

STEP THERAPY PA REQUEST FORM - Proton Pump Inhibitors, Statins, Anti-migraine, Topical Acne Agents and Cytokine & CAM Antagonists[This and other pharmacy PA forms are available at www.ctdssmap.com and can be accessed by clicking on the pharmacy icon]**PA Criteria for Step Therapy Drug Products**

- The Pharmacy team will validate the client's history for the use of preferred agent(s) before approving a non-preferred agent. Non-Preferred drug approvals require documented evidence that the patient has tried and failed, is intolerant to, or has a contraindication to a normal course of therapy with at least one preferred drug in the class.
- For clients new to Medicaid, a pharmacy profile history showing previously failed preferred products, outcomes and compliance with the medication regimen length shall be provided with the non-preferred product request form.
- Clinical prior authorization must be obtained for any non-preferred step therapy drug **using this form only, not the standard drug PA form.**
- A copy of your filed [FDA 3500 Med Watch Form](#) is required if patients have experienced significant adverse effect

Prescriber and Member Information**Please Print:****Note - Incomplete requests will not be granted.**

1. Prescriber's Name (Last, First)	5. Member's Name (Last, First)
2. Prescriber's NPI	6. Member's ID
3. Prescriber's Phone	7. Member's Date of Birth (MM/DD/CCYY)
4. Prescriber's Fax	8. Pharmacy Name & Fax
9. Drug & Dosage Form (print)	
10. Route <input type="checkbox"/> Oral <input type="checkbox"/> Topical <input type="checkbox"/> Inhalation <input type="checkbox"/> Injectable	
11. Strength	12. Quantity
	13. Frequency of Dosing

Medical History**Note - Incomplete requests will be denied.**Please explain why the patient cannot be treated with a preferred alternative. You MUST indicate which preferred product has been utilized in the past, circle a reason for the failure (listed below), AND supply a specific written clinical explanation.

14. Preferred Product Trial (Name & Daily Dose)	15. Reason	16. Clinical Explanation (including length of therapy, date commenced, and outcome)
	1 2 3 4	
1. Use of the preferred alternative is contraindicated. 2. The patient has experienced <u>significant</u> adverse effects from the preferred alternative, Completed FDA 3500 MedWatch form attached and filed with the FDA. 3. Use of the preferred alternative has resulted in therapeutic failure <u>after the normal course of treatment.</u> 4. Pediatric patient (younger than 12 years of age).		

Please Note: Pharmacies should not be contacting prescribers to provide pre-signed PA forms or submitting pre-signed forms for PA, nor should prescribing providers be requesting that pharmacies perform PA activities for them. PA requests must originate from the prescriber, and only the prescriber should sign the form at the time of PA submission.

I certify that documentation is maintained in my files and the information given is true and accurate for the medication requested, subject to penalty under Connecticut Gen. Stat. Section 17b-99 and Regulations of Conn. State Agencies Sections 17-83k-1-3 and 4a-7, inclusive. I certify that this member is under my clinic's/practice's ongoing care.

17. Signature of Prescriber* _____ 18. Date (MM/DD/CCYY) _____

* **Mandatory (others may not sign for prescriber)** In accordance with mandates set forth in the Affordable Care Act (ACA), providers who order, prescribe, or refer clients for services must be enrolled in the Connecticut Medical Assistance Program (CMAP). Effective 10/1/2013, any prescriptions or services provided by a non-enrolled provider shall no longer be considered/covered by CMAP.

This form (and attachments) contains protected health information (PHI) for Gainwell Technologies and is covered by the Electronic Communications Privacy Act, 18 U.S.C. § 2510-2521 and the Standards for Privacy of Individually Identifiable Health Information, 45 CFR Parts 160 and 164, which is intended only for the use of prior authorization. Any unintended recipient is hereby notified that the information is privileged and confidential, and any use, disclosure, or reproduction of this information is prohibited. Any unintended recipient should contact Gainwell Technologies by telephone at (860)255-3900 or by e-mail immediately and destroy the original message.

(Direction Sheet) Informational Only

No.	Name	Description
1.	Prescriber's Name (Last, First)	Enter the prescribing practitioner's last name and first name
2.	Prescriber's NPI	Enter the prescribing practitioner's National Provider Identification (NPI) number
3.	Prescriber's Phone	Enter the prescribing practitioner's phone number where a PA customer service representative can contact the practitioner for additional information or clarification, if necessary
4.	Prescriber's Fax	Enter the prescribing practitioner's fax number where a PA customer service representative can contact the practitioner for additional information or clarification, if necessary
5.	Member's Name (Last, First)	Enter the member's name as it appears on the member's CONNECT Card or as obtained from the Automated Eligibility Verification System (AEVS)
6.	Member's ID	Enter the member's 9-digit identification number as it appears on the member's CONNECT Card or as obtained from the Automated Eligibility Verification System (AEVS)
7.	Member's Date of Birth (MMDDCCYY)	Enter the member's date of birth in MM/DD/CCYY format
8.	Pharmacy's Name & Fax (optional)	Enter the pharmacy's name and fax number, if known
9.	Drug & Dosage Form	Print the drug info for which the Prior Authorization is being requested
10.	Route	Select the route of the drug being requested
11.	Strength	Enter the strength of the drug in milligrams
12.	Quantity	Enter the quantity of the drug being prescribed
13.	Frequency of Dosing	Enter the dosing frequency
14.	Preferred Product	Indicate which preferred drug the patient has tried and failed in the past including the dosage per day. Preferred Drug List
15.	Reason	Circle the number on the form which corresponds to the type of failure experienced, and submit any required documentation.
16.	Clinical Explanation	Provide a written clinical explanation of the indicated failure to a preferred product including length of therapy, outcome and when commenced.
17.	Signature of Prescriber	The prescribing practitioner must sign the PA form; agent's signature is not acceptable
18.	Date (MMDDCCYY)	Enter the date the form was completed, signed, and submitted in MM/DD/CCYY format

STEP THERAPY CATEGORIES

ACNE AGENTS, TOPICAL		
Preferred Step Therapy Agents	Non-Preferred Agents Requiring Step Therapy PA Request	
DIAGNOSIS CODE REQUIRED ON SELECT AGENTS - SEE NOTATION BELOW		
ACNE MEDICATION 5% & 10% GEL (OTC BENZOYL PEROXIDE)	ACANYA	DAPSONE
ACNE MEDICATION LOTION (OTC BENZOYL PEROXIDE)	ACNE MEDICATION 2.5% GEL (OTC BENZOYL	ERY 2% PADS, ERYGEL
ADAPALENE/BENZOYL PEROXIDE 0.1-2.5% (EPIDUO) (DX CODE REQ.)	ADAPALENE 0.1% CREAM (DX CODE REQ.)	ERYTHROMYCIN 2% GEL, PLEDGETS
ADAPALENE 0.1% GEL (OTC) (DX CODE REQ.)	ADAPALENE 0.3% GEL, GEL PUMP (DX CODE REQ.)	ERYTHROMYCIN-BENZOYL GEL
BENZOYL PEROXIDE 2.5%, 5%, 10% GEL (OTC)	ADAPALENE-BENZOYL PEROX 0.3-2.5% (DX CODE REQ)	EVOCOLIN
BENZOYL PEROXIDE 5%, 10% WASH (OTC)	ALTRENO (DX CODE REQ.)	FABIOR
CLINDAMYCIN PH 1% PLEGET	ARAZLO (DX CODE REQ.)	KLARON
CLINDAMYCIN PH 1% SOLUTION (not GEL or LOTION)	ATRALIN (DX CODE REQ.)	NEUAC
CLINDAMYCIN / BENZOYL PEROXIDE 1.2%-5% (DUAC)	AVAR, AVAR-E	ONEXTON
ERYTHROMYCIN 2% SOLUTION (not GEL)	AVITA (DX CODE REQ.)	OVACE
PANOXYL 10% ACNE FOAMING WASH (OTC)	BENZAMYCIN	RETIN-A MICRO (DX CODE REQ.)
RETIN-A CREAM (DX CODE REQ.)	BENZOYL PEROXIDE FOAM	ROSULA
RETIN-A GEL (not MICRO) (DX CODE REQ.)	BPO FOAMING CLOTHS	SODIUM SULFACETAMIDE
	CABTREO (DX CODE REQ.)	SODIUM SULFACETAMIDE-SULFUR
	CLEOCIN T	SSS CLEANSER, CREAM, FOAM
	CLINDACIN FOAM, KIT	SUMADAN, SUMAXIN
	CLINDAGEL	TAZAROTENE
	CLINDAMYCIN PH GEL, FOAM, LOTION	TRETINOIN, TRETINOIN GEL MICRO (DX CODE REQ.)
	CLIND PH-BENZOYL PEROX 1.2-2.5% GEL PUMP	TRETIN-X (DX CODE REQ.)
	CLIND PH-BENZOYL PEROX 1.2-3.75% GEL PUMP	WINLEVI
	CLIND PH-BENZOYL PEROX 1-5% GEL, GEL PUMP	ZIANA
	CLINDAMYCIN-TRETINOIN	ZMA CLEAR 9%-4.5% SUSPENSION
CYTOKINE/CAM ANTAGONISTS		
Preferred Step Therapy Agents	Non-Preferred Agents Requiring Step Therapy PA Request	
ENBREL DISP SYRINGE, KIT, PEN	ABRILADA	KINERET
ENBREL MINI CARTRIDGE	ACTEMRA	OLUMIANT
ENBREL VIAL	ADALIMUMAB-AACF, -ADAZ, -ADBM, -FKJP	OMVOH
HUMIRA KIT, PEN INJ KIT	AMJEVITA	ORENCIA
INFLIXIMAB VIAL *	ARCALYST	REMICADE
OTEZLA STARTER PACK, TABLET	AVSOLA	RENFLEXIS
XELJANZ IR TABLET (not XR or SOLUTION) *	BIMZELX	RINVOQ ER
	CIBINQO	SILIQ
	CIMZIA	SIMPONI
	COSENTYX	SKYRIZI
	CYLTEZO	SOTYKTU
	ENSPRYNG	SPEVIGO
	ENTYVIO	STELARA
	HADLIMA	TALTZ
	HULIO	TREMFYA
	HYRIMOZ	UPLIZNA
	IDACIO	VELSIPITY
	ILARIS	XELJANZ SOLUTION
	ILUMYA	XELJANZ XR
	INFLECTRA	YUFLYMA
	KEVZARA	YUSIMRY
ANTIMIGRAINE AGENTS, TRIPTAN		
Preferred Step Therapy Agents	Non-Preferred Agents Requiring Step Therapy PA Request	
IMITREX NASAL SPRAY	ALMOTRIPTAN	SUMATRIPTAN CARTRIDGE, INJECT, NASAL SPRAY
RELPAK TABLET	ELETRIPTAN	SUMATRIPTAN-NAPROXEN
RIZATRIPTAN ODT	FROVA & FROVATRIPTAN	TOSYMRA
RIZATRIPTAN TABLET	IMITREX CARTRIDGES, PEN INJECT, TABLET	ZEMBRACE
SUMATRIPTAN TABLET	MAXALT, MAXALT MLT	ZOLMITRIPTAN
SUMATRIPTAN VIAL (not CARTRIDGE)	NARATRIPTAN	ZOMIG
LIPOTROPICS, STATINS		
Preferred Step Therapy Agents	Non-Preferred Agents Requiring Step Therapy PA Request	
ATORVASTATIN TABLET	ALTOPREV	LESCOL XL
LOVASTATIN TABLET	AMLODIPINE-ATORVASTATIN	LIPITOR
PRAVASTATIN TABLET	ATORVALIQ	LIVALO
ROSUVASTATIN TABLET	CADUET	PITAVASTATIN
SIMVASTATIN TABLET	CRESTOR	VYTORIN
	EZALLOR SPRINKLE	ZOCOR
	EZETIMZBE-SIMVASTATIN	ZYPITAMAG
	FLUVASTATIN, FLUVASTATIN ER	
PROTON PUMP INHIBITORS		
Preferred Step Therapy Agents	Non-Preferred Agents Requiring Step Therapy PA Request	
DEXILANT CAPSULE	ACIPHEX	OMEPRAZOLE (OTC VERSIONS)
ESOMEPRAZOLE 20MG CAPSULE (OTC & RX)	DEXLANSOPRAZOLE CAPSULES	OMEPRAZOLE-SODIUM BICARBONATE
ESOMEPRAZOLE 40MG CAPSULE	ESOMEPRAZOLE PACKET	PANTOPRAZOLE 40 MG SUSPENSION PKT
NEXIUM PACKET SUSPENSION (not CAPSULE)	ESOMEPRAZOLE (OTC TABLET)	PREVACID
OMEPRAZOLE 10MG, 20MG, 40MG CAPSULE (Rx ONLY)	KONVOMEF	PRILOSEC
PANTOPRAZOLE TABLET	LANSOPRAZOLE (OTC VERSIONS)	PROTONIX TABLET
PROTONIX SUSPENSION	LANSOPRAZOLE CAPSULE, ODT	RABEPRAZOLE
	NEXIUM CAPSULE	ZEGERID

(DX Code Req) notation for agents that require ICD-10 code

Preferred Brand Name medications are listed in **BOLD**

*Updated 1/1/2024