

Provider Help Desk 1.866.399.0928

Request for Prior Authorization CYSTIC FIBROSIS AGENTS, ORAL

(PLEASE PRINT – ACCURACY IS IMPORTANT)

| IA Medicaid Member ID # | Patient name | DOB | |
|-------------------------------------|---|-------------------------|-------------|
| Patient address | | | |
| Provider NPI | Prescriber name | Phone | |
| Prescriber address | | Fax | |
| Pharmacy name | Address | Phone | |
| Prescriber must complete all inform | ation above. It must be legible, correct, and c | omplete or form will be | e returned. |
| Pharmacy NPI | Pharmacy fax | NDC | |
| | | | |

Prior authorization (PA) is required for oral cystic fibrosis agents. Payment will be considered for patients when the following criteria are met:

- 1) Patient meets the FDA approved age; and
- 2) Patient has a diagnosis of cystic fibrosis (CF); and
- 3) Patient has a mutation in the cystic fibrosis transmembrane conductance regulator (CFTR) gene confirmed by an FDA-cleared CF mutation test (attach test results) for which the requested drug is indicated; and
- 4) Prescriber is a CF specialist or pulmonologist; and
- 5) Baseline liver function tests (AST, ALT, and bilirubin) are provided; and
- 6) Requests for Trikafta will not be considered for patients with severe hepatic impairment (Child-Pugh Class C); and
- 7) Will not be used with other CFTR modulator therapies.

If the criteria for coverage are met, an initial authorization will be given for 6 months. Additional approvals will be granted if the following criteria are met:

- 1) Adherence to oral cystic fibrosis therapy is confirmed; and
- 2) Liver function tests (AST, ALT, and bilirubin) are assessed every 3 months during the first year of treatment and annually thereafter.

| Non-Preferre | <u>d</u> | | | | | | | |
|---------------|-----------|-----------|-----------------------------------|------------|-----------|----------|-------------|---|
| Kalydeco | Or | kambi 🗌 |] Symdeko | 🗌 Trikat | fta | | | |
| | Strength | Dos | age Instructior | ıs Q | uantity | Days Sup | oly | |
| • | | | red CF mutatio on test (AST/AL | | | | - | _ |
| | | | | , , , | - | | | |
| Prescriber Sp | pecialty: | CF Specia | ilist 🛛 Pulmo | nologist [| Other (sp | pecify): | | _ |
| Rev. 11/20 | | | | | | | Page 1 of 2 | |



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| Request for Prior Authorization | |
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| IVACAFTOR (KALYDECO™) | |

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| Will requested medication be used with other CFTR modulator therapies? No Yes |
|--|
| Trifakta Requests: |
| Does patient have severe hepatic impairment (Child-Pugh Class C)? No Ves |
| Renewal Requests: |
| Patient is adherent to oral cystic fibrosis therapy: Yes No |
| Liver function tests (AST/ALT/bilirubin) are assessed every 3 months during first year of treatment and annually thereafter: |

Attach lab results and other documentation as necessary.

| Prescriber signature (Must match prescriber listed above.) | Date of submission | | |
|---|--------------------|--|--|
| | | | |
| IMPORTANT NOTE: In evaluating requests for prior authorization the consultant will consider the treatment from the standpoint of medical necessity only. If approval of this request is granted, this does not indicate that the member continues to be eligible for | | | |
| Medicaid. It is the responsibility of the provider who initiates the request for prior authorization to establish by inspection of the member's Medicaid eligibility card and, if necessary by contact with the county Department of Human Services, that the member | | | |

continues to be eligible for Medicaid.