

Clinical Policy: Tazarotene (Arazlo, Fabior, Tazorac)

Reference Number: DE.PMN.244

Effective Date: 01.23 Last Review Date: 01.23 Line of Business: Medicaid

Revision Log

See <u>Important Reminder</u> at the end of this policy for important regulatory and legal information.

Description

Tazarotene lotion (Arazlo[™]), foam (Fabior[®]), cream and gel (Tazorac[®]) are topical retinoids.

FDA Approved Indication(s)

Tazarotene is indicated for the topical treatment of:

- Plaque psoriasis (*Tazorac cream and gel 0.05% and 0.1%*)
- Acne vulgaris:
 - That is mild to moderate (*Tazorac cream and gel 0.1%*)
 - o In patients 9 years of age and older (*Arazlo lotion*)
 - o In patients 12 years of age or older (Fabior foam)

Limitation(s) of use: The safety of Tazorac gel use on more than 20% body surface area has not been established.

Policy/Criteria

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

It is the policy of health plans affiliated with Centene Corporation[®] that Arazlo, Fabior, and Tazorac are **medically necessary** when the following criteria are met:

I. Initial Approval Criteria

- A. Plaque Psoriasis (must meet all):
 - 1. Request is for tazarotene cream or gel;
 - 2. Diagnosis of plaque psoriasis with body surface area involvement of $\leq 20\%$;
 - 3. Request does not exceed 1 tube per month.

Approval duration: 12 months

B. Acne Vulgaris (must meet all):

- 1. Diagnosis of acne vulgaris;
- 2. For Arazlo and Fabior requests only, member meets all of the following (a, b, and c):
 - a. Member meets one of the following (i or ii):
 - i. For Arazlo: age ≥ 9 years;
 - ii. For Fabior: age ≥ 12 years;
 - b. Failure of generic formulary topical tretinoin, unless clinically significant adverse effects are experienced or contraindicated;
- 3. Request does not exceed 1 tube (Arazlo, Tazorac) or 1 can (Fabior) per month.

Approval duration: 12 months

C. Other diagnoses/indications

1. Refer to the off-label use policy for the relevant line of business if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized): CP.PMN.53 for Medicaid.

II. Continued Therapy

A. All Indications in Section I (must meet all):

- 1. Currently receiving medication via Centene benefit or member has previously met initial approval criteria;
- 2. Member is responding positively to therapy;
- 3. If request is for a dose increase, new request does not exceed 1 tube (Arazlo, Tazorac) or 1 can (Fabior) per month.

Approval duration: 12 months

B. Other diagnoses/indications (must meet 1 or 2):

1. Currently receiving medication via Centene benefit and documentation supports positive response to therapy.

Approval duration: Duration of request or 12 months (whichever is less); or

2. Refer to the off-label use policy for the relevant line of business if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized): CP.PMN.53 for Medicaid.

III. Diagnoses/Indications for which coverage is NOT authorized:

A. Non-FDA approved indications, which are not addressed in this policy, unless there is sufficient documentation of efficacy and safety according to the off label use policies – CP.PMN.53 for Medicaid, or evidence of coverage documents.

IV. Appendices/General Information

Appendix A: Abbreviation/Acronym Key FDA: Food and Drug Administration

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
tretinoin (Retin-A®)	Acne Vulgaris 0.025% gel, 0.05% cream, 0.1% cream: Apply once daily	Not applicable
adapalene (Differin®)	Acne Vulgaris Lotion, Cream: 0.1%; Gel: 0.1%, 0.3% Apply topically QD	Not applicable

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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):
 - Pregnancy
 - o Tazorac: Individuals who have known hypersensitivity to any of its components
- Boxed warning(s): none reported

V. Dosage and Administration

Drug Name	Indication	Dosing Regimen	Maximum Dose
Tazarotene	Plaque	Apply gel or cream, 0.05% with strength	2 mg/cm ² /day
(Tazorac)	psoriasis	increased to 0.1% if tolerated and	
cream and		medically indicated, qPM to psoriatic	
gel 0.05%		lesions, using enough (2 mg/cm ²) to cover	
and 0.1%		only the lesion with a thin film.	
		*Do not cover more than 20% of body	
		surface area with the gel formulation.	
Tazarotene	Acne	Apply a thin film (2 mg/cm ²) of gel or	2 mg/cm ² /day
(Tazorac)		cream 0.1% qPM, to the skin where acne	
cream and		lesions appear.	
gel 0.1%			
Tazarotene	Acne	Apply a thin layer to the affected areas	Once daily
(Arazlo)		once daily. Avoid the eyes, mouth,	application
lotion		paranasal creases and mucous membranes.	
0.045%		Not for oral, ophthalmic or intravaginal	
		use.	
Tazarotene	Acne	Apply a thin layer to the entire affected	Once daily
(Fabior)		areas of the face and/or upper trunk once	application
foam 0.1%		daily in the evening. Avoid the eyes, lips,	
		and mucous membranes.	

VI. Product Availability

Drug Name	Availability
Tazarotene (Tazorac)	Cream (30 g and 60 g tube): 0.05%, 0.1% (<i>generic available</i>)
	Gel (30 g and 100 g tube): 0.05%, 0.1%
Tazarotene (Arazlo)	Lotion (45 g tube): 0.045%
Tazarotene (Fabior)	Foam (50 g and 100 g can): 0.1%

VII. References

1. Arazlo Prescribing Information. Bridgewater, NJ: Bausch Health US LLC, May 2021. Available at: https://pi.bauschhealth.com/globalassets/BHC/PI/arazlo-pi.pdf. Accessed August 15, 2022. Fabior Prescribing Information. Greenville, NC: Mayne Pharmaceuticals, June 2018. Available at www.dailymed.nlm.nih.gov. Accessed August 15, 2022.

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- 2. Tazorac Cream Prescribing Information. Irvine, CA: Allergan, Inc., August 2019. Available at https://www.tazorachcp.com/static/tazorac-cream-pi-54115b39c03f775e241400f432307541.pdf. Accessed August 15, 2022.
- 3. Tazorac Gel Prescribing Information. Irvine, CA: Allergan, Inc., August 2019. Available at https://www.almirall.us/pdf/Tazorac_Gel_USPI_final_Aug2019.pdf . Accessed August 15, 2022.
- 4. Clinical Pharmacology. Tampa, FL: Gold Standard; 2021. Available at www.clinicalpharmacology.com. Accessed August 15, 2022.
- 5. Zaenglein AL, Pathy AL, Schlosser BJ et al. Guidelines of care for the management of acne vulgaris. J Am Acad Dermatol. 2016 May;74(5):945-73.e33. doi:10.1016/j.jaad.2015.12.037.

Reviews, Revisions, and Approvals	Date	P&T Approval Date
Policy created	01.23	01.23

Important Reminder

This clinical policy has been developed by appropriately experienced and licensed health care professionals based on a review and consideration of currently available generally accepted standards of medical practice; peer-reviewed medical literature; government agency/program approval status; evidence-based guidelines and positions of leading national health professional organizations; views of physicians practicing in relevant clinical areas affected by this clinical policy; and other available clinical information. The Health Plan makes no representations and accepts no liability with respect to the content of any external information used or relied upon in developing this clinical policy. This clinical policy is consistent with standards of medical practice current at the time that this clinical policy was approved. "Health Plan" means a health plan that has adopted this clinical policy and that is operated or administered, in whole or in part, by Centene Management Company, LLC, or any of such health plan's affiliates, as applicable.

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This clinical policy is effective as of the date determined by the Health Plan. The date of posting may not be the effective date of this clinical policy. This clinical policy may be subject to applicable legal and regulatory requirements relating to provider notification. If there is a discrepancy between the effective date of this clinical policy and any applicable legal or regulatory requirement, the requirements of law and regulation shall govern. The Health Plan retains the right to change, amend or withdraw this clinical policy, and additional clinical policies may be developed and adopted as needed, at any time.

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This clinical policy does not constitute medical advice, medical treatment or medical care. It is not intended to dictate to providers how to practice medicine. Providers are expected to exercise professional medical judgment in providing the most appropriate care, and are solely responsible for the medical advice and treatment of members. This clinical policy is not intended to recommend treatment for members. Members should consult with their treating physician in connection with diagnosis and treatment decisions.

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Note: For Medicaid members, when state Medicaid coverage provisions conflict with the coverage provisions in this clinical policy, state Medicaid coverage provisions take precedence. Please refer to the state Medicaid manual for any coverage provisions pertaining to this clinical policy.

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