**HIGHLIGHTS OF PRESCRIBING INFORMATION**

These highlights do not include all the information needed to use LONHALA® MAGNIR® safely and effectively. See full prescribing information for LONHALA MAGNIR.

**LONHALA MAGNIR (glycopyrrolate) inhalation solution, for oral inhalation use**

**Initial U.S. Approval: 1961**

**INDICATIONS AND USAGE**

LONHALA MAGNIR is an anticholinergic indicated for the long-term, maintenance treatment of airflow obstruction in patients with chronic obstructive pulmonary disease (COPD). (1)

- **For oral inhalation only. Do not swallow LONHALA solution. Only use LONHALA vials with MAGNIR.** (2)
- **Maintenance treatment of COPD: The contents of one LONHALA vial twice-daily.** (2)

LONHALA Inhalation Solution is supplied as a sterile solution for inhalation in a unit-dose single-use low-density polyethylene (LDPE) vial. Each 1 mL vial contains 25 mcg of glycopyrrolate. (3)

**CONTRAINDICATIONS**

LONHALA MAGNIR is contraindicated in patients with a hypersensitivity to glycopyrrolate or any of the ingredients. (4)

**WARNINGS AND PRECAUTIONS**

- **Do not initiate in acutely deteriorating COPD or to treat acute symptoms.** (5.1)
- **If paradoxical bronchospasm occurs, discontinue LONHALA MAGNIR immediately and institute alternative therapy.** (5.2)
- **Worsening of narrow-angle glaucoma may occur. Use with caution in patients with narrow-angle glaucoma and instruct patients to contact a physician immediately if symptoms occur.** (5.4)
- **Worsening of urinary retention may occur. Use with caution in patients with prostatic hyperplasia or bladder neck obstruction and instruct patients to consult a physician immediately if symptoms occur.** (5.5)

**ADVERSE REACTIONS**

- **Most common adverse reactions (incidence greater than or equal to 2.0% and higher than placebo) are dyspnea and urinary tract infection.** (6.1)

To report SUSPECTED ADVERSE REACTIONS, contact Sunovion Pharmaceuticals Inc. at 1-877-737-7226 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

**DRUG INTERACTIONS**

Anticholinergics: May interact additively with concomitantly used anticholinergic medications. Avoid administration of LONHALA MAGNIR with other anticholinergic-containing drugs. (7.2)

**USE IN SPECIFIC POPULATIONS**

Use in patients with severe renal impairment should be considered if the potential benefit of the treatment outweighs the risk. (8.6)

See 17 for PATIENT COUNSELING INFORMATION and FDA-approved patient labeling.

Revised: 06/2019

---

**FULL PRESCRIBING INFORMATION: CONTENTS**

1. **INDICATIONS AND USAGE**
2. **DOSE AND ADMINISTRATION**
3. **DOSE FORMS AND STRENGTHS**
4. **CONTRAINDICATIONS**
5. **WARNINGS AND PRECAUTIONS**
6. **ADVERSE REACTIONS**
7. **DRUG INTERACTIONS**
8. **USE IN SPECIFIC POPULATIONS**
9. **OVERDOSAGE**
10. **DESCRIPTION**
11. **CLINICAL PHARMACOLOGY**
12. **CLINICAL PHARMACOLOGY**
13. **NONCLINICAL TOXICOLOGY**
14. **CLINICAL STUDIES**
15. **HOW SUPPLIED/STORAGE AND HANDLING**
16. **PATIENT COUNSELING INFORMATION**

*Sections or subsections omitted from the full prescribing information are not listed.*
5.3 Immediate Hypersensitivity Reactions

Immediate hypersensitivity reactions may occur after administration of LONHALA MAGNAIR. If signs suggesting allergic reactions occur, in particular, angioedema (including difficulties in breathing or swallowing, swelling of the tongue, lips, and face), urticaria, or skin rash, LONHALA MAGNAIR should be discontinued immediately and alternative therapy instituted.

5.4 Worsening of Narrow-Angle Glaucoma

LONHALA MAGNAIR should be used with caution in patients with narrow-angle glaucoma. Prescribers and patients should be alert for signs and symptoms of acute narrow-angle glaucoma (e.g., eye pain or discomfort, blurred vision, visual halos or colored images in association with red eyes from conjunctival congestion and corneal edema). Instruct patients to consult a physician immediately should any of these signs or symptoms develop.

5.5 Worsening of Urinary Retention

LONHALA MAGNAIR should be used with caution in patients with urinary retention. Prescribers and patients should be alert for signs and symptoms of urinary retention (e.g., difficulty passing urine, painful urination), especially in patients with prostatic hyperplasia or bladder-neck obstruction. Instruct patients to consult a physician immediately should any of these signs or symptoms develop.

6. ADVERSE REACTIONS

The following adverse reactions are described in greater detail in other sections:

- Paradoxical bronchospasm [see Warnings and Precautions (5.2)]
- Immediate hypersensitivity reactions [see Warnings and Precautions (5.3)]
- Worsening of narrow-angle glaucoma [see Warnings and Precautions (5.4)]
- Worsening of urinary retention [see Warnings and Precautions (5.5)]

6.1 Clinical Trials Experience

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

The LONHALA MAGNAIR safety database included 2379 subjects with COPD in two 12-week efficacy studies and one 48-week long-term safety study. A total of 431 subjects received treatment with LONHALA MAGNAIR 25 mcg twice-daily (BID). The safety data described below are based on the two 12-week trials and the one 48-week trial.

12-Week Trials

LONHALA MAGNAIR was studied in two 12-week placebo-controlled trials in subjects with COPD. In these trials, 431 subjects were treated with LONHALA MAGNAIR at the recommended dose of 25 mcg twice daily. The population had a mean age of 63 years (ranging from 40 to 87 years), with 56% males, 90% Caucasian, and a mean post-bronchodilator forced expiratory volume in one second (FEV1) percent predicted of 52% of predicted normal value (20%-80%) at study entry. The study population also included subjects with pre-existing cardiovascular disease as well as subjects with continued use of stable long-acting bronchodilator (LABA) +/- inhaled corticosteroid (ICS) and ipratropium bromide background therapy. Subjects with unstable cardiac disease, narrow-angle glaucoma, or symptomatic prostatic hypertrophy or bladder outlet obstruction were excluded from these studies.

Table 1 shows the most common adverse reactions incidence greater than or equal to 2.0% in the LONHALA MAGNAIR group and higher than placebo in the two 12-week placebo-controlled trials.

The proportion of subjects who discontinued treatment due to adverse reactions was 5% for the LONHALA MAGNAIR-treated subjects and 9% for placebo-treated subjects.

<table>
<thead>
<tr>
<th>Table 1. Adverse Reactions with LONHALA MAGNAIR</th>
<th>≥2.0% Incidence and Higher than Placebo</th>
</tr>
</thead>
<tbody>
<tr>
<td>Placebo</td>
<td>LONHALA MAGNAIR 25 mcg BID</td>
</tr>
<tr>
<td></td>
<td>(N=430)</td>
</tr>
<tr>
<td>Dyspnea</td>
<td>13 (3.0)</td>
</tr>
<tr>
<td>Urinary Tract Infection</td>
<td>6 (1.4)</td>
</tr>
</tbody>
</table>

Other adverse reactions defined as events with an incidence of ≥1.0% but less than 2.0% with LONHALA MAGNAIR but more common than with placebo included the following: wheezing, upper respiratory tract infection, nasopharyngitis, oedema peripheral, and fatigue.

48-Week Trial

In a long-term open-label safety trial, 1086 subjects were treated for up to 48-weeks with LONHALA MAGNAIR 50 mcg twice-daily (N=620) or tiotropium (N=466). The demographic and baseline characteristics of the long-term safety trial were similar to those of the placebo-controlled efficacy studies described above.

The adverse reactions reported in the long-term safety trial were consistent with those observed in the placebo-controlled studies of 12-weeks. Adverse reactions that occurred at a frequency greater than that seen in either active treatment dose in the pooled 12-week placebo controlled studies and ≥ 2.0% were: diarrhea, edema peripheral, bronchitis, nasopharyngitis, pneumonia, sinusitis, upper respiratory tract infection, urinary tract infection, back pain, headache, Chronic Obstructive Pulmonary Disease, cough, dyspnea, ophryopathy, pain, and hypertension.

7. DRUG INTERACTIONS

7.1 Sympathomimetics and Steroids

In clinical studies, concurrent administration of glycopyrrolate and other drugs commonly used in the treatment of COPD including sympathomimetics (long and short-acting beta2 agonists), anticholinergics (short-acting anti-muscarinic antagonists) and oral and inhaled steroids showed no increases in adverse drug reactions.

7.2 Anticholinergics

There is a potential for an additive interaction with concomitantly used anticholinergic medications. Therefore, avoid unnecessary co-administration of LONHALA MAGNAIR with other anticholinergic-containing drugs as this may lead to an increase in anticholinergic effects [see Warnings and Precautions (5.4, 5.5) and Adverse Reactions (6)].

8. USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

There are no adequate and well-controlled studies in pregnant women. LONHALA MAGNAIR should only be used during pregnancy if the expected benefit to the patient outweighs the potential risk to the fetus. Women should be advised to contact their physician if they become pregnant while taking LONHALA MAGNAIR. In animal reproduction studies, there were no teratogenic effects in Wistar rats and New Zealand White rabbits at inhaled doses approximating 1521 and 580 times, respectively, the maximum recommended human daily inhalation dose (MRHDID) based on an AUC comparison (see Data).

The estimated background risk of major birth defects and miscarriage for the indicated population is unknown. In the U.S. general population, the estimated background risk of major birth defects and miscarriage in clinically recognized pregnancies is 2-4% and 15-20%, respectively.

Labor or Delivery

The potential effect of LONHALA MAGNAIR on labor and delivery is unknown. LONHALA MAGNAIR should be used during labor and delivery only if the potential benefit to the patient justifies the potential risk to the fetus.

Data

Animal Data

Developmental studies in Wistar rats and New Zealand White rabbits in which glycopyrrolate was administered by inhalation during the period of organogenesis did not result in evidence of teratogenicity at exposures approximately 1521 and 580 times, respectively, the MRHDID of LONHALA MAGNAIR based on a comparison of plasma AUC levels (maternal doses up to 3.8 mg/kg/day in rats and 4.4 mg/kg/day in rabbits).

Glycopyrrolate had no effects on peri-natal and post-natal development in rats following subcutaneous exposure of approximately 1137 times the MRHD of LONHALA MAGNAIR based on an AUC comparison (at a maternal dose of up to 1.885 mg/kg/day).

8.2 Lactation

Risk Summary

There are no data on the presence of glycopyrrolate or its metabolites in human milk, the effects on the breastfed infant, or the effects on milk production. However, in a study of lactating rats, glycopyrrolate was present in the milk [see Data]. The developmental and health benefits of breastfeeding should be considered along with the mother’s clinical need for LONHALA MAGNAIR and any potential adverse effects on the breastfed infant from LONHALA MAGNAIR or from the underlying maternal condition.

Data

Glycopyrrolate (and its metabolites) was detected in the milk of lactating rats following a single intravenous injection of 4 mg/kg of radiolabeled glycopyrrolate.

8.4 Pediatric Use

LONHALA MAGNAIR is not indicated for use in children. The safety and efficacy of LONHALA MAGNAIR in pediatric patients have not been established.

8.5 Geriatric Use

Based on available data, no adjustment of the dosage of LONHALA MAGNAIR in geriatric patients is warranted. LONHALA MAGNAIR can be used at the recommended dose in elderly patients 75 years of age and older.
Of the total number of subjects in clinical studies of LONHALA MAGNAIR, 41% were aged 65 and older, while 8% were aged 75 and older. No overall differences in safety or effectiveness were observed between these subjects and younger subjects, and other reported clinical experience has not identified differences in responses between the elderly and younger patients, but greater sensitivity of some older individuals cannot be ruled out.

8.6 Renal Impairment
No dose adjustment is required for patients with mild and moderate renal impairment. The effects of renal impairment on the pharmacokinetics of glycopyrrolate have not been studied [see Clinical Pharmacology (12.3)].

8.7 Hepatic Impairment
No dose adjustment is required for patients with hepatic impairment. The effects of hepatic impairment on the pharmacokinetics of glycopyrrolate have not been studied [see Clinical Pharmacology (12.3)].

10. OVERDOSAGE
An overdose of glycopyrrolate may lead to anticholinergic signs and symptoms such as nausea, vomiting, dizziness, lightheadedness, blurred vision, increased intraocular pressure (causing pain, vision disturbances, or reddening of the eye), obtipation or difficulties in voiding.

In COPD patients, orally inhaled administration of LONHALA MAGNAIR at a total daily dose of 200 mcg for 28 consecutive days (maximum of 1 mg) was well tolerated. Pharmacokinetic results from several studies conducted in COPD patients showed that a single, well-tolerated dose of 1000 mcg had a Cmax of 1534 pg/mL and AUC0-inf of 5271 pg*hr/mL. These values are approximately 44 fold and 21 fold higher, respectively, than the estimated daily Cmax of 34.5 pg/mL and AUC0-inf of 255 pg*hr/mL for a 25 mcg BID dose regimen at steady-state.

11. DESCRIPTION
LONHALA MAGNAIR consists of LONHALA vials and a MAGNAIR nebulizer system. LONHALA (glycopyrrolate) Inhalation Solution is a sterile, clear, colorless, aqueous solution for oral inhalation.

Glycopyrrolate USP, the active component of LONHALA Inhalation Solution, is chemically described as (3RS)-3-[(2SR)-(2-cyclopentyl-2-hydroxy-2-penylacetyl)oxy]-1,1-dimethylpyrrolidinium bromide. Glycopyrrolate is a synthetic quaternary ammonium compound that acts as a competitive antagonist at muscarinic acetylcholine receptors, also referred to as an anticholinergic. Glycopyrrolate, C31H42BrN2O2, is a white, odorless, crystalline powder that is soluble in water and in alcohol. It has a molecular mass of 398.33. The structural formula is:

The inactive ingredients in LONHALA are: citric acid monohydrate, sodium chloride, sodium hydroxide and water for injection.

Like all other nebulized treatments, the amount delivered to the lungs will depend upon patient factors. Under standardized in vitro testing per USP<1601>- adult breathing pattern (500 mL tidal volume, 15 breaths per minute, and inhalation: exhalation ratio of 1:1), the mean delivered dose from the mouthpiece was approximately 14.2 mcg of glycopyrrolate (500 mL tidal volume, 15 breaths per minute, and inhalation: exhalation ratio of 1:1). The mean nebulization time was approximately 2 to 3 minutes.

12. CLINICAL PHARMACOLOGY
12.1 Mechanism of Action
Glycopyrrolate is a long-acting muscarinic antagonist, which is often referred to as an anticholinergic. It has similar affinity to the subtypes of muscarinic receptors M1 to M5. In the airways, it exhibits pharmacological effects through inhibition of M3 receptor at the smooth muscle leading to bronchodilation. The competitive and reversible nature of antagonism was shown with human and animal origin receptors and isolated organ preparations. In preclinical in vivo as well as in vivo studies, prevention of methacholine and acetylcholine induced bronchoconstrictive effects was dose-dependent and lasted longer than 24 hours. The clinical relevance of these findings is unknown. The bronchodilation following inhalation of glycopyrrolate is predominantly a site-specific effect.

12.2 Pharmacodynamics
Cardiac Electrophysiology:
In the dose ranging and confirmatory clinical studies, the administration of LONHALA MAGNAIR did not demonstrate any clinically relevant changes in cardiac function including: vital signs (heart rate, blood pressure), electrocardiograms (including QTc) and Holter monitoring. In addition, no major adverse cardiovascular events (MACE) were reported following the administration of LONHALA MAGNAIR 25 mcg in any clinical study.

12.3 Pharmacokinetics
Absorption
Following oral inhalation using MAGNAIR, glycopyrrolate was rapidly absorbed and reached peak plasma levels <20 minutes post dose. In patients with COPD, pharmacokinetic steady-state plasma levels of glycopyrrolate were reached within one week of the start of treatment. A twice daily dose regimen leads to approximately 2-3 fold accumulation of systemic glycopyrrolate exposure at steady-state.

Distribution
The in vitro human plasma protein binding of glycopyrrolate was 38% to 41% at concentrations of 1 to 10 ng/mL.

Metabolism
In vitro metabolism studies show glycopyrrolate hydroxylation resulting in a variety of mono- and bis-hydroxylated metabolites and direct hydrolysis resulting in the formation of a carboxylic acid derivative (M9). Further in vitro investigations showed that multiple CYP isoenzymes contribute to the oxidative biotransformation of glycopyrrolate and the hydrolysis to M9 is likely to be catalyzed by members from the cholinesterase family pre-systemically and/or via first pass metabolism from the swallowed dose fraction of orally inhaled glycopyrrolate.

Elimination
After intravenous administration of [3H]-labelled glycopyrrolate to humans, the mean urinary excretion of radioactivity in 48 hours amounted to 85% of the dose. A further 5% of the dose was found in the bile.

Renal elimination of parent drug accounts for about 60 to 70% of total clearance of systemically available glycopyrrolate whereas non-renal clearance processes account for about 30 to 40%. Biliary clearance contributes to the non-renal clearance, but the majority of non-renal clearance is thought to be due to metabolism.

Drug Interactions
In vitro inhibition studies demonstrated that glycopyrrolate has no relevant capacity to inhibit CYP1A2, CYP2A6, CYP2B6, CYP2C9, CYP2D6, CYP2E1 or CYP3A4/5, the efflux transporters MDRI, MRPI or MRP2, and the uptake transporters OATP1B1, OATP1B3, OAT1, OAT3, OCT1 or OCT2. In vitro enzyme induction studies did not indicate a clinically relevant induction by glycopyrrolate for cytochrome P450 isoenzymes, or for UGT1A1 and the transporters MDRI and MRPI.

There is potential for additive interaction with concomitantly used anticholinergic medications. Therefore, avoid coadministration of LONHALA MAGNAIR with other anticholinergic-containing drugs as this may lead to an increase in anticholinergic effects [see Warnings and Precautions (5.4, 5.5) and Adverse Reactions (6)].

Specific Populations
A population pharmacokinetic analysis of data in COPD patients indicated no clinically relevant effect of age (41 to 80 years) or body weight (40.1 to 154.8 kg) on systemic exposure to glycopyrrolate. In addition, there was no evidence of clinically significant ethnic/race effect.

Renal Impairment
The effects of renal impairment on the pharmacokinetics of glycopyrrolate have not been studied [see Use in Specific Populations (8.6)].

Hepatic Impairment
The effects of hepatic impairment on the pharmacokinetics of glycopyrrolate have not been studied. Glycopyrrolate is cleared predominantly from systemic circulation by renal excretion [see Use in Specific Populations (8.7)].

13. NONCLINICAL TOXICOLOGY
13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility
Carcinogenicity studies of glycopyrrolate did not result in an increase in the incidence of tumors in a 2-year inhalation study of glycopyrrolate in Wistar rats at doses up to 0.56 mg/kg/day, approximately 143 times the MRHDID of LONHALA MAGNAIR in adults on an AUC basis. Also, no evidence of tumorigenicity occurred in a 26-week oral (gavage) study in male and female TgrasH2 mice that received glycopyrrolate at doses up to 93.8 and 125.1 mg/kg/day, respectively, approximately 66 times the MRHDID of LONHALA MAGNAIR.

For oral inhalation.

LONHALA Inhalation Solution is supplied in low-density polyethylene (LDPE) unit dose vials, each containing 1.0 mL of the solution. Each unit-dose vial contains 25 mcg of glycopyrrolate in a sterile, isotonic saline solution, pH-adjusted to 4.0 with citric acid and sodium hydroxide.

Like all other nebulized treatments, the amount delivered to the lungs will depend upon patient factors. Under standardized in vitro testing per USP<1601>- adult breathing pattern (500 mL tidal volume, 15 breaths per minute, and inhalation: exhalation ratio of 1:1), the mean delivered dose from the mouthpiece was approximately 14.2 mcg of glycopyrrolate (equivalent to 11.4 mcg glycopyronium and 56.8% label claim). The mass median aerodynamic diameter (MMAD) of the nebulized aerosol particles/droplets is 3.71 μm with a geometric standard deviation of 2.92 - 4.49 μm as determined using the Next Generation Impactor (NGI) method.

Aerosol delivered to the lungs accounts for about 30 to 40%. Biliary clearance contributes to the non-renal clearance, but the majority of non-renal clearance is thought to be due to metabolism.

Drug Interactions
In vitro inhibition studies demonstrated that glycopyrrolate has no relevant capacity to inhibit CYP1A2, CYP2A6, CYP2B6, CYP2C9, CYP2D6, CYP2E1 or CYP3A4/5, the efflux transporters MDRI, MRPI or MRP2, and the uptake transporters OATP1B1, OATP1B3, OAT1, OAT3, OCT1 or OCT2. In vitro enzyme induction studies did not indicate a clinically relevant induction by glycopyrrolate for cytochrome P450 isoenzymes, or for UGT1A1 and the transporters MDRI and MRPI.

There is potential for additive interaction with concomitantly used anticholinergic medications. Therefore, avoid coadministration of LONHALA MAGNAIR with other anticholinergic-containing drugs as this may lead to an increase in anticholinergic effects [see Warnings and Precautions (5.4, 5.5) and Adverse Reactions (6)].

Specific Populations
A population pharmacokinetic analysis of data in COPD patients indicated no clinically relevant effect of age (41 to 80 years) or body weight (40.1 to 154.8 kg) on systemic exposure to glycopyrrolate. In addition, there was no evidence of clinically significant ethnic/race effect.

Renal Impairment
The effects of renal impairment on the pharmacokinetics of glycopyrrolate have not been studied [see Use in Specific Populations (8.6)].

Hepatic Impairment
The effects of hepatic impairment on the pharmacokinetics of glycopyrrolate have not been studied. Glycopyrrolate is cleared predominantly from systemic circulation by renal excretion [see Use in Specific Populations (8.7)].
Glycopyrrolate was not mutagenic in the following genotoxicity assays: the *in vitro* Ames assay, *in vitro* human lymphocyte chromosomal aberration assay, and *in vivo* rat bone marrow micronucleus assay.

Impairment of fertility was observed in male and female Wistar rats at a subcutaneous glycopyrrolate dose of 1.88 mg/kg/day (approximately and 2035 and 1136 times, respectively, the MRHD of LONHALA MAGNAIR on an AUC basis) based upon findings of decreased implantation sites and corresponding reduction of live fetuses. No effects on fertility and reproductive performance occurred in male and female rats at a subcutaneous glycopyrrolate dose of 0.63 mg/kg/day, approximately 384 times the MRHD of LONHALA MAGNAIR on an AUC basis.

14. CLINICAL STUDIES

The safety and efficacy of LONHALA MAGNAIR were evaluated in 2 dose-ranging studies, 2 placebo-controlled confirmatory studies (12-week studies), and a 48-week long-term safety study. The efficacy of LONHALA MAGNAIR is based primarily on the dose-ranging studies in 378 subjects with COPD and the 2 placebo-controlled confirmatory studies in 1293 subjects with COPD.

14.1 Dose Ranging Studies

Dose selection for the confirmatory COPD studies for LONHALA MAGNAIR was supported by two studies. Study A was a randomized, double-blind, placebo-controlled, parallel arm study with a 28-day treatment period. The study included LONHALA MAGNAIR doses of placebo, 12.5 mcg, 25 mcg, 50 mcg, and 100 mcg twice daily. The Study demonstrated a dose-response effect on peak and trough FEV₁ over 24-hour dosing period in subjects treated with LONHALA MAGNAIR twice daily (Figure 1 (Day 1) and Figure 2 (Day 28)). The LS mean differences in trough FEV₁ from baseline after 28 days compared to placebo for the 12.5 mcg, 25 mcg, 50 mcg, and 100 mcg twice daily doses were 0.117 L (95% CI: 0.037, 0.197); 0.128 L (95% CI: 0.048, 0.209), 0.146 L (95% CI: 0.067, 0.226), and 0.177 L (95% CI: 0.099, 0.255), respectively. In Study A, all subjects in each treatment group (N=282) had FEV₁, AUC_Co, Co, and serial spirometry assessments while a subset of subjects (N=125; shown in Figure 1 and Figure 2 below) had extended FEV₁, AUC_24h, assessments on Days 1 and 28.

Study B was a randomized, six-way, crossover study with 7-day treatment periods separated by 5-7-day washout periods. Study B included LONHALA MAGNAIR doses of placebo, 3 mcg, 6.25 mcg, 12.5 mcg, and 50 mcg twice daily with aclidinium bromide 400 mcg BID as an active control.

The dose-ranging results from Study A and Study B supported the evaluation of LONHALA MAGNAIR 25 mcg and 50 mcg twice-daily in the confirmatory COPD trials. The results of Study A are reported in Figure 1 below.

14.2 Confirmatory Studies

There were 2 confirmatory studies (Study 1 and Study 2) for LONHALA MAGNAIR. Both studies were randomized, double-blind, placebo-controlled, parallel-group 12-week studies in subjects with COPD designed to evaluate the efficacy of LONHALA MAGNAIR on lung function. These studies treated subjects who had a clinical diagnosis of COPD, were 40 years of age or older, had a history of smoking greater than or equal to 10 pack-years, a post-bronchodilator FEV₁ less than or equal to 80% of predicted, and an FEV₁/FVC ratio less than 0.7. Subjects also had pre-existing or concurrent cardiovascular disease and stable, background LABA ± ICS and SAMA use were permitted. Subjects in Study 1 and Study 2 had a mean age of 63 years, were primarily male (56%), Caucasian (90%), and 53% were current smokers. At screening, the post-bronchodilator percent predicted FEV₁ was 52% (range: 20% to 80%), the mean post-bronchodilator percent reversibility was 18% (range: -33% to 86%).

Study 1 and Study 2 evaluated LONHALA MAGNAIR (glycopyrrolate) 25 mcg and 50 mcg twice-daily and placebo twice-daily. The primary endpoint was the change from baseline in trough FEV₁ at Day 84 compared with placebo. LONHALA MAGNAIR twice-daily demonstrated a larger increase in LS mean change from baseline in trough FEV₁ compared to placebo. Compared to LONHALA MAGNAIR 25 mcg twice daily, LONHALA MAGNAIR 50 mcg twice daily did not provide sufficient additional benefit on a variety of endpoints, including FEV₁, to support use of higher doses. Table 2 presents the results from Studies 1 and 2 for LONHALA MAGNAIR 25 mcg twice-daily.

<table>
<thead>
<tr>
<th>Treatment</th>
<th>N</th>
<th>Change from baseline LS Mean (SE)</th>
<th>Comparison</th>
<th>Treatment Difference LS Mean (SE)</th>
<th>95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Study 1</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>LONHALA MAGNAIR 25 mcg BID</td>
<td>217</td>
<td>0.089 (0.014)</td>
<td>LONHALA MAGNAIR - Placebo</td>
<td>0.096 (0.019)</td>
<td>0.059, 0.133</td>
</tr>
<tr>
<td>Placebo</td>
<td>218</td>
<td>-0.008 (0.014)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Study 2</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>LONHALA MAGNAIR 25 mcg BID</td>
<td>214</td>
<td>0.092 (0.014)</td>
<td>LONHALA MAGNAIR - Placebo</td>
<td>0.081 (0.020)</td>
<td>0.042, 0.120</td>
</tr>
<tr>
<td>Placebo</td>
<td>212</td>
<td>0.011 (0.015)</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*Study results are from a treatment policy strategy which analyzes all collected data, including data for some patients who discontinued study treatment prior to Week 12 and may have received other COPD treatment but were followed. Analyses of efficacy data measured only while on randomized blinded study treatment showed similar results.*
In Study 1, serial spirometric evaluations throughout the 12-hour dosing interval were performed in a subset of subjects on Day 1 and Day 84. The spirometric curves from Study 1 on Day 1 and Day 84 are displayed in Figure 3 and Figure 4.

In Study 1, the SGRQ responder rate (defined as an improvement in score of 4 or more as a threshold) for the LONHALA MAGNAIR 25 mcg treatment arm was 51% compared to 31% for placebo [Odds Ratio: 1.65; 95% CI: 1.15, 2.36]. In Study 2, the SGRQ responder rate for the LONHALA MAGNAIR 25 mcg treatment arm was 46% compared to 33% for placebo [Odds Ratio: 1.39; 95% CI: 0.93, 2.08].

The peak FEV₁ was defined as the highest postdose FEV₁ within the first 12 hours after morning dosing for each subject on Days 1 and 84, respectively, for the substudy population. The mean peak FEV₁ improvement from baseline for LONHALA MAGNAIR on Day 1 and on Day 84 in the subset of subjects was 0.228 L and 0.214 L (Study 1) respectively.

The St. George's Respiratory Questionnaire (SGRQ) was assessed in Studies 1 and 2. Over Time on Day 1 and Day 84 in the subset of subjects was 0.228 L and 0.214 L (Study 1) respectively. The peak FEV₁ within the first 12 hours after morning dosing for each subject on Days 1 and 84.

16. HOW SUPPLIED/STORAGE AND HANDLING

16.1 How Supplied
LONHALA MAGNAIR is supplied as a 1 mL sterile, clear, colorless, aqueous solution in low-density polyethylene (LDPE) unit-dose vials overwrapped in foil. LONHALA MAGNAIR is available in a Starter Kit containing 60 unit-dose vials packaged with one MAGNAIR, and FDA approved patient labeling. LONHALA MAGNAIR is also supplied in a Refill Kit containing 60 unit-dose vials packaged with a MAGNAIR Replacement Handset and FDA approved patient labeling.

<table>
<thead>
<tr>
<th>Package Configuration</th>
<th>Dosage Strength</th>
<th>NDC</th>
</tr>
</thead>
<tbody>
<tr>
<td>Starter Kit with 30 day supply (30 foil pouches with 2 vials per pouch) and complete MAGNAIR Nebulizer System</td>
<td>25 mcg</td>
<td>NDC: 63402-201-00</td>
</tr>
<tr>
<td>Refill Kit with 30 day supply (30 foil pouches with 2 vials per pouch) and MAGNAIR Replacement Handset</td>
<td>25 mcg</td>
<td>NDC: 63402-301-01</td>
</tr>
</tbody>
</table>

16.2 Storage and Handling
Store LONHALA Inhalation Solution in the protective foil pouch at 20°-25°C (68°-77°F) [see USP Controlled Room Temperature].
- LONHALA vials should be used with MAGNAIR only. Do not use MAGNAIR with any other vials.
- Store LONHALA vials in the protective foil pouch. After opening the foil pouch, unused unit-dose vials should be returned to, and stored in, the foil pouch. Once a foil pouch is opened, discard the vials if not used within 7 days. An opened unit-dose vial should be used right away. Discard any unit-dose vial if the solution is not colorless. Always use the MAGNAIR Replacement Handset parts that come with each LONHALA MAGNAIR refill prescription. Keep out of the reach of children.

17. PATIENT COUNSELING INFORMATION
Advising the patient to read the FDA-approved patient labeling (Patient Information and Instructions for Use).

Not for Acute Symptoms: Inform patients that LONHALA MAGNAIR is not meant to relieve acute symptoms of COPD and extra doses should not be used for that purpose. Advise them to treat acute symptoms with a rescue inhaler such as albuterol. Provide patients with such medicine and instruct them in how it should be used [see Warnings and Precautions (5.1)].

Instruct patients to seek medical attention immediately if they experience any of the following:
- Symptoms get worse
- Need for more inhalations than usual of their rescue inhaler

Patients should not stop therapy with LONHALA MAGNAIR without physician/provider guidance since symptoms may recur after discontinuation.

Paradoxical Bronchospasm: Inform patients that LONHALA MAGNAIR can cause paradoxical bronchospasm. If paradoxical bronchospasm occurs, instruct patients to discontinue LONHALA MAGNAIR.

Worsening of Narrow-Angle Glaucoma: Instruct patients to be alert for signs and symptoms of acute narrow-angle glaucoma (e.g., eye pain or discomfort, blurred vision, visual halos or colored images in association with red eyes from conjunctival congestion and corneal edema). Instruct patients to consult a physician immediately if any of these signs or symptoms develop.

Instruct patients to consult a physician immediately if any of these signs or symptoms develop.

Worsening of Urinary Retention: Instruct patients to be alert for signs and symptoms of urinary retention (e.g., difficulty passing urine, painful urination). Instruct patients to consult a physician immediately if any of these signs or symptoms develop.

Instruct patients not to inject or swallow the LONHALA solution. Instruct patients to use one inhalation of LONHALA MAGNAIR orally twice daily (1 vial in the morning and 1 vial in the evening) at the same time every day.

Instructions for Administering LONHALA MAGNAIR
It is important for patients to understand how to correctly administer LONHALA vials using MAGNAIR [see Instructions for Use], instruct patients that LONHALA vials should only be administered via MAGNAIR and MAGNAIR should not be used for administering other medications. Patients should be instructed not to inject or swallow the LONHALA solution.

Instruct patients to store LONHALA vials in the sealed foil pouch and to only open the foil pouch to remove a LONHALA vial immediately before use. Instruct patients that unopened vials should be returned to the opened foil pouch for use at their next treatment and discarded if not used within 7 days or if it may not be as effective.

Instruct patients to use one inhalation of LONHALA MAGNAIR orally twice daily (1 vial in the morning and 1 vial in the evening) at the same time every day.

Inform patients that they may miss a dose of LONHALA MAGNAIR, they should use their next vial at the usual time. Instruct patients to not use 2 vials at one time and to not use more than 2 vials in a day. Patients should throw the plastic dispensing vials away immediately after use. Due to their small size, the vials pose a danger of choking to small children.
Inform patients treated with LONHALA MAGNAIR that a Refill Kit will be provided to them on a monthly basis. The Refill Kit will include foil pouches containing 60 vials of LONHALA (2 vials of LONHALA in each pouch; 1 vial per dose), and 1 MAGNAIR Replacement Handset (containing only these replacement parts: Medication cap, Handset body, Mouthpiece, and Aerosol head; Manufacturer’s Instructions for Use booklet).

Important: Instruct patients to throw away the old Handset parts after using 60 vials of LONHALA and use the replacement Handset parts with the next 60 vials of LONHALA.

PATIENT INFORMATION

LONHALA MAGNAIR (lon-HAH-luh MAGG-nair)
(glycopyrrolate) inhalation solution, for oral inhalation use

Important: For oral inhalation only. Do not inject or swallow the LONHALA medicine. LONHALA vials are used only with the MAGNAIR device. Do not use MAGNAIR with any other medicine.

Read this Patient Information that comes with LONHALA MAGNAIR before you start using it and each time you get a refill. There may be new information. This Patient Information does not take the place of talking to your healthcare provider about your medical condition or treatment.

What is LONHALA MAGNAIR?

LONHALA MAGNAIR is an anticholinergic medicine known as glycopyrrolate.

- Anticholinergic medicines such as LONHALA MAGNAIR help the muscles around the airways in your lungs stay relaxed to prevent symptoms such as wheezing, coughing, chest tightness, and shortness of breath. This makes it hard to breathe.
- LONHALA MAGNAIR is used for maintenance treatment of Chronic Obstructive Pulmonary Disease (COPD). COPD is a long-term (chronic) lung disease that includes chronic bronchitis, emphysema, or both.
- LONHALA MAGNAIR is for long-term use and should be taken 2 times each day to improve symptoms of COPD for better breathing.
- LONHALA MAGNAIR is not used to treat sudden symptoms of COPD. Always have a short-acting beta₂-agonist medicine (rescue inhaler) with you to treat sudden symptoms of COPD. If you do not have a rescue inhaler, contact your healthcare provider to have one prescribed for you.
- LONHALA MAGNAIR should not be used in children. It is not known if LONHALA MAGNAIR is safe and effective in children younger than 18 years of age.

Do not use LONHALA MAGNAIR if you:

- are allergic to glycopyrrolate, or any of the ingredients in LONHALA MAGNAIR. Ask your healthcare provider if you are not sure. See “What are the ingredients in LONHALA MAGNAIR?” at the end of this Patient Information leaflet for a complete list of ingredients in LONHALA MAGNAIR.

Before using LONHALA MAGNAIR, tell your healthcare provider about all of your medical conditions, including if you:

- have kidney problems.
- have eye problems such as glaucoma. LONHALA MAGNAIR may make your glaucoma worse.
- have prostate or bladder problems, or problems passing urine. LONHALA MAGNAIR may make these problems worse.
- are pregnant or plan to become pregnant. It is not known if LONHALA MAGNAIR can harm your unborn baby.
- are breastfeeding or plan to breastfeed. It is not known if the medicine in LONHALA MAGNAIR passes into your breast milk and if it can harm your baby. You and your healthcare provider should decide if you will take LONHALA MAGNAIR or breastfeed.
- are allergic to LONHALA MAGNAIR or any of its ingredients or any other medicines.

Tell your healthcare provider about all the medicines you take, including prescription medicines and over-the-counter medicines, vitamins, and herbal supplements. LONHALA MAGNAIR may affect the way other medicines work, and other medicines can affect how LONHALA MAGNAIR works. Using LONHALA MAGNAIR with other medicines may cause serious side effects.

Especially tell your healthcare provider if you take anticholinergics (including umeclidinium, tiotropium, ipratropium, aclidinium, glycopyrrolate).

Know the medicines you take. Keep a list of your medicines with you and show it to your healthcare provider and pharmacist when you get a new medicine.

How should I use LONHALA MAGNAIR?

Read the step-by-step instructions for using LONHALA MAGNAIR at the end of this Patient Information leaflet and the Manufacturer’s Instructions for Use booklet. The Manufacturer’s Instructions for Use booklet provides complete information about how to put together (assemble), prepare, use, care for, and troubleshoot your MAGNAIR nebulizer system.

- Do not use LONHALA MAGNAIR unless your healthcare provider has taught you how to use the device and you understand how to use it correctly.
- Use LONHALA MAGNAIR exactly as your healthcare provider tells you to use it. Do not use LONHALA MAGNAIR more often than prescribed for you.
- Only use LONHALA vials with the MAGNAIR device.
- Do not inject or swallow the LONHALA medicine.
- Inhale the medicine in 1 LONHALA vial through the MAGNAIR device 2 times each day (1 vial in the morning and 1 vial in the evening) at the same time each day.
- If you miss a dose of LONHALA MAGNAIR, take your next dose at your usual time.
  - Do not use 2 vials at 1 time.
  - Do not use more than 2 vials in a day.
- Do not stop using LONHALA MAGNAIR or other medicines to control or treat your COPD unless told to do so by your healthcare provider because your symptoms might get worse. Your healthcare provider will change your medicines as needed.
- Call your healthcare provider or get emergency medical care right away if your breathing problems worsen with LONHALA MAGNAIR, you need to use your rescue medicine more often than usual, or your rescue inhaler medicine does not work as well for you at relieving your symptoms.
What are the possible side effects of LONHALA MAGNAIR?

LONHALA MAGNAIR can cause serious side effects, including:

- **sudden shortness of breath immediately after use of LONHALA MAGNAIR. Sudden shortness of breath may be life-threatening.** If you have sudden breathing problems immediately after inhaling your medicine, stop taking LONHALA MAGNAIR and call your healthcare provider or go to the nearest hospital emergency room right away.

- **serious allergic reactions.** Stop using LONHALA MAGNAIR and call your healthcare provider or get emergency medical care right away if you get any of the following symptoms of a serious allergic reaction:
  - rash
  - swelling of the tongue, lips, and face
  - difficulty breathing
  - hives
  - or swallowing

- **new or worsened eye problems including acute narrow-angle glaucoma.** Acute narrow-angle glaucoma can cause permanent loss of vision if not treated. Symptoms of acute narrow-angle glaucoma may include:
  - eye pain or discomfort
  - red eyes
  - blurred vision
  - seeing halos or bright colors around lights
  - nausea or vomiting
  - difficulty breathing
  - or swallowing

If you have any of these symptoms, stop taking LONHALA MAGNAIR and call your healthcare provider right away before using another dose.

- **new or worsened problems emptying your bladder (urinary retention).** People who use LONHALA MAGNAIR may develop new or worsened urinary retention. Urinary retention can be caused by a blockage in your bladder. Urinary retention can also happen in men who have a larger than normal prostate. Symptoms of urinary retention may include:
  - difficulty urinating
  - painful urination
  - urinating frequently
  - urination in a weak stream or drips
  - or swallowing

If you have any of these symptoms, stop taking LONHALA MAGNAIR and call your healthcare provider right away before taking another dose.

**Common side effects of LONHALA MAGNAIR include shortness of breath, and urinary tract infections.**

These are not all of the possible side effects of LONHALA MAGNAIR.

**Call your doctor or pharmacist for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.**

How should I store LONHALA MAGNAIR?

- **Store LONHALA vials in the protective foil pouch at room temperature between 68°F and 77°F (20°C and 25°C).**
- **LONHALA vials should be used with the MAGNAIR device only. Do not use MAGNAIR with any other medicine.**

After opening the protective foil pouch, unused LONHALA vials should be returned to, and stored in, the opened foil pouch. Once a foil pouch is opened, discard the vials if not used within 7 days.

- **An opened LONHALA vial should be used right away.**
- **Throw away the LONHALA vial right away after use.**
- **The medicine in the LONHALA vial should be colorless. Throw away the LONHALA vial if the medicine is not colorless.**
- **Always use the MAGNAIR Replacement Handset parts that come with each LONHALA MAGNAIR refill prescription.**
- **Keep LONHALA MAGNAIR and all medicines out of the reach of children.**

General information about the safe and effective use of LONHALA MAGNAIR.

Medicines are sometimes prescribed for purposes other than those listed in a Patient Information leaflet. Do not use LONHALA MAGNAIR for a condition for which it was not prescribed. Do not give LONHALA MAGNAIR to other people, even if they have the same symptoms you have. It may harm them.

You can ask your healthcare provider or pharmacist for information about LONHALA MAGNAIR that is written for healthcare professionals.

What are the ingredients in LONHALA MAGNAIR?

**Active ingredient:** glycopyrrolate

**Inactive ingredients:** citric acid monohydrate, sodium chloride, sodium hydroxide and water for injection.

**Manufactured for:** Sunovion Respiratory Development Inc., a wholly-owned subsidiary of Sunovion Pharmaceuticals Inc. Made in Germany.

©2019 Sunovion Pharmaceuticals Inc. All rights reserved.

For more information, go to www.LonhalaMagnair.com or call Sunovion Customer Service at 1-888-394-7377.

This Patient information has been approved by the U.S. Food and Drug Administration

Revised: June 2019
INSTRUCTIONS FOR USE
LONHALA MAGNAIR (lon-HAH-luh MAGG-nair) (glycopyrrolate)
inhalation solution, for oral inhalation use

Read this Instructions for Use leaflet and the Manufacturer’s Instructions for Use booklet before you start using LONHALA MAGNAIR and each time you get a refill. There may be new information. This information does not take the place of talking to your healthcare provider about your medical condition or treatment. If you have any questions, ask your healthcare provider or pharmacist. The Manufacturer’s Instructions for Use booklet provides complete information about how to put together (assemble), prepare, use, care for, and troubleshoot your MAGNAIR nebulizer system.

Your LONHALA MAGNAIR:
MAGNAIR is a nebulizer system to be used by the patient, caregiver, or healthcare provider to deliver the medicine LONHALA. LONHALA MAGNAIR consists of both the MAGNAIR nebulizer system and the medicine LONHALA.
The following supplies come with your LONHALA MAGNAIR:
Starter Kit: Foil pouches containing 60 vials of LONHALA (2 vials of LONHALA in each pouch; 1 vial per dose), Instructional video, and 1 MAGNAIR Nebulizer System with carrying bag (including Manufacturer’s Instructions for Use booklet and Quick Reference Guide) (see the figure below).
Refill Kit: Foil pouches containing 60 vials of LONHALA (2 vials of LONHALA in each pouch; 1 vial per dose), and 1 MAGNAIR refill Handset (containing only these replacement parts: Medication cap, Handset body, Mouthpiece, and Aerosol head; Manufacturer’s Instructions for Use booklet).

Important: Throw away the old Handset parts after using 60 vials of LONHALA and use the replacement Handset parts in the Refill Kit with the next 60 vials of LONHALA.

MAGNAIR Nebulizer System

Controller
Medication cap
Aerosol head
AC adapter
Connection cord
Handset body
Mouthpiece
Carrying bag
4 AA batteries

Steps for Using Batteries with MAGNAIR
Step 1: Open the battery door on the Controller. Place your thumb on the black tab of the battery door and firmly push the tab to open the door.
Step 2: Put 4 AA batteries in the Controller as shown.
Step 3: Close the battery door. You may hear a “Click”.
Important: Make sure to have an extra set of batteries with you at all times if you choose not to use the AC adapter.

Steps for Using the AC Adapter with MAGNAIR
Step 1: Plug the AC adapter into the Inlet on the battery door of the Controller.
Step 2: Plug the AC adapter into the wall outlet.

Assembling Your MAGNAIR
Step 1: Wash your hands.
Step 2: Open the top of the Handset body by lifting the clasp.
Step 3: Insert the Aerosol head into the Handset body as shown. Do not touch the center of the Aerosol head. Notice that the Aerosol head has a small tab on the side. Align the small tab with the matching notch in the Handset body.
Step 4: Close the Handset body. You may hear a “Click”.

Important: Check to make sure that your MAGNAIR nebulizer system is working properly before you use LONHALA MAGNAIR for the first time. See the Manufacturer’s Instructions for Use that come with your MAGNAIR nebulizer system.
### Using LONHALA MAGNAIR

**Step 1:** Open the foil pouch, enough to remove the 2 LONHALA vials and separate them. Return 1 vial to the opened foil pouch and store in the carrying bag to be used at the next treatment. Discard the vial if not used within 7 days.

**Step 2:** Insert one LONHALA vial into the bottom of the Medication cap until it “Clicks”.

**Step 3:** Make sure the Aerosol head is installed before attaching the Medication cap because your medicine could leak and you will not get your full treatment. Place the Medication cap with LONHALA vial on the top of the Handset body.

**Step 4:** To attach the Medication cap to the Handset body, turn the Medication cap in a clockwise direction as shown, until you hear a “Click”. The notch in the Medication cap (at the base of the opening) should line up with the blue line on the Handset body.

**Step 5:** Insert the Mouthpiece into your mouth. **Important: Do not tilt the Handset, loosen or remove the Medication cap, or unclasp the Handset body until the treatment is complete because you will not get your full treatment.**

**Step 6:** Press the On/Off button to turn on the Controller as shown, and start your treatment.

**Step 7:** Breathe in (inhale) and breathe out (exhale) normally through the Mouthpiece. At the end of your treatment, you will hear a beeping sound and the Controller will automatically shut off. Your treatment should take about 2 to 3 minutes.

### Cleaning the Handset

**Step 1:** Disconnect the Handset from the Connection cord.

**Step 2:** Turn the Medication cap in a counterclockwise direction as shown, to remove it from the Handset body.

**Step 3:** Place the top of the Medication cap into the palm of your hand and push up as shown to remove the LONHALA vial. Throw away the LONHALA vial into the wastebasket.

**Step 4:** Remove the Mouthpiece from the Handset body by giving it a gentle twist and pull to separate from the Handset body.

**Step 5:** Carefully loosen the Blue valve from the slot in the Mouthpiece. Make sure the valve is still attached on one side to the Mouthpiece.

**Step 6:** Remove the Aerosol head from the Handset body by lifting the clasp on the side of the Handset body as shown. **Do not** touch the center of the Aerosol head.
Step 7: Set aside the Aerosol head to be cleaned separately (see Step 10). Rinse all Handset parts well with warm running water for about 10 seconds.

Step 8: Wash all Handset parts in warm soapy water (water and clear liquid dishwashing soap) for about 10 seconds.

Step 9: Rinse the Handset parts well with warm running water for about 10 seconds to remove all of the soap.

Step 10: Clean the Aerosol head using the instructions in Steps 7 through 10.

10A. Rinse each side of the Aerosol head well with warm running water for about 10 seconds.

10B. Hold the Aerosol head by the handle and swish it back and forth in the warm soapy water for about 10 seconds.

10C. Then, rinse both sides of the Aerosol head well with warm running water for about 10 seconds on each side.

Rinsing the Aerosol head well helps prevent clogging and ensure proper operation.

Step 11: Inspect all Handset parts to make sure they are completely clean. If any Handset parts are still dirty, soak the parts in warm soapy water for 5 more minutes. Rinse well with warm running water until the Handset parts are clean. Shake Handset parts to remove excess water. Air-dry all Handset parts on a lint-free towel. Do not put the Handset parts back together until ready to use again for your next treatment of LONHALA.

Step 12: Store the Handset parts in the carrying bag provided.

Warnings and Precautions:
Failure to follow the Warnings and Precautions below could cause serious injury or may lead to death in some cases:

- Check all parts of your LONHALA MAGNAIR to make sure that they are clean and not damaged.
- Clean the Handset before the first use and after every use. If you do not clean the Handset after every use, your treatment could take more than 3 minutes.
- Do not leave the Aerosol head in your Handset.
- Do not wash the Controller, Connection cord, or AC adapter.
- Only use clear liquid dishwashing soap to wash the Handset parts. Do not use any other type of soap.
  - Do not use antibacterial soap. Antibacterial soap can damage the Aerosol head.
- Do not use a microwave oven to dry any parts of your LONHALA MAGNAIR.
- Allow all parts of your LONHALA MAGNAIR to air dry completely.

If you have any questions, contact Sunovion Customer Service at 1-888-394-7377.

This Instructions for Use has been approved by the U.S. Food and Drug Administration.

LONHALA and are registered trademarks of Sunovion Pharmaceuticals Inc. MAGNAIR is a registered trademark of PARI Pharma GmbH, used under license.

SUNOVION and are registered trademarks of Sumitomo Dainippon Pharma Co., Ltd. Sunovion Pharmaceuticals Inc. is a U.S. subsidiary of Sumitomo Dainippon Pharma Co., Ltd.

PARI and are registered trademarks of PARI GmbH. eFlow is a registered trademark of PARI Pharma GmbH. Made under license of The Technology Partnership plc.

Manufactured for: Sunovion Respiratory Development Inc., a wholly-owned subsidiary of Sunovion Pharmaceuticals Inc. Made in Germany.

©2019 Sunovion Pharmaceuticals Inc. All rights reserved.
Revised: June 2019
10140-02-MKT2