

**CENTER FOR DRUG EVALUATION AND RESEARCH**

**Approval Package for:**

**Application Number: 20822**

**Trade Name: CELEXA 20MG, 40MG, and 60MG  
TABLETS**

**Generic Name: CITALOPRAM HYDROBROMIDE**

**Sponsor: FOREST LABORATORIES, INC.**

**Approval Date: 07/17/98**

**Indication(s): TREATMENT OF DEPRESSION**

# CENTER FOR DRUG EVALUATION AND RESEARCH

**APPLICATION: 20822**

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**CENTER FOR DRUG EVALUATION AND RESEARCH**

**Application Number: 20822**

**APPROVAL LETTER**

NDA 20-822

Forest Laboratories, Inc.  
Attention: Kathryn Bishburg, Pharm.D.  
Associate Director, Regulatory Affairs  
909 Third Avenue  
New York, New York 10022-4731

Dear Dr. Bishburg:

Please refer to your New Drug Application dated May 7, 1997, received May 12, 1997, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act for Celexa™ (citalopram hydrobromide) 20 mg, 40 mg, and 60 mg Tablets.

Reference is also made to an agency approvable letter dated May 12, 1998.

We acknowledge receipt of your submissions dated May 22, June 15, June 22, July 6, July 16, and July 17, 1998. Your submission dated May 22, and received May 26, 1998 constituted a full response to our May 12, 1998 action letter. The user fee goal date for this application is July 26, 1998.

This new drug application provides for the use of Celexa™ to treat depression.

We have completed the review of this application, as amended, and have concluded that adequate information has been presented to demonstrate that the drug product is safe and effective for use as recommended in the enclosed labeling text. Accordingly, the application is approved effective on the date of this letter.

Accompanying this letter (ATTACHMENT) is the labeling, including the revisions agreed to, that should be used for marketing this drug product. These revisions are terms of the NDA approval. Marketing the product before making the agreed upon revisions in the product's labeling may render the product misbranded and an unapproved new drug.

Additionally, we note that you do not intend to market, at this time, the 60 mg strength and, as such, references to this strength have been removed from the attached labeling.

We have the following additional comments:

#### **Phase 4 Commitments**

We remind you of your Phase 4 commitments specified in your submissions dated May 22, July 6, July 16, and July 17, 1998. These commitments, along with any completion dates agreed upon, are listed below.

1. (b)(4)(CC)-----

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314.82(b)(2)(vii), we request that you include a status summary of each commitment in your annual report to this NDA. The status summary should include the number of patients entered in each study, expected completion and submission dates, and any changes in plans since the last annual report. For administrative purposes, all submissions, including labeling supplements, relating to these Phase 4 commitments must be clearly designated "Phase 4 Commitments."

## MANUFACTURING AND CONTROLS

### 1. Expiration Date

As requested in your correspondence dated May 22, 1998, the Agency is approving an initial 24-month expiration dating period based upon your currently available 18-month data.

### 2. Nomenclature

We again note that the established name, citalopram hydrobromide, has not been adopted by the USAN Council. We acknowledge your commitment to submit a copy of the USAN approval letter to the Agency when granted.

### 3. Methods Validation

Validation of the regulatory methods has not been completed. At the present time, it is the policy of the Center not to withhold approval because the methods are being validated. Nevertheless, we expect your continued cooperation to resolve any problems that may be identified.

## BIOPHARMACEUTICS

We note your agreement to the following dissolution method and specification for all tablet strengths:

Apparatus:	USP Apparatus 1 (basket)
Speed:	100 RPM
Medium:	800(b)(4)ffer (pH = 1.5) at 37 °C
Specification:	Q =(CC)--t 30 minutes

Please submit 20 copies of the FPL as soon as it is available, in no case more than 30 days after it is printed. Please individually mount ten of the copies on heavy-weight paper or similar material. For administrative purposes, this submission should be designated "FPL for approved NDA 20-822." Approval of this submission by FDA is not required before the labeling is used.

In addition, please submit three copies of the introductory promotional materials that you propose to use for this product. All proposed materials should be submitted in draft or mock-up form, not final print. Please send one copy to the Neuropharmacological Drug Products and two copies of both the promotional materials and the package insert directly to:

Division of Drug Marketing, Advertising, and Communications, HFD-40  
Food and Drug Administration  
5600 Fishers Lane

Rockville, Maryland 20857

Please submit one market package of the drug product when it is available.

We remind you that you must comply with the requirements for an approved NDA set forth under 21 CFR 314.80 and 314.81.

If you have any questions, contact Paul David, R.Ph., Regulatory Project Manager, at (301) 594-5530.

Sincerely yours,

APPEARS THIS WAY  
ON ORIGINAL

Robert Temple, M.D.  
Director  
Office of Drug Evaluation I  
Center for Drug Evaluation and Research

ATTACHMENT

APPEARS THIS WAY  
ON ORIGINAL

**CENTER FOR DRUG EVALUATION AND RESEARCH**

**APPLICATION NUMBER:20822**

**APPROVABLE LETTER**

MAY 12 1998

NDA 20-822

Forest Laboratories, Inc.  
Attention: Kathryn Bishburg, Pharm.D.  
Associate Director, Regulatory Affairs  
909 Third Avenue  
New York, New York 10022-4731

Dear Dr. Bishburg:

Please refer to your New Drug Application dated May 7, 1997, submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act for Celexa™ (citalopram hydrobromide) 10 mg, 20 mg, 40 mg, and 60 mg Tablets.

We also acknowledge receipt of the following additional communications:

July 7, 1997	July 9, 1997	July 15, 1997	July 16, 1997
July 29, 1997	July 30, 1997	July 31, 1997	August 14, 1997
August 21, 1997(2)	August 28, 1997	September 4, 1997	September 5, 1997
October 9, 1997	October 14, 1997	October 24, 1997	October 28, 1997
November 7, 1997	November 18, 1997	November 21, 1997	November 26, 1997
December 12, 1997	December 18, 1997	January 2, 1998	<del>January 16, 1998</del>
January 19, 1998	January 21, 1998	February 6, 1998	February 11, 1998
February 23, 1998	February 26, 1998	March 25, 1998	March 26, 1998
March 30, 1998	April 1, 1998	April 2, 1998	April 15, 1998

We have completed the review of this application as submitted with draft labeling, and it is approvable. Before the application may be approved, however, it will be necessary for you to respond to the following items:

## CLINICAL

### 1. Labeling

Accompanying this letter (ATTACHMENT) is the Agency's proposal for the labeling of Celexa. We believe it presents a fair summary of the information available on the benefits and risks of citalopram. Please use the proposed text verbatim.

We have proposed a number of changes to the draft labeling submitted in your original May 7, 1997 submission, and explanations for these changes are provided in the bracketed comments embedded within the proposed text. Division staff would be happy to discuss these proposed changes in detail, and we would be happy to meet with you to discuss any disagreements you might have with any part of the proposed labeling format or content.

2. Safety Update

Our assessment of the safety of citalopram is based on our review of all safety information provided in your original submission. This original review was based on an integrated database for studies with a cutoff date of October 1, 1996 and postmarketing data with a cutoff date of October 31, 1996. Please provide a final safety update focusing on deaths, serious adverse events, and dropouts for adverse events in clinical trials. In addition, please provide an updated report on the postmarketing experience with citalopram with a more recent cutoff date.

3. Regulatory Status Update

Please provide any new information on the regulatory status of citalopram worldwide. We require a review of the status of all actions with regard to this drug, either taken or pending before foreign regulatory authorities. Approval actions can be noted, but we ask that you describe in detail any and all actions taken that have been negative, supplying a full explanation of the views of all parties and the resolution of the matter. In addition, we ask that you provide us any current foreign labeling for citalopram, if appropriate, along with English translations when needed. It is only necessary to provide information that is more recent than that provided in your original May 7, 1997 submission.

4. World Literature Update

Prior to the approval of citalopram, we require an updated report on the world's archival literature pertaining to the safety of citalopram. This report should include only literature not covered in your previous submissions. We need your warrant that you have reviewed this literature systematically, and in detail, and that you have discovered no finding that would adversely affect conclusions about the safety of citalopram. The report should also detail how the literature search was conducted, by whom (their credentials) and whether it relied on abstracts or full texts (including translations) of articles. The report should emphasize clinical data, but new findings in preclinical reports of potential significance should also be described. Should any report or finding be judged important, a copy (translated as required) should be submitted for our review.

5.

6.

7. Cardiovascular Risk

Since the initial findings of sudden unexpected deaths in the 1 year chronic dog toxicology study, there have been concerns about the cardiovascular risks posed by citalopram (CT) to humans. We are mindful that you have conducted studies in both animals and in humans to evaluate the potential for this risk and that these findings provide some reassurance. Unfortunately, some residual concerns persist.

Despite your efforts, the precise mechanism(s) causing sudden death in dogs remains unknown. While didemethylcitalopram (DDCT) may have an important role, other factors might be important. For example, the possible role of another metabolite of citalopram, demethylcitalopram (DCT), has not been directly examined.

At doses in the range recommended, QT interval prolongation of a clinically important degree has been observed infrequently; such changes were not observed in studies 91206 and 92104, for example. On the other hand, there are some signals of concern. In the pool of 5 placebo controlled studies, 1.1% of the 797 patients exposed to citalopram compared to only 0.4% of the 241 patients randomized to placebo developed a QTc interval exceeding 500 msec.

Labeling will need to describe the findings of the 1 year oral and acute intravenous dog toxicology studies and the uncertainties involved with their interpretation *vis a vis* human risk (see our draft labeling proposal). Whether further information concerning citalopram's effects on cardiovascular function and cardiac repolarization needs to be developed and the specific information that would be useful bears further discussion.

Potentially critical, for example, is a better understanding of the extent to which DCT and DDCT plasma concentrations may vary under recommended dosing and conditions of actual use. If DDCT is, as you suspect, a potent QT prolonger, it will be important to learn whether there are subgroups in the population that regularly develop higher levels of this metabolite. Similarly, it may be useful to assess the *in vitro* effects of CT, DCT, and DDCT on the potassium channel currents.

Given the many issues to be resolved and the many approaches that might be used to

resolve them, we believe it best to meet to discuss available options. In advance of such a meeting, it would be helpful for you to submit a plan describing your proposed approach to dealing with these issues.

8. Valvulopathy

Recent concern about pulmonary hypertension and/or valvulopathy in association with the drugs dexfenfluramine and d,l-fenfluramine has raised questions about other drugs that affect serotonergic function. In order to assist us in following up this issue, we ask that you respond to the following questions:

- Are you aware of data from your clinical trials, from spontaneous reports, or from any other source that are suggestive of an association between the use of citalopram and pulmonary hypertension and/or valvulopathy? If so, please submit a report on any such findings as well as the extent of information that bears on the question.
- Do you have possession of, or are you aware of, any echocardiographic data for patients exposed to citalopram? If so, please submit a report on any such findings. In addition, please comment on any plans you may have to obtain echocardiographic data on patients exposed to citalopram.
- Do you have possession of, or are you aware of, any gross anatomic or microscopic evidence from diseased cardiac valves from patients exposed to citalopram? If so, please submit a report on any such findings. In addition, please comment on any plans you may have to obtain anatomic data on patients exposed to citalopram. Sections may be available for deceased patients with valvular lesions for whom autopsies were performed or for patients requiring valve replacements.

9.

10. Maximum Recommended Dose/Highest Tablet Strength

Given the failure of study 91206 to distinguish between the 40 and 60 mg doses, we do not feel there is a basis for recommending doses above 40 mg/day. Consequently, we have changed the maximum recommended dose in Dosage and Administration to 40 mg/day, and we have deleted all references to the 60 mg dosage strength from labeling.

**MANUFACTURING AND CONTROLS**1. Expiration Date

Please note that, at this time, we are only approving an 18-month expiry date for the drug product.

2. Nomenclature

The established name, citalopram hydrobromide, has not been adopted by the USAN Council. We request that you submit this name to the council expeditiously.

**BIOPHARMACEUTICS**

1. Considering that carbamazepine (CBZ) is a hepatic enzyme inducer, the pharmacokinetics of citalopram would be expected to be altered during the continuous administration of CBZ. Although this expectation is not supported by the trough citalopram plasma concentration observed over time, the plasma concentrations of citalopram metabolites, demethylcitalopram and didemethylcitalopram, almost doubled between day 26 and 36.
2. Given that the pharmacokinetic profiles of citalopram isomers are different, it is necessary that you provide data from a dose proportionality study on the individual isomers, in accordance with FDA's policy statement for the development of new stereoisomeric drugs (May 1992). We note that you have already conducted a dose proportionality study showing that citalopram is linear over the dose range of 10 to 60 mg.
3. In order to characterize the dissolution profile of citalopram, we note that you have taken samples from numerous batches at 10, 15, 20 and 30 minutes. The samples have not been

taken consistently, i.e. different batches have different sampling times. For example, if a sample was taken at 10 minutes, then 15 minutes sample has not been taken but a 20 minutes sample was collected. Please provide a rationale for this sampling scheme.

4. In Vivo Drug Interaction Study

5. Dissolution Specification

We ask that you agree to the following recommendation by the Office of Clinical Pharmacology and Biopharmaceutics for a dissolution method and specification for all tablet strengths:

Apparatus:	USP Apparatus 1 (basket)
Speed:	100 RPM
Medium:	(pH = 1.5) at 37°C
Specification:	Q = at 30 minutes

Please submit twenty copies of the printed labeling, ten of which are individually mounted on heavy weight paper or similar material.

In addition, please submit three copies of the introductory promotional material that you propose to use for this product. All proposed materials should be submitted in draft or mock-up form, not final print. Please submit one copy to this Division and two copies of both the promotional material and the package insert directly to:

Food and Drug Administration  
Division of Drug Marketing, Advertising and Communications,  
HFD-40  
5600 Fishers Lane  
Rockville, Maryland 20857

Within 10 days after the date of this letter, you are required to amend the application, notify us of your intent to file an amendment, or follow one of your other options under 21 CFR 314.110. In the absence of such action FDA may take action to withdraw the application. Should you have any questions, please contact:

Paul David, R.Ph.  
Project Manager  
Telephone: (301) 594-5530

The drug may not be legally marketed until you have been notified in writing that the application is approved.

Sincerely yours,

/S/

5/12/98

Robert Temple, M.D.  
Director  
Office of Drug Evaluation I  
Center for Drug Evaluation and Research

ATTACHMENT

**CENTER FOR DRUG EVALUATION AND RESEARCH**

**APPLICATION NUMBER: 20822**

**FINAL PRINTED LABELING**

**ATTACHMENT**  
**FINAL LABELING**

**Celexa™**  
**(citalopram hydrobromide)**

APPEARS THIS WAY  
ON ORIGINAL

**DESCRIPTION**

Celexa™ (citalopram HBr) is an orally administered selective serotonin reuptake inhibitor (SSRI) with a chemical structure unrelated to that of other SSRI's or of tricyclic, tetracyclic, or other available antidepressant agents. Citalopram HBr is a racemic bicyclic pthalane derivative designated (±)-1-(3-dimethylaminopropyl)-1-(4-fluorophenyl)-1,3-dihydroisobenzofuran-5-carbonitrile, HBr with the following structural formula:

[Structural formula here]

APPEARS THIS WAY  
ON ORIGINAL

The molecular formula is C<sub>20</sub>H<sub>22</sub>BrFN<sub>2</sub>O and its molecular weight is 405.35.

Citalopram HBr occurs as a fine white to off-white powder. Citalopram HBr is sparingly soluble in water and soluble in ethanol.

Celexa™ is a film coated, oval, scored tablet containing citalopram HBr in strengths equivalent to 20 mg or 40 mg citalopram base.

Inactive Ingredients: Copolyvidone, Corn Starch, Crosscarmellose Sodium, Glycerin, Lactose Monohydrate, Magnesium Stearate, Hydroxypropyl Methyl Cellulose, Microcrystalline Cellulose, Polyethylene Glycol. Titanium Dioxide and Iron Oxides are used as coloring agents in the pink (20 mg) tablets.

**CLINICAL PHARMACOLOGY**

APPEARS THIS WAY  
ON ORIGINAL

**Pharmacodynamics**

The mechanism of action of citalopram HBr as an antidepressant is presumed to be linked to potentiation of serotonergic activity in the central nervous system resulting from its inhibition of CNS neuronal reuptake of serotonin (5-HT). *In vitro* and *in vivo* studies in animals suggest that citalopram is a highly selective serotonin reuptake inhibitor (SSRI) with minimal effects on norepinephrine (NE) and dopamine (DA) neuronal reuptake. Tolerance to the inhibition of 5-HT uptake is not induced by

long term (14 day) treatment of rats with citalopram. Citalopram is a racemic mixture (50/50), and the inhibition of 5-HT reuptake by citalopram is primarily due to the (S)-enantiomer. Citalopram has no or very low affinity for 5-HT<sub>1A</sub>, 5-HT<sub>2A</sub>, dopamine D<sub>1</sub> and D<sub>2</sub>,  $\alpha_1$ -,  $\alpha_2$ -, and  $\beta$ -adrenergic, histamine H<sub>1</sub>, gamma aminobutyric acid (GABA), muscarinic cholinergic, and benzodiazepine receptors. Antagonism of muscarinic, histaminergic and adrenergic receptors has been hypothesized to be associated with various anticholinergic, sedative and cardiovascular effects of other psychotropic drugs.

### **Pharmacokinetics**

The single- and multiple-dose pharmacokinetics of citalopram are linear and dose-proportional in a dose range of 10-60 mg/day. Biotransformation of citalopram is mainly hepatic, with a mean terminal half-life of about 35 hours. With once daily dosing, steady state plasma concentrations are achieved within approximately one week. At steady state, the extent of accumulation of citalopram in plasma, based on the half-life, is expected to be 2.5 times the plasma concentrations observed after a single dose.

### **Absorption and Distribution**

Following a single oral dose (40 mg tablet) of citalopram, peak blood levels occur at about 4 hours. The absolute bioavailability of citalopram was about 80% relative to an intravenous dose and absorption is not affected by food. The volume of distribution of citalopram is about 12 L/kg and the binding of citalopram (CT), demethylcitalopram (DCT) and didemethylcitalopram (DDCT) to human plasma proteins is about 80%.

### **Metabolism and Elimination**

Following intravenous administrations of citalopram, the fraction of drug recovered in the urine as citalopram and DCT was about 10% and 5%, respectively. The systemic clearance of citalopram was 330 mL/min, with approximately 20% of that due to renal clearance.

Citalopram is metabolized to demethylcitalopram (DCT), didemethylcitalopram (DDCT), citalopram-N-oxide and a deaminated propionic acid derivative. In humans, unchanged citalopram is the predominant compound in plasma. At steady state, the concentrations of citalopram's metabolites, DCT and DDCT, in plasma are approximately one-half and one-tenth, respectively, that of the parent drug. *In vitro* studies show that citalopram is at least 8 times more potent than its metabolites in the inhibition of serotonin reuptake, suggesting that the metabolites evaluated do not likely contribute significantly to the antidepressant actions of citalopram.

*In vitro* studies using human liver microsomes indicated that CYP3A4 and CYP2C19 are the primary isozymes involved in the N-demethylation of citalopram.

### **Population Subgroups**

Age - Citalopram pharmacokinetics in subjects  $\geq 60$  years of age were compared to younger subjects in two normal volunteer studies. In a single dose study, citalopram AUC and half-life were increased in the elderly subjects by 30% and 50%, respectively, whereas in a multiple dose study they were increased by 23% and 30% respectively. 20 mg is the recommended dose for most elderly patients (see Dosage and Administration).

Gender - In three pharmacokinetic studies (total N=32), citalopram AUC in women was one and a half to two times that in men. This difference was not observed in five other pharmacokinetic studies (total N=114). In clinical studies, no differences in steady state serum citalopram levels were seen between men (N=237) and women (N=388). There were no gender differences in the pharmacokinetics of DCT and DDCT. No adjustment of dosage on the basis of gender is recommended.

Reduced hepatic function - Citalopram oral clearance was reduced by 37% and half-life was doubled in patients with reduced hepatic function compared to normal subjects. 20 mg is the recommended dose for most hepatically impaired patients (see Dosage and Administration).

Reduced renal function - In patients with mild to moderate renal function impairment, oral clearance of citalopram was reduced by 17% compared to normal subjects. No adjustment of dosage for such patients is recommended. No information is available about the pharmacokinetics of citalopram in patients with severely reduced renal function (creatinine clearance  $< 20$  ml/min).

### Drug-Drug Interactions

*In vitro* enzyme inhibition data did not reveal an inhibitory effect of citalopram on CYP3A4, but did suggest that it is a weak inhibitor of CYP-1A2, -2D6, and -2C19. Citalopram would be expected to have little inhibitory effect on *in vivo* metabolism mediated by these cytochromes. However, *in vivo* data to address this question are very limited.

Since CYP3A4 and 2C19 are the primary enzymes involved in the metabolism of citalopram, it is expected that potent inhibitors of 3A4, e.g., ketoconazole, itraconazole, and macrolide antibiotics, and potent inhibitors of CYP2C19, e.g., omeprazole, might decrease the clearance of citalopram. Citalopram steady state levels were not significantly different in poor metabolizers and extensive 2D6 metabolizers after multiple dose administration of Celexa<sup>TM</sup>, suggesting that coadministration, with Celexa<sup>TM</sup>, of a drug that inhibits CYP2D6, is unlikely to have clinically significant effects on citalopram metabolism. See Drug Interactions under Precautions for more detailed information on available drug interaction data.

### **Clinical Efficacy Trials**

The efficacy of Celexa<sup>TM</sup> as a treatment for depression was established in two placebo-controlled studies (of 4 to 6 weeks in duration) in adult outpatients (ages 18-66) meeting DSM-III or DSM-III-R criteria for major depression. Study 1, a 6-week trial in which patients received fixed Celexa<sup>TM</sup>

doses of 10, 20, 40, and 60 mg/day, showed that Celexa™ at doses of 40 and 60 mg/day was effective as measured by the Hamilton Depression Rating Scale (HAMD) total score, the HAMD depressed mood item (Item 1), the Montgomery Asberg Depression Rating Scale, and the Clinical Global Impression (CGI) Severity scale. This study showed no clear effect of the 10 and 20 mg/day doses, and the 60 mg/day dose was not more effective than the 40 mg/day dose. In study 2, a 4-week, placebo-controlled trial in depressed patients, of whom 85% met criteria for melancholia, the initial dose was 20 mg/day, followed by titration to the maximum tolerated dose or a maximum dose of 80 mg/day. Patients treated with Celexa™ showed significantly greater improvement than placebo patients on the HAMD total score, HAMD item 1, and the CGI Severity score. In three additional placebo-controlled depression trials, the difference in response to treatment between patients receiving Celexa™ and patients receiving placebo was not statistically significant, possibly due to high spontaneous response rate, smaller sample size, or, in the case of one study, too low a dose.

In two long-term studies, depressed patients who had responded to Celexa™ during an initial 6 or 8 weeks of acute treatment (fixed doses of 20 or 40 mg/day in one study and flexible doses of 20-60 mg/day in the second study) were randomized to continuation of Celexa™ or to placebo. In both studies, patients receiving continued Celexa™ treatment experienced significantly lower relapse rates over the subsequent 6 months compared to those receiving placebo. In the fixed dose study, the decreased rate of depression relapse was similar in patients receiving 20 or 40 mg/day of Celexa™.

Analyses of the relationship between treatment outcome and age, gender, and race did not suggest any differential responsiveness on the basis of these patient characteristics.

#### Comparison of Clinical Trial Results

Highly variable results have been seen in the clinical development of all antidepressant drugs. Furthermore, in those circumstances when the drugs have not been studied in the same controlled clinical trial(s), comparisons among the results of studies evaluating the effectiveness of different antidepressant drug products are inherently unreliable. Because conditions of testing (e.g., patient samples, investigators, doses of the treatments administered and compared, outcome measures, etc.) vary among trials, it is virtually impossible to distinguish a difference in drug effect from a difference due to one of the confounding factors just enumerated.

#### **INDICATIONS AND USAGE**

Celexa™ (citalopram HBr) is indicated for the treatment of depression.

The efficacy of Celexa™ in the treatment of depression was established in 4-6 week controlled trials of outpatients whose diagnosis corresponded most closely to the DSM-III and DSM-III-R category of major depressive disorder (see Clinical Pharmacology).

A major depressive episode (DSM-IV) implies a prominent and relatively persistent (nearly every day for at least 2 weeks) depressed or dysphoric mood that usually interferes with daily functioning, and

includes at least five of the following nine symptoms: depressed mood, loss of interest in usual activities, significant change in weight and/or appetite, insomnia or hypersomnia, psychomotor agitation or retardation, increased fatigue, feelings of guilt or worthlessness, slowed thinking or impaired concentration, a suicide attempt or suicidal ideation.

The antidepressant action of Celexa™ in hospitalized depressed patients has not been adequately studied.

The efficacy of Celexa™ in maintaining an antidepressant response for up to 24 weeks following 6 to 8 weeks of acute treatment was demonstrated in two placebo-controlled trials (see Clinical Pharmacology). Nevertheless, the physician who elects to use Celexa™ for extended periods should periodically re-evaluate the long-term usefulness of the drug for the individual patient.

## **CONTRAINDICATIONS**

Concomitant use in patients taking monoamine oxidase inhibitors (MAOI's) is contraindicated (see Warnings).

Celexa™ is contraindicated in patients with a hypersensitivity to citalopram or any of the inactive ingredients in Celexa™.

## **WARNINGS**

### **Potential for Interaction with Monoamine Oxidase Inhibitors**

**In patients receiving serotonin reuptake inhibitor drugs in combination with a monoamine oxidase inhibitor (MAOI), there have been reports of serious, sometimes fatal, reactions including hyperthermia, rigidity, myoclonus, autonomic instability with possible rapid fluctuations of vital signs, and mental status changes that include extreme agitation progressing to delirium and coma. These reactions have also been reported in patients who have recently discontinued SSRI treatment and have been started on a MAOI. Some cases presented with features resembling neuroleptic malignant syndrome. Furthermore, limited animal data on the effects of combined use of SSRI's and MAOI'S suggest that these drugs may act synergistically to elevate blood pressure and evoke behavioral excitation. Therefore, it is recommended that Celexa™ should not be used in combination with a MAOI, or within 14 days of discontinuing treatment with a MAOI. Similarly, at least 14 days should be allowed after stopping Celexa™ before starting a MAOI.**

## **PRECAUTIONS**

### **General**

### Hyponatremia

Several cases of hyponatremia and SIADH (syndrome of inappropriate antidiuretic hormone secretion) have been reported in association with Celexa™ treatment. All patients with these events have recovered with discontinuation of Celexa™ and/or medical intervention.

### Activation of Mania/Hypomania

In placebo-controlled trials of Celexa™, some of which included patients with bipolar disorder, activation of mania/hypomania was reported in 0.2% of 1063 patients treated with Celexa™ and in none of the 446 patients treated with placebo. Activation of mania/hypomania has also been reported in a small proportion of patients with major affective disorders treated with other marketed antidepressants. As with all antidepressants, Celexa™ should be used cautiously in patients with a history of mania.

### Seizures

Although anticonvulsant effects of citalopram have been observed in animal studies, Celexa™ has not been systematically evaluated in patients with a seizure disorder. These patients were excluded from clinical studies during the product's premarketing testing. In clinical trials of Celexa™, seizures occurred in 0.3% of patients treated with Celexa™ (a rate of one patient per 98 years of exposure) and 0.5% of patients treated with placebo (a rate of one patient per 50 years of exposure). Like other antidepressants, Celexa™ should be introduced with care in patients with a history of seizure disorder.

### Suicide

The possibility of a suicide attempt is inherent in depression and may persist until significant remission occurs. Close supervision of high risk patients should accompany initial drug therapy. Prescriptions for Celexa™ should be written for the smallest quantity of tablets consistent with good patient management, in order to reduce the risk of overdose.

### Interference with Cognitive and Motor Performance

In studies in normal volunteers, Celexa™ in doses of 40 mg/day did not produce impairment of intellectual function or psychomotor performance. Because any psychoactive drug may impair judgement, thinking, or motor skills, however, patients should be cautioned about operating hazardous machinery, including automobiles, until they are reasonably certain that Celexa™ therapy does not affect their ability to engage in such activities.

### Use in Patients with Concomitant Illness

APPEARS THIS WAY  
ON ORIGINAL

Clinical experience with Celexa™ in patients with certain concomitant systemic illnesses is limited. Caution is advisable in using Celexa™ in patients with diseases or conditions that produce altered metabolism or hemodynamic responses.

Celexa™ has not been systematically evaluated in patients with a recent history of myocardial infarction or unstable heart disease. Patients with these diagnoses were generally excluded from clinical studies during the product's premarketing testing. However, the electrocardiograms of 1116 patients who received Celexa™ in clinical trials were evaluated and the data indicate that Celexa™ is not associated with the development of clinically significant ECG abnormalities.

In subjects with hepatic impairment, citalopram clearance was decreased and plasma concentrations were increased. The use of Celexa™ in hepatically impaired patients should be approached with caution and a lower maximum dosage is recommended (see Dosage and Administration).

Because citalopram is extensively metabolized, excretion of unchanged drug in urine is a minor route of elimination. Until adequate numbers of patients with severe renal impairment have been evaluated during chronic treatment with Celexa™, however, it should be used with caution in such patients (see Dosage and Administration).

### **Information for Patients**

Physicians are advised to discuss the following issues with patients for whom they prescribe Celexa™:

Although in controlled studies Celexa™ has not been shown to impair psychomotor performance, any psychoactive drug may impair judgment, thinking or motor skills, and so patients should be cautioned about operating hazardous machinery, including automobiles, until they are reasonably certain that Celexa™ therapy does not affect their ability to engage in such activities.

Patients should be told that, although Celexa™ has not been shown in experiments with normal subjects to increase the mental and motor skill impairments caused by alcohol, the concomitant use of Celexa™ and alcohol in depressed patients is not advised.

Patients should be advised to inform their physician if they are taking, or plan to take, any prescription or over-the-counter drugs, as there is a potential for interactions.

Patients should be advised to notify their physician if they become pregnant or intend to become pregnant during therapy.

Patients should be advised to notify their physician if they are breast feeding an infant.

While patients may notice improvement with Celexa™ therapy in 1 to 4 weeks, they should be advised to continue therapy as directed.

### **Laboratory Tests**

There are no specific laboratory tests recommended.

### **Drug Interactions**

**CNS Drugs** - Given the primary CNS effects of citalopram, caution should be used when it is taken in combination with other centrally acting drugs.

**Alcohol** - Although citalopram did not potentiate the cognitive and motor effects of alcohol in a clinical trial, as with other psychotropic medications, the use of alcohol by depressed patients taking Celexa™ is not recommended.

**Monoamine Oxidase Inhibitors (MAOI's)** - See Contraindications and Warnings.

**Cimetidine** - In subjects who had received 21 days of 40 mg/day Celexa™, combined administration of 400 mg/day cimetidine for 8 days resulted in an increase in citalopram AUC and C<sub>max</sub> of 43% and 39%, respectively. The clinical significance of these findings is unknown.

**Digoxin** - In subjects who had received 21 days of 40 mg/day Celexa™, combined administration of Celexa™ and digoxin (single dose of 1 mg) did not significantly affect the pharmacokinetics of either citalopram or digoxin.

**Lithium** - Coadministration of Celexa™ (40 mg/day for 10 days) and lithium (30 mmol/day for 5 days) had no significant effect on the pharmacokinetics of citalopram or lithium. Nevertheless, plasma lithium levels should be monitored with appropriate adjustment to the lithium dose in accordance with standard clinical practice. Because lithium may enhance the serotonergic effects of citalopram, caution should be exercised when Celexa™ and lithium are coadministered.

**Warfarin** - Administration of 40 mg/day Celexa™ for 21 days did not affect the pharmacokinetics of warfarin, a CYP3A4 substrate. Prothrombin time was increased by 5%, the clinical significance of which is unknown.

**Carbamazepine** - Combined administration of Celexa™ (40 mg/day for 14 days) and carbamazepine (titrated to 400 mg/day for 35 days) did not significantly affect the pharmacokinetics of carbamazepine, a CYP3A4 substrate. Although trough citalopram plasma levels were unaffected, given the enzyme inducing properties of carbamazepine, the possibility that carbamazepine might increase the clearance of citalopram should be considered if the two drugs are coadministered.

**CYP3A4 and 2C19 Inhibitors** - In vitro studies indicated that CYP3A4 and 2C19 are the primary enzymes involved in the metabolism of citalopram. As data are not available from clinical pharmacokinetic studies, the possibility that the clearance of citalopram will be decreased when citalopram is administered with a potent inhibitor of CYP3A4 (e.g., ketoconazole, itraconazole, fluconazole, or erythromycin), or a potent inhibitor of CYP2C19 (e.g., omeprazole), should be considered.

Metoprolol - Administration of 40 mg/day Celexa™ for 22 days resulted in a two-fold increase in the plasma levels of the beta-adrenergic blocker metoprolol. Increased metoprolol plasma levels have been associated with decreased cardioselectivity. Coadministration of Celexa™ and metoprolol had no clinically significant effects on blood pressure or heart rate.

Imipramine and Other Tricyclic Antidepressants (TCAs) - *In vitro* studies suggest that citalopram is a relatively weak inhibitor of CYP2D6. Coadministration of Celexa™ (40 mg/day for 10 days) with the tricyclic antidepressant imipramine (single dose of 100 mg), a substrate for CYP2D6, did not significantly affect the plasma concentrations of imipramine or citalopram. However, the concentration of the imipramine metabolite desipramine was increased by approximately 50%. The clinical significance of the desipramine change is unknown. Nevertheless, caution is indicated in the coadministration of TCAs with Celexa™.

Electroconvulsive Therapy (ECT) - There are no clinical studies of the combined use of electroconvulsive therapy (ECT) and Celexa™.

## **Carcinogenesis, Mutagenesis, Impairment of Fertility**

### Carcinogenesis

Citalopram was administered in the diet to NMRI/BOM strain mice and COBS WI strain rats for 18 and 24 months, respectively. There was no evidence for carcinogenicity of citalopram in mice receiving up to 240 mg/kg/day, which is equivalent to 20 times the maximum recommended human daily dose (MRHD) of 60 mg on a surface area (mg/m<sup>2</sup>) basis. There was an increased incidence of small intestine carcinoma in rats receiving 8 or 24 mg/kg/day, doses which are approximately 1.3 and 4 times the MRHD, respectively, on a mg/m<sup>2</sup> basis. A no-effect dose for this finding was not established. The relevance of these findings to humans is unknown.

### Mutagenesis

Citalopram was mutagenic in the *in vitro* bacterial reverse mutation assay (Ames test) in 2 of 5 bacterial strains (Salmonella TA98 and TA1537) in the absence of metabolic activation. It was clastogenic in the *in vitro* Chinese hamster lung cell assay for chromosomal aberrations in the presence and absence of metabolic activation. Citalopram was not mutagenic in the *in vitro* mammalian forward gene mutation assay (HPRT) in mouse lymphoma cells or in a coupled *in vitro/in vivo* unscheduled DNA synthesis (UDS) assay in rat liver. It was not clastogenic in the *in vitro* chromosomal aberration assay in human lymphocytes or in two *in vivo* mouse micronucleus assays.

### Impairment of Fertility

When citalopram was administered orally to male and female rats prior to and throughout mating and gestation at doses of 16/24 (males/females), 32, 48, and 72 mg/kg/day, mating was decreased at all doses, and fertility was decreased at doses  $\geq 32$  mg/kg/day, approximately 5 times the maximum

recommended human dose (MRHD) of 60 mg/day on a body surface area ( $\text{mg}/\text{m}^2$ ) basis. Gestation duration was increased at 48 mg/kg/day, approximately 8 times the MRHD.

## **Pregnancy**

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### Pregnancy Category C

In animal reproduction studies, 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 citalopram (32, 56, or 112 mg/kg/day) to pregnant animals during the period of organogenesis resulted in decreased embryo/fetal growth and survival and an increased incidence of fetal abnormalities (including cardiovascular and skeletal defects) at the high dose, which is approximately 18 times the maximum recommended human dose (MRHD) of 60 mg/day on a body surface area ( $\text{mg}/\text{m}^2$ ) basis. This dose was also associated with maternal toxicity (clinical signs, decreased BW gain). The developmental no effect dose of 56 mg/kg/day is approximately 9 times the MRHD on a  $\text{mg}/\text{m}^2$  basis. In a rabbit study, no adverse effects on embryo/fetal development were observed at doses of up to 16 mg/kg/day, or approximately 5 times the MRHD on  $\text{mg}/\text{m}^2$  basis. Thus, teratogenic effects were observed at a maternally toxic dose in the rat and were not observed in the rabbit.

When female rats were treated with 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, which is approximately 5 times the MRHD on a  $\text{mg}/\text{m}^2$  basis. The no effect dose of 12.8 mg/kg/day is approximately 2 times the MRHD on a  $\text{mg}/\text{m}^2$  basis. Similar effects on offspring mortality and growth were seen when dams were treated throughout gestation and early lactation at doses  $\geq 24$  mg/kg/day, approximately 4 times the MRHD on a  $\text{mg}/\text{m}^2$  basis. A no effect dose was not determined in that study.

There are no adequate and well-controlled studies in pregnant women; therefore, citalopram should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus.

## **Labor and Delivery**

The effect of Celexa<sup>TM</sup> on labor and delivery in humans is unknown.

## **Nursing Mothers**

As has been found to occur with many other drugs, citalopram is excreted in human breast milk. There have been two reports of infants experiencing excessive somnolence, decreased feeding, and weight loss in association with breast feeding from a citalopram-treated mother; in one case, the infant was reported to recover completely upon discontinuation of citalopram by its mother. The decision

whether to continue or discontinue either nursing or Celexa™ therapy should take into account the risks of citalopram exposure for the infant and the benefits of Celexa™ treatment for the mother.

### **Pediatric Use**

Safety and effectiveness in pediatric patients have not been established.

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### **Geriatric Use**

Of 4422 patients in clinical studies of Celexa™, 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 greater sensitivity of some older individuals cannot be ruled out. Most elderly patients treated with Celexa™ in clinical trials received daily doses between 20 and 40 mg (see Dosage and Administration).

In two pharmacokinetic studies, citalopram AUC was increased by 23% and 30%, respectively, in elderly subjects as compared to younger subjects, and its half-life was increased by 30% and 50%, respectively. (see Clinical Pharmacology).

20 mg/day is the recommended dose for most elderly patients (see Dosage and Administration).

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### **ADVERSE REACTIONS**

The premarketing development program for Celexa™ included citalopram exposures in patients and/or normal subjects from 3 different groups of studies: 429 normal subjects in clinical pharmacology/pharmacokinetic studies; 4422 exposures from patients in controlled and uncontrolled clinical trials, corresponding to approximately 1370 patient exposure years. There were, in addition, over 19,000 exposures from mostly open-label, European postmarketing studies. The conditions and duration of treatment with Celexa™ varied greatly and included (in overlapping categories) open-label and double-blind studies, inpatient and outpatient studies, fixed-dose and dose-titration studies, and short-term and long-term exposure. Adverse reactions were assessed by collecting adverse events, results of physical examinations, vital signs, weights, laboratory analyses, ECGs, and results of ophthalmologic examinations.

Adverse events during exposure were obtained primarily by general inquiry and recorded by clinical investigators using terminology of their own choosing. Consequently, it is not possible to provide a meaningful estimate of the proportion of individuals experiencing adverse events without first grouping similar types of events into a smaller number of standardized event categories. In the tables and tabulations that follow, standard World Health Organization (WHO) terminology has been used to classify reported adverse events.

The stated frequencies of adverse events represent the proportion of individuals who experienced, at least once, a treatment-emergent adverse event of the type listed. An event was considered treatment emergent if it occurred for the first time or worsened while receiving therapy following baseline evaluation.

## **Adverse Findings Observed in Short-Term, Placebo-Controlled Trials**

### Adverse Events Associated with Discontinuation of Treatment

Among 1063 depressed patients who received Celexa™ at doses ranging from 10 to 80 mg/day in placebo-controlled trials of up to 6 weeks in duration, 16% discontinued treatment due to an adverse event, as compared to 8% of 446 patients receiving placebo. The adverse events associated with discontinuation and considered drug-related (i.e., associated with discontinuation in at least 1% of Celexa™-treated patients at a rate at least twice that of placebo) are shown in Table 1. It should be noted that one patient can report more than one reason for discontinuation and be counted more than once in this table.

**TABLE 1**  
**Adverse Events Associated with Discontinuation of Treatment in Short-Term, Placebo-Controlled, Depression Trials**

**Percentage of Patients Discontinuing Due to Adverse Event**

	<b>Citalopram (n=1063)</b>	<b>Placebo (n=446)</b>
<b>Body System/Adverse Event</b>		
<b>General</b>		
Asthenia	1%	<1%
<b>Gastrointestinal Disorders</b>		
Nausea	4%	0%
Dry Mouth	1%	<1%
Vomiting	1%	0%
<b>Central and Peripheral Nervous System Disorders</b>		
Dizziness	2%	<1%
<b>Psychiatric Disorders</b>		

Insomnia	3%	1%
Somnolence	2%	1%
Agitation	1%	<1%

Adverse Events Occurring at an Incidence of 2% or More Among Celexa™-Treated Patients

Table 2 enumerates the incidence, rounded to the nearest percent, of treatment emergent adverse events that occurred among 1063 depressed patients who received Celexa™ at doses ranging from 10 to 80 mg/day in placebo-controlled trials of up to 6 weeks in duration. Events included are those occurring in 2% or more of patients treated with Celexa™ and for which the incidence in patients treated with Celexa™ was greater than the incidence in placebo-treated patients.

The prescriber should be aware that these figures cannot be used to predict the incidence of adverse events in the course of usual medical practice where patient characteristics and other factors differ from those which prevailed in the clinical trials. Similarly, the cited frequencies cannot be compared with figures obtained from other clinical investigations involving different treatments, uses, and investigators. The cited figures, however, do provide the prescribing physician with some basis for estimating the relative contribution of drug and non-drug factors to the adverse event incidence rate in the population studied.

The only commonly observed adverse event that occurred in Celexa™ patients with an incidence of 5% or greater and at least twice the incidence in placebo patients was ejaculation disorder (primarily ejaculatory delay) in male patients (see Table 2).

**Table 2**

**Treatment-Emergent Adverse Events:  
Incidence in Placebo-Controlled Clinical Trials\***

**(Percentage of Patients Reporting Event)**

<b><u>Body System/Adverse Event</u></b>	<b><u>Celexa™</u> (N=1063)</b>	<b><u>Placebo</u> (N=446)</b>
<b>Autonomic Nervous System Disorders</b>		
Dry Mouth	20%	14%
Sweating Increased	11%	9%
<b>Central &amp; Peripheral Nervous System Disorders</b>		
Tremor	8%	6%
<b>Gastrointestinal Disorders</b>		
Nausea	21%	14%
Diarrhea	8%	5%
Dyspepsia	5%	4%
Vomiting	4%	3%

Abdominal Pain	3%	2%
<b>General</b>		
Fatigue	5%	3%
Fever	2%	<1%
<b>Musculoskeletal System Disorders</b>		
Arthralgia	2%	1%
Myalgia	2%	1%
<b>Psychiatric Disorders</b>		
Somnolence	18%	10%
Insomnia	15%	14%
Anxiety	4%	3%
Anorexia	4%	2%
Agitation	3%	1%
Dysmenorrhea <sup>1</sup>	3%	2%
Libido Decreased	2%	<1%
Yawning	2%	<1%
<b>Respiratory System Disorders</b>		
Upper Respiratory Tract Infection	5%	4%
Rhinitis	5%	3%
Sinusitis	3%	<1%
<b>Urogenital</b>		
Ejaculation Disorder <sup>2,3</sup>	6%	1%
Impotence <sup>3</sup>	3%	<1%

\*Events reported by at least 2% of patients treated with Celexa™ are reported, except for the following events which had an incidence on placebo  $\geq$  Celexa™ : headache, asthenia, dizziness, constipation, palpitation, vision abnormal, sleep disorder, nervousness, pharyngitis, micturition disorder, back pain.

<sup>1</sup>Denominator used was for females only (N=638 Celexa™; N=252 placebo).

<sup>2</sup>Primarily ejaculatory delay.

<sup>3</sup>Denominator used was for males only (N=425 Celexa™; N=194 placebo).

#### Dose Dependency of Adverse Events

The potential relationship between the dose of Celexa™ administered and the incidence of adverse events was examined in a fixed dose study in depressed patients receiving placebo or Celexa™ 10, 20, 40, and 60 mg. Jonckheere's trend test revealed a positive dose response ( $p < 0.05$ ) for the following adverse events: fatigue, impotence, insomnia, sweating increased, somnolence, and yawning.

#### Male and Female Sexual Dysfunction with SSRIs

While sexual dysfunction is often part of depression and other psychiatric disorders, there is increasing evidence that treatment with selective serotonin reuptake inhibitors (SSRIs) may induce

sexual side effects. This is a difficult area to study because patients may not spontaneously report symptoms of this nature, and therefore, it is thought that sexual side effects with the SSRIs may be underestimated. In placebo-controlled clinical trials (Table 2), the reported incidence of decreased libido, ejaculation disorder (primarily ejaculatory delay), and impotence in male depressed patients receiving Celexa™ (n=425) was 3.8%, 6.1%, and 2.8%, respectively. In female depressed patients receiving Celexa™ (n=638), the reported incidence of decreased libido and anorgasmia was 1.3% and 1.1%, respectively. The reported incidence of each of these adverse events was  $\leq 1.0\%$  among male and female depressed patients receiving placebo.

While it is difficult to know the precise risk of sexual dysfunction associated with the use of SSRIs, physicians should routinely inquire about such possible side effects.

### Vital Sign Changes

Celexa™ and placebo groups were compared with respect to (1) mean change from baseline in vital signs (pulse, systolic blood pressure, and diastolic blood pressure) and (2) the incidence of patients meeting criteria for potentially clinically significant changes from baseline in these variables. These analyses did not reveal any clinically important changes in vital signs associated with Celexa™ treatment. In addition, a comparison of supine and standing vital sign measures for Celexa™ and placebo treatments indicated that Celexa™ treatment is not associated with orthostatic changes.

### Weight Changes

Patients treated with Celexa™ in controlled trials experienced a weight loss of about 0.5 kg compared to no change for placebo patients.

### Laboratory Changes

Celexa™ and placebo groups were compared with respect to (1) mean change from baseline in various serum chemistry, hematology, and urinalysis variables and (2) the incidence of patients meeting criteria for potentially clinically significant changes from baseline in these variables. These analyses revealed no clinically important changes in laboratory test parameters associated with Celexa™ treatment.

### ECG Changes

Electrocardiograms from Celexa™ (n=802) and placebo (n=241) groups were compared with respect to (1) mean change from baseline in various ECG parameters and (2) the incidence of patients meeting criteria for potentially clinically significant changes from baseline in these variables. The only statistically significant drug-placebo difference observed was a decrease in heart rate for Celexa™ of 1.7 bpm compared to no change in heart rate for placebo. There were no observed differences in QT or other ECG intervals.

## Other Events Observed During the Premarketing Evaluation of Celexa™ (citalopram HBr)

Following is a list of WHO terms that reflect treatment-emergent adverse events, as defined in the introduction to the ADVERSE REACTIONS section, reported by patients treated with Celexa™ at multiple doses in a range of 10 to 80 mg/day during any phase of a trial within the premarketing database of 4422 patients. All reported events are included except those already listed in Table 2 or elsewhere in labeling, those events for which a drug cause was remote, those event terms which were so general as to be uninformative, and those occurring in only one patient. It is important to emphasize that, although the events reported occurred during treatment with Celexa™, they were not necessarily caused by it.

Events are further categorized by body system and listed in order of decreasing frequency according to the following definitions: frequent adverse events are those occurring on one or more occasions in at least 1/100 patients; infrequent adverse events are those occurring in less than 1/100 patients but at least 1/1000 patients; rare events are those occurring in fewer than 1/1000 patients.

**Cardiovascular** - *Frequent:* tachycardia, postural hypotension, hypotension. *Infrequent:* hypertension, bradycardia, edema (extremities), angina pectoris, extrasystoles, cardiac failure, flushing, myocardial infarction, cerebrovascular accident myocardial ischemia. *Rare:* transient ischemic attack, phlebitis, atrial fibrillation, cardiac arrest, bundle branch block.

**Central and Peripheral Nervous System Disorders** - *Frequent:* paresthesia, migraine. *Infrequent:* hyperkinesia, vertigo, hypertonia, extrapyramidal disorder, leg cramps, involuntary muscle contractions, hypokinesia, neuralgia, dystonia, abnormal gait, hypesthesia, ataxia. *Rare:* abnormal coordination, hyperesthesia, ptosis, stupor.

**Endocrine Disorders** - *Rare:* hypothyroidism, goiter, gynecomastia.

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**Gastrointestinal Disorders** - *Frequent:* saliva increased, flatulence. *Infrequent:* gastritis, gastroenteritis, stomatitis, eructation, hemorrhoids, dysphagia, teeth grinding, gingivitis, esophagitis. *Rare:* colitis, gastric ulcer, cholecystitis, cholelithiasis, duodenal ulcer, gastroesophageal reflux, glossitis, jaundice, diverticulitis, rectal hemorrhage, hiccups.

**General** - *Infrequent:* hot flushes, rigors, alcohol intolerance, syncope, influenza-like symptoms. *Rare:* hayfever.

**Hemic and Lymphatic Disorders** - *Infrequent:* purpura, anemia, epistaxis, leukocytosis, leucopenia, lymphadenopathy. *Rare:* pulmonary embolism, granulocytopenia, lymphocytosis, lymphopenia, hypochromic anemia, coagulation disorder, gingival bleeding.

**Metabolic and Nutritional Disorders** - *Frequent*: decreased weight, increased weight. *Infrequent*: increased hepatic enzymes, thirst, dry eyes, increased alkaline phosphatase, abnormal glucose tolerance. *Rare*: bilirubinemia, hypokalemia, obesity, hypoglycemia, hepatitis, dehydration.

**Musculoskeletal System Disorders** - *Infrequent*: arthritis, muscle weakness, skeletal pain. *Rare*: bursitis, osteoporosis.

**Psychiatric Disorders** - *Frequent*: impaired concentration, amnesia, apathy, depression, increased appetite, aggravated depression, suicide attempt, confusion. *Infrequent*: increased libido, aggressive reaction, paranoia, drug dependence, depersonalization, hallucination, euphoria, psychotic depression, delusion, paranoid reaction, emotional lability, panic reaction, psychosis. *Rare*: catatonic reaction, melancholia.

**Reproductive Disorders/Female\*** - *Frequent*: amenorrhea. *Infrequent*: galactorrhea, breast pain, breast enlargement, vaginal hemorrhage.

\*% based on female subjects only: 2955

**Respiratory System Disorders** - *Frequent*: coughing. *Infrequent*: bronchitis, dyspnea, pneumonia. *Rare*: asthma, laryngitis, bronchospasm, pneumonitis, sputum increased.

**Skin and Appendages Disorders** - *Frequent*: rash, pruritus. *Infrequent*: photosensitivity reaction, urticaria, acne, skin discoloration, eczema, alopecia, dermatitis, skin dry, psoriasis. *Rare*: hypertrichosis, decreased sweating, melanosis, keratitis, cellulitis, pruritus ani.

**Special Senses** - *Frequent*: accommodation abnormal, taste perversion. *Infrequent*: tinnitus, conjunctivitis, eye pain. *Rare*: mydriasis, photophobia, diplopia, abnormal lacrimation, cataract, taste loss.

**Urinary System Disorders** - *Frequent*: polyuria. *Infrequent*: micturition frequency, urinary incontinence, urinary retention, dysuria. *Rare*: facial edema, hematuria, oliguria, pyelonephritis, renal calculus, renal pain.

#### **Other Events Observed During the Non-US Postmarketing Evaluation of Celexa™ (citalopram HBr)**

It is estimated that approximately 8 million patients have been treated with Celexa™ since market introduction. Although no causal relationship to Celexa™ treatment has been found, the following adverse events have been reported to be temporally associated with Celexa™ treatment in at least 3 patients (unless otherwise noted) and are not described elsewhere in labeling: angioedema, choreoathetosis, epidermal necrolysis (3 cases), erythema multiforme, hepatic necrosis (2 cases), neuroleptic malignant syndrome, pancreatitis, serotonin syndrome, spontaneous abortion, thrombocytopenia, ventricular arrhythmia, Torsades de pointes, priapism, and withdrawal syndrome.

## **DRUG ABUSE AND DEPENDENCE**

### **Controlled Substance Class**

Celexa™ (citalopram HBr) is not a controlled substance.

### **Physical and Psychological Dependence**

Animal studies suggest that the abuse liability of Celexa™ is low. Celexa™ has not been systematically studied in humans for its potential for abuse, tolerance, or physical dependence. The premarketing clinical experience with Celexa™ 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 Celexa™ 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**

Although there were no reports of fatal citalopram overdose in clinical trials involving overdoses of up to 2000 mg, postmarketing reports of drug overdoses involving citalopram have included 12 fatalities, 10 in combination with other drugs and/or alcohol and 2 with citalopram alone (3920 mg and 2800 mg), as well as non-fatal overdoses of up to 6000 mg. Symptoms most often accompanying citalopram overdose, alone or in combination with other drugs and/or alcohol, included dizziness, sweating, nausea, vomiting, tremor, somnolence, and sinus tachycardia. In more rare cases, observed symptoms included amnesia, confusion, coma, convulsions, hyperventilation, cyanosis, rhabdomyolysis, and ECG changes (including QTc prolongation, nodal rhythm, ventricular arrhythmia, and one possible case of Torsades de pointes).

### **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 citalopram, forced diuresis, dialysis, hemoperfusion, and exchange transfusion are unlikely to be of benefit. There are no specific antidotes for Celexa™.

In managing overdosage, consider the possibility of multiple drug involvement. The physician should consider contacting a poison control center for additional information on the treatment of an overdose.

## **DOSAGE AND ADMINISTRATION**

### **Initial Treatment**

Celexa™ (citalopram HBr) should be administered at an initial dose of 20 mg once daily, generally with an increase to a dose of 40 mg/day. Dose increases should usually occur in increments of 20 mg at intervals of no less than one week. Although certain patients may require a dose of 60 mg/day, the only study pertinent to dose response for effectiveness did not demonstrate an advantage for the 60 mg/day dose over the 40 mg/day dose; doses above 40 mg are therefore not ordinarily recommended.

Celexa™ should be administered once daily, in the morning or evening, with or without food.

### **Special Populations**

20 mg/day is the recommended dose for most elderly patients and patients with hepatic impairment, with titration to 40 mg/day only for nonresponding patients.

No dosage adjustment is necessary for patients with mild or moderate renal impairment. Celexa™ should be used with caution in patients with severe renal impairment.

### **Maintenance Treatment**

It is generally agreed that acute episodes of depression require several months or longer of sustained pharmacologic therapy. Systematic evaluation of Celexa™ in two studies has shown that its antidepressant efficacy is maintained for periods of up to 24 weeks following 6 or 8 weeks of initial treatment (32 weeks total). In one study, patients were assigned randomly to placebo or to the same dose of Celexa™ (20-60 mg/day) during maintenance treatment as they had received during the acute stabilization phase, while in the other study, patients were assigned randomly to continuation of Celexa™ 20 or 40 mg/day, or placebo, for maintenance treatment. In the latter study, the rates of relapse to depression were similar for the two dose groups (see Clinical Trials under Clinical Pharmacology). Based on these limited data, it is not known whether the dose of citalopram needed to maintain euthymia is identical to the dose needed to induce remission. If adverse reactions are bothersome, a decrease in dose to 20 mg/day can be considered.

### **Switching Patients To or From a Monoamine Oxidase Inhibitor**

At least 14 days should elapse between discontinuation of an MAOI and initiation of Celexa™ therapy. Similarly, at least 14 days should be allowed after stopping Celexa™ before starting a MAOI (see Contraindications and Warnings).

## HOW SUPPLIED

20 mg Tablets	Bottle of 30	NDC # 0456-4020-30
	Bottle of 100	NDC # 0456-4020-01
	Bottle of 500	NDC # 0456-4020-05
	10 x 10 Unit Dose	NDC # 0456-4020-63

Pink, oval, scored film coated:  
Imprint on scored side with "F" on the left side and "P" on the right side.  
Imprint on the non-scored side with "20 mg".

40 mg Tablets	Bottle of 30	NDC # 0456-4040-30
	Bottle of 100	NDC # 0456-4040-01
	Bottle of 500	NDC # 0456-4040-05
	10 x 10 Unit Dose	NDC # 0456-4040-63

White, oval, scored film coated.  
Imprint on scored side with "F" on the left side and "P" on the right side.  
Imprint on the non-scored side with "40 mg".

Store at 25°C (77°F); excursions permitted to 15 - 30°C (59-86°F).

## ANIMAL TOXICOLOGY

### Retinal Changes in Rats

Pathologic changes (degeneration/atrophy) were observed in the retinas of albino rats in the 2-year carcinogenicity study with citalopram. There was an increase in both incidence and severity of retinal pathology in both male and female rats receiving 80 mg/kg/day (13 times the maximum recommended daily human dose of 60 mg on a mg/m<sup>2</sup> basis). Similar findings were not present in rats receiving 24 mg/kg/day for two years, in mice treated for 18 months at doses up to 240 mg/kg/day or in dogs treated for one year at doses up to 20 mg/kg/day (4, 20 and 10 times, respectively, the maximum recommended daily human dose on a mg/m<sup>2</sup> basis).

Additional studies to investigate the mechanism for this pathology have not been performed, and the potential significance of this effect in humans has not been established.

### Cardiovascular Changes in Dogs

In a one-year toxicology study, 5 of 10 beagle dogs receiving oral doses of 8 mg/kg/day (4 times the maximum recommended daily human dose of 60 mg on a mg/m<sup>2</sup> basis) died suddenly between weeks 17 and 31 following initiation of treatment. Although appropriate data from that study are not available to directly compare plasma levels of citalopram (CT) and its metabolites, demethylcitalopram (DCT) and didemethylcitalopram (DDCT), to levels that have been achieved in humans, pharmacokinetic data indicate that the relative dog to human exposure was greater for the metabolites than for citalopram. Sudden deaths were not observed in rats at doses up to 120 mg/kg/day, which produced plasma levels of CT, DCT and DDCT similar to those observed in dogs at doses of 8 mg/kg/day. A subsequent intravenous dosing study demonstrated that in beagle dogs, DDCT caused QT prolongation, a known risk factor for the observed outcome in dogs. This effect occurred in dogs at doses producing peak DDCT plasma levels of 810 to 3250 nM (39-155 times the mean steady state DDCT plasma level measured at the maximum recommended human daily dose of 60 mg. In dogs, peak DDCT plasma concentrations are approximately equal to peak CT plasma concentrations, whereas in humans, steady state DDCT plasma concentrations are less than 10% of steady state CT plasma concentrations. Assays of DDCT plasma concentrations in 2,020 citalopram treated individuals demonstrated that DDCT levels rarely exceeded 70 nM; the highest measured level of DDCT in human overdose was 138 nM. While DDCT is ordinarily present in humans at lower levels than in dogs, it is unknown whether there are individuals who may achieve higher DDCT levels. The possibility that DCT, a principal metabolite in humans, may prolong the QT interval in the dog has not been directly examined because DCT is rapidly converted to DDCT in that species.

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