

PA Criteria	Criteria Details						
Covered Uses (FDA approved indication)	Fasenra is an interleukin-5 (IL-5) antagonist indicated for severe eosinophilic asthma add-on therapy and for the treatment of adult patients with eosinophilic granulomatosis with polyangiitis (EGPA).						
Exclusion Criteria	Must not be used in combination with other biologic drugs.						
Required Medical Information	<p>For initial coverage of severe eosinophilic asthma:</p> <p>Medical records supporting the request, including documentation of prior therapies and responses to treatment must be provided.</p> <p>Must have an elevated eosinophil level greater than or equal to 150 cells/mcL within six weeks (prior to the immediate start of treatment with Fasenra) - OR - greater than or equal to 300 cells/mcL in the previous 12 months.</p> <p>Must try and fail one ICS/LABA inhaler drug in the past six months (fail is defined as an intolerance or inability to improve the condition on required therapy for at least four weeks).</p> <p>For initial coverage of eosinophilic granulomatosis with polyangiitis (EGPA): Medical records supporting the request must be provided and include documentation that the patient has non-severe EGPA (defined as absence of life or organ-threatening manifestations).</p> <p>For reauthorization requests for severe eosinophilic asthma: (1) Medical records supporting the request must be provided - (2) Must have documentation of clinical benefit (e.g., decrease in exacerbations, improvement in symptoms, decrease in oral steroid use).</p> <p>For reauthorization requests for EGPA: (1) Medical records supporting the request must be provided - (2) Must have documentation of clinical benefit (e.g., decrease in exacerbations, improvement in symptoms, decrease in oral steroid use).</p>						
Age Restriction	None.						
Prescriber Restrictions	Prescriber is a specialist or has consulted with a specialist for the condition being treated.						
Coverage Duration	Initial: One year; reauthorization: two years. Dose will be approved according to the FDA approved labeling or within accepted standards of medical practice.						
Other Criteria/Information	<p>Refer to the Gold Coast Health Plan Medicare Part B Reference and Summary of Evidence document.</p> <table border="1"> <thead> <tr> <th>HCPCS</th> <th>Description</th> <th>Billing Units/How Supplied</th> </tr> </thead> <tbody> <tr> <td>J0517</td> <td>Fasenra (benralizumab) prefilled syringe</td> <td>Billing unit: 1 mg 30 mg/mL SD syringe</td> </tr> </tbody> </table>	HCPCS	Description	Billing Units/How Supplied	J0517	Fasenra (benralizumab) prefilled syringe	Billing unit: 1 mg 30 mg/mL SD syringe
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STATUS	DATE REVISED	REVIEW DATE	APPROVED/REVIEWED BY	EFFECTIVE DATE
Created	3/26/2025	3/26/2025	Dawn Shojai, PharmD, Senior Pharmacy Benefit Consultant (PSG)	N/A
Approved	N/A	8/21/2025	Pharmacy & Therapeutics (P&T) Committee	8/21/2025