

# PhRMA, Senate Panel Reach Deal on Part D

BY JOYCE FRIEDEN

Major pharmaceutical firms have agreed to offer drug discounts to Medicare beneficiaries trapped in the Part D “doughnut hole,” President Obama has announced.

The president endorsed an agreement reached between the Pharmaceutical Research and Manufacturers of America (PhRMA) and Sen. Max Baucus (D-

Mont.), chairman of the Senate Finance Committee. Mr. Obama explained that “as part of the health care reform I expect Congress to enact this year, Medicare beneficiaries whose spending falls within this gap will now receive a discount on prescription drugs of at least 50% from the negotiated price their plan pays. It’s a reform that will make prescription drugs more affordable for millions of seniors, and restore a measure of

fairness to Medicare Part D.” The estimated cost of the discount program, which applies only to brand-name drugs, is \$80 billion over the next decade.

Medicare Part D enrollees who are in the doughnut hole will receive their discounts at the pharmacy and will not have to fill out any additional paperwork. They also will receive credit for the full cost of a drug against their spending obligation in the doughnut hole, even though they are actually paying half that amount.

President Obama noted that under the Medicare Part D prescription drug benefit, “Medicare covers up to \$2,700 in yearly prescription costs and then stops, and the coverage starts back up when the costs exceed \$6,100. [That] means between \$2,700 and \$6,100, folks are out of luck. And this gap in coverage has placed a crushing burden on many older Americans who live on fixed incomes and can’t afford thousands of dollars in out-of-pocket expenses.”

At the White House event, Barry Rand, CEO of AARP, which endorsed the agreement, called the deal “an early win for reform and a major step forward.”

Mr. Rand said, “Too many Americans who fall into the coverage gap stop taking their medications because they simply cannot afford them. They will now have a new opportunity to lead a healthier life.”

Billy Tauzin, president and CEO of PhRMA, noted in a statement that “even

though we have had policy disagreements in the past [with AARP], this is an historic coming-together moment. AARP, the largest advocacy organization on behalf of American seniors, clearly recognizes the importance of innovative, cutting-edge medicines to the lives of patients everywhere.

Sen. Baucus noted in a statement that when it was created, the Part D benefit “helped address the problem of skyrocketing prescription drug prices for millions of seniors. [With this agreement] we helped fill the gap in coverage and finished the job. ... This benefit is part of our continued commitment to seniors and our ongoing effort to reform health care by lowering health care costs and ensuring all Americans have access to the quality, affordable health care coverage they deserve.”

The Medicare Rights Center, a consumer group that advocates improved Medicare benefits, expressed cautious optimism about the agreement. “As always, the devil is in the details,” center president Joe Baker said in a statement. “We look forward to working with President Obama and the Congress to making the promised discount most useful.” He added that the discount complements the health reform proposal from the chairmen of three House committees to phase out the Part D doughnut hole. “Full coverage of both brand-name and generic drugs is the best way to ensure people with Medicare can afford the medicines they need,” Mr. Baker said.

Combined administration of racemic citalopram (40 mg) and ketoconazole (200 mg), a potent CYP3A4 inhibitor, decreased the C<sub>max</sub> and AUC of ketoconazole by 21% and 10%, respectively, and did not significantly affect the pharmacokinetics of citalopram. **Ritonavir**-Combined administration of a single dose of ritonavir (600 mg), both a CYP3A4 substrate and a potent inhibitor of CYP3A4, and escitalopram (20 mg) did not affect the pharmacokinetics of either ritonavir or escitalopram. **CYP3A4 and -2C19 Inhibitors**-*In vitro* studies indicated that CYP3A4 and -2C19 are the primary enzymes involved in the metabolism of escitalopram. However, coadministration of escitalopram (20 mg) and ritonavir (600 mg), a potent inhibitor of CYP3A4, did not significantly affect the pharmacokinetics of escitalopram. Because escitalopram is metabolized by multiple enzyme systems, inhibition of a single enzyme may not appreciably decrease escitalopram clearance. **Drugs Metabolized by Cytochrome P4502D6**-*In vitro* studies did not reveal an inhibitory effect of escitalopram on CYP2D6. In addition, steady state levels of racemic citalopram were not significantly different in poor metabolizers and extensive CYP2D6 metabolizers after multiple-dose administration of citalopram, suggesting that coadministration, with escitalopram, of a drug that inhibits CYP2D6, is unlikely to have clinically significant effects on escitalopram metabolism. However, there are limited *in vivo* data suggesting a modest CYP2D6 inhibitory effect for escitalopram, i.e., coadministration of escitalopram (20 mg/day for 21 days) with the tricyclic antidepressant desipramine (single dose of 50 mg), a substrate for CYP2D6, resulted in a 40% increase in C<sub>max</sub> and a 100% increase in AUC of desipramine. The clinical significance of this finding is unknown. Nevertheless, caution is indicated in the coadministration of escitalopram and drugs metabolized by CYP2D6. **Metoprolol**-Administration of 20 mg/day Lexapro for 21 days in healthy volunteers resulted in a 50% increase in C<sub>max</sub> and 82% increase in AUC of the beta-adrenergic blocker metoprolol (given in a single dose of 100 mg). Increased metoprolol plasma levels have been associated with decreased cardioselectivity. Coadministration of Lexapro and metoprolol had no clinically significant effects on blood pressure or heart rate. **Electroconvulsive Therapy (ECT)**-There are no clinical studies of the combined use of ECT and escitalopram.

**USE IN SPECIFIC POPULATIONS: Pregnancy**-Pregnancy Category C-In a rat embryo/fetal development study, oral administration of escitalopram (56, 112, or 150 mg/kg/day) to pregnant animals during the period of organogenesis resulted in decreased fetal body weight and associated delays in ossification at the two higher doses (approximately ≥ 56 times the maximum recommended human dose [MRHD] of 20 mg/day on a body surface area [mg/m<sup>2</sup>] basis). Maternal toxicity (clinical signs and decreased body weight gain and food consumption), mild to 56 mg/kg/day, was present at all dose levels. The developmental no-effect dose of 56 mg/kg/day is approximately 28 times the MRHD on a mg/m<sup>2</sup> basis. No teratogenicity was observed at any of the doses tested (as high as 75 times the MRHD on a mg/m<sup>2</sup> basis). When female rats were treated with escitalopram (6, 12, 24, or 48 mg/kg/day) during pregnancy and through weaning, slightly increased offspring mortality and growth retardation were noted at 48 mg/kg/day which is approximately 24 times the MRHD on a mg/m<sup>2</sup> basis. Slight maternal toxicity (clinical signs and decreased body weight gain and food consumption) was seen at this dose. Slightly increased offspring mortality was also seen at 24 mg/kg/day. The no-effect dose was 12 mg/kg/day which is approximately 6 times the MRHD on a mg/m<sup>2</sup> basis. In animal reproduction studies, racemic citalopram has been shown to have adverse effects on embryo/fetal and postnatal development, including teratogenic effects, when administered at doses greater than human therapeutic doses. In two rat embryo/fetal development studies, oral administration of racemic citalopram (52, 56, or 112 mg/kg/day) to pregnant animals during the period of organogenesis resulted in decreased embryo/fetal growth and survival and increased incidence of fetal abnormalities (including cardiovascular and skeletal defects) at the high dose. This dose was also associated with maternal toxicity (clinical signs, decreased body weight gain). The developmental no-effect dose was 56 mg/kg/day. In a rabbit study, no adverse effects on embryo/fetal development were observed at doses of racemic citalopram of up to 16 mg/kg/day. Thus, teratogenic effects of racemic citalopram were observed at a maternally toxic dose in the rat and were not observed in the rabbit. When female rats were treated with racemic citalopram (4.8, 12.8, or 32 mg/kg/day) from late gestation through weaning, increased offspring mortality during the first 4 days after birth and persistent offspring growth retardation were observed at the highest dose. The no-effect dose was 12.8 mg/kg/day. Similar effects on offspring mortality and growth were seen when dams were treated throughout gestation and early lactation at doses ≥ 24 mg/kg/day. A no-effect dose was not determined in that study. There are no adequate and well-controlled studies in pregnant women; therefore, escitalopram should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus. **Pregnancy-Nonteratogenic Effects**-Neonates exposed to Lexapro and other SSRIs or SNRIs, late in the third trimester, have developed complications requiring prolonged hospitalization, respiratory support, and tube feeding. Such complications can arise immediately upon delivery. Reported clinical findings have included respiratory distress, cyanosis, apnea, seizures, temperature instability, feeding difficulty, vomiting, hypoglycemia, hypotonia, hyperreflexia, tremor, jitteriness, irritability, and constant crying. These features are consistent with either a direct toxic effect of SSRIs and SNRIs or, possibly, a drug discontinuation syndrome. It should be noted that, in some cases, the clinical picture is consistent with serotonin syndrome [see *Warnings and Precautions*]. Infants exposed to SSRIs in late pregnancy may have an increased risk for persistent pulmonary hypertension of the newborn (PPHN). PPHN occurs in 1-2 per 1000 live births in the general population and is associated with substantial neonatal morbidity and mortality. In a retrospective, case-control study of 377 women whose infants were born with PPHN and 836 women whose infants were born healthy, the risk for developing PPHN was approximately six-fold higher for infants exposed to SSRIs after the 20th week of gestation compared to infants who had not been exposed to antidepressants during pregnancy. There is currently no corroborative evidence regarding the risk for PPHN following exposure to SSRIs in pregnancy; this is the first study that has investigated the potential risk. The study did not include enough cases with exposure to individual SSRIs to determine if all SSRIs posed similar levels of PPHN risk. When treating a pregnant woman with Lexapro during the third trimester, the physician should carefully consider both the potential risks and benefits of treatment [see *Dosage and Administration*]. Physicians should note that in a prospective longitudinal study of 201 women with a history of major depression who were authentic at the beginning of pregnancy, women who discontinued antidepressant medication during pregnancy were more likely to experience a relapse of major depression than women who continued antidepressant medication. **Labor and Delivery**-The effect of Lexapro on labor and delivery in humans is unknown. **Nursing Mothers**-Escitalopram is excreted in human breast milk. Limited data from women taking 10-20 mg escitalopram showed that exclusively breast-fed infants receive approximately 3.9% of the maternal weight-adjusted dose of escitalopram and 1.7% of the maternal weight-adjusted dose of desmethylcitalopram. There were two reports of infants experiencing excessive somnolence, decreased feeding, and weight loss in association with breastfeeding from a racemic citalopram-treated mother; in one case, the infant was reported to recover completely upon discontinuation of racemic citalopram by its mother and, in the second case, no follow-up information was available. Caution should be exercised and breast-feeding infants should be observed for adverse reactions when Lexapro is administered to a nursing woman. **Pediatric Use**-Safety and effectiveness of Lexapro has not been established in pediatric patients (less than 12 years of age) with Major Depressive Disorder. Safety and effectiveness of Lexapro has been established in adolescents (12 to 17 years of age) for the treatment of major depressive disorder [see *Clinical Studies*]. Although maintenance efficacy in adolescent patients with Major Depressive Disorder has not been systematically evaluated, maintenance efficacy can be extrapolated from adult data along with comparisons of escitalopram pharmacokinetic parameters in adults and adolescent patients. Safety and effectiveness of Lexapro has not been established in pediatric patients less than 18 years of age with Generalized Anxiety Disorder. **Geriatric Use**-Approximately 6% of the 1144 patients receiving escitalopram in controlled trials of Lexapro in major depressive disorder and GAD were 60 years of age or older; elderly patients in these trials received daily doses of Lexapro between 10 and 20 mg. The number of elderly patients in these trials was insufficient to adequately assess for possible differential efficacy and safety measures on the basis of age. Nevertheless, greater sensitivity of some elderly individuals to effects of Lexapro cannot be ruled out. SSRIs and SNRIs, including Lexapro, have been associated with cases of clinically significant hyponatremia in elderly patients, who may be at greater risk for this adverse event [see *Hyponatremia*]. In two pharmacokinetic studies, escitalopram half-life was increased by approximately 50% in elderly subjects as compared to young subjects and C<sub>max</sub> was unchanged [see *Clinical Pharmacology*]. 10 mg/day is the recommended dose for elderly patients [see *Dosage and Administration*]. Of 4422 patients in clinical studies of racemic citalopram, 1357 were 60 and over, 1034 were 65 and over, and 457 were 75 and over. No overall differences in safety or effectiveness were observed between these subjects and younger subjects, and other reported clinical experience has not identified differences in responses between the elderly and younger patients, but again, greater sensitivity of some elderly individuals cannot be ruled out.

**DRUG ABUSE AND DEPENDENCE: Abuse and Dependence**-Physical and Psychological Dependence-Animal studies suggest that the abuse liability of racemic citalopram is low. Lexapro has not been systematically studied in humans for its potential for abuse, tolerance, or physical dependence. The postmarketing clinical experience with Lexapro did not reveal any drug-seeking behavior. However, these observations were not systematic and it is not possible to predict on the basis of this limited experience the extent to which a CNS-active drug will be misused, diverted, and/or abused once marketed. Consequently, physicians should carefully evaluate Lexapro patients for history of drug abuse and follow such patients closely, observing them for signs of misuse or abuse (e.g., development of tolerance, incrementations of dose, drug-seeking behavior).

**OVERDOSAGE: Human Experience**-In clinical trials of escitalopram, there were reports of escitalopram overdose, including overdoses of up to 600 mg, with no associated fatalities. During the postmarketing evaluation of escitalopram, Lexapro overdoses involving overdoses of over 1000 mg have been reported. As with other SSRIs, a fatal outcome in a patient who has taken an overdose of escitalopram has been rarely reported. Symptoms most often accompanying escitalopram overdose, alone or in combination with other drugs and/or alcohol, included convulsions, coma, dizziness, hypotension, insomnia, nausea, vomiting, sinus tachycardia, somnolence, and ECG changes (including QT prolongation and very rare cases of torsade de pointes). Acute renal failure has been very rarely reported accompanying overdose. **Management of Overdose**-Establish and maintain an airway to ensure adequate ventilation and oxygenation. Gastric evacuation by lavage and use of activated charcoal should be considered. Careful observation and cardiac and vital sign monitoring are recommended, along with general symptomatic and supportive care. Due to the large volume of distribution of escitalopram, forced diuresis, dialysis, hemoperfusion, and exchange transfusion are unlikely to be of benefit. There are no specific antidotes for Lexapro. In managing overdose, consider the possibility of multiple-drug involvement. The physician should consider contacting a poison control center for additional information on the treatment of any overdose.

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## FYI

### Booklet on Mental Illness in Women

The Substance Abuse & Mental Health Service Administration has created a booklet, “Women’s Mental Health: What It Means to You” to provide information on symptoms, and suggestions for support, and solutions to address women’s mental health issues. The booklet can be downloaded from [mentalhealth.samhsa.gov](http://mentalhealth.samhsa.gov).

### Mapping Out the Teen Years

The Substance Abuse and Mental Health Services Administration is offering a free, high-quality CD called “The Teen Years: A Road Map for Parents,” that provides expert advice on adolescence and guidance aimed at helping adolescents through this transitional time. For more information about the booklet, visit <http://ncadistore.samhsa.gov/catalog/ProductDetails.aspx?ProductID=17459>.

### Report on Women’s Mental Health

A report from the Substance Abuse and Mental Health Services Administration, “Action Steps for Improving Women’s Mental Health,” compiles the latest research and resources on women’s mental health, and provides strategies for policy action and research. For online access to the report, visit [mentalhealth.samhsa.gov](http://mentalhealth.samhsa.gov).

### Preventing Teen Drug Abuse

The Substance Abuse and Mental Health Services Administration is offering a free booklet, “Keeping Your Kids Drug Free: A Family Guide,” which has been developed to help parents keep their kids away from marijuana and other illicit drugs. Versions aimed at Spanish speakers and African Americans also are available. For more information, visit <http://ncadistore.samhsa.gov/catalog/ProductDetails.aspx?ProductID=17293>.

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