

**OLMESARTAN MEDOXOMIL - olmesartan medoxomil tablet, coated**  
**Ascend Laboratories, LLC**

**HIGHLIGHTS OF PRESCRIBING INFORMATION**

These highlights do not include all the information needed to use Olmesartan Medoxomil Tablets safely and effectively. See full prescribing information for Olmesartan Medoxomil Tablets.

**OLMESARTAN Medoxomil Tablets, for oral use**  
**Initial U.S. Approval: 2002**

**WARNING:FETAL TOXICITY**

*See full prescribing information for complete boxed warning.*

- When pregnancy is detected, discontinue Olmesartan Medoxomil Tablets as soon as possible (5.1).
- Drugs that act directly on the renin-angiotensin system can cause injury and death to the developing fetus (5.1).

**RECENT MAJOR CHANGES**

Warnings and Precautions (5.6)

11/2016

**INDICATIONS AND USAGE**

Olmesartan Medoxomil Tablets is an angiotensin II receptor blocker (ARB) indicated for the treatment of hypertension, alone or with other antihypertensive agents, to lower blood pressure. Lowering blood pressure reduces the risk of fatal and nonfatal cardiovascular events, primarily strokes and myocardial infarctions. (1)

**DOSAGE AND ADMINISTRATION**

Indication	Starting dose	Dose Range
Adult Hypertension (2.1)	20 mg once daily	20 to 40 mg once daily
Pediatric Hypertension (6 to 16 years) (2.2)	20 to <35 kg 10 mg once daily ≥35 kg 20 mg once daily	20 to <35 kg 10 to 20 mg once daily ≥35 kg 20 to 40 mg once daily

- Olmesartan Medoxomil Tablets may be administered with or without food. (2)
- If blood pressure is not controlled by Olmesartan Medoxomil Tablets alone, a diuretic may be added. Olmesartan Medoxomil Tablets may be administered with other antihypertensive agents. (2)

**DOSAGE FORMS AND STRENGTHS**

Tablets: 5 mg, 20 mg, and 40 mg (3)

**CONTRAINDICATIONS**

Do not co-administer aliskiren with Olmesartan Medoxomil Tablets in patients with diabetes. (4)

**WARNINGS AND PRECAUTIONS**

- Avoid fetal (in utero) exposure (5.1).
- Children <1 year of age must not receive Olmesartan Medoxomil Tablets for hypertension (5.2).
- Observe for signs and symptoms of hypotension in volume- or salt-depleted patients with treatment initiation (5.3).
- Monitor for worsening renal function in patients with renal impairment (5.4).
- Sprue-like enteropathy has been reported. Consider discontinuation of Olmesartan Medoxomil Tablets in cases where no other etiology is found (5.5).

**ADVERSE REACTIONS**

The most common adverse reaction in adults was dizziness (3%) (6.1).

**To report SUSPECTED ADVERSE REACTIONS, contact Ascend Laboratories, LLC at 1-877-ASC-RX01 (877-272-7901) or FDA at 1-800-332-1088 or [www.fda.gov/medwatch](http://www.fda.gov/medwatch).**

**DRUG INTERACTIONS**

- NSAID use may lead to increased risk of renal impairment and loss of antihypertensive effect (7).
- Dual inhibition of the renin-angiotensin system: Increased risk of renal impairment, hypotension, and hyperkalemia (7).
- Colesevelam hydrochloride: Consider administering olmesartan at least 4 hours before colesevelam hydrochloride dose (7)
- Lithium: Increases in serum lithium concentrations and lithium toxicity (7).

----- **USE IN SPECIFIC POPULATIONS** -----

- Nursing mothers: Choose to discontinue nursing or drug (8.3).
- In patients with an activated renin-angiotensin system, such as volume- or salt-depletion, renin-angiotensin-aldosterone system (RAAS) blockers such as olmesartan medoxomil can cause excessive hypotension. In susceptible patients, e.g., with renal artery stenosis, RAAS blockers can cause renal failure (5.3, 5.4).
- Geriatrics: No overall difference in efficacy or safety vs. younger adult patients, but greater sensitivity of some older individuals cannot be ruled out (8.5).

See 17 for **PATIENT COUNSELING INFORMATION**.

Revised: 3/2017

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**FULL PRESCRIBING INFORMATION: CONTENTS\***

**WARNING: FETAL TOXICITY**

**1 INDICATIONS & USAGE**

**2 DOSAGE & ADMINISTRATION**

2.1 Adult Hypertension

2.2 Pediatric Hypertension (6 to 16 years of age)

**3 DOSAGE FORMS & STRENGTHS**

**4 CONTRAINDICATIONS**

**5 WARNINGS AND PRECAUTIONS**

5.1 Fetal toxicity

5.2 Morbidity in Infants

5.3 Hypotension in Volume or Salt Depleted Patients

5.4 Impaired Renal Function

5.5 Sprue-like Enteropathy

5.6 Electrolyte Imbalances

**6 ADVERSE REACTIONS**

6.1 Clinical Trials Experience

6.2 Post-Marketing Experience

**7 DRUG INTERACTIONS**

**8 USE IN SPECIFIC POPULATIONS**

8.1 Pregnancy

8.3 Nursing Mothers

8.4 Pediatric Use

8.5 Geriatric Use

8.6 Hepatic Impairment

8.7 Renal Impairment

8.8 Black Patients

**10 OVERDOSAGE**

**11 DESCRIPTION**

**12 CLINICAL PHARMACOLOGY**

12.1 Mechanism of Action

12.2 Pharmacodynamics

12.3 Pharmacokinetics

**13 NONCLINICAL TOXICOLOGY**

13.1 Carcinogenesis & Mutagenesis & Impairment Of Fertility

13.2 Animal Pharmacology & OR Toxicology

**14 CLINICAL STUDIES**

14.1 Adult Hypertension

14.2 Pediatric Hypertension

**16 HOW SUPPLIED/STORAGE AND HANDLING**

**17 PATIENT COUNSELING INFORMATION**

\* Sections or subsections omitted from the full prescribing information are not listed.

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## FULL PRESCRIBING INFORMATION

### WARNING: FETAL TOXICITY

- **When pregnancy is detected, discontinue Olmesartan Medoxomil Tablets as soon as possible (5.1)**
- **Drugs that act directly on the renin-angiotensin system can cause injury and death to the developing fetus (5.1)**

## 1 INDICATIONS & USAGE

Olmesartan Medoxomil Tablets are indicated for the treatment of hypertension, to lower blood pressure. Lowering blood pressure reduces the risk of fatal and nonfatal cardiovascular events, primarily strokes and myocardial infarctions. These benefits have been seen in controlled trials of antihypertensive drugs from a wide variety of pharmacologic classes including the class to which this drug principally belongs. There are no controlled trials demonstrating risk reduction with Olmesartan Medoxomil Tablets.

Control of high blood pressure should be part of comprehensive cardiovascular risk management, including, as appropriate, lipid control, diabetes management, antithrombotic therapy, smoking cessation, exercise, and limited sodium intake. Many patients will require more than one drug to achieve blood pressure goals. For specific advice on goals and management, see published guidelines, such as those of the National High Blood Pressure Education Program's Joint National Committee on Prevention, Detection, Evaluation, and Treatment of High Blood Pressure (JNC).

Numerous antihypertensive drugs, from a variety of pharmacologic classes and with different mechanisms of action, have been shown in randomized controlled trials to reduce cardiovascular morbidity and mortality, and it can be concluded that it is blood pressure reduction, and not some other pharmacologic property of the drugs, that is largely responsible for those benefits. The largest and most consistent cardiovascular outcome benefit has been a reduction in the risk of stroke, but reductions in myocardial infarction and cardiovascular mortality also have been seen regularly.

Elevated systolic or diastolic pressure causes increased cardiovascular risk, and the absolute risk increase per mmHg is greater at higher blood pressures, so that even modest reductions of severe hypertension can provide substantial benefit. Relative risk reduction from blood pressure reduction is similar across populations with varying absolute risk, so the absolute benefit is greater in patients who are at higher risk independent of their hypertension (for example, patients with diabetes or hyperlipidemia), and such patients would be expected to benefit from more aggressive treatment to a lower blood pressure goal.

Some antihypertensive drugs have smaller blood pressure effects (as monotherapy) in black patients, and many antihypertensive drugs have additional approved indications and effects (e.g., on angina, heart failure, or diabetic kidney disease). These considerations may guide selection of therapy.

It may be used alone or in combination with other antihypertensive agents.

## 2 DOSAGE & ADMINISTRATION

## 2.1 Adult Hypertension

Dosage must be individualized. The usual recommended starting dose of Olmesartan Medoxomil Tablets is 20 mg once daily when used as monotherapy in patients who are not volume-contracted. For patients requiring further reduction in blood pressure after 2 weeks of therapy, the dose of Olmesartan Medoxomil Tablets may be increased to 40 mg. Doses above 40 mg do not appear to have greater effect. Twice-daily dosing offers no advantage over the same total dose given once daily.

No initial dosage adjustment is recommended for elderly patients, for patients with moderate to marked renal impairment (creatinine clearance <40 mL/min) or with moderate to marked hepatic dysfunction [see Warnings and Precautions (5.4), Use in Specific Populations (8.5,8.6,8.7) and Clinical Pharmacology (12.3)]. For patients with possible depletion of intravascular volume (e.g., patients treated with diuretics, particularly those with impaired renal function), initiate Olmesartan Medoxomil Tablets under close medical supervision and give consideration to use of a lower starting dose [see *Warnings and Precautions* (5.3)].

Olmesartan Medoxomil Tablets may be administered with or without food.

If blood pressure is not controlled by Olmesartan Medoxomil Tablets alone, a diuretic may be added. Olmesartan Medoxomil Tablets may be administered with other antihypertensive agents.

## 2.2 Pediatric Hypertension (6 to 16 years of age)

Dosage must be individualized. For children who can swallow tablets, the usual recommended starting dose of Olmesartan Medoxomil Tablets is 10 mg once daily for patients who weigh 20 to <35 kg (44 to 77 lb), or 20 mg once daily for patients who weigh ≥35 kg. For patients requiring further reduction in blood pressure after 2 weeks of therapy, the dose of Olmesartan Medoxomil Tablets may be increased to a maximum of 20 mg once daily for patients who weigh <35 kg or 40 mg once daily for patients who weigh ≥35 kg.

Children <1 year of age must not receive Olmesartan Medoxomil Tablets for hypertension.

For children who cannot swallow tablets, the same dose can be given using an extemporaneous suspension as described below [see *Clinical Pharmacology* (12.3)]. Follow the suspension preparation instructions below to administer Olmesartan Medoxomil Tablets as a suspension.

Preparation of Suspension (for 200 mL of a 2 mg/mL suspension)

Add 50 mL of Purified Water to an amber polyethylene terephthalate (PET) bottle containing twenty Olmesartan Medoxomil Tablets 20 mg and allow to stand for a minimum of 5 minutes. Shake the container for at least 1 minute and allow the suspension to stand for at least 1 minute. Repeat 1-minute shaking and 1-minute standing for four additional times. Add 100 mL of Ora-Sweet®\* and 50 mL of Ora-Plus®\* to the suspension and shake well for at least 1 minute. The suspension should be refrigerated at 2 to 8°C (36 to 46°F) and can be stored for up to 4 weeks. Shake the suspension well before each use and return promptly to the refrigerator.

\* Ora-Sweet® and Ora-Plus® are registered trademarks of Paddock Laboratories, Inc.

## 3 DOSAGE FORMS & STRENGTHS

- 5 mg yellow coloured round shape film coated tablets debossed with OLM on one side & 5 on other side.
- 20 mg white to off-white coloured, round shape film coated tablets debossed with OLM on one side & 20 on other side.
- 40 mg white to off-white coloured, oval shape film coated tablets debossed with OLM on one side & 40 on other side.

## 4 CONTRAINDICATIONS

Do not co-administer aliskiren with Olmesartan Medoxomil Tablets in patients with diabetes [See *Drug Interactions (7)*].

## 5 WARNINGS AND PRECAUTIONS

### 5.1 Fetal toxicity

#### Pregnancy Category D

Use of drugs that act on the renin-angiotensin system during the second and third trimesters of pregnancy reduces fetal renal function and increases fetal and neonatal morbidity and death. Resulting oligohydramnios can be associated with fetal lung hypoplasia and skeletal deformations. Potential neonatal adverse effects include skull hypoplasia, anuria, hypotension, renal failure, and death. When pregnancy is detected, discontinue Olmesartan Medoxomil Tablets as soon as possible [see *Use in Specific Populations (8.1)*].

### 5.2 Morbidity in Infants

Children <1 year of age must not receive Olmesartan Medoxomil Tablets for hypertension. Drugs that act directly on the renin-angiotensin aldosterone system (RAAS) can have effects on the development of immature kidneys [see *Use in Specific Populations (8.4)*].

### 5.3 Hypotension in Volume or Salt Depleted Patients

In patients with an activated renin-angiotensin aldosterone system, such as volume- and/or salt-depleted patients (e.g., those being treated with high doses of diuretics), symptomatic hypotension may be anticipated after initiation of treatment with Olmesartan Medoxomil Tablets. Initiate treatment under close medical supervision. If hypotension does occur, place the patient in the supine position and, if necessary, give an intravenous infusion of normal saline [see *Dosage and Administration (2.1)*]. A transient hypotensive response is not a contraindication to further treatment, which usually can be continued without difficulty once the blood pressure has stabilized.

### 5.4 Impaired Renal Function

As a consequence of inhibiting the renin-angiotensin-aldosterone system, changes in renal function may be anticipated in susceptible individuals treated with Olmesartan Medoxomil Tablets. In patients whose renal function may depend upon the activity of the renin-angiotensin aldosterone system (e.g., patients with severe congestive heart failure), treatment with angiotensin converting enzyme (ACE) inhibitors and angiotensin receptor antagonists has been associated with oliguria and/or progressive azotemia and rarely with acute renal failure and/or death. Similar results may be anticipated in patients treated with Olmesartan Medoxomil Tablets [see *Dosage and Administration (2.1)*, *Drug Interactions (7)*, *Use in Specific Populations (8.7)* and *Clinical Pharmacology (12.3)*].

In studies of ACE inhibitors in patients with unilateral or bilateral renal artery stenosis, increases in serum creatinine or blood urea nitrogen (BUN) have been reported. There has been no long-term use of Olmesartan Medoxomil Tablets in patients with unilateral or bilateral renal artery stenosis, but similar results may be expected.

### 5.5 Sprue-like Enteropathy

Severe, chronic diarrhea with substantial weight loss has been reported in patients taking olmesartan months to years after drug initiation. Intestinal biopsies of patients often demonstrated villous atrophy. If a patient develops these symptoms during treatment with olmesartan, exclude other etiologies. Consider

discontinuation of Olmesartan Medoxomil Tablets in cases where no other etiology is identified.

## 5.6 Electrolyte Imbalances

Olmesartan Medoxomil Tablet contains olmesartan, a drug that inhibits the renin-angiotensin system (RAS). Drugs that inhibit the RAS can cause hyperkalemia. Monitor serum electrolytes periodically.

## 6 ADVERSE REACTIONS

### 6.1 Clinical Trials Experience

Because clinical studies are conducted under widely varying conditions, adverse reaction rates observed in the clinical studies of a drug cannot be directly compared to rates in the clinical studies of another drug and may not reflect the rates observed in practice.

#### Adult Hypertension

Olmesartan Medoxomil Tablets has been evaluated for safety in more than 3825 patients/subjects, including more than 3275 patients treated for hypertension in controlled trials. This experience included about 900 patients treated for at least 6 months and more than 525 for at least 1 year. Treatment with Olmesartan Medoxomil Tablets was well tolerated, with an incidence of adverse reactions similar to placebo. Events generally were mild, transient and had no relationship to the dose of Olmesartan Medoxomil Tablets .

The overall frequency of adverse reactions was not dose-related. Analysis of gender, age and race groups demonstrated no differences between Olmesartan Medoxomil Tablets and placebo-treated patients. The rate of withdrawals due to adverse reactions in all trials of hypertensive patients was 2.4% (i.e., 79/3278) of patients treated with Olmesartan Medoxomil Tablets and 2.7% (i.e., 32/1179) of control patients. In placebo-controlled trials, the only adverse reaction that occurred in more than 1% of patients treated with Olmesartan Medoxomil Tablets and at a higher incidence versus placebo was dizziness (3% vs. 1%).

The following adverse reactions occurred in placebo-controlled clinical trials at an incidence of more than 1% of patients treated with Olmesartan Medoxomil Tablets , but also occurred at about the same or greater incidence in patients receiving placebo: back pain, bronchitis, creatine phosphokinase increased, diarrhea, headache, hematuria, hyperglycemia, hypertriglyceridemia, influenza-like symptoms, pharyngitis, rhinitis and sinusitis.

The incidence of cough was similar in placebo (0.7%) and Olmesartan Medoxomil Tablets (0.9%) patients.

Other potentially important adverse reactions that have been reported with an incidence of greater than 0.5%, whether or not attributed to treatment, in the more than 3100 hypertensive patients treated with Olmesartan Medoxomil Tablets monotherapy in controlled or open-label trials are listed below.

*Body as a Whole:* chest pain, peripheral edema

*Central and Peripheral Nervous System:* vertigo  
*Gastrointestinal:* abdominal pain, dyspepsia, gastroenteritis, nausea

*Heart Rate and Rhythm Disorders:* tachycardia  
*Metabolic and Nutritional Disorders:* hypercholesterolemia, hyperlipemia, hyperuricemia  
*Musculoskeletal:* arthralgia, arthritis, myalgia

*Skin and appendages:* rash

Facial edema was reported in five patients receiving Olmesartan Medoxomil Tablets . Angioedema has been reported with angiotensin II antagonists.

*Laboratory Test Findings:* In controlled clinical trials, clinically important changes in standard laboratory parameters were rarely associated with administration of Olmesartan Medoxomil Tablets .

*Hemoglobin and Hematocrit:* Small decreases in hemoglobin and hematocrit (mean decreases of approximately 0.3 g/dL and 0.3 volume percent, respectively) were observed.

*Liver Function Tests:* Elevations of liver enzymes and/or serum bilirubin were observed infrequently. Five patients (0.1%) assigned to Olmesartan Medoxomil Tablets and one patient (0.2%) assigned to placebo in clinical trials were withdrawn because of abnormal liver chemistries (transaminases or total bilirubin). Of the five Olmesartan Medoxomil Tablets patients, three had elevated transaminases, which were attributed to alcohol use, and one had a single elevated bilirubin value, which normalized while treatment continued.

## **Pediatric Hypertension**

No relevant differences were identified between the adverse experience profile for pediatric patients aged 1 to 16 years and that previously reported for adult patients.

## **6.2 Post-Marketing Experience**

The following adverse reactions have been reported in post-marketing experience. Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to reliably estimate their frequency or establish a causal relationship to drug exposure.

*Body as a Whole:* Asthenia, angioedema, anaphylactic reactions

*Gastrointestinal:* Vomiting, sprue-like enteropathy [see *Warnings and Precautions (5.6)*]

*Metabolic and Nutritional Disorders:* Hyperkalemia

*Musculoskeletal:* Rhabdomyolysis

*Urogenital System:* Acute renal failure, increased blood creatinine levels

*Skin and Appendages:* Alopecia, pruritus, urticaria

Data from one controlled trial and an epidemiologic study have suggested that high-dose olmesartan may increase cardiovascular (CV) risk in diabetic patients, but the overall data are not conclusive. The randomized, placebo-controlled, double-blind ROADMAP trial (Randomized Olmesartan And Diabetes MicroAlbuminuria Prevention trial, n=4447) examined the use of olmesartan, 40 mg daily, vs. placebo in patients with type 2 diabetes mellitus, normoalbuminuria, and at least one additional risk factor for CV disease. The trial met its primary endpoint, delayed onset of microalbuminuria, but olmesartan had no beneficial effect on decline in glomerular filtration rate (GFR). There was a finding of increased CV mortality (adjudicated sudden cardiac death, fatal myocardial infarction, fatal stroke, revascularization death) in the olmesartan group compared to the placebo group (15 olmesartan vs. 3 placebo, HR 4.9, 95% confidence interval [CI], 1.4, 17), but the risk of non-fatal myocardial infarction was lower with olmesartan (HR 0.64, 95% CI 0.35, 1.18).

The epidemiologic study included patients 65 years and older with overall exposure of > 300,000 patient-years. In the sub-group of diabetic patients receiving high-dose olmesartan (40 mg/d) for > 6 months, there appeared to be an increased risk of death (HR 2.0, 95% CI 1.1, 3.8) compared to similar patients taking other angiotensin receptor blockers. In contrast, high-dose olmesartan use in non-diabetic patients appeared to be associated with a decreased risk of death (HR 0.46, 95% CI 0.24, 0.86) compared to similar patients taking other angiotensin receptor blockers. No differences were observed between the groups receiving lower doses of olmesartan compared to other angiotensin blockers or those receiving therapy for < 6 months.

Overall, these data raise a concern of a possible increased CV risk associated with the use of high-dose olmesartan in diabetic patients. There are, however, concerns with the credibility of the finding of increased CV risk, notably the observation in the large epidemiologic study for a survival benefit in non-diabetics of a magnitude similar to the adverse finding in diabetics.

## **7 DRUG INTERACTIONS**

No significant drug interactions were reported in studies in which Olmesartan Medoxomil Tablets was coadministered with digoxin or warfarin in healthy volunteers.

The bioavailability of olmesartan was not significantly altered by the co-administration of antacids [Al(OH)<sub>3</sub>/Mg(OH)<sub>2</sub>].

Olmesartan medoxomil is not metabolized by the cytochrome P450 system and has no effects on P450 enzymes; thus, interactions with drugs that inhibit, induce, or are metabolized by those enzymes are not expected.

#### *Non-Steroidal Anti-Inflammatory Agents including Selective Cyclooxygenase-2 Inhibitors (COX-2 Inhibitors)*

In patients who are elderly, volume-depleted (including those on diuretic therapy), or with compromised renal function, co-administration of NSAIDs, including selective COX-2 inhibitors, with angiotensin II receptor antagonists, including olmesartan medoxomil, may result in deterioration of renal function, including possible acute renal failure. These effects are usually reversible. Monitor renal function periodically in patients receiving olmesartan medoxomil and NSAID therapy.

The antihypertensive effect of angiotensin II receptor antagonists, including olmesartan medoxomil may be attenuated by NSAIDs including selective COX-2 inhibitors.

#### *Dual Blockade of the Renin-Angiotensin System (RAS)*

Dual blockade of the RAS with angiotensin receptor blockers, ACE inhibitors, or aliskiren is associated with increased risks of hypotension, hyperkalemia, and changes in renal function (including acute renal failure) compared to monotherapy. Most patients receiving the combination of two RAS inhibitors do not obtain any additional benefit compared to monotherapy. In general, avoid combined use of RAS inhibitors. Closely monitor blood pressure, renal function and electrolytes in patients on Benicar and other agents that affect the RAS.

Do not co-administer aliskiren with Olmesartan Medoxomil Tablets in patients with diabetes [see Contraindications (4)]. Avoid use of aliskiren with Olmesartan Medoxomil Tablets in patients with renal impairment (GFR <60 ml/min).

#### *Colesevelam hydrochloride*

Concurrent administration of bile acid sequestering agent colesevelam hydrochloride reduces the systemic exposure and peak plasma concentration of olmesartan. Administration of olmesartan at least 4 hours prior to colesevelam hydrochloride decreased the drug interaction effect. Consider administering olmesartan at least 4 hours before the colesevelam hydrochloride dose [see *Clinical Pharmacology* (12.3)].

#### *Lithium*

Increases in serum lithium concentrations and lithium toxicity have been reported during concomitant administration of lithium with angiotensin II receptor antagonists, including Olmesartan Medoxomil Tablets. Monitor serum lithium levels during concomitant use.

## **8 USE IN SPECIFIC POPULATIONS**

### **8.1 Pregnancy**

#### **Pregnancy Category D**

Use of drugs that act on the renin-angiotensin system during the second and third trimesters of



pregnancy reduces fetal renal function and increases fetal and neonatal morbidity and death. Resulting oligohydramnios can be associated with fetal lung hypoplasia and skeletal deformations. Potential neonatal adverse effects include skull hypoplasia, anuria, hypotension, renal failure, and death. When pregnancy is detected, discontinue Olmesartan Medoxomil Tablets as soon as possible. These adverse outcomes are usually associated with use of these drugs in the second and third trimester of pregnancy. Most epidemiologic studies examining fetal abnormalities after exposure to antihypertensive use in the first trimester have not distinguished drugs affecting the renin-angiotensin system from other antihypertensive agents. Appropriate management of maternal hypertension during pregnancy is important to optimize outcomes for both mother and fetus.

In the unusual case that there is no appropriate alternative to therapy with drugs affecting the renin-angiotensin system for a particular patient, apprise the mother of the potential risk to the fetus. Perform serial ultrasound examinations to assess the intra-amniotic environment. If oligohydramnios is observed, discontinue Olmesartan Medoxomil Tablets, unless it is considered lifesaving for the mother. Fetal testing may be appropriate, based on the week of pregnancy. Patients and physicians should be aware, however, that oligohydramnios may not appear until after the fetus has sustained irreversible injury. Closely observe infants with histories of in utero exposure to Olmesartan Medoxomil Tablets for hypotension, oliguria, and hyperkalemia [see *Use in Specific Populations (8.4)*].

### **8.3 Nursing Mothers**

It is not known whether olmesartan is excreted in human milk, but olmesartan is secreted at low concentration in the milk of lactating rats. Because of the potential for adverse effects on the nursing infant, a decision should be made whether to discontinue nursing or discontinue the drug, taking into account the importance of the drug to the mother.

### **8.4 Pediatric Use**

Neonates with a history of in utero exposure to Olmesartan Medoxomil Tablets :

If oliguria or hypotension occurs, direct attention toward support of blood pressure and renal perfusion. Exchange transfusions or dialysis may be required as a means of reversing hypotension and/or substituting for disordered renal function.

The antihypertensive effects of Olmesartan Medoxomil Tablets were evaluated in one randomized, double-blind clinical study in pediatric patients 1 to 16 years of age [see *Clinical Studies (14.2)*]. The pharmacokinetics of Olmesartan Medoxomil Tablets were evaluated in pediatric patients 1 to 16 years of age [see *Clinical Pharmacology (12.3)*]. Olmesartan Medoxomil Tablets was generally well tolerated in pediatric patients, and the adverse experience profile was similar to that described for adults.

Olmesartan Medoxomil Tablets has not been shown to be effective for hypertension in children <6 years of age.

Children <1 year of age must not receive Olmesartan Medoxomil Tablets for hypertension [see *Warnings and Precautions (5.2)*]. The renin-angiotensin aldosterone system (RAAS) plays a critical role in kidney development. RAAS blockade has been shown to lead to abnormal kidney development in very young mice. Administering drugs that act directly on the renin-angiotensin aldosterone system (RAAS) can alter normal renal development.

### **8.5 Geriatric Use**

Of the total number of hypertensive patients receiving Olmesartan Medoxomil Tablets in clinical studies, more than 20% were 65 years of age and over, while more than 5% were 75 years of age and older. No overall differences in effectiveness or safety were observed between elderly patients and younger patients. 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 [see *Dosage and Administration (2.1)* and *Clinical Pharmacology (12.3)*].

## 8.6 Hepatic Impairment

Increases in AUC<sub>0-a</sub> and C<sub>max</sub> were observed in patients with moderate hepatic impairment compared to those in matched controls, with an increase in AUC of about 60%. No initial dosage adjustment is recommended for patients with moderate to marked hepatic dysfunction [see *Dosage and Administration (2.1) and Clinical Pharmacology (12.3)*].

## 8.7 Renal Impairment

Patients with renal insufficiency have elevated serum concentrations of olmesartan compared to subjects with normal renal function. After repeated dosing, the AUC was approximately tripled in patients with severe renal impairment (creatinine clearance < 20 mL/min). No initial dosage adjustment is recommended for patients with moderate to marked renal impairment (creatinine clearance < 40 mL/min) [see *Dosage and Administration (2.1), Warnings and Precautions (5.4) and Clinical Pharmacology (12.3)*].

## 8.8 Black Patients

The antihypertensive effect of Olmesartan Medoxomil Tablets was smaller in black patients (usually a low-renin population), as has been seen with ACE inhibitors, beta-blockers and other angiotensin receptor blockers.

## 10 OVERDOSAGE

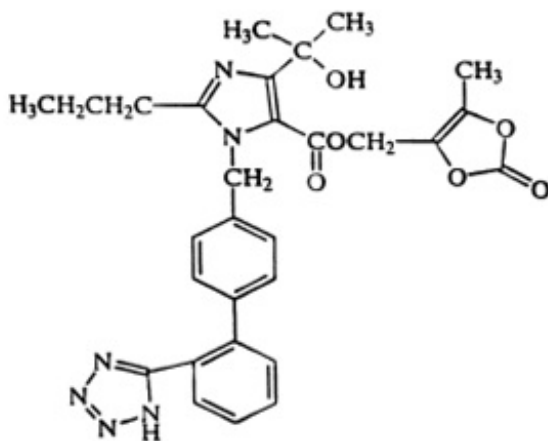
Limited data are available related to overdosage in humans. The most likely manifestations of overdosage would be hypotension and tachycardia; bradycardia could be encountered if parasympathetic (vagal) stimulation occurs. If symptomatic hypotension occurs, initiate supportive treatment. The dialyzability of olmesartan is unknown.

## 11 DESCRIPTION

Olmesartan medoxomil, USP a prodrug, is hydrolyzed to olmesartan during absorption from the gastrointestinal tract. Olmesartan is a selective AT<sub>1</sub> subtype angiotensin II receptor antagonist.

Olmesartan medoxomil is described chemically as 2,3-dihydroxy-2-butenyl 4-(1 hydroxy-1-methylethyl)-2-propyl-1-[p-(o-1H-tetrazol-5-ylphenyl)benzyl]imidazole-5-carboxylate, cyclic 2,3-carbonate.

Its empirical formula is C<sub>29</sub>H<sub>30</sub>N<sub>6</sub>O<sub>6</sub> and its structural formula is:



Olmesartan medoxomil, USP is a white to off-white crystalline powder with a molecular weight of 558.59. It is practically insoluble in water and in heptanes, slightly soluble in ethanol (96%), sparingly soluble in methanol. Olmesartan Medoxomil Tablets is available for oral use as film-coated tablets

containing 5 mg, 20 mg, or 40 mg of olmesartan medoxomil, USP and the following inactive ingredients: lactose monohydrate, microcrystalline cellulose, low-substituted hydroxypropyl cellulose, hydroxypropyl cellulose, stearic acid, magnesium stearate, talc, titanium dioxide, and (5 mg only) yellow iron oxide.

## 12 CLINICAL PHARMACOLOGY

### 12.1 Mechanism of Action

Angiotensin II is formed from angiotensin I in a reaction catalyzed by angiotensin converting enzyme (ACE, kininase II). Angiotensin II is the principal pressor agent of the renin-angiotensin system, with effects that include vasoconstriction, stimulation of synthesis and release of aldosterone, cardiac stimulation and renal reabsorption of sodium. Olmesartan blocks the vasoconstrictor effects of angiotensin II by selectively blocking the binding of angiotensin II to the AT1 receptor in vascular smooth muscle. Its action is, therefore, independent of the pathways for angiotensin II synthesis.

An AT2 receptor is found also in many tissues, but this receptor is not known to be associated with cardiovascular homeostasis. Olmesartan has more than a 12,500-fold greater affinity for the AT1 receptor than for the AT2 receptor.

Blockade of the renin-angiotensin system with ACE inhibitors, which inhibit the biosynthesis of angiotensin II from angiotensin I, is a mechanism of many drugs used to treat hypertension. ACE inhibitors also inhibit the degradation of bradykinin, a reaction also catalyzed by ACE. Because olmesartan medoxomil does not inhibit ACE (kininase II), it does not affect the response to bradykinin. Whether this difference has clinical relevance is not yet known.

Blockade of the angiotensin II receptor inhibits the negative regulatory feedback of angiotensin II on renin secretion, but the resulting increased plasma renin activity and circulating angiotensin II levels do not overcome the effect of olmesartan on blood pressure.

### 12.2 Pharmacodynamics

Olmесartan Medoxomil Tablets doses of 2.5 mg to 40 mg inhibit the pressor effects of angiotensin I infusion. The duration of the inhibitory effect was related to dose, with doses of Olmesartan Medoxomil Tablets >40 mg giving >90% inhibition at 24 hours.

Plasma concentrations of angiotensin I and angiotensin II and plasma renin activity (PRA) increase after single and repeated administration of Olmesartan Medoxomil Tablets to healthy subjects and hypertensive patients. Repeated administration of up to 80 mg Olmesartan Medoxomil Tablets had minimal influence on aldosterone levels and no effect on serum potassium.

### 12.3 Pharmacokinetics

#### Absorption

Olmесartan medoxomil is rapidly and completely bioactivated by ester hydrolysis to olmesartan during absorption from the gastrointestinal tract.

Olmесartan Medoxomil Tablets and the suspension formulation prepared from Olmesartan Medoxomil Tablets are bioequivalent [*see Dosage and Administration (2.2)*].

The absolute bioavailability of olmesartan is approximately 26%. After oral administration, the peak plasma concentration (C<sub>max</sub>) of olmesartan is reached after 1 to 2 hours. Food does not affect the bioavailability of olmesartan.

#### Distribution

The volume of distribution of olmesartan is approximately 17 L. Olmesartan is highly bound to plasma proteins (99%) and does not penetrate red blood cells. The protein binding is constant at plasma

olmesartan concentrations well above the range achieved with recommended doses.

In rats, olmesartan crossed the blood-brain barrier poorly, if at all. Olmesartan passed across the placental barrier in rats and was distributed to the fetus. Olmesartan was distributed to milk at low levels in rats.

### **Metabolism and Excretion**

Following the rapid and complete conversion of olmesartan medoxomil to olmesartan during absorption, there is virtually no further metabolism of olmesartan. Total plasma clearance of olmesartan is 1.3 L/h, with a renal clearance of 0.6 L/h. Approximately 35% to 50% of the absorbed dose is recovered in urine while the remainder is eliminated in feces via the bile.

Olmesartan appears to be eliminated in a biphasic manner with a terminal elimination half-life of approximately 13 hours. Olmesartan shows linear pharmacokinetics following single oral doses of up to 320 mg and multiple oral doses of up to 80 mg. Steady-state levels of olmesartan are achieved within 3 to 5 days and no accumulation in plasma occurs with once-daily dosing.

### **Geriatric**

The pharmacokinetics of olmesartan were studied in the elderly ( $\geq 65$  years). Overall, maximum plasma concentrations of olmesartan were similar in young adults and the elderly. Modest accumulation of olmesartan was observed in the elderly with repeated dosing; AUC<sub>ss</sub>,  $\tau$  was 33% higher in elderly patients, corresponding to an approximate 30% reduction in CLR [see *Dosage and Administration (2.1) and Use in Specific Populations (8.5)*].

### **Pediatric**

The pharmacokinetics of olmesartan were studied in pediatric hypertensive patients aged 1 to 16 years. The clearance of olmesartan in pediatric patients was similar to that in adult patients when adjusted by the body weight [see *Use in Specific Populations (8.4)*].

Olmesartan pharmacokinetics have not been investigated in pediatric patients less than 1 year of age [see *Warnings and Precautions (5.2) and Use in Specific Populations (8.4)*].

### **Gender**

Minor differences were observed in the pharmacokinetics of olmesartan in women compared to men. AUC and C<sub>max</sub> were 10-15% higher in women than in men.

### **Hepatic Insufficiency**

Increases in AUC<sub>0-a</sub> and C<sub>max</sub> were observed in patients with moderate hepatic impairment compared to those in matched controls, with an increase in AUC of about 60% [see *Dosage and Administration (2.1) and Use in Specific Populations (8.6)*].

### **Renal Insufficiency**

In patients with renal insufficiency, serum concentrations of olmesartan were elevated compared to subjects with normal renal function. After repeated dosing, the AUC was approximately tripled in patients with severe renal impairment (creatinine clearance  $< 20$  mL/min). The pharmacokinetics of olmesartan in patients undergoing hemodialysis has not been studied [see *Dosage and Administration (2.1), Warnings and Precautions (5.4) and Use in Specific Populations (8.7)*].

### **Drug Interaction**

#### **Bile acid sequestering agent colesvelam**

Concomitant administration of 40 mg olmesartan medoxomil and 3750 mg colesvelam hydrochloride in healthy subjects resulted in 28% reduction in C<sub>max</sub> and 39% reduction in AUC of olmesartan. Lesser effects, 4% and 15% reduction in C<sub>max</sub> and AUC respectively, were observed when olmesartan

medoxomil was administered 4 hours prior to colessevelam hydrochloride [see Drug Interactions (7)].

## 13 NONCLINICAL TOXICOLOGY

### 13.1 Carcinogenesis & Mutagenesis & Impairment Of Fertility

Olmесartan medoxomil was not carcinogenic when administered by dietary administration to rats for up to 2 years. The highest dose tested (2000 mg/kg/day) was, on a mg/m<sup>2</sup> basis, about 480 times the maximum recommended human dose (MRHD) of 40 mg/day. Two carcinogenicity studies conducted in mice, a 6-month gavage study in the p53 knockout mouse and a 6-month dietary administration study in the Hras2 transgenic mouse, at doses of up to 1000 mg/kg/day (about 120 times the MRHD), revealed no evidence of a carcinogenic effect of olmesartan medoxomil.

Both olmesartan medoxomil and olmesartan tested negative in the *in vitro* Syrian hamster embryo cell transformation assay and showed no evidence of genetic toxicity in the Ames (bacterial mutagenicity) test. However, both were shown to induce chromosomal aberrations in cultured cells *in vitro* (Chinese hamster lung) and tested positive for thymidine kinase mutations in the *in vitro* mouse lymphoma assay. Olmesartan medoxomil tested negative *in vivo* for mutations in the MutaMouse intestine and kidney and for clastogenicity in mouse bone marrow (micronucleus test) at oral doses of up to 2000 mg/kg (olmesartan not tested).

Fertility of rats was unaffected by administration of olmesartan medoxomil at dose levels as high as 1000 mg/kg/day (240 times the MRHD) in a study in which dosing was begun 2 (female) or 9 (male) weeks prior to mating.

### 13.2 Animal Pharmacology & OR Toxicology

#### Reproductive Toxicology Studies

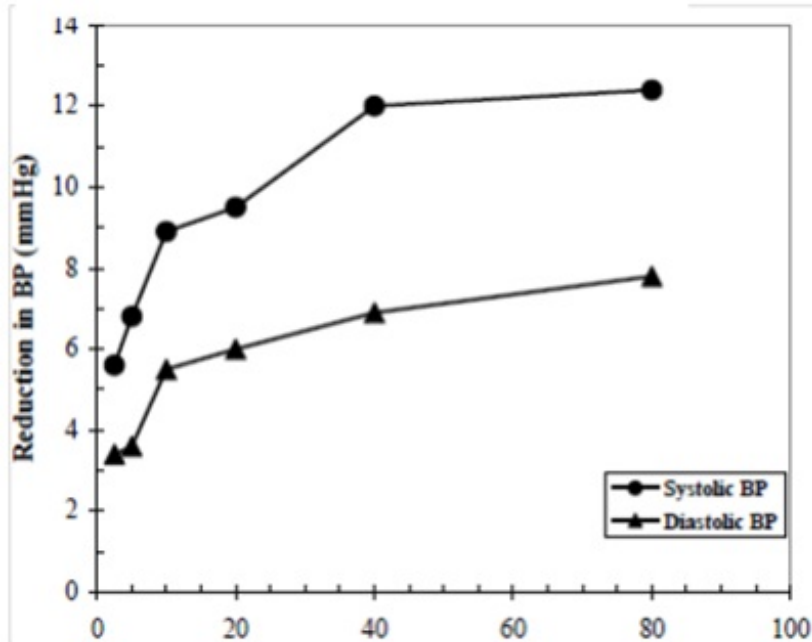
No teratogenic effects were observed when olmesartan medoxomil was administered to pregnant rats at oral doses up to 1000 mg/kg/day (240 times the maximum recommended human dose [MRHD] of olmesartan medoxomil on a mg/m<sup>2</sup> basis) or pregnant rabbits at oral doses up to 1 mg/kg/day (half the MRHD on a mg/m<sup>2</sup> basis; higher doses could not be evaluated for effects on fetal development as they were lethal to the does). In rats, significant decreases in pup birth weight and weight gain were observed at doses  $\geq 1.6$  mg/kg/day, and delays in developmental milestones (delayed separation of ear auricula, eruption of lower incisors, appearance of abdominal hair, descent of testes, and separation of eyelids) and dose-dependent increases in the incidence of dilation of the renal pelvis were observed at doses  $\geq 8$  mg/kg/day. The no observed effect dose for developmental toxicity in rats is 0.3 mg/kg/day, about one-tenth the MRHD of 40 mg/day.

## 14 CLINICAL STUDIES

### 14.1 Adult Hypertension

The antihypertensive effects of Olmesartan Medoxomil Tablets have been demonstrated in seven placebo-controlled studies at doses ranging from 2.5 mg to 80 mg for 6 to 12 weeks, each showing statistically significant reductions in peak and trough blood pressure. A total of 2693 patients (2145 Olmesartan Medoxomil Tablets ; 548 placebo) with essential hypertension were studied. Olmesartan Medoxomil Tablets once daily lowered diastolic and systolic blood pressure. The response was dose-related, as shown in the following graph. A Olmesartan Medoxomil Tablets dose of 20 mg daily produces a trough sitting BP reduction over placebo of about 10/6 mmHg and a dose of 40 mg daily produces a trough sitting BP reduction over placebo of about 12/7 mmHg. Olmesartan Medoxomil Tablets doses greater than 40 mg had little additional effect. The onset of the antihypertensive effect occurred within 1 week and was largely manifest after 2 weeks.

## Olmesartan Medoxomil Tablets Dose Response Placebo-Adjusted Reduction in Blood Pressure (mmHg)



### Olmesartan Medoxomil Tablets Dose (mg)

Data above are from seven placebo-controlled studies (2145 Olmesartan Medoxomil Tablets patients, 548 placebo patients). The blood pressure lowering effect was maintained throughout the 24-hour period with Olmesartan Medoxomil Tablets once daily, with trough-to-peak ratios for systolic and diastolic response between 60 and 80%.

The blood pressure lowering effect of Olmesartan Medoxomil Tablets, with and without hydrochlorothiazide, was maintained in patients treated for up to 1 year. There was no evidence of tachyphylaxis during long-term treatment with Olmesartan Medoxomil Tablets or rebound effect following abrupt withdrawal of olmesartan medoxomil after 1 year of treatment.

The antihypertensive effect of Olmesartan Medoxomil Tablets was similar in men and women and in patients older and younger than 65 years. The effect was smaller in black patients (usually a low-renin population), as has been seen with ACE inhibitors, beta-blockers and other angiotensin receptor blockers. Olmesartan Medoxomil Tablets had an additional blood pressure lowering effect when added to hydrochlorothiazide.

There are no trials of Olmesartan Medoxomil Tablets demonstrating reductions in cardiovascular risk in patients with hypertension, but at least one pharmacologically similar drug has demonstrated such benefits.

### 14.2 Pediatric Hypertension

The antihypertensive effects of Olmesartan Medoxomil Tablets in the pediatric population were evaluated in a randomized, double-blind study involving 302 hypertensive patients aged 6 to 16 years. The study population consisted of an all black cohort of 112 patients and a mixed racial cohort of 190 patients, including 38 blacks. The etiology of the hypertension was predominantly essential hypertension (87% of the black cohort and 67% of the mixed cohort). Patients who weighed 20 to <35 kg were randomized to 2.5 or 20 mg of Olmesartan Medoxomil Tablets once daily and patients who weighed  $\geq 35$  kg were randomized to 5 or 40 mg of Olmesartan Medoxomil Tablets once daily. At the end of 3 weeks, patients were re-randomized to continuing Olmesartan Medoxomil Tablets or to taking placebo for up to 2 weeks. During the initial dose-response phase, Olmesartan Medoxomil Tablets

significantly reduced both systolic and diastolic blood pressure in a weight-adjusted dose-dependent manner. Overall, the two dose levels of Olmesartan Medoxomil Tablets (low and high) significantly reduced systolic blood pressure by 6.6 and 11.9 mmHg from the baseline, respectively. These reductions in systolic blood pressure included both drug and placebo effect. During the randomized withdrawal to placebo phase, mean systolic blood pressure at trough was 3.2 mmHg lower and mean diastolic blood pressure at trough was 2.8 mmHg lower in patients continuing Olmesartan Medoxomil Tablets than in patients withdrawn to placebo. These differences were statistically different. As observed in adult populations, the blood pressure reductions were smaller in black patients.

In the same study, 59 patients aged 1 to 5 years who weighed  $\geq 5$  kg received 0.3 mg/kg of Olmesartan Medoxomil Tablets once daily for three weeks in an open label phase and then were randomized to receiving Olmesartan Medoxomil Tablets or placebo in a double-blind phase. At the end of the second week of withdrawal, the mean systolic/diastolic blood pressure at trough was 3/3 mmHg lower in the group randomized to Olmesartan Medoxomil Tablets ; this difference in blood pressure was not statistically significant (95% C.I. -2 to 7/-1 to 7).

## 16 HOW SUPPLIED/STORAGE AND HANDLING

Olmesartan Medoxomil Tablets is supplied as yellow coloured round shape film coated tablets containing 5 mg of olmesartan medoxomil USP, as white to off-white coloured, round shape film coated tablets containing 20 mg of olmesartan medoxomil USP, and as white to off-white coloured, oval shape film coated tablets containing 40 mg of olmesartan medoxomil USP. Tablets are debossed with OLM on one side and 5 ,20 & 40 on the other side of the 5, 20, and 40 mg tablets, respectively.

	5mg	20mg	40mg
Bottle of 30's	67877-445-30	67877-446-30	67877-447-30
Bottle of 90's	67877-445-90	67877-446-90	67877-447-90
Bottle of 500's	67877-445-05	67877-446-05	67877-447-05
Bottle of 1000's	67877-445-10	67877-446-10	67877-447-10
Carton pack of 10's (1x10's)	--	67877-446-33	--
Carton pack of 30's (3x10's)	--	67877-446-84	--
Carton pack of 50's (5x10's)	--	67877-446-74	--
Carton pack of 100's (10x10's)	--	67877-446-38	--

### Storage

Store at 20 to 25°C (68 to 77°F) [see USP Controlled Room Temperature].

## 17 PATIENT COUNSELING INFORMATION

**Pregnancy:** Female patients of childbearing age should be told about the consequences of exposure to Olmesartan Medoxomil Tablets during pregnancy. Discuss treatment options with women planning to become pregnant. Patients should be asked to report pregnancies to their physicians as soon as possible.

### Manufactured by:

Alkem Laboratories Ltd.

Mumbai - 400 013, INDIA

**Distributed by:**

Ascend Laboratories, LLC

Parsippany, NJ 07054

Revised: 03/2017

Rx Only

PT 2254-01

**PACKAGE LABEL.PRINCIPAL DISPLAY PANEL**

**Rx only**

**NDC 67877-445-30**

Olmesartan Medoxomil Tablets

5 mg

30 Tablets

Each tablet contains:  
Active ingredient: Olmesartan medoxomil, USP 5 mg.

Store at 20 to 25°C (68 to 77°F)  
[see USP Controlled Room Temperature]

**DOSAGE AND USE: SEE PACKAGE INSERT**

M. L. No.: MNB/05/105

 **Manufactured by:** Alkem Laboratories Ltd.,  
Mumbai - 400 013, INDIA.  
**Distributed by:** Ascend Laboratories, LLC Parsippany, NJ 07054

NDC 67877-445-30

**Olmesartan  
Medoxomil Tablets**

**5 mg**

Rx Only 30 Tablets



PL 9787-01

Unvarnished area  
40 x 20 mm (LXH)  
Rest label should be  
with UV Varnish

3 67877 44530 4

**Rx only**

**NDC 67877-446-84**

Olmesartan Medoxomil Tablets 20 mg

30 (3 x 10) Unit Dose Tablets





**Rx only**  
**NDC 67877-446-90**  
**Olmesartan Medoxomil Tablets 20 mg**  
**90 Tablets**



**Rx only**

**NDC 67877-447-30**

**Olmесartan Medoxomil Tablets 40 mg**

**30 Tablets**

Each tablet contains:  
Active ingredient: Olmesartan medoxomil, USP 40 mg.

Store at 20 to 25°C (68 to 77°F)  
[see USP Controlled Room Temperature]

**DOSAGE AND USE: SEE PACKAGE INSERT**

M. L. No.: MNB/05/105

**Manufactured by:** Alkem Laboratories Ltd.,  
Mumbai - 400 013, INDIA.  
**Distributed by:** Ascend Laboratories, LLC Parsippany, NJ 07054

NDC 67877-447-30

**Olmесartan Medoxomil Tablets**

**40 mg**

Rx Only 30 Tablets

PL 9795-01

Unvarnished area  
40 x 20 mm (LXH)  
Rest label should be  
with UV Varnish

3 67877 44730 8



## OLMESARTAN MEDOXOMIL

olmesartan medoxomil tablet, coated

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:67877-445
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OLMESARTAN MEDOXOMIL (UNII: 6M97XTV3HD) (OLMESARTAN - UNII:8W1IQP3U10)	OLMESARTAN MEDOXOMIL	5 mg

### Inactive Ingredients

Ingredient Name	Strength
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
HYDROXYPROPYL CELLULOSE, LOW SUBSTITUTED (UNII: 2165RE0K14)	
HYDROXYPROPYL CELLULOSE (TYPE L) (UNII: UKE75GEA7F)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
HYPROMELLOSE 2910 (6 MP.A.S) (UNII: 0WZ8WG20P6)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	

### Product Characteristics

<b>Color</b>	YELLOW	<b>Score</b>	no score
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<b>Shape</b>	ROUND (round shape)	<b>Size</b>	7mm
<b>Flavor</b>		<b>Imprint Code</b>	OLM;5
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67877-445-30	30 in 1 BOTTLE; Type 0: Not a Combination Product	04/24/2017	
2	NDC:67877-445-90	90 in 1 BOTTLE; Type 0: Not a Combination Product	04/24/2017	
3	NDC:67877-445-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	04/24/2017	
4	NDC:67877-445-10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	04/24/2017	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA206763	04/24/2017	

## OLMESARTAN MEDOXOMIL

olmesartan medoxomil tablet, coated

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:67877-446
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>OLMESARTAN MEDOXOMIL</b> (UNII: 6M97XTV3HD) (OLMESARTAN - UNII:8W1IQP3U10)	OLMESARTAN MEDOXOMIL	20 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>LACTOSE MONOHYDRATE</b> (UNII: EWQ57Q8I5X)	
<b>CELLULOSE, MICROCRYSTALLINE</b> (UNII: OP1R32D61U)	
<b>HYDROXYPROPYL CELLULOSE, LOW SUBSTITUTED</b> (UNII: 2165RE0K14)	
<b>HYDROXYPROPYL CELLULOSE (TYPE L)</b> (UNII: UKE75GEA7F)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>HYPROMELLOSE 2910 (6 MPA.S)</b> (UNII: 0WZ8WG20P6)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	

### Product Characteristics

<b>Color</b>	WHITE (white to off-white)	<b>Score</b>	no score
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<b>Shape</b>	ROUND (round shape)	<b>Size</b>	9mm
<b>Flavor</b>		<b>Imprint Code</b>	OLM;20
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67877-446-30	30 in 1 BOTTLE; Type 0: Not a Combination Product	04/24/2017	
2	NDC:67877-446-90	90 in 1 BOTTLE; Type 0: Not a Combination Product	04/24/2017	
3	NDC:67877-446-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	04/24/2017	
4	NDC:67877-446-10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	04/24/2017	
5	NDC:67877-446-33	1 in 1 CARTON	04/24/2017	
5		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
6	NDC:67877-446-84	3 in 1 CARTON	04/24/2017	
6		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
7	NDC:67877-446-74	5 in 1 CARTON	04/24/2017	
7		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
8	NDC:67877-446-38	10 in 1 CARTON	04/24/2017	
8		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA206763	04/24/2017	

## OLMESARTAN MEDOXOMIL

olmesartan medoxomil tablet, coated

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:67877-447
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>OLMESARTAN MEDOXOMIL</b> (UNII: 6M97XTV3HD) (OLMESARTAN - UNII:8W1IQP3U10)	OLMESARTAN MEDOXOMIL	40 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>LACTOSE MONOHYDRATE</b> (UNII: EWQ57Q8I5X)	
<b>CELLULOSE, MICROCRYSTALLINE</b> (UNII: OP1R32D61U)	
<b>HYDROXYPROPYL CELLULOSE, LOW SUBSTITUTED</b> (UNII: 2165RE0K14)	
<b>HYDROXYPROPYL CELLULOSE (TYPE L)</b> (UNII: UKE75GEA7F)	

<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>HYPROMELLOSE 2910 (6 MPA.S)</b> (UNII: 0WZ8WG20P6)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	

### Product Characteristics

<b>Color</b>	WHITE (white to off-white)	<b>Score</b>	no score
<b>Shape</b>	OVAL (oval shape)	<b>Size</b>	15mm
<b>Flavor</b>		<b>Imprint Code</b>	OLM;40
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67877-447-30	30 in 1 BOTTLE; Type 0: Not a Combination Product	04/24/2017	
2	NDC:67877-447-90	90 in 1 BOTTLE; Type 0: Not a Combination Product	04/24/2017	
3	NDC:67877-447-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	04/24/2017	
4	NDC:67877-447-10	1000 in 1 BOTTLE; Type 0: Not a Combination Product	04/24/2017	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA206763	04/24/2017	

**Labeler** - Ascend Laboratories, LLC (141250469)

### Establishment

Name	Address	ID/FEI	Business Operations
Alkem Laboratories Limited		677605851	MANUFACTURE(67877-445, 67877-446, 67877-447)

Revised: 4/2017

Ascend Laboratories, LLC