

Poteligeo

Poteligeo (mogamulizumab-kpkc) J9204 Prior Authorization Request Medicare Part B Form

Instructions: * Indicates required information – Form may be returned if required information is not provided. Please fax this request to the appropriate fax number listed at the bottom of the page.

| □ Standard Request– (72 Hours) | | | | | ☐ Urgent Request (standard time frame could place the member's life, health or ability in serious jeopardy) | | | | | | | | | |
|--|--|---|-------------|----|--|-----------|---------------|----|---|-----------|---------------------------------------|--|--|--|
| | Date Req | uested | , | | | | | | | | | | | |
| | Requesto | | Phone / Fax | | | | | | | | | | | |
| | MEMBER INFORMATION | | | | | | | | | | | | | |
| *Na | me: | *!! | D#: *DOB: | | | | | | | | | | | |
| | PRESCRIBER INFORMATION | | | | | | | | | | | | | |
| *Name: | | | | | | | | | | | · · · · · · · · · · · · · · · · · · · | | | |
| *Add | dress: | | *Fax: | | | | | | | | | | | |
| | DISPENSING PROVIDER / ADMINISTRATION INFORMATION | | | | | | | | | | | | | |
| *Name: Phone: | | | | | | | | | | | | | | |
| *Address: Fax: | | | | | | | | | | | | | | |
| | | PROCEDURE / F | PROD | DU | CT INI | FOR | MATIO | NC | | I | I | | | |
| нс | PC Code | Name of Drug | Dos | se | (Wt: _ | | kg H | t: |) | Frequency | End Date if known | | | |
| | | | | | | | | | | | | | | |
| □ Self-administered □ Provider-administered □ Home Infusion | | | | | | | | | | | | | | |
| □ Chart notes attached. Other important information: | | | | | | | | | | | | | | |
| Diagnosis: ICD10: Description: | | | | | | | | | | | | | | |
| □ Provider attests the diagnosis provided is an FDA-Approved indication for this drug | | | | | | | | | | | | | | |
| | | | | | | | | | | | | | | |
| □ New Start or Initial Request: (Clinical documentation required for all requests) □ Patient has a diagnosis of Mycosis Fungoides (MF)/Sezary Syndrome □ Patient has relapsed or refractory disease; AND □ Patient has received at least one previous systemic therapy (note: topical and/or photochemotherapy cannot be considered systemic therapies); □ OR □ Used as primary treatment as systemic therapy (excluding use in stage IA mycosis fungoides) □ Patient has a diagnosis of Adult T-Cell Leukemia/Lymphoma □ Poteligeo is being used as subsequent therapy in patients with acute or lymphoma subtypes which did not respond to first-line therapy | | | | | | | | | | | | | | |
| | □ Patien | tion Requests: (Clinical docume t had an <u>adequate response</u> or <u>significal</u> please provide clinical rationale for contin | cant | im | nprove | - emer | <u>nt</u> whi | | - | • | | | | |

| ACKNOWLEDGEMENT | | | | | | | | | | | |
|---|---------------|------------|--------|--|--|--|--|--|--|--|--|
| Request By (Signature Required):D |)ate: | _/ | _/ | | | | | | | | |
| 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. THIS AUTHORIZATION IS NOT A GUARANTEE OF PAYMENT. PAYMENT IS BASED ON BENE SERVICE, MEMBER ELIGIBILITY AND MEDICAL NECESSITY. | FITS IN EFFEC | T AT THE T | TME OF | | | | | | | | |



Prior Authorization Group - Poteligeo Drug PA

Drug Name(s):

POTELIGEO

MOGAMULIZUMAB-KPKC

Criteria for approval of Non-Formulary/Preferred Drug:

- 1. Prescribed for an approved FDA diagnosis (as listed below):
- 2. Member does not have any clinically relevant contraindications, or CMS/Plan exclusions, to the requested drug.
- If the member meets all these criteria, they may be approved by the Plan for the requested drug.
- Quantity limits and Tiering will be determined by the Plan.

Exclusion Criteria:

N/A

Prescriber Restrictions:

Dermatopathologist, Hematopathologist or another related specialist

Coverage Duration:

Initial Approval will be for 6 months

Continuation will be approved for 12 months

FDA Indications:

Poteligeo

- Mycosis fungoides, Relapsed or refractory disease after at least 1 prior systemic therapy
- Sezary's disease, Relapsed or refractory disease after at least 1 prior systemic therapy

Off-Label Uses:

N/A

Age Restrictions:

Safety and effectiveness have not been established in pediatric patients

Other Clinical Consideration:

N/A

Resources:

https://www.micromedexsolutions.com/micromedex2/librarian/CS/20D942/ND_PR/evidencexpert/ND_P/evidencexpert/DUPLICATIONSHIELDSYNC/CA3F98/ND_PG/evidencexpert/ND_B/evidencexpert/ND_AppProduct/evidencexpert/ND_T /evidencexpert/PFActionId/evidencexpert.GoToDashboard?docId=932521&contentSetId=100&title=Mogamulizumab-kpkc&servicesTitle=Mogamulizumab-kpkc&brandName=Poteligeo&UserMdxSearchTerm=Poteligeo&=null#