

	Authorization Form		
•	to: 877.974.4411 toll free, o		Individual (Ontimized)
	<ul><li>☐ Commercial (Traditional)</li><li>☒ Medicaid</li></ul>	Commercial	individuai (Optimized)
This request is:	☐ <b>Urgent</b> (life threatening) ☐	Non-Urgent (standar	d review)
	Urgent means the standard review time material to regain maximum function.	ay seriously jeopardize the life	e or health of the patient or the patient's ability
	_		
Naproxen s	uspension		
Member			
Last Name:		First Name:	
		DOB:	Gender:
Primary Care Physician:			
Requesting Provider:		Prov. Phone:	Prov. Fax:
Provider Address:			
Provider NPI:		Contact Name:	
Provider Signature:		Date:	
Product Information			
☐ New request ☐ Con	tinuation request		
☐ New request ☐ Con	unuation request		
Drug product: 🔲 Naproxe	en 125 mg/5 mL suspension	•	f next dose):
			applicable):
		Dosing frequency:	
Precertification Requir	ements		
Before this drug is covered,	the patient must meet all of the fol	lowing requirements:	
Must be age 12 years a	nd vounger		
<ol> <li>Must be age 12 years a</li> <li>Must be prescribed by a</li> </ol>			
	gnosis of: Rheumatoid Arthritis, Os	steoarthritis, Ankylosing	Spondylitis, or Juvenile Rheumatoid
Arthritis.  4. Must have had a trial ar	nd failure of ibuprofen suspension	or a clinical reason ibup	profen suspension cannot be used.
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	s, dosing, or a route of administration not a ex, AHFS, U.S. Pharmacopeia, and also C		ug Administration (FDA) or recognized in CMS- plogy indications only) require supporting
	ovide two published peer-reviewed literatu e used for the identified indication.	re articles supporting the appr	ropriateness of the drug, the dosing of the drug,
New request Priority Health Precerti	ification Documentation		
•			
A. What condition is this	s drug being requested for?		
B. Is prescriber a rheum	atologist?		
☐ Yes ☐ No			



C.	Has patient had a trial and failure of ibuprofen suspension?  Yes, list dose tried, dates, and outcomes:
	□ No, clinical rationale: