

MEDICARE FORM Alpha 1 – Antitrypsin Inhibitor Therapy **Medication Precertification Request**

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(All fields must be completed and legible for precertification review.)

Note: Aralast NP and Glassia are non-preferred. The preferred products are Prolastin-C and Please indicate: Start of treatment: Start date / / Zemaira. ☐ Continuation of therapy: Date of last treatment / / Precertification Requested By: _ Phone: __ Fax: ___ A. PATIENT INFORMATION First Name: Last Name: Address: Citv: State: ZIP: Home Phone: Work Phone: Cell Phone: DOB: Allergies: Email: Current Weight: Ibs or kgs Height: _____ inches or ____ cms B. INSURANCE INFORMATION Aetna Member ID #: Does patient have other coverage? ☐ Yes ☐ No If yes, provide ID#: _____ Carrier Name: ____ Group #: _____ Insured: Insured: **Medicare**: ☐ Yes ☐ No If yes, provide ID #: **Medicaid**: ☐ Yes ☐ No If yes, provide ID #: C. PRESCRIBER INFORMATION Last Name: First Name: (Check One): M.D. D.O. N.P. P.A. State: ZIP: Address: City: St Lic #: NPI #: DEA #: UPIN: Phone: Fax: Provider Email: Office Contact Name: Phone: Specialty (Check one): Pulmonologist Other: D. DISPENSING PROVIDER/ADMINISTRATION INFORMATION Place of Administration: Dispensing Provider/Pharmacy: ☐ Outpatient Dialysis Center ☐ Physician's Office ☐ Self-administered ☐ Physician's Office ☐ Home Outpatient Infusion Center Phone: ☐ Retail Pharmacy ☐ Specialty Pharmacy Center Name: ____ ☐ Mail Order Other: Home Infusion Center Phone: Agency Name: _ Administration code(s) (CPT): City: _____ State: ____ ZIP: _____ City: ______ State: _____ ZIP: _____ Phone: _____ Fax: _____ Phone: _____ Fax: _____ TIN: _____ PIN: ____ **TIN:** ______ PIN: _____ NPI: NPI: _____ E. PRODUCT INFORMATION Request is for: ☐ Aralast NP ☐ Glassia ☐ Prolastin-C ☐ Zemaira Dose: Frequency: F. DIAGNOSIS INFORMATION – Please indicate primary ICD Code and specify any other where applicable. Primary ICD Code: Secondary ICD Code: Other ICD Code: G. CLINICAL INFORMATION - Required clinical information must be completed in its entirety for all precertification requests. For All Requests: (clinical documentation required for all requests) Note: Aralast NP and Glassia are non-preferred. The preferred products are Prolastin-C and Zemaira. Yes No Has the patient had prior therapy with Aralast NP or Glassia within the last 365 days?

☐ Yes ☐ No Has the patient had a trial and failure, intolerance, or contraindication to Prolastin-C or Zemaira? Please explain if there are any other medical reason(s) that the patient cannot use Prolastin-C or Zemaira:

Continued on next page

For Medicare Advantage Part B:

FAX: <u>1-844-268-7263</u>

Please use other form.

For other lines of business:

Phone: 1-866-503-0857 (TTY: 711)



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For Medicare Advantage Part B: Phone: <u>1-866-503-0857</u> (TTY: <u>711)</u> FAX: <u>1-844-268-7263</u>

For other lines of business: Please use other form.

Note: Aralast NP and Glassia are non-preferred. The preferred products are Prolastin-C and Zemaira.

| Patient First Name | Patient Last Name | Patient Phone | Patient DOB |
|--|--|--|---|
| G. CLINICAL INFORMATION (continued) – | Required clinical information must be comp | leted in its entirety for all p | recertification requests. |
| interventions immediately Yes No Does the para outpatient he para infusion there were not para infusion there with the para infusion there were not para infusion to para infusion there were not para infusion to para infusion there were not para infusion to para infusion the para infusion to para inf | utpatient hospital setting? ient experienced an adverse event with the s (e.g., acetaminophen, steroids, diphenhyr after an infusion? atient have laboratory confirmed IgA antiboratient have severe venous access issues th ospital setting? atient have significant behavioral issues and rapy AND the patient does not have access vide a description of the behavioral issue on at medically unstable which may include res bility to tolerate a large volume or load or p an alternate setting without appropriate me | requested product that had dramine, fluids, other pre-ndies? at require the use of special dramine are consistive in a caregiver? r impairment: spiratory, cardiovascular, or redispose the member to a edical personnel and equip | s not responded to conventional medications or slowing of infusion rate) or all interventions only available in the mpairment that would impact the safety of the renal conditions that may limit the a severe adverse event that cannot be |
| , and the second | | | |
| | ☐ Rena | al: | _ |
| Other: | | | |
| ☐ Yes ☐ No Has the patient been diagnosed with alpha 1-antitrypsin (AAT) deficiency? ☐ Yes ☐ No Does the patient have a documented diagnosis of emphysema due to alpha 1-antitrypsin (AAT) deficiency? | | | |
| For Initiation of Therapy: | | | |
| Yes No Is this request for Aralast NP o | or Glassia? | | |
| 1 1 | ent had an intolerance or an ineffective resp | oonse to Prolastin-C or Zer | maira? |
| ☐ Yes ☐ No Does the patient have a contraindication to Prolastin-C or Zemaira? | | | |
| Yes No Is the patient's pretreatment post-bronchodilation FEV1 (forced expiratory volume 1 second) greater than or equal to 25 percent and less than | | | |
| or equal to 80 percent of the predicted value? | | | |
| Please provide the patient's pretreatment alpha 1-antitrypsin (AAT) serum concentration: specify result: mg/dL, uM/L, g/L, or µmol/L | | | |
| Please specify the alpha 1-antitrypsin (AAT) protein phenotype: PiZZ PiZ (null) Pi (null, null) PiMZ PiMS Other phenotype associated with serum AAT concentrations of less than 11 micromol/L | | | |
| | <u> </u> | imunodiffusion or 50 mg/dl | |
| For Continuation of Therapy: | | | |
| Yes No Is the patient currently receiving Yes No Is the patient experiencing ben | | manufacturer's patient as: | sistance program? |
| H. ACKNOWLEDGEMENT | | | |
| Request Completed By (Signature Require | red): | | Date: / / |
| | · · | procedure or service with | |
| Any person who knowingly files a request for authorization of coverage of a medical procedure or service with the intent to injure, defraud or deceive any insurance company by providing materially false information or conceals material information for the purpose of misleading, commits a fraudulent | | | |

insurance act, which is a crime and subjects such person to criminal and civil penalties. The plan may request additional information or clarification, if needed, to evaluate requests.