CENTER FOR DRUG EVALUATION AND RESEARCH

APPLICATION NUMBER:

206143Orig1s000

LABELING
CORLANOR (ivabradine) tablets, for oral use
Initial U.S. Approval: 2015

INDICATIONS AND USAGE
Corlanor (ivabradine) is a hyperpolarization-activated cyclic nucleotide-gated
channel blocker indicated to reduce the risk of hospitalization for worsening
heart failure in patients with stable, symptomatic chronic heart failure with left
ventricular ejection fraction \( \leq 35\% \), who are in sinus rhythm with resting heart
rate \( \geq 70 \) beats per minute and either are on maximally tolerated doses of beta-
blockers or have a contraindication to beta-blocker use. (1)

DOSAGE AND ADMINISTRATION
• Starting dose is 5 mg twice daily. After 2 weeks of treatment, adjust dose
  based on heart rate. The maximum dose is 7.5 mg twice daily. (2)
• In patients with conduction defects or in whom bradycardia could lead to
  hemodynamic compromise, initiate dosing at 2.5 mg twice daily. (2)

DOSAGE FORMS AND STRENGTHS
Tablets: 5 mg, 7.5 mg (3)

CONTRAINDICATIONS
• Acute decompensated heart failure (4)
• Blood pressure less than 90/50 mmHg (4)
• Sick sinus syndrome, sinoatrial block or 3rd degree AV block, unless a
  functioning demand pacemaker is present (4)
• Resting heart rate less than 60 bpm prior to treatment (4)
• Severe hepatic impairment (4)
• Pacemaker dependence (heart rate maintained exclusively by the
  pacemaker) (4)

ADVERSE REACTIONS
Most common adverse reactions occurring in \( \geq 1\% \) of patients are
bradycardia, hypertension, atrial fibrillation and luminous phenomena
(phosphenes). (6)

DRUG INTERACTIONS
• CYP3A4 inhibitors increase Corlanor plasma concentrations and
  CYP3A4 inducers decrease Corlanor plasma concentrations. (7.1)
• Negative chronotropes: Increased risk of bradycardia, monitor heart rate.
  (7.2)
• Pacemakers: Not recommended for use with demand pacemakers set to
  rates \( \geq 60 \) beats per minute. (7.3)

USE IN SPECIFIC POPULATIONS
• Lactation: Breastfeeding not recommended. (8.2)

See 17 for PATIENT COUNSELING INFORMATION and Medication
Guide.

Revised: 04/2015

Reference ID: 3732763
1. INDICATIONS AND USAGE
Corlanor is indicated to reduce the risk of hospitalization for worsening heart failure in patients with stable, symptomatic chronic heart failure with left ventricular ejection fraction ≤ 35%, who are in sinus rhythm with resting heart rate ≥ 70 beats per minute and either are on maximally tolerated doses of beta-blockers or have a contraindication to beta-blocker use.

2. DOSAGE AND ADMINISTRATION
The recommended starting dose of Corlanor is 5 mg twice daily with meals. Assess patient after two weeks and adjust dose to achieve a resting heart rate between 50 and 60 beats per minute (bpm) as shown in Table 1. Thereafter, adjust dose as needed based on resting heart rate and tolerability. The maximum dose is 7.5 mg twice daily.

In patients with a history of conduction defects, or other patients in whom bradycardia could lead to hemodynamic compromise, initiate therapy at 2.5 mg twice daily before increasing the dose based on heart rate [see Warnings and Precautions (5.3)].

<table>
<thead>
<tr>
<th>Heart Rate</th>
<th>Dose Adjustment</th>
</tr>
</thead>
<tbody>
<tr>
<td>&gt; 60 bpm</td>
<td>Increase dose by 2.5 mg (given twice daily) up to a maximum dose of 7.5 mg twice daily</td>
</tr>
<tr>
<td>50-60 bpm</td>
<td>Maintain dose</td>
</tr>
<tr>
<td>&lt; 50 bpm or signs and symptoms of bradycardia</td>
<td>Decrease dose by 2.5 mg (given twice daily); if current dose is 2.5 mg twice daily, discontinue therapy*</td>
</tr>
</tbody>
</table>

*[see Warnings and Precautions (5.3).]*

3. DOSAGE FORMS AND STRENGTHS
Corlanor 5 mg: salmon-colored, oval-shaped, film-coated tablet, scored on both edges, debossed with “5” on one face and bisected on the other face. The tablet is scored and can be divided into equal halves to provide a 2.5 mg dose.
Corlanor 7.5 mg: salmon-colored, triangular-shaped, film-coated tablet debossed with “7.5” on one face and plain on the other face.
4. **CONTRAINDICATIONS**

Corlanor is contraindicated in patients with:

- Acute decompensated heart failure
- Blood pressure less than 90/50 mmHg
- Sick sinus syndrome, sinoatrial block, or 3rd degree AV block, unless a functioning demand pacemaker is present
- Resting heart rate less than 60 bpm prior to treatment [see Warnings and Precautions (5.3)]
- Severe hepatic impairment [see Use in Specific Populations (8.6)]
- Pacemaker dependence (heart rate maintained exclusively by the pacemaker) [see Drug Interactions (7.3)]
- Concomitant use of strong cytochrome P450 3A4 (CYP3A4) inhibitors [see Drug Interactions (7.1)]

5. **WARNINGS AND PRECAUTIONS**

5.1 **Fetal Toxicity**

Corlanor may cause fetal toxicity when administered to a pregnant woman based on findings in animal studies. Embryo-fetal toxicity and cardiac teratogenic effects were observed in fetuses of pregnant rats treated during organogenesis at exposures 1 to 3 times the human exposures (AUC\textsubscript{0-24hr}) at the maximum recommended human dose (MRHD) [see Use in Specific Populations (8.1)]. Advise females to use effective contraception when taking Corlanor [see Use in Specific Populations (8.3)].

5.2 **Atrial Fibrillation**

Corlanor increases the risk of atrial fibrillation. In SHIFT, the rate of atrial fibrillation was 5.0% per patient-year in patients treated with Corlanor and 3.9% per patient-year in patients treated with placebo [see Clinical Studies (14)]. Regularly monitor cardiac rhythm. Discontinue Corlanor if atrial fibrillation develops.

5.3 **Bradycardia and Conduction Disturbances**

Bradycardia, sinus arrest, and heart block have occurred with Corlanor. The rate of bradycardia was 6.0% per patient-year in patients treated with Corlanor (2.7% symptomatic; 3.4% asymptomatic) and 1.3% per patient-year in patients treated with placebo. Risk factors for bradycardia include sinus node dysfunction, conduction defects (e.g., 1\textsuperscript{st} or 2\textsuperscript{nd} degree atrioventricular block, bundle branch block), ventricular dyssynchrony, and use of other negative chronotropes (e.g., digoxin, diltiazem, verapamil, amiodarone). Concurrent use of verapamil or diltiazem will increase Corlanor exposure, may themselves contribute to heart rate lowering, and should be avoided [see Clinical Pharmacology (12.3)]. Avoid use of Corlanor in
patients with 2nd degree atrioventricular block, unless a functioning demand pacemaker is present [see Contraindications (4) and Dosage and Administration (2)].

6. ADVERSE REACTIONS

Clinically significant adverse reactions that appear in other sections of the labeling include:

- Fetal Toxicity [see Warnings and Precautions (5.1)]
- Atrial Fibrillation [see Warnings and Precautions (5.2)]
- Bradycardia and Conduction Disturbances [see Warnings and Precautions (5.3)]

6.1 Clinical Trials Experience

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

In the Systolic Heart failure treatment with the I<sub>r</sub> inhibitor ivabradine Trial (SHIFT), safety was evaluated in 3260 patients treated with Corlanor and 3278 patients given placebo. The median duration of Corlanor exposure was 21.5 months.

The most common adverse drug reactions in the SHIFT trial are shown in Table 2 [see also Warnings and Precautions (5.2), (5.3)].

### Table 2. Adverse Drug Reactions with Rates ≥ 1.0% Higher on Ivabradine than Placebo occurring in > 1% on Ivabradine in SHIFT

<table>
<thead>
<tr>
<th></th>
<th>Ivabradine (N=3260)</th>
<th>Placebo (N=3278)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bradycardia</td>
<td>10%</td>
<td>2.2%</td>
</tr>
<tr>
<td>Hypertension, blood pressure increased</td>
<td>8.9%</td>
<td>7.8%</td>
</tr>
<tr>
<td>Atrial fibrillation</td>
<td>8.3%</td>
<td>6.6%</td>
</tr>
<tr>
<td>Phosphenes, visual brightness</td>
<td>2.8%</td>
<td>0.5%</td>
</tr>
</tbody>
</table>

Reference ID: 3732763
Luminous Phenomena (Phosphenes)
Phosphenes are phenomena described as a transiently enhanced brightness in a limited area of the visual field, halos, image decomposition (stroboscopic or kaleidoscopic effects), colored bright lights, or multiple images (retinal persistency). Phosphenes are usually triggered by sudden variations in light intensity. Corlanor can cause phosphenes, thought to be mediated through Corlanor’s effects on retinal photoreceptors [see Clinical Pharmacology (12.1)]. Onset is generally within the first 2 months of treatment, after which they may occur repeatedly. Phosphenes were generally reported to be of mild to moderate intensity and led to treatment discontinuation in < 1% of patients; most resolved during or after treatment.

6.2 Postmarketing Experience
Because these reactions are reported voluntarily from a population of uncertain size, it is not always possible to estimate their frequency reliably or establish a causal relationship to drug exposure.

The following adverse reactions have been identified during post-approval use of Corlanor: syncope, hypotension, angioedema, erythema, rash, pruritus, urticaria, vertigo, diplopia, and visual impairment.

7. DRUG INTERACTIONS
7.1 Cytochrome P450-Based Interactions
Corlanor is primarily metabolized by CYP3A4. Concomitant use of CYP3A4 inhibitors increases ivabradine plasma concentrations, and use of CYP3A4 inducers decreases them. Increased plasma concentrations may exacerbate bradycardia and conduction disturbances.

The concomitant use of strong CYP3A4 inhibitors is contraindicated [see Contraindications (4) and Clinical Pharmacology (12.3)]. Examples of strong CYP3A4 inhibitors include azole antifungals (e.g., itraconazole), macrolide antibiotics (e.g., clarithromycin, telithromycin), HIV protease inhibitors (e.g., nelfinavir), and nefazodone.

Avoid concomitant use of moderate CYP3A4 inhibitors when using Corlanor. Examples of moderate CYP3A4 inhibitors include diltiazem, verapamil, and grapefruit juice [see Warnings and Precautions (5.3) and Clinical Pharmacology (12.3)].

Avoid concomitant use of CYP3A4 inducers when using Corlanor. Examples of CYP3A4 inducers include St. John’s wort, rifampicin, barbiturates, and phenytoin [see Clinical Pharmacology (12.3)].
7.2 Negative Chronotropes
Most patients receiving Corlanor will also be treated with a beta-blocker. The risk of bradycardia increases with concomitant administration of drugs that slow heart rate (e.g., digoxin, amiodarone, beta-blockers). Monitor heart rate in patients taking Corlanor with other negative chronotropes.

7.3 Pacemakers
Corlanor dosing is based on heart rate reduction, targeting a heart rate of 50 to 60 beats per minute [see Dosage and Administration (2)]. Patients with demand pacemakers set to a rate \( \geq 60 \) beats per minute cannot achieve a target heart rate < 60 beats per minute, and these patients were excluded from clinical trials [see Clinical Studies (14)]. The use of Corlanor is not recommended in patients with demand pacemakers set to rates \( \geq 60 \) beats per minute.

8. USE IN SPECIFIC POPULATIONS
8.1 Pregnancy
Risk Summary
Based on findings in animals, Corlanor may cause fetal harm when administered to a pregnant woman. There are no adequate and well-controlled studies of Corlanor in pregnant women to inform any drug-associated risks. In animal reproduction studies, oral administration of ivabradine to pregnant rats during organogenesis at a dosage providing 1 to 3 times the human exposure (AUC\(_{0-24\text{hr}}\)) at the MRHD resulted in embryo-fetal toxicity and teratogenicity manifested as abnormal shape of the heart, interventricular septal defect, and complex anomalies of primary arteries. Increased postnatal mortality was associated with these teratogenic effects in rats. In pregnant rabbits, increased post-implantation loss was noted at an exposure (AUC\(_{0-24\text{hr}}\)) 5 times the human exposure at the MRHD. Lower doses were not tested in rabbits. The background risk of major birth defects for the indicated population is unknown. The estimated background risk of major birth defects in the U.S. general population is 2 to 4%, however, and the estimated risk of miscarriage is 15 to 20% in clinically recognized pregnancies. Advise a pregnant woman of the potential risk to the fetus.

Clinical Considerations
Disease-associated maternal and/or embryo/fetal risk
Stroke volume and heart rate increase during pregnancy, increasing cardiac output, especially during the first trimester. Pregnant patients with left ventricular ejection fraction less than 35% on maximally tolerated doses of beta-blockers may be particularly heart-rate dependent for augmenting cardiac output. Therefore, pregnant patients who are started on Corlanor, especially during the first trimester, should be
followed closely for destabilization of their congestive heart failure that could result from heart rate slowing.

Monitor pregnant women with chronic heart failure in 3rd trimester of pregnancy for preterm birth.

Data
Animal Data
In pregnant rats, oral administration of ivabradine during the period of organogenesis (gestation day 6-15) at doses of 2.3, 4.6, 9.3, or 19 mg/kg/day resulted in fetal toxicity and teratogenic effects. Increased intrauterine and post-natal mortality and cardiac malformations were observed at doses ≥ 2.3 mg/kg/day (equivalent to the human exposure at the MRHD based on AUC$_{0-24}$h). Teratogenic effects including interventricular septal defect and complex anomalies of major arteries were observed at doses ≥ 4.6 mg/kg/day (approximately 3 times the human exposure at the MRHD based on AUC$_{0-24}$h).

In pregnant rabbits, oral administration of ivabradine during the period of organogenesis (gestation day 6-18) at doses of 7, 14, or 28 mg/kg/day resulted in fetal toxicity and teratogenicity. Treatment with all doses ≥ 7 mg/kg/day (equivalent to the human exposure at the MRHD based on AUC$_{0-24}$h) caused an increase in post-implantation loss. At the high dose of 28 mg/kg/day (approximately 15 times the human exposure at the MRHD based on AUC$_{0-24}$h), reduced fetal and placental weights were observed, and evidence of teratogenicity (ectrodactylyia observed in 2 of 148 fetuses from 2 of 18 litters) was demonstrated.

In the pre- and postnatal study, pregnant rats received oral administration of ivabradine at doses of 2.5, 7, or 20 mg/kg/day from gestation day 6 to lactation day 20. Increased postnatal mortality associated with cardiac teratogenic findings was observed in the F1 pups delivered by dams treated at the high dose (approximately 15 times the human exposure at the MRHD based on AUC$_{0-24}$h).

### 8.2 Lactation

Risk Summary
There is no information regarding the presence of ivabradine in human milk, the effects of ivabradine on the breastfed infant, or the effects of the drug on milk production. Animal studies have shown, however, that ivabradine is present in rat milk [see Data]. Because of the potential risk to breastfed infants from exposure to Corlanor, breastfeeding is not recommended.
Lactating rats received daily oral doses of $[^{14}\text{C}]-\text{ivabradine} \ (7 \text{ mg/kg})$ on post-parturition days 10 to 14; milk and maternal plasma were collected at 0.5 and 2.5 hours post-dose on day 14. The ratios of total radioactivity associated with $[^{14}\text{C}]-\text{ivabradine}$ or its metabolites in milk vs. plasma were 1.5 and 1.8, respectively, indicating that ivabradine is transferred to milk after oral administration.

### 8.3 Females and Males of Reproductive Potential

#### Contraception

**Females**

Corlanor may cause fetal harm, based on animal data. Advise females of reproductive potential to use effective contraception during Corlanor treatment [see Use in Specific Populations (8.1)].

### 8.4 Pediatric Use

Safety and effectiveness in pediatric patients have not been established.

### 8.5 Geriatric Use

No pharmacokinetic differences have been observed in elderly (≥ 65 years) or very elderly (≥ 75 years) patients compared to the overall population. However, Corlanor has only been studied in a limited number of patients ≥ 75 years of age.

### 8.6 Hepatic Impairment

No dose adjustment is required in patients with mild or moderate hepatic impairment. Corlanor is contraindicated in patients with severe hepatic impairment (Child-Pugh C) as it has not been studied in this population and an increase in systemic exposure is anticipated [see Contraindications (4) and Clinical Pharmacology (12.3)].

### 8.7 Renal Impairment

No dosage adjustment is required for patients with creatinine clearance 15 to 60 mL/min. No data are available for patients with creatinine clearance below 15 mL/min [see Clinical Pharmacology (12.3)].

### 10. OVERDOSAGE

Overdose may lead to severe and prolonged bradycardia. In the event of bradycardia with poor hemodynamic tolerance, temporary cardiac pacing may be required. Supportive treatment, including intravenous (IV) fluids, atropine, and intravenous beta-stimulating agents such as isoproterenol, may be considered.
11. DESCRIPTION

Corlanor (ivabradine) is a hyperpolarization-activated cyclic nucleotide-gated channel blocker that reduces the spontaneous pacemaker activity of the cardiac sinus node by selectively inhibiting the I_{f}-current (I_f), resulting in heart rate reduction with no effect on ventricular repolarization and no effects on myocardial contractility.

The chemical name for ivabradine is 3-(3-\{(((7S)-3,4-Dimethoxybicyclo[4.2.0]octa-1,3,5-trien-7-yl)methyl] methyl amino\} propyl)-1,3,4,5-tetrahydro-7,8-dimethoxy-2H-3-benzazepin-2-one, hydrochloride. The molecular formula is C_{27}H_{36}N_{2}O_{5}, HCl, and the molecular weight (free base + HCl) is 505.1 (468.6 + 36.5). The chemical structure of ivabradine is shown in Figure 1.

Figure 1. Chemical Structure of Ivabradine

Corlanor tablets are formulated as salmon-colored, film-coated tablets for oral administration in strengths of 5 mg and 7.5 mg of ivabradine as the free base equivalent.

Inactive Ingredients

Core
Lactose monohydrate, maize starch, maltodextrin, magnesium stearate, colloidal silicon dioxide

Film Coating
Hypermellose, titanium dioxide, glycerol, magnesium stearate, polyethylene glycol 6000, yellow iron oxide, red iron oxide

12. CLINICAL PHARMACOLOGY

12.1 Mechanism of Action

Corlanor blocks the hyperpolarization-activated cyclic nucleotide-gated (HCN) channel responsible for the cardiac pacemaker I_{f}-current, which regulates heart rate. In clinical electrophysiology studies, the cardiac effects were most pronounced in the sinoatrial (SA) node, but prolongation of the AH interval has
occurred on the surface ECG, as has PR interval prolongation. There was no effect on ventricular repolarization and no effects on myocardial contractility [see Clinical Pharmacology (12.2)].

Corlanor can also inhibit the retinal current $I_h$. $I_h$ is involved in curtailing retinal responses to bright light stimuli. Under triggering circumstances (e.g., rapid changes in luminosity), partial inhibition of $I_h$ by Corlanor may underlie the luminous phenomena experienced by patients. Luminous phenomena (phosphenes) are described as a transient enhanced brightness in a limited area of the visual field [see Adverse Reactions (6.1)].

12.2 Pharmacodynamics

Corlanor causes a dose-dependent reduction in heart rate. The size of the effect is dependent on the baseline heart rate (i.e., greater heart rate reduction occurs in subjects with higher baseline heart rate). At recommended doses, heart rate reduction is approximately 10 bpm at rest and during exercise. Analysis of heart rate reduction vs. dose indicates a plateau effect at doses $\geq 20$ mg twice daily. In a study of subjects with preexisting conduction system disease (first- or second-degree AV block or left or right bundle branch block) requiring electrophysiologic study, IV ivabradine (0.20 mg/kg) administration slowed the overall heart rate by approximately 15 bpm, increased the PR interval (29 msec), and increased the AH interval (27 msec).

Corlanor does not have negative inotropic effects. Ivabradine increases the uncorrected QT interval with heart rate slowing but does not cause rate-corrected prolongation of QT.

12.3 Pharmacokinetics

Absorption and Bioavailability

Following oral administration, peak plasma ivabradine concentrations are reached in approximately 1 hour under fasting conditions. The absolute oral bioavailability of ivabradine is approximately 40% because of first-pass elimination in the gut and liver.

Food delays absorption by approximately 1 hour and increases plasma exposure by 20% to 40%. Corlanor should be taken with meals [see Dosage and Administration (2)].

Ivabradine is approximately 70% plasma protein bound, and the volume of distribution at steady state is approximately 100 L.
**Metabolism and Excretion**

The pharmacokinetics of ivabradine are linear over an oral dose range of 0.5 mg to 24 mg. Ivabradine is extensively metabolized in the liver and intestines by CYP3A4-mediated oxidation. The major metabolite is the N-desmethylated derivative (S 18982), which is equipotent to ivabradine and circulates at concentrations approximately 40% that of ivabradine. The N-desmethylated derivative is also metabolized by CYP3A4. Ivabradine plasma levels decline with a distribution half-life of 2 hours and an effective half-life of approximately 6 hours.

The total clearance of ivabradine is 24 L/h, and renal clearance is approximately 4.2 L/h, with ~ 4% of an oral dose excreted unchanged in urine. The excretion of metabolites occurs to a similar extent via feces and urine.

**Drug Interactions**

The effects of coadministered drugs (CYP3A4 inhibitors, substrates, inducers, and other concomitantly administered drugs) on the pharmacokinetics of Corlanor were studied in several single- and multiple-dose studies. Pharmacokinetic measures indicating the magnitude of these interactions are presented in Figure 2.
Digoxin exposure did not change when concomitantly administered with ivabradine. No dose adjustment is required when ivabradine is concomitantly administered with digoxin.

**Effect of Ivabradine on Metformin Pharmacokinetics**

Ivabradine, dosed at 10 mg twice daily to steady state, did not affect the pharmacokinetics of metformin (an organic cation transporter [OCT2] sensitive substrate). The geometric mean (90% confidence interval [CI]) ratios of $C_{\text{max}}$ and $\text{AUC}_{\text{inf}}$ of metformin, with and without ivabradine were 0.98 [0.83–1.15] and 1.02 [0.86–1.22], respectively. No dose adjustment is required for metformin when administered with Corlanor.

**Specific Populations**

**Age**

No pharmacokinetic differences (AUC or $C_{\text{max}}$) have been observed between elderly ($\geq 65$ years) or very elderly ($\geq 75$ years) patients and the overall patient population [see Use in Specific Populations (8.5)].
Hepatic Impairment
In patients with mild (Child-Pugh A) and moderate (Child-Pugh B) hepatic impairment, the pharmacokinetics of Corlanor were similar to that in patients with normal hepatic function. No data are available in patients with severe hepatic impairment (Child-Pugh C) [see Contraindications (4)].

Renal Impairment
Renal impairment (creatinine clearance from 15 to 60 mL/min) has minimal effect on the pharmacokinetics of Corlanor. No data are available for patients with creatinine clearance below 15 mL/min.

Pediatrics
The pharmacokinetics of Corlanor have not been investigated in patients < 18 years of age.

13. NONCLINICAL TOXICOLOGY
13.1 Carcinogenesis, Mutagenesis, Impairment of Fertility
There was no evidence of carcinogenicity when mice and rats received ivabradine up to 104 weeks by dietary administration. High doses in these studies were associated with mean ivabradine exposures of at least 37 times higher than the human exposure (AUC_{0-24hr}) at the MRHD.

Ivabradine tested negative in the following assays: bacterial reverse mutation (Ames) assay, in vivo bone marrow micronucleus assay in both mouse and rat, in vivo chromosomal aberration assay in rats, and in vivo unscheduled DNA synthesis assay in rats. Results of the in vitro chromosomal aberration assay were equivocal at concentrations approximately 1,500 times the human C_{max} at the MRHD. Ivabradine tested positive in the mouse lymphoma assays and in vitro unscheduled DNA synthesis assay in rat hepatocytes at concentrations greater than 1,500 times the human C_{max} at the MRHD.

Reproduction toxicity studies in animals demonstrated that ivabradine did not affect fertility in male or female rats at exposures 46 to 133 times the human exposure (AUC_{0-24hr}) at the MRHD.

13.2 Animal Toxicology and/or Pharmacology
Reversible changes in retinal function were observed in dogs administered oral ivabradine at total doses of 2, 7, or 24 mg/kg/day (approximately 0.6 to 50 times the human exposure at the MRHD based on AUC_{0-24hr}) for 52 weeks. Retinal function assessed by electroretinography demonstrated reductions in cone system responses, which reversed within a week post-dosing, and were not associated with damage to ocular structures as evaluated by light microscopy. These data are consistent with the pharmacological
effect of ivabradine related to its interaction with hyperpolarization-activated \( I_h \) currents in the retina, which share homology with the cardiac pacemaker \( I_f \) current.

14. CLINICAL STUDIES

SHIFT
The Systolic Heart failure treatment with the \( I_f \) inhibitor ivabradine Trial (SHIFT) was a randomized, double-blind trial comparing Corlanor and placebo in 6558 adult patients with stable NYHA class II to IV heart failure, left ventricular ejection fraction \( \leq 35\% \), and resting heart rate \( \geq 70 \) bpm. Patients had to have been clinically stable for at least 4 weeks on an optimized and stable clinical regimen, which included maximally tolerated doses of beta-blockers and, in most cases, ACE inhibitors or ARBs, spironolactone, and diuretics, with fluid retention and symptoms of congestion minimized. Patients had to have been hospitalized for heart failure within 12 months prior to study entry.

The underlying cause of CHF was coronary artery disease in 68\% of patients. At baseline, approximately 49\% of randomized subjects were NYHA class II, 50\% were NYHA class III, and 2\% were NYHA class IV. The mean left ventricular ejection fraction was 29\%. All subjects were initiated on Corlanor 5 mg (or matching placebo) twice daily and the dose was increased to 7.5 mg twice daily or decreased to 2.5 mg twice daily to maintain the resting heart rate between 50 and 60 bpm, as tolerated. The primary endpoint was a composite of the first occurrence of either hospitalization for worsening heart failure or cardiovascular death.

Most patients (89\%) were taking beta-blockers, with 26\% on guideline-defined target daily doses. The main reasons for not receiving the target beta-blocker doses at baseline were hypotension (45\% of patients not at target), fatigue (32\%), dyspnea (14\%), dizziness (12\%), history of cardiac decompensation (9\%), and bradycardia (6\%). For the 11\% of patients not receiving any beta-blocker at baseline, the main reasons were chronic obstructive pulmonary disease, hypotension, and asthma. Most patients were also taking ACE inhibitors and/or angiotensin II antagonists (91\%), diuretics (83\%), and anti-aldosterone agents (60\%). Few patients had an implantable cardioverter-defibrillator (ICD) (3.2\%) or a cardiac resynchronization therapy (CRT) device (1.1\%). Median follow-up was 22.9 months. At 1 month, 63\%, 26\%, and 8\% of Corlanor-treated patients were taking 7.5, 5, and 2.5 mg BID, whereas 3\% had withdrawn from the drug, primarily for bradycardia.

SHIFT demonstrated that Corlanor reduced the risk of the combined endpoint of hospitalization for worsening heart failure or cardiovascular death based on a time-to-event analysis (hazard ratio: 0.82, 95\% confidence interval [CI]: 0.75, 0.90, \( p < 0.0001 \)) (Table 3). The treatment effect reflected only a reduction

Reference ID: 3732763
in the risk of hospitalization for worsening heart failure; there was no favorable effect on the mortality component of the primary endpoint. In the overall treatment population, Corlanor had no statistically significant benefit on cardiovascular death.

**Table 3. SHIFT – Incidence of the Primary Composite Endpoint and Components**

<table>
<thead>
<tr>
<th>Endpoint</th>
<th>Corlanor (N = 3241)</th>
<th>Placebo (N = 3264)</th>
<th>Hazard Ratio [95% CI]</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>n</td>
<td>%</td>
<td>% PY</td>
<td>n</td>
</tr>
<tr>
<td>Primary composite endpoint: Time to first hospitalization for worsening heart failure or cardiovascular death$^a$</td>
<td>793</td>
<td>24.5</td>
<td>14.5</td>
<td>937</td>
</tr>
<tr>
<td>Hospitalization for worsening heart failure</td>
<td>505</td>
<td>15.6</td>
<td>9.2</td>
<td>660</td>
</tr>
<tr>
<td>Cardiovascular death as first event</td>
<td>288</td>
<td>8.9</td>
<td>4.8</td>
<td>277</td>
</tr>
<tr>
<td>Subjects with events at any time</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hospitalization for worsening heart failure$^b$</td>
<td>514</td>
<td>15.9</td>
<td>9.4</td>
<td>672</td>
</tr>
<tr>
<td>Cardiovascular death$^b$</td>
<td>449</td>
<td>13.9</td>
<td>7.5</td>
<td>491</td>
</tr>
</tbody>
</table>

$^a$ Subjects who died on the same calendar day as their first hospitalization for worsening heart failure are counted under cardiovascular death.

$^b$ Analyses of the components of the primary composite endpoint were not prospectively planned to be adjusted for multiplicity.

N: number of patients at risk; n: number of patients having experienced the endpoint; %: incidence rate = (n/N) x 100; % PY: annual incidence rate = (n/number of patient-years) x 100; CI: confidence interval

The Kaplan-Meier curve (Figure 3) shows time to first occurrence of the primary composite endpoint of hospitalization for worsening heart failure or cardiovascular death in the overall study.
A wide range of demographic characteristics, baseline disease characteristics, and baseline concomitant medications were examined for their influence on outcomes. Many of these results are shown in Figure 4. Such analyses must be interpreted cautiously, as differences can reflect the play of chance among a large number of analyses.

Most of the results show effects consistent with the overall study result. Corlanor's benefit on the primary endpoint in SHIFT appeared to decrease as the dose of beta-blockers increased, with little if any benefit demonstrated in patients taking guideline-defined target doses of beta-blockers.
Figure 4. Effect of Treatment on Primary Composite Endpoint in Subgroups

<table>
<thead>
<tr>
<th>% of Total Population</th>
<th>Conforan n(%)</th>
<th>Placebo n(%)</th>
<th>HR (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&gt;65 years</td>
<td>31.9%</td>
<td>13.1%</td>
<td>0.85 (0.77-1.03)</td>
</tr>
<tr>
<td>≥75 years</td>
<td>11.1%</td>
<td>15.5%</td>
<td>0.75 (0.65-0.88)</td>
</tr>
<tr>
<td>Age quartile</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>18 to ≤35</td>
<td>30.6%</td>
<td>14.5%</td>
<td>0.74 (0.64-0.87)</td>
</tr>
<tr>
<td>&gt;35 to ≤60</td>
<td>23.9%</td>
<td>17.8%</td>
<td>0.80 (0.71-0.90)</td>
</tr>
<tr>
<td>&gt;60 to ≤69</td>
<td>22.1%</td>
<td>22.2%</td>
<td>0.90 (0.79-1.02)</td>
</tr>
<tr>
<td>&gt;69</td>
<td>25.1%</td>
<td>20.5%</td>
<td>0.84 (0.76-0.94)</td>
</tr>
<tr>
<td>Sex</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>76.4%</td>
<td>66.8%</td>
<td>0.84 (0.76-0.93)</td>
</tr>
<tr>
<td>Female</td>
<td>23.6%</td>
<td>33.2%</td>
<td>0.74 (0.65-0.85)</td>
</tr>
<tr>
<td>Race</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Caucasian</td>
<td>89.7%</td>
<td>83.2%</td>
<td>0.75 (0.69-0.82)</td>
</tr>
<tr>
<td>Black</td>
<td>10.3%</td>
<td>16.8%</td>
<td>0.79 (0.67-0.93)</td>
</tr>
<tr>
<td>Other/unknown</td>
<td>0.9%</td>
<td>0.0%</td>
<td>0.03 (0.00-0.53)</td>
</tr>
<tr>
<td>Cause of heart failure</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Non-ischemic</td>
<td>78.1%</td>
<td>73.2%</td>
<td>0.73 (0.66-0.80)</td>
</tr>
<tr>
<td>Ischemic</td>
<td>21.9%</td>
<td>26.8%</td>
<td>0.73 (0.66-0.80)</td>
</tr>
<tr>
<td>Weight quartile</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>≤59.4</td>
<td>28.0%</td>
<td>27.6%</td>
<td>0.93 (0.85-1.02)</td>
</tr>
<tr>
<td>&gt;59.4 to ≤79.6</td>
<td>25.6%</td>
<td>26.0%</td>
<td>0.92 (0.85-1.01)</td>
</tr>
<tr>
<td>&gt;79.6 to ≤91</td>
<td>23.9%</td>
<td>23.3%</td>
<td>0.91 (0.84-0.99)</td>
</tr>
<tr>
<td>&gt;91</td>
<td>22.5%</td>
<td>21.1%</td>
<td>0.89 (0.81-0.98)</td>
</tr>
<tr>
<td>Baseline heart rate</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;75</td>
<td>25.3%</td>
<td>24.8%</td>
<td>0.89 (0.81-0.98)</td>
</tr>
<tr>
<td>≥75 to &lt;77</td>
<td>23.5%</td>
<td>23.9%</td>
<td>0.92 (0.84-1.01)</td>
</tr>
<tr>
<td>≥77 to &lt;84</td>
<td>22.5%</td>
<td>22.4%</td>
<td>0.89 (0.81-0.98)</td>
</tr>
<tr>
<td>≥84</td>
<td>24.3%</td>
<td>25.9%</td>
<td>0.86 (0.79-0.94)</td>
</tr>
<tr>
<td>NYHA Class</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>NYHA Class I</td>
<td>40.7%</td>
<td>39.7%</td>
<td>0.94 (0.87-1.02)</td>
</tr>
<tr>
<td>NYHA Class II</td>
<td>30.2%</td>
<td>30.2%</td>
<td>0.95 (0.89-1.01)</td>
</tr>
<tr>
<td>NYHA Class III</td>
<td>29.0%</td>
<td>30.0%</td>
<td>0.95 (0.89-1.01)</td>
</tr>
<tr>
<td>Baseline LV ejection</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>fraction quartile</td>
<td>≤25</td>
<td>25.6%</td>
<td>0.75 (0.65-0.86)</td>
</tr>
<tr>
<td>&gt;25 to ≤30</td>
<td>25.6%</td>
<td>25.6%</td>
<td>0.75 (0.65-0.86)</td>
</tr>
<tr>
<td>&gt;30 to ≤33</td>
<td>30.8%</td>
<td>30.4%</td>
<td>0.74 (0.65-0.84)</td>
</tr>
<tr>
<td>&gt;33</td>
<td>28.2%</td>
<td>27.6%</td>
<td>0.79 (0.69-0.90)</td>
</tr>
<tr>
<td>Baseline beta blocker</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>use quartile</td>
<td>≤25</td>
<td>25.4%</td>
<td>0.82 (0.70-0.95)</td>
</tr>
<tr>
<td>&gt;25 to ≤10</td>
<td>25.4%</td>
<td>25.4%</td>
<td>0.82 (0.70-0.95)</td>
</tr>
<tr>
<td>&gt;10 to ≤30</td>
<td>24.8%</td>
<td>24.8%</td>
<td>0.82 (0.70-0.95)</td>
</tr>
<tr>
<td>&gt;30</td>
<td>25.2%</td>
<td>25.2%</td>
<td>0.82 (0.70-0.95)</td>
</tr>
<tr>
<td>Baseline aldosterone</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>antagonist use quartile</td>
<td>≤25</td>
<td>25.4%</td>
<td>0.81 (0.70-0.95)</td>
</tr>
<tr>
<td>&gt;25 to ≤10</td>
<td>25.4%</td>
<td>25.4%</td>
<td>0.81 (0.70-0.95)</td>
</tr>
<tr>
<td>&gt;10 to ≤30</td>
<td>24.8%</td>
<td>24.8%</td>
<td>0.81 (0.70-0.95)</td>
</tr>
<tr>
<td>&gt;30</td>
<td>25.2%</td>
<td>25.2%</td>
<td>0.81 (0.70-0.95)</td>
</tr>
<tr>
<td>Diabetes</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>No history of diabetes</td>
<td>95.0%</td>
<td>95.0%</td>
<td>1.00 (0.89-1.12)</td>
</tr>
<tr>
<td>History of diabetes</td>
<td>5.0%</td>
<td>5.0%</td>
<td>1.00 (0.89-1.12)</td>
</tr>
<tr>
<td>Hypertension</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>No history of hypertension</td>
<td>92.9%</td>
<td>92.9%</td>
<td>1.00 (0.89-1.12)</td>
</tr>
<tr>
<td>History of hypertension</td>
<td>7.1%</td>
<td>7.1%</td>
<td>1.00 (0.89-1.12)</td>
</tr>
<tr>
<td>All</td>
<td>100.0%</td>
<td>99.0%</td>
<td>1.00 (0.89-1.12)</td>
</tr>
</tbody>
</table>

Note: The figure above presents effects in various subgroups, all of which are baseline characteristics. The 95% confidence limits that are shown do not take into account the number of comparisons made, and may not reflect the effect of a particular factor after adjustment for all other factors. Apparent homogeneity or heterogeneity among groups should not be over-interpreted.

BEAUTIFUL and SIGNIFY: No benefit in stable coronary artery disease with or without stable heart failure

BEAUTIFUL was a randomized, double-blind, placebo-controlled trial in 10,917 adult patients with coronary artery disease, impaired left ventricular systolic function (ejection fraction < 40%) and resting heart rate ≥ 60 bpm. Patients had stable symptoms of heart failure and/or angina for at least 3 months, and were receiving conventional cardiovascular medications at stable doses for at least 1 month. Beta-blocker therapy was not required, nor was there a protocol mandate to achieve any specific dosing targets for
patients who were taking beta-blockers. Patients were randomized 1:1 to Corlanor or placebo at an initial
dose of 5 mg twice daily with the dose increased to 7.5 mg twice daily depending on resting heart rate and
tolerability. The primary endpoint was the composite of time to first cardiovascular death, hospitalization
for acute myocardial infarction, or hospitalization for new-onset or worsening heart failure. Most patients
were NYHA class II (61.4%) or class III (23.2%) - none were class IV. Through a median follow-up of
19 months, Corlanor did not significantly affect the primary composite endpoint (HR 1.00, 95% CI =
0.91, 1.10).

SIGNIFY was a randomized, double-blind trial administering Corlanor or placebo to 19,102 adult patients
with stable coronary artery disease but without clinically evident heart failure (NYHA class I). Beta-
blocker therapy was not required. Corlanor was initiated at a dose of 7.5 mg twice daily and the dose
could be increased to as high as 10 mg twice daily or down-titrated to 5.0 mg twice daily to achieve a
target heart rate of 55 to 60 bpm. The primary endpoint was a composite of the first occurrence of either
cardiovascular death or myocardial infarction. Through a median follow-up of 24.1 months, Corlanor did
not significantly affect the primary composite endpoint (HR 1.08, 95% CI = 0.96, 1.20).

16. HOW SUPPLIED/STORAGE AND HANDLING
Corlanor 5 mg tablets are formulated as salmon-colored, oval-shaped, film-coated tablets scored on both
dges, marked with “5” on one face and bisected on the other face. They are supplied as follows:
• Bottles of 60 tablets (NDC 55513-800-60)
• Bottles of 180 tablets (NDC 55513-800-80)

Corlanor 7.5 mg tablets are formulated as salmon-colored, triangular-shaped, film-coated tablets debossed
with “7.5” on one face and plain on the other face. They are supplied as follows:
• Bottles of 60 tablets (NDC 55513-810-60)
• Bottles of 180 tablets (NDC 55513-810-80)

Storage
Store at 25°C (77°F); excursions permitted to 15° - 30°C (59° - 86°F) [see USP Controlled Room
Temperature].

17. PATIENT COUNSELING INFORMATION
Advise the patient to read the FDA-approved patient labeling (Medication Guide).
• Fetal Toxicity
  Advise pregnant women of the potential risks to a fetus.
Advise females of reproductive potential to use effective contraception and to notify their healthcare provider with a known or suspected pregnancy [see Warnings and Precautions (5.1) and Use in Specific Populations (8.1, 8.3)].

- **Low Heart Rate**
  Advise patients to report significant decreases in heart rate or symptoms such as dizziness, fatigue, or hypotension [see Warnings and Precautions (5.3)].

- **Atrial fibrillation**
  Advise patients to report symptoms of atrial fibrillation, such as heart palpitations or racing, chest pressure, or worsened shortness of breath [see Warnings and Precautions (5.2)].

- **Phosphenes**
  Advise patients about the possible occurrence of luminous phenomena (phosphenes). Advise patients to use caution if they are driving or using machines in situations where sudden changes in light intensity may occur, especially when driving at night. Advise patients that phosphenes may subside spontaneously during continued treatment with Corlanor [see Adverse Reactions (6.1)].

- **Drug Interactions**
  Advise patients to avoid ingestion of grapefruit juice and St. John’s wort [see Drug Interactions (7.1)].

- **Intake with Food**
  Advise patients to take Corlanor twice daily with meals [see Dosage and Administration (2)].
Read this Medication Guide before you start taking Corlanor and each time you get a refill. There may be new information. This information does not take the place of talking with your doctor about your medical condition or your treatment.

What is the most important information I should know about Corlanor?

Corlanor may cause serious side effects, including:

- **Harm to your unborn baby.** You should not become pregnant while taking Corlanor. Women who are able to get pregnant must use birth control when taking Corlanor. If you become pregnant while taking Corlanor, tell your doctor right away.

- **Increased risk of irregular heartbeat (atrial fibrillation or heart rhythm problems).** Tell your doctor if you have symptoms of an irregular heartbeat, such as feeling that your heart is pounding or racing (palpitations), chest pressure, or worsened shortness of breath.

- **Low heart rate (bradycardia).** Tell your doctor if you have a slowing of your heart rate or if you have symptoms of a low heart rate such as dizziness, fatigue, lack of energy, or have low blood pressure. Low heart rate is a common side effect of Corlanor and can be serious.

What is Corlanor?

Corlanor is a prescription medicine that is used to reduce the risk of hospitalization for worsening heart failure in people who have chronic heart failure.

- It is not known if Corlanor is safe and effective in children.

Who should not take Corlanor?

Do not take Corlanor if:

- You have symptoms of heart failure that recently worsened

- You have low blood pressure (less than 90/50 mmHg)

- You have a certain heart condition called sick sinus syndrome, sinoatrial block, or 3rd degree atrioventricular block

- You have a slow resting heart rate (less than 60 beats per minute) before treatment with Corlanor

- You are taking medicines that can change how much Corlanor gets into your body after it is swallowed. Your doctor can advise you if you are taking a medicine that should not be used with Corlanor.

What should I tell my doctor before taking Corlanor?

Reference ID: 3732763
Before you take Corlanor, tell your doctor about all of your medical conditions, including if you:

- Have any other heart problems, including heart rhythm problems, a slow heart rate, or a heart conduction problem.

- Are breastfeeding or plan to breastfeed. It is not known if Corlanor passes into your breast milk. You and your doctor should decide if you will take Corlanor or breastfeed. You should not do both.

Tell your doctor about all the medicines you take, including prescription and over-the-counter medicines, vitamins, and herbal supplements. Corlanor may affect the way other medicines work, and other medicines may affect how Corlanor works, and could cause serious side effects.

How should I take Corlanor?

- Take Corlanor exactly as your doctor tells you to take it.

- Your doctor may change your dose of Corlanor during treatment.

- If you take too much Corlanor, call your doctor or go to the nearest emergency room right away.

- If you forget to take a dose of Corlanor, take the next dose at the usual time. Do not take a double dose to make up for the forgotten dose.

What should I avoid while taking Corlanor?

Avoid drinking grapefruit juice and taking St. John’s wort during treatment with Corlanor. These can affect the way Corlanor works and may cause serious side effects.

What are the possible side effects of Corlanor?

See “What is the most important information I should know about Corlanor?”

The most common side effects of Corlanor are:

- Increased blood pressure.

- Temporary brightness in your field of vision, usually caused by sudden changes in light (luminous phenomena). This brightness usually happens within the first 2 months of treatment with Corlanor and usually goes away during or after treatment with Corlanor. Use caution when driving or operating machinery where sudden changes in light can happen, especially when driving at night.

These are not all the side effects of Corlanor. Ask your doctor or pharmacist for more information.

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

How should I store Corlanor?

- Store Corlanor at room temperature between 68°F to 77°F (20°C to 25°C).

- Safely throw away medicine that is out of date or no longer needed.

Keep Corlanor and all medicines out of the reach of children.
General information about the safe and effective use of Corlanor
Medicines are sometimes prescribed for purposes other than those listed in a Medication Guide.
Do not use Corlanor for a condition for which it was not prescribed. Do not give Corlanor to other people, even if they have the same symptoms that you have. It may harm them. You can ask your doctor or pharmacist for information about Corlanor that is written for health professionals.

What are the ingredients in Corlanor?
Active ingredient: ivabradine
Inactive ingredients:
Core: Lactose monohydrate, maize starch, maltodextrin, magnesium stearate, colloidal silicon dioxide
Film Coating: Hypromellose, titanium dioxide, glycerol, magnesium stearate, polyethylene glycol 6000, yellow iron oxide, red iron oxide

Corlanor® (ivabradine)
Manufactured for:
Amgen Inc.
One Amgen Center Drive
Thousand Oaks, California 91320-1799

Patent: http://pat.amgen.com/Corlanor/

For more information, go to www.Corlanor.com or call 1-800-772-6436.
Medication Guide has been approved by the U.S. Food and Drug Administration.
Issued: April/2015

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v1
Corlanor® (ivabradine) tablets

Each tablet contains 5 mg ivabradine equivalent to 5.39 mg ivabradine as hydrochloride.
Oral Use.
Store at 25°C, excursions permitted to 15°C - 30°C.
Keep out of the sight and reach of children.

Rx Only

14 Film-coated tablets

Reference ID: 3732763
Corlanor (ivabradine) tablets

Each tablet contains 5 mg ivabradine equivalent to 5.39 mg ivabradine as hydrochloride.

Oral Use.
Store at 25°C, excursions permitted to 15°C - 30°C.
Keep out of the sight and reach of children.

Rx Only

180 Film-coated tablets

NDC 55513-800-80

Reference ID: 3732763
Corlanor®
(ivabradine) tablets

Each tablet contains 7.5 mg ivabradine equivalent to 8.085 mg ivabradine as hydrochloride.

Oral Use.
Store at 25°C, excursions permitted to 15°C - 30°C.
Keep out of the sight and reach of children.

Rx Only
14 Film-coated tablets

Reference ID: 3732763
Corlanor® (ivabradine) tablets

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

Manufactured for Amgen Inc.
One Amgen Center Drive
Thousand Oaks, CA 91320-1799

Rx Only

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Manufactured for Amgen Inc.
One Amgen Center Drive
Thousand Oaks, CA 91320-1799

7.5 mg

Rx Only

Reference ID: 3732763
This is a representation of an electronic record that was signed electronically and this page is the manifestation of the electronic signature.

/s/

ELLIS F UNGER
04/15/2015