

Administrative Policy Statement ARKANSAS PASSE				
Policy Name		Policy Number	Date Effective	
Medical Necessity – Off Label		PAD-0899-AR-PASSE	01/01/2022	
Policy Type				
Medical	ADMINISTRATIVE	Pharmacy	Reimbursement	

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Table of Contents

Adn	ninistrative Policy Statement	. 1
<u>A.</u>	Subject.	. 2
<u>B.</u>	Background	. 2
<u>C.</u>	Definitions	. 2
<u>D.</u>	Policy	. 2
<u>E.</u>	Conditions of Coverage	. 3
<u>F.</u>	Related Policies/Rules	. 3
<u>G.</u>	Review/Revision History	3
<u>H.</u>	References	3



Medical Necessity – Off Label Arkansas PASSE PAD-0899-AR-PASSE Effective Date: 01/01/2022

B. Background

The U.S. Food and Drug Administration (FDA) approves drugs for specific indications included in the drug's product information label. Off-label or "unlabeled" drug use is the utilization of an FDA approved drug for uses other than those listed in the FDA approved labeling or in treatment regimens or populations that are not included in approved labeling. Many off-label uses are effective, well documented in the peer-reviewed literature, and widely used even though the manufacturer has not pursued the additional indications.

The FDA advises physician's use of off-label or "unlabeled" drugs must be done in a wellinformed manner in conjunction with firm scientific rationale and medical evidence. CareSource will employ, at its discretion, drug utilization management programs (i.e., prior authorization) to ensure appropriate and safe use of medications.

NOTE: The Introduction section is for your general knowledge and is not to be construed as policy coverage criteria. The rest of the policy uses specific words and concepts familiar to medical professionals and is intended for providers. A provider can be a person, such as a doctor, nurse, psychologist, or dentist. A provider can also be a place where medical care is given, like a hospital, clinic or lab. This policy informs providers about when a service may be covered.

C. Definitions

- **FDA Approved medication**: Is the official description of a drug product which includes indication; who should take it; adverse events; instructions for uses in pregnancy; children, and other populations; and safety information for the patient. Labels are often found inside drug product packaging.
- **Off-label or "unlabeled" drug use**: Is the use of a drug approved by the U.S. Food and Drug Administration (FDA) for other uses that are not included in approved labeling. The FDA approves drugs for specific indications that are included in the drug's labeling. When a drug is used for an indication other than those specifically included in the labeling, it is referred to as an off-label use. Many off-label uses are effective, well documented in the literature, and widely used.

D. Policy

CareSource will review prior authorization requests for coverage based on medical necessity. This policy will not supersede drug-specific criteria developed and approved by the CareSource Pharmacy and therapeutics Committee (P&T).

Requests for off-label uses of a drug will be considered for approval according to the following criteria:

- I. Documentation must be submitted showing the member has tried and failed the existing FDA approved and/or clinical guideline recommended therapies unless contraindicated or not tolerated; AND
- II. The prescribed use must be supported by <u>one or more</u> of the following:
 - a. Narrative information from American Hospital Formulary Service Drug Information (AHFS) or Clinical Pharmacology





Medical Necessity – Off Label Arkansas PASSE PAD-0899-AR-PASSE Effective Date: 01/01/2022

- b. Lexicomp: Evidence level A
- c. Micromedex: Recommendation class I, IIa, or IIb
- d. Evidence from <u>at least two</u> published studies from major scientific or medical peer reviewed journals demonstrates safety and efficacy for the specified condition in a comparable population (i.e. age group, level of disease severity, etc.) *If applicable clinical trial is yet to be published but interim results are supportive, this may be taken into consideration by the clinician reviewer.*

NOTE: For off-label use of <u>oncology</u> drugs, please refer to the policy titled "Oncology Treatment Regimen Review" accessible from the CareSource website.

E. Conditions of Coverage

AUTHORIZATION PERIOD

Approved authorizations are designated an appropriate authorization period. Continued treatment may be considered when the member has shown tolerability and a positive clinical response.

F. Related Policies/Rules

Oncology Treatment Regimen Review

G. Review/Revision History

	DATES	ACTION
Date Issued	10/29/2021	
Date Revised	10/29/2021	Policy created
Date Effective	01/01/2022	
Date Archived		

H. References

 Medical Benefit Policy Manual. CMS website. Updated August 7, 2020. Accessed February 12, 2021. <u>https://www.cms.gov/Regulations-and-</u> Guidance/Guidance/Manuals/Downloads/bp102c15.pdf

This guideline contains custom content that has been modified from the standard care guidelines and has not been reviewed or approved by MCG Health, LLC.

The Administrative Policy Statement detailed above has received due consideration as defined in the Administrative Policy Statement Policy and is approved.

