For oral inhalation only.

**INDICATIONS AND USAGE**

Inhalation Aerosol, for oral inhalation use

**Important limitations:**

- Adrenergic agonist indicated for:
  - Asthma
  - Chronic obstructive pulmonary disease (COPD)
  - Acute episodes of asthma
  - Treatment of status asthmaticus

**Dosage and Administration**

- Treatment of asthma in patients 6 years of age and older. (1.1)
- Maintenance treatment of airflow obstruction and reducing exacerbations in patients with chronic obstructive pulmonary disease (COPD) including chronic bronchitis and/or emphysema. (1.2)

**Dosage Forms and Strengths**

- Metered-dose inhaler containing a combination of budesonide (80 or 160 mcg) and formoterol (4.5 mcg) as an inhalation aerosol. (3)

**Contraindications**

- Hypersensitivity to any of the ingredients in SYMBICORT (4)

**Warnings and Precautions**

- Serious asthma-related events: Long-acting beta₂-adrenergic agonists as monotherapy increase the risk. (5.1)
- Deterioration of disease and acute episodes: Do not initiate in acutely deteriorating asthma or COPD or to treat acute symptoms. (5.2)
- Use with additional long-acting beta₂-agonist: Do not use in combination because of risk of overuse. (5.3)
- Localized infections: Candida albicans infection of the mouth and throat may occur. Monitor patients periodically for signs of adverse effects on the oral cavity. Advise the patient to rinse his/her mouth with water without swallowing after inhalation to help reduce the risk. (5.4)
- Pneumonia: Increased risk in patients with COPD. Monitor patients for signs and symptoms of pneumonia and other potential lung infections. (5.5)

**Adverse Reactions**

- Most common adverse reactions (incidence ≥3%):
  - Asthma: nasopharyngitis, headache, upper respiratory tract infection, pharyngolaryngeal pain, sinusitis, influenza, back pain, nasal congestion, stomach discomfort, vomiting, and oral candidiasis. (6.1)
  - COPD: nasopharyngitis, oral candidiasis, bronchitis, sinusitis, upper respiratory tract infections. (6.2)

**Drug Interactions**

- Strong cytochrome P450 3A4 inhibitors (e.g., ritonavir): Use with caution. May cause increased systemic corticosteroid effects. Exercise caution when used with SYMBICORT. (5.9)

**Use in Specific Populations**

- Hepatic impairment: Monitor patients for signs of increased drug exposure. (8.6)

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

**Revised:** 07/2019
If asthma symptoms arise in the period between doses, an inhaled, short-acting beta-agonist (LABA) may be used if the patient is known to be more likely to experience adverse effects with higher doses of formoterol. Patients using LABA twice daily (morning and evening) should be instructed to use the full dose of SYMBICORT as needed. SYMBICORT is NOT indicated for the relief of acute bronchospasm.

2.1 Administration Information
SYMBICORT should be administered as 2 inhalations twice daily (morning and evening), approximately 12 hours apart, every day by the orally inhaled route only. After inhalation, the patient should rinse the mouth with water without swallowing.
Prime SYMBICORT before using for the first time by releasing two test sprays into the air away from the face, shaking well for 5 seconds before each spray. In cases where the inhaler has not been used for more than 7 days or when it has been dropped, prime the inhaler again by shaking well before each spray and releasing two test sprays into the air away from the face.

More frequent administration or a higher number of inhalations (more than 2 inhalations twice daily) of the prescribed strength of SYMBICORT is not recommended as some patients are more likely to experience adverse effects with higher doses of formoterol. Patients using LABA should not use additional LABA for any reason [see Warnings and Precautions (5.3, 5.12)].

2.2 Asthma
If asthma symptoms arise in the period between doses, an inhaled, short-acting beta-agonist should be taken for immediate relief.

Adult and Adolescent Patients 12 Years of Age and Older
For patients 12 years of age and older, the dose is 2 inhalations of SYMBICORT 80/4.5 or SYMBICORT 160/4.5 twice daily.
The recommended starting dosages for SYMBICORT for patients 12 years of age and older are based upon patients' asthma severity or level of control of asthma symptoms, and risk of exacerbations on current inhaled corticosteroids.
The maximum recommended dosage in adult and adolescent patients 12 years of age and older is SYMBICORT 160/4.5, two inhalations daily. Improvement in asthma control following inhaled administration of SYMBICORT can occur within 15 minutes of beginning treatment, although maximum benefit may not be achieved for 2 weeks or longer after beginning treatment. Individual patients will experience a variable time to onset and degree of symptom relief.
For patients who do not respond adequately to the starting dose after 1-2 weeks of therapy with SYMBICORT 80/4.5, replacement with SYMBICORT 160/4.5 may provide additional asthma control.

If a previously effective dosage regimen of SYMBICORT fails to provide adequate control of asthma, the therapeutic regimen should be re-evaluated and additional therapeutic options, (e.g., replacing the lower strength of SYMBICORT with the higher strength, adding additional inhaled corticosteroid, or initiating oral corticosteroids) should be considered.

Pediatric Patients Aged 6 to Less than 12 Years
For patients 6 to less than 12 years of age, the dosage is 2 inhalations of SYMBICORT 80/4.5 twice daily.

2.3 Chronic Obstructive Pulmonary Disease
For patients with COPD the recommended dose is SYMBICORT 160/4.5, two inhalations twice daily.
If shortness of breath occurs in the period between doses, an inhaled, short-acting beta-agonist should be taken for immediate relief.

3. DOSAGE FORMS AND STRENGTHS
SYMBICORT is available as a metered-dose inhaler containing a combination of budesonide (80 or 160 mcg) and formoterol (4.5 mcg) as an inhalation aerosol in the following two strengths: 80/4.5 and 160/4.5. Each dosage strength contains 60 or 120 actuations per/canister. Each strength of SYMBICORT is supplied with a red plastic actuator with a gray dust cap.

3.1 CONTRAINDICATIONS
The use of SYMBICORT is contraindicated in the following conditions:
- Primary treatment of status asthmaticus or other acute episodes of asthma or COPD where intensive measures are required.
- Hypersensitivity to any of the ingredients in SYMBICORT.

3.2 WARNINGS AND PRECAUTIONS
5.1 Serious Asthma-Related Events – Hospitalizations, Intubations and Death
Use of LABA as monotherapy (without ICS) for asthma is associated with an increased risk of asthma-related death [see Salmeterol Multicenter Asthma Research Trial (SMART)]. Available data from controlled clinical trials also suggest that use of LABA as monotherapy increases the risk of asthma-related hospitalization in pediatric and adolescent patients. These findings are considered a class effect of LABA. When LABA are used in fixed-dose combination with ICS, data from large clinical trials do not show a significant increase in the risk of serious asthma-related events (hospitalizations, intubations, death) compared to ICS alone (see Serious Asthma-Related Events with ICS/LABA).

Serious Asthma-Related Events with ICS/LABA
Four large, 26-week, randomized, blinded, active-controlled clinical safety trials were conducted to evaluate the risk of serious asthma-related events when LABA were used in fixed-dose combination with ICS compared to ICS alone in patients with asthma. Three trials included adult and adolescent patients aged ≥12 years: one trial compared budesonide/formoterol (SYMBICORT) to budesonide [see Clinical Studies (14.1)]; one trial compared fluticasone propionate/salmeterol inhalation powder to fluticasone propionate inhalation powder; and one trial compared mometasone furoate/formoterol to mometasone furoate. The fourth trial included pediatric patients 4 to 11 years of age and compared fluticasone propionate/salmeterol inhalation powder to fluticasone propionate inhalation powder. The primary safety endpoint for all four trials was serious asthma-related events (hospitalizations, intubations and death). A blinded adjudication committee determined whether events were asthma-related.

The three adult and adolescent trials were designed to rule out a risk margin of 2.0, and the pediatric trial was designed to rule out a risk of 2.7. Each individual trial met its pre-specified objective and demonstrated non-inferiority of ICS/LABA to ICS alone. A meta-analysis of the three adult and adolescent trials did not show a significant increase in risk of a serious asthma-related event with ICS/LABA fixed-dose combination compared with ICS alone (Table 1). These trials were not designed to rule out all risk for serious asthma-related events with ICS/LABA compared with ICS.

<table>
<thead>
<tr>
<th>Event (endotracheal)</th>
<th>ICS/LABA vs ICS</th>
<th>Hazard ratio (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Asthma-related death</td>
<td>2.0</td>
<td>1.2</td>
</tr>
<tr>
<td>Asthma-related intubation</td>
<td>1.0</td>
<td>1.2</td>
</tr>
<tr>
<td>Asthma-related hospitalization</td>
<td>2.0 (24-hour stay)</td>
<td>1.2</td>
</tr>
</tbody>
</table>

ICS = inhaled Corticosteroid, LABA = Long-acting Beta-agonist
1. Randomized patients who had taken at least 1 dose of study drug. Planned treatment used for analysis.
2. Estimated using a Cox proportional hazards model of time to first event with baseline hazards stratified by each of the 3 trials.
3. Number of patients with event that occurred within 6 months after the first use of study drug or 7 days after the last date of study drug, whichever was later. Patients can have one or more events, but only the first event was counted for analysis. A single, blinded, independent adjudication committee determined whether events were asthma-related.

The pediatric safety trial included 6208 pediatric patients 4 to 11 years of age who received ICS/LABA (fluticasone propionate/salmeterol inhalation powder) or ICS (fluticasone propionate inhalation powder). In this trial, 27/3107 (0.9%) patients randomized to ICS/LABA and 21/3101 (0.7%) patients randomized to ICS experienced a serious asthma-related event. There were no asthma-related deaths or intubations. ICS/LABA did not show a significantly increased risk of a serious asthma-related event compared to ICS based on the pre-specified risk margin (2.7), with an estimated hazard ratio of time to first event of 1.29 (95% CI: 0.73, 2.27).

Salmeterol Multicenter Asthma Research Trial (SMART)
A 28-week, placebo-controlled U.S. trial that compared the safety of salmeterol with placebo, each added to usual asthma therapy, showed an increase in asthma-related deaths in patients receiving salmeterol (13/13,176 in patients treated with salmeterol vs. 3/13,179 in patients treated with placebo; relative risk: 4.37 [95% CI 1.25, 15.34]). Use of background ICS was not required in SMART. The increased risk of asthma-related death is considered a class effect of LABA monotherapy.

Formoterol Monotherapy Studies
Clinical studies with formoterol used as monotherapy suggested a higher incidence of serious asthma exacerbation in patients who received formoterol than in those who received placebo. The sizes of these studies were not adequate to precisely quantify the difference in serious asthma exacerbations between treatment groups.

5.2 Deterioration of Disease and Acute Episodes
SYMBICORT should not be initiated in patients during rapidly deteriorating or potentially life-threatening episodes of asthma or COPD. SYMBICORT has not been studied in patients with acutely deteriorating asthma or COPD. The initiation of SYMBICORT in this setting is not appropriate.

Increasing use of inhaled, short-acting beta-agonists is a marker of deteriorating asthma. In this situation, the patient requires immediate re-evaluation with reassessment of the treatment regimen, giving special consideration to the possible need for replacing the current strength of SYMBICORT with a higher strength, adding additional inhaled corticosteroid, or initiating systemic corticosteroids. Patients should not use more than 2 inhalations twice daily (morning and evening) of SYMBICORT.

SYMBICORT should not be used for the relief of acute symptoms, i.e., as rescue therapy for the treatment of acute episodes of bronchospasm. An inhaled, short-acting beta-agonist, not SYMBICORT, should be used to relieve acute symptoms such as shortness of breath.

When beginning treatment with SYMBICORT, patients who have been taking oral or inhaled, short-acting beta-agonists on a regular basis (e.g., 4 times a day) should be instructed to discontinue the regular use of these drugs.
5.3 Excessive Use of SYMBOICORT and Use with Other Long-Acting Beta-2-Agonists
As with other inhaled drugs containing beta2-adrenergic agents, SYMBOICORT should not be used more often than recommended, at higher doses than recommended, or in conjunction with other medications containing LABA, as an overdose may result. Clinically significant cardiovascular effects and fatalities have been reported in association with excessive use of inhaled sympathomimetic drugs. Patients using SYMBOICORT should not use an additional LABA (e.g., salmeterol, formoterol fumarate, arformoterol tartrate) for any reason, including prevention of exercise-induced bronchospasm (EIB) or the treatment of asthma or COPD.

5.4 Local Effects
In clinical studies, the development of localized infections of the mouth and pharynx with Candida species has occurred in patients treated with SYMBOICORT. When such an infection develops, it should be treated with appropriate local or systemic (i.e., oral antifungal) therapy while treatment with SYMBOICORT continues, but at times therapy with SYMBOICORT may need to be interrupted. Advise the patient to rinse his/her mouth with water without swallowing following inhalation to help reduce the risk of oropharyngeal candidiasis.

5.5 Pneumonia and Other Lower Respiratory Tract Infections
Physicians should remain vigilant for the possible development of pneumonia in patients with COPD as the clinical features of pneumonia and exacerbations frequently overlap. Lower respiratory tract infections, including pneumonia, have been reported following the inhaled administration of corticosteroids.

In a 6-month lung function study of 1704 patients with COPD, there was a higher incidence of lung infections other than pneumonia (e.g., bronchitis, viral lower respiratory tract infections, etc.) in patients receiving SYMBOICORT 160/4.5 (7.6%) than in those receiving SYMBOICORT 80/4.5 (3.2%), formoterol 4.5 mcg (4.6%) or placebo (3.8%). Pneumonia did not occur with greater incidence in the SYMBOICORT 160/4.5 group (1.1%) compared with placebo (1.3%). In a 12-month lung function study of 1964 patients with COPD, there was also a higher incidence of lung infections other than pneumonia in patients receiving SYMBOICORT 160/4.5 (9.1%) than in those receiving SYMBOICORT 80/4.5 (8.9%), formoterol 4.5 mcg (7.1%) or placebo (6.2%). Similar to the 6-month study, pneumonia did not occur with greater incidence in the SYMBOICORT 160/4.5 group (4.0%) compared with placebo (5.0%).

5.6 Immunosuppression
Patients who are on drugs that suppress the immune system are more susceptible to infection than healthy individuals. Chicken pox and measles, for example, can have a more serious or even fatal course in susceptible children or adults using corticosteroids. In such children or adults who have had chicken pox or measles and who are then exposed to varicella-zoster virus, the disease may be more severe than in those without a prior history of chicken pox or measles. Patients who are on drugs that suppress the immune system (e.g., corticosteroids, cyclophosphamide) should avoid contact with individuals who have chicken pox or measles. If exposure to chicken pox or measles occurs, appropriate treatment should be used.

5.7 Transferring Patients From Systemic Corticosteroid Therapy
Particular care is needed for patients who have been transferred from systemically active corticosteroids to inhaled corticosteroids because deaths due to adrenal insufficiency have occurred in patients with asthma during and after transfer from systemic corticosteroids to less systemically available inhaled corticosteroids. After withdrawal from systemic corticosteroids, a number of months are required for recovery of hypothalamic-pituitary-adrenal (HPA) function. Patients who have been previously maintained on 20 mg or more per day of prednisone (or its equivalent) may be most susceptible, particularly when their systemic corticosteroids have been almost completely withdrawn. During this period of HPA suppression, patients may exhibit signs and symptoms of adrenal insufficiency when exposed to trauma, surgery, or infection (particularly gastroenteritis) or other conditions associated with severe electrolyte loss. Although SYMBOICORT may provide control of asthma symptoms during these episodes, in recommended doses it supplies less than normal physiological amounts of glucocorticoid systemically and does NOT provide the mineralocorticoid activity that is necessary for coping with these emergencies. During periods of stress, a severe asthma attack or a severe COPD exacerbation, patients who have been withdrawn from systemic corticosteroids should be instructed to resume oral corticosteroids (in large doses) immediately and to contact their physicians for further instruction. These patients should also be instructed to carry a warning card indicating that they may need supplementary systemic corticosteroids during periods of stress, a severe asthma attack, or a severe COPD exacerbation.

Patients requiring oral corticosteroids should be weaned slowly from systemic corticosteroid use after transferring to SYMBOICORT. Prednisone reduction can be accomplished by reducing the daily prednisone dose by 2.5 mg on a weekly basis during therapy with SYMBOICORT. Lung function (mean forced expiratory volume in 1 second [FEV1] or morning peak expiratory flow [PEF]), beta-agonist use, and asthma or COPD symptoms should be carefully monitored during withdrawal of oral corticosteroids. In addition, patients should be observed for signs and symptoms of adrenal insufficiency, such as fatigue, lassitude, weakness, nausea and vomiting, and impaired wound healing.

Transfer of patients from systemic corticosteroid therapy to inhaled corticosteroids or SYMBOICORT may unmask conditions previously suppressed by the systemic corticosteroid therapy (e.g., rhinitis, conjunctivitis, eczema, arthritis, eosinophilic conditions). Some patients may experience symptoms of systemically active corticosteroid withdrawal (e.g., joint and/or muscular pain, lassitude, depression) despite maintenance or even improvement of respiratory function.

5.8 Hypercorticism and Adrenal Suppression
Budesonide, a component of SYMBOICORT, will often help control asthma and COPD symptoms with less suppression of HPA function than therapeutically equivalent oral doses of prednisone. Since budesonide is absorbed into the circulation and can be systemically active at higher doses, the beneficial effects of SYMBOICORT in minimizing HPA dysfunction may be expected only when recommended dosages are not exceeded and individual patients are titrated to the lowest effective dose.

Because of the possibility of systemic absorption of inhaled corticosteroids, patients treated with SYMBOICORT should be observed carefully for any evidence of systemic corticosteroid effects. Particular care should be taken in observing patients postoperatively or during periods of stress for evidence of inadequate adrenal response.

It is possible that systemic corticosteroid effects such as hypercorticism and adrenal suppression (including adrenal crisis) may appear in a small number of patients, particularly when budesonide is administered at higher than recommended doses over prolonged periods of time. If such effects occur, the dosage of SYMBOICORT should be reduced slowly, consistent with accepted procedures for reducing systemic corticosteroids and for management of asthma symptoms.

5.9 Drug Interactions With Strong Cytochrome P450 3A4 Inhibitors
Caution should be exercised when considering the coadministration of SYMBOICORT with ketoconazole, and other known strong CYP3A4 inhibitors (e.g., ritonavir, atazanavir, clarithromycin, indinavir, itraconazole, nefazodone, nelfinavir, saquinavir, telithromycin) because adverse effects related to increased systemic exposure to budesonide may occur [see Drug Interactions (7.1) and Clinical Pharmacology (12.3)].

5.10 Paradoxical Bronchospasm and Upper Airway Symptoms
As with other inhaled medications, SYMBOICORT can produce paradoxical bronchospasm, which may be life threatening. If paradoxical bronchospasm occurs following dosing with SYMBOICORT, it should be treated immediately with an inhaled, short-acting bronchodilator, SYMBOICORT should be discontinued immediately, and alternative therapy should be instituted.

5.11 Immediate Hypersensitivity Reactions
Immediate hypersensitivity reactions may occur after administration of SYMBOICORT, as demonstrated by cases of urticaria, angioedema, rash, and bronchospasm.

5.12 Cardiovascular and Central Nervous System Effects
Excessive beta-adrenergic stimulation has been associated with seizures, angina, hypertension or hypotension, tachycardia with rates up to 200 beats/min, arrhythmias, nervousness, headache, tremor, palpitation, nausea, dizziness, fatigue, malaise, and insomnia [see Overdosage (10)]. Therefore, SYMBOICORT, like all products containing sympathomimetic amines, should be used with caution in patients with cardiovascular disorders, especially coronary insufficiency, cardiac arrhythmias, and hypertension.

Formoterol, a component of SYMBOICORT, can produce a clinically significant cardiovascular effect in some patients as measured by pulse rate, blood pressure, and/or symptoms. Although such effects are uncommon after administration of formoterol at recommended doses, if they occur, the drug may need to be discontinued. In addition, beta-agonists have been reported to produce ECG changes, such as flattening of the T wave, prolongation of the QTc interval, and ST segment depression. The clinical significance of these findings is unknown. Fatalities have been reported in association with excessive use of inhaled sympathomimetic drugs.

5.13 Reduction in Bone Mineral Density
Decreases in bone mineral density (BMD) have been observed with long-term administration of inhaled corticosteroids. The clinical significance of small changes in BMD with regard to long-term consequences such as fracture is unknown. Patients with major risk factors for decreased bone mineral content, such as prolonged immobilization, family history of osteoporosis, postmenopausal status, tobacco use, advanced age, poor nutrition, or chronic use of drugs that can reduce bone mass (e.g., anticonvulsants, oral corticosteroids) should be monitored and treated with established standards of care. Since patients with COPD often have multiple risk factors for reduced BMD, an assessment of BMD should be considered prior to initiating SYMBOICORT and periodically thereafter. If significant reductions in BMD are seen, SYMBOICORT is still considered medically important for that patient's COPD therapy, use of medication to treat or prevent osteoporosis should be strongly considered.
The incidence of common adverse events in Table 2 below is based upon pooled data from three 12-week, double-blind, placebo-controlled clinical studies in which 401 adult and adolescent patients (148 males and 253 females) aged 12 years and older were treated with 2 inhalations of SYMBICORT 80/4.5 or SYMBICORT 160/4.5 twice daily. The SYMBICORT group was composed of mostly Caucasian (84%) patients with a mean age of 38 years, and a mean percent predicted FEV1 at baseline of 76 and 68 for the 80/4.5 mcg and 160/4.5 mcg treatment groups, respectively. Control arms for comparison included 2 inhalations of budesonide HFA metered dose inhaler (MDI) 80 or 160 mcg, formoterol dry powder inhaler (DPI) 4.5 mcg, or placebo (MDI and DPI) twice daily. Table 2 includes all adverse events that occurred at an incidence of ≥3% in any one SYMBICORT group and more commonly than in the placebo group with twice-daily dosing. In considering these data, the increased average duration of patient exposure for SYMBICORT patients should be taken into account, as incidences are not adjusted for an imbalance of treatment duration.

Table 2  Adverse reactions occurring at an incidence of ≥3% and more commonly than placebo in the SYMBICORT groups: pooled data from three 12-week, double-blind, placebo-controlled clinical asthma trials in patients 12 years and older

<table>
<thead>
<tr>
<th>Treatment</th>
<th>SYMBICORT 80/4.5</th>
<th>SYMBICORT 160/4.5</th>
<th>Budesonide (pMDI 80mcg)</th>
<th>Formoterol (DPI 4.5mcg)</th>
<th>Placebo</th>
</tr>
</thead>
<tbody>
<tr>
<td>Adverse Event</td>
<td>N = 277</td>
<td>N = 124</td>
<td>N = 101</td>
<td>N = 109</td>
<td>N = 237</td>
</tr>
<tr>
<td>Nasopharyngitis</td>
<td>10.5</td>
<td>9.7</td>
<td>14.0</td>
<td>11.0</td>
<td>10.1</td>
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<tr>
<td>Headache</td>
<td>6.5</td>
<td>11.6</td>
<td>11.6</td>
<td>12.8</td>
<td>8.9</td>
</tr>
<tr>
<td>Upper respiratory tract infection</td>
<td>7.6</td>
<td>10.5</td>
<td>8.3</td>
<td>9.2</td>
<td>7.6</td>
</tr>
<tr>
<td>Pharyngolaryngeal pain</td>
<td>6.1</td>
<td>8.9</td>
<td>5.0</td>
<td>7.3</td>
<td>3.0</td>
</tr>
<tr>
<td>Sinusitis</td>
<td>5.8</td>
<td>4.8</td>
<td>5.8</td>
<td>2.8</td>
<td>6.3</td>
</tr>
<tr>
<td>Influenza</td>
<td>3.2</td>
<td>2.4</td>
<td>6.6</td>
<td>0.9</td>
<td>3.0</td>
</tr>
<tr>
<td>Back pain</td>
<td>3.2</td>
<td>1.6</td>
<td>2.5</td>
<td>5.5</td>
<td>2.1</td>
</tr>
<tr>
<td>Nasal congestion</td>
<td>2.5</td>
<td>3.2</td>
<td>2.5</td>
<td>3.7</td>
<td>1.3</td>
</tr>
<tr>
<td>Stomach discomfort</td>
<td>1.1</td>
<td>6.5</td>
<td>2.5</td>
<td>4.6</td>
<td>1.3</td>
</tr>
<tr>
<td>Vomiting</td>
<td>1.4</td>
<td>3.2</td>
<td>0.8</td>
<td>2.8</td>
<td>1.7</td>
</tr>
<tr>
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<td>1.4</td>
<td>3.2</td>
<td>0</td>
<td>0</td>
<td>0.8</td>
</tr>
</tbody>
</table>

Average Duration of Exposure (days) 77.7 73.8 77.0 71.4 62.4 55.9

1. All treatments were administered as 2 inhalations twice daily.

Long-term safety - asthma clinical trials in patients 12 years and older

Long-term safety studies in adolescent and adult patients 12 years and older, treated for up to 1 year at doses up to 1200/80 mcg/day (64/12/18 mcg twice daily), revealed neither clinically important changes in the incidence nor new types of adverse events emerging after longer periods of treatment. Similarly, no significant or unexpected patterns of abnormalities were observed for up to 1 year in safety measures including chemistry, hematology, ECG, Holter monitor, and HPA-axis assessments.

Pediatric Patients 6 to Less Than 12 Years of Age

The safety data for pediatric patients aged 6 to less than 12 years is based on 1 trial of 12 weeks treatment duration. Patients (79 female and 105 male) receiving inhaled corticosteroid at trial entry were randomized to SYMBICORT 80/4.5 (n=92) or budesonide pMDI 80 mcg (n=92), 2 inhalations twice daily. The overall safety profile of these patients was similar to that observed in patients 12 years of age and older who received SYMBICORT 80/4.5 twice daily in studies of similar design. Common adverse reactions that occurred in patients treated with SYMBICORT 80/4.5 with a frequency of ≥3% and more frequently than patients treated only with budesonide pMDI 80 mcg included upper respiratory tract infection, pharyngitis, headache, and rhinitis.

6.2 Clinical Trials Experience in Chronic Obstructive Pulmonary Disease

The safety data described below reflect exposure to SYMBICORT 160/4.5 in 1783 patients. SYMBICORT 160/4.5 was studied in two placebo-controlled lung function studies (6 and 12 months in duration), and two active-controlled exacerbation studies (6 and 12 months in duration) in patients with COPD.

The incidence of common adverse events in Table 3 below is based upon pooled data from two double-blind, placebo-controlled lung function clinical studies (6 and 12 months in duration) in which 771 adult COPD patients (496 males and 275 females) 40 years of age and older were treated with SYMBICORT 160/4.5, 2 inhalations twice daily. Of these patients 651 were treated for 6 months and 366 were treated for 12 months. The SYMBICORT group was composed of mostly Caucasian (83%) patients with a mean age of 63 years, and a mean percent predicted FEV1 at baseline of 33%. Control arms for comparison included 2 inhalations of budesonide HFA (MDI) 160 mcg, formoterol (DPI) 4.5 mcg or placebo (MDI and DPI) twice daily. Table 3 includes all adverse events that occurred at an incidence of ≥3% in the SYMBICORT group and more commonly than in the placebo group. In considering these data, the increased average duration of patient exposure to SYMBICORT should be taken into account, as incidences are not adjusted for an imbalance of treatment duration.

Table 3  Adverse reactions occurring at an incidence of ≥3% and more commonly than placebo in the SYMBICORT groups: pooled data from two 12-month, double-blind, placebo-controlled clinical asthma trials in patients 12 years and older

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<thead>
<tr>
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<th>Budesonide (pMDI 80mcg)</th>
<th>Formoterol (DPI 4.5mcg)</th>
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<td>3.2</td>
<td>2.4</td>
<td>6.6</td>
<td>0.9</td>
<td>3.0</td>
</tr>
<tr>
<td>Back pain</td>
<td>3.2</td>
<td>1.6</td>
<td>2.5</td>
<td>5.5</td>
<td>2.1</td>
</tr>
<tr>
<td>Nasal congestion</td>
<td>2.5</td>
<td>3.2</td>
<td>2.5</td>
<td>3.7</td>
<td>1.3</td>
</tr>
<tr>
<td>Stomach discomfort</td>
<td>1.1</td>
<td>6.5</td>
<td>2.5</td>
<td>4.6</td>
<td>1.3</td>
</tr>
<tr>
<td>Vomiting</td>
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<td>3.2</td>
<td>0.8</td>
<td>2.8</td>
<td>1.7</td>
</tr>
<tr>
<td>Oral Candidiasis</td>
<td>1.4</td>
<td>3.2</td>
<td>0</td>
<td>0</td>
<td>0.8</td>
</tr>
</tbody>
</table>

Average Duration of Exposure (days) 77.7 73.8 77.0 71.4 62.4 55.9

6.1 Clinical Trials Experience in Asthma

Adolescent and Younger Patients 12 Years of Age and Older

The overall safety data in adults and adolescents are based upon 10 active- and placebo-controlled clinical trials in which 3393 patients ages 12 years and older (2052 females and 1341 males) with asthma of varying severity were treated with SYMBICORT 80/4.5 or 160/4.5 taken 2 inhalations once or twice daily for 12 to 52 weeks. In these trials, the patients on SYMBICORT had a mean age of 38 years and were predominantly Caucasian (82%).

The safety data for pediatric patients aged 6 to less than 12 years is based on 1 trial of 12 weeks treatment duration. Patients (79 female and 105 male) receiving inhaled corticosteroid at trial entry were randomized to SYMBICORT 80/4.5 (n=92) or budesonide pMDI 80 mcg (n=92), 2 inhalations twice daily. The overall safety profile of these patients was similar to that observed in patients 12 years of age and older who received SYMBICORT 80/4.5 twice daily in studies of similar design. Common adverse reactions that occurred in patients treated with SYMBICORT 80/4.5 with a frequency of ≥3% and more frequently than patients treated only with budesonide pMDI 80 mcg included upper respiratory tract infection, pharyngitis, headache, and rhinitis.

6.2 Clinical Trials Experience in Chronic Obstructive Pulmonary Disease

The safety data described below reflect exposure to SYMBICORT 160/4.5 in 1783 patients. SYMBICORT 160/4.5 was studied in two placebo-controlled lung function studies (6 and 12 months in duration), and two active-controlled exacerbation studies (6 and 12 months in duration) in patients with COPD.

The incidence of common adverse events in Table 3 below is based upon pooled data from two double-blind, placebo-controlled lung function clinical studies (6 and 12 months in duration) in which 771 adult COPD patients (496 males and 275 females) 40 years of age and older were treated with SYMBICORT 160/4.5, 2 inhalations twice daily. Of these patients 651 were treated for 6 months and 366 were treated for 12 months. The SYMBICORT group was composed of mostly Caucasian (83%) patients with a mean age of 63 years, and a mean percent predicted FEV1 at baseline of 33%. Control arms for comparison included 2 inhalations of budesonide HFA (MDI) 160 mcg, formoterol (DPI) 4.5 mcg or placebo (MDI and DPI) twice daily. Table 3 includes all adverse events that occurred at an incidence of ≥3% in the SYMBICORT group and more commonly than in the placebo group. In considering these data, the increased average duration of patient exposure to SYMBICORT should be taken into account, as incidences are not adjusted for an imbalance of treatment duration.
Vascular disorders:

Skin and subcutaneous tissue disorders:

In clinical studies, concurrent administration of SYMBICORT and other drugs, such as short-acting beta-agonists, intranasal corticosteroids, and antihistamines/decongestants has not resulted in an increased frequency of adverse reactions. No formal drug interaction studies have been performed with SYMBICORT.

Cardiac disorders: angina pectoris, tachycardia, atrial and ventricular tachyarrhythmias, atrial fibrillation, extrasystoles, palpitations

Endocrine disorders: hypercorticism, growth velocity reduction in pediatric patients

Eye disorders: cataract, glaucoma, increased intraocular pressure

Gastrointestinal disorders: oropharyngeal candidiasis, nausea

Immune system disorders: immediate and delayed hypersensitivity reactions, such as anaphylactic reaction, angioedema, bronchospasm, urticaria, exanthema, dermatitis, pruritus

Metabolic and nutrition disorders: hyperglycemia, hypokalemia

Musculoskeletal, connective tissue, and bone disorders: muscle cramps

Nervous system disorders: tremor, dizziness

Psychiatric disorders: behavior disturbances, sleep disturbances, nervousness, agitation, depression, restlessness

Respiratory, thoracic, and mediastinal disorders: dysphonia, cough, throat irritation

Skin and subcutaneous tissue disorders: skin bruising

Vascular disorders: hypotension, hypertension

7 DRUG INTERACTIONS

In clinical studies, concurrent administration of SYMBICORT and other drugs, such as short-acting beta-agonists, intranasal corticosteroids, and antihistamines/decongestants has not resulted in an increased frequency of adverse reactions. No formal drug interaction studies have been performed with SYMBICORT.

7.1 Inhibitors of Cytochrome P4503A4

The main route of metabolism of corticosteroids, including budesonide, a component of SYMBICORT, is via cytochrome P450 (CYP) isoenzyme 3A4 (CYP3A4). After oral administration of ketoconazole, a strong inhibitor of CYP3A4, the mean plasma concentration of orally administered budesonide increased. Concomitant administration of CYP3A4 may inhibit the metabolism of, and increase the systemic exposure to, budesonide. Caution should be exercised when considering the coadministration of SYMBICORT with long-term ketoconazole and other known strong CYP3A4 inhibitors (e.g., ritonavir, azatavir, clarithromycin, indinavir, iraconazole, nefazodone, neflavin, saquinavir, telithromycin) [See Warnings and Precautions (5.9)].

7.2 Monoamine Oxidase Inhibitors and Tricyclic Antidepressants

SYMBICORT should be administered with caution to patients being treated with monoamine oxidase inhibitors or tricyclic antidepressants, or within 2 weeks of discontinuation of such agents, because the action of a tricyclic antidepressant in patients with asthma. In this setting, cardioselective beta-blockers could be considered, although they should be administered with caution.

7.3 Beta-Adrenergic Receptor Blocking Agents

Beta-blockers (including eye drops) may not only block the pulmonary effect of beta-agonists, such as formoterol, a component of SYMBICORT, but may produce severe bronchospasm in patients with asthma. Therefore, patients with asthma should not normally be treated with beta-blockers. However, under certain circumstances, there may be no acceptable alternatives to the use of beta-adrenergic blocking agents in patients with asthma. In this setting, cardioselective beta-blockers could be considered, although they should be administered with caution.

7.4 Diuretics

The ECG changes and/or hypokalemia that may result from the administration of non-potassium-sparing diuretics (such as loop or thiazide diuretics) can be acutely worsened by beta-agonists, especially when the recommended dose of the beta-agonist is exceeded. Although the clinical significance of these effects is not known, caution is advised in the coadministration of SYMBICORT with non-potassium-sparing diuretics.

8 USE IN SPECIFIC POPULATIONS

8.1 Pregnancy

Risk Summary

There are no adequate and well-controlled studies of SYMBICORT or one of its individual components, formoterol fumarate, in pregnant women; however studies are available for the other component budesonide. In animal reproduction studies, SYMBICORT, administered by the inhalation route, was teratogenic, embryocidal, and reduced fetal weights in rats at less than the maximum recommended human daily inhalation dose (MRHID) on a mcg/m² basis. Budesonide alone, administered by the subcutaneous route, was teratogenic, embryocidal, and reduced fetal weights in rats and rabbits at less than the MRHID. These effects were not seen in rats that received inhaled doses up to 4 times the MRHID. Studies of pregnant women have not shown that inhaled budesonide alone increases the risk of abnormalities when administered during pregnancy. Experience with oral corticosteroids suggests that rodents are more prone to teratogenic effects from corticosteroid exposure than humans. Formoterol fumarate alone, administered by the oral route, was teratogenic in rats and rabbits at 1600 and 65,000 times the MRHID, respectively. Formoterol fumarate was also embryocidal, increased pup loss at birth and during lactation, and decreased pup weight in rats at 110 times the MRHID. These adverse effects generally occurred at large multiples of the MRHID when formoterol fumarate was administered by the oral route to achieve high systemic exposures. No teratogenic, embryocidal, or developmental effects were seen in rats that received inhalation doses up to 375 times the MRHID.

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

Clinical Considerations

Disease-Associated Maternal and/or Embryo/Fetal risk

In women with poorly or moderately controlled asthma, there is an increased risk of several perinatal adverse outcomes such as preclampsia in the mother and prematurity, low birth weight, and small for gestational age in the neonate. Pregnant women with asthma should be closely monitored and medication adjusted as necessary to maintain optimal asthma control.

Labor or Delivery

There are no well-controlled human studies that have investigated the effects of SYMBICORT during labor and delivery. Because of the potential for beta-agonist interference with uterine contractility, use of SYMBICORT during labor should be restricted to those patients in whom the benefits clearly outweigh the risk.

Data

Human Data

Studies of pregnant women have not shown that inhaled budesonide increases the risk of abnormalities when administered during pregnancy. The results from a large population-based prospective cohort epidemiological study reviewing data from three Swedish registries covering approximately 95% of the pregnancies from 1995-1997 (i.e., Swedish Medical Birth Registry, Registry of Congenital Malformations; Child Cardiology Registry) indicate no increased risk for congenital malformations from the use of inhaled budesonide during early pregnancy. Congenital malformations were studied in 2014 infants born to mothers reporting the use of inhaled budesonide for asthma in early pregnancy (usually 10-12 weeks after the last menstrual period), the period when most major organ malformations occur. The rate of recorded congenital malformations was similar compared to the general population rate (3.8% vs. 2.5%, respectively). In addition, after exposure to inhaled budesonide, the number of infants born with oral clefts was similar to the expected number in the normal population (4 children vs. 3.3, respectively). These same data were utilized in a second study bringing the total to 2534 infants whose mothers were exposed to inhaled budesonide. In this study, the rate of congenital malformations among infants whose mothers were exposed to inhaled budesonide during early pregnancy was not different from the rate for all newborn babies during the same period (3.6%).

Animal Data

SYMBICORT

In an embryo-fetal development study in pregnant rats dosed during the period of organogenesis from gestation days 6-16, SYMBICORT produced umbilical hernia in fetuses at doses less than the MRHID (on a mcg/m² basis at maternal inhaled doses of 12.0/6.8 mcg/kg/day and above). Fetal weights were reduced at approximately 5 and 3 times the MRHID, respectively (on an AUC basis at a maternal inhaled dose of 80/4.4 mcg/kg [budesonide/formoterol]). No teratogenic or embryocidal effects were detected at doses less than the MRHID (on a mcg/m² basis at a maternal inhaled dose of 2.5/0.14 mcg/kg/day).

Budesonide

In a fertility and reproduction study, male rats were subcutaneously dosed for 9 weeks and females for 2 weeks prior to pairing and throughout the mating period. Females were dosed up until weaning of their offspring. Budesonide caused a decrease in prenatal viability and weight gain at birth and during lactation, along with a decrease in maternal body-weight gain, at doses less than the MRHID (on a mcg/m² basis at maternal subcutaneous doses of 20 mcg/kg/day and above). No such effects were noted at a dose less than the MRHID (on a mcg/m² basis at a maternal subcutaneous dose of 5 mcg/kg/day).
In an embryo-fetal development study in pregnant rabbits dosed during the period of organogenesis from gestation days 6-18, budesonide produced fetal loss, decreased fetal weight, and skeletal abnormalities at doses less than the MRHDID (on a mcg/m² basis at a maternal subcutaneous dose of 25 mcg/kg/day). In an embryo-fetal development study in pregnant rats dosed during the period of organogenesis from gestation days 6-15, budesonide produced similar adverse fetal effects at doses approximately 8 times the MRHDID (on a mcg/m² basis at a maternal subcutaneous dose of 500 mcg/kg/day). In another embryo-fetal development study in pregnant rats, no teratogenic or embryocidal effects were seen at doses up to 4 times the MRHDID (on a mcg/m² basis at maternal inhalation doses up to 250 mcg/kg/day).

In a peri- and post-natal development study, rats dosed from gestation day 15 to postpartum day 21, budesonide had no effects on delivery, but did have an effect on growth and development of offspring. Offspring survival was reduced and surviving offspring had decreased mean body weights at birth and during lactation at doses less than the MRHDID and higher (on a mcg/m² basis at maternal subcutaneous doses of 20 mcg/kg/day and higher). These findings occurred in the presence of maternal toxicity.

Formoterol

In a fertility and reproductive study, male rats were orally dosed for 9 weeks and females for 2 weeks prior to pairing and throughout the mating period. Females were either dosed up to gestation day 19 or up until weaning of their offspring. Males were dosed up to 25 weeks. Umbilical hernia was observed in rat fetuses at oral doses 1600 times greater than the MRHDID (on a mcg/m² basis at maternal oral doses of 3000 mcg/kg/day and higher). Brachygnathia was observed in rat fetuses at a dose 8000 times the MRHDID (on a mcg/m² basis at a maternal oral dose of 15,000 mcg/kg/day). Pregnancy was prolonged at a dose 8000 times the MRHDID (on a mcg/m² basis at a maternal oral dose of 15,000 mcg/kg/day). Fetal and pup deaths occurred at doses approximately 1600 times the MRHDID and higher (on a mcg/m² basis at oral doses of 3000 mcg/kg/day and higher) during gestation.

In an embryo-fetal development study in pregnant rats dosed during the period of organogenesis from gestation days 6-15, no teratogenic, embryocidal or developmental effects were seen at doses up to 375 times the MRHDID (on a mcg/m² basis with maternal inhalation doses up to 690 mcg/kg/day).

In an embryo-fetal development study in pregnant rabbits dosed during the period of organogenesis from gestation days 6-18, subcapsular cysts on the liver were observed in the fetuses at a dose 65,000 times the MRHDID (on a mcg/m² basis with a maternal oral dose of 60,000 mcg/kg/day). No teratogenic effects were observed at doses up to 3800 times the MRHDID (on a mcg/m² basis at maternal oral doses up to 3500 mcg/kg/day).

In a pre- and post-natal development study, pregnant female rats received formoterol at oral doses of 0, 210, 840, and 3400 mcg/kg/day from gestation day 6 through the lactation period. Pup survival was decreased from birth to postpartum day 26 at doses 110 times the MRHDID and higher (on a mcg/m² basis at maternal oral doses of 210 mcg/kg/day and higher), although there was no evidence of a dose-response relationship. There were no treatment-related effects on the physical, functional, and behavioral development of rat pups.

8.2 Lactation

Risk Summary

There are no available data on the effects of SYMBICORT, budesonide or formoterol fumarate on the breastfed child or on milk production. Budesonide, like other inhaled corticosteroids, is secreted in human milk. Formoterol fumarate is present in rat milk. Data for SYMBICORT indicate that the potential for oral absorption of budesonide and formoterol fumarate in breast milk is trivial since only minute quantities of the budesonide available to the infant in breast milk, as a percentage of the maternal dose, would be expected. There is no clinical evidence that SYMBICORT therapy has an adverse effect on the breastfed infant from SYMBICORT or from the underlying maternal condition.

Data

Human data with budesonide delivered via dry powder inhaler indicates that the total daily oral dose of budesonide available in breast milk to the infant is approximately 0.3% to 1% of the dose ingested by the mother [see Clinical Pharmacology (12.3)]. For SYMBICORT, the dose of budesonide available to the infant in breast milk, as a percentage of the maternal dose, would be expected to be similar.

In the fertility and reproduction study in rats, plasma levels of formoterol were measured in pups on post-natal day 15 [see Use in Specific Populations (8.1)]. It was estimated that the maximum plasma concentration that the pups received from the maternal animal, at the highest dose of 15 mg/kg, after nursing was 4.4% (0.24 nmol/L for a litter vs. 5.5 nmol/L for the mother).

8.4 Pediatric Use

Safety and effectiveness of SYMBICORT in asthma patients 12 years of age and older have been established in studies up to 12 months. In the two 12-week, double-blind, placebo-controlled US pivotal studies 25 patients 12 to 17 years of age were treated with SYMBICORT twice daily [see Clinical Studies (14.1)]. Efficacy results in this age group were similar to those observed in patients 18 years and older. There were no obvious differences in the type or frequency of adverse events reported in this age group compared with patients 18 years of age and older.

The safety and effectiveness of SYMBICORT 80/4.5 in asthma patients 6 to less than 12 years of age have been established in studies of up to 12-week duration [see Clinical Studies (14.1)]. The safety profile in these patients was consistent to that observed in patients 12 years of age and older, and other reported clinical experience has not identified differences in responses between the elderly and younger patients.

With other products containing beta₂-agonists, special caution should be observed when using SYMBICORT in geriatric patients who have concomitant cardiovascular disease that could be adversely affected by beta₂-agonists.

8.5 Geriatric Use

The number of asthma patients treated with SYMBICORT twice daily in two 12-week studies and a 26-week postmarketing study, 751 were 65 years of age or older, of whom 141 were 75 years of age or older.

In the COPD studies of 6 to 12 months duration, 810 patients treated with SYMBICORT 160/4.5, two inhalations twice daily were 65 years old and above and of those, 177 patients were 75 years of age and older. No overall differences in safety or effectiveness were observed between these patients and younger patients, and other reported clinical experience has not identified differences in responses between the elderly and younger patients.

8.6 Hepatic Impairment

Formal pharmacokinetic studies using SYMBICORT have not been conducted in patients with hepatic impairment. However, since both budesonide and formoterol fumarate are predominantly cleared by hepatic metabolism, impairment of liver function may lead to accumulation of budesonide and formoterol fumarate in plasma. Therefore, patients with hepatic disease should be closely monitored.

8.7 Renal Impairment

Formal pharmacokinetic studies using SYMBICORT have not been conducted in patients with renal impairment.

10 OVERDOSAGE

SYMBICORT

SYMBICORT contains both budesonide and formoterol, therefore, the risks associated with overdosage for the individual components described below apply to SYMBICORT. In pharmacokinetic studies, single doses of 960/54 mcg (12 actuations of SYMBICORT 80/4.5) and 1280/36 mcg (8 actuations of 160/4.5) were administered to patients with COPD. A total of 1920/54 mcg (12 actuations of SYMBICORT 160/4.5) was administered as a single dose to both healthy subjects and patients with asthma. In a long-term active-controlled safety study in adolescent and adult asthma patients 12 years of age and older, SYMBICORT 160/4.5 was administered for up to 12 months at doses up to twice the highest recommended daily dose. There were no clinically significant adverse reactions observed in any of these studies.

Budesonide

The potential for acute toxic effects following overdosage of budesonide is low. If used at excessive doses for prolonged periods, systemic corticosteroid effects such as hypercorticism may occur [see Warnings and Precautions (5)]. Budesonide at five times the highest recommended dose (3200 mcg daily) administered to humans for 6 weeks caused a significant reduction (27%) in the plasma cortisol response to a 6-hour infusion of ACTH compared with placebo (+1%). The corresponding effect of 10 mg prednisone daily was a 35% reduction in the plasma cortisol response to ACTH.

Formoterol

An overdose of formoterol would likely lead to an exaggeration of effects that are typical for beta₂-agonists, which include bronchospasm. There is insufficient evidence to determine if dialysis is beneficial for overdosage of formoterol. Cardiac monitoring is recommended in cases of overdosage.
SYMBICORT® (budesonide/formoterol fumarate dihydrate) Inhalation Aerosol

11 DESCRIPTION

SYMBICORT 90/4.5 and SYMBICORT 180/4.5 each contain micronized budesonide and micronized formoterol fumarate dihydrate for oral inhalation only. Each SYMBICORT 90/4.5 and SYMBICORT 180/4.5 canister is formulated as a hydrofluoroalkane (HFA 227, 1,1,1,3,3,3-heptafluoroisopropane)-propelled pressurized metered dose inhaler containing either 60 or 120 actuations [see Dosage Forms and Strengths (3) and How Supplied/Storage and Handling (16)]. After priming, each actuation meter either 91.5/1. mcg or 181.5/1.5 mcg from the valve and delivers either 80/4.5 mcg, or 160/4.5 mcg (budesonide micronized/formoterol fumarate dihydrate micronized) from the actuator. The actual amount of drug delivered to the lung may depend on patient factors, such as the coordination between actuator and device and inspiration through the delivery system. SYMBICORT also contains povivone K25 USP as a suspending agent and polyethylene glycol 1000 NF as a lubricant.

SYMBICORT should be primed before using for the first time by releasing two test sprays into the air away from the face, shaking well for 5 seconds before each spray. In cases where the inhaler has not been used for more than 7 days or when it has been dropped, prime the Inhaler again by shaking well for 5 seconds before each spray and releasing two test sprays into the air away from the face.

One active component of SYMBICORT is budesonide, a corticosteroid designated chemically as (RS)-11, 16c, 17α-Tetrahydroxyprogna-1,4-diene-3,20-dione cyclic 16,17-acetal with butyraldehyde. Budesonide is provided as a mixture of two epimers (22R and 22S). The empirical formula of budesonide is C25H34O6 and its molecular weight is 430.9. Its structural formula is:

Formoterol fumarate dihydrate is a powder which is slightly soluble in water. Its octanol-water partition coefficient at pH 7.4 is 2.6. The pKa of formoterol fumarate dihydrate at 25°C is 7.9 for the phenolic group and 9.2 for the amino group.

12 CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

SYMBICORT contains both budesonide and formoterol; therefore, the mechanisms of action described below for the individual components apply to SYMBICORT. These drugs represent two classes of medications (a synthetic corticosteroid and a long-acting selective beta2-adrenoceptor agonist) that have different effects on clinical, physiological, and inflammatory indices of COPD and asthma.

Budesonide

Budesonide is an anti-inflammatory corticosteroid that exhibits potent glucocorticoid activity and weak mineralocorticoid activity. In standard in vitro and animal models, budesonide has approximately a 200-fold higher affinity for the glucocorticoid receptor and a 1000-fold higher topical anti-inflammatory potency than cortisol (rat corion oil ear edema assay). As a measure of systemic activity, budesonide is 40 times more potent than cortisol when administered subcutaneously and 25 times more potent when administered orally in the rat thymus involution assay.

In glucocorticoid receptor affinity studies, the 2α form of budesonide was two times as active as the 22S epimer. In vitro studies indicated that the two forms of budesonide do not interconvert.

Inflammation is an important component in the pathogenesis of COPD and asthma. Corticosteroids have a wide range of inhibitory activities against multiple cell types (e.g., mast cells, eosinophils, neutrophils, macrophages, and lymphocytes) and mediators (e.g., histamine, eosinoids, leukotrienes, and cytokines) involved in allergic and non-allergic-mediated inflammation. These anti-inflammatory actions of corticosteroids may contribute to their efficacy in COPD and asthma.

Studies in asthmatic patients have shown a favorable ratio between topical anti-inflammatory activity and systemic corticosteroid effects over a wide range of doses of budesonide. This is explained by a combination of a relatively high local anti-inflammatory effect, extensive first-pass hepatic degradation of orally absorbed drug (85%-95%), and the low potency of formed metabolites.

Formoterol

Formoterol fumarate is a long-acting selective beta2-adrenergic agonist (beta2-agonist) with a rapid onset of action. Inhaled formoterol fumarate acts locally in the lung as a bronchodilator. In vitro studies have shown that formoterol has more than 200-fold greater agonist activity at beta2-receptors than at beta1-receptors. The in vitro binding selectivity to beta2- over beta1-adrenoceptors is higher for formoterol than for albuterol (5 times), whereas salmeterol has a higher (3 times) beta2-selectivity ratio than formoterol.

Although beta2-receptors are the predominant adrenergic receptors in bronchial smooth muscle and beta1-receptors are the predominant receptors in the heart, there are also beta2-receptors in the human heart comprising 10% to 50% of the total beta-adrenergic receptors. The precise function of these receptors has not been established, but they raise the possibility that even highly selective beta2-agonists may have cardiac effects.

The pharmacologic effects of beta2-adrenoceptor agonists, including formoterol, are at least in part attributable to stimulation of intracellular adenyl cyclase, the enzyme that catalyzes the conversion of adenosine triphosphate (ATP) to cyclic-3′, 5′-adenosine monophosphate (cyclic AMP). Increased cyclic AMP levels cause relaxation of bronchial smooth muscle and inhibition of release of mediators of immediate hypersensitivity from cells, especially from mast cells. In vitro tests show that formoterol is an inhibitor of the release of mast cell mediators, such as histamine and leukotrienes, from the human lung. Formoterol also inhibits histamine-induced plasma albumin extravasation in anesthetized guinea pigs and inhibits allergen-induced eosinophil influx in dogs with airway hyperresponsiveness. The relevance of these in vitro and animal findings to humans is unknown.

12.2 Pharmacodynamics

Asthma

Cardiovascular effects: In a single-dose cross-over study involving 201 patients with persistent asthma, single-dose treatments of 4.5, 9, and 18 mcg of formoterol in combination with 320 mcg of budesonide delivered via SYMBICORT were compared to budesonide 320 mcg alone. Dose-related improvements in FEV1 were demonstrated when compared with budesonide. ECGs and blood samples for glucose and potassium were obtained post-dose. For SYMBICORT, small mean increases in serum glucose and decreases in serum potassium (+0.44 mmol/L and -0.25 mmol/L, respectively) were observed with increasing doses of formoterol, compared to budesonide. In ECGs, SYMBICORT produced small dose-related mean increases in heart rate (approximately 3 bpm at the highest dose), and QTc intervals (3-6 msec) compared to budesonide alone. No subject had a QT or QTc value ≥500 msec.

In the United States, five 12-week, active- and placebo-controlled studies and one 6-month active-controlled study evaluated 2976 patients aged 6 years and older with asthma. Systemic pharmacodynamic effects of formoterol (heart/pulse rate, blood pressure, QT interval, potassium, and glucose) were similar in patients treated with SYMBICORT, compared with patients treated with formoterol dry inhalation powder 4.5 mcg; 2 inhalations twice daily. No patient had a QT or QTc value ≥500 msec during treatment.

In three placebo-controlled studies in adolescents and adults with asthma, aged 12 years and older, a total of 1232 patients (553 patients in the SYMBICORT group) had evaluable continuous 24-hour electrocardiographic monitoring. Overall, there were no important differences in the occurrence of ventricular or supraventricular ectopy and no evidence of increased risk for clinically significant dysrhythmia in the SYMBICORT group compared to placebo.

Cardiovascular effects: In two COPD lung function studies, 6 months and 12 months in duration including 3668 COPD patients, no clinically important differences were seen in pulse rate, blood pressure, potassium, and glucose between SYMBICORT, the individual components of SYMBICORT, and placebo [see Clinical Studies (14.2)].

EGGs recorded at multiple clinic visits on treatment in both studies showed no clinically important differences for heart rate, PR interval, QRS duration, heart rate, signs of cardiac ischemia or arrhythmias between SYMBICORT 160/4.5 the monoproducts and placebo, all administered as 2 inhalations twice daily. Based on ECGs, 6 patients treated with SYMBICORT 160/4.5, 6 patients treated with formoterol 4.5 mcg, and 6 patients in the placebo group experienced atrial fibrillation or flutter that was not present at baseline. There were no cases of nonsustained ventricular tachycardia in the SYMBICORT 160/4.5, formoterol 4.5 mcg, or placebo group.

In the 12-month study, 502 patients had evaluable continuous 24-hour ECG (Holter) monitoring prior to the first dose and after approximately 1 and 4 months on treatment. No clinically important differences in ventricular or supraventricular arrhythmias, ventricular or supraventricular ectopic beats, or heart rate were observed among the groups treated with SYMBICORT 160/4.5, formoterol or placebo taken as 2 inhalations twice daily. Based on ECG (Holter) monitoring, one patient on SYMBICORT 160/4.5, no patients on formoterol 4.5 mcg, and three patients in the placebo group experienced atrial fibrillation or flutter that was not present at baseline.

Cardiovascular effects: Twenty-four hour urinary cortisol measurements were collected in a pooled subset (n=161) of patients from two COPD lung function studies. The data indicated approximately 30% lower mean 24-hour urinary free cortisol values following chronic administration (> 6 months) of SYMBICORT relative to placebo. SYMBICORT appeared to exhibit comparable cortisol suppression to budesonide 160 mcg alone or coadministration of budesonide 160 mcg and formoterol 4.5 mcg. For patients treated with SYMBICORT or formoterol up to 12 months, the percentage of patients who shifted from normal to low for this measure were generally comparable.

Other Budesonide Products

To confirm that systemic absorption is not a significant factor in the clinical efficacy of inhaled budesonide, a clinical study in patients with asthma was performed comparing...
Peak steady-state plasma concentrations of budesonide administered by DPI in adults with asthma averaged 0.6 and 1.6 nmol/L at doses of 180 mcg and 360 mcg twice daily, respectively. In asthmatic patients, budesonide showed a linear increase in AUC and C_max with increasing dose after both single and repeated dosing of inhaled budesonide.

**COPD Patients:** In a single-dose study, 12 inhalations of SYM-BICORT 80/4.5 (total dose 960/54 mcg) were administered to patients with COPD. Mean budesonide peak plasma concentration of 3.3 nmol/L occurred at 30 minutes following dosing. Budesonide systemic exposure was comparable between SYM-BICORT pMDI and coadministration of budesonide via dry powder inhaler and formoterol via a dry powder inhaler (budesonide 960 mcg and formoterol 54 mcg). In the same study, an open-label group of moderate asthma patients also received the same higher dose of SYM-BICORT. For budesonide, COPD patients exhibited 12% greater AUC and 10% lower C_max compared to asthma patients.

In the 6-month pivotal lung function clinical study, steady-state pharmacokinetic data of budesonide was obtained in a subset of COPD patients with treatment arms of SYM-BICORT pMDI 160/4.5. SYM-BICORT pMDI 80/4.5, budesonide 160 mcg, budesonide 160 mcg and formoterol 4.5 mcg given together, all administered as 2 inhalations twice daily. Budesonide peak plasma concentration of 1.2 nmol/L occurred at 21 minutes in healthy subjects (AUC and C_max with doses from 80 mcg to 160 mcg and was generally similar between the 3 treatment groups receiving the same dose of budesonide (SYM-BICORT pMDI 160/4.5, budesonide 160 mcg, budesonide 160 mcg and formoterol 4.5 mcg administered together).

Formoterol

Inhaled formoterol is rapidly absorbed; peak plasma concentrations are typically reached at the first plasma sampling time, within 5-10 minutes after dosing. As with many drug products for oral inhalation, it is likely that the majority of the inhaled formoterol delivered is swallowed and then absorbed from the gastrointestinal tract.

**Healthy Subjects:** Following administration of SYM-BICORT (160/4.5, two or four inhalations twice daily) for 5 days in healthy subjects, plasma concentration of formoterol generally increased in proportion to dose. The accumulation index for the group that received 2 inhalations twice daily was 1.77 for formoterol.

**Asthma patients:** In a single-dose study, higher than recommended doses of SYM-BICORT (12 inhalations of SYM-BICORT 160/4.5) were administered to patients with moderate asthma. Peak plasma concentration for formoterol of 136 pmol occurred at 10 minutes following dosing. Approximately 8% of the delivered dose of formoterol was recovered in the urine as unchanged drug.

In a repeat dose study, the highest recommended dose of SYM-BICORT (160/4.5, two inhalations twice daily) was administered to patients with moderate asthma. Peak formoterol plasma concentration occurred at 10 minutes after dosing. Formoterol exposure was slightly greater (-16-18%) from SYM-BICORT pMDI compared to coadministration of budesonide via a metered-dose inhaler and formoterol via a dry powder inhaler (total dose of budesonide 960 mcg and formoterol 54 mcg). In the same study, an open label group of moderate asthma patients received the same dose of SYM-BICORT. COPD patients exhibited 12-15% greater AUC and C_max for formoterol compared to asthma patients.

In the 6-month pivotal lung function clinical study, steady-state pharmacokinetic data of formoterol was obtained in a subset of COPD patients with treatment arms of SYM-BICORT pMDI 160/4.5, SYM-BICORT pMDI 80/4.5, formoterol 4.5 mcg, budesonide 160 mcg and formoterol 4.5 mcg given together, all administered as 2 inhalations twice daily. The systemic exposure of formoterol as evidenced by AUC, was about 30% and 16% higher from SYM-BICORT pMDI compared to formoterol alone treatment arm and coadministration of individual components of budesonide and formoterol treatment arm, respectively.

**Distribution**

**Budesonide:** The volume of distribution of budesonide was approximately 3 L/kg. It was 85%-90% bound to plasma proteins. Protein binding was constant over the concentration range (1-100 nmol/L) achieved with, and exceeding, recommended inhaled doses. Budesonide showed little or no binding to corticosteroid binding globulin. Budesonide rapidly equilibrated with red blood cells in a concentration independent manner with a blood plasma ratio of about 0.8.

**Formoterol:** Over the concentration range of 10-500 nmol/L, plasma protein binding for the RR and SS enantiomers of formoterol was 40% and 58%, respectively. The concentrations of formoterol used to assess the plasma protein binding were higher than those achieved in plasma following inhalation of a single 54 mcg dose.

**Metabolism**

**Budesonide:** In vitro studies with human liver homogenates have shown that budesonide was rapidly and extensively metabolized. Two major metabolites formed via cytochrome P450 (CYP) isoenzyme 3A4 (CYP3A4) catalyzed biotransformation have been isolated and identified as 16β-hydroxybudesonide and 6β-hydroxybudesonide. The corticosteroid activity of each of these two metabolites was less than 1% that of the parent compound. No qualitative differences between the in vitro and in vivo metabolic patterns were detected. Negligible metabolic inactivation was observed in human lung and serum preparations.

**Formoterol:** The primary metabolism of formoterol is by direct glucuronidation and by O-demethylation followed by conjugation to inactive metabolites. Secondary metabolic pathways include deamination and sulfate conjugation. CYP2D6 and CYP2C have been identified as being primarily responsible for O-demethylation.
Elimination

Budesonide: Budesonide was excreted in urine and feces in the form of metabolites. Approximately 60% of an intravenous radiolabeled dose was recovered in the urine. No unchanged budesonide was detected in the urine. The 22R form of budesonide was preferentially cleared by the liver with systemic clearance of 1.4 L/min vs. 1.0 L/min for the 22S form. The terminal half-life, 2 to 3 hours, was the same for both epimers and was independent of dose.

Formoterol: The excretion of formoterol was studied in four healthy subjects following simultaneous administration of radiolabeled formoterol via the oral and IV routes. In that study, 82% of the radiolabeled formoterol was excreted in the urine while 24% was eliminated in the feces.

Special Populations

Geriatric

The pharmacokinetics of SYMBICORT in geriatric patients have not been specifically studied.

Pediatric

Plasma concentrations of budesonide were measured following administration of four inhalations of SYMBICORT 160/4.5 in a single-dose study in pediatric patients with asthma, 6 to less than 12 years of age. Peak budesonide concentrations of 1.4 nmol/L occurred at 20 minutes post-dose. This study also demonstrated that the total systemic exposure to budesonide from SYMBICORT was approximately 39% lower than from inhaled budesonide via a dry powder inhaler that was also evaluated at the same delivered dose. The dose-normalized Cmax and AUCinf of budesonide following single dose inhalation in children 6 to less than 12 years of age were numerically lower than that observed in adults.

Following 2 inhalations of SYMBICORT 160/4.5 twice daily treatment, formoterol Cmax and AUCinf at steady state in children 6 to less than 12 years of age were comparable to that observed in adults.

Gender/Race

Specific studies to examine the effects of gender and race on the pharmacokinetics of SYMBICORT have not been conducted. Population PK analysis of the SYMBICORT data indicates that gender does not affect the pharmacokinetics of budesonide and formoterol. No conclusions can be drawn on the effect of race due to the low number of non-Caucasians evaluated for PK.

Nursing Mothers

The disposition of budesonide when delivered by inhalation from a dry powder inhaler at doses of 200 or 400 mcg twice daily for at least 3 months was studied in eight lactating women with asthma from 1 to 6 months postpartum. Systemic exposure to budesonide in these women appears to be comparable to that in non-lactating women with asthma from other studies. Breast milk obtained over eight hours post-dose revealed that the maximum concentration of budesonide for the 400 and 200 mcg total daily doses was 0.39 and 0.78 nmol/L, respectively, and occurred within 45 minutes after dosing. The estimated oral daily dose of budesonide from breast milk to the infant is approximately 0.007 and 0.014 mcg/kg/day for the two dose regimens used in this study, which represents approximately 0.3% to 1% of the dose ingested by the mother. Budesonide levels in plasma samples obtained from five infants at about 90 minutes after breastfeeding (and about 140 minutes after drug administration to the mother) were below quantifiable levels (<0.02 nmol/L in four infants and <0.04 nmol/L in one infant) [see Use in Specific Populations (8.2)].

Renal or Hepatic Insufficiency

There are no data regarding the specific use of SYMBICORT in patients with hepatic or renal impairment. Reduced liver function may affect the elimination of corticosteroids. Budesonide pharmacokinetics was affected by compromised liver function as evidenced by a doubled systemic availability after oral ingestion. The intravenous budesonide pharmacokinetics was, however, similar in cirrhotic patients and in healthy subjects. Specific data with formoterol is not available, but because formoterol is primarily eliminated via hepatic metabolism, an increased exposure can be expected in patients with severe liver impairment.

Drug-Drug Interactions

A single-dose crossover study was conducted to compare the pharmacokinetics of eight inhalations of the following: budesonide, formoterol, and budesonide plus formoterol administered concurrently. The results of the study indicated that there was no evidence of a pharmacokinetic interaction between the two components of SYMBICORT.

Inhibitors of Cytchrome P450 Enzymes

Ketoconazole: Ketoconazole, a strong inhibitor of cytochrome P450 (CYP) isoenzyme 3A4 (CYP3A4), the main metabolic enzyme for corticosteroids, increased plasma levels of orally ingested budesonide.

Cimetidine: At recommended doses, cimetidine, a non-specific inhibitor of CYP enzymes, had a slight but clinically insignificant effect on the pharmacokinetics of oral budesonide.

Specific drug-drug interaction studies with formoterol have not been performed.

13 NONCLINICAL TOXICOLOGY

13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility

Budesonide

Long-term studies were conducted in rats and mice using oral administration to evaluate the carcinogenic potential of budesonide. In a 2-year study in Sprague-Dawley rats, budesonide caused a statistically significant increase in the incidence of gliomas in male rats at an oral dose of 50 mcg/kg (approximately equivalent to the MRHDID in adults and children on a mcg/m² basis). No tumorigenicity was seen in male and female rats at respective oral doses up to 25 and 50 mcg/kg (approximately equivalent to the MRHDID in adults and children on a mcg/m² basis). In two additional 2-year studies in male Fischer and Sprague-Dawley rats, budesonide caused no gliomas at an oral dose of 50 mcg/kg (approximately equivalent to the MRHDID in adults and children on a mcg/m² basis). However, in the male Sprague-Dawley rats, budesonide caused a statistically significant increase in the incidence of hepatocellular tumors at an oral dose of 50 mcg/kg (approximately equivalent to the MRHDID in adults and children on a mcg/m² basis). The concurrent reference corticosteroids (methylprednisolone and triamcinolone acetonide) in these two studies showed similar findings.

In a 91-week study in mice, budesonide caused no treatment-related carcinogenicity at oral doses up to 200 mcg/kg (approximately 2 times the MRHDID in adults and children on a mcg/m² basis).

Budesonide was not mutagenic or clastogenic in six different test systems: Ames Salmonella/microsome plate test, mouse micronucleus test, mouse lymphoma test, chromosome aberration test in human lymphocytes, sex-linked recessive lethal test in Drosophila melanogaster, and DNA repair analysis in rat hepatocyte culture.

Fertility and reproductive performance were unaffected in rats at subcutaneous doses up to 80 mcg/kg (approximately equal to the MRHDID on a mcg/m² basis). However, it caused a decrease in prenatal viability and viability in the pups at birth and during lactation, along with a decrease in maternal body-weight gain, at subcutaneous doses of 20 mcg/kg and above (less than the MRHDID on a mcg/m² basis). No such effects were noted at 5 mcg/kg (less than the MRHDID on a mcg/m² basis).

Formoterol

Long-term studies were conducted in mice using oral administration and rats using inhalation administration to evaluate the carcinogenic potential of formoterol fumarate.

In a 24-month carcinogenicity study in CD-1 mice, formoterol at oral doses of 100 mcg/kg and above (approximately 30 and 15 times the MRHDID in adults and children, respectively, on a mcg/m² basis) caused a dose-related increase in the incidence of uterine leiomyomas.

In a 24-month carcinogenicity study in Sprague-Dawley rats, an increased incidence of mesonephric (uterine) leiomyomas and uterine leiomyosarcoma was observed at the mesonephric dose of 130 mcg/kg (approximately 70 and 35 times the MRHDID in adults and children, respectively, on a mcg/m² basis), no tumors were seen at 22 mcg/kg (approximately 12 and 8 times the MRHDID in adults and children, respectively, on a mcg/m² basis).

Other beta-agonist drugs have similarly demonstrated increases in leiomyomas of the genital tract in female rodents. The relevance of these findings to human use is unknown.

Formoterol was not mutagenic or clastogenic in Ames Salmonella/microsome plate test, mouse lymphoma test, chromosome aberration test in human lymphocytes, and rat micronucleus test. A reduction in fertility and/or reproductive performance was identified in male rats treated with formoterol at an oral dose of 15,000 mcg/kg (approximately 2200 times the MRHDID in adults and children, and 8600 times the MRHDID in children on a mcg/m² basis). No such effects were observed in the pups (approximately 1600 times the MRHDID on a mcg/m² basis). In a separate study with male rats treated with an oral dose of 15,000 mcg/kg (approximately 8000 times the MRHDID on a mcg/m² basis), there were findings of testicular tubular atrophy and spermatogenic debris in the testes and oesogephrasia in the epididymides. No effect on fertility was detected in female rats at doses up to 15,000 mcg/kg (approximately 1100 times the MRHDID on an AUC basis).

13.2 Animal Toxicology and/or Pharmacology

Preclinical: Studies in laboratory animals (minipigs, rodents, and dogs) have demonstrated the occurrence of cardiac arrhythmias and sudden death (with histologic evidence of myocardial necrosis) when beta-agonists and methylxanthines are administered concurrently. The clinical significance of these findings is unknown.

14 CLINICAL STUDIES

14.1 Asthma

Patients with asthma 12 years of age and older

In two clinical studies comparing SYMBICORT with the individual components, improvements in most efficacy end points were greater with SYMBICORT than with the use of either budesonide or formoterol alone. In addition, one clinical study showed similar results between SYMBICORT and the concurrent use of budesonide and formoterol at corresponding doses from separate inhalers. The safety and efficacy of SYMBICORT were demonstrated in two randomized, double-blind, placebo-controlled US clinical studies involving 1076 patients 12 years of age and older. Fixed SYMBICORT dosages of 160/0.9 mcg, and 320/0.9 mcg twice daily (each dose administered as 2 inhalations of the 80/4.5 and 160/4.5 mcg strengths, respectively) were compared with the monocomponents (budesonide and formoterol) and placebo to provide information about appropriate dosing to cover a range of asthma severity.

Study 1: Clinical Study with SYMBICORT 160/4.5

This 12-week study evaluated 596 patients 12 years of age and older by comparing SYMBICORT 160/4.5, the free combination of budesonide 160 mcg plus formoterol 4.5 mcg in separate inhalers; budesonide 162 mcg formoterol 4.5 mcg; and placebo; each administered as 2 inhalations twice daily. The study included a 2-week run-in period with budesonide 80 mcg, 2 inhalations twice daily. Most patients had moderate to severe asthma and were using moderate to high doses of inhaled corticosteroids prior to study entry. Randomization was stratified by previous inhaled corticosteroid treatment (71.8% on moderate- and 28.4% on high-dose inhaled corticosteroid). Mean percent predicted FEV1 at baseline was 68.1% and was similar across treatment groups. The co-primary efficacy end points were 12-hour-average post-dose FEV1, at week 2, and pre-dose FEV1, averaged over the course of the study. The study
also required that patients who satisfied a predefined asthma-worsening criterion be withdrawn. The predefined asthma-worsening criteria were a clinically important decrease in FEV1 or PEF, increase in rescue albuterol use, nighttime awakening due to asthma, emergency intervention or hospitalization due to asthma, or requirement for asthma medication not allowed by the protocol. For the criterion of nighttime awakening due to asthma, patients were allowed to remain in the study at the discretion of the investigator if none of the other asthma-worsening criteria were met. The percentage of patients withdrawing due to or meeting predefined criteria for worsening asthma is shown in Table 4.

Table 4 The number and percentage of patients withdrawing due to or meeting predefined criteria for worsening asthma (Study 1)

<table>
<thead>
<tr>
<th>Efficacy Variable</th>
<th>SYMBICORT 160/4.5</th>
<th>Budesonide 160 mcg</th>
<th>Formoterol 4.5 mcg</th>
<th>Placebo n = 125</th>
</tr>
</thead>
<tbody>
<tr>
<td>Patients withdrawn due to predefined asthma event¹</td>
<td>13 (10.5)</td>
<td>13 (11.3)</td>
<td>22 (20.2)</td>
<td>44 (35.8)</td>
</tr>
<tr>
<td>Patients with a predefined asthma event²</td>
<td>37 (29.8)</td>
<td>24 (20.9)</td>
<td>48 (44.0)</td>
<td>68 (55.3)</td>
</tr>
<tr>
<td>Decrease in FEV₁ in the treatment group</td>
<td>4 (3.2)</td>
<td>8 (7.0)</td>
<td>7 (6.4)</td>
<td>15 (12.2)</td>
</tr>
<tr>
<td>Rescue medication use</td>
<td>2 (1.6)</td>
<td>0</td>
<td>3 (2.8)</td>
<td>3 (2.4)</td>
</tr>
<tr>
<td>Decrease in AM PEF</td>
<td>2 (1.6)</td>
<td>5 (4.3)</td>
<td>5 (4.6)</td>
<td>17 (13.8)</td>
</tr>
<tr>
<td>Nighttime awakenings³</td>
<td>24 (19.4)</td>
<td>11 (9.6)</td>
<td>29 (26.6)</td>
<td>32 (26.0)</td>
</tr>
<tr>
<td>Clinical exacerbation</td>
<td>7 (5.6)</td>
<td>6 (5.2)</td>
<td>5 (4.6)</td>
<td>17 (13.8)</td>
</tr>
</tbody>
</table>

1. These criteria were assessed on a daily basis irrespective of the timing of the clinic visit, with the exception of FEV1, which was assessed at each clinic visit.
2. Individual criteria are shown for patients meeting any predefined asthma event, regardless of withdrawal status.
3. For the criterion of nighttime awakening due to asthma, patients were allowed to remain in the study at the discretion of the investigator if none of the other criteria were met.

Mean percent change from baseline in FEV1, measured immediately prior to dosing at each clinic visit, was a differential withdrawal rate in the treatment groups, pre-dose over 12 weeks is displayed in Figure 1. Because this study used predefined withdrawal criteria for pre-dose FEV1 at the end of treatment (0.19 L, 9.4%), compared with budesonide 160 mcg receiving SYMBICORT 160/4.5 had significantly greater mean improvements from baseline in pre-dose FEV1, which was assessed at each clinic visit. Patients receiving SYMBICORT 160/4.5 had clinically meaningful improvement in overall asthma-specific quality of life, as defined by a mean difference between treatment groups of >0.5 points in change from baseline in overall AQoL score (difference in AQoL score of 0.70 [95% CI 0.47, 0.93], compared to placebo).

The subjective impact of asthma on patients’ health-related quality of life was evaluated through the use of the standardized Asthma Quality of Life Questionnaire (AQoL(S)) (based on a 7-point scale where 1 = maximum impairment and 7 = no impairment). Patients receiving SYMBICORT 160/4.5 had a significantly greater mean improvement in overall asthma-specific quality of life, as defined by a mean difference between treatment groups of >0.5 points in change from baseline in overall AQoL score (difference in AQoL score of 0.70 [95% CI 0.47, 0.93], compared to placebo).

Table 5 Mean values for selected secondary efficacy variables (Study 1)

<table>
<thead>
<tr>
<th>Efficacy Variable</th>
<th>SYMBICORT 160/4.5 (n=124)</th>
<th>Budesonide 160 mcg plus Formoterol 4.5 mcg (n=109)</th>
<th>Budesonide 160 mcg (n=107)</th>
<th>Placebo (n=123)</th>
</tr>
</thead>
<tbody>
<tr>
<td>AM PEF (L/min) Baseline</td>
<td>341</td>
<td>338</td>
<td>342</td>
<td>339</td>
</tr>
<tr>
<td>Change from Baseline</td>
<td>9</td>
<td>-9</td>
<td>9</td>
<td>-18</td>
</tr>
<tr>
<td>PM PEF (L/min) Baseline</td>
<td>351</td>
<td>348</td>
<td>357</td>
<td>369</td>
</tr>
<tr>
<td>Change from Baseline</td>
<td>7</td>
<td>-7</td>
<td>7</td>
<td>-18</td>
</tr>
<tr>
<td>Albuterol rescue use</td>
<td>2.1</td>
<td>2.3</td>
<td>2.7</td>
<td>5</td>
</tr>
<tr>
<td>Baseline</td>
<td>-1.0</td>
<td>-1.5</td>
<td>-0.8</td>
<td>-0.3</td>
</tr>
<tr>
<td>Average symptom score/ day (0–3 scale) Baseline</td>
<td>0.99</td>
<td>1.03</td>
<td>1.04</td>
<td>1.04</td>
</tr>
<tr>
<td>Change from Baseline</td>
<td>-0.28</td>
<td>-0.32</td>
<td>-0.14</td>
<td>-0.05</td>
</tr>
</tbody>
</table>

1. Number of patients (n) varies slightly due to the number of patients for whom data were available for each variable. Results shown are based on last available data for each variable.

This 12-week study was similar in design to Study 1, and included 480 patients 12 years of age and older. This study compared SYMBICORT 80/4.5, budesonide 80 mcg, formoterol 4.5 mcg, and placebo; each administered as 2 inhalations twice daily. The study included a 2-week placebo run-in period. Most patients had mild to moderate asthma and were using low to moderate doses of inhaled corticosteroids prior to study entry. Mean percent predicted FEV1 at baseline was 71.3% and was similar across treatment groups. Efficacy variables and end points were identical to those in Study 1.

The percentage of patients withdrawing due to or meeting predefined criteria for worsening asthma is shown in Table 6. The method of assessment and criteria used were identical to that in Study 1.

Table 6 The number and percentage of patients withdrawing due to or meeting predefined criteria for worsening asthma (Study 2)

<table>
<thead>
<tr>
<th>Efficacy Variable</th>
<th>SYMBICORT 80/4.5 (n=122)</th>
<th>Budesonide 80 mcg (n=121)</th>
<th>Formoterol 4.5 mcg (n=114)</th>
<th>Placebo (n=122)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Patients withdrawn due to predefined asthma event¹</td>
<td>9 (7.3)</td>
<td>8 (6.6)</td>
<td>21 (18.4)</td>
<td>40 (32.8)</td>
</tr>
<tr>
<td>Patients with a predefined asthma event²</td>
<td>23 (18.7)</td>
<td>26 (21.5)</td>
<td>48 (42.1)</td>
<td>69 (56.6)</td>
</tr>
<tr>
<td>Decrease in FEV₁ in the treatment group</td>
<td>3 (2.4)</td>
<td>3 (2.5)</td>
<td>11 (9.6)</td>
<td>9 (7.4)</td>
</tr>
<tr>
<td>Rescue medication use</td>
<td>1 (0.8)</td>
<td>3 (2.5)</td>
<td>1 (0.9)</td>
<td>3 (2.5)</td>
</tr>
<tr>
<td>Decrease in AM PEF</td>
<td>3 (2.4)</td>
<td>1 (0.8)</td>
<td>8 (7.0)</td>
<td>14 (11.5)</td>
</tr>
<tr>
<td>Nighttime awakenings³</td>
<td>17 (13.8)</td>
<td>20 (16.5)</td>
<td>31 (27.2)</td>
<td>52 (42.6)</td>
</tr>
<tr>
<td>Clinical exacerbation</td>
<td>1 (0.8)</td>
<td>3 (2.5)</td>
<td>5 (4.4)</td>
<td>20 (16.4)</td>
</tr>
</tbody>
</table>

1. These criteria were assessed on a daily basis irrespective of the timing of the clinic visit, with the exception of FEV1, which was assessed at each clinic visit.
2. Individual criteria are shown for patients meeting any predefined asthma event, regardless of withdrawal status.
3. For the criterion of nighttime awakening due to asthma, patients were allowed to remain in the study at the discretion of the investigator if none of the other criteria were met.

Mean percent change from baseline in pre-dose FEV1 over 12 weeks is displayed in Figure 2.

The effect of SYMBICORT 160/4.5 two inhalations twice daily on selected secondary efficacy variables, including morning and evening PEF, albuterol rescue use, and asthma symptoms over 24 hours on a 0-3 scale is shown in Table 5.
Efficacy results for other secondary end points, including quality of life, were similar to those observed in Study 1.

Onset and Duration of Action and Progression of Improvement in Asthma Control
The onset of action and progression of improvement in asthma control were evaluated in the two pivotal clinical studies. The median time to onset of clinically significant bronchodilatation (>15% improvement in FEV1) was seen within 15 minutes. Maximum improvement in FEV1 occurred within 3 hours, and clinically significant improvement was maintained over 12 hours. Figures 3 and 4 show the percent change from baseline in post-dose FEV1 over 12 hours on the day of randomization and on the last day of treatment for Study 1.

Reduction in asthma symptoms and in albuterol rescue use, as well as improvement in morning and evening PEF, occurred within 1 day of the first dose of SYMBICORT; improvement in these variables was maintained over the 12 weeks of therapy.

Following the initial dose of SYMBICORT, FEV1 improved markedly during the first 2 weeks of treatment, continued to show improvement at the Week 6 assessment, and was maintained through Week 12 for both studies.

No diminution in the 12-hour bronchodilator effect was observed with either SYMBICORT 80/4.5 treatment, continued to show improvement at the Week 6 assessment, and was maintained compared to placebo for the primary endpoint of FEV1 to less than 12 years of age with asthma. Results showed a dose response of formoterol 9 mcg formoterol were administered in combination with budesonide in 54 pediatric patients available visit.

Figure 3 Mean Percent Change From Baseline in FEV1 on Day of Randomization (Study 1)

Figure 4 Mean Percent Change From Baseline in FEV1 at End of Treatment (Study 1)

Patients with asthma 6 to less than 12 years of age
The clinical program to support the efficacy of SYMBICORT 80/4.5 in children 6 to less than 12 years of age included the following: 1) a budesonide dose confirmatory study, 2) a formoterol dose finding study, and 3) an efficacy and safety study of the SYMBICORT combination product. The selection of budesonide 80 mcg is supported by a 6-week, randomized, double-blind, multicenter study in which SYMBICORT 80/4.5 was compared with budesonide pMDI 80 mcg, each administered as 2 inhalations twice daily over 26 weeks (NCT01444430). The primary safety objective was to evaluate whether the addition of formoterol to budesonide therapy (SYMBICORT) was non-inferior to budesonide in terms of the risk of serious asthma-related events (asthma-related hospitalization, endotracheal intubation, and death). The study was designed to rule out a pre-defined risk margin of serious asthma-related events of 2.0. A blinded adjudication committee determined whether events were asthma-related.

This study enrolled patients who were 12 years of age and older, had a clinical diagnosis of asthma for at least 1 year, and had at least 1 asthma exacerbation requiring treatment with systemic corticosteroids or an asthma-related hospitalization in the previous year. Patients were stratified to one of the two dose levels of SYMBICORT or budesonide based on assessment of asthma control and ongoing asthma therapy. Patients with a history of life-threatening asthma were excluded. The study included 11,693 patients (5846 receiving SYMBICORT 80/4.5 or 160/4.5) and 5847 receiving budesonide (80 or 160 mcg)), whose mean age was 44 years, and of whom 66% were female and 68% were Caucasian. SYMBICORT was non-inferior to budesonide in terms of time to first serious asthma-related events based on the pre-specified risk margin, with an estimated hazard ratio of 1.07 [95% CI: 0.70, 1.65] (Table 7).

Table 7 Serious Asthma-Related Events (Postmarketing Safety and Efficacy study)

<table>
<thead>
<tr>
<th>Event</th>
<th>SYMBICORT (N = 5846)</th>
<th>Budesonide (N = 5847)</th>
<th>SYMBICORT vs. Budesonide Hazard ratio (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Asthma-related death</td>
<td>2 (0.03)</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>Asthma-related endotracheal intubation</td>
<td>1 (0.01)</td>
<td>0</td>
<td></td>
</tr>
<tr>
<td>Asthma-related hospitalization</td>
<td>42 (0.7)</td>
<td>40 (0.7)</td>
<td>1.07 (0.70, 1.65)</td>
</tr>
</tbody>
</table>

1. N = number of patients
2. n = number of patients with the event
3. The hazard ratio for time to first event was based on a non-stratified Cox proportional hazard model with covariates of treatment (SYMBICORT vs. budesonide) and inhaled corticosteroid dose level (160 mcg vs. 80 mcg), as randomized. If the resulting upper 95% CI estimate for the relative risk was < 2.0, then non-inferiority was concluded.

The primary endpoint was asthma exacerbations, defined as a deterioration of asthma that led to use of systemic corticosteroids for at least 3 days, or a hospitalization, or an emergency room visit that required systemic corticosteroids. The estimated hazard ratio for time to first asthma exacerbation rate for SYMBICORT relative to budesonide was 0.84 [95% CI: 0.75, 0.94]. This outcome was primarily driven by a reduction in systemic corticosteroid use.
14.2 Chronic Obstructive Pulmonary Disease

Long Function

The efficacy of Symbicort 80/4.5 and Symbicort 160/4.5 in the maintenance treatment of airflow obstruction in COPD patients was evaluated in two randomized, double-blind, placebo-controlled multinational studies, conducted over 6 months (Study 1) and 12 months (Study 2), in a total of 3668 patients (2146 males and 1522 females). The majority of patients (93%) were Caucasian. All patients were required to be at least 40 years of age, with a FEV1 of less than or equal to 50% predicted, a clinical diagnosis of COPD with symptoms for at least 2 years, and a smoking history of at least 10 pack years, prior to entering the trial. The mean predose bronchodilator FEV1 at baseline of the patients enrolled in the study was 34% predicted. Forty-eight percent of the patients enrolled were on inhaled corticosteroids and 52.7% of patients were on short-acting anticholinergic bronchodilators during run-in. On randomization, inhaled corticosteroids were discontinued, and ipratropium bromide was allowed at a stable dose for those patients previously treated with short-acting anticholinergic bronchodilators. The co-primary efficacy variables in both studies were the change from baseline in average pre-dose and 1-hour post-dose FEV1 over the treatment period. The results of both studies 1 and 2 are described below.

Study 1

This was a 6-month, placebo-controlled study of 1704 COPD patients (mean % predicted FEV1 ranging from 33.5% to 37.4%) conducted to demonstrate the efficacy and safety of Symbicort in the treatment of airflow obstruction in COPD. The patients were randomized to one of the following treatment groups: Symbicort 160/4.5 (n=277), Symbicort 80/4.5 (n=281), budesonide 160 mcg + formoterol 4.5 mcg (n=278), budesonide 160 mcg (n=273), formoterol 4.5 mcg (n=284), or placebo (n=300). Patients receiving Symbicort 160/4.5, two inhalations twice daily, had significantly greater mean improvements from baseline in pre-dose FEV1 averaged over the treatment period (0.08 L, 10.7%) compared with formoterol 4.5 mcg (0.04 L, 6.9%) and placebo (0.01 L, 2.2%) (see Figure 6). Patients receiving Symbicort 80/4.5, two inhalations twice daily, did not have significantly greater improvement from baseline in the pre-dose FEV1, averaged over the treatment period compared with formoterol 4.5 mcg.

Figure 6 Mean Percent Change From Baseline in Pre-dose FEV1 over 6 Months (Study 1)

Patients receiving Symbicort 160/4.5, two inhalations twice daily, had significantly greater mean improvements from baseline in 1-hour post-dose FEV1, averaged over the treatment period [0.20 L, 22.6%], compared with budesonide 160 mcg [0.03 L, 4.9%] and placebo [0.03 L, 4.1%] (see Figure 7).

Figure 7 Mean Percent Change From Baseline in 1-hour Post-dose FEV1 Over 6 months (Study 1)

Study 2

This was a 12-month, placebo-controlled study of 1964 COPD patients (mean % predicted FEV1 at baseline ranging from 33.7% to 35.5%) conducted to demonstrate the efficacy and safety of Symbicort in the treatment of airflow obstruction in COPD. The patients were randomized to one of the following treatment groups: Symbicort 160/4.5 (n=494), Symbicort 80/4.5 (n=494), formoterol 4.5 mcg (n=495), or placebo (n=481). Patients receiving Symbicort 160/4.5, two inhalations twice daily, had significantly greater improvements from baseline in mean pre-dose FEV1, averaged over the treatment period [0.10 L, 10.8%] compared with formoterol 4.5 mcg (0.06 L, 7.3%) and placebo (0.01 L, 2.8%). Patients receiving Symbicort 80/4.5, two inhalations twice daily, did not have significantly greater improvements from baseline in the mean pre-dose FEV1, averaged over the treatment period compared to formoterol. Patients receiving Symbicort 160/4.5, two inhalations twice daily, also had significantly greater mean improvements from baseline in 1-hour post-dose FEV1, averaged over the treatment period [0.21 L, 24.0%] compared with placebo [0.02 L, 9.2%].

Serial FEV1, measures over 12 hours were obtained in a subset of patients in Study 1 (n=99) and Study 2 (n=121). The median time to onset of bronchodilation, defined as an FEV1 increase of 15% or greater from baseline, occurred at 5 minutes post-dose. Maximum improvement (calculated as the average change from baseline at each timepoint) in FEV1 occurred at approximately 2 hours post-dose.

In both Studies 1 and 2, improvements in secondary endpoints of morning and evening peak expiratory flow and reduction in rescue medication use were supportive of the efficacy of Symbicort 160/4.5.

Exacerbations

Studies 3 and 4 were primarily designed to evaluate the effect of Symbicort 160/4.5 on COPD exacerbations.

Study 3

This was a 6-month, active-control study conducted to evaluate the effect of Symbicort 160/4.5 compared to formoterol 4.5 mcg, each administered as 2 inhalations twice daily, on the rate of moderate and severe COPD exacerbations. COPD exacerbations were defined as worsening of 2 or more major symptoms (dyspnea, sputum volume, sputum color/purulence) or worsening of any 1 major symptom together with at least 1 of the minor symptoms: sore throat, colds (nasal discharge and/or nasal congestion), fever without other cause, increased cough or increased wheeze for at least 2 consecutive days. COPD exacerbations were considered of moderate severity if treatment of symptoms with systemic corticosteroids (≥3 days) and/or antibiotics were required, and were considered severe if hospitalization was required. The study randomized 1219 subjects to Symbicort 160/4.5 (606) and formoterol 4.5 mcg (613) of which 57% were male and 92% were Caucasian. They had a mean age of 64 years and a median smoking history of 39 pack years, with 46% identified as current smokers. At run-in, the mean post-bronchodilator % predicted normal FEV1 was 48.7% (range: 16.0% to 78.1%), and patients had a history of at least 1 COPD exacerbation in the previous year treated with systemic corticosteroids and/or hospitalization. All subjects were treated with Symbicort 160/4.5, two inhalations twice daily during a 4-week run-in period prior to being assigned trial treatment.

Study 4

This was a 12-month, active-control study which included 811 subjects treated with Symbicort 160/4.5 or formoterol 4.5 mcg, each administered as 2 inhalations twice daily. The study was conducted to evaluate for COPD exacerbation reduction in patients with COPD. COPD exacerbations were defined as worsening of COPD that required a course of oral steroids for treatment and/or hospitalization. This study randomized 407 subjects to Symbicort 160/4.5 and 404 to formoterol 4.5 mcg of which 61% were male and 83% were Caucasian. They had a mean age of 63 years and a median smoking history of 45 pack years, with 36% identified as current smokers. At run-in, the mean post-bronchodilator % predicted normal FEV1 was 37.8% (range: 11.75% to 76.50%), and a history of at least 1 COPD exacerbation in the previous year treated with systemic corticosteroids and/or antibiotics.

In Study 3, subjects treated with Symbicort 160/4.5, two inhalations twice daily had a significantly lower annual rate of moderate/severe COPD exacerbations compared with formoterol 4.5 mcg with a reduction of 26% (95% CI: 9%, 39%). In Study 4, a significantly lower annual rate of exacerbations was also observed in subjects treated with Symbicort 160/4.5 compared with formoterol 4.5 mcg with a reduction of 35% (95% CI: 20%, 47%) (Table 8).

<table>
<thead>
<tr>
<th>Treatment</th>
<th>n</th>
<th>Annual Rate Estimate</th>
<th>Rate ratio Symbicort 160/4.5 vs. Formoterol 4.5 mcg</th>
</tr>
</thead>
<tbody>
<tr>
<td>Study 3</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Symbicort 160/4.5</td>
<td>606</td>
<td>0.94</td>
<td>0.74</td>
</tr>
<tr>
<td>Formoterol 4.5 mcg</td>
<td>613</td>
<td>1.27</td>
<td></td>
</tr>
<tr>
<td>Study 4</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Symbicort 160/4.5</td>
<td>404</td>
<td>0.68</td>
<td>0.65</td>
</tr>
<tr>
<td>Formoterol 4.5 mcg</td>
<td>403</td>
<td>1.05</td>
<td></td>
</tr>
</tbody>
</table>

n = Number of patients included in efficacy analysis set.
Health-related quality of life was measured using the St. George’s Respiratory Questionnaire (SGRQ) in both COPD exacerbation clinical studies. In Study 3, the SGRQ responder rates at 6-months (defined as an improvement in score of 4 or more as a threshold) were 40% and 33% for SYMBICORT 160/4.5 vs. formoterol 4.5 mcg, respectively, with an odds ratio of 1.5 (95% CI: 1.0, 2.0) for SYMBICORT 160/4.5 vs. formoterol 4.5 mcg. In Study 4, the responder rates at 12-months were 50% and 49% for SYMBICORT 160/4.5 and formoterol 4.5 mcg, respectively, with an odds ratio of 1.0 (95% CI: 0.8, 1.4) for SYMBICORT 160/4.5 vs. formoterol 4.5 mcg.

### 16 HOW SUPPLIED/STORAGE AND HANDLING

SYMBICORT is available in two strengths and is supplied in the following package sizes:

<table>
<thead>
<tr>
<th>Package Size</th>
<th>NDC</th>
</tr>
</thead>
<tbody>
<tr>
<td>SYMBICORT Inhalations 80/4.5</td>
<td>0186-0372-20</td>
</tr>
<tr>
<td>SYMBICORT inhalations (institutional pack) 80/4.5</td>
<td>0186-0372-28</td>
</tr>
<tr>
<td>SYMBICORT Inhalations 160/4.5</td>
<td>0186-0370-20</td>
</tr>
<tr>
<td>SYMBICORT inhalations (institutional pack) 160/4.5</td>
<td>0186-0370-28</td>
</tr>
</tbody>
</table>

Each strength is supplied as a pressurized aluminum canister with an attached counting device, a red plastic actuator body with a white mouthpiece, and attached gray dust cap. Each 120 inhalation canister has a net fill weight of 10.2 grams and each 60 inhalation canister has a net fill weight of 6.9 grams (SYMBICORT 80/4.5) or 6 grams (SYMBICORT 160/4.5). Each canister is packaged in a foil overwrap pouch with desiccant sachet and placed into a carton. Each carton contains one canister and a Patient Information leaflet.

The SYMBICORT canister should only be used with the SYMBICORT actuator, and the SYMBICORT actuator should not be used with any other inhalation drug product.

The correct amount of medication in each inhalation cannot be ensured after the labeled number of inhalations from the canister have been used, even though the inhaler may not feel completely empty and may continue to operate. The inhaler should be discarded when the labeled number of inhalations have been used or within 3 months after removal from the foil pouch. Never immerse the canister into water to determine the amount remaining in the canister (“float test”).

Store at controlled room temperature 20°C to 25°C (68°F to 77°F) [see USP]. Store the inhaler with the mouthpiece down.

For best results, the canister should be at room temperature before use. Shake well for 5 seconds before using.

Keep out of the reach of children. Avoid spraying in eyes.

### CONTENTS UNDER PRESSURE.

Do not puncture or incinerate. Do not store near heat or open flame. Exposure to temperatures over 120°F may cause bursting. Never throw container into fire or incinerator.

### 17 PATIENT COUNSELING INFORMATION

**Serious Asthma-Related Events:** Inform patients with asthma that LABA when used alone increases the risk of asthma-related hospitalization or asthma-related death. Available data show that when ICS and LABA are used together, such as with SYMBICORT, there is not a significant increase in risk of these events.

**Not for Acute Symptoms:** Inform patients that SYMBICORT is not meant to relieve acute symptoms of asthma or COPD and extra doses should not be used for that purpose. Advise patients to treat acute symptoms with an inhaled, short-acting beta-agonist such as albuterol. Provide patients with such medication and instruct the patient in how it should be used.

**Instruct patients to seek medical attention immediately if they experience any of the following:**
- Decreasing effectiveness of inhaled, short-acting beta-agonists
- Need for more inhalations than usual of inhaled, short-acting beta-agonists
- Significant decrease in lung function as outlined by the physician

**Tell patients they should not stop therapy with SYMBICORT without physician/provider guidance since symptoms may recur after discontinuation.**

**Do Not Use Additional Long-Acting Beta-Agonists:** Instruct patients not to use other LABA for asthma and COPD.

**Local Effects:** Inform patients that localized infections with Candida albicans occurred in the mouth and pharynx in some patients. If oropharyngeal candidiasis develops, it should be treated with appropriate local or systemic (i.e., oral) antifungal therapy while still continuing therapy with SYMBICORT, but at times therapy with SYMBICORT may need to be temporarily interrupted under close medical supervision. Rinsing the mouth with water without swallowing after inhalation is advised to reduce the risk of thrush.

**Pneumonia:** Patients with COPD have a higher risk of pneumonia; instruct them to contact their healthcare provider if they develop symptoms of pneumonia.

**Immunosuppression:** Warn patients who are on immunosuppressant doses of corticosteroids to avoid exposure to chicken pox or measles and, if exposed, to consult their physician without delay. Inform patients of potential worsening of existing tuberculosis, fungal, bacterial, viral, or parasitic infections, or ocular herpes simplex.

**Hypercorticism and Adrenal Suppression:** Advise patients that SYMBICORT may cause systemic corticosteroid effects of hypercorticism and adrenal suppression. Additionally, inform patients that deaths due to adrenal insufficiency have occurred during and after transfer from systemic corticosteroids. Patients should taper slowly from systemic corticosteroids if transferring to SYMBICORT.

**Reduction in Bone Mineral Density:** Advise patients who are at an increased risk for decreased BMD that the use of corticosteroids may pose an additional risk.

**Reduced Growth Velocity:** Inform patients that orally inhaled corticosteroids, a component of SYMBICORT, may cause a reduction in growth velocity when administered to pediatric patients. Physicians should closely follow the growth of children and adolescents taking corticosteroids by any route.

**Ocular Effects:** Long-term use of inhaled corticosteroids may increase the risk of some eye problems (cataracts or glaucoma); consider regular eye examinations.

**Risks Associated With Beta-Agonist Therapy:** Inform patients of adverse effects associated with beta-agonists, such as palpitations, chest pain, rapid heart rate, tremor, or nervousness.

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Manufactured for: AstraZeneca Pharmaceuticals LP, Wilmington, DE 19850

By: AstraZeneca Dunkerque Production, Dunkerque, France

Product of France

Rev. 07/2019 US-31454 8/19
**Patient Information**

**SYMBCORT (SIM-bi-kort)**  
(budesonide 80 mcg and formoterol fumarate dihydrate 4.5 mcg)  
Inhalation Aerosol

**SYMBCORT (SIM-bi-kort)**  
(budesonide 160 mcg and formoterol fumarate dihydrate 4.5 mcg)  
Inhalation Aerosol

### What is SYMBCORT?

SYMBCORT combines an inhaled corticosteroid medicine (ICS), budesonide and a long-acting beta₂-adrenergic agonist (LABA) medicine, formoterol.

- Inhaled corticosteroids help to decrease inflammation in the lungs. Inflammation in the lungs can lead to breathing problems.
- LABA medicines are used in people with chronic obstructive pulmonary disease (COPD) and asthma. LABA medicines help the muscles around the airways in your lungs stay relaxed to prevent symptoms such as wheezing, cough, chest tightness, and shortness of breath. These symptoms can happen when the muscles around the airways tighten. This makes it hard to breathe. In severe cases, wheezing can stop your breathing and may lead to death if not treated right away.

**SYMBCORT is not used to relieve sudden breathing problems** and will not replace a rescue inhaler.

### SYMBCORT is used for asthma and COPD as follows:

- **Asthma:** SYMBCORT is used to control symptoms of asthma, and prevent symptoms such as wheezing in adults and children ages 6 and older. SYMBCORT contains formoterol. LABA medicines such as formoterol when used alone increase the risk of death and hospitalizations from asthma problems. SYMBCORT contains an ICS and a LABA. When an ICS and LABA are used together, there is not a significant increased risk in hospitalizations and death from asthma problems. SYMBCORT is not for adults and children with asthma who are well controlled with an asthma-control medicine, such as a low to medium dose of an ICS. SYMBCORT is for adults and children with asthma who need both an ICS and LABA medicine. It is not known if SYMBCORT is safe and effective in children less than 6 years of age with asthma.

- **COPD:** COPD is a long-term (chronic) lung disease that includes chronic bronchitis, emphysema, or both. SYMBCORT 160/4.5 mcg is used long-term, as 2 inhalations 2 times each day, to improve symptoms of COPD for better breathing and to reduce the number of flare-ups (the worsening of your COPD symptoms for several days).

### Do not use SYMBCORT:

- to treat sudden severe symptoms of asthma or COPD.
- if you are allergic to any of the ingredients in SYMBCORT. See the end of this leaflet for a list of ingredients in SYMBCORT.

### Before you use SYMBCORT, tell your healthcare provider about all of your medical conditions, including if you:

- have heart problems.
- have high blood pressure.
- have seizures.
- have thyroid problems.
- have diabetes.
- have liver problems.
- have osteoporosis.
- have an immune system problem.
- have eye problems such as increased pressure in the eye, glaucoma, or cataracts.
- are allergic to any medicines.
- have any type of viral, bacterial, fungal, or parasitic infection.
- are exposed to chicken pox or measles.
- are pregnant or plan to become pregnant. It is not known if SYMBCORT may harm your unborn baby.
- are breastfeeding. Budesonide, one of the active ingredients in SYMBCORT, passes into breast milk. You and your healthcare provider should decide if you will take SYMBCORT while breast-feeding.

Tell your healthcare provider about all the medicines you take including prescription and over-the-counter medicines, vitamins, and herbal supplements. SYMBCORT and certain other medicines may interact with each other. This may cause serious side effects. Especially tell your healthcare provider if you take antifungal or anti-HIV medicines.

Know all the medicines you take. Keep a list and show it to your healthcare provider and pharmacist each time you get a new medicine.
How should I use SYMBICORT?
See the step-by-step instructions for using SYMBICORT at the end of this Patient Information leaflet. Do not use SYMBICORT unless your healthcare provider has taught you and you understand everything. Ask your healthcare provider or pharmacist if you have any questions.

- Use SYMBICORT exactly as prescribed. Do not use SYMBICORT more often than prescribed. SYMBICORT comes in 2 strengths. Your healthcare provider has prescribed the strength that is best for you. Note the differences between SYMBICORT and your other inhaled medications, including the differences in prescribed use and physical appearance.
- Children should use SYMBICORT with an adult’s help, as instructed by the child’s healthcare provider.
- SYMBICORT should be taken every day as 2 puffs in the morning and 2 puffs in the evening about 12 hours apart.
- If you miss a dose of SYMBICORT, you should take your next dose at the same time you normally do.
- Rinse your mouth with water and spit the water out after each dose (2 puffs) of SYMBICORT. Do not swallow the water. This will help to lessen the chance of getting a fungus infection (thrush) in the mouth and throat.
- If you take too much SYMBICORT, call your healthcare provider or go to the nearest hospital emergency room right away if you have any unusual symptoms, such as worsening shortness of breath, chest pain, increased heart rate, or shakiness.
- Do not spray SYMBICORT in your eyes. If you accidentally get SYMBICORT in your eyes, rinse your eyes with water, and if redness or irritation persists, consult your healthcare provider.
- Do not change or stop any medicines used to control or treat your breathing problems. Your healthcare provider will change your medicines as needed.
- While you are using SYMBICORT 2 times each day, do not use other medicines that contain a LABA for any reason. Ask your healthcare provider or pharmacist if any of your other medicines are LABA medicines.
- SYMBICORT does not relieve sudden symptoms. Always have a rescue inhaler medicine with you to treat sudden symptoms. If you do not have a rescue inhaler, call your healthcare provider to have one prescribed for you.
- Call your healthcare provider or get medical care right away if:
  - your breathing problems worsen with SYMBICORT.
  - you need to use your rescue inhaler medicine more often than usual.
  - your rescue inhaler medicine does not work as well for you at relieving symptoms.
  - your peak flow meter results decrease. Your healthcare provider will tell you the numbers that are right for you.
  - your symptoms do not improve after using SYMBICORT regularly for 1 week.

What are the possible side effects of SYMBICORT?
SYMBICORT may cause serious side effects, including:

- Using too much of a LABA medicine may cause:
  - chest pain
  - a fast and irregular heartbeat
  - tremor
  - increased blood pressure
  - headache
  - nervousness

- Fungal infection in your mouth or throat (thrush). Rinse your mouth with water without swallowing after using SYMBICORT to help reduce your chance of getting thrush.

- Pneumonia and other lower respiratory tract infections. People with COPD have a higher chance of getting pneumonia and other lung infections. Inhaled corticosteroids may increase the chance of getting pneumonia. Call your healthcare provider if you notice any of these symptoms:
  - increase in mucus (sputum) production
  - fever
  - increased cough
  - change in mucus color
  - chills
  - increased breathing problems

- Immune system effects and a higher chance for infections. Tell your healthcare provider about any signs of infection such as:
  - fever
  - body aches
  - feeling tired
  - vomiting
  - pain
  - chills
  - nausea

- Adrenal insufficiency. Adrenal insufficiency is a condition in which the adrenal glands do not make enough steroid hormones. This can happen when you stop taking oral corticosteroid medicines and start inhaled corticosteroid medicine.

- Increased wheezing right after taking SYMBICORT. Always have a rescue inhaler with you to treat sudden wheezing.

- Serious allergic reactions including rash, hives, swelling of the face, mouth, and tongue, and breathing problems. Call your healthcare provider or get emergency medical care if you get any symptoms of a serious allergic reaction.

- Lower bone mineral density. This can happen in people who have a high chance for low bone mineral density (osteoporosis). Your healthcare provider should check you for this during treatment with SYMBICORT.

- Slowed growth in children. A child’s growth should be checked regularly while using SYMBICORT.

- Eye problems including glaucoma and cataracts. You should have regular eye exams while using SYMBICORT.

- Swelling of your blood vessels. This can happen in people with asthma. Tell your healthcare provider right away if you have:
  - a feeling of pins and needles or numbness of your arms or legs
  - rash
  - pain and swelling of the sinuses

- Decreases in blood potassium levels (hypokalemia).
- Increases in blood sugar levels (hyperglycemia).
The most common side effects of SYMBICORT include:

**People with asthma:**
- throat irritation
- upper respiratory tract infection
- inflammation of mucous membranes of the sinuses (sinusitis)
- back pain
- stomach discomfort
- thrush in the mouth and throat. Rinse your mouth with water without swallowing after use to help prevent thrush

**People with COPD:**
- throat irritation.
- thrush in the mouth and throat. Rinse your mouth with water without swallowing after use to help prevent thrush.
- infection and inflammation of the mucous membranes of the bronchial tubes (bronchitis).
- inflammation of mucous membranes in the sinuses (sinusitis).
- upper respiratory tract infection.

Tell your healthcare provider about any side effect that bothers you or that does not go away.

These are not all the possible side effects of SYMBICORT.

Tell your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.
You may also report side effects to AstraZeneca at 1-800-236-9933.

### How should I store SYMBICORT?
- Store SYMBICORT at room temperature between 68°F to 77°F (20°C to 25°C).
- Store SYMBICORT with the mouthpiece down.
- The contents of your SYMBICORT canister are under pressure. Do not puncture or throw the canister into a fire or incinerator. Do not use or store it near heat or open flame. Storage above 120°F may cause the canister to burst.
- Throw away SYMBICORT when the counter reaches zero (“0”) or 3 months after you take SYMBICORT out of its foil pouch, whichever comes first.
- Keep SYMBICORT and all medicines out of the reach of children.

### General Information about the safe and effective use of SYMBICORT.
Medicines are sometimes prescribed for purposes other than those listed in a Patient Information leaflet. Do not use SYMBICORT for a condition for which it was not prescribed. Do not give SYMBICORT to other people, even if they have the same symptoms that you have. It may harm them.

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

### What are the ingredients in SYMBICORT?
Active ingredients: micronized budesonide and micronized formoterol fumarate dihydrate
Inactive ingredients: hydrofluoroalkane (HFA 227), povidone K25 USP, and polyethylene glycol 1000 NF

Manufactured for: AstraZeneca Pharmaceuticals LP, Wilmington, DE 19850
By: AstraZeneca Dunkerque Production, Dunkerque, France  Product of France
For more information, call 1-800-236-9933 or go to www.MySymbicort.com.
How to Use SYMBICORT

Follow the instructions below for using SYMBICORT. If you have any questions, ask your doctor or pharmacist.

Preparing your SYMBICORT inhaler for use

1. Take your SYMBICORT out of the moisture-protective foil pouch before you use it for the first time and throw the foil away. Write the date that you open the foil pouch on the box.

2. A counter is attached to the top of the metal canister. The counter will count down each time you release a puff of SYMBICORT. The arrow points to the number of inhalations (puffs) left in the canister. The counter will stop counting at zero (“0”).

3. Use the SYMBICORT canister only with the red SYMBICORT inhaler supplied with the product. Parts of the SYMBICORT inhaler should not be used with parts from any other inhalation product.

4. Shake your SYMBICORT inhaler well for 5 seconds right before each use. Remove the mouthpiece cover by squeezing gently at both sides, then pulling out (see Figure 2). Check the mouthpiece for foreign objects before use.

5. Priming your SYMBICORT inhaler

Before you use SYMBICORT for the first time, you will need to prime it. To prime SYMBICORT, hold it in the upright position. See Figure 1. Shake the SYMBICORT inhaler well for 5 seconds. Hold your SYMBICORT inhaler facing away from you and press down firmly and fully on the top of the counter on the SYMBICORT inhaler to release a test spray. Then shake it again for 5 seconds and release a second test spray. Your SYMBICORT inhaler is now primed and ready for use.

If you do not use your SYMBICORT inhaler for more than 7 days or if you drop it, you will need to prime again.

Ways to hold the SYMBICORT inhaler for use

Using your SYMBICORT inhaler

6. Shake your SYMBICORT inhaler well for 5 seconds. Remove the mouthpiece cover. Check the mouthpiece for foreign objects.

7. Breathe out fully (exhale). Hold the SYMBICORT inhaler up to your mouth. Place the white mouthpiece fully into your mouth and close your lips around it. Make sure that the SYMBICORT inhaler is upright and that the opening of the mouthpiece is pointing towards the back of your throat (see Figure 5).

8. Breathe in (inhale) deeply and slowly through your mouth. Press down firmly and fully on the top of the counter on the SYMBICORT inhaler to release the medicine (see Figures 3 and 4).

9. Continue to breathe in (inhale) and hold your breath for about 10 seconds, or for as long as is comfortable. Before you breathe out (exhale), release your finger from the top of the counter. Keep the SYMBICORT inhaler upright and remove from your mouth.

10. Shake the SYMBICORT inhaler again for 5 seconds and repeat steps 7 to 9.
After using your SYMBICORT inhaler

11. After use, close the mouthpiece cover by pushing until it clicks in place.

12. After you finish taking SYMBICORT (2 puffs), rinse your mouth with water. Spit out the water. Do not swallow it.

Reading the counter

- The arrow on the counter on the top of the SYMBICORT inhaler points to the number of inhalations (puffs) left in your inhaler.
- The counter will count down each time you release a puff of medicine (either when priming your SYMBICORT inhaler or when taking the medicine).
- When the arrow on the counter approaches 20, you will notice the beginning of a yellow area letting you know that it is time to call your healthcare provider for a refill.
- It is important that you pay attention to the number of inhalations (puffs) left in your SYMBICORT inhaler by reading the counter. Throw away SYMBICORT when the counter shows zero (“0”) or 3 months after you take your SYMBICORT inhaler out of its foil pouch, whichever comes first. Your SYMBICORT inhaler may not feel empty and it may continue to operate, but you will not get the right amount of medicine if you keep using it. Use a new SYMBICORT inhaler and follow the instructions for priming (see instruction 5 above).

How to clean your SYMBICORT inhaler

Clean the white mouthpiece of your SYMBICORT inhaler every 7 days. To clean the mouthpiece:

- Remove the grey mouthpiece cover
- Wipe the inside and outside of the white mouthpiece opening with a clean, dry cloth
- Replace the mouthpiece cover
- **Do not put the SYMBICORT inhaler into water**
- **Do not try to take apart your SYMBICORT inhaler**