

STANDARD COMMERCIAL DRUG FORMULARY PRIOR AUTHORIZATION GUIDELINES

TRAMADOL

Generic	Brand	HICL	GCN	Medi-Span	Exception/Other
TRAMADOL HCL	QDOLO, TRAMADOL HCL		48598	GPI-14 (65100095102005)	

GUIDELINES FOR USE

- 1. Is the request for the management of pain and the patient meets **ALL** of the following criteria?
 - The patient is 18 years of age or older
 - The patient's pain is severe enough to require an opioid analgesic and alternative treatments are inadequate
 - The patient had a trial of or contraindication to generic tramadol IR tablet or a generic tramadol with acetaminophen product
 - The patient is unable to take oral solid formulations of tramadol or tramadol with acetaminophen (e.g., difficulty swallowing)

If yes, approve for 6 months by GPID or GPI-14 with a quantity limit of #80mL per day. If no, do not approve.

DENIAL TEXT: *Some terms are already pre-defined in parenthesis. Please use these definitions if the particular text you need to use does not already have definition(s) in it.

Our guideline named **TRAMADOL** (**Qdolo**) requires the following rule(s) be met for approval:

- A. The request is for the management of pain
- B. You are 18 years of age or older
- C. Your pain is severe enough to require an opioid analgesic (type of pain medication) and alternative treatments are inadequate
- D. You had a trial of or contraindication (harmful for) to generic tramadol immediate-release (IR) tablet or a generic tramadol with acetaminophen product
- E. You are unable to take oral solid formulations of tramadol or tramadol with acetaminophen (such as with difficulty swallowing)

Your doctor told us [INSERT PT SPECIFIC INFO PROVIDED]. We do not have information showing you [INSERT UNMET CRITERIA]. This is why your request is denied. Please work with your doctor to use a different medication or get us more information if it will allow us to approve this request.

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RATIONALE

For further information, please refer to the Prescribing Information and/or Drug Monograph for Qdolo.

REFERENCES

Qdolo [Prescribing Information]. Athens, GA: Athena Bioscience, LLC; September 2021.

Library	Commercial	NSA
Yes	Yes	No

Part D Effective: N/A Created: 02/21

Commercial Effective: 03/14/22 Client Approval: 02/22 P&T Approval: 01/21

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