

PRIOR AUTHORIZATION CRITERIA

BRAND NAME*
(generic)

LI DODERM
(lidocaine patch 5%)

ZTLi DO
(lidocaine topical system)

Status: CVS Caremark Criteria

Type: Initial Prior Authorization with Quantity Limit

Ref # 125-C

* Drugs that are listed in the target drug box include both brand and generic and all dosage forms and strengths unless otherwise stated. OTC products are not included unless otherwise stated.

FDA-APPROVED INDICATIONS

Li doder m

Li doder m is indicated for relief of pain associated with post-herpetic neuralgia. It should be applied only to **intact skin**.

ZTLi do

ZTLi do (lidocaine topical system) 1.8% is indicated for the relief of pain associated with post-herpetic neuralgia (PHN).

Compended Uses

Pain associated with diabetic neuropathy^{4,5,8}

Pain associated with cancer-related neuropathy^{4,6,7}

COVERAGE CRITERIA

The requested drug will be covered with prior authorization when the following criteria are met:

- The requested drug is being prescribed for any of the following: A) Pain associated with post-herpetic neuralgia, B) Pain associated with diabetic neuropathy, C) Pain associated with cancer-related neuropathy (including treatment-related neuropathy [e.g. neuropathy associated with radiation treatment or chemotherapy])

Quantity limits apply.

RATIONALE

The intent of the criteria is to provide coverage consistent with product labeling, FDA guidance, standards of medical practice, evidence-based drug information, and/or published guidelines. Li doder m is indicated for relief of pain associated with post-herpetic neuralgia. ZTLi do (lidocaine topical system) 1.8% is indicated for the relief of pain associated with post-herpetic neuralgia (PHN).

Because of the difference in bioavailability of ZTLi do compared with Li doder m (lidocaine 5% patch), a different dosage strength is required to be administered to the patient. One ZTLi do (lidocaine topical system) 1.8% provides equivalent lidocaine exposure to one Li doder m (lidocaine patch 5%). In a single-dose, crossover study conducted in 53 healthy volunteers, ZTLi do (lidocaine topical system) 1.8% demonstrated equivalent exposure (AUC) and peak concentration (C_{max}) of lidocaine to Li doder m (lidocaine patch 5%).²

Li doder m ZTLi do 125-C 09-2019a

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Li docaine patches 5% were well tolerated and reduced pain in patients with diabetic neuropathy in an open-label, flexible dosing, 3-week study with a 5-week extension with 56 patients with clinically defined painful diabetic polyneuropathy of longer than a 3 months duration. Results determined that up to four 5% li docaine patches for up to 18 hours per day are well tolerated in patients with painful diabetic polyneuropathy, significantly improve pain and quality-of-life ratings, and may allow tapering of concomitant analgesic therapy.⁵ Additionally, the American Academy of Neurology recommends that the Lidoderm patch may be considered for the treatment of painful diabetic neuropathy.⁸ Since ZTLi do patches provide an equivalent dose of li docaine as the Lidoderm patches, coverage is available for Lidoderm (li docaine patches) 5% and ZTLi do (li docaine topical system) 1.8% for pain associated with diabetic neuropathy.

Neuropathic cancer pain (NCP) may be cancer-related, namely resulting from nervous system tumor invasion, surgical nerve damage during tumor removal, radiation-induced nerve damage and chemotherapy-related neuropathy, or may be of benign origin unrelated to cancer.⁷ Additional analgesic medications or therapies may be necessary in patients with cancer-related pain, particularly when opioid analgesics are ineffective or produce inadequate pain relief. According to National Comprehensive Cancer Network (NCCN) guidelines for adult cancer pain, topical agents such as Lidocaine can be used as an adjunctive treatment option in neuropathic pain.^{4,6}

For PHN, Lidoderm or ZTLi do should be applied to intact skin to cover the most painful area with application of up to three patches, only once for up to 12 hours within a 24-hour period. Patches may be cut into smaller sizes with scissors prior to removal of the release liner. Excessive dosing by applying more than the recommended quantity or applying for longer than the recommended wearing time could result in increased absorption of li docaine and high blood concentrations, leading to serious adverse effects.¹⁻⁴

REFERENCES

1. Lidoderm [package insert]. Chadds Ford, PA: Endo Pharmaceuticals Inc.; November 2018.
2. ZTLi do [package insert]. San Diego, CA: Solix Pharmaceuticals Inc.; November 2018.
3. Lexi-Comp Online, AHFS DI (Adult and Pediatric) Online. Hudson, OH: Wolters Kluwer Clinical Drug Information, Inc. <http://online.lexi.com>. Accessed August 2019.
4. Micromedex (electronic version). Truven Health Analytics, Greenwood Village, Colorado, USA. <http://www.micromedexsolutions.com>. Accessed September 2018.
5. Barbano RL, Herrmann DN, Hart-Goulet S, et al: Effectiveness, tolerability, and impact on quality of life of the 5% li docaine patch in diabetic polyneuropathy. *Arch Neurol* 2004; 61: 914-918.
6. National Comprehensive Cancer Network: Adult Cancer Pain V3. 2019. National Comprehensive Cancer Network. Fort Washington, PA 2008. Available from URL: http://www.nccn.org/professional/s/physi_dan_g/s/PDF/pain.pdf. Accessed August 2019.
7. Vadalouca A, Raptis E, Moka E, et al. Pharmacological Treatment of Neuropathic Cancer Pain: A Comprehensive Review of Current Literature. *World Institute of Pain Pain Practice* 2011; 12(3): 219-251.
8. Bril V, Engel J, Franklin GM, et al. Evidence-based guideline: Treatment of painful diabetic neuropathy. *Neurology* 2011; 76: 1758. Available at www.neurology.org. Accessed August 2019.
9. Derry S, Wiffen PJ, Moore RA, et al. Topical li docaine for neuropathic pain in adults (Review). *Cochrane Database Syst Rev* 2014. doi: 10.1002/14651858.CD010958.

Written by:	UM Development (JG)
Date Written:	09/2002
Revised:	03/2005; (NB) 02/2006, 02/2007, 01/2008, 01/2009; (CY) 12/2009, 02/2011; (SE) 10/2011 operational clarification; (CY) 02/2012; (RP) 01/2013, 01/2014, (JH) 01/2015, 01/2016 (no clinical changes); (SF) 01/2017, 09/2017 (no clinical changes), 09/2018 (added ZTLi do); (DFW) 09/2019 (no clinical changes)
Reviewed:	Medical Affairs (MM) 03/2005, 02/2006; (WF) 02/2007, 01/2008, 01/2009, 12/2009; (KP) 02/2011, 02/2012; (LS) 01/2013; (DNC) 01/2014; (KQ) 01/2015; (LMS) 04/2015; (CHART) 09/26/19 External Review: 05/2005, 06/2006, 04/2007, 04/2008, 04/2009, 5/2010, 06/2011, 06/2012, 06/2013, 06/2014, 04/2015, 04/2016, 04/2017, 02/2018, 02/2019, 02/2020

CRITERIA FOR APPROVAL

- | | | | |
|---|--|-----|----|
| 1 | Is the requested drug being prescribed for any of the following: A) Pain associated with post-herpetic neuralgia, B) Pain associated with diabetic neuropathy, C) Pain associated with cancer-related neuropathy (including treatment-related neuropathy [e.g. neuropathy associated with radiation treatment or chemotherapy])? | Yes | No |
| 2 | Does the patient require more than the plan allowance of 90 patches per month?
[Rph Note: If yes, then deny and enter a partial approval for 90 patches per 25 days or 270 patches per 75 days of lidoderm or ZTLi do.] | Yes | No |

Guidelines for Approval

Duration of Approval 36 Months

Quantity for Approval

90 patches/ 25 days* or 270 patches/ 75 days*

Set 1

Yes to question(s)

1

No to question(s)

2

*The duration of 25 days is used for a 30-day fill period and 75 days is used for a 90-day fill period to allow time for refill processing.

Mapping Instructions

	Yes	No	DENIAL REASONS – DO NOT USE FOR MED CARE PART D
1	Go to 2	Deny	Coverage for this medication is denied for the following reason(s). We reviewed the information we received about your condition and circumstances. We used the policy (INSERT CRITERIA NAME) when making this decision. The policy states that this medication may be approved when being used for any of the following: - Pain associated with post-herpetic neuralgia - Pain associated with diabetic neuropathy - Pain associated with cancer-related neuropathy (including cancer treatment-related neuropathy) Based on the policy and the information we have, the request is denied. The information provided to us indicates that the requested medication will not be used for one of the uses listed above. [Short Description: Diagnosis]
2	Deny	Approve, 36 months, 90 patches/ 25 days* 270 patches/ 75 days*	You do not meet the requirements of your plan. You have requested more than the maximum quantity allowed by your plan. Current plan approved criteria cover up to 90 patches/ month of the requested drug. You have been approved for the maximum quantity that your plan covers for a duration of 36 months. Your request for additional quantities of the requested drug and strength has been denied. [Short Description: Over max quantity]

*The duration of 25 days is used for a 30-day fill period and 75 days is used for a 90-day fill period to allow time for refill processing.