtaining and processing the information needed to make comparisons. Members of the intervention group received a one page cover letter showing (1) the individual's current plan and its predicted annual cost conditional on their personalized drug profile, (2) the lowest-cost plan and its

¹¹ In addition, a second major pharmacy chain offered an internet service in conjunction with a technology partner specializing in decision support systems. A code was developed to trigger the import of individual medications into the partner's Medicare Part D decision tool. Customers and pharmacy staff were able to produce personalized Medicare Part D Plan comparisons by entering these codes into the tool.

predicted annual cost, (3) the potential savings from switching to the lowest-cost plan, and (4) the date of the end of open enrollment. They also received a printout from the Medicare Plan Finder including costs and other data on all available plans. The comparison group received a general letter referring them to the Medicare website. Both groups received an informational booklet on how to use the site. (For examples of these letters and the booklet, see the Appendix). The intervention included a recommended default option (the lowest-cost plan), a clear statement of that option's benefits (potential savings), and a deadline. It neither contained difficult to acquire information nor reduced the effort required to change plans.

VI.A. Baseline Characteristics

At the time of the baseline interview, participants reported regularly using an average of five and half medications. The study participants were all from Wisconsin, nearly all white, with an average age of 75. About two-thirds were women, about two-thirds were married, and about half were college graduates (see Table II). Relative to the national population of seniors, study participants were typical in terms of age and gender but were more likely to be married and were substantially better educated.

The potential savings from changing plans, as a share of current expenditure, was similar in our intervention sample and in our entirely separate sample of pharmacy claims data suggesting that the study did not disproportionately attract those who stood to benefit financially from changing plans. Specifically, predicted consumer cost could be reduced an average of 30 percent in our intervention sample by switching to the lowest-cost plan. The corresponding reduction was between 24 and 41 percent in the pharmacy claims sample. (The reason for the range is that among plans offered by a particular plan sponsor, we could not determine the

specific plan currently covering an individual in many instances. The smaller potential savings is based on a calculation using the lowest-cost plan among those offered by a sponsor, and the larger potential savings is based on imputation from the sponsor's highest-cost plan.)

Although the proportional potential savings was similar, the level of expenditure on prescription drugs was substantially higher in our intervention sample than samples more representative of the general population, including our pharmacy claims and national samples. For example, Domino et al. (2008) projected costs under Part D for a nationally representative sample deriving medication usage from the Medicare Expenditure Panel Survey (MEPS). The average 2006 predicted cost in the lowest-cost plan for individuals in the MEPS was \$1114, which was 30 percent lower than the corresponding 2007 predicted cost of \$1593 for the lowest-cost plans in our intervention sample. That difference is probably due to the intervention sample being drawn from a list of patients with recent clinical visits (and tending to have more health problems and higher levels of prescription drug use), and also to being from a more recent year.

Individual characteristics for the 406 individuals with complete data from both the baseline survey in 2006 and the 2007 follow-up survey were similar for those assigned to the intervention and comparison groups, although the intervention group had a higher fraction age 75 or older and a higher fraction whose satisfaction with their 2006 plan was fair or poor.

VI.B. Percentile Rank in Cost of Chosen Plans

As context for understanding the potential savings from switching plans during open enrollment, we examined the percentile rank in cost of chosen plans in the distribution of available plans, as calculated based on the medication usage reported in our baseline survey. There were 54 Medicare prescription drug plans available to beneficiaries in our Wisconsin

sample. The baseline plans initially enrolled in for 2006 by the individuals in our sample were nearer the median-cost plan than the lowest-cost plan among all those offered: the average rank was at the 39th percentile.

To see how the baseline plans compared to other plans of similar benefit type, we grouped the plans into three different types. "Basic" included: plans with a deductible, 25 percent cost sharing, then a coverage gap, and then catastrophic coverage (known as defined standard plans); actuarially equivalent plans with the same deductible as a defined standard plan but a different cost sharing structure (known as actuarially equivalent standard plans); and actuarially equivalent plans with a reduced or eliminated deductible and a different cost sharing structure (known as basic alternative plans). "Enhanced without gap" included plans with actuarial value exceeding the defined standard plan and no cost sharing in the coverage gap. "Enhanced with gap" included plans with actuarial value exceeding the defined standard plan and with cost sharing in the coverage gap (only for generic drugs, except for one plan).

There is considerable variation in the consumer costs among basic plans, which are identical or actuarially equivalent to the standard plan in terms of insurance value. One plan was the lowest cost of the basic plans for about half the individuals in our sample, but fourteen different plans were the lowest-cost basic plan for others–depending on the drugs they took. On average, the baseline plan was at the 38th percentile of basic plans (among those who reported enrollment in a basic plan for 2006 in the baseline survey). For individuals with enhanced plans without gap coverage in 2006, the baseline plan was at the 43rd percentile of predicted consumer costs among plans of that type. For those having enhanced plans with gap coverage in 2006, the baseline plan was at the 50th percentile of costs among that type of plan. (For analysis of the differences in

dollars of cost between baseline plans and the lowest-cost plans by benefit type, see the Appendix.)

In addition to analysis of percentile rank of predicted costs based on drugs taken in 2006, we also examined costs based on the drugs actually taken throughout 2007 as reported in the 2008 follow-up interview. That percentile was similar—37th for actual costs versus 39th for predicted costs—when plans of all benefit types were compared. For actual costs among basic plans (among people who had that type plan in 2006), the percentile rank of 35th was slightly higher than for predicted costs. Among enhanced plans without and with gap coverage (again, among people who had that type of plan in 2006), the percentile ranks were 37th and 49th, respectively. Thus, it appears that both ex-ante and ex-post there were numerous options for reducing consumer costs among all plans and among plans of the same benefit type.

VI.C. Intervention Impacts

Switching in 2007. 28 percent of those in the group receiving the letter intervention switched plans between 2006 and 2007, compared to 17 percent in the comparison group.¹² The difference of 11.5 percentage points is found in a simple comparison of means (see Table III).

We also estimated the effect of the intervention (Z) on plan switching (D) using linear regression and controlling for covariates (X) known at the time of random assignment including the age and plan rating variables where there were some differences between the comparison and intervention groups as discussed above—as in equation (4).

(4)
$$D_i = Z_i \pi + X_i \beta + \varepsilon_i$$

¹² The switching rate in the comparison group is more than twice as high as the national average, which is likely related to the higher rates of drug utilization and the higher plan dissatisfaction in our sample.

After regression adjustment, the estimated difference is 9.8 percentage points. The probability of such a large difference occurring by chance under the null hypothesis of no effect of the intervention is very small, with p-values less than 0.02 for both specifications. People rating their baseline plan as fair or poor were ten percentage points more likely switch plans, holding other factors constant. The regression-adjusted impact of the intervention on switching is slightly lower than the simple comparison of means primarily because of the interaction between those marginal effects and the higher baseline prevalence of those low satisfaction ratings in the intervention group than in the comparison group.

The average time spent on all aspects of plan consideration and possible switching was 3 hours in the comparison group. Exploring seniors' choice process and knowledge, we found that several of the differences between the two groups supported the notion that the intervention worked through cognitive channels. These included statistically significantly greater percentages of intervention group members later reporting that they remembered receiving the materials, that they read them, and that they found them helpful.

Predicted 2007 Costs. To estimate effects on costs, we used the same approach as in equation (4) except with change in cost as the dependent variable. Specifically, the change in cost $(Y^s - Y^b)$ is the 2007 predicted consumer cost of the plan selected for 2007 minus that cost for the baseline plan that had been selected in 2006. The average regression-adjusted decrease in predicted cost for the entire intervention group versus the comparison group was \$103 (see Table III). Expressed in terms of the change relative to 2006, again estimated using the same approach as in equation (4) but with log of the relative change in cost $[\ln(Y^s/Y^b)]$ as the dependent variable, this decrease was an average of 0.064 log points. Again, the probability of such a large difference occurring by chance under the null hypothesis was less than 0.005. (The average cost change for

the entire intervention group versus the comparison group averages over people who were not affected by the intervention and those who potentially were affected. Estimates for those affected are discussed in the Appendix.)

Covariates other than the intervention indicator had little power to explain the changes in consumer cost between the baseline 2006 plan and the plan selected in 2007. Holding other factors constant, the indicator for seven or more medications was associated with a reduction in predicted consumer cost of \$112, indicating that people with higher levels of medication use chose plans in 2007 resulting in larger reductions in predicted consumer costs than people with less use.

Potential Variability of 2007 Costs. Plans differed in the extent to which costs could be higher or lower if medication use were to change in the future. To create a measure of that potential variability, we used our data on the predicted consumer cost of every plan offered for each of the 406 individuals in our sample. For each plan, we then calculated the difference between the 90th and 10th percentiles of the predicted 2007 consumer cost among individuals that reported taking a similar number of medications in the baseline survey. (Specifically, the percentiles were calculated within three subsamples: 0-3 medications, 4-6 medications, and 7 or more medications.) This approach—similar to that used by Abaluck and Gruber (2011)— implicitly assumes that the experiences of other members of our sample with similar medication use represent the range of potential variability for each individual. Because of the small samples sizes used, however, that assumption only holds very roughly, and that measure is also imprecise.

The sample average for that measure of potential variability was about \$2900 for the 2006 plans used at the time of the baseline survey. In analysis of the change in that measure between

the 2006 plan and the 2007 plan for each individual, using the same approach as in equation (4), potential variability for the intervention group was \$10 less than the comparison group (with a standard error of \$36). Thus, we did not find evidence that the intervention caused individuals to switch to plans that had greater potential variability according to this measure.¹³

Actual 2007 Costs. In terms of the impacts on consumer cost as measured for the respondents to the 2008 follow-up survey, the predicted 2007 cost was \$111 lower for the intervention group and the actual cost was \$137 lower—although the standard error was 4 times larger for the impact on actual cost (see Table IV).¹⁴ In addition to allowing more precise estimation, we focused our primary analysis on the predicted 2007 consumer cost of the plan chosen for 2007 rather than the actual 2007 cost because the predicted cost uses the information set available to individuals when they were making their plan choice during open enrollment prior to 2007, which corresponds to the predictable component of costs discussed in section III. Also, the predicted 2007 cost has less attrition, since it is based on our 2007 follow-up survey.

Along with having similar impacts for actual and predicted costs, those two measures were fairly similar at the individual level. The correlation between the actual cost and the predicted cost was 0.68; excluding the three most extreme differences, the correlation between actual and predicted costs was 0.79.¹⁵ Comparing the actual and predicted costs of the 2007 plan selected by

¹³ We also used other measures of dispersion, such as the standard deviation and interquartile range, and found effects that were also statistically insignificant.

¹⁴ The actual 2007 cost is based on drug list information collected in 2008 (as is the predicted 2008 and 2009 cost), whereas predicted 2007 cost uses the 2006 baseline drug list. Since the baseline data is used in the calculation of the predicted 2007 cost of both the 2007 and 2006 plans and their difference is used in the estimation, the estimate of the effect of the intervention on that outcome is much more precise that the estimates of the effect on the actual 2007 cost. That is, the outcomes derived from the information collected in 2008 have much more variability in cost that is not removed by subtracting the predicted 2007 cost of the 2006 plan. The impacts in log points estimated based on the 2008 follow-up survey data also show more instability in magnitude and sign that those based on the 2007 follow-up survey, consistent with the imprecise nature of those estimates.

¹⁵ In a regression of actual realized cost on predicted consumer cost for the 2007 plan selected (based on drugs taken in 2006), adding the demographic characteristics gathered in 2006 (gender, marital status, education, age, number of medications, plan satisfaction) to the model increased the R-squared from 0.47 to 0.52, with significant coefficients on having seven or more medications and having low plan satisfaction.

the individual, the actual cost was an average of \$354 higher than predicted cost among respondents to our 2008 follow-up survey. In the distribution of differences between actual and potential costs, the actual cost was \$1872 higher at the 90th percentile, \$51 higher at the median, and \$663 lower at the 10th percentile.¹⁶

Quality in 2007. Our 2008 follow-up survey also collected self-reported information on experiences in the plan during 2007. There were no statistically significant differences in satisfaction with non-cost features or in overall plan ratings, although the point estimates go in the direction of relatively more dissatisfaction with non-cost features and less dissatisfaction overall for the intervention group (see Table IV). Thus, it is possible that individuals chose lower-cost plans that had lower quality; we do not have sufficient statistical power to reject a hypothesis of small reductions in quality. Analysis of other measures of administrative quality at the plan sponsor level showed essentially no impact.

Switching in 2008. In another assessment of choices from our 2008 follow-up survey, we examined whether individuals were sufficiently satisfied with their choices in 2007 to keep them for 2008 after receiving another opportunity to switch plans. 23 percent of the comparison group switched in 2008, and 20 percent of the intervention group switched—a statistically insignificant difference—implying that the intervention group was at least as satisfied as the comparison group overall in terms of their revealed preferences.

Predicted 2008 and 2009 Costs. The impacts on 2008 and 2009 predicted costs were of roughly the same magnitude as both the predicted and actual costs for 2007 (see Table IV). Like

¹⁶ When calculated as an average over all possible plans, rather than the 2007 plan selected by the individual, the actual cost was \$339 higher than predicted cost. In the distribution of differences, the actual cost was \$1888 higher at the 90th percentile, \$50 higher at the median, and \$838 lower at the 10th percentile. In terms of the ratio of the actual to the predicted costs, at the 90th percentile the actual was about 90 percent higher, and at the 10th percentile the actual was about 30 percent lower. The correlation of the within-person ranks of 2007 actual and predicted cost was 0.70.

those actual costs, they were imprecisely estimated. We interpret these results as being consistent with continued savings over time due to the intervention, but we also could not reject a null hypothesis of no impact at conventional levels of statistical significance.

VI.D. Role of the Lowest-Cost Plan

We found some evidence that more aspects of the intervention mattered for decision-making than simply the identification of the lowest-cost plan. The intervention letter sent in the fall of 2006 named the plan with the lowest predicted consumer cost in 2007, based on reported prescription drug use, and gave the predicted cost and calculated the difference in cost relative to the 2006 plan. An attachment to the intervention letter also showed the predicted cost of each plan. We found that 9 percent of the intervention group switched specifically to the lowest-cost plans while 20 percent switched to a different plan; in the comparison group these percentages were 2 percent (statistically significantly different from 9 percent) and 15 percent (not statistically significantly different from 20 percent). This result is consistent with the idea that the intervention specifically caused seniors to consider the lowest-cost plan, and also that seniors gave additional consideration to the personalized cost of plans other than the lowest-cost plan.

As a complement to the analysis of the impact of the intervention on switching rates and average predicted costs and to give more structure to the estimated effects, we also examined differences between the intervention and comparison groups in discrete choice models of plan selection. As a point of departure for this analysis, consider selecting a plan at random, which is equivalent to a discrete choice model with coefficients of zero on explanatory variables. The probability of plan selection from among 54 plans would be 1/54 = 0.019. To examine the probability of selecting plans of different prices, we formulated a conditional logit model for

individual *i* and plan *j* estimated using comparison group data only and controlling for individual fixed effects (α_i), predicted cost (P_{ij}), and predicted cost squared, based on the indirect utility function in equation (5), with utility (u_{ii}^*) and an error term (ε_{ii}).¹⁷

(5)
$$u_{ij}^* = P_{ij}\beta_1 + P_{ij}^2\beta_2 + \alpha_i + \varepsilon_{ij}$$

Results using that model estimate the predicted probability of choosing a plan in 2007 with the same price as that actually selected was 0.025, indicating some sensitivity to price.

We then enriched this basic model, to examine any effect of being the lowest-cost plan (beyond what would be predicted by cost alone) and to analyze differences between the intervention and comparison groups in the sensitivity of plan selection to cost in general and to the lowest-cost plan in particular. The enriched model added an intervention group indicator (Z_i) and interactions of that indicator with predicted cost and predicted cost squared, an indicator for being the lowest-cost plan for that individual (L_{ij}), and the interaction of lowest-cost plan with the intervention group indicator—as well as 2006 baseline plan choice (B_{ij}) and plan fixed effects (θ_j), which improve precision of the estimates and also cause plans selected by fewer than 2 individuals in the sample to drop out of this analysis. That model is based on the indirect utility function in equation (6); all explanatory variables in the model were known at the time of random assignment.

(6) $u_{ij}^* = P_{ij}\beta_1 + P_{ij}^2\beta_2 + L_{ij}\beta_3 + Z_iP_{ij}\beta_4 + Z_iP_{ij}^2\beta_5 + Z_iL_{ij}\beta_6 + B_{ij}\beta_7 + \theta_j + \alpha_i + \varepsilon_{ij}$ We conducted three versions of the analysis based on this equation: plan choices in 2007 for the full sample of 2007 follow-up survey respondents, plan choices in 2007 for the sample of 2008

¹⁷ We also experimented with modeling the effects of price more and less flexibly. The quadratic specification was selected because of its parsimony, fit to the data, and robustness to outliers.

follow-up survey respondents, and plan choices in 2008 for the full sample of 2008 follow-up survey respondents.

The results for all three versions indicate that the intervention group is significantly more sensitive to cost than the comparison group (see Table V). For the intervention group, the estimates for the first version of the analysis imply that a twenty-five percent decrease in predicted cost (say from \$2120 to \$1590, or from the 2007 average cost of the plan chosen in 2006 to roughly the average of the lowest-cost plans in 2007—with marginal effects calculated as the sum of 1000 changes of 51.1 cents each) increased the odds of plan selection by 2.7. That is, it increased the probability of selection from 0.025 to 0.070. If that lower cost plan was also the lowest-cost plan, the estimated odds ratio was 8.2, and the probability of selection further rose to 0.27. In the comparison group, a twenty-five percent decrease in predicted cost increased the probability of plan selection only from 0.025 to 0.040. If the lower cost plan was also the lowest-cost plan, the probability of selection further rose only from 0.040 to 0.062 in the comparison group. The test of whether the coefficient on the interaction term of the differential effect of the lowest-cost plan in the intervention group relative to the comparison group was equal to zero generated a p-value of 0.09. A joint test of whether the coefficients on the cost and cost-squared interactions terms were equal to zero also yielded a p-value of 0.09, while the joint test of whether the coefficients on all three cost interactions terms were zero yielded a p-value of less than 0.005. This evidence is consistent with the effect of changes in the choice environment working through both increased sensitivity to the entire vector of costs for all plans and in particular to the lowest-cost plan.

The estimates for prediction of 2008 plan selection in the third version of the analysis show that the interactions of cost with the intervention were somewhat smaller and the impact of the

lowest-cost plan was somewhat larger than for 2007 plan selection. These results imply that a twenty-five percent decrease in predicted cost in 2007 and being the lowest-cost plan in 2007 increased the estimated odds ratio to 9.9, i.e. the probability of selection in 2008 rose from 0.025 to 0.40. These results are very similar to those from the second version of the analysis, for 2007 plan selection limited to individuals for whom we observe 2008 data, where a twenty-five percent decrease in predicted cost and being the lowest-cost plan increased the probability of selection from 0.025 to 0.38. That is, the 2007 cost information provided in the intervention continued to have an effect of essentially the same magnitude on 2008 plan selection.

VI.E. Impacts on Subgroups

As discussed earlier, our sample was more educated, less satisfied with their baseline plans, and had higher dollar value of potential saving from switching to the lowest-cost plan than a national sample would have had. To examine the sensitivity of our results to these factors, we examined impacts within subgroups by education, plan satisfaction, and potential savings. We also examined the subgroups of people in baseline plans with small and large insurer market shares, and in baseline plans with basic versus enhanced coverage.¹⁸

Education. Our sample is quite highly educated, but estimated impacts for non-college graduates are actually larger than for college graduates, and for both subgroups the null hypothesis of no effect can be rejected at the 5 percent level of significance (see Table VI). These results are consistent with the notion that any limits in comprehending information by less-educated groups are offset by the marginal value of information to these groups.

¹⁸ These analyses were exploratory. Under the null hypothesis of no impact of the intervention on consumer cost, the probability that the maximum t-statistic among the 20 comparisons examined in this section would be 1.96 or higher in absolute value is much greater than 5 percent. The hypothesis testing throughout this paper treats each comparison separately, and does not adjust for multiple comparisons.

Plan Satisfaction. The proportion of our sample that rated their 2006 baseline plan as fair or poor is much higher than that in national samples (including our telephone survey and data from Medicare) who say they were neither satisfied or dissatisfied, somewhat dissatisfied, or very dissatisfied with their plans. Our sample probably had a high proportion dissatisfied with their plans because they were the people willing to volunteer to participate in our study about drug plan choice. The impacts are larger for the more dissatisfied subgroup (although imprecisely estimated), but quite substantial (and statistically significantly different from zero) even for those who rated their 2006 plan good or better—indicating that the results were not driven primarily by the dissatisfaction of participants with their plans.

Potential Savings. The potential savings from switching to the lowest-cost plan was greater in our sample in dollars and similar in percentage terms to a national sample. The impacts in dollars for those with potential savings less than \$400 were much smaller (although statistically significantly different from zero) than for those with greater potential savings. The impacts in relative terms were 0.052 and 0.075 log points for those two groups, respectively. These results suggest that the impact in percentage terms for a sample with nationally representative potential savings would probably have been only slightly smaller than that estimated for the full 2007 Wisconsin follow-up survey sample.

Insurer Market Share. We had initially speculated that individuals with relatively low knowledge of drug plans and drug costs might have placed a high weight on name-recognition and popularity, as potential signals of quality, and had chosen insurers with high enrollment in their plans in 2006. (For example, the plan with the highest national enrollment in 2006 was cobranded by AARP, formerly the American Association of Retired Persons.) We hypothesized that when the intervention made personalized cost information available to individuals in these

plans, they would be relatively more likely to switch plans. We found the opposite result. Individuals in plans with insurer market share of less than 15 percent were more likely to respond to the intervention by switching plans and enjoyed greater cost savings. Ex-post, the results are more consistent with the idea that large market share plans attracted members who highly valued a trusted brand or other non-cost attributes and were relatively less sensitive to personalized cost information.

Benefit Type. The impact on switching by benefit type was essentially the same—about 11 percentage points. The rate of switching from enhanced plans to enhanced plans was similar in the intervention and comparison groups, but the rate of switching from enhanced to basic plans was much higher in the intervention group. The impact on predicted consumer cost in log points was similar for those with basic and enhanced plans at baseline. The higher level of predicted consumer cost among people with enhanced plans at baseline translated into larger impacts in dollars. In sum, the intervention resulted in lower costs among people with both benefit types at baseline. For those with enhanced plans at baseline, those lower costs appear to have been primarily the result of switching to basic plans.

VI.F. Reduction of Comparison Friction and Stated Preferences

To obtain supplemental evidence about how individuals respond to a reduction in comparison friction, we presented seniors with several sets of plan characteristics including those of the plan they had chosen for themselves and asked them to indicate which they preferred. Following a technique developed by Bernartzi and Thaler (2002), our 2008 follow-up survey asked seniors to evaluate the choice between several pairs of unnamed drug plans based on cost measures, plan size, and Medicare quality ratings. In these questions, the cost information was personalized using the information they had provided about medication use in the 2006 baseline survey and was similar to the information that the intervention group had received via the Medicare printout; the enrollment and quality information were new.

When seniors in the comparison group compared their 2007 plan to their 2006 plan (among those who had changed plans during those years), 61 percent did not select their 2007 plan. When seniors who had not chosen the lowest-cost plan in 2007 were asked to compare their 2007 plan to the lowest-cost plan at that time, 63 percent of the comparison group did not select their 2007 plan. This evidence shows how a reduction in comparison friction (that is, providing personalized information about the unnamed plans) shifted stated preferences away from the actual choices, which is consistent with the substantial impact such a reduction had on actual consumer choices as observed in our field experiment. For the intervention group the analogous results were that 52 percent did not select their 2007 plan over their 2006 plan (among those who switched plans), indicating that the shift in stated preferences away from the actual choices was smaller among the intervention group for which comparison friction had been previously reduced during 2006 open enrollment when actual choices were made.

VII. DISCUSSION

We interpret the results of our field experiment to indicate that an intervention which reduced comparison friction had a substantial impact on consumer choices, as it increased the percentage who switched plans from 17 percent to 28 percent and reduced predicted consumer cost by about \$100 per person in our Wisconsin sample. Our examination of the choice environment found that information to facilitate comparisons was accessible at quite low cost (say, by calling 1-800-

Medicare), but that only 18 percent of individuals nationally had ever used personalized cost information. Why didn't people seek out and use the available information?

One potential reason people may not have used this information is that the gains are not as large as they appear. Suppose that individuals face high costs from the act of switching plans. The net gains from switching are then smaller than the cost savings alone. This means that the benefits of undertaking comparative research will be lower. Put simply, high switching costs would make it less valuable to investigate options than our cost savings would imply. However, consider the implications if it were the case that essentially all the potential savings from the intervention were offset by switching costs. Say that 17 percent of individuals would have switched if they had been assigned to the comparison group, and therefore would have saved enough to compensate them for switching costs without the intervention; thus, in this case where potential savings are essentially offset by switching costs there would be no effect on potential savings for these individuals because the intervention did not cause them to switch or increase switching costs. Then the intervention caused only 10 percent of individuals to switch plans. In this case then the overall effect of \$103 per person in potential savings from receiving a letter would have been a combination of no effect on 90 percent of individuals and \$1030 per person caused to switch. Since the act of switching itself could be accomplished in a phone call, this case seems implausible and we conclude such switching costs were very unlikely to have fully offset the potential savings from the intervention. Switching costs less than \$100 per person caused to switch seem more plausible.

Individuals may have expected the costs of understanding the forms and adjusting to the procedures of a new plan to be higher than the costs directly related to the act of switching. That uncertainty is a form of comparison friction that our intervention—which focused on premiums

plus out-of-pocket expenditures—did not reduce. If individuals had greater knowledge of these factors for their current plan and did not have an effective way to learn about them for other plans, then again the net benefits of alternative choice would have been lower and comparative research would have been less likely to be worthwhile. These factors probably contributed to the low use of personalized cost information.

In our view, a key reason people did not seek personalized comparative information was that they had biased expectations about how much they could save from switching plans. We asked participants in the comparison group during our 2007 follow-up interview how much they thought they could save if they had chosen the least expensive plan. Of those who could give an estimate, more than 70 percent gave an underestimate, and the average underestimate was more than \$400. Because they thought the value of comparative research was going to be low, they did not undertake it.

Biased expectations about costs may have combined with confirmation and status-quo biases (the tendency to stick with one's existing opinions and choices), procrastination, limited attention, and small transaction costs to generate high rates of reported satisfaction and low rates of change. Our intervention, while modest, challenged these tendencies by altering price and market perceptions, countering confirmation bias (by showing the savings available), and providing an alternative default (the lowest-cost plan). Our results suggest that the mechanisms underlying the intervention impact increased sensitivity to plan cost in general, and to the lowest-cost plan highlighted in the letter in particular.

VIII. DIRECTIONS FOR FURTHER RESEARCH

This study highlights four areas for further research. One is very concrete work on the design of clear, actionable information about Medicare drug plans or other health insurance coverage choices. Our work shows the potential for information to have an effect, although the study intervention incorporated multiple features including partnership with a trusted hospital, the priming effect of an in-person interview, a behaviorally sensitive letter, the full Medicare printout, and a mailing that both communicated personalized information about potential savings and raised general awareness about the potential for savings and the nature of the variation among plans. Additional work could unbundle these effects, with potential implications for the design of larger scale programs, and could explore the effects of quality as well as cost information. Tools for creating more sophisticated price information could also be developed that would incorporate, for example, forecasts of changes in drug use, rather than simply assume that next year's use will be the same as the previous year's use.

Another area for further research is the role of product and information markets in reducing comparison friction. It is striking that, despite the apparent value of personalized comparative information, few third parties emerged to provide it, or even to highlight its potential value and steer seniors towards Medicare and its local partners. The actual provision of information may have been impeded by CMS regulations that constrained the role of third parties and by the effort involved in working with seniors one-on-one, although third parties with access to drug histories can provide personalized information relatively efficiently. Among the challenges in facilitating an expanded role for third parties would be the need to minimize the potential for plans to capture the market for advice, to respect individual privacy, to provide information that balanced cost and other considerations, and to hold beneficiaries' well-being as the greatest value.

Possibilities to explore could be one-on-one counseling and the ability for beneficiaries and their advisors to manually update an automatically generated drug list.

A third area involves the potential response of insurance firms to broader provision of personalized price information. For example, if the information provided assumed last year's drug use is the same as next year's drug use, then firms would have strong incentives to cut prices on drugs used for short periods and increase prices on drugs used for long periods to encourage individuals to perceive their costs to be lower than they would actually be. In contexts of increased price salience, there would also be greater incentives for firms to cut costs which could lead to lower overall quality of service.

A fourth area for more conceptual research is the interaction between comparison friction and various forms of market failure at both the theoretical and the more practical level. In the case of Medicare drug plans, the private and public optima may differ, and comparison friction may actually counteract market failure by reducing the extent of adverse selection and contributing to the success of the voluntary insurance market. Market functioning could be harmed if all plans with more than basic coverage attract only those for whom those plans are least costly (with these plans then becoming too expensive and being dropped), or if all individuals chose one low-cost provider who then obtained enough market power to keep out new entrants and also set monopolistic prices in future periods.

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TABLE I	
Information on Choices from a Nationally Representative Sample, 20	07

A. Knowledge of plans Knew different plans were better for different people 0.82 Knew one could only change plans during open enrollment 0.74 Knew that not all plans have a deductible 0.37 0.55 Knew plans have different co-payments for generics B. Plan satisfaction and switching At least somewhat satisfied with 2006 plan 0.85 Switched plans from 2006 to 2007 0.10 C. Information source about plans Read at least some of Annual Notice of Change 0.57 Ever reviewed mailings for plan choice 0.53 Ever had in-person contact for plan choice 0.14 Ever had phone contact for plan choice 0.07 Ever used internet for plan choice 0.04 Ever reviewed side-by-side comparison for choice 0.34 Ever reviewed personalized information for choice 0.18

Sample size	348
-------------	-----

Notes. Data are from a phone survey of Medicare beneficiaries over age 65 who were enrolled in stand-alone Medicare prescription drug plans, as described in section IV.A. The Annual Notice of Change is a mailing from the current plan describing changes in that plan for the coming year.

Baseline Characteristics for 2007 Wiscon	onsin Follow-up Survey Respondents				
	Comparison	Intervention	Difference		
	(1)	(2)	(3)		
Female	0.63	0.64	0.01		
			(0.05)		
Married	0.67	0.63	-0.04		
			(0.05)		
High school graduate	0.94	0.95	0.01		
			(0.02)		
College graduate	0.47	0.48	0.01		
			(0.05)		
Post-college graduate	0.20	0.16	-0.05		
			(0.04)		
Age 70+	0.78	0.85	0.07		
			(0.04)		
Age 75+	0.45	0.56	0.11*		
			(0.05)		
4+ Medications	0.61	0.66	0.04		
			(0.05)		
7+ Medications	0.29	0.33	0.05		
			(0.05)		
2006 plan fair or poor	0.26	0.35	0.09*		
			(0.05)		
Predicted consumer cost of baseline plan in 2007	2126	2113	-12		
			(175)		
Cost of lowest-cost plan minus baseline plan in 2007	-520	-533	-13		
			(62)		
Average percentile rank of baseline plan in predicted	37	41	4		
2007 consumer cost			(3)		

 TABLE II

 Baseline Characteristics for 2007 Wisconsin Follow-up Survey Respondents

Notes. Data are from a sample of University of Wisconsin Hospital patients, as described in section IV.D. Sample size is 197 in the comparison group and 209 in intervention group. Standard errors are in parentheses. * = p-value < 0.05

		Switched plan for 2007		$Y^{s} - Y^{b} \qquad \qquad \ln(Y^{s}/Y^{b})$		(X^{s}/Y^{b})
	(1)	(2)	(3)	(4)	(5)	(6)
Intervention	0.115* (0.041)	0.098* (0.041)	-116* (37)	-103* (37)	-0.065* (0.017)	-0.064* (0.017)
Female		-0.023 (0.045)		-34 (41)		0.008 (0.019)
Married		0.107* (0.045)		-72 (41)		-0.013 (0.019)
High school graduate		-0.044 (0.093)		-45 (84)		-0.005 (0.039)
College graduate		0.048 (0.048)		20 (43)		0.001 (0.020)
Post-college graduate		-0.084 (0.062)		37 (56)		0.029 (0.026)
Age 70+		-0.039 (0.060)		41 (54)		0.054* (0.025)
Age 75+		0.079 (0.048)		-49 (43)		-0.032 (0.020)
4+ Medications		-0.054 (0.050)		-23 (45)		-0.000 (0.021)
7+ Medications		0.116* (0.052)		-112* (47)		-0.008 (0.022)
2006 plan fair or poor		0.097* (0.045)		-55 (41)		-0.010 (0.019)

TABLE III Regression Coefficients In Models of Plan Switching and Consumer Cost for 2007 Wisconsin Follow-up Survey Respondents

Notes. Estimates are from linear regression based on equation (4), as described in the text. Y^s is the 2007 predicted consumer cost of the plan selected for 2007. Y^b is that cost for the baseline plan that had been selected in 2006. Sample size is 406. Standard errors are in parentheses. * = p-value < 0.05

A. Consumer costs in dollars	
Predicted 2007 consumer cost (2007 plan)	-111*
	(48)
Actual 2007 consumer cost (2007 plan)	-137
	(217)
Predicted 2008 consumer cost (2008 plan)	-93
	(169)
Predicted 2009 consumer cost (2008 plan)	-159
	(170)
B. Quality	
Proportion dissatisfied with quality, non-cost features	0.032
	(0.033)
Proportion rating overall plan as fair or poor	-0.026
	(0.041)

 TABLE IV

 Intervention Impacts for 2008 Wisconsin Follow-up Survey Respondents, by Outcome

Notes. All estimates are of the coefficient on the intervention indicator from linear regression based on equation (4), as described in the text, using the same vector of covariates as in Table III. Costs are differences from Y^b , the predicted 2007 consumer cost of the 2006 baseline plan. Predicted 2007 costs are based on drugs reported in the baseline 2006 survey. Predicted 2008 and 2009 costs are based on drugs reported in the 2008 follow-up survey. Sample size is 285 in rows 1-3, 241 in row 4, and 302 in rows 5-6. Standard errors are in parentheses. * = p-value < 0.05

Conditional Logit Analysis of Plan Selection for	t Analysis of Plan Selection for Wisconsin Follow-up Survey Respondents				
	2007 Plan 2007 Plan 2008 Pla				
	(1)	(2) -1.78	(3)		
β_1 : 2007 consumer cost/1,000	-1.18	-1.78	-1.63		
	(0.75)	(1.05)	(0.88)		
β_2 : 2007 (consumer cost/10,000) ²	7.69	13.85	14.52		
	(6.99)	(8.90)	(7.63)		
β ₃ : 2007 lowest-cost plan	0.46	0.34	-0.46		
	(0.60)	(0.80)	(0.73)		
β_4 : 2007 consumer cost/1,000 * intervention	-1.46	-1.10	0.11		
	(0.94)	(1.25)	(1.11)		
β_5 : 2007 (consumer cost/10,000) ² * intervention	6.92	4.91	-10.33		
	(8.53)	(10.68)	(10.95)		
β_6 : 2007 lowest-cost plan * intervention	1.16	1.71	3.04*		
	(0.68)	(0.89)	(0.82)		
Ν	12719	7101	7101		
Sample limited to 2008 follow-up survey respondents	No	Yes	Yes		
p-value on H_0 : $\beta_4 = \beta_5 = 0$	0.09	0.41	0.15		
p-value on H ₀ : $\beta_4 = \beta_5 = \beta_6 = 0$	0.00	0.02	0.00		

 TABLE V

 Conditional Logit Analysis of Plan Selection for Wisconsin Follow-up Survey Respondents

Notes. Conditional logit models estimated with individual fixed effects, as in equation (6). Additional coefficients not shown in the table were estimated for the 2006 choice and for each plan; the estimation sample is therefore limited to plans selected by at least 2 individuals in the sample and to individuals choosing a plan selected by at least two individuals. 7 observations with predicted consumer costs > \$20k were dropped. Column 1 uses the 2007 follow-up survey respondents, with 401 individuals, 32 plans, and 113 missing cost observations. Columns 2 and 3 use the 2008 follow-up survey respondents, with 265 individuals, 27 plans, and 54 missing cost observations. Standard errors are in parentheses. * = p-value < 0.05.

	Switching	nnahahilitu	Impost on r	Impact on predicted cost		
	Comparison	probability Intervention	Dollars	Log points	<u>N</u>	
	(1)	(2)	(3)	(4)	(5)	
	(1)	(2)	(3)	(+)	(3)	
A. Education						
Not college graduate	0.154	0.284~	-129*	-0.064*	213	
0.0			(64)	(0.025)		
College graduate	0.183	0.280	-91*	-0.068*	193	
			(37)	(0.024)		
B. Satisfaction rating of 2006 plan	0.005	0.070	1.60	0.115*	10.4	
Fair, poor, or unknown	0.235	0.370	-168	-0.115*	124	
~	0.4.4.4	A A A F	(103)	(0.038)	• • •	
Good, very good, or excellent	0.144	0.235~	-73*	-0.046*	282	
			(32)	(0.019)		
C. Dollar potential savings						
≤ \$400	0.145	0.217	-32*	-0.052*	216	
_ +	01110	01217	(16)	(0.023)	-10	
> \$400	0.195	0.350~	-176*	-0.075*	190	
· • • •	0.170	0.000	(77)	(0.026)	170	
D. Insurer share of sample in 2006						
≤ 0.15	0.141	0.333~	-265*	-0.154*	142	
			(96)	(0.031)		
> 0.15	0.180	0.252	-28	-0.031	264	
			(28)	(0.020)		
E Bonafit type in 2006 hassiling						
E. Benefit type in 2006 baseline	0.144	0.257~	-85	-0.061*	224	
Basic	0.144	0.237~			224	
Enhanced	0.198	0.312	(44) -139*	(0.026) -0.078*	182	
	0.190	0.312	(65)	(0.022)	102	
(with or without gap coverage)			(03)	(0.022)		

 TABLE VI

 Impacts for 2007 Wisconsin Follow-up Survey Respondents, by Subgroup

Notes. All subgroups are defined on characteristics known prior to random assignment. Dollar potential savings = predicted 2007 consumer cost of 2006 plan - predicted consumer cost of least expensive plan in 2007. \sim = p-value < 0.05 on difference between columns 1 and 2. Columns 3 and 4 are estimated with regression adjustment based on equation (4) using the models in Table III (columns 4 and 6, respectively) estimated separately for each subgroup. Standard errors are in parentheses.* = p-value < 0.05.

Appendix

for

COMPARISON FRICTION:

EXPERIMENTAL EVIDENCE FROM MEDICARE DRUG PLANS

Jeffrey R. Kling, Sendhil Mullainathan, Eldar Shafir, Lee Vermeulen, and Marian V. Wrobel

August 2011

This appendix consists of three parts:

- A. Potential cost savings from plan switching by benefit type
- B. Intervention impacts assuming some participants were not affected
- C. Examples of materials used in the intervention

A. Potential cost savings from plan switching by benefit type

Using data from the baseline survey on drugs taken in 2006, the average predicted consumer savings from switching to the lowest-cost plan in 2007 of any benefit type was \$527. Those savings from switching to the lowest-cost plan of the same benefit type as the individual had in 2006 ranged from \$386 to \$464 (see Table A-1).

We also examined information on drugs actually taken in 2007 from our 2008 follow-up survey. The average actual savings from switching to the lowest-cost plan in 2007 of any benefit type was \$487. Those savings from switching to the lowest-cost plan of the same benefit type as the individual had in 2006 ranged from \$337 to \$392.

The average consumer cost among all 2007 basic plans was \$2334. In comparison to the lowest-cost basic plan for each individual, the 27 other basic plans had predicted consumer costs \$705 greater on average.

	Dollars		Percenti	le Rank
	Potential (1)	Actual (2)	Potential (3)	Actual (4)
Among Basic Plans	-386	-337	38	35
Among Enhanced Plans Without Gap Coverage	-429	-391	43	37
Among Enhanced Plans With Gap Coverage	-464	-392	50	49
Among All Plans	-527	-487	39	37

TABLE A-1. Average Difference in Cost between Baseline Plan and Lowest-Cost 2007 Plan

Notes. The baseline plan is the plan in which the individual was enrolled in 2006. The dollar difference is the cost in 2007 of lowest-cost plan minus that of the baseline plan. The percentile rank the rank of the cost of the lowest-cost plan minus the baseline plan (where zero provided the most savings and 100 provided the least savings). The sample for rows 1-3 is limited to plans of the same type as the baseline plan. The sample size for potential costs was 224, 73, 109, and 406 for rows 1-4 respectively in columns 1 and 3; for actual costs it was 155, 58, 85 and 298 for those rows in columns 2 and 4.

B. Intervention impacts assuming some participants were not affected

The intervention impacts on predicted consumer cost discussed in the main text are intent-totreat estimates which compare the outcomes for all members of the intervention group to those of the comparison group. However, the change in predicted consumer cost is zero by definition for individuals who did not switch plans. Thus, the intent-to-treat estimates average together impacts on those potentially affected by the intervention with a large proportion of zeros for those not affected. This section explores how different assumptions about the proportion of the sample not affected scales up the point estimates and standard errors of those who were potentially affected.

Define A as an indicator of being potentially affected by the intervention, where A involves the counterfactual and cannot be directly observed. Define D as an observed indicator for switching plans, and Z as an indicator for assignment to the intervention group. Define Y as the difference in predicted consumer cost of the plan selected for 2007 and the baseline plan in 2006, Y_1 as the potential outcome if an individual were assigned to the intervention group, Y_0 as the potential outcome if an individual were assigned to the comparison group. The causal effect of the intervention is then Y_1 - Y_0 .

There was a causal effect for any individual who would have chosen a plan with a different predicted consumer cost in the intervention group than in the comparison group. These situations included having the intervention cause someone to switch to a lower cost plan ($Y_1 < 0$; $Y_0=0$), having the intervention cause someone who was going to choose a more expensive plan to not switch ($Y_1=0$; $Y_0>0$), and other cases (anytime $Y_1 \neq Y_0$). A special case was when someone would not switch plans regardless of the intervention, so there was no effect on cost. The upper bound on the probability of this special case occurred when everyone who switched plans in one group would have switched if assigned to the other group (1- max{E[D | Z=1], E[D | Z=0]}). The lower bound on the probability of this special case occurred when no one who switched plans in one group would have switched if assigned to the other group (1- {E[D | Z=1] + E[D | Z=0]}). Intuitively, we can use the lower bound on the fraction of zeros included in the estimate of the average cost change for the entire intervention group versus the comparison group to calculate a lower bound on the average cost change for those who potentially were affected by the intervention. This bound is based on the derivation in equation (B1).¹⁹

(B1)
$$E[Y_1 | Z=1] - E[Y_0 | Z=0]$$

- $= E[Y_1 Y_0]$
- $= E[Y_1-Y_0 | A=1]Pr(A=1) + E[Y_1-Y_0 | A=0]Pr(A=0)$
- = $E[Y_1-Y_0 | A=1]Pr(A=1) + 0$
- $\leq E[Y_1-Y_0 | A=1] \{ E[D | Z=1] + E[D | Z=0] \}$

This approach is similar to that used by Imbens and Angrist (1994) to estimate a local average treatment effect (LATE), where those who did not comply and take up the treatment offer are assumed to have been unaffected. However, the approach used here is less restrictive. The exclusion restriction required for LATE, but not needed for (B1), is that people who would have switched plans even without the intervention would not be affected; relaxing this assumption is sometimes described as allowing effects on "always-takers." LATE also involves an assumption of monotonicity, not needed for (B1), where the intervention only encourages switching; relaxation of this assumption is sometimes referred to as allowing "defiers." We can now calculate an expression based for a lower bound on the average cost change for those who were potentially affected by the intervention, shown in equation (B2).

¹⁹ The first line of equation uses the definition of potential outcomes. The second line uses the independence of potential outcomes from randomly assigned groups. The third line uses the definition of conditional expectation. The fourth line uses the definition of A, where Y_1 - Y_0 = 0 when A=0. The fifth line uses the lower bound described in the text, where Pr(A=0) = 1- $Pr(A=1) \le 1$ - {E[D | Z=1] + E[D | Z=0]}.

(B2)
$$E[Y_1-Y_0|A=1] \ge \{E[Y|Z=1] - E[Y|Z=0]\} / \{E[D|Z=1] + E[D|Z=0]\}$$

In this paper's application, the lower bound point estimates and standard errors simply rescale the intent-to-treat estimates by $1/{E[D | Z=1] + E[D | Z=0]}$, or 2.2 (see Table B-1). There is a small amount of negative covariance between the estimation of average cost differences and switching rates, and accounting for this slightly reduces the standard errors; for simplicity, this adjustment is not included in the results.

Our intuition is that the exclusion restriction does not hold in this application but monotonicity probably does. The exclusion restriction would be violated if those in the comparison group who would have switched without the intervention nevertheless had their predicted consumer cost affected by the intervention. Monotonicity would be violated if the intervention caused some people to not switch who would have otherwise switched. If we did not impose the exclusion restriction but assume monotonicity holds—that is, allow effects on "always-takers" but assume no "defiers"—then then we would obtain treatment-on-treated results from rescaling by 1/E[D | Z=1] instead of $1/{E[D | Z=1] + E[D | Z=0]}$, as in equation (B3).

(B3)
$$E[Y_1-Y_0|A=1] = \{E[Y | Z=1] - E[Y | Z=0]\} / E[D | Z=1]$$

Estimates of treatment-on-treated effects based on (B3) generate point estimates about 3.6 times larger than the lower bound on the potentially affected group based on (B2).

The regression-adjusted estimate of the treatment-on-treated effect for the full sample is \$369 (see Table B-1). That turns out to be very similar to the non-experimental estimate that compares the savings of those who switched plans in the intervention group (who had potential savings that averaged \$469) to those who switched plans in the comparison group (who had potential savings

that averaged \$97), where the difference is \$372. Because the potential savings for non-switchers is zero by construction, if there were no regression adjustment and the switching rates in the two groups were the same then treatment-on-treated estimate would exactly equal the difference between switchers in potential savings; in this application, the regression adjustment and lower switching rate in the comparison group roughly offset each other.

Reference

Imbens, Guido. W., and Joshua. D. Angrist, "Identification and Estimation of Local Average Treatment Effects," *Econometrica*, 62:2 (March 1994), 467-476.

	Lower bound impact on predicted cost		Treatment	-On-Treated	
			impact on p	Ν	
	Dollars	Log points	Dollars	Log points	
	(1)	(2)	(3)	(4)	(5)
A. Education					
Not college graduate	-295*	-0.146*	-454*	-0.224*	213
	(147)	(0.057)	(226)	(0.088)	
College graduate	-196*	-0.146*	-325*	-0.242*	193
	(80)	(0.053)	(132)	(0.087)	
B. Satisfaction rating of 2006 plan					
Fair, poor, or unknown	-278	-0.189*	-455	-0.310*	124
	(171)	(0.062)	(279)	(0.102)	
Good, very good, or excellent	-192*	-0.122*	-310*	-0.196*	282
	(85)	(0.050)	(137)	(0.081)	
C. Dollar potential savings					
\leq \$400	-88*	-0.143*	-148*	-0.238*	216
	(45)	(0.064)	(75)	(0.106)	
> \$400	-323*	-0.137*	-503*	-0.214*	190
	(141)	(0.048)	(220)	(0.075)	
D. Insurer share of sample in 2006					
≤ 0.15	-559*	-0.324*	-795*	-0.461*	142
	(203)	(0.065)	(289)	(0.093)	
> 0.15	-65	-0.073	-112	-0.125	264
	(64)	(0.047)	(110)	(0.081)	
E. Benefit type in 2006 baseline					
Basic	-211	-0.152*	-330	-0.237*	224
	(111)	(0.065)	(173)	(0.101)	
Enhanced	-272*	-0.153*	-445*	-0.249*	182
(with or without Gap Coverage)	(127)	(0.044)	(208)	(0.072)	
F. All					
Full sample	-229*	-0.143*	-365*	-0.228*	406
_	(83)	(0.038)	(132)	(0.061)	

TABLE B-1. Intervention Impacts in the Wisconsin 2007 Follow-up Survey
Assuming Some Participants Were Not Affected by Subgroups

Notes. Estimates are with regression adjustment using the models in Table III, columns 4 and 6, estimated separately for each subgroup, and modified as follows: columns (1) and (2) are based on equation (B2), and multiply the estimates by 1/(0.282+0.168); columns (3) and (4) are based on equation (B3), and multiply the estimates by 1/0.282. All subgroups are defined on characteristics known prior to random assignment. Dollar potential savings = predicted 2007 consumer cost of 2006 plan - predicted consumer cost of least expensive plan in 2007. Standard errors are in parentheses. * = p-value < 0.05.

C. Examples of materials used in the intervention

- Exhibit C1: Comparison Group Letter
- Exhibit C2: Intervention Group Letter
- Exhibit C3: Booklet provided *The New Medicare Prescription Drug Coverage: Using the Medicare Prescription Drug Plan Finder*
- Exhibit C4: Medicare Prescription Drug Plan Finder Printout



Exhibit C1: Comparison Group Letter

December 2006

Dear UWHC Patient:

Thank you for participating in our Medicare Part D prescription drug plan study. I hope the information you received on the phone recently was helpful to you.

As you were told, you can find additional information regarding the plans available to you by accessing the Medicare Prescription Drug Plan Finder web site at: <u>http://www.medicare.gov/MPDPF/Public/Include/DataSection/Questions/SearchOptions.asp</u>.

You can use the enclosed document from AARP to assist you in using the web site.

Please remember that later this coming spring, we will call again to find out what plan you chose and how satisfied you are with your choice; that call will take about 10 minutes.

If you have any questions or concerns about this study or the information you have received, please feel free to contact me at 608-262-7537. Thank you for your kind consideration!

Sincerely,

Lee Vermeulen, R.Ph., M.S., FCCP Director, Center for Drug Policy UW Hospital and Clinics Clinical Associate Professor UW- Madison School of Pharmacy



Exhibit C2: Intervention Group Letter

December 2006

«First_Name» «Last_Name» «Street» «City», «State» «Zip»

Dear UWHC Patient:

Thank you for participating in our Medicare Part D prescription drug plan study. I hope the information you received on the phone recently was helpful to you.

Please find enclosed a summary of the information that you received during the interview. As you can see from the Medicare web site, you may have an opportunity to save on your prescription drug costs by changing plans for 2007.

The plan you reported being in for 2006:	«M_06_PLAN»
The estimated cost of that plan for 2007:	\$«ANNUALOF_PTS_CURRENT_PLAN_06»
The lowest cost plan available to you for 2007:	«LEAST_EXPENSIVE_PLAN_07»
The estimated cost of that plan for 2007:	\$«ANNUAL_07»
Your potential savings for 2007 if you choose	\$«POTENTIAL_SAVINGS»
the lowest cost plan:	
Comments:	«COMMENTS_TO_SUBJECTS»

Note that even if you have already chosen a plan for 2007, you can still change your mind and choose a different plan until December 31, 2006! If you do choose to change plans, you can access the Medicare web site at:

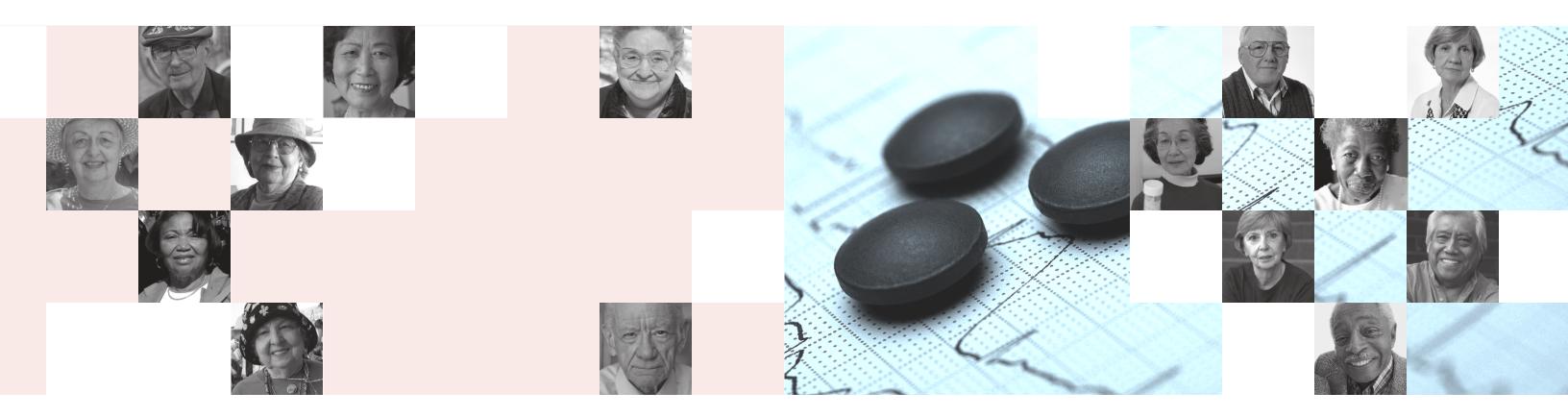
http://www.medicare.gov/MPDPF/Public/Include/DataSection/Questions/SearchOptions.asp

Please remember that later this coming spring, we will call again to find out what plan you chose and how satisfied you are with your choice; that call will take about 10 minutes. If you have any questions or concerns about this study or the information you have received, please feel free to contact me at 608-262-7537. Thank you for your kind consideration!

Sincerely,

Lee Vermeulen, R.Ph., M.S., FCCP Director, Center for Drug Policy UW Hospital and Clinics Clinical Associate Professor UW- Madison School of Pharmacy

The New Medicare Prescription Drug Coverage Using the Medicare Prescription Drug Plan Finder





For more information about Medicare's prescription drug coverage, visit AARP's website at www.aarp.org/medicarerx, or call 1-888-OUR-AARP (1-888-687-2277).



The power to make it better.™

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List of Medications for Medicare Plan Finder Search

11

Introduction

In this booklet you will find information about how to use the tools available to help you make a decision about a Medicare prescription drug plan. For those who have access to a computer and/or assistance from a computer user, Medicare has developed the Medicare Prescription Drug Plan Finder. In this booklet you will find a step-by-step guide on how to do a "General Search" of prescription drug plans. There are various other kinds of searches that can be done, but our goal was to familiarize you with the tool itself—enabling you to move on to other searches yourself later. For those who do not have access to a computer and/or who are uncomfortable working with this kind of a tool, there is also information about how to enroll in a Medicare prescription drug plan using the telephone. Either way, we hope you will explore the new Medicare prescription drug coverage options available to you—to see if it will help you save on your drug costs. Finally, if you need more information before making a final decision, see the "Other resource information" column on page 9.

Using the Medicare **Prescription Drug** Plan Finder

Preparing for Your Drug Plan Search

Before using the Medicare Prescription Drug Plan Finder, you should collect the following information (see worksheet on page 11):

1. A List of the Prescription Drugs You Take:

- Name of each drug*
- Dosage or strength of the drug you take (e.g., 20 mg)
- Number of times you take each drug each day or per week

*Prescription drugs are dispensed in a variety of ways—tablets, capsules, patches, time-released, etc. Often this information is entered in coded language on your pill container. The names of some drugs may also be abbreviated because they are so long. Thus, your pill container may not give you everything you need for your drug list. It therefore might be a good idea to talk with your pharmacist when making your list—and if you get all your drugs from one place, your pharmacist may be willing to provide you with a printout of your drug list.

2. Information about Your Current Insurance Coverage (if any) such as a Medicare supplement, retiree coverage, Medicare Advantage (managed care) plan, VA, TRICARE, or FEHB.

For Help on the Computer:

- Ask a family member or friend to help you.
- Contact the State Health Insurance Assistance Program (SHIP). Call 1-800-633-4227 to learn the number for your state's SHIP. SHIP trains volunteers to help people with their Medicare issues. Most are computer savvy. Or,

Current coverage

If you have VA, TRICARE, or FEHB, your drug coverage may already be as good as the new Medicare Rx coverage. Your medical plan provider should have sent you information by now about whether your coverage is as good as Medicare's.

If you were in a Medicare Advantage Plan with prescription drug coverage prior to 2006, your plan should have informed you that you would be enrolled in their Medicare prescription drug plan.

If you had retiree drug coverage through a former employer or union, or a Medicare supplemental plan with drug benefits, prior to 2006, you should have been notified by your plan provider if your coverage is as good as Medicare's drug coverage.

Tips on using the **Medicare Speech Automated System**

It will ask you a series of questions about what kind of information you need. The basic questions you will need to answer are as follows:

- 1. If you are calling about the new Medicare drug coverage, say: "Drug Coverage."
- 2. For information on how the new prescription drug plans work, say: "Plan Choices."
- 3. Do you have your Medicare Card?
- 4. Please tell me your Medicare Claim Number, including any letters (it's on your Medicare card).
- 5. May I have your last name, without spelling it?
- 6. What is your date of birth?
- 7. What was the starting date of your Medicare coverage? (This is on your Medicare card.)
- 8. Our records show that you are... (enrolled or not enrolled in Medicare prescription drug plan).
- 9. If would like to speak with an agent, you will need your drug list and Medicare card. Say: "Agent." This will bring a customer service representative from Medicare on to the line.

• Call your local Area Agency on Aging at 1-800-677-**1116** to learn which office is the closest to you.

If You Don't Have Computer Access:

• Call MEDICARE at 1-800-633-4227. Customer service representatives will walk you through the Plan Finder. If you request it, they will send you a "Customized Print on Demand" booklet of the prescription drug plans that might work best for you.

Tips for calling Medicare:

- Medicare's toll-free number is currently available 24 hours a day, 7 days a week. Try to call during non-peak hours such as evenings or weekends, and try to avoid Monday mornings.
- You will need to have your list of drugs and dosages, your Medicare card, your date of birth, and the date you originally enrolled in Medicare (the date is on your Medicare card) available when you call.
- Write down any specific questions you have before calling, and check your questions off as they are answered when you call.
- When you make the call, you may first get a "Speech Automated System" (a computer) that will instruct you to say your answers. If you decide to use this system, you will need to speak slowly, distinctly, and loudly enough for the computer to understand you. Also, try to minimize any background noises that might make it hard for you and the computer to hear.
- If the response from the Medicare automated system is that you are not enrolled in a plan, it will ask you to please check back later because Medicare is updating its records daily. So if you have enrolled already, wait about a week and call Medicare back. If you haven't enrolled yet, and want to get Medicare drug coverage, you will need to make a choice and enroll in a plan. Say: "Enroll."

• If you dislike or have difficulty using a speech automated system, you can press zero (0) on your telephone to bypass the automated system. This may or may not get you to a customer service operator right away-depending upon how busy they are. It's important to know, however, that calls are taken in the order they are received—so hanging up and calling back only puts you at the back of the line!

If You Do Have Computer Access:

Once you have gathered your drug information, you are ready to begin your prescription drug plan search, using the Medicare Prescription Drug Plan Finder.

To keep your initial search simple, this tool focuses on "Stand Alone" prescription drug plans associated with the traditional Medicare fee-for-service program. By working through a "General Search," you will become better acquainted with the information available through the Plan Finder.

To start your prescription drug plan search: 1. On the computer, go to the Internet, type: www.medicare.gov, hit enter, and log onto the Medicare website.

- 2. Click on Compare Medicare Prescription Drug Plans.
- 3. Scroll down to where you see (in blue) Where Would You Like to Begin? Then, where you see, Find a Medicare Prescription Drug Plan, click on the orange arrow to the right.
- 4. Scroll down to the lower half of the page until you see a button that says General Search and click on that button.
- 5. Enter your **Zip Code** in the box that is provided. **Do** not hit enter.

Note: In some cases, when you enter your zip code, you will get another box that asks you to select your county. Click on the drop down arrow and select your county, then click on Continue.

Extra Help for those with limited income

Extra Help is available if your income and assets are below these figures:

- Income eligibility for Extra Help is \$14,355 (single) or \$19,245 (couple).*
- Asset eligibility is \$11,500 (single) or \$23,000 (couple).

If it looks like your income is less than these figures, even if you have some doubts, it is a good idea to apply for this benefit, since it could provide:

- Unlimited drug coverage (no coverage gap).
- Greatly reduced (or even zero) monthly premiums.
- Greatly reduced (or even zero) annual deductible.
- Co-pays of \$1 to \$5 or 15% of the cost of each drug depending on income and asset levels.
- *Income figures will go up for 2006.

Generic check box

When you enter a drug name, the computer automatically checks to see if a generic drug is available.

Generics will automatically come up on your list of drugs

В.

Review you	ır drug list					
	Drug Name					
	Biaxin	CLARI				
	Se lower cost generic	drugs v				
	Add Additional Drugs					

—unless you uncheck the box first (below).

3. Review y	our drug list	
	Drug Name	
	Biaxin	CLARI
	Use lower cost gen	ieric drugs v
	Add Additional D	rugs

Because generics are usually less expensive than their brand name equivalents, the computer will also factor the generics listed into the plan costs. Therefore, the cost quotes might be misleading, if you intend to stay with a brand name as opposed to a generic.

You will have another chance to do this kind of cost comparison later, so we suggest you uncheck the box at this point in your search.

- 6. On the same page, scroll down and click on the box that best describes your Current Prescription Coverage. If you don't know, click "None of the Above."
- 7. On that same page, scroll down and select "yes" or "no" to the question Eligible or Qualify For Additional Help, then scroll down and click the Continue button. (If you are not sure if you will qualify for the Extra Help, check the eligibility information in the box in the shaded area on page 3.)
- 8. This will bring up the next window **Decide on Your** Plan Options. At the bottom of the page, click on Choose a Drug Plan Type.
- 9. Near the bottom of the page, click on the gray button Search for Medicare Prescription Drug Plans.
- 10. Scroll down to Enter Your Medications, and click on the gray button. Using the list of drugs you made earlier, type the name of your drug in the box that appears, and click Search for Drug. Scroll down to **Review Your Drug List** to see that drug name has been entered. If you have more than one drug to enter, click on Add Additional Drugs and enter the next drug. Repeat for each drug used, clicking **Add Selected Drug** to My List each time after typing in the name of your drug. Don't panic when the screen appears to go blank—the computer is just doing its work.

Note 1: If you type the drug name incorrectly or just type in the first few letters, the program will give you a drop-down screen with a list of drugs to choose from that are close to what you've entered—select the correct drug and then click on Add the Drug to My List. You can also enter just the first letter of the drug name (such as "L" for Lovastatin) and you will get a complete listing of all drugs that start with that letter. If you find your drug on this list, you then need to select it and click on Add the Drug to My List.

Note 2: When all your drugs are listed, click on the little box below the list. This will remove the check mark (see explanation in the shaded box) from the box that says: "Use lower cost generics when available."

Note 3: If a particular drug does not appear on the Medicare Plan Finder list, it is probably not covered by Medicare and you will have to pay for the drug yourself-unless you qualify for another program such as Medicaid or a state pharmacy assistance program

11. When you have completed entering your list of medications, click on **Continue with Selected Drugs**.

- 12. Next click on Change/Update My Drug Dosage. This is one of the most important steps in the whole **process!** You will get a more precise plan comparison if you take the time to enter exact dosages and quantities. For example, if two pills are taken per day, change the 30-day quantity to 60. To change the dosage, click the drop-down arrow to the right of the drug name, and then click Update Dosage/Quantity to make sure all your changes have been registered. You next need to click on Continue with Selected Drugs.
- 13. Scroll down to Choose How You Want to View Your Plan. Click on Continue to Plan List rather than Select My Preferred Pharmacy. You will have another chance to do this later. You will then see a screen titled **Find a** Medicare Prescription Drug Plan. Scroll down to the Prescription Drug Plan Comparison multi-colored chart that gives you plans for the zip code you entered, starting with the one that is estimated to be least expensive overall calculated for 30-day supplies from local "preferred" pharmacies (those that are in the plan's network). You still need to compare plans, however, because there are other factors to think about.

Note: In order to view the information about the various plans, you will click on the words in the column that are underlined (Plan Information, Plan Name, # of Pharmacies, etc.). This will bring up more detailed information about each plan. In order to move back and forth between these columns, you should scroll down to the bottom of the screen and click on the buttons that may say Return to Personalized Search, Close Window, or Return to Previous Page. You can also use your "back" button on your computer in most cases to move back and forth between the various windows.

Getting the answers vou need from the Medicare Rx Plan Finder

What will my detailed cost be under this plan? See View Cost Details.

What are my 90-day mail order options? See View Cost Details.

Are there ways of paying less under a prescription drug plan? See Lower My Cost Share.

How can I tell whether all my drugs are covered? See View Cost Details.

How can I tell what my drug costs will be in the "gap in coverage?" See View Cost Details/Show Details or Plan Name/Show Details (if it says Hide Details click on it to bring up the Show Details).

Note: A Double Asterisk (**)to the right of a drug's name (in **Show Details**) means it is not part of a plan's preferred drug list, or formulary, and therefore will be more costly to you.

How can I tell whether I can get my prescription drugs in other parts of the country under this plan, if I'm traveling? See View Plan Notes.

Which pharmacies can I go to under this plan? Click on the number in the **# of** Pharmacies Column.

14. Scroll down about halfway until you see a multi-colored chart. Look for the heading More About this Plan.

Then, starting in the **Plan Information** column on the topmost plan, click on the Select Below arrow that will provide a drop-down menu with the following selections:

- Enroll in this Plan
- View Cost Details: This gives you what you need to compare plans properly. Some of the things it shows include: (see box to the right also)
 - > An estimate of your total annual costs (including premiums, deductibles and co-pays) for all of 2006.
 - > Your fixed costs —monthly premiums and annual deductible (if any).
 - > Your co-pays in the initial coverage period.
 - > If or when you might reach the "gap in coverage" and what your drugs will cost you when you do.
- Lower My Cost Share: This is where you can check to see what savings might be available through the use of a similar drug or generic equivalent.
- View Notes: This will give you information about whether the plan is regional or national, as well as information about where you might obtain your drugs (pharmacy networks, etc.)
- 15. In the **Plan Name Column**, click on the name of the plan to get detailed information about your specific drug costs, including:
 - Your monthly premium and deductible (if any).
 - Your specific co-pay or co-insurance for each drug you listed earlier.
 - The "tier" your drug(s) are in for that plan (remember tiers won't necessarily be the same for all plans).

- The cost for a 90-day mail order supply.
- If and when your costs are estimated to reach the "gap in coverage" or catastrophic level, and your estimated monthly cost during the "gap" period.
- 16. In the Select to Compare column, click in the boxes to the left of the plan names (up to three at a time) you are interested in, and then scroll down and click the button that says **Compare Three Plans**. Repeat for more comparisons.
- 17. In the **# of Pharmacies column**, click on the **#** to get a list of the pharmacies that are preferred by that particular plan.

By now you should be getting a feel for how the Medicare Prescription Drug Plan Finder works. There is a lot of information available at your fingertips—but it takes a little practice to become comfortable using it—so take your time and don't be afraid to explore some of the other search possibilities available. There will be times when "traffic" is heavy on the Plan Finder. You may have to try at another non-peak time, but don't get discouraged. As the "traffic jam" begins to break up, you'll have access when vou want it.

Some other things to be aware of when using the Medicare Prescription Drug Plan Finder:

- 1. **Restrictions on certain drugs**—Watch for a single asterisk (*) to the right of a drug name in the "View Cost Details" page. It means there are special rules that apply, such as:
 - **Prior Authorization**: This means that you or your doctor must obtain the plan's approval before it will cover a particular drug (often a high-priced one). The physician generally has to document why this specific medication is needed.
 - **Step Therapy**: This is a variation of prior authorization. It requires the physician to use a similar but

Exploring similar drugs to treat a particular condition

AARP has developed an online consumer guide to help you find the most effective and affordable drugs. The guide provides unbiased information on drug safety, effectiveness, and cost. It's based upon an independent review and assessment of the available medical evidence.

The guide shows how the most expensive drugs are not necessarily the best, and that consumers willing to consult with their doctor or pharmacist, can often find similar safe and effective lower cost drugs. To learn more about AARP's prescription drug consumer's guide, visit the AARP website listed below. www.aarp.org/ comparedrugs

Other types of searches

Formulary Search:

Each plan has a formulary, or its preferred list of drugs.

To do a Formulary Search, go to www.medicare.gov, click on Formulary Search. Enter your state, then your list of drugs. When the list of plans is displayed, click on the name of each plan to see in which tier each of your drugs falls. This won't give you copays, however, so you'll need to go to "View Cost Details" (see page 6) for that.

Personalized Search:

Start at www.medicare.gov again. Click on **Compare Medicare Prescription Drug** Plans. Click on Find a Medicare Drug Plan. Enter your personal information in the boxes, then click on **Personalized Search**. From there, follow the same basic steps as for the "General Search."

. Personalized plan search	
Personal information	
Medicare claim number:	
Last name:	
Date of birth:	Month 🗘Day 🗘Year 🗘
Effective date for Medicare Part A or B:	Hospital Part A
	Month 🛟Year 🛟
ZIP code:	

less expensive "preferred" drug to treat a condition before being allowed to use the one originally prescribed.

- 2. Quantity Limits: This does NOT mean that your supply of drugs will be cut off after a certain timeor restricted to a particular number of prescriptions per year. It does mean that your doctor must follow your plan's guidelines when prescribing drugs for a current condition. If, for instance, your doctor writes a prescription that falls outside your drug plan's quantity limit, he or she must get prior authorization from your plan before the plan will agree to pay for the prescription. For example, prescriptions for sleeping pills need to be monitored very closely by your doctor and therefore most will have a quantity limit associated with them. In other words, if the normal course of treatment calls for a 10-day supply of pills, then your doctor can't write the prescription for more than that—unless he or she has prior approval from the drug plan to do SO.
- 3. **Co-payment/Coinsurance**: This is the amount you pay for each prescription after you have paid the deductible. In some plans, you pay the same co-payment (a set amount) or coinsurance (a percentage of the cost) for any prescription. In other plans, there might be different levels or "tiers," with different costs, depending on which tier a particular drug is assigned. Some plans may have co-payments for some drugs and coinsurance for others.
- "Tiered" co-payment levels: Typically, less expensive drugs are assigned to lower tiers, with lower co-pay amounts. Brand-name drugs may be further divided into "preferred" and "non-preferred" tiers as well. Further, some plans may dedicate another tier for less common and/or very expensive drugs. With these kinds of variations, it's a good idea to compare not only the monthly premium and annual deductible-but also co-payments-when calculating your total costs.

5. **Premiums and Deductibles**: Keep in mind that some plans will not require a premium or a deductible-but you still need to compare total costs for the plan as a whole to see which plan makes the most dollar sense. While your premiums and deductible (if any) cannot increase in the calendar year, your co-pays or co-insurance may. Such changes, though not expected to happen frequently, can raise or lower your estimated annual drug costs.

6. Types of Medicare Plans:

- "Stand alone" plans that offer only drug coverage. Such a plan may be for you if you wish to stay in the traditional Medicare fee-for-service program for other medical coverage.
- Medicare Advantage (MA) plans that cover both medical services and prescription drugs. Such a plan may be for you if you can accept restrictions on your choice of health care providers, and you prefer managed care.
- 7. Call and Confirm Plan Information: Prior to enrolling in any prescription drug plan, it's important to call the Prescription Drug Plan sponsor (or check their website) to confirm the information that you have gotten from the Plan Finder. The plans are able to make changes on a daily basis—and many have, as they try to be more competitive with the plans they only saw for the first time on November 15, 2005.
- 8. Changing Plans: Medicare beneficiaries who qualify for Extra Help have the option of changing their prescription drug plan at any time. If you do not qualify for Extra Help you can change your plan once between now and May 16, 2006. Thereafter, changes can be made during the annual enrollment period which is typically November 15 to December 31 of each year.

Other resource information

On the AARP Medicare website www.aarp.org/ medicarerx you will find a host of helpful information (in English and Spanish), including:

The New Medicare Prescription Drug Coverage: What You Need to Know

The New Medicare Prescription Drug Coverage: Extra Help for People with Limited Incomes

The New Medicare Prescription Drug Coverage: A Glossary of Terms

The New Medicare Prescription Drug Coverage: Drug Formularies, Exceptions, & Appeals

The New Medicare Prescription Drug Coverage: More Information for People with Medigap

The New Medicare Prescription Drug Coverage: More Information for People with *Retiree Coverage*

The New Medicare Prescription Drug Coverage: How to Ask the Right Questions & Get the Most from Your Medicines (includes My Medication Record)

The Medicare website www.medicare.gov, features more than the plan finder. On their home page, click on Other Resources for fact sheets, things to consider, etc.

Don't **Give Out**

Social Security Number

Bank Account Number

Credit Card Information

Final Note: Protect yourself against fraud

- **Applying is free**. The forms are free, and there's plenty of free help in your community to fill them out.
- Don't give out bank or credit card account numbers. You will have to give some personal information about your income and resources if you apply for Extra Help, but not your banking, checking or credit card information!
- Talk to the right person. If you have sent information to Social Security to apply for Extra Help, they may call you, but if the caller asks for your Social Security number, hang up! They probably aren't from the Social Security Administration!
- By law, Medicare plans cannot come to your door to sell their product, unless you have invited them. If you experience this kind of activity, contact your state attorney general. Look in the blue pages of your telephone directory for state government listings, then look for "Attorney General."
- **Don't be pressured**. Companies can call you to tell you about their drug plans, but they can't sign you up over the phone. In addition, organizations must:
 - Comply with the National-Do-Not-Call Registry
- Honor "do not call again" requests, and
- Abide by federal and state calling hours.
- Take your time. You have until May 15 of this year to enroll without incurring a late penalty.

List of Medications for Medicare Plan Finder Search	What I pay now monthly for the drugs I take	Plan 1	Plan 2	Plan 3
Monthly Premium	\$	\$	\$	\$
Deductible	\$	\$	\$	\$

#1			
#2			
#3			
#4			
#5			
#6			
#7			
#8			
#9			
#10			
#11			
#12			
Total Monthly Cost			
Does this plan have a gap in	coverage?		
If so, when does the ga	p begin?		
When does the gap end	!?		
Can I use the plan at the pha have to go to a different pha			
Does the plan give me discou drugs I currently take?			
If I currently buy a 90-day supp price difference? Do I have to			
If I live in another state for p can I get my drugs in that sta			

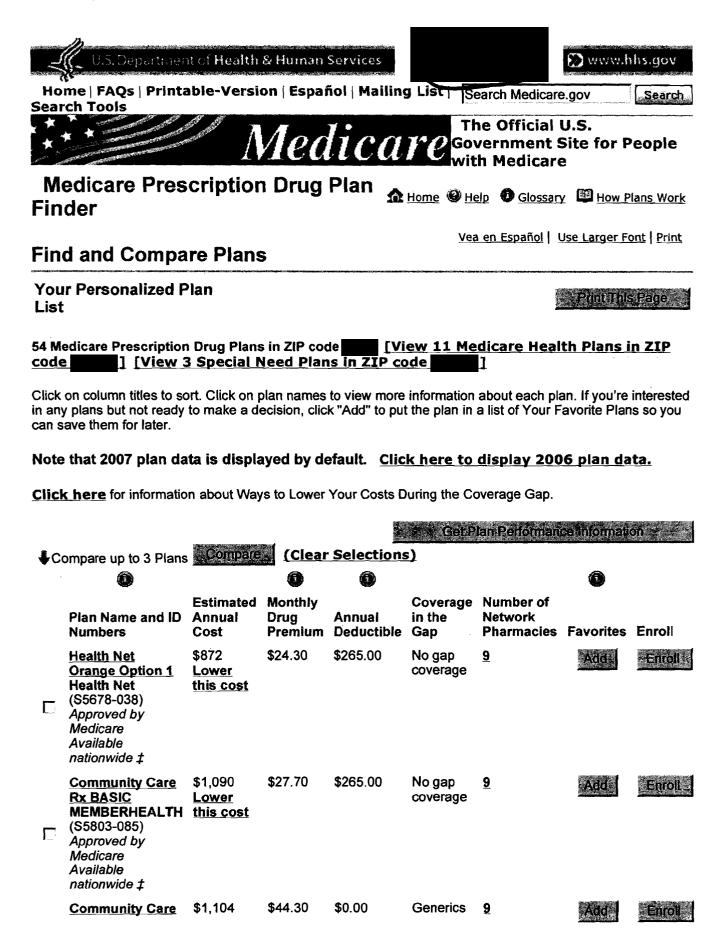
Name of the drug (including strength/numbers; e.g., Lipitor 20mg, 30 tab or Metformin HCL tab 500mg)

List of Medications for Medicare Plan Finder Search	What I pay now monthly for the drugs I take	Plan 1	Plan 2	Plan 3
Monthly Premium	\$	\$	\$	\$
Deductible	\$	\$	\$	\$

Name of the drug (including strength/numbers; e.g., Lipitor 20mg, 30 tab or Metformin HCL tab 500mg)

#1					
#2					
#3					
#4					
#5					
#6					
#7					
#8					
#9					
#10					
#11					
#12					
Total Monthly Cost					
Does this plan have a gap in	coverage?				
If so, when does the ga	p begin?				
When does the gap end	! ?				
	Can I use the plan at the pharmacy? Do I have to go to a different pharmacy?				
Does the plan give me discou drugs I currently take?					
If I currently buy a 90-day supp price difference? Do I have to u					
If I live in another state for p can I get my drugs in that sta					

AARP is a nonprofit, nonpartisan membership organization that helps people 50+ have independence, choice and control in ways that are beneficial and affordable to them and society as a whole. We produce AARP The Magazine, published bimonthly; AARP Bulletin, our monthly newspaper; AARP Segunda Juventud, our bimonthly magazine in Spanish and English; NRTA Live & Learn, our quarterly newsletter for 50+ educators; and our website, www.aarp.org. AARP Foundation is an affiliated charity that provides security, protection, and empowerment to older persons in need with support from thousands of volunteers, donors, and sponsors. We have staffed offices in all 50 states, the District of Columbia, Puerto Rico, and the U.S. Virgin Islands.



F	Rx GOLD MEMBERHEALTH (S5803-233) Approved by Medicare Available nationwide ‡	<u>Lower</u> this cost						
Γ	DeanCare Rx Enhanced Dean Health Insurance, Inc. (S5954-005) Approved by Medicare	\$1,193 <u>Lower</u> this cost	\$48.90	\$0.00	Generics and Preferred Brands	<u>9</u>	Add	Enroll
Г	Sterling Rx Plus Sterling Life Insurance Company (S4802-049) Approved by Medicare	\$1,215 <u>Lower</u> <u>this cost</u>	\$57.00	\$100.00	Generics	<u>9</u>	Add -	Enjol
	SilverScript Complete SilverScript Insurance Company (S5601-087) Approved by Medicare Available nationwide ‡	\$1,228 <u>Lower</u> this cost	\$35.80	\$0.00	Generics	<u>9</u>	Add	AFICUL
	First Health Select First Health Part D (S5768-063) Approved by Medicare	\$1,237 <u>Lower</u> <u>this cost</u>	\$35.30	\$0.00	No gap coverage	<u>9</u>	Add	Entol
—	CIGNATURE Rx Value Plan CIGNATURE Rx (S5617-078) Approved by Medicare Available nationwide ‡	\$1,238 <u>Lower</u> this cost	\$28.60	\$265.00	No gap coverage	<u>8</u>	Add	Enroll
Г	Community Care Rx CHOICE MEMBERHEALTH (S5803-153) Approved by Medicare Available nationwide ‡	\$1,248 <u>Lower</u> <u>this_cost</u>	\$35.70	\$0.00	No gap coverage	<u>9</u>	Add	Entoll
	<u>HealthSpring</u>	\$1,258	\$26.50	\$265.00	No gap	<u>9</u>	Add	Enroll

Ē	Prescription Drug Plan-Reg 16 HealthSpring Prescription Drug Plan (S5932-015) Approved by Medicare Available nationwide ‡	<u>Lower</u> this cost			coverage			
C	DeanCare Rx Classic Dean Health Insurance, Inc. (S5954-004) Approved by Medicare	\$1,268 <u>Lower</u> thi <u>s cost</u>	\$25.90	\$250.00	No gap coverage	<u>9</u>	Add	Enrol
F	UnitedHealth Rx Basic UnitedHealthcare (S5921-072) Approved by Medicare Available nationwide ‡	\$1,284 <u>Lower</u> t <u>his cost</u>	\$27.60	\$0.00	No gap coverage	<u>9</u>	Acc	Enrol
Γ	Aetna Medicare Rx Plus Aetna Medicare (S5810-152) Approved by Medicare Available nationwide ‡	\$1,292 <u>Lower</u> this cost	\$42.70	\$0.00	No gap coverage	<u>8</u>	Add	Enrol
r_	WPS MedicareRx Plan 2 WPS Health Insurance (S5753-007) Approved by Medicare	\$1,301 <u>Lower</u> this cost	\$42.50	\$0.00	Generics	<u>9</u>	Add	Enroll
Γ	SAMAscript SAMAscript (S7950-016) Approved by Medicare Available nationwide ‡	\$1,306 <u>Lower</u> this cost	\$44.60	\$265.00	No gap coverage	<u>8</u> .	Ada	<u>Contact</u> <u>Plan to</u> <u>Enroll</u>
С	Medco YOURx PLAN Medco YOURx PLAN (S5660-016) Approved by Medicare	\$1,328 <u>Lower</u> <u>this cost</u>	\$35.80	\$100.00	No gap coverage	<u>9</u>	Add	SEATON &
	Prescription	\$1,330	\$43.50	\$0.00	Generics	<u>9</u>	Add	Enroll

	Pathway Platinum Plan Reg 16 Pennsylvania Life Insurance Company (S5597-213) Approved by Medicare	<u>Lower</u> this cost						
	WPS MedicareRx Plan 1 WPS Health Insurance (S5753-006) Approved by Medicare	\$1,339 <u>Lower</u> this cost	\$37.30	\$0.00	No gap coverage	<u>9</u>	Add	Enrol
Γ	CIGNATURE Rx Plus Plan CIGNATURE Rx (S5617-080) Approved by Medicare Available nationwide ‡	\$1,339 <u>Lower</u> this cost	\$37.50	\$0.00	No gap coverage	<u>8</u>	Add	Enroll
	Health Net Orange Option 2 Health Net (S5678-037) Approved by Medicare Available nationwide ‡	\$1,349 <u>Lower</u> <u>this cost</u>	\$29.00	\$0.00	No gap coverage	<u>9</u>	Add	Enroll
.	AARP MedicareRx Plan - Saver UnitedHealthcare (S5921-071) Approved by Medicare Available nationwide ‡	\$1,357 <u>Lower</u> <u>this cost</u>	\$20.80	\$265.00	No gap coverage	<u>9</u>	Add	Enroll
٢	AdvantraRx Premier Coventry AdvantraRx (S5670-082) Approved by Medicare	\$1,359 <u>Lower</u> this cost	\$34.20	\$0.00	No gap coverage	<u>9</u>	Add	Enroll
F	<u>SilverScript Plus</u> SilverScript Insurance Company (S5601-033)	\$1,376 <u>Lower</u> <u>this cost</u>	\$31.70	\$0.00	No gap coverage	<u>9</u>	Acid	Enroll

Company (S5601-033) Approved by Medicare Available

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	nationwide ‡							
	CIGNATURE Rx Complete Plan CIGNATURE Rx (S5617-186) Approved by Medicare Available nationwide ‡	\$1,381 <u>Lower</u> <u>this cost</u>	\$48.90	\$0.00	Generics	<u>8</u>	XACC	<u>Contact</u> <u>Plan to</u> <u>Enroll</u>
[Aetna Medicare Rx Premier Aetna Medicare (S5810-186) Approved by Medicare Available nationwide ‡	\$1,386 <u>Lower</u> this cost	\$72.20	\$0.00	Generics	<u>8</u>	Add	Enroll
	Prescription Pathway Gold Plan Reg 16 Pennsylvania Life Insurance Company (S5597-048) Approved by Medicare	\$1,394 <u>Lower</u> this cost	\$23.00	\$0.00	No gap coverage	<u>9</u>	6116	Enrol
Г	SilverScript SilverScript Insurance Company (S5601-032) Approved by Medicare Available nationwide ‡	\$1,426 <u>Lower</u> <u>this cost</u>	\$23.40	\$265.00	No gap coverage	<u>9</u>	Add	Entol
	UnitedHealth Rx Extended UnitedHealthcare (S5820-119) Approved by Medicare	\$1,436 <u>Lower</u> <u>this cost</u>	\$40.60	\$0.00	No gap coverage	<u>9</u>	Add	Enroll
Γ	AdvantraRx Premier Plus Coventry AdvantraRx (S5670-084) Approved by Medicare	\$1,453 <u>Lower</u> <u>this cost</u>	\$48.30	\$0.00	Generics	<u>9</u>	Add	Enrol
Г	Health Net Orange Option 3 Health Net (S5678-088) Approved by Medicare Available	\$1,461 <u>Lower</u> <u>this cost</u>	\$44.10	\$0.00	Generics	<u>9</u>	Add	Enroll

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	nationwide ‡							
E	UA Medicare Part D Prescription Drug Cov United American Insurance Company (S5755-019) Approved by Medicare	\$1,464 <u>Lower</u> <u>this cost</u>	\$39.90	\$0.00	No gap coverage	<u>9</u>	Add	Enroll
	AARP MedicareRx Plan UnitedHealthcare (S5820-015) Approved by Medicare	\$1,484 <u>Lower</u> <u>this cost</u>	\$26.10	\$0.00	No gap coverage	<u>9</u>	Add	Enroll
Γ	AdvantraRx Value Coventry AdvantraRx (S5670-081) Approved by Medicare	\$1,502 <u>Lower</u> <u>this cost</u>	\$23.40	\$0.00	No gap coverage	<u>9</u>	Add	Enroll
	Sterling Rx Sterling Life Insurance Company (S4802-027) Approved by Medicare	\$1,503 <u>Lower</u> this cost	\$31.70	\$100.00	No gap coverage	<u>9</u>	Add	Enroll
Г	UA Medicare Part D Rx Covg - Silver Plan United American Insurance Company (S5755-054) Approved by Medicare	\$1,555 <u>Lower</u> <u>this cost</u>	\$30.60	\$265.00	No gap coverage	<u>9</u>	AGC	Enroll
—	EnvisionRxPlus Gold EnvisionRx Plus (S7694-050) Approved by Medicare Available nationwide ‡	\$1,570 <u>Lower</u> <u>this cost</u>	\$69.00	\$0.00	Generics	<u>8</u>	Add	Enroll
Γ	AARP MedicareRx Plan - Enhanced UnitedHealthcare (S5921-073) Approved by Medicare Available	\$1,597 <u>Lower</u> <u>this cost</u>	\$44.00	\$0.00	Generics	<u>9</u>	Acc	Entol

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	nationwide ‡							
Γ	Blue MedicareRx Plus Blue Cross Blue Shield of Wisconsin (S5596-022) Approved by Medicare	\$1,600 <u>Lower</u> this cost	\$34.60	\$0.00	No gap coverage	<u>9</u>	Add	Enroll
.	Blue MedicareRx Value Blue Cross Blue Shield of Wisconsin (S5596-021) Approved by Medicare	\$1,607 <u>Lower</u> <u>this cost</u>	\$27.50	\$250.00	No gap coverage	<u>9</u>	Add	Enroll
L	Prescription Pathway Bronze Plan Reg 16 Pennsylvania Life Insurance Company (S5597-081) Approved by Medicare	\$1,609 <u>Lower</u> <u>this_cost</u>	\$25.00	\$265.00	No gap coverage	<u>9</u>	Add	Enrol
	Blue MedicareRx Premier Blue Cross Blue Shield of Wisconsin (S5596-023) Approved by Medicare	\$1,626 <u>Lower</u> <u>this cost</u>	\$45.40	\$0.00	Generics	<u>9</u>	Add	Enroll
Γ	MedicareRx Rewards Premier Unicare (S5960-086) Approved by Medicare Available nationwide ‡	\$1,650 <u>Lower</u> <u>this cost</u>	\$47.40	\$0.00	Generics	<u>9</u>	Add	Enroll
٢	Humana PDP Complete S5884- 044 Humana Insurance Company (S5884-044) Approved by Medicare	\$1,801 <u>Lower</u> this cost	\$80.30	\$0.00	Generics	<u>9</u>	Add	Enrol
	<u>NMHC Medicare</u> <u>PDP Gold</u> NMHC Group Solutions	\$1,824 <u>Lower</u> this cost	\$29.30	\$0.00	No gap coverage	7	Add	Enroll

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Г	(S8841-016) Approved by Medicare Available nationwide ‡							
Г .,	WellCare Signature WellCare (S5967-050) Approved by Medicare Available nationwide ‡	\$1,833 <u>Lower</u> this cost	\$24.50	\$0.00	No gap coverage	<u>9</u>	Add	Enroll
Γ	Humana PDP Enhanced S5884- 014 Humana Insurance Company (S5884-014) Approved by Medicare	\$1,848 <u>Lower</u> this cost	\$23.50	\$0.00	No gap coverage	<u>9</u>	Add	Enroll
Γ	Aetna Medicare Rx Essentials Aetna Medicare (S5810-050) Approved by Medicare Available nationwide ‡	\$1,900 <u>Lower</u> <u>this_cost</u>	\$29.20	\$200.00	No gap coverage	<u>8</u>	Add	Enrol
F actor	MedicareRx Rewards Value Unicare (S5960-016) Approved by Medicare Available nationwide ‡	\$1,920 <u>Lower</u> this cost	\$26.20	\$265.00	No gap coverage	<u>9</u>	Add	Enroll
Γ	WellCare Classic WellCare (S5967-153) Approved by Medicare Available nationwide ‡	\$2,051 <u>Lower</u> <u>this cost</u>	\$15.90	\$265.00	No gap coverage	<u>9</u>	Add	Enroll
	WellCare Complete WellCare (S5967-084) Approved by Medicare Available nationwide ‡	\$2,085 <u>Lower</u> <u>this cost</u>	\$45.50	\$0.00	Generics	<u>9</u>	Ado	Enroli
	<u>Humana PDP</u> Standard S5884-	\$2,134 <u>Lower</u>	\$14.80	\$265.00	No gap coverage	<u>9</u>	Add	Enfol

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L	074 Humana Insurance Company (S5884-074) Approved by Medicare	<u>this cost</u>					•	
	Advantage Freedom Plan by RxAmerica RxAmerica (S5644-177) Approved by Medicare Available nationwide ‡	\$2,415 <u>Lower</u> this_cost	\$34.00	\$265.00	No gap coverage	<u>6</u>	Add	Enroll
	Advantage Star Plan by RxAmerica RxAmerica (S5644-191) Approved by Medicare Available nationwide ‡	\$2,461 <u>Lower</u> this cost	\$29.30	\$265.00	No gap coverage	<u>6</u>	Add .	Enroll
٢	EnvisionRxPlus Standard EnvisionRx Plus (S7694-016) Approved by Medicare Available nationwide ‡	\$3,136 <u>Lower</u> this cost	\$45.00	\$265.00	No gap coverage	<u>8</u>	And	Enroll
	Plans per page: All						ge: All 🔀	

Compare up to 3 Plans Compare (Clear Selections)

‡The organization that offers this plan offers at least one plan in all fifty states and the District of Columbia. Note: The specific plan(s) offered by this organization may differ from state to state.

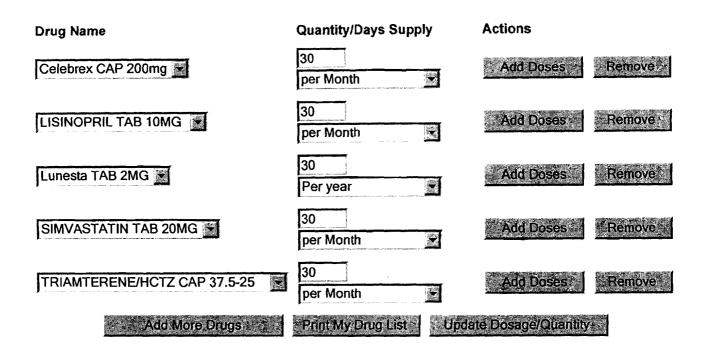
The drug costs displayed above are estimates based on your current drug use and preferred pharmacies. These estimates may vary based on the specific quantity, strength, and/or dosage of each medication, the order in which you purchase your prescriptions, and the pharmacy you use. You may wish to revise your Drug & Pharmacy List below in order to get the most accurate cost estimates.

My Drug & Pharmacy List

Review the dosage and quantity information displayed below for each of your drugs and update if necessary. If you take more than one dose of the same drug, click "Add Doses."

Note that if you change the strength of a drug using a dropdown menu in the "Drug Name" column, you then need to make sure the information in the "Quantity & Days Supply" column is still correct.

My Drugs



My Pharmacies

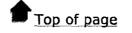
No pharmacies selected. Click the Change Pharmacy Selection button to add pharmacies.

Selected Pharmacies

Change Pharmacy Selection

Remove

Page Last Updated: October 31, 2006





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Centers for Medicare & Medicaid Services | U.S. Department of Health and Human Services

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