



FLORIDA MEDICAID PRIOR AUTHORIZATION MULTI-SOURCE BRAND DRUG

Note: Form must be completed in full.
An incomplete form may be returned.

Request for Multi-Source Brand Drug Due to Adverse Effects or Ineffectiveness of Generic

Note to Prescribing Physician: THIS FORM MUST BE SUBMITTED ALONG WITH A MISCELLANEOUS PRIOR AUTHORIZATION FORM AND COPY OF THE PRESCRIPTION IF A REQUEST IS BEING MADE TO DISPENSE A BRAND PRODUCT DUE TO ADVERSE EFFECTS OR INEFFECTIVENESS OF A GENERIC.

It is very important that physician's prescribe generic drugs whenever possible. Most FDA-approved generics are bioequivalent and therapeutically equivalent to the brand name drug. This request form is only to be used if your patient has experienced an adverse medical reaction to the generic drug or if you can document that your patient has had better medical results when taking the multi-source brand drug, as opposed to its generic substitute.

Recipient's Medicaid ID#	Date of Birth (MM/DD/YYYY)
<input type="text"/>	<input type="text"/>

Recipient's Full Name
<input type="text"/>

Prescriber's Full Name
<input type="text"/>

Prescriber License # (ME, OS, ARNP, PA)
<input type="text"/>

Prescriber Phone Number	Prescriber Fax Number
<input type="text"/>	<input type="text"/>

GENERIC PRODUCT (Give labeled strength & mfr/labeler, if known)	REQUESTED BRAND PRODUCT (Give labeled strength & mfr/labeler, if known)
Name: _____ Manufacturer: _____ NDC#: _____ Strength: _____ Dose, Frequency, & Route Used: _____ Therapy Dates (if unknown, give duration) from/to (or best estimate): _____ Diagnosis for Use (Indication): _____	Name: _____ Manufacturer: _____ NDC#: _____ Strength: _____ Dose, Frequency, & Route Used: _____ Diagnosis for Use (Indication): _____

ADVERSE EVENT	BENEFITS OF BRAND PRODUCT
Describe event or problem with generic: (Must provide medical record documentation describing adverse event)	Describe how brand will alleviate problem: (Must provide medical record documentation describing adverse event)