

Duexis® & Vimovo® Prior Authorization Request Form (Page 1 of 2)

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Member Information <small>(required)</small>			Provider Information <small>(required)</small>		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:		Specialty:
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:
Medication Information <small>(required)</small>					
Medication Name:			Strength:		Dosage Form:
<input type="checkbox"/> Check if requesting brand			Directions for Use:		
<input type="checkbox"/> Check if request is for continuation of therapy					
Clinical Information <small>(required)</small>					
Select the diagnosis below: <input type="checkbox"/> Ankylosing spondylitis [Vimovo only] <input type="checkbox"/> Osteoarthritis <input type="checkbox"/> Rheumatoid arthritis <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____					
Clinical information: Does the patient have a history of peptic ulcer disease/gastrointestinal (GI) bleed? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient have one additional risk factor for gastrointestinal adverse events (e.g., use of anticoagulants, chronic corticosteroids)? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient have a history of asthma or urticaria after taking aspirin or other NSAIDs? <input type="checkbox"/> Yes <input type="checkbox"/> No					
For Duexis requests, please also answer the following: Has the patient had a 30 day trial of a preferred generic H2-receptor blocker (e.g., famotidine, cimetidine, ranitidine, nizatidine) AND a generic NSAID within the last 180 days? <input type="checkbox"/> Yes <input type="checkbox"/> No					
For Vimovo requests, please also answer the following: Has the patient had a 30 day trial of a preferred generic proton pump inhibitor (e.g., omeprazole, lansoprazole, pantoprazole) AND a generic NSAID within the last 180 days? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Quantity limit requests: What is the quantity requested per DAY? _____					
What is the reason for exceeding the plan limitations? <input type="checkbox"/> Titration or loading dose purposes <input type="checkbox"/> Patient is on a dose-alternating schedule (e.g., one tablet in the morning and two tablets at night, one to two tablets at bedtime) <input type="checkbox"/> Requested strength/dose is not commercially available <input type="checkbox"/> Other: _____					

This document and others if attached contain information that is privileged, confidential and/or may contain protected health information (PHI). The Provider named above is required to safeguard PHI by applicable law. The information in this document is for the sole use of OptumRx. Proper consent to disclose PHI between these parties has been obtained. If you received this document by mistake, please know that sharing, copying, distributing or using information in this document is against the law. **If you are not the intended recipient, please notify the sender immediately.**

Office use only: Duexis-Vimovo_SouthDakotaMedicaid_2017May-P

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Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?

Please note: This request may be denied unless all required information is received.
For urgent or expedited requests please call 1-855-401-4262.
This form may be used for non-urgent requests and faxed to 1-844-403-1029.