



Entresto® Prior Authorization Request Form (Page 1 of 2)
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Member Information (required)			Provider Information (required)		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:	Specialty:	
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:
Medication Information (required)					
Medication Name:			Strength:	Dosage Form:	
<input type="checkbox"/> Check if requesting brand			Directions for Use:		
<input type="checkbox"/> Check if request is for continuation of therapy					
Clinical Information (required)					
Select the diagnosis below:					
<input type="checkbox"/> Heart failure (with or without hypertension)					
<input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____					
Continuation of therapy:					
Is the requested medication a continuation of therapy initiated during an inpatient stay? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Prescriber's specialty:					
Was the requested medication prescribed by or in consultation with a cardiologist? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Clinical information:					
Does the patient have an ejection fraction that is less than or equal to 40 percent? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Is the patient's heart failure classified as one of the following: New York Heart Association Class II, III or IV? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Is the patient on a stabilized dose and receiving concomitant therapy with one of the following beta blockers: Bisoprolol, carvedilol, or metoprolol? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Does the patient have a contraindication or intolerance to beta-blocker therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Does the patient have a history of angioedema? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Will the patient discontinue any use of concomitant angiotensin-converting enzyme (ACE) inhibitors or angiotensin-II receptor blockers (ARB) before initiating treatment with Entresto? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Please note: ACE inhibitors must be discontinued at least 36 hours prior to initiation of Entresto					
Is the patient receiving concomitant aliskiren (Tekturna) therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Reauthorization:					
If this is a reauthorization request, answer the following:					
Has the dose of Entresto been titrated to a dose of 97mg/103mg twice daily, or to a maximum dose as tolerated by the patient? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Is there documentation of positive clinical response to therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Quantity limit requests:					
What is the quantity requested per DAY? _____					
What is the reason for exceeding the plan limitations?					
<input type="checkbox"/> Titration or loading dose purposes					
<input type="checkbox"/> Patient is on a dose-alternating schedule (e.g., one tablet in the morning and two tablets at night, one to two tablets at bedtime)					
<input type="checkbox"/> Requested strength/dose is not commercially available					
<input type="checkbox"/> Other: _____					



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Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

Please note: This request may be denied unless all required information is received.
For urgent or expedited requests please call 1-800-711-4555.
This form may be used for non-urgent requests and faxed to 1-800-527-0531.