# SPECIALTY GUIDELINE MANAGEMENT

# **KEVEYIS** (dichlorphenamide)

### **POLICY**

#### I. INDICATIONS

The indications below including FDA-approved indications and compendial uses are considered a covered benefit provided that all the approval criteria are met and the member has no exclusions to the prescribed therapy.

## FDA-Approved Indication<sup>1</sup>

Treatment of primary hyperkalemic periodic paralysis, primary hypokalemic periodic paralysis, and related variants.

All other indications are considered experimental/investigational and are not a covered benefit.

### **II. CRITERIA FOR INITIAL APPROVAL**

## A. Primary Hypokalemic Periodic Paralysis<sup>1-8</sup>

Authorization of 60 days may be granted to members who are initiating Keveyis therapy when the following criteria is met:

- 1. The diagnosis was supported by at least one of the following:
  - a. Genetic test results or
  - b. Patient has a family history of primary hypokalemic periodic paralysis, or
  - c. Patient's attacks are associated with hypokalemia AND both Andersen-Tawil syndrome and thyrotoxic periodic paralysis have been ruled out.
- 2. Trial with suboptimal response to treatment with acetazolamide

## B. Primary Hyperkalemic Periodic Paralysis<sup>1-8</sup>

Authorization of 60 days may be granted to members who are initiating Keveyis therapy when the following criteria is met:

- 1. The diagnosis was supported by at least one of the following:
  - a. Genetic test results, or
  - b. Patient has a family history of primary hyperkalemic periodic paralysis, or
  - Patient's attacks are associated with hyperkalemia AND Andersen-Tawil syndrome has been ruled out.
- 2. Trial with suboptimal response to treatment with acetazolamide

## **III. CONTINUATION OF THERAPY**

Authorization of 12 months may be granted to members who have demonstrated a response to Keveyis therapy as demonstrated by an improvement in their condition (e.g. decrease in the number or severity of attacks).

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#### IV. REFERENCE

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