

Please complete this **entire** form and fax it to: **866-940-7328**. If you have questions, please call **800-310-6826**.  
**This form may contain multiple pages. Please complete all pages to avoid a delay in our decision.**  
**Allow at least 24 hours for review.**

**Section A – Member Information**

First Name:	Last Name:	Member ID:
Address:		
City:	State:	ZIP Code:
Phone:	DOB:	Allergies:
Primary Insurance Information (if any):		
Is the requested medication: <input type="checkbox"/> New or <input type="checkbox"/> Continuation of Therapy? If continuation, list start date: _____		
Is this patient currently hospitalized? <input type="checkbox"/> Yes <input type="checkbox"/> No If recently discharged, list discharge date: _____		

**Section B - Provider Information**

First Name:	Last Name:	M.D./D.O.
Address:	City:	State: ZIP code:
NPI #:	Phone:	Fax: Specialty:
Office Contact Name / Fax attention to:		

**Section C - Medical Information**

Medication:	Strength:
Directions for use:	Quantity:
Diagnosis (Please be specific & provide as much information as possible):	ICD-10 CODE:
Is this member pregnant? <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, what is this member's due date? _____	

**Section D – Previous Medication Trials**

Medication Name	Strength	Directions	Dates of Therapy	Reason for failure / discontinuation

**Section E – Additional information and Explanation of why preferred medications would not meet the patient's needs:  
Please refer to the patient's PDL for a list of preferred alternatives**

<b>Member First name:</b>	<b>Member Last name:</b>	<b>Member DOB:</b>
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**Clinical and Drug Specific Information**

**ALL REQUESTS:**

- **What is the patient's diagnosis? (check which applies)**

- Post-Herpetic Neuralgia
- Neuropathic Pain
- Other. **List diagnosis:** \_\_\_\_\_

- **Does the patient have a history of failure, contraindication, or intolerance to ALL of the following:**

- Tricyclic anti-depressant (e.g., amitriptyline)**
- SNRI anti-depressant (e.g., duloxetine, venlafaxine)**
- Gabapentin**

(If yes, complete Section D above with medication information, including dose, date of trial, and reason for discontinuation)

**Requests for CONTINUATION OF THERAPY:**

- **Does the patient have a documented positive clinical response to lidocaine patch therapy?**  **Yes**  **No**

**If yes, list response:** \_\_\_\_\_

**Provider Signature:** \_\_\_\_\_ **Date:** \_\_\_\_\_

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