MEDICARE FORM
Ilumya™ (tildrakizumab-asmn) Injectable Medication Precertification Request

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

For Medicare Advantage Part B:
For other lines of business:
Please use commercial form.

Note: Ilumya is non-preferred. Preferred products vary based on plan type. See section G below.

Would you like to use electronic prior authorization? Consider using **Availity**, our electronic prior authorization portal. Learn more about **Availity** from the links in the table below.

For phone or fax requests, refer to the table below for routing information. To determine which box to use, refer to the patient's Aetna ID card. State specific special needs and Medicare-Medicaid Plans may be designated on the front of the ID card or in the website URL on the back of the card. If you don't see your specific plan listed, call the number on the back of the member's ID card to confirm routing information.

For **Aetna Medicare Advantage** and **Allina Health Aetna Medicare** members send request to: **Phone:** 1-866-503-0857 **(TTY:** 711**)** **Fax:** 1-844-268-7263 **Availity:** https://www.aetna.com/health-care-professionals/resource-center/availity.html
For Aetna Medicare Advantage **Virginia Dual Eligible Special Needs Plans** (HMO D-SNP) send request to: **Phone:** 1-855-463-0933 **Fax:** 1-833-280-5224 **Availity:** https://www.aetnabetterhealth.com/virginia-hmosnp/providers/portal
For Aetna Assure Premier Plus Medicare Advantage **New Jersey Dual Eligible Special Needs Plans** (HMO D-SNP) send request to: **Phone:** 1-844-362-0934 **Fax:** 1-833-322-0034 **Availity:** https://www.aetnabetterhealth.com/new-jersey-hmosnp/providers/portal.html
For Aetna Better Health of **Illinois Premier Medicare Medicaid Plan** (MMP) send request to: **Phone:** 1-866-600-2139 **FAX:** 1-855-320-8445 **Availity:** https://www.aetnabetterhealth.com/illinois/providers/portal
For Aetna Better Health of **Ohio Premier Medicare Medicaid Plan** (MMP) send request to: **Phone:** 1-855-364-0974 **Fax:** 1-855-734-9389 **Availity:** https://www.aetnabetterhealth.com/ohio/providers/portal
For Aetna Better Health of **Michigan Premier Medicare Medicaid Plan** (MMP) send request to: **Phone:** 1-855-676-5772 **Fax:** 1-844-241-2495 **Availity:** https://www.aetnabetterhealth.com/michigan/providers/portal.html

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Please indicate: ☐ Start of treatment: Start date ____/____/____
☐ Continuation of therapy: Date of last treatment ____/____/____

Precertification Requested By: _____ Phone: _____ Fax: _____

A. PATIENT INFORMATION

First Name:	Last Name:		
Address:	City:	State:	ZIP:
Home Phone:	Work Phone:	Cell Phone:	
DOB:	Allergies:	E-mail:	

Current Weight: _____ lbs or _____ kgs Height: _____ inches or _____ cms

B. INSURANCE INFORMATION

Aetna Member ID #: _____
Group #: _____
Insured: _____

Does patient have other coverage? ☐ Yes ☐ No
If yes, provide ID#: _____ Carrier Name: _____
Insured: _____

C. PRESCRIBER INFORMATION

First Name:	Last Name:		*(Check One):* ☐ M.D. ☐ D.O. ☐ N.P. ☐ P.A.		
Address:		City:	State:	ZIP:	
Phone:	Fax:	St Lic #:	NPI #:	DEA #:	UPIN:
Provider Email:	Office Contact Name:		Phone:		

D. DISPENSING PROVIDER/ADMINISTRATION INFORMATION

Place of Administration:
☐ Self-administered ☐ Physician's Office
☐ Outpatient Infusion Center Phone: _____
Center Name: _____
☐ Home Infusion Center Phone: _____
Agency Name: _____
☐ Administration code(s) (CPT): _____
Address: _____
City: _____ State: _____ ZIP: _____
Phone: _____ Fax: _____
TIN: _____ PIN: _____
NPI: _____

Dispensing Provider/Pharmacy:
☐ Physician's Office ☐ Retail Pharmacy
☐ Specialty Pharmacy ☐ Other _____
Name: _____
Address: _____
City: _____ State: _____ ZIP: _____
Phone: _____ Fax: _____
TIN: _____ PIN: _____
NPI: _____

E. PRODUCT INFORMATION

Request is for: Ilumya (tildrakizumab-asmn): 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 Initiation Requests (clinical documentation required for all requests):

Note: Ilumya is non-preferred. Inflectra and Renflexis are preferred for MA plans. Cosentyx SC, Enbrel, Humira, Idacio, Skyrizi, Sotyktu, Stelara and Tremfya are preferred for MAPD plans.

☐ Yes ☐ No Has the patient had prior therapy with Ilumya (tildrakizumab-asmn) within the last 365 days?

☐ No Has the patient had a trial and failure of any of the following? (if yes, select all that apply below)
- ☐ Inflectra (infliximab-dyyb) ☐ Renflexis (infliximab-abda)
- When was the member's trial and failure of the preferred drug? _____
- Please describe the nature of the failure of the preferred drug _____

☐ No Has the patient had an adverse reaction to any of the following? (if yes, select all that apply below)
- ☐ Inflectra (infliximab-dyyb) ☐ Renflexis (infliximab-abda)
- When was the member's adverse reaction to the preferred drug? _____
- Please describe the nature of the adverse reaction to the preferred drug _____

☐ No Has the patient had a trial and failure of any of the following? (if yes, select all that apply below)
- ☐ Cosentyx SC (secukinumab) ☐ Enbrel (etanercept) ☐ Humira (adalimumab) ☐ Idacio (adalimumab-aacf)
- ☐ Skyrizi (risankizumab-rzaa) ☐ Sotyktu (deucravacitinib) ☐ Stelara (ustekinumab) ☐ Tremfya (guselkumab)
- When was the member's trial and failure of the preferred drug? _____
- Please describe the nature of the failure of the preferred drug _____

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Patient First Name	Patient Last Name	Patient Phone	Patient DOB

G. CLINICAL INFORMATION *(continued)* - Required clinical information must be completed in its entirety for all precertification requests.

For Initiation Requests *continued* (clinical documentation required for all requests):

☐ No Has the patient had an adverse reaction to any of the following? (if yes, select all that apply below)
☐ Cosentyx SC (secukinumab) ☐ Enbrel (etanercept) ☐ Humira (adalimumab) ☐ Idacio (adalimumab-aacf)
☐ Skyrizi (risankizumab-rzaa) ☐ Sotyktu (deucravacitinib) ☐ Stelara (ustekinumab) ☐ Tremfya (guselkumab)
→ When was the member's adverse reaction to the preferred drug? _____
→ Please describe the nature of the adverse reaction to the preferred drug _____

Please explain if there are any contraindications or other medical reason(s) that the patient cannot use any of the following preferred products when indicated for the patient's diagnosis (select all that apply):
☐ Inflectra (infliximab-dyyb) ☐ Renflexis (infliximab-abda)

_____

_____

Please explain if there are any contraindications or other medical reason(s) that the patient cannot use any of the following preferred products when indicated for the patient's diagnosis (select all that apply):
☐ Cosentyx SC (secukinumab) ☐ Enbrel (etanercept) ☐ Humira (adalimumab) ☐ Idacio (adalimumab-aacf) ☐ Skyrizi (risankizumab-rzaa)
☐ Sotyktu (deucravacitinib) ☐ Stelara (ustekinumab) ☐ Tremfya (guselkumab)

_____

_____

Plaque Psoriasis:
Please indicate the severity of the patient's disease: ☐ mild ☐ moderate ☐ severe
☐ Yes ☐ No Is there evidence that the disease is active?
☐ Yes ☐ No Is there clinical documentation of chronic disease?
☐ Yes ☐ No Is the patient a candidate for systemic therapy or phototherapy?
Please select: ☐ phototherapy ☐ systemic therapy ☐ phototherapy and systemic therapy
Please provide the patient's Psoriasis Area and Severity Index (PASI) score: _____
Please indicate the percentage of body surface area affected by plaque psoriasis: _____%
☐ Yes ☐ No Does the plaque psoriasis involve sensitive areas? ***If yes***, please select: ☐ hands ☐ feet ☐ face ☐ genitals
☐ Yes ☐ No Was the trial with systemic conventional DMARD(s) (e.g., methotrexate, acetretin, or cyclosporine) ineffective?
→ ☐ Yes ☐ No Was the trial with systemic conventional DMARD(s) not tolerated?
→ ☐ Yes ☐ No Are systemic conventional DMARDs contraindicated?
→ Please select: ☐ acetretin ☐ cyclosporine ☐ methotrexate ☐ mycophenolate ☐ None of the above
Please indicate the length of the medication trial: ☐ Less than 1 month ☐ 1 month ☐ 2 months ☐ 3 months or greater
☐ Yes ☐ No Was the trial with phototherapy ineffective?
→ ☐ Yes ☐ No Was the trial with phototherapy not tolerated?
→ ☐ Yes ☐ No Is phototherapy contraindicated?
→ Please check all that apply: ☐ Psoralens (methoxsalen, trioxsalen) with UVA light (PUVA)
☐ UVB with coal tar or dithranol
☐ UVB (standard or narrow band)
☐ Home UVB
☐ None of the above
Please indicate the length of trial: ☐ Less than 1 month ☐ 1 month ☐ 2 months ☐ 3 months or greater

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Patient First Name	Patient Last Name	Patient Phone	Patient DOB

G. CLINICAL INFORMATION *(continued)* - Required clinical information must be completed in its entirety for all precertification requests.

For Continuation of Therapy (clinical documentation required for all requests):

Please indicate the length of time on Ilumya (tildrakizumab-asmn): _____

☐ Yes ☐ No Is this continuation request a result of the patient receiving samples of Ilumya (tildrakizumab-asmn)?

☐ Yes ☐ No Will Ilumya (tildrakizumab-asmn) be used concomitantly with apremilast, tofacitinib, or other biologic DMARDs (e.g., adalimumab, certolizumab)?

☐ Yes ☐ No Is there clinical documentation supporting disease stability?

☐ Yes ☐ No Is there clinical documentation supporting disease improvement?

☐ Yes ☐ No Does the patient have any risk factors for TB?

→ ☐ Yes ☐ No Has the patient had a TB test within the past year?

→ (check all that apply): ☐ PPD test ☐ interferon-gamma assay (IGRA) ☐ chest x-ray

Please enter the results of the TB test: ☐ positive ☐ negative ☐ unknown

☐ Yes ☐ No Has the patient received Ilumya (tildrakizumab-asmn) within the past 6 months?

→ ☐ 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 home or office setting?

Please indicate the severity of the disease at baseline (pretreatment with Ilumya (tildrakizumab-asmn)): ☐ mild ☐ moderate ☐ severe

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.