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United Healthcare Community Plan

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) / /

Recipient's Full Name

Prescriber's Full Name

Prescriber's NPI

Prescriber Phone Number - -

Prescriber Fax Number - -

GENERIC PRODUCT (Give labeled strength & mfr/labeler, if known)	**REQUESTED BRAND PRODUCT** (Give labeled strength & mfr/labeler, if known)
Name:	**Name:**
Manufacturer:	**Manufacturer:**
NDC#:	**NDC#:**
Strength:	**Strength:**
Dose, Frequency, & Route Used:	**Dose, Frequency, & Route Used:**
Therapy Dates (if unknown, give duration) from/to (or best estimate):	**Diagnosis for Use (Indication):**
Diagnosis for Use (Indication):	
ADVERSE EVENT	**BENEFITS OF BRAND PRODUCT**
Describe event or problem with generic:	**Describe how brand will alleviate problem:**
(Must provide medical record documentation describing adverse event)	(Must provide medical record documentation describing adverse event)

Signature: _____ Date: _____

Fax this form to 1-866-940-7328

Pharmacy PA Call Center:
1-800-310-6826

02.01.2025