

MEDICARE FORM

Lemtrada® (alemtuzumab) **Medication Precertification Request**

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Note: Lemtrada is non-(All fields must be completed and legible for precertification review.) preferred. The preferred product Please indicate: is Tysabri for MA plans and ☐ Start of treatment: Start date / / Kesimpta for MAPD plans. ☐ Continuation of therapy: Date of last treatment ____ / __/ Phone: ___ Precertification Requested By: _ Fax: ___ A. PATIENT INFORMATION Last Name: First Name: Address: City: State: 7IP: Work Phone: Home Phone: Cell Phone: DOB: Allergies: E-mail: Current Weight: ___ _ lbs or _____ kgs Height: _____ inches or ___ cms **B. INSURANCE INFORMATION** Aetna Member ID #: Does patient have other coverage? ☐ Yes ☐ No Group #: _____ If yes, provide ID#: _____ Carrier Name: ____

Insured: C. PRESCRIBER INFORMATION Last Name: First Name: (Check One): M.D. D.O. N.P. P.A. State: ZIP: Address: City: NPI#: St Lic #: DEA #: UPIN: Phone: Fax: Office Contact Name: Phone: Provider Email: D. DISPENSING PROVIDER/ADMINISTRATION INFORMATION Place of Administration: Dispensing Provider/Pharmacy: ☐ Physician's Office ☐ Self-administered ☐ Physician's Office ☐ Retail Pharmacy Outpatient Infusion Center Phone: ☐ Specialty Pharmacy ☐ Mail Order ☐ Other: _____ Center Name: __ Phone: Name: _____ ☐ Home Infusion Center Address: Agency Name: Phone: _____ Fax: __ Administration code(s) (CPT): Address: TIN: PIN: E. PRODUCT INFORMATION Request is for Lemtrada: Dose: ___ Frequency: ___ **HCPCS Code:** 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 Note: Lemtrada is non-preferred. The preferred product is Tysabri for MA plans and Kesimpta for MAPD plans. ☐ Yes ☐ No Has the patient had prior therapy with Lemtrada (alemtuzumab) within the last 365 days? ☐ Yes ☐ No Has the patient had a trial and failure, intolerance, or contraindication to Tysabri (natalizumab)? Please explain if there are any other medical reason(s) that the patient cannot use Tysabri (natalizumab). ☐ Yes ☐ No Has the patient had a trial and failure, intolerance, or contraindication to Kesimpta (ofatumumab)? Please explain if there are any other medical reason(s) that the patient cannot use Kesimpta (ofatumumab). Please indicate the type of multiple sclerosis the patient has been diagnosed with: ☐ Relapsing-remitting (RRMS) ☐ Secondary-progressive MS (SPMS) ☐ Primary-progressive MS (PPMS) ☐ Progressive-relapsing MS (PRMS) ☐ Yes ☐ No Has the patient discontinued other medications used for treating MS (not including Ampyra)? ☐ Yes ☐ No Will a maximum of two courses of Lemtrada be utilized? Please indicate the patient's HIV status:

Positive

Negative

Unknown For Continuation requests: ☐ Yes ☐ No Is this continuation request a result of the patient receiving samples of Lemtrada? ☐ Yes ☐ No Does the patient have a documented severe and/or potentially life threatening adverse event that occurred during or following the previous infusion? → ☐ Yes ☐ No Could the adverse reaction be managed through pre-medication in the office setting?

For Illinois MMP:

FAX: 1-855-320-8445 PHONE: 1-866-600-2139

Please use other form.

For other lines of business:



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Patient First Name	Patient Last Name	Patient Phone	Patient DOB
H. ACKNOWLEDGEMENT			
Request Completed By (Signature Required):			Date: /
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.