

# CORLANOR® ORAL SOLUTION PRODUCT FACT SHEET



## INDICATIONS



### Heart Failure in Adult Patients

Corlanor® is indicated to reduce the risk of hospitalization for worsening heart failure in adult 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.

### Heart Failure in Pediatric Patients

Corlanor® is indicated for the treatment of stable symptomatic heart failure due to dilated cardiomyopathy (DCM) in pediatric patients aged 6 months and older, who are in sinus rhythm with an elevated heart rate.

## ORAL SOLUTION PRODUCT INFORMATION

NDC	Description	Quantity
55513-813-28 	Corlanor® 5 mg/5 mL (1 mg/mL) oral solution is a colorless liquid in an opaque, low density polyethylene (LDPE) plastic ampule containing 5 mg of Corlanor® in 5 mL of liquid	Box of 28 ampules individually packaged in a child-resistant foil pouch 

## ORAL SOLUTION PRODUCT DISTRIBUTION

For out-patient use, Corlanor® oral solution is distributed exclusively by Optum Specialty Pharmacy, 855-312-9074.

For VA/DOD, 340B, or hospital in-patient use, please refer to Amgen's USA Authorized Wholesaler listing on Amgen's website, <https://www.amgen.com/partners/wholesalers/>

## STORAGE AND HANDLING REQUIREMENTS

Store at 25°C (77°F); excursions permitted to 15° - 30°C (59° - 86°F)  
Protect Corlanor® oral solution ampule from light by storing in the foil pouch until use.

## PRODUCT EXPIRATION

Expiration date is printed on the carton for oral solution.

## SUPPLIED AND MARKETED BY

Amgen USA Inc.  
Phone 1-800-28-AMGEN (1-800-282-6436) • Fax 1-800-29-AMGEN (1-800-292-6436)  
[www.amgen.com](http://www.amgen.com) [www.corlanorhcp.com](http://www.corlanorhcp.com)

## PRODUCT RETURNS

For information and instructions regarding product returns, please contact Amgen Trade Operations at 1-800-28-Amgen (1-800-282-6436). Credit for returns is subject to Amgen's current Returned Goods Policy.

## PRODUCT INFORMATION

Medical Information: 1-800-77-AMGEN (1-800-772-6436)

## COVERAGE INFORMATION

1-844-6CORLANOR (1-844-626-7526)

## IMPORTANT SAFETY INFORMATION

- **Contraindications:** Corlanor® is contraindicated in patients with acute decompensated heart failure, clinically significant hypotension, sick sinus syndrome, sinoatrial block, 3rd degree atrioventricular block (unless a functioning demand pacemaker is present), clinically significant bradycardia, severe hepatic impairment, pacemaker dependence (heart rate maintained exclusively by the pacemaker), and concomitant use of strong cytochrome P450 3A4 (CYP3A4) inhibitors.

*Please see additional Important Safety Information on next page.*

[www.corlanorhcp.com](http://www.corlanorhcp.com)

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- **Fetal Toxicity:** Corlanor<sup>®</sup> may cause fetal toxicity when administered to a pregnant woman based on embryo-fetal toxicity and cardiac teratogenic effects observed in animal studies. Advise females of reproductive potential to use effective contraception when taking Corlanor<sup>®</sup>.
- **Atrial Fibrillation:** Corlanor<sup>®</sup> increases the risk of atrial fibrillation. The rate of atrial fibrillation in patients treated with Corlanor<sup>®</sup> compared to placebo was 5% vs. 3.9% per patient-year, respectively. Regularly monitor cardiac rhythm. Discontinue Corlanor<sup>®</sup> if atrial fibrillation develops.
- **Bradycardia and Conduction Disturbances:**

### Adult Heart Failure Patients

Bradycardia, sinus arrest and heart block have occurred with Corlanor<sup>®</sup>. The rate of bradycardia in patients treated with Corlanor<sup>®</sup> compared to placebo was 6% (2.7% symptomatic; 3.4% asymptomatic) vs. 1.3% per patient-year, respectively. Risk factors for bradycardia include sinus node dysfunction, conduction defects, ventricular dyssynchrony, and use of other negative chronotropes. Bradycardia may increase the risk of QT prolongation which may lead to severe ventricular arrhythmias, including torsades de pointes, especially in patients with risk factors such as use of QTc prolonging drugs.

Concurrent use of verapamil or diltiazem also increases Corlanor<sup>®</sup> exposure, contributes to heart rate lowering, and should be avoided. Avoid use of Corlanor<sup>®</sup> in patients with 2nd degree atrioventricular block unless a functioning demand pacemaker is present.

### Pediatric Heart Failure Patients

Bradycardia and first-degree heart block were observed in pediatric patients treated with Corlanor<sup>®</sup>. Asymptomatic and symptomatic bradycardia were observed in 6.8% and 4.1% of pediatric patients treated with Corlanor<sup>®</sup>, respectively. In the placebo treatment arm, 2.4% of pediatric patients had asymptomatic bradycardia, but none had symptomatic bradycardia. Bradycardia was managed through dose titration but did not result in study drug discontinuation.

- **Adverse Reactions:**

### Adult Heart Failure Patients

The most common adverse drug reactions reported at least 1% more frequently with Corlanor<sup>®</sup> than placebo and that occurred in more than 1% of patients treated with Corlanor<sup>®</sup> were bradycardia (10% vs. 2.2%), hypertension or increased blood pressure (8.9% vs. 7.8%), atrial fibrillation (8.3% vs. 6.6%), and luminous phenomena (phosphenes) or visual brightness (2.8% vs. 0.5%).

In postmarketing experience, torsades de pointes has been observed.

### Pediatric Heart Failure Patients

Bradycardia (symptomatic and asymptomatic) occurred at rates similar to those in adults. Phosphenes were observed in pediatric patients treated with Corlanor<sup>®</sup>.

***Please see full Prescribing Information and Medication Guide***