

# Pharmacy Prior Authorization Form

Fax completed form to: 877.974.4411 toll free, or 616.942.8206

This form applies to:  Commercial (Traditional)  Commercial (Individual/Optimized)  
 Medicaid

This request is:  Urgent (life threatening)  Non-Urgent (standard review)

Urgent means the standard review time may seriously jeopardize the life or health of the patient or the patient's ability to regain maximum function.

## Entresto<sup>®</sup> (sacubitril/valsartan)

### Member

Last Name: \_\_\_\_\_ First Name: \_\_\_\_\_  
 ID #: \_\_\_\_\_ DOB: \_\_\_\_\_ Gender: \_\_\_\_\_  
 Primary Care Physician: \_\_\_\_\_  
 Requesting Provider: \_\_\_\_\_ Prov. Phone: \_\_\_\_\_ Prov. Fax: \_\_\_\_\_  
 Provider Address: \_\_\_\_\_  
 Provider NPI: \_\_\_\_\_ Contact Name: \_\_\_\_\_  
 Provider Signature: \_\_\_\_\_ Date: \_\_\_\_\_

### Product Information

New request  Continuation request

Drug product:  Entresto 24-26 mg tablet  Entresto 49-51 mg tablet  Entresto 97-103 mg tablet  
 Start date (or date of next dose): \_\_\_\_\_  
 Date of last dose (if applicable): \_\_\_\_\_  
 Dosing frequency: \_\_\_\_\_

### Precertification Requirements

Before this drug is covered, the patient must meet all of the following requirements:

1. Must have chronic New York Heart Association class II to IV heart failure
2. Must have left ventricular ejection fraction less than or equal to 35%
3. Patient is tolerating an ACEI or ARB at HIGH doses (equivalent to at least enalapril 10mg BID) for at least 4 weeks (Entresto will replace the ACEI and/or ARB, after 36 hour washout)
4. Patient is currently on spironolactone or other diuretic for at least 4 weeks AND a beta blocker at a MAXIMAL tolerated dose for at least 4 weeks (or provide clinical reasoning as to why a maximal dosed beta blocker is inappropriate) and has not achieved improvement functional class or has worsening symptoms.
5. Systolic BP  $\geq$  100mmHg
6. eGFR  $\geq$  30mL/min/1.73m<sup>2</sup>
7. No history of angioedema or severe hepatic impairment (Child Pugh Class C)

For continuation, patient must have met the following requirements:

1. Must be adherent to all heart failure medications

**Note:** Authorization for indications, dosing, or a route of administration not approved by the Food and Drug Administration (FDA) or recognized in CMS-accepted compendia (e.g. DrugDex, AHFS, U.S. Pharmacopeia, and also Clinical Pharmacology for oncology indications only) require supporting evidence for coverage. Please provide two published peer-reviewed literature articles supporting the appropriateness of the drug, the dosing of the drug, or the route of administration to be used for the identified indication.

**New request**  
**Priority Health Precertification Documentation**

**A. What condition is this drug being requested for?**

- Chronic heart failure  
 Other – the patient's condition is: \_\_\_\_\_  
 Rationale for use: \_\_\_\_\_

**B. What NYHA functional classification of heart failure does this patient have?**

- I    II    III    IV

**C. Does the patient have a reduced ejection fraction?**

- Yes, the ejection fraction is \_\_\_\_\_%.  
 No – rationale for use: \_\_\_\_\_

**D. Is the patient currently tolerating an ACEI or ARB at HIGH doses (equivalent to at least enalapril 10mg BID) for at least 4 weeks (Entresto will replace the ACEI and/or ARB, after 36 hour washout)?**

- Yes  
 No – rationale for use: \_\_\_\_\_

**E. Is the patient currently on spironolactone or other diuretic for at least 4 weeks without improvement in functional class or with worsening symptoms?**

- Yes  
 No – rationale for use: \_\_\_\_\_

**F. Is the patient currently on a beta blocker at a MAXIMAL dose for at least 4 weeks without improvement in functional class or with worsening symptoms?**

- Yes  
 No. *What is the clinical reason why a maximally dosed beta blocker is not appropriate?* \_\_\_\_\_  
 \_\_\_\_\_

**G. Systolic BP  $\geq$  100mgHg?**

- Yes  
 No – rationale for use: \_\_\_\_\_

**H. Does the patient have an eGFR  $\geq$  30mL/min/1.73m<sup>2</sup>?**

- Yes  
 No – rationale for use: \_\_\_\_\_

**I. History of angioedema?**

- No  
 Yes – rationale for use: \_\_\_\_\_

**J. Severe hepatic impairment (Child Pugh Class C)?**

- No  
 Yes – rationale for use: \_\_\_\_\_

**Continuation request**  
**Priority Health Precertification Documentation**

**A. Has the patient been adherent to all heart failure medications (e.g., beta blocker, diuretic, Entresto)?**

- Yes  
 No – rationale for use: \_\_\_\_\_

**Additional information**

**Note:** When approved, the quantity of Entresto is limited to 60 tablets every 30 days.

**Comparison of ACCF/AHA Stages of HF and NYHA Functional Classifications.<sup>9</sup>**

ACCF/AHA HF Stage	ACCF/AHA Definition	NYHA Functional Class	NYHA Functional Class Definition
A	At high risk of HF but without structural heart disease or symptoms of HF.	None	---
B	Structural heart disease but without signs or symptoms of HF.	I	No limitation of physical activity. Ordinary physical activity does not cause symptoms of HF.
C	Structural heart disease with prior or current symptoms of HF.	I	No limitation of physical activity. Ordinary physical activity does not cause symptoms of HF.
	---	II	Slight limitation of physical activity. Comfortable at rest, but ordinary physical activity results in symptoms of HF.
	---	III	Marked limitation of physical activity. Comfortable at rest, but less than ordinary activity causes symptoms of HF.
	---	IV	Unable to carry on any physical activity without symptoms of HF, or symptoms of HF at rest.
D	Refractory HF requiring specialized interventions.	IV	Unable to carry on any physical activity without symptoms of HF, or symptoms of HF at rest.

ACCF – American College of Cardiology Foundation; AHA – American Heart Association; HF – Heart failure; NYHA – New York Heart Association; --- – Not applicable.