Last Review Date: 10/03/2022 Date of Origin: 12/12/2017 Dates Reviewed: 12/2017, 03/2018, 06/2018, 10/2018, 11/2019, 01/2020, 10/2020, 03/2021, 08/2021, 10/2022

I. Length of Authorization

Coverage is provided for 6 months may be renewed.

II. Dosing Limits

A. Quantity Limit (max daily dose) [NDC Unit]:

- <u>Fasenra 30 mg single-dose prefilled syringe</u>
 - Load: 1 syringe every 28 days for 3 doses
 - Maintenance: 1 syringe every 56 days
- Fasenra Pen 30 mg single-dose autoinjector
 - \circ Load: 1 autoinjector every 28 days for 3 doses
 - Maintenance: 1 autoinjector every 56 days

B. Max Units (per dose and over time) [HCPCS Unit]:

- Load: 30 billable units every 28 days for 3 doses
- Maintenance: 30 billable units every 56 days

III. Initial Approval Criteria¹

Note: For Medicaid members, please refer to the <u>Medicaid specific criteria</u>. (Not applicable to self-administered formulation)

Coverage is provided in the following conditions:

• Patient is at least 12 years of age; AND

Universal Criteria¹

- Will not be used in combination with other anti-IgE, anti-IL4, or anti-IL5 monoclonal antibody (e.g., omalizumab, mepolizumab, reslizumab, dupilumab, etc.); **AND**
- Must NOT be used for either of the following:
 - Treatment of other eosinophilic conditions (e.g., allergic bronchopulmonary aspergillosis/mycosis, Churg-Strauss syndrome, hypereosinophilic syndrome, etc.)

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• Relief of acute bronchospasm or status asthmaticus; AND

Severe Asthma † 1,2,5,7,8,9,11

- Patient must have severe* disease; **AND**
- Patient must have asthma with an eosinophilic phenotype indicated by blood eosinophils $\geq 150 \text{ cells}/\mu L$ within 6 weeks of dosing; AND
- Must be used for add-on maintenance treatment in patients <u>regularly</u> receiving BOTH of the following:
 - Medium to high-dose inhaled corticosteroids; AND
 - An additional controller medication (e.g., long-acting beta agonist, leukotriene modifiers, etc.); **AND**
- Patient must have two or more exacerbations in the previous year requiring daily oral corticosteroids for at least 3 days (in addition to the regular maintenance therapy defined above); **AND**
- Baseline measurement of at least one of the following for assessment of clinical status:
 - Use of systemic corticosteroids
 - Use of inhaled corticosteroids
 - Number of hospitalizations, ER visits, or unscheduled visits to healthcare provider due to condition
 - \circ Forced expiratory volume in 1 second (FEV₁)

*Components of severity for classifying asthma as severe may include any of the following (not all inclusive):^{2,9}

- Symptoms throughout the day
- Nighttime awakenings, often 7x/week
- SABA use for symptom control occurs several times per day
- Extremely limited normal activities
- Lung function (percent predicted FEV₁) <60%
- Exacerbations requiring oral systemic corticosteroids are generally more frequent and intense relative to moderate asthma

FDA-approved indication(s); Compendia recommended indication(s); Orphan Drug

IV. Renewal Criteria ^{1,7,8}

- Patient continues to meet the universal and other indication-specific relevant criteria identified in section III; **AND**
- Absence of unacceptable toxicity from the drug. Examples of unacceptable toxicity include: parasitic (helminth) infection, severe hypersensitivity reactions (e.g., anaphylaxis, angioedema, urticaria, rash), etc.; **AND**
 - Improvement in asthma symptoms or asthma exacerbations as evidenced by a decrease in one or more of the following:
 - Use of systemic corticosteroids

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- Two-fold or greater decrease in inhaled corticosteroid use for at least 3 days
- Hospitalizations
- ER visits
- Unscheduled visits to healthcare provider; **OR**
- Improvement from baseline in forced expiratory volume in 1 second (FEV₁)

V. Dosage/Administration¹

| Indication | Dose |
|------------------------|---|
| Severe Asthma with | Administer 30 mg subcutaneously every 4 weeks for the first three doses |
| eosinophilic phenotype | and then once every 8 weeks thereafter. |

VI. Billing Code/Availability Information

HCPCS Code:

• J0517 – Injection, benralizumab, 1 mg: 1 billable unit = 1 mg

NDC:

- Fasenra 30 mg/mL single-dose prefilled syringe: 00310-1730-xx
- Fasenra 30 mg/mL single-dose autoinjector FASENRA PEN: 00310-1830-xx

VII. References

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- Global Initiative for Asthma (GINA). Global Strategy for Asthma Management and Prevention. 2019 Update. Available from: http://www.ginasthma.org. Accessed September 2020.
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- Goldman M, Hirsch I, Zangrilli JG, et al. The association between blood eosinophil count and benralizumab efficacy for patients with severe, uncontrolled asthma: subanalyses of the Phase III SIROCCO and CALIMA studies. Curr Med Res Opin. 2017 Sep;33(9):1605-1613. Doi: 10.1080/03007995.2017.1347091. Epub 2017 Jul 19.
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- 7. Chung KF, Wenzel SE, Brozek JL, et al. International ERS/ATS Guidelines on Definition, Evaluation, and Treatment of Severe Asthma. Eur Respir J 2014; 43: 343-373.
- Holguin F, Cardet JC, Chung KF, et al. Management of severe asthma: a European Respiratory Society/American Thoracic Society guideline. Eur Respir J 2020; 55: 1900588 [https://doi.org/10.1183/13993003.00588-2019].
- 9. National Asthma Education and Prevention Program (NAEPP). 2020 Focused Updates to the Asthma Management Guidelines: A Report from the National Asthma Education and Prevention Program Coordinating Committee Expert Panel Working Group. Bethesda, MD: National Institutes of Health (NIH), National Heart, Lung, and Blood Institute (NHLBI); December 2020.
- 10. Global Initiative for Asthma (GINA). Global Strategy for Asthma Management and Prevention. 2021 Update. Available from: <u>http://www.ginasthma.org</u>. Accessed June 2021.
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Appendix 1 – Covered Diagnosis Codes

| ICD-10 | ICD-10 Description | |
|--------|--|--|
| J45.50 | Severe persistent asthma, uncomplicated | |
| J82.81 | Eosinophilic pneumonia, NOS | |
| J82.82 | Acute eosinophilic pneumonia | |
| J82.83 | Eosinophilic asthma | |
| J82.89 | Other pulmonary eosinophilia, not elsewhere classified | |

Appendix 2 – Centers for Medicare and Medicaid Services (CMS)

Medicare coverage for outpatient (Part B) drugs is outlined in the Medicare Benefit Policy Manual (Pub. 100-2), Chapter 15, §50 Drugs and Biologicals. In addition, National Coverage Determination (NCD), Local Coverage Determinations (LCDs), and Local Coverage Articles (LCAs) may exist and compliance with these policies is required where applicable. They can be found at: http://www.cms.gov/medicare-coverage-database/search.aspx. Additional indications may be covered at the discretion of the health plan.

Medicare Part B Covered Diagnosis Codes (applicable to existing NCD/LCD/LCA): N/A

| Medicare Part B Administrative Contractor (MAC) Jurisdictions | | | | |
|---|--|---|--|--|
| Jurisdiction | Applicable State/US Territory | Contractor | | |
| E (1) | CA, HI, NV, AS, GU, CNMI | Noridian Healthcare Solutions, LLC | | |
| F (2 & 3) | AK, WA, OR, ID, ND, SD, MT, WY, UT, AZ | Noridian Healthcare Solutions, LLC | | |
| 5 | KS, NE, IA, MO | Wisconsin Physicians Service Insurance Corp (WPS) | | |

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| Medicare Part B Administrative Contractor (MAC) Jurisdictions | | | | |
|---|---|---|--|--|
| Jurisdiction | Applicable State/US Territory | Contractor | | |
| 6 | MN, WI, IL | National Government Services, Inc. (NGS) | | |
| H (4 & 7) | LA, AR, MS, TX, OK, CO, NM | Novitas Solutions, Inc. | | |
| 8 | MI, IN | Wisconsin Physicians Service Insurance Corp (WPS) | | |
| N (9) | FL, PR, VI | First Coast Service Options, Inc. | | |
| J (10) | TN, GA, AL | Palmetto GBA, LLC | | |
| M (11) | NC, SC, WV, VA (excluding below) | Palmetto GBA, LLC | | |
| L (12) | DE, MD, PA, NJ, DC (includes Arlington & Fairfax counties and the city of Alexandria in VA) | Novitas Solutions, Inc. | | |
| K (13 & 14) | NY, CT, MA, RI, VT, ME, NH | National Government Services, Inc. (NGS) | | |
| 15 | KY, OH | CGS Administrators, LLC | | |

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