

Clinical Policy Title:	nitroglycerin
Policy Number:	RxA.156
Drug(s) Applied:	GoNitro™
Original Policy Date:	02/07/2020
Last Review Date:	09/14/2020
Line of Business Policy Applies to:	All lines of business

Background

Nitroglycerin (GoNitro™) is an organic nitrate that is a vasodilator which has effects on both arteries and veins. It is indicated for acute relief of an attack or prophylaxis of angina pectoris due to coronary artery disease.

Dosing Information

Drug Name	Indication	Dosing Regimen	Maximum Dose
Nitroglycerin (GoNitro™)	Angina	<p>At the onset of an attack, administer one or two packets (400 mcg each) under the tongue. One additional packet may be administered every 5 minutes as needed. No more than three total packets (1200 mcg) are recommended within a 15minute period.</p> <p>If chest pain persists after three packets, seek prompt medical attention.</p> <p>May be used prophylactically 5 to 10 minutes prior to engaging in activities that might precipitate an acute attack.</p>	1,200 mcg within 15 minutes

Dosage Forms

- Sublingual powder: 400 mcg/packet

Clinical Policy

Provider must submit documentation (such as office chart notes, lab results or other clinical information) supporting that member has met all approval criteria.

This clinical policy has been developed to authorize, modify, or determine coverage for individuals with similar conditions. Specific care and treatment may vary depending on individual need and benefits covered by the plan. This policy is not intended to dictate to providers how to practice medicine, nor does it constitute a contract or guarantee regarding payment or results. This document may contain prescription brand name drugs that are trademarks of pharmaceutical manufacturers that are not affiliated with RxAdvance.

I. Initial Approval Criteria

A. Angina (must meet all):

1. Diagnosis of coronary artery disease requiring angina prophylaxis;
2. Documentation supports inability to use generic sublingual nitroglycerin tablets (generic Nitrostat®).

Approval Duration

Commercial: 6 months

Medicaid: 6 months

II. Continued Therapy Approval

A. Angina (must meet all):

1. Currently receiving medication that has been authorized by Rxadvance or member has previously met initial approval criteria listed in this policy.

Approval Duration

Commercial: 6 months

Medicaid: 6 months

III. Appendices

APPENDIX A: Abbreviation/Acronym Key

FDA: Food and Drug Administration

PDE-5: Phosphodiesterase type 5

sGC: Soluble Guanylate Cyclase

APPENDIX B: Therapeutic Alternatives

This table provides a listing of preferred alternative therapy recommended in the approval criteria. The drugs listed here may not be a formulary agent for all relevant lines of business and may require prior authorization.

Drug Name	Dosing Regimen	Dose Limit/ Maximum Dose
nitroglycerin sublingual tablets (Nitrostat®)	0.3 to 0.6 mg every 5 minutes for a maximum of 3 tablets in 15 minutes; may also use prophylactically 5 to 10 minutes prior to activities which may provoke an attack.	1.8 mg within 15 minutes

Therapeutic alternatives are listed as Brand name® (generic) when the drug is available by brand name only and generic (Brand name®) when the drug is available by both brand and generic.

APPENDIX C: Contraindications/Boxed Warnings

- Contraindication(s):
 - Use of phosphodiesterase type 5 (PDE-5) inhibitors, such as avanafil, sildenafil, tadalafil, or vardenafil, or soluble guanylate cyclase (sGC) stimulators;
 - Severe anemia;
 - Increased intracranial pressure;
 - Hypersensitivity to GoNitro or to other nitrates or nitrates or any excipient;
 - Circulatory failure and shock.

- Boxed Warning(s):
 - None reported

APPENDIX D: General Information

Not Applicable

References

1. GoNitro Prescribing Information. Jacksonville, FL: Espero Pharmaceuticals, Inc.; June 2018. Available at: <https://dailymed.nlm.nih.gov/dailymed/>. Accessed July 8, 2020.
2. Micromedex® Healthcare Series [Internet database]. Greenwood Village, Colo: Thomson Healthcare. Updated periodically. Accessed July 8, 2020.

Review/Revision History	Review/Revised Date	P&T Approval Date
Policy established.	01/2020	02/07/2020
Policy was reviewed: 1. Clinical Policy Title was updated 2. Drug(s) Applied was updated 3. Line of Business Policy Applies to was updated 4. Continued Therapy criteria II.A.1 was rephrased to "Currently receiving medication that has been authorized by RxAdvance..." 5. Commercial approval duration and Medicaid approval duration updated. 6. Updated APPENDIX A: Abbreviation/Acronym Key to include PDE-5: Phosphodiesterase type 5 and sGC: Soluble Guanylate Cyclase 7. References were updated	07/08/2020	09/14/2020