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THE MEDICARE DRUG BENEFIT:

ARE PRIVATE INSURERS GETTING

GOOD DISCOUNTS FOR THE TAXPAYER?

Thursday, July 24, 2008

House of Representatives,

Committee on Oversight and

Government Reform,

Washington, D.C.

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## **Committee Hearings**

of the

## U.S. HOUSE OF REPRESENTATIVES



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- 1 RPTS KESTERSON
- 2 DCMN MAYER
- 3 THE MEDICARE DRUG BENEFIT:
- 4 ARE PRIVATE INSURERS GETTING
- 5 GOOD DISCOUNTS FOR THE TAXPAYER?
- 6 Thursday, July 24, 2008
- 7 House of Representatives,
- 8 Committee on Oversight and
- 9 Government Reform,
- 10 Washington, D.C.

- The committee met, pursuant to call, at 10:10 a.m., in
  Room 2154, Rayburn House Office Building, Hon. Henry A.
- Waxman [chairman of the committee] presiding.
- 14 Present: Representatives Waxman, Cummings, Kucinich,
- 15 Tierney, Watson, Higgins, Yarmuth, Braley, Van Hollen, Murphy
- of Connecticut, Sarbanes, Speier, Davis, Burton, Shays,
- 17 Platts, Issa, Marchant, McHenry, Foxx, Bilbray, and Jordan.

Staff Present: Kristin Amerling, General Counsel; Caren 18 Auchman, Press Assistant; Phil Barnett, Staff Director and 19 20 Chief Counsel; Jen Berenholz, Deputy Clerk; Brian Cohen, Senior Investigator & Policy Advisor; Miriam Edelman, Special 21 Assistant; Earley Green, Chief Clerk; Ella Hoffman, Press 22 23 Assistant, Karen Lightfoot, Communications Director and Senior Policy Advisor; Karen Nelson, Health Policy Director; 24 25 Jennifer Owens, Special Assistant; Andy Schneider, Chief Health Counsel; Leneal Scott, Information Systems Manager; 26 27 Mitch Smiley, Special Assistant; John Williams, Deputy Chief 28 Investigative Counsel; Lawrence Halloran, Minority Staff Director; Jennifer Safavian, Minority Chief Counsel for 29 Oversight and Investigations; Ali Ahmad, Minority Deputy 30 Press Secretary; Larry Brady, Minority Senior Investigator & 32 Policy Advisor; Patrick Lyden, Minority Parliamentarian & 33 Member Services Coordinator; Brian McNicoll, Minority Communications Director; John Ohly, Minority Professional Staff Member; Jill Schmaltz, Minority Senior Professional Staff Member; and Molly Boyl, Minority Professional Staff 36 Member.

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Chairman WAXMAN. Good morning. The committee will please come to order.

Today, the committee is holding another hearing in our series of how to make government work better. Our subject is the Medicare Part D program that provides a prescription drug benefit to seniors and individuals with disabilities.

Providing drug coverage to seniors and disabled is essential, but it is also expensive. Over the next decade, the benefit will cost taxpayers hundreds of billions of dollars. We need to make sure this money is spent responsibly and with good value for the taxpayers.

This committee has been investigating Medicare Part D for 18 months. During our investigation, we have conducted the only in-depth oversight of the Part D program. GAO and the Congressional Budget Office have been unable to review how well the program is working because the Centers for Medicare and Medicaid Services won't give them the data; and CMS, which does have access to data, refuses to acknowledge fundamental flaws in the program.

Last October, I and other members of the committee released the staff report that examined the administrative costs of Medicare Part D. We found that the private insurers that delivered the Medicare benefit are charging taxpayers and beneficiaries \$4.6 billion in administrative costs annually. In percentage terms, that is over six times more

than it costs to run traditional Medicare. And we found that the Part D program is exceptionally lucrative for private health insurers. They made a billion dollars in profit last year alone.

Today, I am joining with 10 members of the committee to release a new staff report, which I ask to be made part of today's hearing record. Without objection.

[The information follows:]

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Chairman WAXMAN. Last year's report looked at the profits of the private insurers. Today's report examines the windfall revenues of the drug manufacturers. In this report, we compare the prices that the drug companies charge the new Medicare Part D program with the prices that the companies charged the Medicaid program.

What we discovered is that the taxpayers are paying far more for drugs under Medicare Part D than they do under Medicaid. In effect, Medicare Part D has given the major drug companies a taxpayer-funded windfall worth billions of dollars.

Our report focuses on the cost to the taxpayer of providing drugs to the 6 million beneficiaries who are enrolled in both Medicare and Medicaid. These are Americans who are old or disabled enough to qualify to be on Medicare, and they are poor enough also to qualify for Medicaid. They are often the oldest and sickest Medicare beneficiaries and their drug coverage is almost fully subsidized by Federal taxpayers. These---"dual eligibles" is what they are called, these dual-eligible beneficiaries account for about half of all drug spending in Medicare Part D.

The multibillion-dollar windfall is a result of a provision in the Medicare Part D law that switched drug coverage for the dual eligibles from Medicaid to Medicare Part D. The transfer took effect 2 years ago. Since then,

the drug manufacturers have been paid billions more for the drugs used by the dual-eligible beneficiaries than they would have been paid if the dual eligibles had continued to receive their drug coverage through Medicaid.

Under Medicare Part D, the 6 million dual-eligible beneficiaries can take the same drugs they got under Medicaid; the only difference is that the Federal taxpayer is now paying 30 percent more. Add it up and it amounts to a drug manufacturer windfall worth at least \$3.7 billion in just the first 2 years of the Medicare Part D program. In fact, the actual windfall could be worth billions more if all drugs used by dual-eligible beneficiaries were taken into account.

Let me describe some examples. Johnson & Johnson earned over \$500 million in additional profits, much of it from just one drug, the antipsychotic medication Risperdal. Bristol Myers earned a windfall of almost \$400 million thanks to the higher prices for the stroke medication Plavix. This is an enormous giveaway, and it—it has absolutely no justification. The drug companies are making the same drugs, they are being used by the same beneficiaries, yet because the drugs are being bought through Medicare Part D instead of Medicaid, the prices paid by the taxpayers have ballooned by billions of dollars.

The privatization of Medicare Part D is a great deal for

the drug companies, And it is a great deal for the private insurers. It is the taxpayers who are taking it on the chin.

The circumstances that led to passage of the Medicare Part D were controversial. The chairman of the House committee that wrote the Part D law now runs PhRMA, the drug manufacturers trade association. The administration's top negotiator left the government to lobby for health insurers and drug companies.

There are allegations of threats and arm-twisting on the House floor, but that is not the focus of today's hearing.

The Medicare drug benefit is providing real help to seniors and the disabled, and it is going to be part of our health care landscape for years to come.

The key question for us is, how we can fix the program so that more of the benefit goes to seniors and the disabled and less winds up in the pockets of the drug companies and insurers.

Medicaid is one proven model for how the government can use its purchasing power to ensure that it gets low prices.

Medicaid is a voluntary program. No drug manufacturers are required to participate. Medicaid gets its low prices by making discounts a condition of manufacturers participating.

The program says that if a manufacturer wants to sell their drugs to Medicaid beneficiaries, they have to offer Medicaid their lowest prices. The manufacturers also have to agree to

protect the taxpayers from price increases that exceed the rate of inflation.

We have well over a decade of operational experience with the Medicaid rebate. It works. It delivers \$10 billion annually in savings to the Federal and State governments. In many ways, this is the exact opposite of what is going on under Medicare Part D. Under Part D, the drug manufacturers can charge essentially what they want. Despite their high administrative costs and billion dollar profits, the private insurers have been unable to stand up for the interest of the taxpayers.

Now, many of our hearings on waste, fraud and abuse identify problems that the executive branch can fix administratively; that is not the case with Medicare Part D. The waste in this program is the direct result of the statutory design of the law. Congress wrote this law and must lead the way to a solution. To start this process, I will soon be introducing legislation that will protect the taxpayer by bringing down the high price--high drug prices in Medicare Part D. This bill will guarantee that Federal taxpayers cannot be charged higher prices for the dual-eligible beneficiaries under Medicare Part D than under Medicaid.

The potential savings to Medicare and the Federal taxpayers are enormous. Passage of reform legislation could

save the taxpayer almost \$90 billion over the next 10 years; even more could be saved if the Federal Government were to authorize to negotiate prices on behalf of all Medicare beneficiaries.

I am looking forward to hearing more about this issue today and working together with the members of this committee to improve the Part D program. I will be introducing our witnesses, who I'm grateful are here today. All of them are here voluntarily.

But before we do that, I want to recognize Mr. Davis for an opening statement.

[Prepared statement of Chairman Waxman follows:]

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Mr. DAVIS OF VIRGINIA. Well, thank you, Mr. Chairman.

The Medicare prescription drug program, known as Part D, has successfully provided needed medicines to millions of American seniors. The proof is in the pudding: Overwhelming number of seniors have opted into this program. It is an optional program that speaks for its success. While only in its third year of operation, Part D continues to come in below initial budget projections.

Nevertheless, even with all of its successes, Medicare
Part D, like any Federal program, could benefit from
thoughtful, evenhanded oversight; and I hope that is our goal
here today. But I'm not convinced there is much constructive
to be learned simply by comparing controlled prices under
Medicaid and market prices under Part D and labeling the
entire difference a windfall.

The majority staff analysis released this morning focuses on dual eligibles, seniors eligible for both Medicare and Medicaid. Before 2006, they received prescription drug insurance through Medicaid which uses statutory price controls. At the request of States and many senior citizen advocates, dual eligibles were included under Part D. Not surprisingly, market-negotiated drug prices for this special population were found to be higher than the legally mandated, below-market Medicaid rates.

But any alleged windfall, however large, tells really

less than half the story. That difference buys dual-eligible seniors access to drugs not available under Medicaid's more restrictive pharmacy rules, and capturing the alleged savings would be short lived and painful. It would come at a very, very high cost as other segments of the health care delivery system, nongovernment segments—we are talking about employer plans, union plans—payments for the uninsured would then absorb the cost shifts that are inevitably generated by price controls.

This is not just a theoretical argument about how free markets work. The Federal Government does have almost 20 years of experience with the implications of prescription drug price controls. The Congressional Budget Office and the Government Accountability Office both have repeatedly found that Medicaid price controls increase prescription drug prices to every other purchaser.

Transplanting Medicaid price controls onto Part D could have other unwanted implications. We should be very concerned about a Federal Government process to set Part D prices that would turn into a political exercise. There would be enormous political pressure to pick winners and losers.

Elsewhere in Medicare, relentless lobbying shifts and shapes reimbursement policies for some services or specialties over others; and it is not a very pretty process.

Just a couple of weeks ago, Medicare physicians almost took a 10 percent reimbursement cut at the hands of a government-run pricing system.

Given the critical role of Medicare in caring for seniors as they age, we should conduct oversight of the program, but it strikes me that this committee's discussion of Part D is stuck in a rut. With every new report and each successive hearing, I understand Yogi Berra's concept of "deja vu all over again." Repeatedly making economically and plausible arguments about the efficiency of government-run drug pricing or plucking artificial windfalls from thin air won't make Part D, a good program, work any better.

It is running well under the original 2003 budget projections, due largely to lower-than-anticipated bids from prescription drug plans. That is what happens in the free, competitive market. And most importantly, opinion surveys report that 85 percent of Part D beneficiaries are happy with the program, the 15 percent obviously on the other side of the aisle here, with the satisfaction rate even higher among the dual eligibles.

Meanwhile, other aspects of the program urgently need scrutiny. We could be talking about Medicare payments for durable medical equipment prescribed by physicians or the serious financial trouble facing Part A, Medicare hospital insurance, which is due to go bankrupt in 11 years.

The bedrock of the program, Part A, is in dismal shape. The Medicare trustees reported this year the Hospital Insurance Trust Fund will be insolvent in 2019. When that happens, payments can no longer be made to cover seniors' hospital care. There is no authority in current law to allow general revenue funding of that shortfall. We obviously--we fund Part B.

I look forward to our oversight hearings on these pressing issues today. Mr. Chairman, I would also ask unanimous consent that the minority staff analysis be submitted for the record.

Chairman WAXMAN. Without objection, that will be the order.

[The information follows:]

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275	Chairman WAXMAN. We are pleased to welcome for our
276	first panel, Dr. Stephen Schondelmeyer, who is a Ph.D. And
277	Professor and Head of the Department of Pharmaceutical Care
278	and Health Systems at the University of Minnesota; Dr. Gerard
279	Anderson, Ph.D., Professor and Director for the Center for
280	Hospital Finance and Management, Bloomberg School of Public
281	Health at Johns Hopkins University; Fiona M. Scott Morton,
282	Ph.D., Professor of Economics, Yale School of Management,
283	Yale University.
284	We are pleased to have the three of you here today. It
285	is the practice of this committee that all witnesses testify
286	under oath. So if you would please stand.
287	Mr. DAVIS OF VIRGINIA. Mr. Chairman, could I just note
288	for the record, Dr. Schondelmeyer is the majority's witness
289	who, 2 weeks ago, was given notice of this; and we have not
290	yet received written testimony from him.
291	Our minority witness has submitted his for the record
292	ahead of time for scrutiny. Thank you.
293	Chairman WAXMAN. Thank you, Mr. Davis. If the three of
294	you would please stand and raise your right hand.
295	[Witnesses sworn.]
296	Chairman WAYMAN The record will indicate that each of

the witnesses answered in the affirmative.

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298 l STATEMENTS OF DR. STEPHEN SCHONDELMEYER, PHARM.D., Ph.D., PROFESSOR AND HEAD, DEPARTMENT OF PHARMACEUTICAL CARE AND 299 HEALTH SYSTEMS, UNIVERSITY OF MINNESOTA; DR. GERARD ANDERSON, 300 301 Ph.D., PROFESSOR AND DIRECTOR, CENTER FOR HOSPITAL FINANCE 302 AND MANAGEMENT, BLOOMBERG SCHOOL OF PUBLIC HEALTH, JOHNS 303 HOPKINS UNIVERSITY; AND FIONA M. SCOTT MORTON, Ph.D., PROFESSOR OF ECONOMICS, YALE SCHOOL OF MANAGEMENT, YALE 304 305 UNIVERSITY 306 Chairman WAXMAN. Dr. Schondelmeyer, we are going to

Chairman WAXMAN. Dr. Schondelmeyer, we are going to start with you, but Mr. Davis made a very good point that we expect witnesses to submit their statements in advance under the rules. Please go ahead.

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Did you submit a statement to us, a written statement?

Mr. SCHONDELMEYER. I have not yet. I can after this
meeting. I do apologize.

Chairman WAXMAN. Turn on the mic. Yes, there is a button on the mic.

Mr. SCHONDELMEYER. I do apologize. I accepted this assignment with many other commitments, and this was a very tight schedule for me, given other commitments. But I was pleased to do so and--

Chairman WAXMAN. We're happy to have you here anyway.

Thanks.

We are going to ask each of you, as we will all of our witnesses, to try to keep within 5 minutes. I think you all have been informed of that in advance. And if you have submitted written statements, they will be part of the record in full. We're going to have a clock that will be green for 4 minutes, yellow for 1 minute and then when the 5 minutes is up, it will turn red. We're not going to be abrupt in stopping you, but I hope that red will be an indication that it is time to get ready-get ready and to conclude.

Thank you. Please go ahead.

331 STATEMENT OF DR. STEPHEN SCHONDELMEYER, PHARM.D., Ph.D.

Mr. SCHONDELMEYER. Thank you, Mr. Chairman, for inviting--

Chairman WAXMAN. Pull your mic a little closer.

Mr. SCHONDELMEYER. Thank you for inviting me and thank you to the rest of the committee. I will skip the normal formalities and broad background descriptions, because you've done that well in your introduction.

The dual eligibles, as was noted, however, represent a large share of the expenditures both under the previous Medicaid program and under the current Medicare Part D program. Just to put that in perspective, in the year 2005, total Medicaid drug expenditures were about \$43 billion a year. In 2006, after those dual eligibles moved from Medicaid over to Medicare, the Medicaid drug expenditures dropped to less than half of that 43 billion, somewhere around \$21 billion. So it is very real that this shift did move dollars from the State-run Medicaid programs to the private, market-run Part D Medicare programs.

At the same time that that shift occurred, also the access to the rebates under the State-run Medicaid programs disappeared.

Let me put in perspective rebates, briefly, under

Medicaid. The Medicaid drug rebate program began back in 1991 and continues to this day. There is a Federal component to the Medicaid drug rebate program which mandates 15.1 percent rebate for all brand-name drugs, and in addition for brand-name drugs, they are subject to a best-price additional rebate and an inflation adjustment rebate that often adds substantially beyond that 15.1 percent for all brand-name drugs. For generic drugs, all generic drugs must provide an 11 percent rebate.

Now, notice in both brand-name and generic drugs, all prescription drugs are subject to rebates. That is not necessarily the case today. Under the Medicare Part D program, not all drugs are subject to rebate; and particularly those drugs that are covered under the must-cover categories, the categories where the Part D plans can't negotiate or opt to cross different drug categories, those don't appear to receive as much rebate, although under the Medicaid program they did receive the same amount of rebate--at a minimum at least--as the other brand-name drugs.

Second, the amount of rebates from 1991--it took a year or two to get the program stabilized. From 1993 to 2000, about 18 to 19-1/2 percent of total drug spending came back to Medicaid programs as rebates. So about 18 to 19-1/2 percent came back.

Beginning in 2000-2001, though, the States woke up and

realized that the Medicaid legislation also authorized States' supplemental rebate programs. In those State supplemental rebate programs, it said States could negotiate on their own rebates above and beyond the Federal rebate, and that has started to grow.

In the early--2000 through 2003, we saw rebates grow to 20-21 percent. And then we saw a dramatic growth; in 2004 rebates grew to 24 percent of the total drug spend, 2005 rebates under Medicaid grew to 28.8 percent of the drug spend.

Unfortunately, to the best of my knowledge, CMS has not released the rebate data for the years 2006 and 2007 under Medicaid, so we can't look to see what the total amount is. As best I can tell from talking with various States out there, however, the number is probably somewhere above 30 to 31 percent total drug spend returned in rebates.

Now, that compares with--this committee did a report a year ago that suggested only about 8 percent of the drug spend under Medicare Part D was coming back as rebates, and that wasn't for all drugs and all classes. So if you compare 28.8 or 30 percent rebates on Medicaid to 8 percent on Medicare--and I understand your new report shows that the number has gone up under Medicare Part D, but it is still less than half of what the rebate amount was under the Medicaid program--it is obvious that if these same dual

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eligibles remained in the Medicaid program, the taxpayers and 404 the beneficiaries themselves would benefit from lower drug spend, as you pointed out, Mr. Chairman, on the same drug, the same people. It--just at a lower price in the marketplace. And those are based on State-negotiated supplemental rebates, not mandated rebates. negotiated with the States above and beyond the Federal rebate.

So it is also important to realize, under the Medicare Part D program, that the dual eligibles and the people on the private side do not receive the benefit of these rebates in lower drug price for most cases. You can find the odd drug, there may be a handful of 10 or 15 drugs where a lower price is actually passed on to the recipients, but for the most part, lower prices are not passed onto the recipient. the coverage gap, the person pays the entire cost of the drug without the benefit of any of the rebate. And for specialty drugs, where they may be paying 50 to 75 percent coinsurance, they're paying the entire cost of the drug without the benefit of the rebates.

In conclusion, it is not just observations of State accountants and academics like myself that say this was a shift in resources. Also Wall Street and corporate annual reports in both 2006 and 2007 noted that drug companies had substantially increased revenues that heretofore had been

429	unexpected due largely, in part, to volume increases under
430	Medicare Part D and the decreased payment for rebates under
431	Part D versus under Medicaid.
432	Thank you very much, Mr. Chairman.
433	Chairman WAXMAN. Thank you, Mr. Schondelmeyer.
434	[Prepared statement of Mr. Schondelmeyer follows:]
435	****** COMMITTEE INSERT ******

436 Chairman WAXMAN. Dr. Anderson.

437 STATEMENT OF DR. GERARD ANDERSON, Ph.D.

Mr. ANDERSON. Thank you, Chairman Waxman. It is a pleasure to return to this committee to talk about the issue of drug pricing.

My testimony can be summarized in two observations and three recommendations.

My first observation is that Part D plans paid even higher prices for drugs than Medicaid programs were paying. My second observation is the United States pays significantly higher prices for prescription drugs than other countries and that, in the United States, the private sector pays generally 20 percent higher prices than the public sector pays for drugs.

These two observations lead me to three recommendations. First, there should be greater price transparency in the pharmaceutical market. Second, drug pricing data should be readily accessible to congressional agencies and academic researchers so they can easily know if Part D plans are paying higher prices than Medicaid. And third of all, all government agencies should be paying the same prices for drugs.

The remainder of my testimony will explain in greater detail the rationale behind these observations and recommendations.

When the responsibility for providing drug coverage for the dual eligibles was transferred in 2005, the expectation, or even the hope, was that Part D plans would be able to obtain lower prices than the Medicaid programs.

Unfortunately, a growing body of data, including the report today, suggest that Part D plans are paying even higher prices than Medicaid programs. Amazingly, all the data seems to confirm that the windfall to the drug companies is about \$2 billion a year.

The first indication of higher prices came from the disclosures by the pharmaceutical companies themselves in their 10-Ks and 10-Qs filed with the Security and Exchange Commission. My written testimony cites specific documents, showing that the pharmaceutical companies were getting higher prices than Part D. Pfizer alone, for example, estimated in its 10-Q an additional \$300 million in profits.

Secondly, in my report, I show how CBO-CMS actuary data estimate using that data that Part D plans were paying 22 percentage points more than Medicaid was paying for the same drug. This committee says 30 percent; the CMS testimony today says 20 percent. So they are all in pretty much the same range.

The third indication was the report by this committee last year. So basically all the different sources--and as a researcher you want to have multiple sources--then, the transfer from the dual eligibles will result in about a \$2 billion annual windfall to the drug companies; and it is currently in line with the report of this committee.

Surprisingly, the Medicare program is not the insurer paying the highest prices for drugs in the United States.

Typically, the private sector pays 20 percent more for drugs . than the Medicare and Medicaid programs.

The fact that Part D plans were unable to obtain substantial discounts from the pharmaceutical companies is surprising to me, given the difficulties that the Medicaid agencies have obtaining actual transaction prices to set their own rates. In a series of recent court decisions, judges and juries have found that this lack of price transparency has made it difficult for the Medicaid agencies to actually set prices.

President Bush has argued that there should be greater price transparency in the health care sector. When the Bush--while the Bush administration has promoted major efforts to increase the level of price transparency in the hospital and physician sectors, surprisingly there has been very little emphasis on price transparency in the pharmaceutical sector.

In order to make greater price transparency, I believe the Secretary of Health and Human Services should determine in the markets are actually working for pharmaceuticals. One way to determine this is to compare the lowest prices that any of the Part D plans are obtaining and compare to the prices that the Medicaid programs, the VA or even Canada are obtaining.

Unfortunately, provisions in the MMA limit disclosure of information on drug prices and drug utilization. This data should be given to CBO, CRS, MedPac and other government agencies to analyze the effectiveness of the Part D program. It should also be given to academic researchers.

My third and final recommendation is that all government programs should pay the same rate for each drug. I cannot think of a compelling reason, either economically or ethically, why one government program, save the VA, should pay a higher price or a lower price through the Medicare program; all the money comes from the taxpayers. Governments in other countries manage to pay one price for drugs. Why not the United States?

Thank you for the opportunity to testify this morning.

Chairman WAXMAN. Thank you very much, Dr. Anderson.

[Prepared statement of Dr. Anderson follows:]

\*\*\*\*\*\* INSERT 1-2 \*\*\*\*\*\*

532 Chairman WAXMAN. Dr. Morton.

## STATEMENT OF FIONA M. SCOTT MORTON

Ms. MORTON. Good morning to the chairman and members of the committee. Thank you very much for inviting me to testify. I just have some short remarks.

The report that was released this morning repeatedly says that manufacturers charge more to Part D than they charge to Medicaid. I just would like everyone to keep in mind that the manufacturers--under Medicaid, they sell to drugstores in the normal way, and then they are required to give a rebate back to the government. And that is how we get a net price; it is not a charged price.

And the size of that rebate is set in law; and the important thing, I think, that we see today, that we didn't see in the early 1990s, was the size of the inflation component of that. And that is not something that Part D can negotiate for. That inflation component is big, and it is mandated under Medicaid.

So I would say that the findings of the report are completely predictable in the sense that we knew that Medicaid was required to get the lowest price, and we knew it had these big rebates. And so, of course, that is going to

be, as Mr. Davis said, the place where you've got the lowest prices, and we wouldn't expect Part D to be able to do as well as that.

So I think if Congress is concerned about just the cost of covering duals, then you should move them back into Medicaid. I mean, that is where you're going to get the lowest prices for these people. It would also reduce confusion for them and plan shifting as the plan they are in becomes too high cost and they're moved to another plan that—I believe that kind of transition is difficult.

Secondly, the report finds that the protected classes in Part D get small discounts. Again, I'm going to take this opportunity to say that when I testified for the Senate in January 2007, I predicted this, because you can't move market share in these groups. The formularies are restricted and the Part D plans have to cover all drugs, essentially; and if you can't bargain with the manufacturer, saying, I'm going to move market share to Drug A from Drug B, you can't get a discount. And I think it is very reasonable then to see that you're not getting discounts in these protected classes.

Again, this is something you could change with respect to the regulation. You could have fewer protected classes, you could loosen the formulary restrictions so that plans can do a bit more shifting of market share from one drug to another; and then you'd expect to see bigger discounts.

Thirdly, we have talked a lot about the windfall that has arisen from moving guys from Medicaid into Medicare. I have some research looking at the opposite effect, which is the movement of the uninsured from paying cash to having coverage under Medicare, and there the windfall appears to have gone in the opposite direction. So the prices that an uninsured, cash-paying person pays are a lot higher than--now, I don't have the same access to information as you do, Mr. Waxman, so I'm inferring it from some less-good data, but it looks like the prices are going down quite drastically.

So we do have success of the program in helping the uninsured get access to drugs at lower prices. But--so I just would like to point that out, since we have the windfall going the other way as well.

Then two--just points that are longer run. First of all, I think this committee might want to return to this question next year because the way the negotiations work is, they happen in February for prices to set in November for the next year. So when you think about the experience with the program, it wasn't until February of 2007 that plans and everybody could watch a whole year of operation of this program. And so it wasn't, therefore, until prices were set for 2008 that you see kind of informed outcomes, as opposed to just guessing what are people going to do and where are

they going to enroll. So I think we can learn more going forward.

And then, lastly, it seems messy and costly to me to try to have a Medicaid rebate applied to some purchases inside Medicare. It seems just--because you get those rebates. The supplemental rebates come from shifting, having a preferred drug; the Medicare Part D rebates come from having a preferred drug. So trying to get a plan to have a Medicaid rebate for a guy who is in their Medicare plan that they are trying to negotiate over with the manufacturers, that seems very complex to me. I think it would be just easier to move them, for the plan.

And I think that—oh, the last thing about the Medicaid rebates is, they are large and they really reduce the profitability, of course, of selling to the Medicaid program. I think that works partly because the Medicaid program is small, so if it is 12, 15, 18 percent of the Nation's drug spending, the manufacturers can afford and should be interested in providing medications at low cost to those poor people who are also sick.

But when you think about Medicare, 40 percent of all prescriptions are doled out to people who are eligible for Medicare. I mean, by the time you add on Medicaid--that's half the market--you're then talking about a very serious change in the market structure of the pharmaceutical

529	industry.
630	Thank you. That's all.
531	Chairman WAXMAN. Thank you very much.
532	[Prepared statement of Ms. Morton follows:
533	****** INSERT 1-3 *****

634 Chairman WAXMAN. I'll start off the question -- 5 minutes 635 of questions. 636 That was an interesting point you just raised about the Medicaid population being so much smaller than the Medicare 637 638 population, but when we talk about dual eligibles, we are 639 talking about half the budget for pharmaceuticals under Part 640 D. 641 You're shaking your head. You acknowledge that fact? 642 Ms. MORTON. Yes, I think not everybody who is Medicare 643 eligible is enrolled in Part D. So the current proportion of 644 duals is quite high relative to all the people who could be 645 signing up for Part D going forward. 646 Chairman WAXMAN. If we paid the Medicaid price for 647 those dual eligibles, there would be a tremendous savings. 648 Do you agree? 649 Oh, there would. Ms. MORTON. Because we used to have them in Medicaid where these regulated prices were below 650 651 market level. Absolutely. 652 Chairman WAXMAN. Do you think that did any harm to the 653 ability of the prescription drug industry to do their 654 research, market their products? 655 Or your statements seem to be, it is a small amount so 656 that the controls on those prices, requirements of discounts -- it did not have an adverse effect? 657 658 Ms. MORTON. It is hard to know what the ideal amount of

research and development is, so I won't tread in that area. But in terms of where we were before with kind of 18 percent of spending in Medicaid, seemed like, you know--if you take that as a benchmark, you know, it seemed not so terrible to me; whereas I feel like half of all spending being subject to these rules is really pretty drastically different and moves us a lot more toward a single payer--you know, national health almost.

Chairman WAXMAN. As I hear the testimony of the three of you, you all seem to agree that our report is accurate. Is that a fair statement?

Start with Dr. Schondelmeyer.

Mr. SCHONDELMEYER. Yes, it is. I think it is quite accurate. And I'm not sure it takes fully into account the effect of State supplemental rebates.

I would point out that your own State of California gets about 40 percent of their total drug spend back in rebates. And those State supplemental rebates are negotiated, not government-set prices. They are negotiated with the drug companies based on movement of market share and the same tools that the private Part D plans have available.

So why is it that States can negotiate up to a 40 percent rebate, an additional 20 percent on top of what the Federal rebate is, and the private Part D plans can only get 8 to 14 percent rebates? I don't know.

Chairman WAXMAN. And, of course, our report was only on the 100 most-prescribed drugs. There are other drugs beyond that, as well, for which there could be a greater savings or that we are paying far more for than we otherwise might have to.

Mr. SCHONDELMEYER. Could--but given what I know about the market and how rebates work, I would be willing to wager that there are even smaller rebates on the rest of the drugs in the market than the 100 you looked at.

Chairman WAXMAN. I see. Okay.

Dr. Anderson, what is your view?

Mr. ANDERSON. I agree that these numbers are quite accurate.

I think you have to look at it from a variety of different perspectives. One is from the 10-Ks and the 10-Qs, and you add up those that they report, you'll get to about \$2 billion. Then you sort of look at the differential in the prices between Medicare and Medicaid, and it is about a 20-25 percent differential. You do those and you get about a \$2 billion number.

So I think, from a variety of sources, we are seeing that your numbers are quite accurate; and I wish we had access, actually, to your numbers so we could look at them. As researchers, I think it is really important.

708 Chairman WAXMAN. And, Dr. Morton, as I heard your

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testimony, you confirmed the committeestaff's findings? You can't tell us exactly that we are correct because you don't have the same data, but you confirmed the fact that we're paying far more under Part D for these dual eligibles?

Ms. MORTON. That's consistent with what I know.

The States get supplemental--can negotiate for supplemental rebates. They get the best price on a brand and there is the inflation component, and Part D can't mimic those latter two. They can mimic the supplemental, but they can't get the inflation piece, for example.

And then, secondly, looking outside the drugs that you examined, I would actually think the rebates would be bigger for Part D. And the reason is--

Chairman WAXMAN. You would agree with Dr. Anderson?

Ms. MORTON. Yes. Because the big drugs for the duals

are largely in the protected classes where, as I said, there
is less ability to negotiate.

Outside the protected classes, you would expect more negotiation, more market share shifting and bigger rebates.

Chairman WAXMAN. These are protected classes because they are drugs that--there is no other alternative to those drugs and they are life saving; is that basically right?

Ms. MORTON. I think there is also a second factor, which is that you're trying to stop Part D plans from engaging in adverse selection, from cream-skimming in taking

healthy people. And if you offer only one HIV drug on your formulary, you're not going to attract the sick people.

Chairman WAXMAN. So we protect those classes of drugs, and it is important that we do so for the well-being of the people.

Ms. MORTON. That's right.

So in some sense that is why I suggest moving these guys back into Medicaid, given--if you're concerned, for this reason, about having a restrictive formulary, then, you know, that going to cost you.

Chairman WAXMAN. Thank you.

745 Mr. Davis.

746 Mr. DAVIS OF VIRGINIA. Thank you.

Of course, the problem is, these folks don't want to go
the back into Medicaid. But that is a political issue the other
that is a political issue the other
that is a political issue the other

Dr. Scott, let me ask you. We keep referring to private sector price controls that would result from Medicaid price regulation being extended to Part D. Can you elaborate on the expected impact of extending price controls to the Part D program on the following groups: employers, employees, unions and uninsured?

Ms. MORTON. Certainly. If you have--the best price provision of the Medicaid rebate rules is the critical thing. So if I, as a manufacturer, offer a low price to any

private-sector buyer, I have to offer that same--effectively, the way the rebate works--I have to offer that same low price to Medicaid. So the bigger--so that gets expensive as the group that gets that forced rebate gets bigger.

So as that group getting bigger and bigger, which it would be if you put in duals or all of Medicare or whatever, then I don't want to give a discount anymore, as a manufacturer, because if I give a discount to even one party, I have to give to the entire portion of the market covered by that best price. And that causes discounts for private-sector employers, for everybody else.

Mr. DAVIS OF VIRGINIA. The extension of the philosophy over there is just, why not just fix prices for everybody; that at the end of the day, if you fix prices, that somehow the drug companies are going to go along and just take it?

What you are arguing is, they make it up somewhere else along the way.

Ms. MORTON. Well, they are going to have an incentive to eliminate those discounts elsewhere in the economy and will move toward a more uniform pricing where everybody pays the same price and nobody can negotiate for a discount.

And that is dangerous, I believe, because the way we run our intellectual property is that these brands have patent protection, and the way to create price competition when two molecules have patent protection is to threaten to substitute

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one for the other and get a discount. If you can't do that because of the best price regulation, then you undermine 785 786 price competition. 787 Mr. DAVIS OF VIRGINIA. One of the problems with 788 Medicaid is that you don't get the same breadth of offerings, 789 isn't that right, that you would get Medicare Part D? 790 Technically, it is supposed to be an open Ms. MORTON. 791 formulary, but I believe the supplemental rebate States are 792 negotiating for depend most now on having a preferred drug 793 and then a list where the physician has to get prior 794 authorization to prescribe the drug, so that effectively 795 you're getting a narrow formulary. That's right. 796 Mr. DAVIS OF VIRGINIA. Dr. Anderson, do you want to 797 comment? 798 I would say, if you would compare the Mr. ANDERSON. 799 formularies between Medicaid and any of the private-sector plans, you would see that Medicaid has a much broader 800 801 formulary than most of the private-sector plans. 802 Mr. SCHONDELMEYER. I would agree with that. 803 Mr. DAVIS OF VIRGINIA. But they limit the number of 804 prescriptions that can be filled at any one time, right? 805 Mr. ANDERSON. Some of the States do have those as ways 806 to control expenditures, yes. But the formularies are quite extensive. 807

Congress essentially mandated that in OBRA '90 and

essentially said that all State Medicaid programs had to offer all drugs and have access provisions in there to make sure that they are available to all communities, all beneficiaries.

So it is quite an open program.

Mr. DAVIS OF VIRGINIA. But does a large formulary matter if you can't fill the prescription?

Mr. ANDERSON. Essentially, that is the problem of the States having not enough money in their Medicaid programs, and so they are making choices here as to how to save money; and I would not do that, but that's the choices that they have, given limited resources.

Mr. DAVIS OF VIRGINIA. Well, I know in Virginia we have gone from Medicaid, 10 years ago, being zero percent of the State budget to, now, 17 percent of the State budget. It has crowded out education and everything else. It is a huge--I wouldn't say completely unfunded Federal--but it is a Federal mandate that carries with it a lot of costs.

And, of course, States have to balance their budgets. We don't. There is just, I think, a huge problem.

Let me ask, long term on price controls; I'll ask each of you. Are you surprised to learn that in the first 4 years after the government mandated Medicaid price controls in order to control prescription drugs spending, that spending actually increased by 40 percent? Does that surprise

anybody?

Dr. Morton.

Ms. MORTON. I think spending on drugs--it doesn't surprise me, but it might be due partially to the best-price legislation that was passed in 1991, but it also might be due to technological change. We invent new drugs, people want to consume them. The population is aging, more people are on the disability rolls; we're just consuming more health care.

Mr. DAVIS OF VIRGINIA. Do prescription drug price controls hold down spending over time? I mean, immediately, obviously, price controls, we know they have an immediate effect; but over time, how does the marketplace reflect that?

Ms. MORTON. One of the things you have to realize when you're engaging in this kind of price regulation is that the manufacturer will have some kind of optimal response. So they will raise prices or alter their mix of drugs or change their forms or whatever, if that is going to get them bigger reimbursement. So that is one thing to keep in mind.

Then the second thing to keep in mind is just the research and development consequences. If we cut by half our spending on pharmaceuticals, then, you know, that's going to help us today, but it has consequences for future generations because we have privately funded R&D. And unless we're willing to think of some other way to do R&D, I think we have to make sure there is some money to be earned for somebody

who develops a novel therapy.

Mr. DAVIS OF VIRGINIA. Of course.

Mr. SCHONDELMEYER. Earlier, you asked all three of us to respond to the question, are we surprised that 40 percent expenditure increase occurred in the first 4 years. That is expenditure increase, not price increase; and the number of recipients increased in that time and a number of other factors unrelated to price.

Also I point out, you ask, do price controls result in lower prices or higher prices over time. I would point out, the other major governments around the world that do have price controls--I'm not saying we have to do that--but do have price controls, do pay lower prices than we do. So price controls for many markets in many governments seem to work.

The last thing I'd point out is, the United

States--today, our government pays for 50 to 60 percent of
all drugs in the U.S. We have become the largest buyer in
the marketplace. Whether you act as a regulator of price or
a prudent buyer in the marketplace, you're going to have an
impact in the marketplace. But I would say our government is
not working as a prudent buyer in a market--in a marketplace.
And there are behaviors that they can undertake that do
facilitate markets, but use the power of a 50-to-60 percent
player in a marketplace.

884 Mr. DAVIS OF VIRGINIA. Governments are rarely prudent 885 buyers is my observation. 886 Mr. SCHONDELMEYER. You guys can change that. Mr. DAVIS OF VIRGINIA. 887 I don't think you want Congress 888 to get involved. 889 Chairman WAXMAN. Thank you, Mr. Davis. 890 Mr. Cummings. 891 Mr. CUMMINGS. I have sat here and I have listened to 892 all of you; and I have got to tell you, I'm confused. 893 Because the bottom line, Dr. Anderson and Dr. Schondelmeyer, as I understand it, is that the government is spending more 894 895 money now, in moving these folks to Medicare Part D, than 896 they were before. Is that the bottom line? 897 Mr. ANDERSON. That's \$2 billion more per year. 898 Mr. SCHONDELMEYER. True. 899 Mr. CUMMINGS. Okay. 900 Now, maybe I'm missing something, but Mr. Davis, whom I 901 have tremendous admiration for, talked about "deja vu, here 902 we go again." But the fact is that Americans, hardworking 903 taxpayers that are watching this right now, are probably 904 sitting there scratching their heads and saying, Okay, what 905 does all this mean? 906 Now, Dr. Morton has given us a few suggestions. 907 I sat here and I listened to the suggestions, this is what I 908 asked myself. I asked myself, what is the problem with her

suggestions? And I want you all to answer.

One of the things she says, we should move the folks that are now on Medicare Part D--correct me if I'm wrong--back to Medicaid. Is that right?

Ms. MORTON. Just the duals. I mean, my understanding is, Mr. Waxman's concern is just the duals.

Mr. CUMMINGS. So that we won't be confused and the public won't be confused--see, what happens here in Washington is, people talk past each other, and so then--but when the bottom-line clears, we are still in the same predicament. And we'll be in the same predicament 10 years from now, but it will be far worse.

Is there something wrong with what she said? Is there an issue with that?

Mr. ANDERSON. Well, I think you could do that. The problem is that you want to have one program really be in charge for the person's health care, and that should be through the Medicare program or the Medicaid program. And by putting--in the past, they have been separate, so drugs have been part of the Medicaid program, and lots of other things have been part of the Medicare program; and that makes it much more difficult to get good, quality care.

So there are pricing reasons why you should follow her ideas, but there are clinical reasons why you might not want to.

Ms. MORTON. Now, can I say, the clinical side is not represented so well by our current system of a PDP and then a set of doctors who aren't part of the same organization.

I agree with you, but I think we could fix it for everybody.

Mr. ANDERSON. We should fix this for everybody, and essentially, potentially having separate payment systems makes it more difficult to solve it, because you want to have one system, one insurer really being responsible for the care of an individual.

Mr. SCHONDELMEYER. I agree that could work, to shift them back to Medicaid; but a downside of that is, markets work also based on the principle of volume, and larger volume should get lower price.

But here we have the government paying 50 to 60 percent of the drugs on the market, and paying a higher price and moving the dual eligibles from Medicare Part D back to Medicaid means that the government is dividing up their pie again to lots of smaller pieces, and essentially Medicare Part D does that. Instead of the government saying, we're going to pay for all Medicare Part D under one pricing system, we're going to let each plan and hundreds of these plans across the country negotiate prices. So we want a whole bunch of small people negotiating instead of one big party negotiating. So we structurally built into Medicare

Part D principles that fight against markets working well in ways that do derive better prices in the marketplace.

So we need to ask, should we keep them in Medicare Part D and find ways to better use the government's role in the marketplace.

Mr. CUMMINGS. Dr. Morton, I'm running out of time.
What was your second most powerful suggestion?

Ms. MORTON. I think that we need to study the protected classes quite carefully. I think what Mr. Waxman said about how these are vulnerable populations that are very sick and need access to correct drugs is absolutely right. However, when you give the plans no tools to shift market share or weak tools, then you are going to have expensive prices.

Mr. CUMMINGS. Dr. Anderson, would you react to that, please?

Mr. ANDERSON. Sure.

Essentially what we did when we passed OBRA '90 was, we said everybody in the Medicaid program had--for all the drugs, and so essentially you took out the ability to do formularies. But then you gave them the ability to do rebates.

So essentially what you'd want to do in these protected classes is to institute either the best price or the rebate system, so that when there is no competition, the Federal Government or the dual eligibles get the best prices.

Mr. CUMMINGS. Thank you, Mr. Chairman.

Chairman WAXMAN. Mr. Marchant.

986 Mr. MARCHANT. Thank you, Mr. Chairman.

Dr. Morton, in your testimony, you explained that expanding Medicaid, the Medicaid best-price requirement, to Part D would make prices more uniform across the board. Dr. Anderson seems to advocate uniform prices.

What would be the implication of a uniform prescription price policy?

Ms. MORTON. The implications are twofold. One is that because the production cost of these drugs is quite low relative to the research and development costs, it is worth giving them--it is worth selling at low prices to people who are poor or who can't pay, because you're still covering your manufacturing costs and you're extending the benefit of the drugs to those people. If you have to charge a uniform price to everybody, then those people can't afford it, they don't buy and you don't get as many people being helped. So it is useful to be able to sell at different prices to different consumers.

Secondly, plans--PBMs and insurers and HMOs--in this country have invested a lot in changing their organizations to be able to shift market share from one molecule to another, and that requires education of doctors and a lot of organizational effort. And that ability to shift market

1009 shares is what drives prices down, because it creates price competition between drugs. I buy A and you buy B. A and B 1010 1011 I get a good price on B; that is why I bought it. 1012 You get a good price on A; that is why you bought it. 1013 So your price on A is low and mine is high because we've 1014 engaged in this kind of bargaining. And if you make 1015 everything uniform, then all of that system of extracting 1016 price concessions is no longer worth doing. Mr. MARCHANT. Thank you. 1017 1018 Dr. Anderson, you seemed to express surprise that Part D 1019 prices are higher than Medicaid. Does any other payer in the 1020 United States get Medicaid prices? 1021 Mr. ANDERSON. Sure. The VA actually gets lower prices, 1022 DOD gets lower prices than Medicaid does in most cases. 1023 Mr. MARCHANT. Does GM get Medicaid prices despite their--the fact that they are a very large purchaser? 1024 I haven't--I don't have access to it. 1025 Mr. ANDERSON. 1026 That's where we need price transparency to know whether or 1027 not GM gets the same prices at Medicaid. We don't, as 1028 researchers, have access. My guess is that they do not, 1029 which is what I'm concerned about, that the marketplace for 1030 drugs does not seem to be working. 1031 All the discussion that Fiona Scott Morton talks about 1032 in terms of the marketplace is resulting in the private sector paying 20 percent more than the public sector. And 1033

why would I want to emulate a system where you're paying 20 1034 1035 percent more? Mr. MARCHANT. Well, it seems to me that someone in 1036 their 20s or 30, that had a disease that they felt like there 1037 1038 was a time horizon available to them for that disease or 1039 that -- to be cured with some kind of a medicine, would hope 1040 that the drug companies would not just flatten their product 1041 line to a price point, but would build something into the product line for profit and R&D, so that there would be some 1042 1043 hope later. And, of course, the government would have that 1044 hope, too. 1045 Mr. ANDERSON. And I would share in that hope. Right 1046 now, however, the pharmaceutical industry is spending 1047 anywhere from 14 to 18 percent of its revenues on R&D. spending 30 percent on marketing and spending 25 percent on 1048 1049 profits. 1050 So I would love them to increase the 1051 percentage--certainly as a researcher, certainly as a 1052 professor at Johns Hopkins--to increase them from 14 percent 1053 to 20 percent or 25 percent. But that is not what has 1054 happened, and as the profits have increased, the percentage 1055 has remained absolutely stable. 1056 Mr. MARCHANT. Ms. Morton, do you see a danger in that 1057 theory?

The marketing expenses of a pharmaceutical

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Ms. MORTON.

firm are all driven toward getting more revenue, which--and 1059 1060 those expenses wouldn't be spent if they weren't worthwhile 1061 in bringing in more revenue, so that increases the incentive 1062 to invent something. The more revenue you can collect from 1063 it, then the more incentive you have to invent it. So the marketing, per se, is not a disaster. 1064 1065 Profitability is very difficult to calculate here because the 1066 percent profit has to be calculated on something--percent of sale, percent of assets, percent of whatever--and typically 1067 1068 we would do it as percent of assets. And R&D is an asset for 1069 these firms, but it is not counted as such when the accountants look at assets. 1070 So pharmaceutical companies look 1071 like they have tiny assets and few factories when, in fact, 1072 they spend millions on R&D. 1073 So I'm just always leery of profit numbers, because they 1074 can--you can calculate them so many different ways. 1075 Mr. MARCHANT. Thank you, Mr. Chairman. 1076 Chairman WAXMAN. Mr. Yarmuth. 1077 Mr. YARMUTH. Thank you, Mr. Chairman. I want to thank 1078 all the witnesses for their testimony.

I think we are in general agreement that the treatment of dual eligibles through Medicare Part D is costing the government and the taxpayers more money than it otherwise would. And the staff report estimates that the savings to the taxpayer down the road, or the additional cost to the

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1084 taxpayer for failure to do something different, would be in the neighborhood of \$85 billion over that 10-year period.

Dr. Schondelmeyer and Dr. Anderson, does that seem like a reasonable estimate to you? Is that possible? Is that understating it?

Mr. SCHONDELMEYER. I think if the program continues as designed, that is a reasonable estimate. But I would point out that it is probably even more than that because the States have gained even more in their supplemental rebates in the last year or two, and I think the savings could be even greater than what that represents.

So it is probably a reasonably accurate estimate if not an underestimate.

Mr. ANDERSON. And I would agree.

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Mr. YARMUTH. And it is possible, because the States have the protection of the inflation cap, essentially, it could be more than that in terms of savings if the inflation rate ended up being significantly higher as it has been in many years.

I think in OBRA '90, that was a very Mr. ANDERSON. smart thing to include in there, to put it in, because when the drug companies increase the prices, then essentially the Medicaid programs gets the advantage of that. And that doesn't exist in Medicare Part D.

Mr. YARMUTH. Dr. Morton, you said in your testimony

1109 that the result of the study, the staff study, the staff 1110 report was predictable given what we're talking about. 1111 Would you say that the impact that we've seen over the 1112 last few years was predictable when the legislation was 1113 passed to create Medicare Part D? 1114 Ms. MORTON. Certainly, the magnitude, I wouldn't have 1115 wanted to speculate on. But the fact that Medicaid has a required best-price provision for brands and then the 1116 inflation component on top of that makes me think that it 1117 1118 would be extremely difficult for a private sector--I mean, it 1119 would be impossible if the Part D plans were included in the 1120 best-price provision. 1121 But actually they are exempted, so you could give Part D 1122 a low price, and it wouldn't trigger a Medicaid rebate. 1123 But having said that, I still think it would be very 1124 difficult to match the Medicaid price. 1125 Mr. CUMMINGS. [presiding.] Mr. Bilbray. 1126 Mr. BILBRAY. Mr. Chairman, with your condolence--I 1127 mean, your support, I'd like to yield my time to the ranking 1128 member. 1129 Mr. DAVIS OF VIRGINIA. He is always happy to give you 1130 his condolences. 1131 Dr. Schondelmeyer, let me ask you. Prior to 2006, dual-eligible seniors who qualify for both Medicare and 1132 1133 Medicaid had prescription drug coverage through Medicaid.

1134 | course, now they're moved into the Part D.

The majority report argues that by moving dual-eligible seniors from Medicaid price controls to Part D market prices, prescription drug companies receive a financial windfall.

Do you disagree with CBO's assessment that mandating Medicaid price controls in Part D would increase the cost of drugs to all other private payers?

Mr. SCHONDELMEYER. I haven't looked recently at CBO's assessment or quantification of that.

I would point out that the Medicaid rebate is partly based on the best price, which comes from a price negotiated in the marketplace. And it means that there are at least one--

Mr. DAVIS OF VIRGINIA. The total marketplace or a restricted marketplace?

Mr. SCHONDELMEYER. In various buyers in the private marketplace.

So there is at least one other buyer in the marketplace that is smaller than Medicaid and smaller than Part D plans that have negotiated a better price. And I find it contradictory that the larger Part D plans can't negotiate similar prices in the private marketplace that the best-price buyer--so I would argue that not all of the prices are regulated.

I would give you that the mandated rebate amounts are

set by government law or regulated, but any rebate above and beyond that is affected by the best price of negotiations in the marketplace.

Mr. DAVIS OF VIRGINIA. Well, I'm going to ask unanimous consent that the Congressional Budget Office's letter to Senator Stabenow, stating that including a best-price requirement in Part D would put upward pressure on prices paid by the VA, Medicaid and private purchasers, be included in the record.

1168	RPTS COCHRAN
1169	DCMN MAYER
1170	[11:10 a.m.]
1171	Mr. CUMMINGS. So ordered.
1172	[The information follows:]
1173	****** COMMITTEE INSERT ******

1174 Mr. DAVIS OF VIRGINIA. Dr. Scott Morton, let me just 1175 ask you. You have to look at the marketplace as a whole; 1176 isn't that right? When you are cutting in one place, don't 1177 costs somehow rise -- the drug companies, or whoever, in their 1178 marketplace are going to make allowances for that? 1179 Ms. MORTON. Yes. 1180 I think we have a problem in our country because, for 1181 our government purchases, we tend not to like to say we will 1182 pay \$2.43 for that pill. We like to say we are going to pay 1183 as much as the private sector pays, or 15 percent less than 1184 the private sector, or we are going to pay as much as Canada 1185 pays. 1186 And then the problem for all those sorts of reference 1187 prices is that industry then would like -- if they can move the 1188 reference price, they can shift how much Medicaid and 1189 Medicare pay for their drugs. 1190 So if we say "average prices," then the private sector 1191 prices are going to go up, because that is what triggers--1192 Mr. DAVIS OF VIRGINIA. It is kind of like everybody 1193 taking the lowest seat price on the airplane. If everybody

Ms. MORTON. They would raise the lowest price. That lowest price price wouldn't be where it was before.

would be in worse shape than they are.

paid the lowest price that somebody pays on an airplane, they

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Mr. DAVIS OF VIRGINIA. And that is basically the

argument here, as I understand. It is economics that I took. 1199 1200 Mr. ANDERSON. But I am not sure why the Federal 1201 Government should pay the highest price. 1202 Mr. DAVIS OF VIRGINIA. Well, they don't in many cases. 1203 Mr. ANDERSON. They don't. But essentially--1204 Mr. DAVIS OF VIRGINIA. Dr. Morton, do you think the 1205 government is paying the highest prices? 1206 They don't pay the highest prices. In fact, Medicare, 1207 Part D, the increases are way below what was initially 1208 estimated as we bring some marketplace into health care. One of the problems today is the Federal Government is such a 1209 1210 large buyer, you don't have basically a market in some of 1211 these places. 1212 Dr. Morton, would you react to that? 1213 Ms. MORTON. I think you said it correctly before, 1214 Gerry, when you said that Medicaid pays the lowest and then Medicare and then the private sector. So I think the private 1215 1216 sector is paying the highest prices, and the danger of having 1217 a best-price provision that extends to a large group of 1218 consumers is that those prices go up. 1219 Mr. DAVIS OF VIRGINIA. So are senior taxpayers paying unfairly high prices for prescription drugs in Part D? 1220 1221 Ms. MORTON. I think--since I am an economist, I am not going to comment on the "unfair" part. I think my own 1222 research shows there is a huge benefit to moving the 1223

1224 cash-paying uninsured into a plan, okay, because then you 1225 have someone larger working on your behalf. Mr. DAVIS OF VIRGINIA. They are the ones that took the 1226 1227 brunt of it, aren't they, before this? Ms. MORTON. Our data show that is a big effect. Moving 1228 into a plan, having been uninsured, means you get access to 1229 1230 much better prices, and of course, your utilization goes up. 1231 Mr. DAVIS OF VIRGINIA. You would agree with that, wouldn't you, that the biggest beneficiaries of this are the 1232 1233 uninsured, the poor, in terms of moving them into Part D, 1234 that they get a great reduction? 1235 Mr. ANDERSON. Oh, absolutely, the same thing as, we 1236 should try to cover the uninsured in the United States. I 1237 mean, we want to cover as many people as possible. 1238 absolutely you want to do that; you just don't want to pay more than you need to pay for services. And I think that is 1239 what this committee's report shows, that you are paying too 1240 1241 much for services. And \$2 billion is \$2 billion. 1242 Mr. DAVIS OF VIRGINIA. Are they saying too much, or are 1243 they saying they are not paying what Medicaid pays, which is 1244 clearly the lowest? I think there is a difference between 1245 "too much" versus what Medicaid pays. If you argue that everything over Medicaid prices is too 1246 much and you put Medicaid prices across the board, it 1247 1248 couldn't happen, could it, economically? Wouldn't it raise

1249 Medicaid prices?

1250 Mr. ANDERSON. I don't think it would raise Medicaid 1251 prices.

Mr. DAVIS OF VIRGINIA. So if you think the drug companies, across the board, charged everybody at Medicaid rates, that life would just go on and there would be no ramifications throughout the system?

If that is your opinion, that is fine.

Mr. ANDERSON. They still would be paying more, the United States would still be paying more than Canada would be paying. You would have to bring the rates down to VA in ordered to get down to Canada or U.K. or French rates.

Mr. DAVIS OF VIRGINIA. One thing we have with the U.K. is you do not have--and a lot of veterans have complained about this--you don't have the choices in VA because not everybody is bring their costs down to those levels.

They can't afford to sell their drugs at that level, isn't that correct?

Mr. ANDERSON. Well, they essentially have a formulary, and within a therapeutic class they will have a one-drug, which is exactly the same thing that the Part D plans have; they don't offer every drug. It is Medicaid that offers every drug.

Mr. DAVIS OF VIRGINIA. I have one more question.

Dr. Scott, when proponents of a national formulary are

confronted with the counterargument that a structure would limit seniors' ability to get drugs, their response is often that seniors can just appeal the decision.

I would ask, are the lower prices on formularies only achieved by the ability to move market share?

Ms. MORTON. My understanding is, that is the main reason why you get a low price, that you can promise to move market share. And if you are a senior and you look at PlanFinder, for example, in the Part D context, you can see which plans have a preferred--have a good price on the drug you are interested in. If it is A versus B, you can see that, and then you can join the plan that has the low price on the one you want.

Mr. DAVIS OF VIRGINIA. On the one you want, you get more choice. Thanks.

Mr. CUMMINGS. Mr. Sarbanes.

Mr. SARBANES. Thank you, Mr. Chairman.

Whenever we have a hearing on the pharmaceutical industry or drug pricing, I feel like I am in a magic show because it is all sleight of hand. I mean, it is incredible, the questions.

When you say, well, if the price is this much higher than you would get in another way, isn't it really lower because of X, Y and Z? I mean, people see that the prices are higher. The report makes it clear that we have spent \$2

1299 | billion or \$3 billion more as taxpayers than we needed to.

By the way, yesterday we were considering trying to get full funding for the LIHEAP program, which is the Low Income Home Energy Assistance Program. The cost of that is about \$2 billion to \$3 billion. Just so people understand, when you lose that much money that the taxpayers have put forward, you can't do other things that we ought to be doing to help people.

To me, this is a classic case of, if it's not broken, why would you fix it? Not only is it a chief criticism of the Medicare Part D program that you didn't take advantage of the opportunity to create a beneficiary pool that could negotiate in a significant way with the pharmaceutical industry directly, but in fact with the dual eligibles, what you did was, you took 6 million people out of a pool that was in a position to negotiate directly with the pharmaceutical industry and you put them into a place where they couldn't.

Not only that, you took a system where you had PhRMA on this side, the pharmaceutical industry on this side, the beneficiaries on the other side, and you interposed the insurance companies and the insurance plans and insurance industry in the middle, which is notoriously inefficient in terms of its administrative costs.

So you took a situation where you were paying 3 to 5 percent overhead administrative costs through the Medicaid

program; you put in the middle of the stream, the dollar stream, a system that has got overhead costs of about 17 to 20 percent, right--which is very inefficient--which is a great result for both the pharmaceutical companies who now get all this interference run between them on the pricing, right, so you can hide the ball very easily, and it is good for the insurance companies, who get to come in here and charge these huge overhead costs.

It is absolutely madness.

So my first question is, what was the reasoning? What possible rationale was offered up to justify taking the dual eligibles and moving them from Medicaid as the payer to Medicare as the payer?

Mr. ANDERSON. Well, I think it was, as I explained to Mr. Cummings, that essentially you wanted to have them in one system, and that would be the Medicare system as being the controlling system for insurance. And what it meant, unfortunately, is a \$2 billion windfall to the pharmaceutical companies.

Mr. SARBANES. That is a neat idea to get them into one system, but you could move them into one system that works or you can move them into one system that doesn't work. So what they did was, they moved them into one system that they made sure wasn't going to work by setting them up in a way that we couldn't negotiate.

Mr. ANDERSON. Well, essentially, you put them into a system with 20 percent higher administrative costs and paying 20 percent higher prices, and then trying to say "provide good care." And that is really hard, because you are down at 40 percent already.

Mr. SARBANES. Isn't central to this the fact that the Medicare Part D program is not a directly administered program? You have Medicare Part A, which is directly administered for hospital benefits. You have Medicare Part B, which is directly administered for physician services. You have got Part C, which is a managed care program, which isn't working so well.

But Part D was not designed that way. Part D is not directly administered. Part D is a subsidy to the commercial industry, which has all of these inefficiencies in it. So why you would want to set it up that way, who can imagine?

Now, on the price control thing, we keep talking about price controls, but you put it better. This is really just about a customer called the U.S. Government that goes into the marketplace and has a lot of bargaining power, presumably.

Do people have to sell? Do insurance plans that provide drugs and prescription drugs, do they have to sell to the government, are they required to sell to the government? Or do they want to sell to them because it is a big pool of

1374 beneficiaries that they can make money on?

Mr. SCHONDELMEYER. They don't have to sell to the government. And I would point out, when we talk about using market share movement under State supplemental rebates or VA, we call it "price controls." When we talk about using market share movement under Part D private plans, we call it "the market." It is the same mechanism.

So you can't call VA's--VA gets a lower price largely because it is a closed system and a very tightly controlled market share movement formulary, and that works. And Medicaid did that because they could do that much better than the Part D plans are right now.

Mr. SARBANES. If we are going to pray at the altar of market economics, we ought to at least bring the basic principles of how you negotiate in the market to the table, right?

1390 Thank you.

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1391 Mr. CUMMINGS. Mr. Issa.

1392 Mr. ISSA. Thank you, Mr. Chairman.

Market distortion is a serious concern, and I think for all three of you, you have been trying to deal with it--perhaps in different ways.

Because, Dr. Morton, none of you are here to make a political statements, I will make a short, simple one to open this up. I come from California, where we mandate prevailing

wage. I come to Congress where we vote back and forth and debate and argue, over partisan lines, prevailing wage.

Now, prevailing wage, in at least this Congressman's opinion, is distorted, so we pay a lot more to build our homes--not our homes, but our schools and our roads, at least in California, than we would pay if the large buyer, this \$150 billion entity called California, went out and went to the low bidder and said, you know, You don't have to pay higher wages to build roads just to please the State of California. So I want to be sensitive here that we don't send that message from the government.

Dr. Morton, I will start with you. VA is a buyer-seller, we want a good price, and we may not buy every drug if it isn't the best price, or we may not dispense two competing drugs as often, if it is more expensive. Would you say that that was, as an buyer, as a government buyer, a fair market relationship as an economist?

Ms. MORTON. Yes, I think it is, and I think it is something that most Americans think that they don't want, that they would like something better than that, because it is a very tight formulary.

Also, there is no retail component. So the VA pulls its truck up to the factory, gets the drugs and brings them to VA hospitals. You can't go down to your local pharmacy and get a VA-dispensed drug.

Mr. ISSA. Dr. Morton, I happen to have Indian health care in my district, quite a bit of it, and they get that rate, and they are thrilled to get it. And my centers, my Native Americans, take advantage of it. And by the way, they also look, in some cases, to buy outside those formularies, and they pay a lot more, but they do it with discretion because of the obvious price advantages.

When we are looking at Medicare Part D, as we are here today, is it fair to say from a pure economic standpoint that if you take VA's advantage of single buying, low administration, back-up-the-truck-to-the-dock, that in fact you're going to spend more when you offer people individual, broad formulary choices and you add the administrative burden, that it is essentially where you are, not where we are? And have you ever calculated that cost?

In other words, if we were to take--because I want to do a reality check on whether or not we are distorting and whether or not we are paying too much. If you take the VA rate and you take those elements, where should you end up as a hypothetical for Medicare Part D and where do you end up?

Ms. MORTON. That is a really good question. I haven't done that calculation, but that is exactly the right way to think about it. And part of what makes this difficult is that I know that there are some components; most of these Federal agencies have some component of mandated discounts

and some component of "we negotiated it because we have a tight formulary." And you would want to just look at the cases where it is negotiated, as opposed to mandated.

Mr. ISSA. Let me ask a question I think for all three of you, because this is of interest to me.

Obviously, when we deal with seniors and we deal with drugs developed only for seniors in America, we are dealing under Medicare Part D, Medicare in general, that is the market.

So my question is, how does the United States

Government, in each of your opinions, ensure that drugs which are geriatric in nature only are fairly priced if there is very little alternative way of buying it, other than our VA seniors? Except for that group for the most part, some of these things have no other market in the U.S.

So each of you, have you thought about how we get the fair interpretation? Because I am here today believing that I can't use Medicaid because it is a distorted market. I can use VA, but I have to add those costs that I mentioned with Dr. Morton. So if that is all true, when I have a drug that is limited in its reach. Other than seniors in VA and Native Americans, how do I fairly make sure that the price is achieved?

Mr. SCHONDELMEYER. Actually, we have at least one drug that falls into the category you described. There is a drug

called Epogen that 80 to 90 percent of the market is the government, and the government is the only payer. So there really is no such thing as a market-based price, because the government is the monopolistic buyer in that market; and the government does set and establish the payment rates for that drug, and they come up with the value of, here is what it is worth.

I think Dr. Morton earlier said the government is afraid to say, here is what we will pay for a drug. But on the one hand, they do try to do that, but any time they do that, we call it "control" rather than "market behavior."

I think we have to look for the line between price regulation and prudent market behavior for government. Let's focus on the prudent market behaviors and try to avoid the regulation that drives up the price. But I think you can do prudent buying as a large buyer government and keep some element of market in place.

Mr. ANDERSON. We have done that. Just to explain in a little more in detail, for SRD and renal disease drugs, I think we have gotten good value for those, and we have essentially with a government-administered price.

The other thing, Mr. Issa, I would suggest, is the United States is not the only place where there are seniors. There are millions, billions of them around--a billion of them around the world, and pharmaceutical companies are not

just selling to the United States, but they are selling to
the U.K. and Canada and other places as well; and we have to
recognize that.

Mr. ISSA. Dr. Morton, quickly.

Ms. MORTON. I would say, one of the things that you will get in Part D is this same substituting and bargaining, and I can shift share from A to B. So if your drug for seniors has substitutes, therapeutic substitutes, then I think you can trust to a PBM or a Part D plan to be able to extract discounts on that drug.

I think the very difficult question, which we aren't facing at the moment so hugely, is what happens if somebody invents a pill that cures Alzheimer's, and it is the only one, or something like that? Then really the government becomes the only buyer, and there is no good substitute, and how are you ever going to get a discount in that circumstance?

But as long as there are therapeutic substitutes, they buy like everybody else buys.

Mr. ISSA. Thank you, Mr. Chairman, for your indulgence. Hopefully the follow-up will be how government gets better if we are going to set prices. Obviously, it hasn't been one of our strengths, but I look forward to working with you on that.

Chairman WAXMAN. [Presiding.] Thank you, Mr. Issa.

1524 Ms. Watson.

Ms. WATSON. Thank you very much, Mr. Chairman, for this hearing. I want the witnesses to know we value your input. I am concerned too about the real cost of these drugs and the increases, so--I have heard you allude to a way we should really model this. Can the three of you explain more how the government can model the Part D drug program so that it really works for seniors?

A big smile there. What does that mean?

Mr. SCHONDELMEYER. Well, first, the point that was brought up by Dr. Anderson: price transparency in the marketplace. The basic issue, that we don't see how much rebates are flowing without having a congressional investigation in the Part D program, to me, tells us that is not a market. We are going to hide behind the black box and do what we want, and you guys pay the bills.

So we need to have price transparency and transparency of the flow of dollars in this marketplace. Markets work with information. When you hide information, markets cease to work properly. So one is that price transparency.

Second, I think, look to the Medicaid programs and especially what States are doing in their State supplemental rebates and obtaining these much larger discounts above the already-mandated Federal Medicaid rebate and say, How can you use those mechanisms or apply those to the Medicare Part D

1549 plans; and ask why--the Part D plans, why aren't you 1550 negotiating the same kind of rebate? If this is a market, 1551 why can't you get the same level of rebate out there? 1552 And then look to see, are there reasons, maybe reverse, 1553 perverse incentives, that keep these Part D plans from 1554 wanting to get more rebates from the drug companies. 1555 I would argue that no one in America is really managing 1556 or regulating prices very well, whether it is government regulated or private regulated. What we do is, we get bigger 1557 1558 discounts and rebates, but the top keeps floating up faster 1559 than inflation by a factor of two to three times the 1560 inflation rate, every year, year after year, no matter what we do. 1561 So prices keep going up no matter what we have done, and 1562 we fool ourselves into thinking getting more rebate dollars 1563 1564 back is saving us money. It really isn't. It is not 1565 controlling the net we pay overall in the first place. Rebates are simply a loan to the drug companies for 9 to 1566 1567 12 months, and then we collect the money back and spend a lot in administrative costs doing so. 1568 1569 Mr. ANDERSON. I think if are going to have a 1570 marketplace, we have to have price transparency. We don't 1571 have price transparency in the pharmaceutical, whereas we are pushing it in the physician market, we are pushing it in the 1572

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hospital market.

But I am not sure that we can ever get good prices when we have given the drug companies substantial reasons not to negotiate prices, and that would be the patents that we have given them for up to 17 years. This essentially takes away their reason for negotiation.

So I think what we have to do is take a look at what other countries are doing in this area. They are paying about half the prices that we are paying for pharmaceuticals, in other countries, and that is why Americans are going to Canada and other places for these things. So one of the things the Medicare program could do--and I know many of you voted on this a year ago--is to have the Medicare program negotiate directly with the pharmaceutical industry in order to get a best price.

The other thing that I would just add to that is, I am not sure why the VA, the Medicare program, the prisons and all the other places don't negotiate. I don't understand why the government pays different prices for exactly the same drugs, depending on whether it is a prisoner who needs it or somebody who is part of the community health center or somebody who is the Medicaid recipient.

The government should be paying one price for drugs.

Ms. MORTON. I am a little less enthusiastic about transparency than my colleagues, because I think in the context of Medicare Part D, if I am a plan and I am

negotiating hard in a particular class and I get a good deal on drug A, I don't really want to publish that for all my competing plans to see. And they might in fact be negotiating on drug B and drug C. So there is going to be differences across us, and the plans are going to be trying to get that lowest price as a way to lower their costs and attract more seniors.

So requiring manufacturers to publish that price is going to lead the manufacturers to be less willing to give those discounts and less willing to price aggressively. So I worry about transparency.

Secondly, I completely agree with Dr. Schondelmeyer in terms of the supplemental rebates the States are getting through Medicaid being a good model, but that actually is what Part D is doing. They are negotiating those rebates based on preferred drugs on a formulary. And what they can't do, which Medicaid can do, is get a best price or an inflation component, which are big parts of the discount that Medicaid gets.

Then, lastly, I would say--Dr. Schondelmeyer said, why can't Part D do some of these supplemental rebates, negotiate for lower prices? Part of the reason Part D can't is because there are protected classes, and in these protected classes, the plan is restricted from making a drug preferred and saying, You have to consume this HIV drug instead of that

other one until there is a medical need for you to switch. 1624 1625 And when you have that kind of restriction, then it is not possible for the plan to negotiate aggressively and get a 1626 1627 discount. 1628 Now, there are good reasons for having those restrictions, but I am just saying those restrictions are 1629 1630 expensive. 1631 That is exactly when government Mr. SCHONDELMEYER. 1632 needs to step in, is when you have on the one hand, the 1633 market should work by negotiating lower prices and preferring 1634 one drug over another, but on the other hand, it is clinically not appropriate. 1635 1636 Government has a role in that, and that is why you are 1637 here, and you do have a role in establishing a mechanism to 1638 deal with something the market can't do effectively. 1639 Chairman WAXMAN. Thank you very much, Ms. Watson. 1640 Mr. Shays. 1641 Mr. SHAYS. Thank you, Mr. Chairman, for having this 1642 hearing. 1643 I am struck by the fact that Medicare Part D is about 1644 \$40 billion and Medicare Part A is about \$220 billion; and we want to save money, but we are having a hearing on the 1645 1646 Medicare Part D program. I think we should, and I think we 1647 should because I think it has worked, frankly, phenomenally 1648 well.

1649 For years, politicians talked about having a 1650 prescription drug program, and in 2003 a Republican Congress, believe it or not, passes a prescription drug program. 1651 1652 program they wanted was going to cost about \$400 billion over 1653 a certain period of time, and the Democratic program was 1654 going to cost \$800 billion. I chose the less expensive plan 1655 because I thought it would cost twice as much when we finally 1656 adopted it, because most programs that we pass under Medicare 1657 turn out to be twice as much as the estimate. 1658

And, believe it or not, it is like one-third less than it was going to be, not twice as much.

Dr. Anderson, when you come in with a beaming face as though you have made this great discovery that those products that are controlled may be less expensive, I say, Whoopie, you are exactly right.

I would like to make a proposal. Do you ever get any Federal grants?

Mr. ANDERSON. I do.

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Mr. SHAYS. I want to save the government money. How much do you get paid as a salary?

Mr. ANDERSON. \$175,000.

Mr. SHAYS. I want you to only accept \$50,000. I am going to tell you that is what you get for that grant. I want to save the Federal Government money. But we don't do that, because we want you to have your talents and we want

1674 you to have your creativity. But we don't control what you 1675 get, at least I don't think we do.

We do it with doctors. That is not negotiation; that is, take it or leave it. They are underpaid; our doctors get less for the service than it costs them, but we act like somehow this is a great program because we have price controls.

Tell me why I shouldn't be grateful that this program costs less than it was supposed to cost, that the seniors who are in it have nine out of ten--excuse me, 85 percent satisfactory rate--and nine out of ten who are part of the dually eligible don't want to go back into the old system nor do the States want them to go back into the old system.

Nobody wants to go back into the old system, But you are using that as a price comparison.

Mr. ANDERSON. First of all, you said the \$400 billion. If you look at Kerry Weems' testimony that he is going to give today and you add up the numbers of the expenditures that are projected, you will see it is \$400 billion. So essentially you talk about a 30 percent reduction; but essentially when you voted on the bill, it was \$400 billion--

Mr. SHAYS. You are talking about a shifting 10-year time frame. Let's talk about the same numbers we were using when we did it, compare apples to apples.

Mr. ANDERSON. Right, I think that is what we have got.

1699 Mr. SHAYS. Sir, you are not.

1700 Mr. ANDERSON. We will take a look at that.

1701 Mr. SHAYS. Dr. Morton, what is your comment?

Ms. MORTON. I just wanted to say that underlying all of this discussion, we should remember that pharmaceuticals are really unusual, because the research and development that was used to produce the drugs we are consuming today occurred 15 or 20 years ago.

So part of the problem is, if you say to a doctor, We are going to reduce your salary from \$200,000 to \$100,000, they can take it or they can drive a taxi. And if they go to drive a taxi, then we have no more doctors left. And that constrains what you do as a body for paying for physicians.

Mr. SHAYS. I know how we can build twice as many bridges. We will just pay the construction workers half the price. But I don't believe in that, and I am for the prevailing wage. But here we have a competitive model that is working.

Ms. MORTON. I am sorry. I just want to say one thing.

So the thing about the drugs is that if I say today, as Congress, I am going to pay half as much as I was paying yesterday, that drug is already invented. It costs a tiny amount to manufacture, so, of course, the drug company is going to sell it at half the price.

Mr. SHAYS. Let me ask you about price controls. I went

to California about 15 years ago, and a company was
developing something to slow the beginning stages of
Alzheimer's. They spent \$800 million.

I checked 2 years later, they had spent about \$200 million more and it failed; they lost \$1 billion. But they told me at the time they wouldn't have spent a darn penny if they had price controls.

And it seems to me this is really a debate on whether we with we have price controls or not; that is what it is really about. And I don't buy into price controls. I think what we will have is less discovery. I think we won't have the drugs that we see today.

And if you disagree, either one, tell me why.

Mr. SCHONDELMEYER. First of all, your statement, or the framing of the issue, isn't exactly correct, because price controls were in effect. If you call Medicaid rebates pricing controls, then they were in effect and they did spend the money, and VA price controls were in effect and they did spend the money.

So I find the statement that if price controls were in place, we wouldn't have spent the money to be a little bit specious of an argument, because there were price controls, by your definition.

Mr. SHAYS. Excuse me, you don't believe that when we tell doctors, this is the payment, that is not a price

1749 control? Do you really think we negotiate with our doctors? 1750 Mr. SCHONDELMEYER. No, and the same with pharmacists and others in California. The States cut the fees. 1751 1752 Mr. SHAYS. Do you think we negotiate with our doctors, 1753 or do you think we basically say, this is it? 1754 No, it is take it or leave it. Mr. SCHONDELMEYER. 1755 Mr. SHAYS. Yes, it is price controls. 1756 Mr. SCHONDELMEYER. But it is different. I would point out, there is not a best-price provision for doctors like 1757 1758 there is in the Medicaid State rebate programs, and there are 1759 not State supplemental rebates like there are. 1760 So there are some aspects of this that are market based 1761 in terms of the prices Medicaid pays. It is not all just to 1762 fix, we will only pay this. Chairman WAXMAN. The gentleman's time has expired. 1763 1764 Ms. Speier. 1765 Thank you, Mr. Chairman. Ms. SPEIER. 1766 This lively debate is interesting to me because I 1767 believe that California is a great example. The Medicaid 1768 system in California is one in which we have historically 1769 negotiated rebates and discounts in the Medicaid system, and 1770 they have been healthy discounts. And the pharmaceutical 1771 companies have flocked to California because it is a great 1772 universe from which to sell their product, and there has been great competition there. 1773

1774 So, I guess my question is -- and I would disagree a 1775 little bit with what my colleague has just said--if you look 1776 at how many dollars are actually spent on R&D, at least 1777 historically, the majority of those dollars have come from the taxpayers of this country and NIH grants, if I am not 1779 mistaken. So it is the government that funds the lion's share of this research that goes on.

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All the other industrialized countries in the world have price controls in effect, and we end up subsidizing the prices of pharmaceuticals in these other countries.

So, I guess my question is, you have spoken a lot about transparency. But in trying to identify which is more important, just lifting the language in the bill that was passed by Congress, it says that the Federal Government can't negotiate.

Isn't that the most important thing we can do in terms of trying to bring the costs of these drugs down?

Mr. ANDERSON. I think it is, in fact, the most important thing, and I would strongly support that as an I mean, it is very close to what the other countries are doing, as you suggest; and I don't understand why we want to be spending twice as much for drugs as other countries are spending.

Mr. SCHONDELMEYER. Also, we can look at both the market and other things that have worked. State supplemental

1799 rebates are negotiated and operated by States on behalf of the entire Medicaid program within the State.

Somebody earlier referred to General Motors or large corporations and their behaviors. I don't see General Motors turning over their drug benefit to each local plant and telling each local plant, you go out and negotiate drug prices on your own.

Hey, centralize it and do it centrally.

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The equivalent of that in terms of Medicare would be for the Federal Government to use State supplemental rebate negotiation tactics on behalf of all Medicare Part D programs and then pass the benefit on to those local Part D plans out there.

So we see in the private market centralized behavior, large prudent buyer behavior and using the market to work. And I think the government can do that and be a prudent buyer and not be a price regulator, per se.

Ms. MORTON. The problem I see with that is, if Health and Human Services negotiates directly with pharmaceutical companies, it depends on your interpretation of the word "negotiate."

If you are going to say, I am a large buyer, I am the Secretary, I mandate you give me 20 percent less, of course, that is going to work. If you say, I would like you to give me 20 percent less, then the question is, why?

A regular plan says, I want you to give me 20 percent less because I am going to consume your competitor if you don't. I am going to consume drug A if you don't give me a price cut on drug B.

The Secretary presumably wants to include all drugs, doesn't want to tell American seniors, you can only have drug A and you can't have drug B. So if the Secretary can't exclude somebody, then I don't quite understand how they negotiate a lower price. I understand how they instruct, you will give us a lower price.

Ms. SPEIER. Well, California has a MediCal medical formulary, and drugs get on or off the formulary, and, you know what? They do make those decisions.

Furthermore, these drug companies want to make sure their drug is on the formulary. So it is not like it is so much an exclusion as much as it is, we want to be on your formulary and we will give you this.

Ms. MORTON. Right.

But California Medicaid is excluding some drugs, and the people in California Medicaid are getting this benefit for free, and they don't really have the ability to complain and say, "I would like a choice of all cholesterol drugs," whereas I think seniors and employed people expect to have more choice in their formulary or choice of cost plans.

Ms. SPEIER. I have a mother on 15 drugs right now. She

doesn't know which cholesterol-busting drug is the best. She is on three or four of them.

So I think it is kind of--it doesn't make a lot of sense to say that these seniors want these drugs. They tend to want the drug that they have been on, as opposed to wanting some drug. And if we didn't have direct-to-consumer marketing, we would have a whole lot better system in this country to start off with.

Mr. SCOTT MORTON. So suppose you have a national formulary. They have been on drug A all the time; they arrive at Medicare, and the Secretary has negotiated a good price on B, and that is it. The question is, what does the person do at that point?

That is a system we could have. That is what the Government of France does.

Ms. SPEIER. You know what it is called? It is called prior authorization. We have done it in California, and it has worked. For that individual who does better on the drug that is no longer on the formulary, you can still have that drug, it just needs prior authorization.

Frankly, that is what we should be doing on the Federal level. It is not like it hasn't already been done. It is done, it is done effectively, and it saves a lot of money.

Ms. MORTON. And Part D plans do that.

Chairman WAXMAN. The gentlewoman's time has expired.

1874 Mr. Burton.

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Mr. BURTON. My first wife, who died 6 years ago, was 1876 taking chemotherapy in Indianapolis. And there were two women sitting there next to her, they all had the needle in their arms taking their chemotherapy. And one of the women was saying -- she was actually complaining because, she said, My Tamoxifen costs so much, I can't afford it; it is \$325 a month.

And the other lady said, I am getting mine for \$50 a month.

And she says, No, that can't be right. And I am sitting there as a legislator, and I said, No, that can't be right.

And the lady said, No, I am getting it from Canada for with about one-sixth the cost of what it was in America.

I held hearings on this when I was chairman of the committee, and I couldn't figure out why, right at the border between Canada and the United States, you can go across the border and get the same pharmaceutical product for one-fifth, one-fourth, one-third, one-half. So I started being supportive of a process called reimportation, and that was because I couldn't figure out why Americans should pay more for pharmaceutical products than people in other parts of the world.

I found out, along with my colleagues, that in Spain, France, Germany, all over the world, the price is one-half,

1899 one-third, one-fourth, one-fifth or one-sixth of what it is
1900 in the United States.

The argument was, well, in the United States we have to do research and development, we have to do advertising and all that other sort of thing.

My problem is, why isn't the rest of the world paying for part of that? Why in the world should the American people have the burden of advertising, research and development and everything, and then pay five or six times what it costs for the same pharmaceutical product someplace else?

So we supported the reimportation program. The pharmaceutical companies went to the FDA and started talking about purity and whether or not there could be tampering and all that sort of thing, and they, in effect, have been able to block reimportation. They have been very effective, so they can protect their margins here and protect their market share. I don't understand that, and I don't think anybody in America who really thinks about it understands that.

We should not be paying more for pharmaceutical products than the rest of the world simply because, you know, we can afford the R&D, and we can afford that and load it on the back of the American people.

So we passed the prescription drug benefit, and we guaranteed in there that there would be no control whatsoever

by the Federal Government in the price of the pharmaceutical products that the government is going to be involved in. So they, once again, are able to block and say, It is going to cost a lot more here in America; and they have been successful in blocking pharmaceuticals from the rest of the world.

We can, with the new technologies, guarantee that drugs coming in are the product that we say they are. We can encapsulate them in plastic. We can put microchips or those mini, very small chips in there, to make sure that the product is the same as it is here in the United States, to guarantee the purity and everything. And yet we can't do that. And we can't do that because the pharmaceutical industry wants to keep the prices at a certain level here while they are able to give discounts way, way down the line, much lower costs, in other parts of the world.

I would like for somebody to explain to me why we can't have a process where the pharmaceutical companies can say, Okay, since you in the United States are going to make sure you are going to get comparable prices, we are going to go out and negotiate or tell the other countries in the world we are not going to allow you to charge this much less.

I sat down with the president of Eli Lilly, a company in my State. I sat down with people from Merck, vice presidents and presidents. And I said, why don't you come up to the

Hill and sit down with us, Members of Congress, and let's try to negotiate some type of solution to this problem so

Americans aren't burdened with a huge price while the rest of the world is getting off relatively scot-free. And they wouldn't do it.

Rather than doing that, they had PhRMA, their organization here in Washington that has tons of lobbyists, some of whom I am sure are here today--they had PhRMA go to the FDA and say, Oh, my gosh, these pharmaceutical products coming in from the rest of the world may not be pure; they may be tampered with, while at the same time they knew full well there were mechanisms we could use to protect those products coming into the country.

In addition, many of the products they are talking about are made in India and other parts of the world and coming in here in bulk anyhow--Viagra being one of them, which is used very widely here in the United States and, I understand in India, which really doesn't need it. It is only costing them about 10 or 12 cents a pill, whereas here, it is costing over 10 bucks.

Anyhow, I would like for you to give me an answer to that problem. Why do Americans pay three, four, five, six times what they are paying in Canada and elsewhere? Why can't we do something about negotiating? And why do we pass a Medicare prescription drug benefit that protects the

pharmaceutical companies from negotiation with our
government? I mean, it just seems to me there ought to be a
question of fairness here.

I want the pharmaceutical industry to make a lot of money. I want them to be very profitable. I am for the free enterprise system. But while I say that, I say, why should Americans bear the burden of all this, while the rest of the world is, in effect, getting off scot-free?

Thank you, Mr. Chairman, for giving me the time.

Chairman WAXMAN. The gentleman's time has expired.

We will give a short opportunity for an answer. I think you answered a question there.

Ms. MORTON. I have a short answer. So, one, I like the way you phrase the question, which is, Why doesn't everybody else pay more?

I mean, we have two choices: One, there is too much R&D, we should pay less, pay the same as France, and we have a new industry that responds to that. Or we think the amount of R&D we want is good right now, or it should be more, in which case everybody else is free riding. They are as rich as we are, and they are not contributing to the cost of R&D.

I think that is a very good question. Designing a regulation to get that to happen, I have some thoughts which I would be happy to share with you. But I think it is quite tricky.

Mr. ANDERSON. Fourteen percent R&D, 30 percent marketing.

Mr. SCHONDELMEYER. And they don't spend as much on marketing in other countries because their systems aren't as open.

Others today have commented, if you do this, if you do that, it will raise prices in the rest of the market. But I would bet most of those people who made that comment weren't talking about prices in the rest of the world.

I think we need to take actions and communicate to drug companies we expect them not only to look at raising prices in the rest of the U.S. market, but the rest of the world market; and they do need to look at other countries also to get back the money for R&D and to subsidize their development.

I would also point out that the drug that was involved in many cancer drugs was actually discovered by the National Institutes of Health. One of the leading cancer companies that has more products I think on the market than any other company, the last time I looked, 3 or 4 years ago, had about 21 cancer drug entities. And how many of those had that company discovered in their own R&D? Zero. The largest company that sells cancer drugs, at least 3 or 4 years ago, hadn't discovered a one; they had come from Federal Government funding.

2024 Chairman WAXMAN. Thank you, Mr. Burton. Your time has expired.

Mr. Tierney.

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Mr. TIERNEY. Thank you, Mr. Chairman.

I am always amused when Mr. Burton and I come down on the same side of an issue here. I was sort of hoping that he had made that passionate plea to his caucus a few years back, and maybe we wouldn't be here discussing what we are discussing today.

Look, I think the manufacturers have a hard time justifying the high prices. I think they have gotten a bit of a windfall out of it. But I know one of the arguments we are going to hear back is just what we are talking about right there, that if you do anything about this, research is going to stop and everybody is going to go to hell and die.

So I really want to knock that out of the box right now. It is nonsense and foolishness, as far as I am concerned.

They reported, what, about \$90 billion of profits last year, up \$20 billion previous to that, or whatever, and I don't for a moment think that a change in the price situation here is going to stop them from doing research.

So let me start with Dr. Anderson, if you would. Would reducing the high prices that they are now charging on the Part D program have an impact on the industry's research and development?

2049 Mr. ANDERSON. It is hard to answer that one analytically, but I don't think so. 2050 2051 Mr. TIERNEY. All right. 2052 Dr. Schondelmeyer, what do you think? Can we reduce 2053 Part D prices without adversely impact the research? 2054 Mr. SCHONDELMEYER. I think you can certainly go back to 2055 the Medicaid prices that you had and not affect research 2056 dramatically, because we were there and they were accepting 2057 those prices and they were living with that. So I think you 2058 can at least go back to that level, without a major effect on 2059 the market. 2060 Mr. TIERNEY. Dr. Morton, do you want to weigh in? 2061 Ms. MORTON. I would more or less agree with that, 2062 although I will say that a lot of these entities are 2063 discovered by venture-capital-funded small firms that are 2064 then bought by the larger firms, and anybody who is in 2065 venture capital or that kind of finance is investing because 2066 they expect a return. So anytime you alter the return, that 2067 goes into the calculation of whether they are going to spend 2068 money in the biopharma area. So I don't think you can ever assume no effect. 2069 2070 just, are we making a small shift of duals? Or are we making 2071 a big shift of everyone who's eligible for Medicare? 2072 Mr. TIERNEY. Thank you. 2073 Let me ask you--Dr. Schondelmeyer, you can start on

this--what is the difference or what is the variation between 2074 2075 how much research is done from government-funded projects 2076 versus what the industry does? And which drugs are involved, 2077 the more commonly used drugs or the less commonly used drugs, 2078 and all of that? 2079 Mr. SCHONDELMEYER. I haven't examined that 2080 systematically in recent years, but the evidence seems to 2081 suggest that drugs for categories that are most critical, 2082 such as cancer, tend to come more from government-funded 2083 research, and that drugs that come from the pharmaceutical 2084 companies tend to be more the lifestyle drugs, the drugs 2085 that--you know, feel good, live-well-type drugs, come from 2086 the drug companies that have broader populations. 2087 So the government tends to fund more critical, 2088 life-threatening drug discovery and drugs for smaller 2089 populations, while the drug companies tend to fund drugs for 2090 broader populations and maybe for more symptomatic or 2091 feel-good purposes. 2092 Mr. TIERNEY. We have all heard the expression of "me 2093 too" drugs out there and the research on that. Do you want 2094 to comment on that a little bit? Mr. SCHONDELMEYER. Well, I would be careful. 2095 There is 2096 an issue of "me too" drugs; I think it is often 2097 misunderstood, too, though.

I do think for a legitimate disease-state category,

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where there is three or four or five companies in the race to find a drug in that category, among those three, four or five, for whatever reason, whether it is regulatory or company performance, one of them is going to come out first.

I wouldn't say that the other four or five that were legitimately in the race are "me too" drugs because they were in the race. And, in fact, those other drugs could--if our market works, which it doesn't work well--could create competition.

Where "me too's" come in is when the company that first discovered it or other companies 15 years later come out with an extended release dosage form, a right-handed or left-handed molecule, those are "me too" drugs and those are kind of ways of extending patent pricing without adding a whole lot of value to the market in most cases.

Mr. ANDERSON. The NIH would suggest that more money is actually being spent by PhRMA than by NIH right now. We would have to take a look in terms of what it is spending it on.

NIH is much more basic research kinds of things. PhRMA is a lot more drug development kind of things. But I think overall, the numbers from NIH would suggest that PhRMA is spending a little more.

Ms. MORTON. I would second that.

I mean, NIH doesn't do the testing. So you can invent a

2124 molecule, but then you have to show that it is safe in 2125 thousands and thousands of people and go through the FDA. All of that is actually quite expensive, and NIH doesn't do 2126 2127 that. 2128 You can also see why the lifestyle drugs wouldn't be 2129 coming out of the government. I mean, I imagine the grant 2130 application to NIH for Viagra would not get funded. 2131 Mr. TIERNEY. You have more confidence than I do. would hope you are right on that. 2132 2133 Thank you, Mr. Chairman. I yield back. 2134 Chairman WAXMAN. Thank you, Mr. Tierney. 2135 Ms. Foxx? Ms. FOXX. Thank you, Mr. Chairman. 2136 I want to make one brief comment. As I have been 2137 2138 sitting here, listening to the comments that you all have 2139 been making -- and I've made this observation on a couple of 2140 other occasions -- I grew up in the mountains of North Carolina 2141 in the late 1940s, early 1950s, in the poorest county in 2142 North Carolina when I was growing up. 2143 My family was extraordinarily poor, yet we could afford 2144 health care. Everybody in our county could afford health 2145 care. In fact, I didn't know many people who had any kind of 2146 really big problems with health care. We had a hospital. had doctors. 2147 2148 And I have thought a lot about why it was that we could

2149 get health care in those days, and we have such a problem now with people, who are much better well off than we were, not getting health care.

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My observation is, it is two things: number one, government involvement, and I think any time you get the Federal Government involved in just about anything, you get more of a problem than you get a solution; and the other is third-party payer, when people are not in charge, I think you create problems.

I would just say that as a statement, because when I hear people say, get the government more involved, the Federal Government, it is just like scraping a fingernail across a blackboard for me, because I think what you are doing is simply creating more problems.

But I want to ask a question of Dr. Scott first, and then I have a general question.

Do you think that pharmacy benefit managers are sophisticated negotiators on behalf of seniors? heard about the problems with getting prices. Tell me what you think about that.

Ms. MORTON. Yes, I think they are sophisticated negotiators. A lot of the Part D plans that have been most successful in the sense of being taken up by many people are run by quite large and sophisticated insurance companies.

Ms. FOXX. Then the other question I have, my

understanding is that under Medicare, some drugs are paid for by federally set prices. They are injectable drugs under Part B. I would ask each member of the panel--and I know we have a limited time--do we set the prices for those drugs well? What is the history of the Federal Government setting those prices? My understanding is that there is a mixed history there; sometimes we have done well, sometimes we have done poorly. 

Relate that to what you are recommending now. Those are the folks on the upper end of the panel who are recommending that primarily.

Mr. SCHONDELMEYER. First, I would comment on, Are PBMs a sophisticated buyer? They are, but they don't have a fiduciary responsibility to act on behalf of the recipient. They act on behalf of their own stockholders and corporate entities, and those are different financial decisions that they make. So they are very sophisticated at taking care of themselves and meeting the requirements that are made of them for the recipients, but not acting in the best financial interest of the recipients.

I would also bet that hospital you had in your area was government subsidized under the Phil Burton program--

Ms. FOXX. No. Well, it may have gotten some, but it was primarily supported by the people who used it.

Would you mind answering the question I asked you to

2199	answer?
2200	Mr. SCHONDELMEYER. Yes. And what was that question?
2201	Remind me.
2202	Mr. ANDERSON. Let me answer. I will get it.
2203	Basically, if you take a look the Medicare program, the
2204	seniors in 1964, only about half of them had health insurance
2205	after Medicare. The other half got
2206	Ms. FOXX. You have just made my point.
2207	Mr. ANDREWS. I did? I thought you said that everybody
2208	had coverage.
2209	Ms. FOXX. I just said I think what created the problems
2210	with our not being able to get health care is third-party
2211	payer and the involvement of the government.
2212	Mr. ANDERSON. Well, I would disagree.
2213	Ms. FOXX. Do you mind answering the question I asked?
2214	Mr. ANDERSON. On the Part B thing, sure, essentially
2215	there was a problem with Part B drugs, that they were
2216	essentially giving serious discounts to doctors, but the
2217	Medicare program did not know those serious discounts, did
2218	not have price transparency, did not know that.
2219	Part of the Medicare Modernization Act of 2003,
2220	hopefully, with the average sales price, solved that problem,
2221	and now the discounts are less.
2222	So I think the Medicare program can learn and solve the
2223	problems.

2224 Ms. FOXX. What kind of learning curve is there for the 2225 people in the program? Mr. SCHONDELMEYER. Well, I would answer your first 2226 2227 question about the ASP and the government buying. First of all, Medicare Part B is a very different 2228 2229 It is primarily through physicians and a totally 2230 different distribution system, and there were incentives for 2231 doctors to actually prescribe more higher-priced drugs. 2232 I would argue, though, similar incentives are in place 2233 in the Medicare Part D program for the very reasons I stated. There is no fiduciary responsibility on behalf of PBMs, and 2234 2235 they can make more money by negotiating rebates from drug 2236 companies, but not passing it on in lower costs to the 2237 recipients. 2238 So I think the problems we had and the learning curve we have hasn't really stuck in Medicare Part D. 2239 2240 Chairman WAXMAN. Mrs. Foxx, your time has expired. 2241 Ms. FOXX. Thank you. 2242 I would like to say for Federal bureaucrats, there is no fiduciary responsibility either. 2243 2244 Chairman WAXMAN. The last word. 2245 I want to thank the three of you very much for your participation. I think that all the members on the committee 2246 2247 and all the people in the audience should get college credit 2248 for this discussion. It was a very high-level one, and I

2249 think a very worthwhile one. Certainly you have been helpful 2250 to us. Mr. DAVIS OF VIRGINIA. Let me just add to that and 2251 2252 thank our panel. It has been very informative. 2253 Chairman WAXMAN. Our next witness is Mr. Kerry Weems. He is Acting Administrator for the Center for Medicare and 2254 2255 Medicaid Services, Department of Health and Human Services. I would like to ask him to come forward. 2256 2257 Before you even sit down, it is the policy of this 2258 committee that all witnesses testify under oath. So if you 2259 would please raise your hand. 2260 [witness sworn.] The record will show that the witness 2261 Chairman WAXMAN. 2262 answered in the affirmative. 2263 We have your prepared statement and it will be part of 2264 the record in its entirety. What we would like to ask you to 2265 do is try to stay within 5 minutes for your oral 2266 presentation. 2267 I think you know the routine; it is green, 4 minutes; 2268 yellow for 1 minute, and when it is red, we would like you to 2269 certainly conclude.

Thank you for being here.

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2271 STATEMENT OF KERRY WEEMS, ACTING ADMINISTRATOR, CENTER FOR
2272 MEDICARE AND MEDICAID SERVICES, U.S. DEPARTMENT OF HEALTH AND
2273 HUMAN SERVICES

Mr. WEEMS. Thank you, Mr. Chairman, and thank you, distinguished members of the committee. It is a pleasure to appear before you today.

The success of the Medicare prescription drug benefit provides strong evidence that competition through private plans has contributed significantly to lowering costs to both the government and beneficiaries. Through Part D, Medicare beneficiaries are extremely satisfied with their current prescription drug coverage and have been given meaningful choices for drug coverage at a cost much lower than originally estimated.

Experience with Part D thus far demonstrates that competition is working for beneficiaries and taxpayers alike. According to the fiscal year 2009 President's budget, the necessary cost of the Medicare Part D program is 40 percent lower than the projections at the time the bill was passed, and beneficiaries are reaping these savings.

Independent surveys have consistently shown that more than 85 percent of Medicare beneficiaries and nearly nine out of ten dual eligibles are satisfied with their Part D

coverage. High satisfaction rates are directly related to the other successes in the Part D program, including meaningful and affordable choices, unprecedented information and transparency for beneficiaries, lower-than-projected costs from effective private sector negotiation, and increased generic utilization.

With the overwhelming success and popularity of Medicare's Part D benefit, we should be vigilant against attempts to use government mechanisms to intervene in the market and move to administered government pricing.

When Congress enacted Part D, the decision was made to move dual eligibles to Part D, which offered the dignity of choice and a market-based approach to the drug benefit structure and pricing. Congressional research agencies like CBO and GAO widely agree that direct government negotiation of prescription drug pricing in Part D is unlikely to lead to lower costs. As the chart demonstrates, simply comparing Medicaid's rebates to Medicare does not capture all the other efficiencies and savings achieved through Part D by encouraged use of generic, lower-cost drugs, lower-cost sharing opportunities for copayments and coinsurance.

23:15	RPTS KESTERSON
2316	DCMN MAGMER
2317	[12:06 p.m.]
2318	Mr. WEEMS. What is more, through drug utilization
2319	management, Part D has improved health outcomes by reducing
2320	the possibility of adverse drug events.
2321	The record from implementation of mandatory price
2322	controls and rebates in Medicaid reveals that these
2323	price-setting policies have the potential to increase costs
2324	in the private sector and others not subject to the
2325	government-imposed price controls.
2326	CBO examined the implementation of the Medicaid drug
2327	rebates on the market and found that, while access to rebates
2328	lowered Medicaid's outpatient drug expenditures, spending on
2329	prescription drugs by non-Medicaid patients may have
2330	increased as a result of the Medicaid rebate program.
2331	Further, GAO found that in the first 2 years of the Medicaid
2332	drug pricing program, the average price for medicines
2333	purchased by HMOs and Group Purchasing Organizations
2334	increased.
2335	With Medicare beneficiaries accounting for nearly 40
2336	percent of prescription drug spending in the United States,
2337	it is not at all unreasonable to expect that a change from
2338	market pricing in Part D to a government-mandated rebate

structure could have an even stronger ripple effect on the

cost of prescription drugs for those not subject to government-imposed price controls.

With a combination of more than 50 percent of the market subject to a statutorily dictated pricing structure, these two Federal programs could eliminate the potential rebates for any other purchasers. More specifically, it could lead to higher prices at the pharmacy, may compromise incentives to move enrollees toward low-cost therapeutic equivalents or generic drugs, or may undermine utilization management activities that the participating plans use for important safety protections as well as cost controls.

The Part D Program has been successful beyond expectations even in its infancy. Beneficiaries have meaningful choices for drug coverage at a cost that is much lower than estimated; and, more importantly, they are satisfied with their coverage.

Thank you for the opportunity to appear before you today. I look forward to your questions.

Chairman WAXMAN. Thank you very much, Mr. Weems.

[Prepared statement of Mr. Weems follows:]

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Chairman WAXMAN. Without objection--I think we've discussed this with the minority--we want to do an initial 10 minutes on each side, 10 controlled by the Chair and 10 controlled by Mr. Davis. And without objection, that will be ordered.

I want to start off my questions with you.

Mr. Weems, we are here today because we want to know whether we can make the Part D program work better for the taxpayers. You testified that the program is highly successful. You told us that beneficiaries are satisfied with the program. They have affordable choices, and they have good information with which to make choices and that they have greater, better access to generic medicines. If that is true, it is good news. And to be honest, after we have spent almost \$100 billion on this program, I would hope that that would be the case.

The issue for us is whether the taxpayers are getting the best value for their \$100 billion, and that is why the findings of the report released this morning are so troubling. The report finds that the prices paid by Part D insurers for the 100 drugs most used by dual eligibles are a lot higher than the prices Medicaid pays. On average, Medicare Part D is paying 30 percent more.

Mr. Weems, the central finding of the report is that Medicare Part D is paying significantly higher prices for

2386 drugs than Medicaid. Do you agree with this finding? 2387 Mr. WEEMS. Mr. Chairman, I had the opportunity to be 2388 briefed on your report; and I appreciate the opportunity for 2389 that. I have not had the opportunity to examine it in depth, but I would find that, for those particular drugs, that a 2390 2391 government-enforced price-setting system likely can produce 2392 lower prices, but that does not take into account the cost that may spread through the rest of the system. Yes, the 2393 2394 prices may be lower in a government-administered pricing 2395 system, but, as a result, they may be higher in the Federal 2396 employees benefits. So I would say that we would need to 2397 perform the rest of the analysis to see where those costs flow to. 2398 2399 Chairman WAXMAN. Well, we had the Medicaid system in 2400 place for 10 years with pharmaceutical rebates. Do you know that--if there is any evidence to show that there was a flow 2401 throughout the whole system of higher drug prices? 2402 Mr. WEEMS. We have evidence that suggests that, 2403 Yes. 2404 yes, costs were higher in the private sector as a result and 2405 also that there was a--2406 Chairman WAXMAN. Can you say that those higher prices were attributed to the Medicaid payment? Or are drugs 2407 2408 getting higher every year? 2409 Mr. WEEMS. Well, I believe there is research that

attributes to that, and it is also no accident that the

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amount of rebates that were available under the best price
began to go away under under the--in the private sector.

Chairman WAXMAN. We have looked at all the research on

Chairman WAXMAN. We have looked at all the research on this subject, and we can't find any studies that substantiate your position. So we would like you to submit that to us for the record.

[The information follows:]

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2419 Chairman WAXMAN. You're in charge of Part D; and what 2420 we see is that, according to this report, taxpayers paid more 2421 than \$3.7 billion over the first 2 years of the program as a 2422 result of the dual eligibles not being given the Medicaid price and now going to the Medicare price. Does that concern 2423 2424 you? 2425 Mr. WEEMS. Again, I think the analysis may be 2426 incomplete. It may be that the prices were--you know, there could be a lower price there, but it is also likely that 2427 2428 those prices would have shown up higher someplace else, 2429 probably in the non-dual part of the Part D program. 2430 Chairman WAXMAN. You have emphasized that Medicare Part D is costing less than projected--2431 2432 Mr. WEEMS. Yes. 2433 Chairman WAXMAN. -- and that is true. But the biggest 2434 reason the costs are less is that fewer seniors have enrolled than projected. It is obvious that if Part D is serving 2435 2436 fewer seniors, it's costs are going to be lower. 2437 On the central issue of drug prices, Part D is 2438 overpaying. Before January, 2006, the 6 million 2439 dual-eligible beneficiaries were getting their drugs through 2440 Medicaid. After January 1, 2006, they started getting their drugs through Medicare Part D. The only thing that changed 2441 is how much the taxpayers have to pay for these drugs. 2442

cost for just 100 popular drugs increased by \$3.7 billion.

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2444 That is indisputable.

Are you putting the interest of the big drug companies ahead of the interests of the taxpayers when your concern is not for the extra costs that we are actually paying for these very same beneficiaries?

Mr. WEEMS. Let me dispute one of your premises, if I might, that the only thing that changed was that the price changed. No, something else changed; and that is that the beneficiaries were moved from a State-run, price-fixing program--in some cases, of States with restricted quantities--into a risk-based insurance product, where they have in many cases, even for the low income, the dignity of choice, which they didn't have in Medicaid, broader access to more drugs and no limits on the--

Chairman WAXMAN. That depends on what plan they joined. Because the plans could restrict the drugs' formulary.

But the Medicaid rebate program, which I helped design--I was around when we adopted it. It is all voluntary. The drug company didn't have to participate. And the drug companies participated on the basis that we would demand the best price for them that they were charging others in exchange for adding all their drugs on the formulary. So the companies benefited by making sure that all their drugs could be available to Medicaid patients.

This wasn't a price fixing--this wasn't a fixed price or

price fixing. It was a negotiation by the government for a lower price for that population. Now we have no negotiation; and, as a result, I believe, we are seeing higher prices. We are definitely paying higher prices. Would you say it is not because we don't negotiate it any longer? It is not because we don't have the Medicaid reimbursement formulary that--for that same population for those same drugs?

Mr. WEEMS. Again, I would say there is only half the analysis; and that is the analysis that, you know, the States pay. You can look at the--you know, the price that is mandated by the rebate. The analysis that needs to be complete is what happens on the other side of the equation, the market equation, when--press down prices here, they are going to go up someplace else. The Federal employees benefit program, private insurer, we've seen it happen.

Chairman WAXMAN. We'd have to see if that is the case. I'm looking forward to see what documentation you have for that.

If we had lower prices in the United States, it would probably lead to higher prices in the other countries.

Should we worry about that?

It just seems to me that for the dual eligibles that we actually provided drugs to under the Medicaid program at a lower cost and the same drugs at a lower cost we are now paying for that same population at a much higher cost and for

2494 that group we are paying a lot more money. I don't think--I 2495 don't see what we're getting for that extra money.

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Mr. WEEMS. If we were to--let's take one of the suggestions that one of the academics made here. And that is if we were to take that dual-eligible population and apply the rebate, the Medicaid rebate, to that population, the most likely initial result would be an increase in Part D for everybody else who is not dually insured. Is that, you know, the consequence that we would like to have? Is, you know, a secular increase in Part D that then spread beyond Part D and other parts of private market?

Chairman WAXMAN. I don't believe that would be an accurate statement of what would happen. I think the drug companies are trying to maximize the amount they can get for their drugs; and if you provide more money for their drugs, they are going to be happy to take it. So I don't see evidence for that statement.

I'm going to reserve the balance of my time, which is 1 minute and 37 seconds and yield to--now 10 minutes to Mr. Davis.

Mr. DAVIS OF VIRGINIA. We just have a fundamental disagreement between us over if you reduce costs in one area, does it raise costs in other areas. Somehow I think the chairman and advocates on that side think that this just comes out of the drug companies' hides and that is the end of

2519 it and it has no effect on research and development or 2520 anything else. And I don't think that is borne out. 2521 In fact, I would ask unanimous consent, Mr. Chairman, 2522 that we put in--you had asked a question earlier about 2523 overpaying, and there is no effective -- on the 2524 overpayment -- this is a CBO paper, How the Medicaid Rebate on 2525 Prescription Drugs Affects Pricing in the Pharmaceutical 2526 Industry. This is a Congressional Budget Office report, and 2527 I would unanimous consent that --2528 Chairman WAXMAN. Without objection, we'll put that in 2529 the record. Mr. DAVIS OF VIRGINIA. Thank you very much. 2530 2531 [The information follows:] 2532 \*\*\*\*\* COMMITTEE INSERT \*\*\*\*\*\*

2533 Mr. DAVIS OF VIRGINIA. The majority staff report found 2534 that Part D rebates are smaller than Medicaid rebates. 2535 You're not surprised by that finding, are you? 2536 Mr. WEEMS. Not at all. 2537 Mr. DAVIS OF VIRGINIA. Is this new information? 2538 Mr. WEEMS. No. 2539 Mr. DAVIS OF VIRGINIA. In Congress, we often lobby to 2540 change reimbursement for different services covered by 2541 Medicare or to expand those services all from political 2542 perspectives. The drug company or somebody could be--or a 2543 manufacturer could be from your district and there is 2544 pressure to slip this in here or slip this in there or expand 2545 services to one needy group over another. 2546 At CMS, we are tasked with creating a national formulary 2547 or setting prices. Do you think the process would be open to 2548 meddling by Congress by disease advocates and drug 2549 manufacturers? 2550 Mr. WEEMS. Absolutely. And, you know, we can see the 2551 evidence of this. You know, if you look at the mail that CMS 2552 receives, we receive virtually no mail--I don't think I'm in 2553 a position to say zero--but virtually no mail about the price 2554 of specific drugs under Part D. We receive huge volumes of 2555 mail about those drugs for which we do administer pricing under Part D. A lot of mail, a lot of pressure and, in some 2556 2557 cases, there is even legislated prices--

2558 Mr. DAVIS OF VIRGINIA. When you say mail, are you 2559 talking about mail from Members of Congress? 2560 Mr. WEEMS. Members of Congress, manufacturers, lobbying organizations, you name it. We receive virtually none of 2561 2562 that under Part D. One of the great success stories of Part 2563 D is it has depoliticized the price of individual drugs. 2564 Mr. DAVIS OF VIRGINIA. What would be--is that one of 2565 the reasons, you think, that the costs that were projected 2566 originally are far and above what has actually taken place? That and the effects of competition. 2567 Mr. WEEMS. 2568 Mr. DAVIS OF VIRGINIA. I mean, there is a fundamental 2569 difference, that some of us believe competition brings down 2570 costs, some of us think that the government is smart enough 2571 to be able to just negotiate the best cost because of our buying power. In fact, there are some formularies that have 2572 2573 greater potential buying power than the Federal Government. The PBMs, the prescription benefit managers, 2574 Mr. WEEMS. the ones that the Part D program use, represent about 240 2575 2576 lives across the Nation. So that is real buying power. 2577 Mr. DAVIS OF VIRGINIA. If CMS--we talk about we are tasked with creating a national formulary, setting prices. 2578 2579 What impact could that have on seniors in Part D? 2580 Mr. WEEMS. If it is a highly restrictive formulary, it 2581 might mean that they don't get the drugs that they need. 2582 Mr. DAVIS OF VIRGINIA. Mr. Weems, you have been a

2583 career employee, haven't you? 2584 Mr. WEEMS. I am a career employee, sir. 2585 Mr. DAVIS. So you are a career employee on there. 2586 weren't some administration lackey or anything else that they 2587 were able to take because you had given contributions to a 2588 campaign or been active in political causes, right? You're a 2589 career employee, and you have worked at this all your life? 2590 Mr. WEEMS. I started my career in 1983 as a junior 2591 budget analyst with the Social Security Administration. 2592 Mr. DAVIS OF VIRGINIA. How does the financial outlook 2593 for Medicare Part D compare to the Part A program which 2594 covers hospital care? 2595 Mr. WEEMS. They are financed entirely differently. 2596 Part A is financed by FICA taxes. Part D is financed by premiums and by general fund transfers. So the financing 2597 2598 schemes are different. 2599 Part A, because of its financing schemes and because of 2600 the rising costs in Part A, is going to go broke in 11 years, 2601 according to the trustee's report. Mr. DAVIS OF VIRGINIA. And you concur with that from 2602 2603 your observations? 2604 Mr. WEEMS. 2605 Mr. DAVIS OF VIRGINIA. And Part D? 2606 Mr. WEEMS. Part D is financed, as I said, from -- it is financed entirely differently, and so it is not subject to 2607

the same sort of constraint that the Part A is. 2608 Mr. DAVIS OF VIRGINIA. But, in fact, the projections on 2609 Part D, are they greater or less than were projected in terms 2610 2611 of the costs to the government? 2612 Mr. WEEMS. In fact, you can see the original cost 2613 estimate is the upper line. Mr. DAVIS OF VIRGINIA. 2614 That is the third chart over? 2615 Mr. WEEMS. That is the third chart over. The lower 2616 line is the most recent cost from the President's budget, 2617 most recent cost estimates. 2618 Mr. DAVIS OF VIRGINIA. So Part A has basically been overruns and Part D has been underruns in terms of--2619 2620 Mr. WEEMS. Again, Part A -- in fact, this year in Part A, 2621 the expenditures of -- in Part A will exceed what we take in in 2622 taxes for Part A. 2623 Mr. DAVIS OF VIRGINIA. Now, in the previous panel we 2624 heard--I think it was Dr. Anderson testified that all Federal 2625 prices for prescription drugs should be uniform. Outside of 2626 prescription drugs, does Medicare, Medicaid, the VA and FEHBP 2627 pay uniform prices for health care services? 2628 Mr. WEEMS. No, they don't. Not as a matter of policy. There might be times when they--2629 2630 Mr. DAVIS OF VIRGINIA. Coincidentally. 2631 Mr. WEEMS. Yeah, by coincidence. 2632 Mr. DAVIS OF VIRGINIA. How do you think an effort to

2633 make prices uniform across these programs to the lowest 2634 denominator would be received by physicians or hospitals? Mr. WEEMS. Well, you know, Mr. Davis, it is an 2635 2636 interesting question. And the question--the answer to that question depends on your philosophy. 2637 2638 If you were to do it through competitive means, you would allocate resources correctly. If you were to turn it 2639 2640 over to CMS with my very well-meaning Federal employees who 2641 fix prices every day for A and B, we likely would not get it 2642 right. 2643 Mr. DAVIS OF VIRGINIA. There is sufficient evidence 2644 that Medicaid price controls increase prescription drug 2645 prices to private payers, which in the United States are 2646 generally employers. These are like GM and Ford who are competing in a global marketplace. Although we may get a 2647 2648 reduction for Medicaid recipients, in effect, I think there is evidence that drives up the costs to these companies that 2649 has an effect downstream in terms of their ability to 2650 2651 compete. 2652 GM and Ford have both cited higher health care costs as 2653 one of the factors affecting their decline in global competitiveness. What do you think would be the impact of 2654 2655 requiring Medicaid prices in Part D on Ford or GM? 2656 Mr. WEEMS. For the entirety of Part D? 2657 Mr. DAVIS OF VIRGINIA. And union pension plans I quess

you could throw into that as well.

Mr. WEEMS. Sure, sure. So Part D, together with Medicaid, represents over half of the pharmaceutical market in the United States. Applying government cost controls to more than half the market and pushing down that half of the market to some specified pricing scheme would definitely--and I say this without reservation--cause cost increases in the rest of the market, which specifically would be the private sector. And, you know, for companies like Ford and GM, it would substantially increase the pharmaceutical costs in every vehicle.

Mr. DAVIS OF VIRGINIA. You don't think the pharmaceutical companies would just say, we're going to continue the same amount on research and development anyway. We're just going to take this out of our bottom line, reduce advertising costs and the like?

Mr. WEEMS. I think that is unlikely, but the next panel will have somebody from pharmaceutical companies on it, and I would invite you to ask them.

Mr. DAVIS OF VIRGINIA. Okay. I happen to agree with you.

Much has been made about the Medicaid coverage of prescription drugs, but prices are only one factor in determining the success of any new benefit. How do you think seniors' access to drugs in Part D compares with Medicaid

2683 recipients' access to drugs?

Mr. WEEMS. They have more access and more choices. The main feature of Part D is the ability to choose a plan that works best for the individual.

Mr. DAVIS OF VIRGINIA. You may have a rare disease or something that is not covered, for example, by Medicaid--

Mr. WEEMS. Correct.

Mr. DAVIS OF VIRGINIA. --that is covered by Part D, and you can choose that particular--

Mr. WEEMS. A lot of it just has to do with choice. You know, what is the level of premium that I want to pay each month? What is the amount of co-pay that I want to be exposed to? Do I want to use my neighborhood pharmacy?

Those are the kinds of things that seniors find extremely agreeable about this program, that it is not a government one-size-fits-all, the government picks winners and losers. It is that there is choice and a lot of choice, and their drugs are available to them in a very convenient way that--where they can get what they want.

When I talk to seniors around this Nation--and I spend a lot of time talking to them--we hear great satisfaction with Part D. And what they say over and over again is don't take this benefit away from us. Make sure you keep this benefit. This benefit is working for us.

Mr. DAVIS OF VIRGINIA. I think that is why you don't

2708 l hear the majority saying let us move these dual eligibles 2709 back to Medicaid. Because it would be politically very, very 2710 unpopular with these groups. And now they'd like to have a 2711 hybrid, it seems to me, of--well, we are going to have 2712 Medicaid pricing in Part D for some items and the like. 2713 Mr. WEEMS. In fact, satisfaction rates for the duals 2714 are higher than those even of the regular population. 2715 one of the first times, they have been given the dignity of 2716 choice from a government program. 2717 Mr. DAVIS OF VIRGINIA. As opposed to a one-size-fits-all, take-it-or-leave-it? 2718 2719 Mr. WEEMS. That's correct. 2720 Mr. DAVIS OF VIRGINIA. The purpose of the Medicaid 2721 price regulations was to control the cost to States and the 2722 Federal Government. That is why they put the price controls 2723 in. Since implementing price controls 18 years ago, do you 2724 have any observations on the cost of prescription drugs in Medicaid? Have they remained flat? Have they gone up? Have 2725 2726 they gone down? 2727 Well, you know, the best price provisions, Mr. WEEMS. 2728 the provisions with respect to rebates, are fixed from a 2729 price. So drug prices continue to go up. You know, they 2730 have been effective in reducing the liability for drugs in 2731 the Medicaid program while increasing the liabilities in 2732 other places and causing market distortions in other places

2733 on the market.

2734 Mr. DAVIS OF VIRGINIA. Okay. My time is up. Thank

2735 you.

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2736 Mr. YARMUTH. [presiding.] We have a series of votes, as

2737 you might have noticed. So we'll at this point recess the

2738 hearing and reconvene at 1:00.

2739 [Recess.]

2740 Chairman WAXMAN. [presiding.] The meeting of the 2741 committee will come to order.

The Chair recognizes Mr. Murphy to pursue questions.

Mr. MURPHY. Thank you very much, Mr. Chairman.

I wanted to make a brief comment off of the chairman's concern, Mr. Weems, over the terminology you used regarding the Medicaid rebate program and that is peppered in your testimony, both written and oral, is the idea that this is price control, that this is price fixing. When it seems to us that it is merely using the market leverage and market power of the Federal Government to do exactly what private industry does, what the HMOs do in negotiating these prices, which is to say, through a choice of a particular pharmaceutical company, that this is the price that we're willing to pay. And if you don't pay it, then you're not going to be part of our plan, which is essentially what the Medicaid rebate program does.

Price control strikes me as something very different.

2758 mean, that is a statutorily imposed price that everyone has 2759 to accept for their product.

This is a voluntary program. I would hope that we'd be a little careful in mixing what is a voluntary rebate program that the pharmaceutical companies pay as a means of selling their drug in a particular plan, the Medicaid plans versus what is traditionally thought of as price controls.

But my question is a little bit different, and that is--your testimony, Mr. Weems, as to the disruption in the delivery of health care that would result from imposing Medicaid rebates on the dually eligible population. And I want to just ask you to elaborate a little bit on that as to what evidence you have that gaining these discounts for taxpayers would lead to this potentially troublesome disruption of the health care delivery system.

Mr. WEEMS. Thank you for the question.

And, you know, I don't mean to get into a semantic battle. But, in my view, a system which fixes a specific rebate amount and fixes it through statute is very different than a negotiation. And the 15.1 percent rebate in Medicaid is fixed and fixed in statute. So I would stand by my terms, sir.

You know, as for the disruptions--I mean, we can--we can see this. You know, it was the GAO report that found that, in the 2 years following the implementation of the Medicaid

2783 best price rebate program, the best price discount for 2784 outpatient drugs purchased by HMOs and PPOs decreased to about 14 or 15 percent, which is approximately the minimum 2785 2786 required by the statute. 2787 CBO found that the best price rebate program, found that 2788 drug purchasers in the private sector, their discounts 2789 weren't as good. Between 1991 and 1994, the best price 2790 discounts that pharmaceutical manufacturers gave off of 2791 wholesale prices fell from 36 percent to 19 percent. 2792 Mr. MURPHY. For private insurers? 2793 Mr. WEEMS. That's correct. 2794 Mr. MURPHY. So you're suggesting that there is a 2795 movement--there is also testimony that you give about we would have a discouraging of employers from continuing to 2796 2797 provide prescription drug coverage at the same level they do 2798 today. Is that--2799 Mr. WEEMS. If it is more costly, we can expect less of 2800 it, yes. 2801 Mr. MURPHY. I guess it strikes me as strange that the testimony here is that we are essentially going to be--that 2802 2803 today we are, in essence, subsidizing privately held plans 2804 purchased through employers? Mr. WEEMS. No, not at all. 2805 2806 Mr. MURPHY. Wouldn't that be the converse of suggesting 2807 that -- if your suggestion is that by the taxpayers paying less

2808 that you're essentially pushing the bubble in somewhere and 2809 it comes out somewhere else, that private employers are going to pay more, wouldn't the suggestion be currently today then 2810 2811 we are subsidizing private employers' purchase of--2812 Mr. WEEMS. Not at all. You need to compare the two If, in fact--if you had a competitive pricing 2813 2814 system on both sides, then you can make a direct comparison. But, in fact, on the Medicaid side, there are mandatory 2815 2816 rebates. The simple hydraulics of supply and demand means that, as you force down those prices, they are going to go up 2817 2818 someplace else. That, in fact, means that the private sector 2819 currently is subsidizing Medicaid. 2820 Mr. MURPHY. And currently, though, currently though, 2821 how does that not lead to an argument that we are currently, 2822 through our inflated prices that we are paying -- and you admit 2823 that the prices we are paying today are not commensurate with 2824 what Medicaid is paying--isn't providing a subsidy on the other side to the private insurers? 2825 2826 Mr. WEEMS. No. The market--the market prices--you're 2827 asking to compare a risk-based market price to a 2828 government-imposed price. They don't compare. Because you

have got the cross subsidy and you're not able to capture the

imposes on the rest of the nongovernment cost-controlled part

cost of forcing down the lower price and the cost that that

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of that sector.

Mr. MURPHY. And I know my time has expired, Mr. Chairman. But to get back to, I think, a fundamental disagreement, I think that the government rebate program is not completely risk independent. I mean, we obviously are setting a price at which we believe that the drug provider will continue to provide the pharmaceutical product. We are incorporating risk because we know if we set the rebate price too high that that pharmaceutical company will no longer sell the product. So it may be different than the negotiation in the back and forth that occurs in the private sector, but it is completely interdependent upon risk. Wouldn't you agree that that is part of the--

Mr. WEEMS. No. I think we're talking about risk in two different ways. When I refer to a risk-based insurance product, that is what we have in Part D where the--the profit, the equity of the firm is in fact at risk for achieving a good bid, for lowering drug prices and for bringing in recipients into their plan. That's the risk. That's a much different kind of risk than the kind you're describing.

Mr. MURPHY. You're right. I am mixing terms.

I guess what I'm suggesting is that the fundamentals of supply and demand that underlie a negotiation between an HMO and a pharmaceutical company are not absent from the determination of what the rebate will be under the Medicaid

2858 program. Because if the rebate again is set too high, then 2859 that drug will not be provided as part of the Medicaid 2860 program. So many of the same economic factors that underlie 2861 those negotiations are present in the determination of the --2862 Chairman WAXMAN. Your time has expired. 2863 If you want to make a comment. Otherwise, we can move 2864 on. 2865 Mr. WEEMS. We can move on, sir. 2866 Chairman WAXMAN. Okay. Mr. Issa. 2867 Mr. ISSA. Thank you. 2868 Are you aware of the history of this best price practice 2869 that Medicaid has where, a year after its implementation, the 2870 Department of Veteran Affairs asked Congress to exempt it 2871 from the calculation of Medicaid's best price because in fact 2872 it was raising their prices? Isn't that true? 2873 Mr. WEEMS. To the best of my knowledge, yes. 2874 Mr. ISSA. So here we have the gold standard to a 2875 certain extent. The Veterans Administration buys selected 2876 drugs at the best possible price, makes decisions, including 2877 formulary decisions, based on the best value for our veterans 2878 and then makes it available to other--certain limited other 2879 government agencies such as Bureau of Indian Affairs and so on for Indian health, and they choose to always take 2880 2881 advantage of it because the prices are good. And they are 2882 saying, when you mandate a discount, you distort the market

2883 | and you distort the likely retail price. Now, isn't that really what we're really talking about? 2884 2885 Mr. WEEMS. Sure. And we've seen that, since the best price mandate, that best prices have gone up, unsurprisingly, 2886 2887 I would say. 2888 Mr. ISSA. So a question I asked the economist earlier 2889 today -- and I will challenge you on this side some -- isn't our 2890 gold standard--the Veterans Administration, it backs up a truck, takes a whole truckload, reduces reliability, 2891 2892 administration, takes the drugs and makes a good price. 2893 Isn't that the gold standard for pretty much as good as you 2894 would do, assuming you don't simply distort the market and 2895 demand a lower price, regardless of merit? 2896 Mr. WEEMS. Well, I might disagree with that 2897 characterization, because I think it -- first of all -- and we 2898 are probably trying to get to the same place here. But, first of all, the Veterans Administration is a government 2899 agency that actually takes custody --2900 2901 Mr. ISSA. And maybe I can clarify. What I'm saying is, when you do all of those things, you get the price maybe 2902 2903 lower than any other plan. 2904 Mr. WEEMS. Quite possibly. 2905 Mr. ISSA. But when we're looking for the lowest 2906 possible price, we should not look to Medicaid with a mandated price, we should look to a bulk buyer buying by 2907

reducing administration and risk to these companies. When they make a buy, they make a big buy; and you just ship it.

Mr. WEEMS. That's right.

Mr. ISSA. Okay. So, earlier today, I said, when we want to evaluate Medicare Part D's performance, shouldn't it be taking, if you will, if possible arithmetically, take the VA, put back in the administrative cost of not buying from a single payer but rather allowing people to get drugs where they want to be, where their doctors and their pharmacies are, rather than going to a VA facility to pick them up. Recognizing there is distribution costs, administrative costs, but that convenience is something our seniors demand because they want that capability. They're not asking us to please have 35 locations around the country they can drive to to get their drugs.

If you add back in those reasonable costs and so on, isn't that the standard where we would like to see Medicare Part D close to? And in your estimation are we, when you add back in those costs, somewhat close?

Because here today it seems like everybody wants to use Medicaid, which is an artificial mandated price, as the gold standard, rather than any other comparison. Or they want to use Canada, where they say if you don't give us a lower price, we'll simply void your patent and knock it off. So that is my real question. Can you progress on how you see it

2933 | should--

Mr. WEEMS. Sure. That makes the comparison more fair. But the thing that—that Part D offers that—you know, is that you need to layer in again here is the choice of plans, you know, the many, many choices that are available to seniors and the way that they can, you know, structure their payments. They can choose, you know, a higher premium level in return for lower structured co-payments, those kinds of things. All of that adds to the value of Part D. And, you know, once you step up from a highly restricted—all the way up to a program that offers considerable choice—

Mr. ISSA. Right. And, look, I have no question at all that my seniors want the features of being able to choose between formularies, to have some choices, to decide sort of good, better and best.

One of the controversial things by some here on the dais is, well, why don't we just have one formulary? Why don't we just have one solution? In a sense, the price that Medicare Part D gets, which is better than originally forecasted, isn't one of the most important parts of that. The fact that independent companies compete based on their formulary and features and by the way offered to pharmaceuticals, do you want to be with us, and will you give you a better price for it, because they are not necessarily taking every therapeutic solution.

2958 Mr. WEEMS. That's absolutely true.

2959 Mr. ISSA. If we come up with one mandated solution,
2960 although we might get a lower price on that, don't we distort
2961 the market for what the seniors want?

Mr. WEEMS. Yes. And I would say that there are two aspects to that. First of all, that a restricted formulary may mean that some people don't get the drugs they need; and, secondly, it puts the government in the position of choosing winners and losers in the marketplace.

Chairman WAXMAN. The gentleman's time has expired.

Ms. Foxx, do you have questions?

Ms. FOXX. Thank you, Mr. Chairman.

I think that last point was really important, that we should not be putting the government in charge of picking winners and losers, especially when it comes to health care.

I have a couple of questions that I'd like to ask you, Mr. Weems; and I would say that I'm not always happy with the way CMS operates. There are things that I disagree with that you all have done, and so there are lots of things that I think could be done better over there, And we'll have another conversation about that sometime after this.

But let me ask you a question. According to the material that you all have produced, Medicare Part D enrollees continue to save about--excuse me. I'm asking the wrong question. You show that Part D costs are lower than

2983 the initial estimates. Can you tell us what accounts for 2984 that?

Mr. WEEMS. Sure. There are a number of things. First of all, that the degree of competition that occurred in the system was more robust than originally estimated; secondly, the price of drugs has not risen as fast as originally estimated; then, lastly, the total population enrolled is somewhat lower than originally estimated.

Ms. FOXX. The second question has three parts to it.

You have been around the Department for a long time, and you probably will remember during the debate about Part D there were a lot of doomsday predictions. I was not here during that debate. I didn't vote on Medicare Part D. But tell me in your opinion which—how these doomsday predictions have worked out.

Number one, did plans refuse to offer drug-only insurance? I'll ask all three of the questions, and then you can respond. Did plans cherry-pick only the healthiest seniors? And you've already mentioned this about drug prices not rising exponentially. If we have time, I would like you to also say something about the price of drugs holding down the cost of health care in other areas.

Mr. WEEMS. You know, clearly, there was a lot of concern at the beginning that there wouldn't be marketplace entry. There has been robust and substantial marketplace

entry. In fact, the complaints are reversed, from nobody is going to get into this to aren't there too many.

As for cherry-picking, that is something that we still remain very, very vigilant about in CMS. Every year when the bids come in, we examine the bids, we examine the formularies to make sure that there are not discriminatory bids as part of that.

You know, as for pricing, you know, if you--73 percent of our enrollees are in plans where the price index did not increase by more than 3 percent; 50 percent are in plans where the price index did not increase more than 2; and 14 percent are in plans where the price actually fell. So we not only see good price stability, we also see that our seniors are able to protect themselves against the risk of higher prices in the plans and also during the plan year by choosing tiered co-payments. Ninety-five percent of our beneficiaries buffer themselves against the risk of payment increases by having set co-payments, rather than percentage co-payments.

Ms. FOXX. Thank you.

Mr. Chairman, I would just like to make a brief comment.

I find it so interesting that, in matters of choice, the majority party here wants choice when it comes to destroying life but not choice for citizens when they have the opportunity to save money and have better health care.

Because it seems to me that one of the things that drives the majority party so crazy about Medicare Part D is that people do have choice. We don't want people to have choice about where to go to school, but, again, we do want them to have choice to kill babies.

The other thing that I think is not recognized that Mr. Shays said earlier is Medicare is in deep trouble; and there is material out all over the place today that the majority party is going to avoid dealing with the trigger, going to sweep that under the rug. We don't want to deal with the big issue of Medicare, but because there is this animus towards the drug companies, it is easy to pick on drug companies and pick on the private sector whenever we possibly can and make them look bad.

So I think we need to be dealing with the real problems that we have, which is the major Medicare program and what has come to be called an entitlement, because that is where our real problems are.

Chairman WAXMAN. The gentlelady's time has expired.

Mr. Weems, I want to ask you some questions. Under the Medicare Part D, people can choose a plan that will offer them some drugs. It doesn't have to be every choice of drugs, but they have their formulary or they can join another plan that will have its formulary. Isn't that the way it works?

3058 Mr. WEEMS. That's correct, yes, sir.

Chairman WAXMAN. So they have a choice, but they may
find one drug on one plan but not on that same plan for
another drug so they have to--they really can't pick and
choose. They can't belong to two plans. They can only
belong to one. So they don't really get the choices of all
the drugs they need.

Under the old Medicaid, they had all the drugs on the list. So I just say that rhetorically when we talk about how much choice we are actually giving people.

Secondly, I want to point out you said with pride that a lot of insurance companies are out there competing and that just shows us it is wonderful and really working. But it also might show that they are making a lot of money; and if they're making a lot of money, why not go into that business? I just say that rhetorically as well.

Then the other thing I want to ask you is, we had 6 million people on Medicaid, and we paid less for them. Now they are on Part D Medicare, and we pay more for them. It is your premise that, if we paid less, the prices would go up in other areas where government spends on drugs; is that right?

Mr. WEEMS. That's correct. Or in the private sector.

I wouldn't just limit it to government, sir.

Chairman WAXMAN. Okay. Now that we've taken 6 million people and we have paid less for them, are we seeing a drop

3083 in what is being paid in other government programs or in the 3084 private sector? 3085 Mr. WEEMS. Again, I think that is a question that bears 3086 examination. The question may be--3087 Chairman WAXMAN. It goes to your argument. 3088 Mr. WEEMS. It bears examination, sir. 3089 Chairman WAXMAN. Have you seen any evidence of the 3090 prices dropping for other government programs? 3091 Mr. WEEMS. One of the reasons that we did not see the 3092 top line on that is prices have not increased in the way or 3093 at the speed that was originally estimated. So I would point 3094 to that as evidence, sir. 3095 Chairman WAXMAN. What prices haven't increased at the 3096 speed of --3097 Mr. WEEMS. Drug prices. 3098 Chairman WAXMAN. Who estimated them? 3099 Mr. WEEMS. The original estimate from the Office of the 3100 Actuary for the--3101 Chairman WAXMAN. Is that the one we were never allowed to see? We still haven't gotten that one, as I understand. 3102 3103 That was--the actuary's life--no, not his life, his job was 3104 threatened if he shared with Congress the cost. 3105 Well, let me go into another question. Let us say we 3106 spent \$3.7 billion for 6 million beneficiaries when they're 3107 under Medicaid--\$3.7 million less, now we're paying \$3.7

3108 million more. Is that the best use of our \$3.7 million? 3109 drug companies like it, but couldn't we use that for other 3110 purposes when we have so many uninsured? 3111 For example, one of the reasons the President said he 3112 vetoed the SCHIP bill was because it cost so much money. 3113 Well, that \$3.7 billion would have covered 3.3 million 3114 uninsured children. Which is a better use of that money, paying it to the drug companies or paying less to the drug 3115 3116 companies and using it for children? 3117 Mr. WEEMS. Again, sir, I think that analysis 3118 ignores -- is only half the equation. It ignores the 3119 distortions that the price setting creates in other parts of 3120 the market. You may--3121 Chairman WAXMAN. We can't be responsible for every 3122 distortion -- you have never given us any evidence of that. 3123 But even if you do, there are always distortions. 3124 I want to ask you one question about this issue of 3125 distortion. Do you think if we charge less--let me put it 3126 this way--if we charge more for drugs that the drug companies 3127 say, well, since I'm making so much money under this Medicare Part D, I'm going to give a break to these other payers of 3128 3129 the private sector? 3130 I can't believe that is the case. They are in business to make money. If they can sell their drugs at a certain 3131 3132 price to the private sector, they'll do it. If they can sell

3133 their drugs to the government at a higher price, they'll do 3134 it. It is when somebody says, no, we're not going to pay the 3135 higher price that they have to realize that they're not going 3136 to make the money they were making before and then make their business calculations. 3137 3138 Mr. WEEMS. And I think you perfectly encapsulated the 3139 problem with government-administered pricing. We know that 3140 in Part A and B we overpay in some areas and underpay in 3141 others, and it creates distortions and costs that, frankly, 3142 we're not able to measure. Chairman WAXMAN. 3143 In Part D? 3144 Mr. WEEMS. A and B. In Part A and B. 3145 government-administered prices program. We know that we 3146 overpay. 3147 Chairman WAXMAN. Would you be surprised if you found 3148 that one plan was paying more for the same drug than another 3149 plan under Part D? 3150 Mr. WEEMS. For the same drug, no. 3151 Chairman WAXMAN. You wouldn't be surprised? 3152 Mr. WEEMS. No. Chairman WAXMAN. Would you be surprised if one plan was 3153 bargaining for lower prices and didn't pass it onto the 3154 3155 consumer but increased their profits? 3156 If it is a rebate, they have to pass it on Mr. WEEMS. 3157 in their premiums, sir.

3158 Chairman WAXMAN. Well, it may not be a rebate. 3159 just negotiated a better price because they did some deals. 3160 That's what we want in the market, right? That's correct. 3161 Mr. WEEMS. 3162 Chairman WAXMAN. Pass on the lower prices to the 3163 Medicare system or beneficiary or does it just simply make 3164 all those companies that to our surprise decided to go into 3165 the business richer? Mr. WEEMS. If they are going to compete for 3166 3167 beneficiaries, they're going to have lower premiums, and that 3168 drives down their profits. 3169 Chairman WAXMAN. Do you think that's the only reason 3170 signs up on one plan as opposed to another, the price? Mr. WEEMS. The price and the coverage of the drugs. 3171 3172 Chairman WAXMAN. Yeah. Okay. Thanks. 3173 The gentleman from North Carolina, Mr. McHenry is 3174 recognized. 3175 Mr. MCHENRY. I appreciate it, and I hope we will still 3176 have the same liberal time policies for me as for you. 3177 know being chairman has its privileges. 3178 Chairman WAXMAN. I went over 30 seconds. If you want 3179 an extra 30 seconds, I'll give you--3180 Mr. MCHENRY. That would be great, but I think you probably just burned it. 3181 3182 So anyway--

3183	Chairman WAXMAN. I can't make you happy any way, huh?
3184	Mr. MCHENRY. Well, actually, you know, your philosophy
3185	is very different and your focus is different here
3186	becausebased on the studies
3187	Chairman WAXMAN. The gentleman's time is just beginning
3188	at 5 minutes.
3189	Mr. MCHENRY. Okay.
3190	Chairman WAXMAN. Take my generosity.
3191	Mr. MCHENRY. I appreciate your generosity.
3192	But in this particular case, I think we do have some
3193	disagreements. Because, based on the studies I have seen,
3194	Mr. Weemsnow, you know, Medicare Part D has cost both less
3195	for consumers that are using the program and for the
3196	taxpayers than the original cost estimate; is that correct?
3197	Mr. WEEMS. Forty percent less, yes.
3198	Mr. MCHENRY. Forty percent less?
3199	Mr. WEEMS. Yes, sir.
3200	Mr. MCHENRY. So market forces arehave been much more
3201	powerful in bringing down the cost than the government
3202	setting an arbitrary dollar amount that they will pay for an
3203	arbitrary drug?
3204	Mr. WEEMS. The power of Part D has been to use market
3205	forces to bring prices down well below those that were
3206	originally estimated.
3207	Mr. MCHENRY. Okay. There is an IMS health report in

2007. Generics--and it said, generics account for 13 of the 3208 3209 15 drugs most prescribed by Medicare Part D. All right? And 3210 also according to this study, generics accounted for 68 3211 percent of all medicines prescribed in Part D. 3212 Mr. WEEMS. Generic usage is in the 60 percentile. 3213 number is about 64 percent. 3214 Mr. MCHENRY. So can you comment on the effect that that 3215 has on the cost for the consumer, the senior and for 3216 taxpayers? 3217 Mr. WEEMS. Sure. And that was one of the points that I 3218 was making earlier. It is not an exact comparison to compare 3219 somebody who is in a price-fixed indemnity program to a 3220 risk-based program that has some additional benefits to it, 3221 you know, such as therapy management, such as therapeutic interchange. I mean, there can be and, you know, we have 3222 3223 seen scenarios where somebody who was in Medicaid came over to Medicare, was able to get more of the drugs, would be able 3224 3225 to get more drugs, the ones that they needed and, in many 3226 cases, to be able to get those at a lower price and have 3227 better health outcomes and avoid costs in the A and B part of 3228 the Medicare program. 3229 Mr. MCHENRY. I have got four questions here in 3230 succession. You can answer them just briefly. 3231 Do Medicare and Medicaid programs generally serve the same type of beneficiaries? Yes or no? 3232

3233 Mr. WEEMS. No. 3234 Mr. MCHENRY. Okay. Are Medicare and Medicaid programs 3235 l financed the same way? 3236 Mr. WEEMS. No, they're financed very differently. 3237 Mr. MCHENRY. Okay. So then is it fair to say that 3238 Medicare and Medicaid are two fundamentally different 3239 programs? 3240 Mr. WEEMS. They are. 3241 Mr. MCHENRY. They serve different beneficiaries and 3242 have different benefit structures and are financed in 3243 different ways? 3244 Mr. WEEMS. Yes. 3245 Mr. MCHENRY. So if you and I understand this correctly--I mean, obviously, by overseeing the program, you 3246 3247 know, you have a depth of knowledge. Do you believe that the price structure of one program would work for the other 3248 3249 program? 3250 Mr. WEEMS. Well, clearly, it would not be wise to move the price structure of the Medicaid program into the Medicare 3251 3252 program where there would essentially be an administered 3253 price-fixing arrangement for, you know, more than half of the 3254 pharmaceutical market in the United States. That would have, 3255 at least in my estimation, you know, considerable effects 3256 that would spill over into the private sector in terms of 3257 higher costs. So I would say that would not be particularly

3258 wise.

Mr. MCHENRY. There are some shortcomings with the program. It is a government program. It is what government does very well. Inefficiency is what government does very well. However, because market forces are involved, it has been better in terms of the cost and the benefits to consumers.

So we have talked about the negative aspects of the program. That's what this whole hearing is about, after all. That is why you have a crowd behind you and the reason why the chairman had it. But can we talk about some successes, and, you know, and answer one general question? Has Medicare Part D shown to improve beneficiary access at a less-than-expected cost?

Mr. WEEMS. Certainly. And beneficiaries are getting the drugs that they need. They are getting it in a way that is convenient to them. It is a real challenge to find any program that has a satisfaction rate of 85 percent on the part of the beneficiaries, and that's what the Medicare Part D program has. Among the low-income beneficiaries, it is 90 percent.

Mr. MCHENRY. Thank you, Mr. Chairman.

Chairman WAXMAN. Thank you, Mr. McHenry.

Mr. Weems, thank you very much for your participation.

3282 I know you're anxious to get back to the work that the

3283 government bureaucracies do so poorly, according to my friends on the other side of the aisle. But I salute you for 3284 3285 the work that you do, and we want to make laws that will make 3286 sure that we protect the taxpayers and the beneficiaries. 3287 Mr. WEEMS. Thank you for the opportunity to appear, 3288 sir. 3289 Chairman WAXMAN. For our next panel, we want to call 3290 forward Mr. Mark Merritt, President and Chief Executive Officer of the Pharmaceutical Care Management Association; 3291 3292 Mr. Rick Smith, Senior Vice President for Policy, Pharmaceutical Research Manufacturers Association, PhRMA; Mr. 3293 3294 Paul Precht, Director of Policy and Communications, Medicare 3295 Rights Center; and Ms. Judith Stein, Executive Director of 3296 the Center For Medicare Advocacy. We are very grateful for all of you coming to our 3297 3298 hearing today, and we thank you for being here. And I want 3299 to make mention of the fact that we're particularly grateful 3300 that you allow us to share Mr. Merritt's birthday with him and to have him here on this special occasion. You wouldn't 3301 3302 have wanted to be anywhere else on your birthday. 3303 Mr. MERRITT. It really is a dream come true. 3304 you. 3305 Chairman WAXMAN. Okay. Well, you said that without 3306 being under oath, but the rest of the testimony you all be 3307 asked to give -- it is the practice of this committee that it

3308 be done under oath. So I'd like to ask you to all stand.

3309 [Witnesses sworn.]

3310	RPTS COCHRAN
3311	DCMN MAGMER
3312	Chairman WAXMAN. The record will indicate that each of
3313	the witnesses answered in the affirmative.
3314	STATEMENTS OF MARK MERRITT, PRESIDENT AND CHIEF EXECUTIVE
3315	OFFICER, PHARMACEUTICAL CARE MANAGEMENT ASSOCIATION; RICK
3316	SMITH, SENIOR VICE PRESIDENT FOR POLICY, PHARMACEUTICAL
3317	RESEARCH AND MANUFACTURERS ASSOCIATION (PhRMA); PAUL PRECHT,
3318	DIRECTOR OF POLICY AND COMMUNICATIONS, MEDICARE RIGHTS
3319	CENTER; AND JUDITH STEIN, EXECUTIVE DIRECTOR, CENTER FOR
3320	MEDICARE ADVOCACY
3321	Chairman WAXMAN. Mr. Merritt, as a birthday gift to
3322	you, we are going to let you start.
3323	I think you all know the rules. Your prepared
3324	statements will be in the record in their entirety. We would
3325	like to ask you to try to limit the oral presentation to 5
3326	minutes. We have the clock

## 3327 | STATEMENT OF MARK MERRITT

Mr. MERRITT. Thank you, Mr. Chairman and Ranking Member Davis, the rest of the members who will be in and out throughout.

My name is Mark Merritt. I am President of the Pharmaceutical Care Management Association. PCMA is a national association representing America's pharmacy benefit managers. PBMs administer prescription drug benefits for more than 200 million Americans with health coverage. Our clients include the Nation's largest public and private purchasers, including labor unions, Fortune 500 companies, FEHBP plans, and, of course, Medicare.

First, I would like to thank you, Chairman Waxman, for your leadership on health care issues. PCMA is appreciative of the opportunity to work with your staff on generic biologics legislation and on ensuring generic competition in the marketplace, and I am pleased to be here today to testify about Medicare Part D and what we do in it.

To begin, PBMs use a number of tools and strategies to maximize value in terms of quality, access and convenience and overall drug spending. First, let's talk about PBMs and discounts and rebates regarding manufacturers. There, PBMs pool the purchasing ability of all our clients and consumers

and encourage certain kinds of utilization to obtain discounts and rebates from brand-name manufacturers.

First, our panels of independent clinical experts, called P&T committees, or pharmacy and therapeutic committees, comprised of independent doctors, pharmacists, academics and others, inform us of which drugs are appropriate for certain therapeutic classes which address particular medical conditions. Then we negotiate with manufacturers who make competing products within that class.

The manufacturer which offers the best discounts and rebates typically has their drugs placed on formularies at lower copays than their competitors. That encourages consumers to choose the more affordable drug, although their physician can, of course, direct them to another, if clinically appropriate.

While discounts on individual drugs can vary widely, overall, manufacturer rebates have decreased drug spending by up to 9 percent in FEHBP, according to their report. And I believe your new report, if I read it correctly--and I just got it, of course--says we save about 14 percent in Part D. But I am not sure about that.

Extracting manufacturer discounts, however, is only one way PBMs deliver savings. The majority of our savings that we generate results from innovative and aggressive management of other components of drug spending.

First, we create more affordable delivery options, such as mail service pharmacy, which can save 10 percent for payors and patients alike. Second, we aggressively negotiate more economical reimbursement and dispensing fees with drugstores in our pharmacy networks. Third, we use formularies, medication, therapy management and other tools to increase generic utilization and create a more affordable and often safer drug mix for patients. Four, we employ drug utilization review programs, or DUR, to inform patients and doctors when we identify unsafe or unnecessarily expensive prescribing patterns. And, five, we are constantly developing new innovative tools, like electronic prescribing, which improve efficiency, safety and savings across the whole system.

Today, we are proud of our accomplishments in Part D.

Costs are lower than expected, premiums are as well, generic utilization is higher and getting better, beneficiaries have broad access to formularies and drugs and have access to over 60,000 pharmacies.

Overall, our savings are comparable to those we generate in the private sector and for FEHBP plans. Most importantly, of course, beneficiaries themselves are highly satisfied with the program; and, of course, that is our marketplace.

There are, however, additional policy options that would further enhance our ability to generate savings that I would

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offer for the committee's consideration, some of which have been mentioned already today.

First, we desperately need to create competition among biologics by pursuing legislation such as your proposal, Mr. Chairman, the Access to Lifesaving Medicines Act. This is the fastest-growing component of drug spend and will reach \$100 billion sometime in the next 10 years. We need more competition in that space.

Second, we would ask policymakers to build on the groundbreaking new e-prescribing incentives that were just passed as part of the physician pay package.

Third, we would ask you to take a closer look at the six classes of clinical concern that have been mentioned earlier in which all drugs from all drug makers are mandated for coverage in certain classes, therapeutic classes. These are specifically important regarding dual eligibles, who are heavy utilizers of these drugs.

And this policy of mandating coverage, again, for all drug companies, all drugs in a certain class, we don't believe it improves access, but it does make it difficult, more difficult, for PBMs to negotiate rebates for drugs in those classes. And, again, they account for about 40 percent or more of the spending of dual-related spending.

In fact, the rebates in the six protected classes, we are only able to generate about half as much--or half of

3425 significant rebates as we are in other classes. Because when 3426 that leverage is taken away from us, it inhibits our ability 3427 to get the right discounts from the pharmaceutical 3428 manufacturers. 3429 In conclusion, though, I appreciate the opportunity to 3430 share with you our progress on my birthday and also look 3431 forward to answering any questions you might have and any 3432 concerns you might have. 3433 Chairman WAXMAN. Thank very much, Mr. Merritt. 3434 [Prepared statement of Mr. Merritt follows:] 3435 \*\*\*\*\*\* INSERT 4-1 \*\*\*\*\*\*

3436 Chairman WAXMAN. Mr. Smith.

## 3437 STATEMENT OF RICK SMITH

Mr. SMITH. Thank you, Mr. Chairman and members of the committee. Thank you for the invitation to participate in today's hearing.

My name is Richard Smith. I am Senior Vice President for Policy and Research at PhRMA, which represents pharmaceutical research companies.

Medicare Part D has greatly improved beneficiaries' access to needed medicines, reduced out-of-pocket costs and retained broad choice among medicines. This has been accomplished at much lower than anticipated cost to beneficiaries and taxpayers, and data show that Part D enrollees are highly satisfied and they are saving money.

Last week, Congress adopted an important PhRMA support improvement allowing more low-income beneficiaries to qualify for enhanced assistance.

The committee requested that I provide information on the nature of financial arrangements between pharmaceutical manufacturers and Part D plans, along with the extent of discounts. As a trade association, PhRMA maintains a strict antitrust compliance policy, so I can neither obtain nor

discuss our members' proprietary information related to prices, negotiations or discount strategies. As a result, my testimony reflects only publicly available information.

Part D was designed to achieve a range of objectives by carefully balancing affordability, access choice and improved use of medicines. This careful balance requires assessing the program on an overall basis, recognizing that its objectives are interrelated.

Part D saves beneficiaries money. Peer-reviewed research and government studies report sizeable reductions in seniors' monthly out-of-pocket costs, and premiums in 2008 are actually below the level initially projected for 2006.

Part D's competitive structure saves taxpayers money.

Both CBO and the Medicare Trustees report costs are far less than anticipated, largely because of vigorous competition.

CBO concludes plans have "secured rebates somewhat larger than the average rebates observed in commercial health plans." And the Trustees report states many brand-name prescription drugs carry substantial rebates, often as much as 20 to 30 percent.

I would also note, in the six classes, plans have an array of tools used to negotiate savings. In these classes, plans have tiers, utilization management and many generics.

Comparing CBO's 2008 and 2006 baseline shows that projected total cost for 2007 through 2016 has dropped by

\$438 billion, or 37 percent. Actual plan bids, the best measure of the program's per person cost, are 12.8 percent lower than they were 2 years ago.

Part D offers beneficiaries choice of medicines through the medicines covered by individual plans and through choice among plans. In fact, two of the largest Part D plans report covering all 100 of the most commonly used drugs; and beneficiaries are picking plans that combine no deductible, lower-than-average premium, and a broad choice of medicines.

While access to medicines has improved as intended under Part D, IMS Health estimates that the program's impact on retail pharmaceutical sales was an increase of about 1 percent in 2006. And a recent academic study reports that, overall, Part D reduced average drug prices, and the trustees have reported that rebates increased in 2008. Moreover, drug costs growth has slowed since Part D's enactment to 3.8 percent in 2007, the lowest rate since 1961.

In assessing the program's cost savings, it is important to consider the full range of populations covered and the full range of cost-saving tools used. For instance, 14 million uninsured or underinsured beneficiaries before Part D did not have discounts and rebates routinely negotiated on their behalf. Now, powerful purchasers representing millions of covered lives each negotiate savings on their behalf.

And plans use a variety of tools, among them discounts,

rebates and incentives, to increase generic use to achieve savings. As was mentioned previously, 13 of the 15 most commonly prescribed drugs in Part D are generic. These tools have produced affordable premiums and are largely responsible for the overall \$438 billion reduction in the program's total projected cost.

In conclusion, Part D has achieved its objectives for beneficiaries who clearly recognize its value. Vigorous competition has driven down costs, both for beneficiaries and taxpayers. Changing Part D's market-based structure would undermine the balanced approach which has produced sizeable cost savings and greatly improved access to needed medicines.

We look forward to working with the committee to enhance the program by building on its successful foundation, and I appreciate the opportunity to testify.

Chairman WAXMAN. Thank you very much.

[Prepared statement of Mr. Smith follows:]

3525 \*\*\*\*\*\*\* INSERT 4-2 \*\*\*\*\*\*

3526 Chairman WAXMAN. Mr. Precht.

## STATEMENT OF PAUL PRECHT

Mr. PRECHT. Thank you, Chairman Waxman, members of this committee, for this opportunity to testify.

I am Paul Precht, Director of Policy and Communications for the Medicare Rights Center.

The Medicare Rights Center is a national consumer service organization with offices in New York and Washington. Our hotline volunteers and caseworkers help older and disabled Americans deal with every conceivable type of problem standing between them and the health care they need.

Before the Part D benefit started in 2006, the most frequent call came from people with Medicare who could not afford to buy the medicines they were prescribed. Today, despite the billions in subsidies provided to the insurance companies and pharmacy benefit managers running Part D, it remains the number one problem we hear.

A typical call comes from someone making less than \$20,000 a year. More than half of the people with Medicare earn less than that amount. They don't have much to live on, but it is still too much to qualify for extra help with their prescription drug costs.

Multiple drugs to treat multiple chronic conditions put this person in the Part D coverage gap, the donut hole, where she--and it is often a widow living alone who calls--must pay both the premiums for her Part D drug coverage and the full price of her drugs. With a drug bill in excess of \$500 per month for months on end, on top of medical and other bills, the options are few. She can try to get free samples from her doctor. She can head for the emergency room. When these strategies fail, too often, she may go without the medicine she needs.

Prescription drug prices are just too high, and Part D plans are not delivering the lower prices that were promised when this benefit was created. They certainly are not providing discounts on par with the prices the VA, State Medicaid programs, or our neighbors in Canada have secured. That is widely acknowledged.

What is less well-known, however, is that the rebates and discounts that the Part D plans have been able to obtain are not passed through to consumers in the form of lower prices. That means each time a diabetic person with Medicare scrapes together the money to buy a \$400 specialty drug, the Part D plan pockets a \$30 or \$40 rebate, based on the averages that this committee has uncovered. That rebate is not used to lower the \$100 coinsurance she paid during the initial benefit period, and it does not bring down the \$400

3573 price she pays during the donut hole.

Plans argue that rebate revenue is used to keep premiums down. In effect, under this system, sick people who need expensive medicine pay a surcharge to keep costs down for their healthier neighbors. It is the opposite of the way insurance is supposed to work.

It is not just brand-name drugs that are too expensive under Part D. People with Medicare are also being overcharged for generics under some plan D plans. This scheme was described in the Wall Street Journal this week. This is how it works.

The Part D plan, an insurance company, pays its pharmacy benefits manager \$60, for example, for each prescription of generic Zocor that it covers. But the drug really costs only \$20. The pharmacy receives \$15 from the PBM and \$5 from the consumer. At the end of the month, the consumer gets a statement from the PBM saying it spent \$55 for the prescription, and the customer is \$60 closer to the donut hole.

Consumers who take a few generic drugs that are subject to these inflated prices can be pushed into the donut hole 2 or 3 months earlier in the year. What happens when consumers hit the donut hole? Do they pay the \$20, the reimbursement rate for the pharmacy? They do not. They pay \$60, and the pharmacy is forced to kick back \$40 to the PBM.

3598 l

PBMs argue this pricing scheme keeps administrative costs down for the insurance companies. But here is the twist: Sometimes the Part D plan and the PBM running this pricing scheme are part of the same company. In our view, prices are being manipulated to gouge both the consumer and Medicare, which pays more for the dual eligibles, since they pay the cost sharing.

We are 2-1/2 years into the Part D drug benefit, and even if the administration follows through on its promise to end this scheme--and they backed off last time they proposed to end it--it will continue through the end of 2009.

When the insurance industry and the PBMs talk about how Part D has marshaled market forces to lower costs, this is the market they are talking about. It is untransparent, it is rigged against consumers, particularly when they fall sick, and it does not deliver the prices consumers could receive if Medicare was negotiating with manufacturers and running the benefit.

People with Medicare should have the choice to receive drug coverage directly through Medicare. A Medicare plan that, for example, could encompass the duals, as a start, would be a good way to deal with these overcharges that we are facing.

Just one last remark. Everybody talks about the satisfaction rates with Part D. But those same polls also

3623	show similar percentages of people want a simpler benefit,
3624	they would like the option to have coverage under Medicare,
3625	and they want the government to be able to negotiate lower
3626	prices.
3627	Thank you.
3628	[Prepared statement of Mr. Precht follows:]
3629	****** INSERT 4-3 ******

3630 Chairman WAXMAN. Ms. Stein.

## 3631 STATEMENT OF JUDITH STEIN

Ms. STEIN. Good afternoon and thank you, Chairman Waxman. Thank you for being here, Mr. McHenry and Congressman Murphy.

I am Judy Stein. I am testifying today on behalf of the Center for Medicare Advocacy, of which I am the founder and Executive Director.

Since 1977, first at Connecticut Legal Services and then when I founded the Center in 1986, I have dedicated my legal career to representing Medicare beneficiaries. At the Center for Medicare Advocacy, we have represented thousands of Medicare beneficiaries and their helpers in Connecticut and across the country to understand and utilize Part D. We hear repeatedly from them about problems that arise from the complexity of the program and its ever-increasing costs. Unfortunately, problems go beyond just the dually eligible population.

There are a myriad of plans, each with varying benefit structures, formularies, out-of-pocket costs, and it makes comparisons all but impossible. Beneficiaries have insufficient information to understand formularies,

coinsurance, copayments and coverage gaps. They lack sufficient information to make sound choices. Indeed, the Center has hired an experienced advocate who dedicates all of her time just to handle the Part D problems just in Connecticut.

I thank you very much, Chairman Waxman for your leadership in investigating prescription drugs and Part D in general and Congressman Murphy for all the work he has done in our home State and now very happily here in Washington to help Medicare beneficiaries across the country.

Over the past several years, the Center has written extensively about the effects on our clients of increased reliance on private insurance plans to provide Medicare coverage. Those plans lack the stability and uniformity of the Medicare program, and they have often decreased, not increased, access to care and increased costs.

Unfortunately, the only way to get Medicare coverage for outpatient prescription drugs is through private plans. Our clients must decide each year which plan to choose from among dozens and dozens with varied cost sharing and coverage rules.

This is the packet my mother had to look through, and she is a relatively well woman who takes only three drugs. It took us hours to go through the decisions for her.

If beneficiaries seek assistance, and if it is

available, they must divulge private information about their health and medications. I don't think this has been thought of at all as one the personal expenses of the program. This information is something that many beneficiaries do not even want to share with their families. And, frankly, I was not aware of the drugs my mother took until I had to help her with Part D; and she would have preferred I didn't. It is also a step beyond to divulge this information to 1-800-MEDICARE representatives or a plan operator, and many people don't want to do that.

As a consequence, the vast majority of beneficiaries, because of these problems and others, do not in fact change plans from year to year, so the whole issue of choice is increasingly becoming a red herring. In fact, 17 percent—only 17 percent of people chose to switch plans this last year, even though it would have been in their best interests oftentimes to do so.

Our clients are subject to the whims of the companies that decide to offer drugs to the Medicare program. They must either bear the increased costs and reduced access to drugs or go through one or another an onerous process, either to choose to appeal a decision or to wait until next year when they may be able to get a better plan. Because if your health changes or the plan changes the drug's pricing or the drugs on its formulary, all of which can happen, you cannot

3702 get into a different Part D plan.

According to an ongoing study by AARP, any savings in drug costs achieved by Part D were achieved through a reduction in the cost of generic drugs. However, the prices for 169 brand-name drugs went up 50.4 percent between 2001, when the first AARP study happened, and 2007.

Higher drug costs mean that beneficiaries reach the coverage gap, or donut hole, sooner. Increased costs are causing a terrible impact on our beneficiaries, especially those who cannot take a generic equivalent, and that includes people with cancer, cardiac problems and other very significant illnesses. No stand-alone drug program offers brand-name drug coverage during the gap.

This week, a woman from California e-mailed us telling us, "I am having terrible problems trying to find a way to pick the medication for my father's chronic illness. He is diabetic, needs chemotherapy for bladder cancer, and has cardiac arrhythmia. Between him and my mother, they have only \$1,900 per month, and my father is already in the donut hole." That was in July. There are 6 more months ahead.

One of our clients in Connecticut, a 52-year-old woman, pays \$6,000 a month for her medications, if she could afford them, which she cannot. She is on Social Security Disability because of her sickle cell anemia. Her prescription drug plan refused to provide coverage for the dose needed by this

woman, even though it was ordered by her physicians, who referred her to the Center, and we appealed outside the plan finally and got coverage.

One woman in Tennessee wrote she can't afford and is therefore not taking her drugs.

In conclusion, the program has untold expenses for beneficiaries, for States who, like Connecticut, are wrapping around and paying for Medicaid beneficiaries and people on their State pharmaceutical assistance plans, and are putting ever-increasing costs of prescription drugs into the prices that taxpayers must pay for Medicare in general.

In summary, we urge the Congress to take the following steps: Include a prescription drug benefit in the traditional Medicare program and authorize the Secretary to negotiate the cost of drugs within that program at least; require drug plans to pass along the fullest extent of their rebates and include beneficiaries while they are—and include those rebates when beneficiaries are paying themselves in the gap; increase transparency by requiring drug plans to make available information about their pricing and rebates; increase oversight of the Medicare Web page, which is often very different from the information given on the plan's Web pages themselves; and require CMS to provide greater oversight of the Part D plans in their oversight.

Chairman WAXMAN. Thank you very much, Ms. Stein. We

I am going to start off the questions.

Chairman WAXMAN.

Our committee for the first time was able to analyze the drug and insurance proprietary data on drug pricing and compare the prices charged to the Medicare Part D program and the prices charged to Medicaid, and the findings reveal that the private Medicare Part D insurers are paying 30 percent more for drugs than the Medicaid program. This has resulted in a windfall of over \$3.7 billion for the drug manufacturers on the sale of drugs to dual-eligible enrollees. 

These elderly and disabled individuals used to get their drugs from Medicaid. They have switched to Medicare Part D, and now their higher drug prices are costing taxpayers billions of dollars.

Mr. Weems argued that if Medicare Part D got the same discounts for drugs that the dual eligibles that Medicaid gets, there would be a negative consequence for other Medicare beneficiaries. Specifically, he said this could lead to higher prices at the pharmacy, compromised incentives to move enrollees to generic drugs, undermine utilization management activities that plans for important safety protections as well as cost controls.

Ms. Stein, what do you think about what Mr. Weems' concerns are that he expressed to us about this issue?

Ms. STEIN. Thank you, Chairman.

Well, one of the things I think is that I added one of

3781 the economists who spoke this morning, the figures on the 3782 bottom line on Mr. Weems' chart, and they came to, I believe, \$400 billion, which I believe was also the original estimate of what the program would cost. So it seems to me that I don't understand where the savings are in that explanation that was given. I think one of the things we often find is that one has to add up the numbers and question where they are coming from.

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What I know is that we have 6,500 calls and thousands of e-mails every year at this Center. I sit in the real world listening to real people. They cannot afford these drugs. They are in the donut hole way earlier than was anticipated, and it is a problem with them. They cannot afford the drugs, and they are not getting the rebate in price when they are in the donut hole. Also, the plans don't cover their drugs, more often than not.

Chairman WAXMAN. Thank you very much.

Mr. Precht, what do you think of the argument that we are really doing a favor for the rest of the Medicare beneficiaries by paying a higher price for the dual eligibles?

Mr. PRECHT. I am not an economist, but it doesn't make any sense to me. It seems that there is money that is going into the pharmaceutical manufacturers, rather than into providing coverage for people with Medicare; and it certainly

seems we could use that money to get more people into the extra health program, for example, so they wouldn't have to pay full price in the donut hole.

It seems to me that if there were competition between the private plans and a Medicare option that negotiated its rates that that would provide some price discipline and it could result in lower prices, both in the Medicare option as well as the private option.

Chairman WAXMAN. Mr. Merritt and Mr. Smith, do you disagree with the report's findings that the manufacturers are charging more for drugs under Medicare Part D for dual eligibles than they are under Medicaid?

Mr. SMITH. Mr. Chairman, I haven't had an opportunity to review the report. It certainly wouldn't surprise me if the type of market-based system we have, with very powerful large purchasers, lots of tools at their disposal--Mr. Merritt described those--negotiated a price that was different than the price that was previously set through the administered pricing system of Medicaid.

I think it is important to recognize that the-Chairman WAXMAN. You say because of all the strong
tools they have they negotiated a price that is higher than
Medicaid?

Mr. SMITH. I am saying there might be a valuation in the marketplace that is different than the valuation through

3831 the administered pricing system of Medicaid.

Chairman WAXMAN. So you think Medicaid is lower priced, and we have moved to a higher price system under Part D through the private plans?

Mr. SMITH. Mr. Chairman, without having had an opportunity to review the report, I am simply saying that I can imagine that private purchasers with lots of tools negotiating come up with different valuations than does an administered pricing system.

And when we look at the entire population, including the 14 million individuals who previously weren't typically having discounts and rebates negotiated on their behalf, I think that we see that there is considerable price pressure.

Chairman WAXMAN. How about just the 6 million that are dual eligibles? With all these tools that the private plans have for negotiating better prices, why are we paying more for that distinct population for their drugs than we were under Medicaid?

Mr. SMITH. Well, I believe, first, that private plans negotiate for entire populations, so average rebates for entire populations may differ than average rebates for a segment of the population. They may also use a different mix of savings mechanisms. They may use more than rebates of savings mechanisms. And, ultimately, I think it is difficult to pull the one population out, look at it separately from

3856 the entirety the population being covered and for which 3857 savings is being negotiated. 3858 Chairman WAXMAN. Would you include the private-sector 3859 coverage for non-Medicare? Would you put them in the overall 3860 picture? 3861 Mr. SMITH. I am not quite sure I understand the 3862 question, Mr. Chairman. 3863 Chairman WAXMAN. I will send you a letter about it 3864 afterwards. 3865 Mr. McHenry. 3866 Mr. MCHENRY. Thank you, Mr. Chairman. 3867 You know, this committee is trying to find efficiency in government, and I appreciate it. It has taken us a while to 3868 3869 actually get to hearings that get to that during this 3870 Congress, but I am glad that we can actually have this 3871 discussion. 3872 I do have a question. Mr. Precht, we are speaking about 3873 Medicare Part D today. But, admittedly, Medicare is a larger 3874 issue that we are concerned about. 3875 Ms. Stein, I appreciate your advocacy and help in this process and helping American seniors get the information they 3876 3877 need to make good decisions about this. But, you know, I 3878 would like to know, because you are concerned about Medicare 3879 rights, Mr. Precht, are you concerned about the financial adequacy of Medicare Part A? 3880

3881 Mr. PRECHT. Yes, sir, very much. Mr. MCHENRY. In terms of the amount of money the 3882 government spends, isn't it far greater in Medicare Part A? 3883 Mr. PRECHT. That is correct. There is more money spent 3884 3885 on hospital care than on prescription drugs. 3886 Mr. MCHENRY. Do you think we should be looking at that 3887. as a Congress? 3888 Mr. PRECHT. Absolutely. 3889 Mr. MCHENRY. Okay. I mean, the price differential 3890 between the two is significant. It is--what--about \$200 3891 billion--\$220 billion for Medicare Part A and about \$50 3892 billion for Medicare Part D. Is that roughly correct? 3893 not trying to put you on the spot. 3894 Mr. PRECHT. I will take your word for it. 3895 I mean, there is certainly more spending. I guess I 3896 don't know. I am not as familiar as I should be with 3897 research that looks at the spending under Part A and whether we could be saving money. But I think probably there are 3898 3899 ways to save money there as well. 3900 Mr. MCHENRY. Ms. Stein, to your comment that 3901 beneficiaries are struggling with ever-increasing 3902 prices -- and, generally speaking, in this time right now of inflation, we are all struggling with high prices--gas 3903 3904 prices, food prices and everything else. It is putting a 3905 pinch on seniors, especially. But in terms of the Medicare

3906 Part D beneficiaries and what they pay in premiums, has that 3907 gone up? 3908 Yes, sir. In fact, my--for instance, Humana Ms. STEIN. 3909 has gone up three times what it was in the first year of the 3910 program. 3911 And, by the way, with regard to Part A, the Center for Medicare Advocacy is extremely concerned about the cost of 3912 3913 Medicare in general, and we do a great deal of work with 3914 regard to those issues. 3915 Mr. MCHENRY. Sure. Back to the point of what the 3916 beneficiaries are paying, according to the CBO, the cost estimate at the beginning of this program was, I believe, \$37 3917 3918 or \$35, and CMS estimated about the same at the beginning of 3919 the program. I think CMS estimated \$37. CBO said \$35. fact, the Democrats had an amendment in committee to set the 3920 3921 price of premiums for seniors at \$35. Well, premiums are 3922 under \$25 right now across the population for all beneficiaries, is that not correct? 3923 3924 Ms. STEIN. For all beneficiaries, the premiums went 3925 down. For plans that people were in, they often went up, and 3926 they didn't switch. So that people were in a plan in the 3927 first year, their premium went up three times in the second year for one of the entities that has the largest population. 3928 3929 Mr. MCHENRY. Sure. But there are other entities by 3930 which they can say, I am done with Humana. I am going over

here. There are enough forces out there--3931 3932 Ms. STEIN. Because of the structure of the program--3933 Mr. MCHENRY. Ma'am, let me finish asking the question. 3934 There are enough in the way of choices out there that 3935 seniors can make an informed decision; and if on average the 3936 premiums have gone down, isn't that a good thing? 3937 Ms. STEIN. It depends, sir. In my mother's case, for 3938 instance, yes, she takes two drugs. She decided to stay in 3939 her plan because it was a lower premium, she thought. But it 3940 didn't cover one of her drugs. So you could choose a premium 3941 that is lower this year but not get your drug coverage. 3942 is as not as simple as that. 3943 Mr. MCHENRY. Because an individual makes a mistake 3944 doesn't mean it is a bad policy or bad program. Mistakes are 3945 made every day. After all, look at the United States 3946 Congress. We have made mistakes. We are all human. 3947 Ms. STEIN. With all due respect, sir, just let me say 3948 this. There is only 17 percent of people that switched 3949 plans. So the fact is that people, for whatever reason--I believe the design of the program -- are not utilizing the 3950 3951 choice option because it is so complex. And the fact is 3952 that, if they do choose based on the lowest-cost premium, 3953 they may well find themselves in the wrong plan. 3954 Mr. MCHENRY. Okay. Thank you. I appreciate your 3955 testimony.

I have one final question for Mr. Smith, if I may, Mr. Chairman.

Overall, we are talking about price negotiation. That is a part of this. And the majority report, the Democrat report from this committee, expresses that there will be a windfall--quote-unquote, windfall to the pharmaceutical industry unless government negotiated the price. Even though what they failed to mention is that private entities, all these different insurers, are negotiating for the price of drugs. So, therefore, they want the government to step in and say all these different insurers have to accept this price.

Okay. If there is a windfall for the pharmaceutical industry, how much has your business gone up? Because the statistic I have, in your testimony, is that prescription drug sales have increased by only 1 percent since Medicare Part D was implemented. Where is the windfall?

Mr. SMITH. Yes, sir. I would, of course, view prices that are set by very powerful purchasers negotiating very aggressively for prices and the resulting prices as not generating a windfall. The basic result has been that, in 2008, prescription drug costs in the United States went up by the lowest rate since 1961, 3.8 percent, and the slowdown in growth continues. IMS Health reports, for the 12 months ended May of this year, the growth rate for prescription

3981 medicines in the United States, the entire cost for the whole 3982 country, was 1 percent. 3983 Mr. MCHENRY. Thank you, Mr. Chairman. 3984 Mr. MURPHY [Presiding.] Thank you, Mr. McHenry. 3985 Smith, I want to get back to follow up on a few of Chairman Waxman's questions. I know he may follow up with 3986 3987 you in written correspondence. 3988 But with regard to the differences between the 3989 negotiations that happened with private plans and the 3990 Medicaid rebate system, your ultimate leverage in a 3991 negotiation with a particular health care plan is to not sell 3992 that drug to that plan, to not be part of their formulary, is 3993 that correct? 3994 Mr. SMITH. Without suggesting proprietary information 3995 about business practices, I think that would generally 3996 accurately characterize the market. 3997 Mr. MURPHY. With regard to the Medicare rebate system, 3998 your ultimate leverage with the Medicaid rebate system is to 3999 voluntarily not sell your drug as a part of the Medicaid 4000 system? 4001 Mr. SMITH. That is correct. On a one-size-fits-all 4002 basis, you are really excluded from a very large portion of 4003 the market entirely, very different from the private sector. 4004 Mr. MURPHY. Because the purchasing pool is so large 4005 from the Medicaid side, because, as you say, it is a

4006 one-size-fits-all, the decision is much harder to not sell 4007 the drug to the Medicaid system. 4008 Mr. SMITH. Well, there is no real opportunity to 4009 reflect value, because there is that statutory formula that 4010 sets the price. So I think that one of the challenges is 4011 that there really is no negotiation in that respect because it is a decision that is generated by a statutory pricing 4012 4013 formula. 4014 Mr. MURPHY. But you are not compelled to sell the drug? 4015 Mr. SMITH. It is either sell at that statutory formula 4016 or be excluded from the entire Medicaid market. 4017 Mr. MURPHY. Ms. Stein, the report that is released today details a 6.6 percent increase in the average cost of a 4018 drug from 2006 to 2007, which is about twice the rate of 4019 4020 inflation. You suggested some of the impacts of this in your 4021 testimony. 4022 But I just wanted to ask you, what is the impact of that 4023 6.6 percent increase in the price of the drug to an average 4024 health care consumer in the Part D system, given I think the 4025 testimony that you have given about the number of people 4026 falling into the donut hole earlier than expected or earlier 4027 than people had hoped for? 4028 Ms. STEIN. Sir, they are very often in the donut hole 4029 earlier. Once they are there, they are paying the full cost 4030 of the drug, not with the rebate. People, as you will see in

my written testimony, are taking less than the full prescription which has been given by their physician, as someone is quoted in my testimony. Particularly people on psychotropic drugs we find are not taking their medications. Many of them don't like to take them in the first place.

So we have a lot of problems with the fact that people aren't taking the medications or taking less than has been prescribed, and they are falling into the donut hole earlier.

I would also like to suggest there are tremendous costs to the States as a consequence, which, as you know in Connecticut, we are also paying--when the people fall into the donut hole, we are paying those coinsurances. And on specialty drugs that can be for the individual as well as for the State up to 33 percent of the cost of that special brand-name drug.

Mr. MURPHY. The last question, just to make this point clear, when an individual falls into the donut hole, when they come to pay for the price at the retail pharmacy, they are not getting the benefit, certainly not of Medicaid, but they are not getting the benefit of the potential discount negotiated by the HMO they were covered which?

Ms. STEIN. That is correct. That is included and helps them get into the donut hole sooner. Once they are in the donut hole, they don't have the benefit of that; and they pay more, therefore.

4056 Mr. MURPHY. Thank you, Ms. Stein. 4057 Thank you very much to the entire panel. We will keep 4058 the record open for further comments and statements. 4059 I would like to add without objection for the record a 4060 statement for today's hearing submitted by America's Health 4061 Insurance Plans. 4062 Without objection, that is entered into the record. 4063 [The information follows:] 4064 \*\*\*\*\* COMMITTEE INSERT \*\*\*\*\*\*

4066 to our previous two panels.  4067 This hearing is adjourned.	4065	Mr. MURPHY. Again, thank you to	o this panel. Thank you	
This hearing is adjourned.	4066	to our previous two panels.		
	4067	This hearing is adjourned.	•	
[Whereupon, at 2:26 p.m., the committee was a	4068	[Whereupon, at 2:26 p.m., the co	ommittee was adjourned.]	