



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 TO AHCA WITH A MISCELLANEOUS PA 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.

PATIENT INFORMATION	PRESCRIBING PHYSICIAN
Full Name: _____ Medicaid ID #: _____ Date of Birth: _____ SSN: _____ Sex: <input type="checkbox"/> Female <input type="checkbox"/> Male Weight: _____ lb.(s)	Name: _____ Address: _____ Phone #: _____ Licence #: _____ Fax #: _____ Signature: _____
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)

REQUIRED FOR REVIEW: Copies of medical records (i.e. diagnostic evaluations and recent chart notes), the original prescription, and the most recent copies of related labs.

The provider must retain copies of all documentation for five years.