

LEUKOTRIENE MODIFIERS PRIOR AUTHORIZATION FORM

- Please submit all requested documentation with this form. Incomplete documentation may delay the processing of this request.
- Prior authorization guidelines for **Leukotriene Modifiers** and **Quantity Limits/Daily Dose Limits** are available on the DHS Pharmacy Services website at <http://www.dhs.pa.gov/provider/pharmacyservices/drugsrequiringclinicalpriorauthorization>.

PRIOR AUTHORIZATION REQUEST INFORMATION		PRESCRIBER INFORMATION	
<input type="checkbox"/> New request	<input type="checkbox"/> Renewal request	total # of pgs: _____	Prescriber name:
Name of office contact:		Specialty:	
Contact's phone number:		State license #:	
LTC facility contact/phone:		NPI:	MA Provider ID#:
RECIPIENT INFORMATION		Street address:	
Recipient Name:		Suite #:	City/state/zip:
Recipient ID#:	DOB:	Phone:	Fax:

CLINICAL INFORMATION

Non-preferred medication requested:	<input type="checkbox"/> Accolate tablet*	<input type="checkbox"/> Singulair tablet*	<input type="checkbox"/> zileuton ER tablet
	<input type="checkbox"/> montelukast granule	<input type="checkbox"/> Singulair chewable*	<input type="checkbox"/> Zyflo tablet*
	<input type="checkbox"/> Singulair granule*	<input type="checkbox"/> zafirlukast tablet	<input type="checkbox"/> Zyflo CR tablet
Strength:	Directions:	Quantity:	Refills:
Diagnosis:		Dx code (required):	
1. Did the Recipient try and fail the preferred Leukotriene Modifier, montelukast tablet or chewable tablet ?		<input type="checkbox"/> Yes – <i>Submit all supporting documentation of drug regimen and therapeutic failure.</i> <input type="checkbox"/> No	
2. Does the Recipient have any contraindications or intolerances to the preferred agent listed in question (1)?		<input type="checkbox"/> Yes – <i>Submit all supporting documentation of medication name(s) and associated intolerances / contraindications.</i> <input type="checkbox"/> No	
3. For non-preferred brand name products with available generics (marked with a * in the above non-preferred medication list), why can't the Recipient take the FDA-approved generic equivalent product? <u>Include reason in space below and submit medical record documentation supporting the brand medically necessary request.</u>			

PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION TO DHS – PHARMACY DIVISION

Prescriber Signature:	Date:
------------------------------	--------------

Confidentiality Notice: The documents accompanying this telecopy may contain confidential information belonging to the sender. The information is intended only for the use of the individual named above. If you are not the intended recipient, you are hereby notified that any disclosure, copying, distribution or taking of any telecopy is strictly prohibited.