

## Pharmacy Prior Approval Request for Entresto

### Beneficiary Information

1. Beneficiary Last Name: \_\_\_\_\_ 2. First Name: \_\_\_\_\_  
3. Beneficiary ID #: \_\_\_\_\_ 4. Beneficiary Date of Birth: \_\_\_\_\_ 5. Beneficiary Gender: \_\_\_\_\_

### Prescriber Information

6. Prescribing Provider NPI #: \_\_\_\_\_  
7. Requester Contact Information - Name: \_\_\_\_\_ Phone #: \_\_\_\_\_ Ext. \_\_\_\_\_

### Drug Information

8. Drug Name: \_\_\_\_\_ 9. Strength: \_\_\_\_\_ 10. Quantity Per 30 Days: \_\_\_\_\_  
11. Length of Therapy (in days):  up to 30 Days  60 Days  90 Days  120 Days  180 Days  365 Days

### Clinical Information

1. Does the beneficiary have a diagnosis of chronic heart failure (NYHA class II-IV) with a left ventricular ejection fraction (EF) less than or equal to 40%?  Yes  No List ejection fraction: \_\_\_\_\_
2. Does the beneficiary have a history of angioedema related to therapy with an ACE inhibitor or ARB?  Yes  No
- 3a. Is the beneficiary currently taking an ACE inhibitor or ARB?  Yes  No
- 3b. If the beneficiary is currently taking an ACE inhibitor or ARB, will Entresto replace that current therapy?  
 Yes  No
- 4a. Does the beneficiary have diabetes?  Yes  No
- 4b. If the beneficiary has diabetes, is the beneficiary taking a medication containing aliskiren (e.g. Tekturna or Tekturna HCT)?  Yes  No

#### For reauthorization, please answer questions 1-5

5. Is documentation attached to this request that indicates the beneficiary is receiving clinical benefit from Entresto such as stabilization of symptoms, improvement?  Yes  No

Signature of Prescriber: \_\_\_\_\_ Date: \_\_\_\_\_

**(Prescriber Signature Mandatory)**

I certify that the information provided is accurate and complete to the best of my knowledge, and I understand that any falsification, omission, or concealment of material fact may subject me to civil or criminal liability.