

## Request for Prior Authorization IVABRADINE (CORLANOR®)

FAX Completed Form To 1 (800) 574-2515 Provider Help Desk 1 (877) 776-1567

(PLEASE PRINT – ACCURACY IS IMPORTANT)

IA Medicaid Member ID #	Patient name		DOB
Patient address			
Patient address			
Provider NPI	Prescriber name		Phone
Prescriber address			Fax
Pharmacy name	Address		Phone
-	ation above. It must be legible, correc	<del>-</del>	rm will be returned.
Pharmacy NPI	Pharmacy fax	NDC	
c) Patient is in sinus rhythm of the distribution of the distribu	of a left ventricular ejection fractivith a resting heart rate of ≥70 beart of blood pressure ≥90/50 mmHg; e symptomatic heart failure (NYHA) as and less than 18 years old; and if a left ventricular ejection fraction that a resting heart rate (HR) defined 05 bpm opm obpm obpm; and with maximally tolerated doses of a trial (e.g., carvedilol 50mg daily, ate dosing for pediatric patients, carblockers; and trial and continued use with a present of the property	ats per minute; and or A/Ross class II to I's and the second of the sec	olocker with proven mortality ate 200mg daily, or bisoprolo cumented intolerance or FDA system blocker at a
Non-Preferred			
Corlanor   Ivabradine			
Strength Dosage	Instructions	Quantity	Days Supply
<b>Diagnosis:</b> ☐ Stable, symptomatic heart fail	ure (NYHA Class II to IV)): NYHA	. Class (≥ 18 years	s of age):
☐ Stable, symptomatic heart fail 18 years of age):NYHA/Ross Cla			omyopathy (6 months to <
Other:			

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Provide left ventricular ejection fraction:	Date obtained:			
Provide resting heart rate in which patient is in sinus rhythm:				
Resting heart rate:	Date obtained:			
For diagnosis of stable, symptomatic heart failure (NYHA Class II, III, or IV) in members ≥ 18 years of age:				
Does patient have blood pressure ≥90/50mmHg?				
☐ No ☐ Yes: Blood pressure:	Date obtained:			
Treatment failure with maximally tolerated dose of beta-blocker with proven mortality benefit in a heart failure clinical trial:				
Drug name & dose:	Trial dates:			
Reason for failure:				
Contraindication:				
Trial and continued use with a preferred angiotensin system blocker at maximally tolerated dose:				
Drug name & dose:	Trial dates:			
Will an angiotensin system blocker be used concomitantly with ivabradine?   No Yes				
Attach lab results and other documentation as necessary.				
Prescriber signature (Must match prescriber listed above	ve.) Date of submission			

**IMPORTANT NOTE:** In evaluating requests for prior authorization the consultant will consider the treatment from the standpoint of medical necessity only. If approval of this request is granted, this does not indicate that the member continues to be eligible for Medicaid. It is the responsibility of the provider who initiates the request for prior authorization to establish by inspection of the member's Medicaid eligibility card and, if necessary by contact with the county Department of Health and Human Services, that the member continues to be eligible for Medicaid.

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