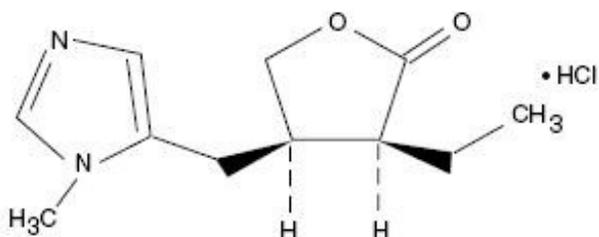


**PILOCARPINE HYDROCHLORIDE- pilocarpine hydrochloride tablet, film coated**  
**Amneal Pharmaceuticals of New York LLC**

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**Pilocarpine Hydrochloride Tablets**  
**(5 mg and 7.5 mg)**  
**Rx only**

**DESCRIPTION**

Pilocarpine hydrochloride tablets contain pilocarpine hydrochloride, a cholinergic agonist for oral use. Pilocarpine hydrochloride is a hygroscopic, odorless, bitter tasting white crystal or powder which is soluble in water and alcohol and virtually insoluble in most non-polar solvents. Pilocarpine hydrochloride, with a chemical name of (3*S*-*cis*)-2(3*H*)-Furanone, 3-ethyl-dihydro-4-[(1-methyl-1*H*-imidazol-5-yl) methyl] monohydrochloride, has a molecular weight of 244.72.



Each 5 mg pilocarpine hydrochloride tablet for oral administration contains 5 mg of pilocarpine hydrochloride. Inactive ingredients in the tablet and the tablet's film coating are: colloidal silicon dioxide, hypromellose Type 2910/ 3cP, 6cP, and 50cP, polyethylene glycol, microcrystalline cellulose, stearic acid, polydextrose, titanium dioxide, triacetate and triacetin.

Each 7.5 mg pilocarpine hydrochloride tablet for oral administration contains 7.5 mg of pilocarpine hydrochloride. Inactive ingredients in the tablet and the tablet's film coating are: colloidal silicon dioxide, FD&C blue #2/indigo carmine aluminum lake, hypromellose type 2910/ 3cP, and 6cP, polyethylene glycol, microcrystalline cellulose, stearic acid, polydextrose, titanium dioxide, triacetate and triacetin.

**CLINICAL PHARMACOLOGY**

**Pharmacodynamics**

Pilocarpine is a cholinergic parasympathomimetic agent exerting a broad spectrum of pharmacologic effects with predominant muscarinic action. Pilocarpine, in appropriate dosage, can increase secretion by the exocrine glands. The sweat, salivary, lacrimal, gastric, pancreatic, and intestinal glands and the mucous cells of the respiratory tract may be stimulated. When applied topically to the eye as a single-dose it causes miosis, spasm of accommodation, and may cause a transitory rise in intraocular pressure followed by a more persistent fall. Dose-related smooth muscle stimulation of the intestinal tract may cause increased tone, increased motility, spasm, and tenesmus. Bronchial smooth muscle tone may increase. The tone and motility of urinary tract, gallbladder, and biliary duct smooth muscle may be enhanced. Pilocarpine may have paradoxical effects on the cardiovascular system. The expected effect of a muscarinic agonist is vasodepression, but administration of pilocarpine may produce hypertension after a brief episode of hypotension. Bradycardia and tachycardia have both been reported with use of pilocarpine.

In a study of 12 healthy male volunteers there was a dose-related increase in unstimulated salivary flow

following single 5 mg and 10 mg oral doses of pilocarpine hydrochloride tablets. This effect of pilocarpine on salivary flow was time-related with an onset at 20 minutes and a peak effect at 1 hour with a duration of 3 to 5 hours (see **Pharmacokinetics** section).

**Head & Neck Cancer Patients:** In a 12 week randomized, double-blind, placebo-controlled study in 207 patients (placebo, N=65; 5 mg, N=73; 10 mg, N=69), increases from baseline (means 0.072 and 0.112 mL/min, ranges -0.690 to 0.728 and -0.380 to 1.689) of whole saliva flow for the 5 mg (63%) and 10 mg (90%) tablet, respectively, were seen 1 hour after the first dose of pilocarpine hydrochloride tablets. Increases in unstimulated parotid flow were seen following the first dose (means 0.025 and 0.046 mL/min, ranges 0 to 0.414 and -0.070 to 1.002 mL/min for the 5 mg and 10 mg dose, respectively). In this study, no correlation existed between the amount of increase in salivary flow and the degree of symptomatic relief.

**Sjogren's Syndrome Patients:** In two 12 week randomized, double-blind, placebo-controlled studies in 629 patients (placebo, N=253; 2.5 mg, N=121; 5 mg, N=255; 5 mg to 7.5 mg, N=114), the ability of pilocarpine hydrochloride tablets to stimulate saliva production was assessed. In these trials using varying doses of pilocarpine hydrochloride tablets (2.5 mg to 7.5 mg), the rate of saliva production was plotted against time. An Area Under the Curve (AUC) representing the total amount of saliva produced during the observation interval was calculated. Relative to placebo, an increase in the amount of saliva being produced was observed following the first dose of pilocarpine hydrochloride tablets and was maintained throughout the duration (12 weeks) of the trials in an approximate dose response fashion (see **Clinical Studies** section).

## **Pharmacokinetics**

In a multiple-dose pharmacokinetic study in male volunteers following 2 days of 5 mg or 10 mg of oral pilocarpine hydrochloride tablets given at 8 a.m., noontime, and 6 p.m., the mean elimination half-life was 0.76 hours for the 5 mg dose and 1.35 hours for the 10 mg dose.  $T_{max}$  values were 1.25 hours and 0.85 hours.  $C_{max}$  values were 15 ng/mL and 41 ng/mL. The AUC trapezoidal values were 33 h(ng/mL) and 108 h(ng/mL), respectively, for the 5 mg and 10 mg doses following the last 6 hour dose.

Pharmacokinetics in elderly male volunteers (N=11) were comparable to those in younger men. In five healthy elderly female volunteers, the mean  $C_{max}$  and AUC were approximately twice that of elderly males and young normal male volunteers.

When taken with a high fat meal by 12 healthy male volunteers, there was a decrease in the rate of absorption of pilocarpine from pilocarpine hydrochloride tablets. Mean  $T_{max}$ 's were 1.47 and 0.87 hours, and mean  $C_{max}$ 's were 51.8 and 59.2 ng/mL for fed and fasted, respectively.

Limited information is available about the metabolism and elimination of pilocarpine in humans. Inactivation of pilocarpine is thought to occur at neuronal synapses and probably in plasma. Pilocarpine and its minimally active or inactive degradation products, including pilocarpic acid, are excreted in the urine. Pilocarpine does not bind to human or rat plasma proteins over a concentration range of 5 to 25,000 ng/mL. The effect of pilocarpine on plasma protein binding of other drugs has not been evaluated.

In patients with mild to moderate hepatic impairment (N=12), administration of a single 5 mg dose resulted in a 30% decrease in total plasma clearance and a doubling of exposure (as measured by AUC). Peak plasma levels were also increased by about 30% and half-life was increased to 2.1 hrs.

There were no significant differences in the pharmacokinetics of oral pilocarpine in volunteer subjects (N=8) with renal insufficiency (mean creatinine clearances 25.4 mL/min; range 9.8 to 40.8 mL/min) compared to the pharmacokinetics previously observed in normal volunteers.

## **CLINICAL STUDIES**

### **Head & Neck Cancer Patients**

A 12 week randomized, double-blind, placebo-controlled study in 207 patients (142 men, 65 women) was conducted in patients whose mean age was 58.5 years with a range of 19 to 77; the racial distribution was Caucasian 95%, Black 4%, and other 1%. In this population, a statistically significant improvement in mouth dryness occurred in the 5 mg and 10 mg pilocarpine hydrochloride tablet treated patients compared to placebo treated patients. The 5 mg and 10 mg treated patients could not be distinguished (see **Pharmacodynamics** section for flow study details).

Another 12 week, double-blind, randomized, placebo-controlled study was conducted in 162 patients whose mean age was 57.8 years with a range of 27 to 80; the racial distribution was Caucasian 88%, Black 10%, and other 2%. The effects of placebo were compared to 2.5 mg three times a day of pilocarpine hydrochloride tablets for 4 weeks followed by adjustment to 5 mg three times a day and 10 mg three times a day. Lowering of the dose was necessary because of adverse events in 3 of 67 patients treated with 5 mg of pilocarpine hydrochloride tablets and in 7 of 66 patients treated with 10 mg of pilocarpine hydrochloride tablets. After 4 weeks of treatment, 2.5 mg of pilocarpine hydrochloride tablets three times a day was comparable to placebo in relieving dryness. In patients treated with 5 mg and 10 mg of pilocarpine hydrochloride tablets, the greatest improvement in dryness was noted in patients with no measurable salivary flow at baseline.

In both studies, some patients noted improvement in the global assessment of their dry mouth, speaking without liquids, and a reduced need for supplemental oral comfort agents.

In the two placebo-controlled clinical trials, the most common adverse events related to drug, and increasing in rate as dose increases, were sweating, nausea, rhinitis, diarrhea, chills, flushing, urinary frequency, dizziness, and asthenia. The most common adverse experience causing withdrawal from treatment was sweating (5 mg t.i.d. ≤ 1%; 10 mg t.i.d.=12%).

### **Sjogren's syndrome Patients**

Two separate studies were conducted in patients with primary or secondary Sjogren's Syndrome. In both studies, the majority of patients best fit the European criteria for having primary Sjogren's Syndrome. ["Criteria for the Classification of Sjogren's Syndrome" (Vitali C, Bombardieri S, Moutsopoulos HM, et al: Preliminary criteria for the classification of Sjogren's Syndrome. *Arthritis Rheum* 1993; 36:340-347.)]

A 12-week, randomized, double-blind, parallel-group, placebo-controlled study was conducted in 256 patients (14 men, 242 women) whose mean age was 57 years with a range of 24 to 85 years. The racial distribution was as follows: Caucasian 91%, Black 6%, and other 3%.

The effects of placebo were compared with those of pilocarpine hydrochloride tablets 5 mg four times a day (20 mg/day) for 6 weeks. At 6 weeks, the patients' dosage was increased from 5 mg pilocarpine hydrochloride tablets q.i.d. to 7.5 mg q.i.d. The data collected during the first 6 weeks of the trial were evaluated for safety and efficacy, and the data of the second 6 weeks of the trial were used to provide additional evidence of safety.

After 6 weeks of treatment, statistically significant global improvement of dry mouth was observed compared to placebo. "Global improvement" is defined as a score of 55 mm or more on a 100 mm visual analogue scale in response to the question, "Please rate your present condition of dry mouth (xerostomia) compared with your condition at the start of this study. Consider the changes to your dry mouth and other symptoms related to your dry mouth that have occurred since you have taken this medication." Patients' assessments of specific dry mouth symptoms such as severity of dry mouth, mouth discomfort, ability to speak without water, ability to sleep without drinking water, ability to swallow food without drinking, and a decreased use of saliva substitutes were found to be consistent with the significant global improvement described.

Another 12 week randomized, double-blind, parallel-group, placebo-controlled study was conducted in 373 patients (16 men, 357 women) whose mean age was 55 years with a range of 21 to 84. The racial distribution was Caucasian 80%, Oriental 14%, Black 2%, and 4% of other origin. The treatment groups

were 2.5 mg pilocarpine tablets, 5 mg pilocarpine hydrochloride tablets, and placebo. All treatments were administered on a four times a day regimen.

After 12 weeks of treatment, statistically significant global improvement of dry mouth was observed at a dose of 5 mg compared with placebo. The 2.5 mg (10 mg/day) group was not significantly different than placebo. However, a subgroup of patients with rheumatoid arthritis tended to improve in global assessments at both the 2.5 mg q.i.d. (9 patients) and 5 mg q.i.d. (16 patients) dose (10 mg/day to 20 mg/day). The clinical significance of this finding is unknown.

Patients' assessments of specific dry mouth symptoms such as severity of dry mouth, mouth discomfort, ability to sleep without drinking water, and decreased use of saliva substitutes were also found to be consistent with the significant global improvement described when measured after 6 weeks and 12 weeks of pilocarpine hydrochloride tablets use.

## **INDICATIONS AND USAGE**

Pilocarpine hydrochloride tablets are indicated for 1) the treatment of symptoms of dry mouth from salivary gland hypofunction caused by radiotherapy for cancer of the head and neck; and 2) the treatment of symptoms of dry mouth in patients with Sjogren's Syndrome.

## **CONTRAINDICATIONS**

Pilocarpine hydrochloride tablets are contraindicated in patients with uncontrolled asthma, known hypersensitivity to pilocarpine, and when miosis is undesirable, e.g., in acute iritis and in narrow-angle (angle closure) glaucoma.

## **WARNINGS**

### **Cardiovascular Disease**

Patients with significant cardiovascular disease may be unable to compensate for transient changes in hemodynamics or rhythm induced by pilocarpine. Pulmonary edema has been reported as a complication of pilocarpine toxicity from high ocular doses given for acute angle-closure glaucoma. Pilocarpine should be administered with caution in and under close medical supervision of patients with significant cardiovascular disease.

### **Ocular**

Ocular formulations of pilocarpine have been reported to cause visual blurring which may result in decreased visual acuity, especially at night and in patients with central lens changes, and to cause impairment of depth perception. Caution should be advised while driving at night or performing hazardous activities in reduced lighting.

### **Pulmonary Disease**

Pilocarpine has been reported to increase airway resistance, bronchial smooth muscle tone, and bronchial secretions. Pilocarpine hydrochloride should be administered with caution to and under close medical supervision in patients with controlled asthma, chronic bronchitis, or chronic obstructive pulmonary disease requiring pharmacotherapy.

## **PRECAUTIONS**

### **General**

Pilocarpine toxicity is characterized by an exaggeration of its parasympathomimetic effects. These may include: headache, visual disturbance, lacrimation, sweating, respiratory distress, gastrointestinal spasm,

nausea, vomiting, diarrhea, atrioventricular block, tachycardia, bradycardia, hypotension, hypertension, shock, mental confusion, cardiac arrhythmia, and tremors.

The dose-related cardiovascular pharmacologic effects of pilocarpine include hypotension, hypertension, bradycardia, and tachycardia.

Pilocarpine should be administered with caution to patients with known or suspected cholelithiasis or biliary tract disease. Contractions of the gallbladder or biliary smooth muscle could precipitate complications including cholecystitis, cholangitis, and biliary obstruction.

Pilocarpine may increase ureteral smooth muscle tone and could theoretically precipitate renal colic (or “ureteral reflux”), particularly in patients with nephrolithiasis.

Cholinergic agonists may have dose-related central nervous system effects. This should be considered when treating patients with underlying cognitive or psychiatric disturbances.

**Hepatic Insufficiency:** Based on decreased plasma clearance observed in patients with moderate hepatic impairment, the starting dose in these patients should be 5 mg twice daily, followed by adjustment based on therapeutic response and tolerability. Patients with mild hepatic insufficiency (Child-Pugh score of 5 to 6) do not require dosage reductions. To date, pharmacokinetic studies in subjects with severe hepatic impairment (Child-Pugh score of 10 to 15) have not been carried out. The use of pilocarpine in these patients is not recommended.

<b>Child-Pugh Scoring System for Hepatic Impairment</b>			
<b>Clinical and Biochemical Measurements</b>	<b>Points Scored for Increasing Abnormality</b>		
	<b>1</b>	<b>2</b>	<b>3</b>
Encephalopathy (grade)*	None	1 and 2	3 and 4
Ascites	Absent	Slight	Moderate
Bilirubin (mg per 100 mL)	1 to 2	2 to 3	> 3
Albumin (g per 100 mL)	3 to 5	2.8 to 3.5	< 2.8
Prothrombin Time (sec. Prolonged)	1 to 4	4 to 6	> 6
For Primary Biliary Cirrhosis:-Bilirubin (mg per 100 mL)	1 to 4	4 to 10	> 10

\*According to grading of Trey C, Burns D, and Saunders S. Treatment of hepatic coma by exchange blood transfusion. *N Engl J Med.* 1966; 274:473-481.

Reference: Pugh RNH, Murray-Lyon IM, Dawson JL Pietroni MC, Williams, R. Transection of the oesophagus for bleeding oesophageal varices, *Brit. J. Surg.* 1973; 60:646-9.

### **Information for Patients**

Patients should be informed that pilocarpine may cause visual disturbances, especially at night, that could impair their ability to drive safely.

If a patient sweats excessively while taking pilocarpine hydrochloride and cannot drink enough liquid, the patient should consult a physician. Dehydration may develop.

### **Drug Interactions**

Pilocarpine should be administered with caution to patients taking beta-adrenergic antagonists because of the possibility of conduction disturbances. Drugs with parasympathomimetic effects administered concurrently with pilocarpine would be expected to result in additive pharmacologic effects. Pilocarpine might antagonize the anticholinergic effects of drugs used concomitantly. These effects should be considered when anticholinergic properties may be contributing to the therapeutic effect of concomitant medication (e.g., atropine, inhaled ipratropium).

While no formal drug interaction studies have been performed, the following concomitant drugs were used in at least 10% of patients in either or both Sjogren's efficacy studies: acetylsalicylic acid, artificial tears, calcium, conjugated estrogens, hydroxychloroquine sulfate, ibuprofen, levothyroxine sodium, medroxyprogesterone acetate, methotrexate, multivitamins, naproxen, omeprazole, paracetamol, and prednisone.

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

Lifetime oral carcinogenicity studies were conducted in CD-1 mice and Sprague-Dawley rats. Pilocarpine did not induce tumors in mice at any dosage studied (up to 30 mg/kg/day, which yielded a systemic exposure approximately 50 times larger than the maximum systemic exposure observed clinically). In rats, a dosage of 18 mg/kg/day, which yielded a systemic exposure approximately 100 times larger than the maximum systemic exposure observed clinically, resulted in a statistically significant increase in the incidence of benign pheochromocytomas in both males and females, and a statistically significant increase in the incidence of hepatocellular adenomas in female rats. The tumorigenicity observed in rats was observed only at a large multiple of the maximum labeled clinical dose, and may not be relevant to clinical use.

No evidence that pilocarpine has the potential to cause genetic toxicity was obtained in a series of studies that included: 1) bacterial assays (*Salmonella* and *E. coli*) for reverse gene mutations; 2) an *in vitro* chromosome aberration assay in a Chinese hamster ovary cell line; 3) an *in vivo* chromosome aberration assay (micronucleus test) in mice; and 4) a primary DNA damage assay (unscheduled DNA synthesis) in rat hepatocyte primary cultures.

Oral administration of pilocarpine to male and female rats at a dosage of 18 mg/kg/day, which yielded a systemic exposure approximately 100 times larger than the maximum systemic exposure observed clinically, resulted in impaired reproductive function, including reduced fertility, decreased sperm motility, and morphologic evidence of abnormal sperm. It is unclear whether the reduction in fertility was due to effects on male animals, female animals, or both males and females. In dogs, exposure to pilocarpine at a dosage of 3 mg/kg/day (approximately 3 times the maximum recommended human dose when compared on the basis of body surface area ( $\text{mg}/\text{m}^2$ ) estimates) for six months resulted in evidence of impaired spermatogenesis. The data obtained in these studies suggest that pilocarpine may impair the fertility of male and female humans. Pilocarpine hydrochloride tablets should be administered to individuals who are attempting to conceive a child only if the potential benefit justifies potential impairment of fertility.

### **Pregnancy: Teratogenic Effects**

**Pregnancy Category C:** Pilocarpine was associated with a reduction in the mean fetal body weight and an increase in the incidence of skeletal variations when given to pregnant rats at a dosage of 90 mg/kg/day (approximately 26 times the maximum recommended dose for a 50 kg human when compared on the basis of body surface area ( $\text{mg}/\text{m}^2$ ) estimates). These effects may have been secondary to maternal toxicity. In another study, oral administration of pilocarpine to female rats during gestation and lactation at a dosage of 36 mg/kg/day (approximately 10 times the maximum recommended dose for a 50 kg human when compared on the basis of body surface area ( $\text{mg}/\text{m}^2$ ) estimates) resulted in an increased incidence of stillbirths; decreased neonatal survival and reduced mean body weight of pups were observed at dosages of 18 mg/kg/day (approximately 5 times the maximum recommended dose for a 50 kg human when compared on the basis of body surface area ( $\text{mg}/\text{m}^2$ ) estimates) and above. There are no adequate and well-controlled studies in pregnant women. Pilocarpine hydrochloride tablets should be used during pregnancy only if the potential benefit justifies the potential risk to the fetus.

### **Nursing Mothers**

It is not known whether this drug is excreted in human milk. Because many drugs are excreted in human milk and because of the potential for serious adverse reactions in nursing infants from pilocarpine hydrochloride tablets, a decision should be made whether to discontinue nursing or to discontinue the

drug, taking into account the importance of the drug to the mother.

### Pediatric Use

Safety and effectiveness in pediatric patients have not been established.

### Geriatric Use

**Head and Neck Cancer Patients:** In the placebo-controlled clinical trials (see **Clinical Studies** section) the mean age of patients was approximately 58 years (range 19 to 80). Of these patients, 97/369 (61/217 receiving pilocarpine) were over the age of 65 years. In the healthy volunteer studies, 15/150 subjects were over the age of 65 years. In both study populations, the adverse events reported by those over 65 years and those 65 years and younger were comparable. Of the 15 elderly volunteers (5 women, 10 men), the 5 women had higher  $C_{max}$ 's and AUC's than the men (see **Pharmacokinetics** section).

**Sjogren's Syndrome Patients:** In the placebo-controlled clinical trials (see **Clinical Studies** section), the mean age of patients was approximately 55 years (range 21 to 85). The adverse events reported by those over 65 years and those 65 years and younger were comparable except for notable trends for urinary frequency, diarrhea, and dizziness (see **ADVERSE REACTIONS** section).

## ADVERSE REACTIONS

### Head & Neck Cancer Patients

In controlled studies, 217 patients received pilocarpine, of whom 68% were men and 32% were women. Race distribution was 91% Caucasian, 8% Black, and 1% of other origin. Mean age was approximately 58 years. The majority of patients were between 50 and 64 years (51%), 33% were 65 years and older and 16% were younger than 50 years of age. The most frequent adverse experiences associated with pilocarpine hydrochloride tablets were a consequence of the expected pharmacologic effects of pilocarpine.

Adverse Event	Pilocarpine Hydrochloride		Placebo (t.i.d.) N=152
	10 mg t.i.d. (30 mg/day) N=121	5 mg t.i.d. (15 mg/day) N=141	
Sweating	68%	29%	9%
Nausea	15	6	4
Rhinitis	14	5	7
Diarrhea	7	4	5
Chills	15	3	< 1
Flushing	13	8	3
Urinary Frequency	12	9	7
Dizziness	12	5	4
Asthenia	12	6	3

In addition, the following adverse events ( $\geq 3\%$  incidence) were reported at dosages of 15 mg/day to 30 mg/day in the controlled clinical trials:

Adverse Event	Pilocarpine Hydrochloride		Placebo (t.i.d.) N=152
	5 mg to 10 mg t.i.d. (15 mg/day to 30 mg/day) N=212		
Headache	11%		8%

Dyspepsia	7	5
Lacrimation	6	8
Edema	5	4
Abdominal Pain	4	4
Amblyopia	4	2
Vomiting	4	1
Pharyngitis	3	8
Hypertension	3	1

The following events were reported with treated head and neck cancer patients at incidences of 1% to 2% at dosages of 7.5 mg/day to 30 mg/day: abnormal vision, conjunctivitis, dysphagia, epistaxis, myalgias, pruritus, rash, sinusitis, tachycardia, taste perversion, tremor, voice alteration.

The following events were reported rarely in treated head and neck cancer patients (< 1%): Causal relation is unknown.

Body as a whole: body odor, hypothermia, mucous membrane abnormality

Cardiovascular: bradycardia, ECG abnormality, palpitations, syncope

Digestive: anorexia, increased appetite, esophagitis, gastrointestinal disorder, tongue disorder

Hematologic: leukopenia, lymphadenopathy

Nervous: anxiety, confusion, depression, abnormal dreams, hyperkinesia, hypesthesia, nervousness, paresthesias, speech disorder, twitching

Respiratory: increased sputum, stridor, yawning

Skin: seborrhea

Special senses: deafness, eye pain, glaucoma

Urogenital: dysuria, metrorrhagia, urinary impairment

In long-term treatment were two patients with underlying cardiovascular disease of whom one experienced a myocardial infarct and another an episode of syncope. The association with drug is uncertain.

### Sjogren's syndrome Patients

In controlled studies, 376 patients received pilocarpine, of whom 5% were men and 95% were women. Race distribution was 84% Caucasian, 9% Oriental, 3% Black, and 4% of other origin. Mean age was 55 years. The majority of patients were between 40 and 69 years (70%), 16% were 70 years and older and 14% were younger than 40 years of age. Of these patients, 161/629 (89/376 receiving pilocarpine) were over the age of 65 years. The adverse events reported by those over 65 years and those 65 years and younger were comparable except for notable trends for urinary frequency, diarrhea, and dizziness. The incidences of urinary frequency and diarrhea in the elderly were about double those in the non-elderly. The incidence of dizziness was about three times as high in the elderly as in the non-elderly. These adverse experiences were not considered to be serious. In the 2 placebo-controlled studies, the most common adverse events related to drug use were sweating, urinary frequency, chills, and vasodilatation (flushing). The most commonly reported reason for patient discontinuation of treatment was sweating. Expected pharmacologic effects of pilocarpine include the following adverse experiences associated with pilocarpine hydrochloride tablets:

Adverse Event	Pilocarpine hydrochloride	Placebo
	5 mg q.i.d. (20 mg/day) N=255	(q.i.d.) N=253

Sweating	40%	7%
Urinary Frequency	10	4
Nausea	9	9
Flushing	9	2
Rhinitis	7	8
Diarrhea	6	7
Chills	4	2
Increased Salivation	3	0
Asthenia	2	2

In addition, the following adverse events ( $\geq 3\%$  incidence) were reported at dosages of 20 mg/day in the controlled clinical trials:

Adverse Event	Pilocarpine hydrochloride 5 mg q.i.d. (20 mg/day) N=255	Placebo (q.i.d.) N=253
Headache	13%	19%
Flu Syndrome	9	9
Dyspepsia	7	7
Dizziness	6	7
Pain	4	2
Sinusitis	4	5
Abdominal Pain	3	4
Vomiting	3	1
Pharyngitis	2	5
Rash	2	3
Infection	2	6

The following events were reported in Sjogren's patients at incidences of 1% to 2% at dosing of 20 mg/day: accidental injury, allergic reaction, back pain, blurred vision, constipation, increased cough, edema, epistaxis, face edema, fever, flatulence, glossitis, lab test abnormalities, including chemistry, hematology, and urinalysis, myalgia, palpitation, pruritus, somnolence, stomatitis, tachycardia, tinnitus, urinary incontinence, urinary tract infection, and vaginitis.

The following events were reported rarely in treated Sjogren's patients ( $< 1\%$ ) at dosing of 10 mg/day to 30 mg/day: Causal relation is unknown.

Body as a whole: chest pain, cyst, death, moniliasis, neck pain, neck rigidity, photosensitivity reaction

Cardiovascular: angina pectoris, arrhythmia, ECG abnormality, hypotension, hypertension, intracranial hemorrhage, migraine, myocardial infarction

Digestive: anorexia, bilirubinemia, cholelithiasis, colitis, dry mouth, eructation, gastritis, gastroenteritis, gastrointestinal disorder, gingivitis, hepatitis, abnormal liver function tests, melena, nausea & vomiting, pancreatitis, parotid gland enlargement, salivary gland enlargement, sputum increased, taste loss, tongue disorder, tooth disorder

Hematologic: hematuria, lymphadenopathy, abnormal platelets, thrombocytopenia, thrombocytopenia, thrombosis, abnormal WBC

Metabolic and Nutritional: peripheral edema, hypoglycemia

Musculoskeletal: arthralgia, arthritis, bone disorder, spontaneous bone fracture, pathological fracture, myasthenia, tendon disorder, tenosynovitis

Nervous: aphasia, confusion, depression, abnormal dreams, emotional lability, hyperkinesia, hypesthesia, insomnia, leg cramps, nervousness, paresthesias, abnormal thinking, tremor

Respiratory: bronchitis, dyspnea, hiccup, laryngismus, laryngitis, pneumonia, viral infection, voice alteration

Skin: alopecia, contact dermatitis, dry skin, eczema, erythema nodosum, exfoliative dermatitis, herpes simplex, skin ulcer, vesiculobullous rash

Special Senses: cataract, conjunctivitis, dry eyes, ear disorder, ear pain, eye disorder, eye hemorrhage, glaucoma, lacrimation disorder, retinal disorder, taste perversion, abnormal vision

Urogenital: breast pain, dysuria, mastitis, menorrhagia, metrorrhagia, ovarian disorder, pyuria, salpingitis, urethral pain, urinary urgency, vaginal hemorrhage, vaginal moniliasis

The following adverse experiences have been reported rarely with ocular pilocarpine: A-V block, agitation, ciliary congestion, confusion, delusion, depression, dermatitis, middle ear disturbance, eyelid twitching, malignant glaucoma, iris cysts, macular hole, shock, and visual hallucination.

**To report SUSPECTED ADVERSE REACTIONS, contact Amneal Pharmaceuticals at 1-877-835-5472 or FDA at 1-800-FDA-1088 or [www.fda.gov/medwatch](http://www.fda.gov/medwatch).**

## MANAGEMENT OF OVERDOSE

Fatal overdosage with pilocarpine has been reported in the scientific literature at doses presumed to be greater than 100 mg in two hospitalized patients. 100 mg of pilocarpine is considered potentially fatal. Overdosage should be treated with atropine titration (0.5 mg to 1 mg given subcutaneously or intravenously) and supportive measures to maintain respiration and circulation. Epinephrine (0.3 mg to 1 mg, subcutaneously or intramuscularly) may also be of value in the presence of severe cardiovascular depression or bronchoconstriction. It is not known if pilocarpine is dialyzable.

## DOSAGE AND ADMINISTRATION

Regardless of the indication, the starting dose in patients with moderate hepatic impairment should be 5 mg twice daily, followed by adjustment based on therapeutic response and tolerability. Patients with mild hepatic insufficiency do not require dosage reductions. The use of pilocarpine in patients with severe hepatic insufficiency is not recommended. If needed, refer to the *Hepatic Insufficiency* subsection of the **Precautions** section of this label for definitions of mild, moderate and severe hepatic impairment.

### Head & Neck Cancer Patients

The recommended initial dose of pilocarpine hydrochloride tablets is 5 mg taken three times a day. Dosage should be titrated according to therapeutic response and tolerability. The usual dosage range is up to 15 mg per day to 30 mg per day. (Not to exceed 10 mg per dose.) Although early improvement may be realized, at least 12 weeks of uninterrupted therapy with pilocarpine hydrochloride tablets may be necessary to assess whether a beneficial response will be achieved. The incidence of the most common adverse events increases with dose. The lowest dose that is tolerated and effective should be used for maintenance.

### Sjogren's syndrome Patients

The recommended dose of pilocarpine hydrochloride tablets is one tablet (5 mg) taken four times a day. Efficacy was established by 6 weeks of use.

## HOW SUPPLIED

Pilocarpine Hydrochloride Tablets, 5 mg—Each white to off-white, film-coated, round convex tablets, debossed with “G” on one side and “592” on the other side. They are available as follows:

Bottles of 100: NDC 0115-5922-01  
Bottles of 1000: NDC 0115-5922-03

Pilocarpine Hydrochloride Tablets, 7.5 mg—Each blue, film-coated, round convex tablets, debossed with “G” on one side and “591” on the other side. They are available as follows:

Bottles of 100 : NDC 0115-5911-01  
Bottles of 1000: NDC 0115-5911-03

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

Dispense in a tightly-closed, light-resistant container (USP).

Manufactured by:  
**Amneal Pharmaceuticals of NY, LLC**  
Brookhaven, NY 11719

Distributed by:  
**Amneal Pharmaceuticals LLC**  
Bridgewater, NJ 08807

Rev. 01-2019-00

## PRINCIPAL DISPLAY PANEL

NDC 0115-5922-01

**Pilocarpine Hydrochloride Tablets, 5 mg**

**Rx Only**

**100 Tablets**

**Amneal Pharmaceuticals LLC**

<p>NDC 0115-5922-01</p> <p><b>Pilocarpine Hydrochloride Tablets</b></p> <p><b>5 mg</b></p> <p>Rx only 100 Tablets</p> 	<p><b>DOSAGE AND ADMINISTRATION:</b> See accompanying outsert for complete prescribing information.</p> <p><b>Store at 20° to 25°C (68° to 77°F) [see USP Controlled Room Temperature].</b></p> <p>Dispense in a tightly-closed, light-resistant container as defined in the USP, with a child-resistant closure, as required.</p> <p>Manufactured by: <b>Amneal Pharmaceuticals of NY, LLC</b> Brookhaven, NY 11719</p> <p>Distributed by: <b>Amneal Pharmaceuticals LLC</b> Bridgewater, NJ 08807</p> <p>Rev. 01-2019-00</p>	 <p>3 01155 92201 9</p>	<p>Non-Varnish Area (For Lot And Exp. Date) 1.125" x 1.063" inch</p>
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## PRINCIPAL DISPLAY PANEL

NDC 0115-5911-01

**Pilocarpine Hydrochloride Tablets, 7.5 mg**

**Rx Only**

100 Tablets

Amneal Pharmaceuticals LLC

NDC 0115-5911-01

# Pilocarpine Hydrochloride Tablets

7.5 mg

Rx only  
100 Tablets



**DOSAGE AND ADMINISTRATION:** See accompanying outsert for complete prescribing information.

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

Dispense in a tightly-closed, light-resistant container as defined in the USP, with a child-resistant closure, as required.

Manufactured by: **Amneal Pharmaceuticals of NY, LLC**  
Brookhaven, NY 11719

Distributed by: **Amneal Pharmaceuticals LLC**  
Bridgewater, NJ 08807

Rev. 01-2019-00



Non-Varnish Area  
(For Lot And Exp. Date)  
0.7 x 1 inch

## PILOCARPINE HYDROCHLORIDE

pilocarpine hydrochloride tablet, film coated

### Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:0115-5922
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PILOCARPINE HYDROCHLORIDE (UNII: 0 WW6D218 XJ) (PILOCARPINE - UNII:01MI4Q9DI3)	PILOCARPINE HYDROCHLORIDE	5 mg

### Inactive Ingredients

Ingredient Name	Strength
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	

### Product Characteristics

Color	WHITE (white to off-white)	Score	no score
Shape	ROUND (round, convex)	Size	6mm
Flavor		Imprint Code	G;592
Contains			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0115-5922-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2007	
2	NDC:0115-5922-03	1000 in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2007	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077248	05/01/2007	

## PILOCARPINE HYDROCHLORIDE

pilocarpine hydrochloride tablet, film coated

### Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:0115-5911
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PILOCARPINE HYDROCHLORIDE (UNII: 0WW6D218XJ) (PILOCARPINE - UNII:01M4Q9D13)	PILOCARPINE HYDROCHLORIDE	7.5 mg

### Inactive Ingredients

Ingredient Name	Strength
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	

### Product Characteristics

Color	BLUE	Score	no score
Shape	ROUND (round, convex)	Size	6mm
Flavor		Imprint Code	G;591
Contains			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0115-5911-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2007	
2	NDC:0115-5911-03	1000 in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2007	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077248	05/01/2007	

**Labeler** - Amneal Pharmaceuticals of New York LLC (123797875)

**Establishment**

Name	Address	ID/FEI	Business Operations
Amneal Pharmaceuticals of New York, LLC		123797875	ANALYSIS(0115-5922, 0115-5911) , MANUFACTURE(0115-5922, 0115-5911) , PACK(0115-5922, 0115-5911)

**Establishment**

Name	Address	ID/FEI	Business Operations
Amneal Pharmaceuticals, LLC		969951594	PACK(0115-5922, 0115-5911) , REPACK(0115-5922, 0115-5911)

Revised: 6/2019

Amneal Pharmaceuticals of New York LLC