

CONTAINS CONFIDENTIAL PATIENT INFORMATION
Ketorolac Tablets & Ketorolac Tromethamine Injection
Prior Authorization of Benefits (PAB) Form

Complete form in its entirety and fax to:
Prior Authorization of Benefits Center at 1-844-512-9004
Provider Help Desk 1-800-454-3730

1. PATIENT INFORMATION
2. PHYSICIAN INFORMATION

Patient Name: _____	Prescribing Physician: _____
Patient ID #: _____	Physician Address: _____
Patient DOB: _____	Physician Phone #: _____
Date of Rx: _____	Physician Fax #: _____
Patient Phone #: _____	Physician Specialty: _____
Patient Email Address: _____	Physician DEA: _____
	Physician NPI #: _____
	Physician Email Address: _____

3. MEDICATION
4. STRENGTH
5. DIRECTIONS
6. QUANTITY PER 30 DAYS

<input type="checkbox"/> Ketorolac Tablets	_____	_____	Specify: _____
<input type="checkbox"/> Ketorolac Tromethamine Injection	_____	_____	

7. DIAGNOSIS: _____

8. APPROVAL CRITERIA: CHECK ALL BOXES THAT APPLY

NOTE: Any areas not filled out are considered not applicable to your patient & MAY AFFECT THE OUTCOME of this request.

Requests for Ketorolac Tromethamine Injection:

- Yes No Documentation of previous trials and therapy failures with at least two preferred non-steroidal anti-inflammatory drugs* at therapeutic doses been provided
- Yes No Patient has a diagnosis indicating moderately severe, acute pain

*The preferred non-steroidal anti-inflammatory drugs are: diclofenac potassium, diclofenac sodium, etodolac tab 400 mg, etodolac tab 500 mg, flurbiprofen, Ibuprofen Susp 100 MG/5ML (OTC), Ibuprofen Tab 200 MG (OTC), ibuprofen tab 400 mg, ibuprofen tab 600 mg, ibuprofen tab 800 mg, indomethacin, ketoprofen, NAPROSYN SUSP, naproxen, naproxen sodium tab 550 mg, sulindac.

Requests for Ketorolac Tablets:

- Yes No Documentation of recent IM/IV ketorolac tromethamine injection including administration date and time, and the total number of injections given is provided
- Yes No Patient has a diagnosis indicating moderately severe, acute pain

Please Note: Documentation may include, but is not limited to, chart notes, prescription claims records, prescription receipts, and laboratory data.

Dosing guidelines: maximum oral dose is 40 mg/day; maximum IV/IM dose is 120 mg/day; combined duration of use of IV/IM and oral is not to exceed five (5) days.



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9. PHYSICIAN SIGNATURE

_____ Prescriber or Authorized Signature	_____ Date
<small><i>Prior Authorization of Benefits is not the practice of medicine or the substitute for the independent medical judgment of a treating physician. Only a treating physician can determine what medications are appropriate for a patient. Please refer to the applicable plan for the detailed information regarding benefits, conditions, limitations, and exclusions. The submitting provider certifies that the information provided is true, accurate, and complete and the requested services are medically indicated and necessary to the health of the patient.</i></small> <small><i>Note: Payment is subject to member eligibility. Authorization does not guarantee payment.</i></small>	
<small>The document(s) accompanying this transmission may contain confidential health information that is legally privileged. This information is intended only for the use of the individual or entity named above. The authorized recipient of this information is prohibited from disclosing this information to any other party unless required to do so by law or regulation. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution, or action taken in reliance on the contents of these documents is strictly prohibited. If you have received this information in error, please notify the sender immediately and arrange for the return or destruction of these documents.</small>	