



# REQUEST FOR WAIVER OF BRAND PENALTY

## Patient Information

Patient Name \_\_\_\_\_ Patient ID \_\_\_\_\_  
 Gender:  Male  Female Date of birth \_\_\_\_/\_\_\_\_/\_\_\_\_

## Prescriber Information

Prescriber Name \_\_\_\_\_ Prescriber Specialty \_\_\_\_\_  
 Address \_\_\_\_\_  
 City \_\_\_\_\_ State \_\_\_\_\_ ZIP \_\_\_\_\_  
 Phone Number (\_\_\_\_) \_\_\_\_\_ Fax Number (\_\_\_\_) \_\_\_\_\_  
 Contact Person at Prescriber's Office \_\_\_\_\_ DEA Number \_\_\_\_\_

## Diagnosis

Diagnosis \_\_\_\_\_ Disease Duration \_\_\_\_\_  
 Diagnosis (ICD) Code(s) \_\_\_\_\_

## Medication Information

Brand Name Medication \_\_\_\_\_  
 Strength \_\_\_\_\_ Frequency \_\_\_\_\_ Quantity \_\_\_\_\_  
 Expected Length of Therapy \_\_\_\_\_

## Form Cannot Be Evaluated Without Required Clinical Information

*Please complete form and fax back to (888) 836-0730*

1. Is "Dispense As Written" indicated by the provider on the prescription?  Yes  No
2. Has the patient tried an AB-rated generic equivalent to the brand prescribed?  Yes  No
3. Does the patient have a documented allergic reaction to an inactive ingredient that is present in the generic formulation, but absent in the brand name equivalent? **(must provide documentation)**  Yes  No
4. Has the patient had a documented life-threatening side effect that required medical intervention to a generic medication that did not occur with the brand name equivalent? **(must provide documentation)**  Yes  No
5. Has the prescriber completed and submitted an FDA MedWatch Adverse Event Reporting Form on behalf of this patient? **(must provide copy)**  Yes  No

I attest that the information provided is accurate and true, and that medical records accurately reflect the information provided.

Provider Signature \_\_\_\_\_ Date \_\_\_\_/\_\_\_\_/\_\_\_\_

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