

# **Keveyis Quantity Limit Program Summary**

Quantity limits apply to Medicaid.

#### POLICY REVIEW CYCLE

Effective Date Date of Origin 7/1/2023 3/1/2018

### FDA APPROVED INDICATIONS AND DOSAGE

See package insert for FDA prescribing information: <a href="https://dailymed.nlm.nih.gov/dailymed/index.cfm">https://dailymed.nlm.nih.gov/dailymed/index.cfm</a>

## **CLINICAL RATIONALE**

#### **REFERENCES**

#### POLICY AGENT SUMMARY OUANTITY LIMIT

Target Brand Agent Name(s)		Strengt h	QL Amount	Dose Form	Day Supply		Addtl QL Info	Allowed Exceptions	Targete d NDCs When Exclusi ons Exist	
Keveyis	Dichlorphenamide Tab 50 MG	50 MG	120	TABS	30	DAYS				

#### CLIENT SUMMARY - QUANTITY LIMITS

Target Brand Agent Name(s)	Target Generic Agent Name(s)	Strength	Client Formulary	
Keveyis	Dichlorphenamide Tab 50 MG	50 MG	Medicaid	

# QUANTITY LIMIT CLINICAL CRITERIA FOR APPROVAL

Module	Clinical Criteria for Approval
QL Standalo	Evaluation
ne	Quantities above the program quantity limit for the <b>Target Agent(s)</b> will be approved when ONE of the following is met:
	<ol> <li>The requested quantity (dose) does NOT exceed the program quantity limit OR</li> <li>ONE of the following:         <ul> <li>BOTH of the following:</li> <li>The requested agent does NOT have a maximum FDA labeled dose for the requested indication AND</li> </ul> </li> </ol>
	<ol> <li>Information has been provided to support therapy with a higher dose for the requested indication OR</li> <li>BOTH of the following:         <ol> <li>The requested quantity (dose) does NOT exceed the maximum FDA labeled dose for the requested indication AND</li> </ol> </li> </ol>

Module	Clinical Criteria for Approval
	<ol> <li>Information has been provided to support why the requested quantity (dose) cannot be achieved with a lower quantity of a higher strength that does NOT exceed the program quantity limit OR</li> </ol>
	C. BOTH of the following:
	<ol> <li>The requested quantity (dose) is greater than the maximum FDA labeled dose for the requested indication AND</li> </ol>
	<ol><li>Information has been provided to support therapy with a higher dose for the requested indication</li></ol>
	Length of Approval: up to 12 months