











Prior Authorization Request Administrative Information

Member Information						
Last name	First name		МІ			
Member ID	Date of birth					
Sex assigned at birth Female Male "X" or Intersex						
Current gender Female Male Transgender male Transgender female Other						
Place of residence						
Race/ethnicity Preferred spoken language Preferred written language						
MassHealth does not exclude people or treat them differently because of race, color, national origin, age, disability, religion, creed, sexual orientation, or sex (including gender identity and gender stereotyping).						
Plan Contact Information						
Please indicate the member's MassHealth Plan and fax or submit this completed and signed form according to the Plan's contact information below.						
MassHealth Fee-For-Service (FFS) Plan, Primary Care Clinician (PCC) Plan, Primary Care Accountable Care Organization (PCACO) Plan, Children's Medical Security Plan, and Health Safety Net Plan						
☐ MassHealth Drug Utilization Review Prog	gram					
Pharmacy: Fax: (877) 208-7428 - Tel: (800) 745-7318					
MassHealth Managed Care Organization	n (MCO) and Acco	untable Care Partnershi	p Plans (ACPP)			
☐ Fallon Health						
Online Prior Authorization: go.covermymeds.com/OptumRx						
Online Prior Authorization: providerportal.s	urescripts.net/Provi	derPortal/optum				
Pharmacy: Fax: (844) 403-1029 - Tel: (844) 720-0033						
☐ Health New England						
Online Prior Authorization: go.covermymeds.com/OptumRx						
Pharmacy: Fax: (800) 550-9246 - Tel: (800) 918-7545						
Online Prior Authorization (Non-Specialty Drugs): go.covermymeds.com/OptumRx						
Online Prior Authorization (Specialty/Medical Drugs): provider.massgeneralbrighamhealthplan.org						
Pharmacy: Fax: (844) 403-1029 - Tel: (800) 711-4555						
☐ Tufts Health Plan						
Online Prior Authorization: point32health.promptpa.com						
Pharmacy: Fax: (617) 673-0939 - Tel: (888) 257-1985						
□ WellSense Health Plan						
Online Prior Authorization: wellsense.org/providers/ma/pharmacy/prior-authorizations						
Pharmacy: Fax: (833) 951-1680 - Tel: (877) 417-1822						

Topical Corticosteroids Prior Authorization Request

MassHealth reviews requests for prior authorization (PA) on the basis of medical necessity only. If MassHealth approves the request, payment is still subject to all general conditions of MassHealth, including current member eligibility, other insurance, and program restrictions. MassHealth will notify the requesting provider and member of its decision. Keep a copy of this form for your records. If faxing this form, please use black ink.

Additional information about these agents, including PA requirements and preferred products, can be found within the MassHealth Drug List at www.mass.gov/druglist.

Medication information Medication requested					
Class I Superpotent products (See Sections I., II., and III.) clobetasol propionate (Impeklo, Olux-E): foam/emollient, lotion, shampoo-kit diflorasone: ointment	☐ halobetasol: foam☐ halobetasol (Bryhali, Ultravate): lotion				
Class II Potent products (See Sections I., II., and III.) betamethasone dipropionate (Sernivo): spray desoximetasone (Topicort): ointment, spray (0.25%), gel (0.05%)	☐ diflorasone (Apexicon-E): cream ☐ halcinonide (Halog): cream, solution				
Class III Upper Mid-Strength Potent products (See Sections amcinonide: cream desoximetasone (Topicort): cream, ointment	I., II., and III.) diflorasone: cream				
Class IV Mid-Strength Potent products (See Sections I., II., a clocortolone (Cloderm): cream fluocinolone (Synalar): ointment-kit flurandrenolide: ointment	Ind III.)				
Class V Lower Mid-Strength Potent products (See Sections desonide: lotion fluocinolone (Synalar): cream-kit flurandrenolide: cream, lotion	I., II., and III.) ☐ fluticasone propionate: lotion ☐ hydrocortisone butyrate: lotion ☐ hydrocortisone butyrate/emollient: cream				
Class VI Mild Potent products (See Sections I., II., and III.) ☐ fluocinolone (Synalar): solution-kit					
Class VII Least Potent products (See Sections I., II., and III.) hydrocortisone: solution					
Combination products betamethasone/calcipotriene (Taclonex): ointment, scalp suspension	☐ halobetasol/tazarotene (Duobrii): lotion☐ neomycin/fluocinolone: cream, cream-kit				
Strength and formulation requested					
Frequency and duration of therapy	Drug NDC (if known)				
Indication(s) or ICD-10 code(s), if applicable PA-66 (Rev. 04/24)	over				

Section I. Please complete for all requests, excluding combination products. Has the member had a trial with all topical corticosteroids of the same formulation and potency range that are available without prior authorization? Yes. Please list the specific drug name, dates/duration of use, and outcomes below*. Drug name, strength, and formulation Dates/duration of use Did the member experience any of the following? Adverse reaction Inadequate response Briefly describe details of adverse reaction or inadequate response. Dates/duration of use Drug name, strength, and formulation Did the member experience any of the following? Adverse reaction Inadequate response Briefly describe details of adverse reaction or inadequate response. Dates/duration of use Drug name, strength, and formulation Did the member experience any of the following? Adverse reaction Inadequate response Briefly describe details of adverse reaction or inadequate response. Drug name, strength, and formulation Dates/duration of use Did the member experience any of the following? Adverse reaction Inadequate response Briefly describe details of adverse reaction or inadequate response. Dates/duration of use Drug name, strength, and formulation Did the member experience any of the following? Adverse reaction Inadequate response Briefly describe details of adverse reaction or inadequate response. No. Please explain contraindication or clinical rationale for not using other topical corticosteroid(s) that are available without prior authorization in this member. Section II. Please complete for foam and shampoo formulations in scalp-related conditions. Has the member had a trial with one topical corticosteroid of a similar formulation and similar or greater potency range that is available without prior authorization? Yes. Please list the specific drug name, dates/duration of use, and outcomes below*. Drug name, strength, and formulation Dates/duration of use Did the member experience any of the following? Adverse reaction Inadequate response Briefly describe details of adverse reaction or inadequate response. oxdot No. Please explain contraindication or clinical rationale for not using other topical corticosteroid(s) that are available without prior authorization for this member.

Ex	xplain medical necessity for the requested formulation.
Sec	ction IV. Please complete for combination products.
1.	Provide medical necessity for the combination product instead of the individual agents.
2.	For Duobrii, has the member had a trial with one superpotent or potent topical corticosteroid? Yes. Please list the specific drug name, dates/duration of use, and outcomes below.*
	Drug name, strength, and formulation Did the member experience any of the following? Adverse reaction Inadequate response Briefly describe details of adverse reaction or inadequate response.
	□ No
*Atta	ach a letter with additional information regarding medication trials as applicable.
1.	Is the alternative drug required under the step therapy protocol contraindicated, or will likely cause an adverse reaction in, or physical or mental harm to the member? Yes No If yes, briefly describe details of contraindication, adverse reaction, or harm.
	Is the alternative drug required under the step therapy protocol expected to be ineffective based on the know clinical characteristics of the member and the known characteristics of the alternative drug regimen? Yes No If yes, briefly describe details of known clinical characteristics of member and alternative drug regimen.
	Has the member previously tried the alternative drug required under the step therapy protocol, or another alternative drug in the same pharmacologic class or with the same mechanism of action, and such alternative drug was discontinued due to lack of efficacy or effectiveness, diminished effect, or an adverse event? Yes No If yes, please provide details for the previous trial.
	Drug name Dates/duration of use Did the member experience any of the following? Adverse reaction Inadequate response Briefly describe details of adverse reaction or inadequate response.
	Is the member stable on the requested prescription drug prescribed by the health care provider, and switching drugs will likely cause an adverse reaction in or physical or mental harm to the member?
]	☐ Yes. Please provide details.

Prior Authorization Request Prescriber and Provider Information

Prescriber Information		
Last name*	First name*	MI
NPI*	Individual MH Provide	er ID
DEA No.	Office Contact Name	
Address	City	State Zip
Email address		
Telephone No.*	Fax No.*	
* Required		
Please also complete for professionally	administered medication	ns, if applicable.
Start date	End date	
Servicing prescriber/facility name		☐ Same as prescribing provider
Servicing provider/facility address		
Servicing provider NPI/tax ID No.		
Name of billing provider		
Billing provider NPI No.		
Is this a request for recertification? Yes] No	
CPT code No. of visits	J code	No. of units
Prescribing provider's attestation, signal certify under the pains and penalties of perjoinformation section of this form. Any attached I certify that the medical necessity information complete, to the best of my knowledge. I under prosecution for any falsification, omission, or	ury that I am the prescribing I statement on my letterhead (per 130 CMR 450.204) on erstand that I may be subject concealment of any material	has been reviewed and signed by me. this form is true, accurate, and to civil penalties or criminal I fact contained herein.
Prescribing provider's signature		_
Printed name of prescribing provider (The form can either be signed by hand and		

(The form can either be signed by hand and then scanned, or it can be signed electronically using DocuSign or Adobe Sign. For electronic signatures, the signer can upload a picture of their wet signature. The typed text of a signature is not an acceptable form of an electronic signature.)