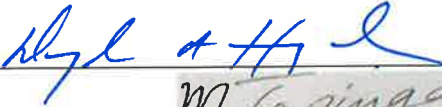





KERN HEALTH SYSTEMS

Policy and Procedure Review/Revision

Policy 13.01-P Drug Utilization and Non-Formulary Treatment Request has been updated and is provided here for your review and approval.

Reviewer	Date	Comment/Signature
Doug Hayward	7/23/21	
Dr. Tasinga	7/21/2021	
Bruce Wearda	7/19/2021	Bruce Wearda

(CEO decision(s))

Board approval required: Yes ___ No ___ QI/UM Committee approval: Yes ___ No ___
Date approved by the KHS BOD: _____ Date of approved by QI: _____
PAC approval: Yes ___ No ___ Date of approval by PAC: _____
Approval for internal implementation: Yes ___ No ___
Provider distribution date: Immediately _____ Quarterly _____

Effective date: _____
DHCS submission: _____
DMHC submission: _____
Provider distribution: _____



KERN HEALTH SYSTEMS					
POLICY AND PROCEDURES					
SUBJECT: Drug Utilization and Non-Formulary Treatment Request				POLICY #: 13.01-P	
DEPARTMENT: Pharmacy					
Effective Date: 08/1997	Review/Revised Date: 07/23/2021	DMHC		PAC	
		DHCS		QI/UM COMMITTEE	
		BOD		FINANCE COMMITTEE	

Douglas A. Hayward
Chief Executive Officer

Date _____

Chief Medical Officer

Date _____

Director of Pharmacy

Date _____

POLICY:

The following applies to pharmacy authorization requests that will be billed on a pharmacy NCPDP claim prior to the launch of Medi-Cal Rx and those medical supplies and devices remaining with the managed care plans outlined by the Medi-Cal Rx Scope document. All NCPDP pharmacy requests for claims for date of service after the launch of Medi-Cal Rx will be directed to Medi-Cal Rx for review. Until that time, these policies and protocols effectuated by Kern Health Systems will remain unchanged and in place. Institutional and professional claims will continue to be processed by the managed care plan and therefore the requests would follow these procedures outlined. Those medical supplies and devices will be reviewed as stated in this policy and the encounters submitted on an 837P file.

All non-formulary medications or formulas require prior authorization. All medically necessary outpatient prescription drugs, except for those specifically excluded from the Medi-Cal contract, shall be available to KHS Medi-Cal members.¹ This determination will be made through the non-formulary treatment request process as outlined in this policy and procedure.

The non-formulary treatment request process will conform to the requirements outlined in the

following statutory, regulatory, and contractual sources:

- Code of Federal Regulations Title 42 §§431.211; 431.213; and 431.214
- California Health and Safety Code §§ 1367.01²; 1367.21; 1367.22; 1367.24
- California Welfare and Institutions Code §14185
- CCR Title 28 §1300.67.24
- CCR Title 22 §§ 51003; 51014.1; 51014.2; 53854; 53894
- DHS Contract Exhibit A - Attachment 5 (3)(F); Exhibit A –Attachment 10 (7)(F)
- DHS MMCD Letters 04006 (November 1, 2004) and 05005 (April 11, 2005), and 08-013 (December 16, 2008).

This document shall be disclosed to the public upon request.³

DEFINITIONS:

Chronic and Seriously Debilitating⁴	Diseases or conditions that require ongoing treatment to maintain remission or prevent deterioration and cause significant long-term morbidity.
Life Threatening⁵	Diseases or conditions (1) where the likelihood of death is high unless the course of the disease is interrupted and/or (2) with potentially fatal outcomes where the endpoint of clinical intervention is survival.

PROCEDURES:

1.0 SUBMISSION OF A NON-FORMULARY DRUG REQUEST

Non-Formulary drug requests can be made by KHS providers electronically via the secure KHS portal or on a *61-211 Form* if one does not have access to the KHS Provider Portal. (See Attachment A). Form should be mailed/faxed to the following location:

KHS Pharmacy Department
2900 Buck Owens Boulevard
Bakersfield, CA 93308
661-664-5191

KHS only requests information reasonably necessary to make a decision regarding the request.⁶ Documentation must be complete and include:

- A. Patient name.
- B. CIN number.
- C. Diagnosis with brief history.
- D. Reason for request/justification including formulary medication failures.
- E. Drug name, strength, directions, and National Drug Code.
- F. Prescriber's name.

2.0 REVIEW OF TAR

Incoming requests are date and time stamped. TAR review includes the actions outlined in

the following table.

Action	Timeline	Comments
Review by Pharmacist or MD		Evaluation for medical necessity denials signed by licensed pharmacist or MD ⁷
Decision (approve or deny)	Within 24 hours of receipt ⁸ .	

Medications and supplies are evaluated on the basis of appropriateness, efficacy, safety, pharmacokinetics and cost effectiveness.

3.0 PRACTITIONER/PROVIDER AND MEMBER NOTIFICATION

Results of the TAR review are communicated by Pharmacy staff to the practitioner/provider and member as outlined in the following table. Notification to providers is provided via portal or facsimile if possible. The notification confirmation is attached to the request. If notice by electronic portal or facsimile is not possible, verbal notice is provided via phone within 24 hours of receipt. In such cases, written notice follows as outlined in the table below.

Result of Review	Practitioner/Provider Notice	Member Notice
Approved	Approved form ⁹ (within 24 hours of receipt). ¹⁰	
Denied	Denied form (within 24 hours of receipt). ¹¹	<i>Notice of Action Documents</i> (within 2 business days of the decision). ¹² Documents include all of the following: ❖ <i>Notice of Action – Denial</i> (Attachments B- D) <i>Your Rights Under Medi-Cal Managed Care & Form to File a State Hearing.</i> Medi-Cal members only.

Notice of Action letters together with the indicated enclosures contain all of the required elements for both provider and member notice of delay, denial, or modification including the following¹³:

- A. The action taken.
- B. A clear and concise explanation of the reason for the decision (including clinical reasons for decisions regarding medical necessity).
- C. A description of the criteria/guidelines used.
- D. A citation of the specific regulations or plan authorization procedures supporting the action¹⁴.
- E. Information on how to file a grievance with KHS including the Plan's name address and phone number.

- F. Information regarding a Medi-Cal member's right to a State Fair Hearing including:
 - 1. The method by which a hearing may be obtained.
 - 2. That the member may either be self- represented or represented by an authorized third party such as legal counsel, relative, friend, or any other person.
 - 3. The time limit for requesting a fair hearing.
 - 4. The toll-free number for obtaining information on legal service organizations for representation.
- G. Nondiscrimination Notice.
- H. Language Assistance Taglines.
- I. Information regarding the member's right to an Independent Medical Review with DMHC.
- J. DMHC required language regarding grievances¹⁵.
- K. Name and telephone number of the pharmacy department.

4.0 DOCUMENTATION

Letters regarding authorization requests, including those sent by KHS to both members and providers, are retained as outlined in *KHS Policy and Procedure #10.51 – Records Retention*.¹⁶

5.0 ALLOWED SUPPLIES OF MEDICATION

Members may receive up to a 30 day supply of medication. Women may receive up to 365 day supply of hormonal contraceptives.

5.1 Emergency Supplies¹⁷

During weekends, holidays and non-business hours a pharmacy may choose to dispense enough medication (72 hours supply maximum) as an emergency supply to the member until the next working day, at the dispensing pharmacist's discretion according to pharmacy policy and procedures as defined by Title 22 section 51056¹⁸. If the medication is not on the Plan Formulary, a request must be submitted for payment processing stating the emergency and relevant clinical information about the member's condition and why they were considered immediately necessary and medication dispensed. A mere statement that an emergency existed is not sufficient. It must be comprehensive enough to support a finding that an emergency existed.¹⁹ TAR approval is not needed for reimbursement before dispensing of 72 hour emergency supply of non-formulary drugs.

6.0 CONTINUITY OF CARE²⁰

Medi-Cal members are allowed continued coverage of a non-formulary single source drug which is part of a prescribed therapy previously approved for coverage by the plan for a medical condition of the member and the provider continues to prescribe the drug for the medical condition provided that drug is appropriately prescribed and is considered safe and effective for treating the member's medical condition. Nothing in this section shall preclude the prescribing provider from prescribing another drug covered by the plan that is medically appropriate²¹ If previously approved by plan immediately prior to the date of enrollment, coverage may be continued until the prescribed therapy is no longer prescribed by a contracting practitioner²². Approval is contingent upon documentation that the patient had

authorization from the previous plan of the medication at the time of enrollment no more than fifteen (15) days beyond the estimated day supply for the last documented pharmacy fill date.²³

KHS does not require a new member to repeat step therapy when the member is already being treated for a medical condition by a prescription drug provided that the drug is appropriately prescribed and is considered safe and effective for the enrollee's condition. For purposes of this section, "step therapy" means a type of protocol that specifies the sequence in which different prescription drugs for a given medical condition and medically appropriate for a particular patient are to be prescribed.

Medi-Cal members are allowed continued coverage of a drug which is removed from the KHS formulary if the drug is part of a prescribed therapy in effect immediately prior to the date of removal until the prescribed therapy is no longer prescribed by a contracting practitioner.

7.0 BRAND NAME MEDICATIONS WHEN EQUIVALENT GENERIC BRAND IS AVAILABLE

If a medication is available or becomes available in an AB rated generic brand, the brand name version will become non-Formulary for KHS.

Unless it is determined to be medically necessary for the patient to continue using the brand name, if a generic brand becomes available during a patient's treatment, the patient will be expected to switch to the generic brand and must fail the generic brand prior to KHS granting authorization for the brand name.

Providers with patients having untoward effects from a generic brand must submit a completed FDA *MedWatch* form to KHS as part of the request for authorization to allow a brand name version instead of a generic brand. (See Attachment F).

Biosimilars and drugs considered as Follow Ons will be treated in the same fashion as if they were a traditional generic of the innovator drug. Per FDA rules, they are not automatically substitutable, but from clinical perspectives they are viewed as a generic version.

8.0 OFF-LABEL USE FOR LIFE THREATENING OR CHRONIC AND SERIOUS CONDITIONS²⁴

8.1 Medi-Cal Product

Section does not apply to the Medi-Cal product.²⁵

8.2 Peer Reviewed Professional Society Endorsed Supporting Documentation If a physician or other provider wishes to prescribe a non-formulary or restricted FDA approved medication for an off-label use for a life threatening or chronic and debilitating condition, he/she may submit a referral or TAR to the Plan for the same. In the referral, the provider must demonstrate the medication is recognized for the treatment of that condition in one of the following sources:

- A. American Hospital Formulary Service's Drug Information.
- B. One of the following compendia, if recognized by the federal Centers for Medicare and Medicaid Services as part of an anticancer

chemotherapeutic regimen:

- (1) The Elsevier Gold Standard's Clinical Pharmacology.
 - (2) The National Comprehensive Cancer Network Drug and Biologics Compendium.
 - (3) The Thomson Micromedex DrugDex.
- C. Two articles from major peer reviewed medical journals that present data supporting the proposed off-label use or uses as generally safe and effective unless there is clear and convincing contradictory evidence presented in a major peer reviewed medical journal.

The provider is responsible for submitting the required documentation to KHS²⁶.

9.0 SAMPLE MEDICATIONS²⁷

Providers are discouraged from providing samples; however, if samples are given to the member, the entire course of therapy must be covered by the samples. Medications provided as samples do not establish a continuity precedent or satisfy step therapy criteria and, therefore, do not obligate coverage by KHS. If providing samples, Providers shall follow the outlined steps in *KHS Policy and Procedure #2.24-P Pharmacy Guidelines*.

10.0 TRIAL PERIOD²⁸

Barring any medically adverse responses from the member, the trial period of a medication shall be determined per the recommended dosing titration guidelines presented to the FDA.

11.0 MONITORING²⁹

The Compliance Department will conduct bi-annual audits to monitor compliance of the contracted emergency departments to provide a sufficient quantity of drugs to Medi-Cal members under emergency circumstances to last until the member can reasonably be expected to have a prescription filled prior to leaving the emergency department. Issues discovered by this monitoring will be brought to the attention of the contracted emergency department and a Corrective Action Plan (CAP) will be required.

ATTACHMENTS:

- ❖ Attachment A - Treatment Authorization Request (TAR) Form (61-211)
- ❖ Attachment B - Notice of Action – Denial letter, Deny – Criteria Not Met
- ❖ Attachment C - Notice of Action – Denial letter, Deny – Incorrect Form
- ❖ Attachment D - Notice of Action – Denial letter, Deny – Member Not Eligible
- ❖ Attachment E - Your Rights Under Medi-Cal Managed Care, How to File a State Hearing, Independent Medical Review, Nondiscrimination and Language Assistance Taglines
- ❖ Attachment F - MedWatch form

REFERENCE:

¹**Revision 2021-07:** Policy revised by Director of Pharmacy regarding changes in pharmacy authorizations and billing. **Revision 2020-02:** **Revision 2017-07:** Policy revised to comply with CMS Final Rule on prior authorization process. Attachments updated. **Revision 2017-03:** Policy reviewed and updated by Director of Pharmacy. New Section 8.2 provides guidelines for prescribing medication for an off-label use for a life threatening or chronic and debilitating condition. **Revision 2014-10:**

Formatting changes to policy, no material changes. Notice of Action letters (NOAs) revised as a result of the DHCS 2013 Medical Audit ending in 2014- CAF-9. “Your Right’s Forms” updated to ensure continued compliance. Translation changes made to comply with MMCD APL 05005. **Revision 2014-04:** Language included in Section 1.0 to add time statement on authorization request. Revised to remove references to Health Families product. **Revision 2013-07:** Reviewed by Director of Pharmacy. Routine revision, updated Section 1.0 regarding submission of treatment authorization request. **Revision 2009-10:** Revision requested by Director of Pharmacy. **Revision 2009-02:** Revised to comply with MMCD Policy Letter 08-013. Notice of Action letters updated with language assistance services notice. **Revision 2007-05:** Revised per DHS/DMHC Medical Audit comment 5/13/2007. **Revision 2007-04:** Created Notice of Action Letters for Healthy Families product line per DHS/DMHC Medical Review Audit (YE 10/31/06). **Revision 2005-07:** Reviewed against MMCD Letters 04006 and 05005. New NOAs. **Revision 2005-04:** Continuity of care processes reviewed and revised. Reviewed against DHS Contract 03-76165 (Effective May 1, 2004). **Revision 2004-05:** Revised per DMHC/DHS Medical Audit YE Oct03; finding 1.2. (Addition of member notice of modifications). New single letter for deferral, modification, or denial. **Revision 2002-03:** Revised per DHS comment 01/30/02. **Revision 2002-01:** DHS CAP Verification Visit Report (Med Rev YE 08/00). **Revision 2001-02:** changes made for 2000 Legislation submission – DMHC and DHS/DMHC Medical Review Audit (YE 08/31/00).

¹ HSC 1367.24. CCR Title 22 §53854(d), CCR Title 28 1300.67.24

² Applicable to pharmacy per Title 28 §1300.67.24(a)(1)

³ HSC §1367.01(b)

⁴ HSC §1367.21(e)

⁵ HSC §1367.21(d)

⁶ HSC §1367.01(g)

⁷ MMCD Policy Letter 08-013

⁸ Social Security Act, Title XIX, Section 1927(a)(5). Per John Kaylen at DHS, allows an extension from 24 hours to 1 business day. Although action must be taken on pain medications for the terminally ill within 72 hours, the Plan must follow the stricter Federal Medicaid 24 hours standard. HSC Section 1367.215. DHS Contract A-5 3(F). Change to comply with CMS Managed Care final Rule Prior Authorization Process (§438.3(s)(6))

⁹ Must include specific service approved (HSC §1367.01(h)(4))

¹⁰ Social Security Act, Title XIX, Section 1927(a)(5). Per John Kaylen at DHS, allows an extension from 24 hours to 1 business day. Although action must be taken on pain medications for the terminally ill within 72 hours, the Plan must follow the stricter Federal Medicaid 24 hours standard. HSC Section 1367.215. HSC §1367.01(h)(3) requires notice to provider within 24 hours of the decision.

¹¹ Social Security Act, Title XIX, Section 1927(a)(5). Per John Kaylen at DHS, allows an extension from 24 hours to 1 business day. Although action must be taken on pain medications for the terminally ill within 72 hours, the Plan must follow the stricter Federal Medicaid 24 hours standard. HSC Section 1367.215. HSC §1367.01(h)(3) requires notice to provider within 24 hours of the decision. Change to comply with CMS Managed Care final Rule Prior Authorization Process (§438.3(s)(6))

¹² HSC §1367.01(h)(3) has the shortest time period for member notice (2 b/days of decision). CCR Title 22 Section 53894(a) and (d)

¹³ HSC §1367.01(h)(4) and (5) and 1367.24(b); CCR Title 22 §53894

¹⁴ Required for member notice only. CCR Title 22 §53894(d)(3)

¹⁵ Required for member notice only. HSC §1367.24(b)

¹⁶ DHS Contract 03-76165 Exhibit A – Attachment 5 (2)(G)

¹⁷ CCR Title 22 §53854(2)

¹⁸ Title 22 51056

¹⁹ Title 22 51056

²⁰ HSC 1367.22

²¹ HSC § 1367.22

²² W&I Code 14185(c)

²³ HSC §1367.22

²⁴ Health and Safety Code §1367.21

²⁵ Plan shall reserve the right to modify this. ²⁶ HSC 1367.21 (b)

²⁷ Section added upon request of the Director of Pharmacy (3/2/05). Language also included in *Policy 2.24 – Pharmaceutical Guidelines*

²⁸ Section added upon request of the Director of Pharmacy (3/2/05).

²⁹ DHS/DMHC Medical Review Audit (YE 10/31/06).

PRESCRIPTION DRUG PRIOR AUTHORIZATION OR STEP THERAPY EXCEPTION REQUEST FORM

Plan/Medical Group Name: _____ Plan/Medical Group Phone#: (_____) _____
 Plan/Medical Group Fax#: (_____) _____ Non-Urgent ☐ Exigent Circumstances ☐

Instructions: Please fill out all applicable sections on both pages completely and legibly. Attach any additional documentation that is important for the review, e.g. chart notes or lab data, to support the prior authorization or step-therapy exception_request. **Information contained in this form is Protected Health Information under HIPAA.**

Patient Information

First Name:	Last Name:	MI:	Phone Number:
Address:		City:	State: Zip Code:
Date of Birth:	<input type="checkbox"/> Male <input type="checkbox"/> Female	Circle unit of measure Height (in/cm): _____ Weight (lb/kg): _____	Allergies:
Patient's Authorized Representative (if applicable):		Authorized Representative Phone Number:	

Insurance Information

Primary Insurance Name:	Patient ID Number:
Secondary Insurance Name:	Patient ID Number:

Prescriber Information

First Name:	Last Name:	Specialty:
Address:		City: State: Zip Code:
Requestor (if different than prescriber):		Office Contact Person:
NPI Number (individual):		Phone Number:
DEA Number (if required):		Fax Number (in HIPAA compliant area):
Email Address:		

Medication / Medical and Dispensing Information

Medication Name:			
<input type="checkbox"/> New Therapy <input type="checkbox"/> Renewal <input type="checkbox"/> Step Therapy Exception Request If Renewal: Date Therapy Initiated: _____ Duration of Therapy (specific dates): _____			
How did the patient receive the medication?			
<input type="checkbox"/> Paid under Insurance Name: _____ Prior Auth Number (if known): _____ <input type="checkbox"/> Other (explain): _____			
Dose/Strength:	Frequency:	Length of Therapy/#Refills:	Quantity:
Administration:			
<input type="checkbox"/> Oral/SL <input type="checkbox"/> Topical <input type="checkbox"/> Injection <input type="checkbox"/> IV <input type="checkbox"/> Other: _____			
Administration Location:		<input type="checkbox"/> Patient's Home <input type="checkbox"/> Long Term Care <input type="checkbox"/> Physician's Office <input type="checkbox"/> Home Care Agency <input type="checkbox"/> Other (explain): _____ <input type="checkbox"/> Ambulatory Infusion Center <input type="checkbox"/> Outpatient Hospital Care _____	

PRESCRIPTION DRUG PRIOR AUTHORIZATION OR STEP THERAPY EXCEPTION REQUEST FORM

Patient Name:

ID#:

Instructions: Please fill out all applicable sections on both pages completely and legibly. Attach any additional documentation that is important for the review, e.g. chart notes or lab data, to support the prior authorization or step therapy exception request.

1. Has the patient tried any other medications for this condition? ☐ YES (if yes, complete below) ☐ NO

Medication/Therapy
(Specify Drug Name and Dosage)

Duration of Therapy
(Specify Dates)

Response/Reason for Failure/Allergy

2. List Diagnoses:

ICD-10:

3. Required clinical information - Please provide all relevant clinical information to support a prior authorization or step therapy exception request review.

Please provide symptoms, lab results with dates and/or justification for initial or ongoing therapy or increased dose and if patient has any contraindications for the health plan/insurer preferred drug. Lab results with dates must be provided if needed to establish diagnosis, or evaluate response. Please provide any additional clinical information or comments pertinent to this request for coverage, including information related to exigent circumstances, or required under state and federal laws.

☐ Attachments

Attestation: I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan, insurer, Medical Group or its designees may perform a routine audit and request the medical information necessary to verify the accuracy of the information reported on this form.

Prescriber Signature or Electronic I.D. Verification: _____ **Date:** _____

Confidentiality Notice: The documents accompanying this transmission contain confidential health information that is legally privileged. 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 (via return FAX) and arrange for the return or destruction of these documents.

Plan/Insurer Use Only: Date/Time Request Received by Plan/Insurer: _____ Date/Time of Decision _____

Fax Number (_____) _____

☐ Approved ☐ Denied Comments/Information Requested: _____



NOTICE OF ACTION
About Your Treatment Request - Deny

Today's date

Member name
 Address
 City, State, Zip

Treating Provider
 Address
 City, State, Zip

Identification Number: KHS ID

RE: [Service requested]

[Name of requesting provider] has asked Kern Family Health Care to approve [Service requested]. This request is denied. This is because [Insert: 1. A clear and concise explanation of the reasons for the decision; 2. A description of the criteria or guidelines used, including a reference to the specific regulations or plan authorization procedures that support the action; and. 3. The clinical reasons for the decision regarding medical necessity]. Based on documentation provided, criteria was not met

You may ask for free copies of all information used to make this decision. This includes a copy of the actual benefit provision, guideline, protocol, or criteria that we based our decision on. To ask for this, please call Kern Family Health Care at (661) 632-1590 inside Bakersfield, or (800) 391-2000 outside of Bakersfield.

You may appeal this decision. The enclosed "Your Rights" information notice tells you how. It also tells you where you can get free help. This also means free legal help. You are encouraged to send in any information that could help your case. The "Your Rights" notice tells you the cut off dates to ask for an appeal.

The State Medi-Cal Managed Care "Ombudsman Office" can help you with any questions. You may call them at 1-888-452-8609. You may also get help from your doctor, or call us at (661) 632-1590 inside Bakersfield, or (800) 391-2000 outside of Bakersfield.

This notice does not affect any of your other Medi-Cal services.

Medical Directors Name

Enclosed: "Your Rights under Medi-Cal Managed Care"



**NOTICE OF ACTION
About Your Treatment Request - Deny**

Today's date

Member name
Address
City, State, Zip

Treating Provider
Address
City, State, Zip

Identification Number: KHS ID

RE: [Service requested]

[Name of requesting provider] has asked Kern Family Health Care to approve [Service requested]. This request is denied. This is because the request was not submitted on the State Mandated 61-211 from.

You may ask for free copies of all information used to make this decision. This includes a copy of the actual benefit provision, guideline, protocol, or criteria that we based our decision on. To ask for this, please call Kern Family Health Care at (661) 632-1590 inside Bakersfield, or (800) 391-2000 outside of Bakersfield.

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This notice does not affect any of your other Medi-Cal services.

Medical Directors Name

Enclosed: "Your Rights under Medi-Cal Managed Care"



**NOTICE OF ACTION
About Your Treatment Request - Deny**

Today's date

Member name
Address
City, State, Zip

Treating Provider
Address
City, State, Zip

Identification Number: KHS ID

RE: [Service requested]

[Name of requesting provider] has asked Kern Family Health Care to approve [Service requested]. This request is denied. This is because no eligibility is recorded for the date of service.

You may ask for free copies of all information used to make this decision. This includes a copy of the actual benefit provision, guideline, protocol, or criteria that we based our decision on. To ask for this, please call Kern Family Health Care at (661) 632-1590 inside Bakersfield, or (800) 391-2000 outside of Bakersfield.

You may appeal this decision. The enclosed "Your Rights" information notice tells you how. It also tells you where you can get free help. This also means free legal help. You are encouraged to send in any information that could help your case. The "Your Rights" notice tells you the cut off dates to ask for an appeal.

The State Medi-Cal Managed Care "Ombudsman Office" can help you with any questions. You may call them at 1-888-452-8609. You may also get help from your doctor, or call us at (661) 632-1590 inside Bakersfield, or (800) 391-2000 outside of Bakersfield.

This notice does not affect any of your other Medi-Cal services.

Medical Directors Name

Enclosed: "Your Rights under Medi-Cal Managed Care"

YOUR RIGHTS UNDER MEDI-CAL MANAGED CARE

IF YOU DO NOT AGREE WITH THE DECISION MADE FOR YOUR MEDICAL TREATMENT, YOU CAN FILE AN APPEAL. THIS APPEAL IS FILED WITH YOUR HEALTH PLAN.

HOW TO FILE AN APPEAL

You have **60 days** from the date of this “Notice of Action” letter to file an appeal. But, **if you are currently getting treatment and you want to continue getting treatment, you must ask for an appeal within 10 days** from the date this letter was postmarked or delivered to you, OR before the date your health plan says services will stop. You must say that you want to keep getting treatment when you file the appeal.

You can file an appeal by phone, in writing, or electronically:

- **By phone:** Contact Kern Family Health Care between 8:00 a.m. to 5:00 p.m. by calling (661) 632-1590 inside Bakersfield, or 1-800-391-2000 outside Bakersfield. Or, if you cannot hear or speak well, please call 711.
- **In writing:** Fill out an appeal form or write a letter and send it to:

Kern Family Health Care
2900 Buck Owens Boulevard
Bakersfield, CA 93308

Your doctor’s office will have appeal forms available. Your health plan can also send a form to you.

- **Electronically:** Visit your health plan’s website. Go to <http://www.kernfamilyhealthcare.com>

You may file an appeal yourself. Or, you can have a relative, friend, advocate, doctor, or attorney file the appeal for you. You can send in any type of information you want your health plan to review. A doctor who is different from the doctor who made the first decision will look at your appeal.

Your health plan has 30 days to give you an answer. At that time, you will get a “Notice of Appeal Resolution” letter. This letter will tell you what the health plan has decided. **If you do not get a letter within 30 days, you can:**

- Ask for an “**Independent Medical Review**” (**IMR**) and an outside reviewer that is not related to the health plan will review your case.
- Ask for a “**State Hearing**” and a judge will review your case

Please read the section below for instructions on how to ask for an IMR or State Hearing.

EXPEDITED APPEALS

If you think waiting 30 days will hurt your health, you might be able to get an answer within 72 hours. When filing your appeal, say why waiting will hurt your health. Make sure you ask for an **“expedited appeal.”**

IF YOU DO NOT AGREE WITH THE APPEAL DECISION

If you filed an appeal and received a “Notice of Appeal Resolution” letter telling you that your health plan will still not provide the services, or **you never received a letter telling you of the decision and it has been past 30 days**, you can:

- Ask for an **“Independent Medical Review” (IMR)** and an outside reviewer that is not related to the health plan will review your case
- Ask for a **“State Hearing”** and a judge will review your case

You can ask for both an IMR and State Hearing at the same time. You can also ask for one before the other to see if it will resolve your problem first. For example, if you ask for an IMR first, but do not agree with the decision, you can still ask for a State Hearing later. However, if you ask for a State Hearing first, but the hearing has already taken place, you cannot ask for an IMR. In this case, the State Hearing has the final say.

You will not have to pay for an IMR or State Hearing.

INDEPENDENT MEDICAL REVIEW (IMR)

If you want an IMR, you must first file an appeal with your health plan. If you do not hear from your health plan within 30 days, or if you are unhappy with your health plan's decision, then you may then request an IMR. You must ask for an IMR within **180 days** from the date of the “Notice of Appeal Resolution” letter.

You may be able to get an IMR right away without filing an appeal first. This is in cases where your health is in immediate danger or the request was denied because treatment is considered experimental or investigational.

The paragraph below will provide you with information on how to request an IMR. Note that the term “grievance” is talking about both “complaints” and “appeals.”

The California Department of Managed Health Care is responsible for regulating health care service plans. If you have a grievance against your health plan, you should first telephone your health plan at 1-800-391-2000 and use your health plan's grievance process before contacting the Department. Utilizing this grievance procedure does not prohibit any potential legal rights or remedies that may be available to you. If you need help with a grievance involving an emergency, a grievance that has not been satisfactorily resolved by your health plan, or a grievance that has remained unresolved for more than 30 days, you may call the Department for assistance. You may also be eligible for an Independent Medical Review (IMR). If you are eligible for an IMR, the IMR process will provide an impartial review of medical decisions made by a health plan related to the medical necessity of a proposed service or treatment, coverage decisions for treatments that are experimental or investigational in nature and payment disputes for emergency or urgent medical services. The Department also has a toll-free telephone number **(1-888-HMO-2219)** and a TDD line **(1-877-688-9891)** for the hearing and speech impaired. The Department's Internet Website (<http://www.hmohelp.ca.gov>) has complaint forms, IMR application forms, and instructions online.

STATE HEARING

If you want a State Hearing, you must ask for one within **120 days** from the date of the "Notice of Appeal Resolution" letter. You can ask for a State Hearing by phone or in writing:

- **By phone:** Call **1-800-952-5253**. This number can be very busy. You may get a message to call back later. If you cannot speak or hear well, please call **TTY/TDD 1-800-952-8349**.
- **In writing:** Fill out a State Hearing form or send a letter to:

**California Department of Social Services
State Hearings Division
P.O. Box 944243, Mail Station 9-17-37
Sacramento, CA 94244-2430**

Be sure to include your name, address, telephone number, Social Security Number, and the reason you want a State Hearing. If someone is helping you ask for a State Hearing, add their name, address, and telephone number to the form or letter. If you need an interpreter, tell us what language you speak. You will not have to pay for an interpreter. We will get you one.

After you ask for a State Hearing, it could take up to 90 days to decide your case and send you an answer. If you think waiting that long will hurt your health, you might be able to get an answer within 3 working days. Ask your doctor or health plan to write a letter for you. The letter must explain in detail how waiting for up to 90 days for your case to be decided will seriously harm your life, your health, or your ability to attain,

maintain, or regain maximum function. Then, make sure you ask for an **“expedited hearing”** and provide the letter with your request for a hearing.

You may speak at the State Hearing yourself. Or, you can have a relative, friend, advocate, doctor, or attorney speak for you. If you want another person to speak for you, then you must tell the State Hearing office that the person is allowed to speak on your behalf. This person is called an “authorized representative.”

LEGAL HELP

You may be able to get free legal help. Call the Greater Bakersfield Legal Assistance at (661) 325-5943. You may also call the local Legal Aid Society in your county at 1-888-804-3536.

**SUS DERECHOS
BAJO ATENCIÓN ADMINISTRADA DE MEDI-CAL**

SI USTED NO ESTÁ DE ACUERDO CON LA DECISIÓN TOMADA PARA SU TRATAMIENTO MÉDICO, USTED PUEDE PRESENTAR UNA APELACIÓN. ESTA APELACIÓN ES PRESENTADA CON SU PLAN DE SALUD.

CÓMO PRESENTAR UNA APELACIÓN

Usted tiene **60 días** de la fecha de esta carta de “Aviso de Acción” para presentar una apelación. Pero, **si usted está actualmente recibiendo tratamiento y desea continuar recibiendo el tratamiento, tiene que solicitar una apelación dentro de los 10 días** de la fecha en que esta carta fue marcada con el sello postal o entregado a usted O antes de la fecha en que su plan de salud dice que los servicios van a terminar. Usted tiene que decir que desea seguir recibiendo tratamiento cuando presente la apelación.

Usted puede presentar una apelación por teléfono, por escrito o electrónicamente.

- **Por teléfono:** contacte a Kern Family Health Care entre las 8:00 a.m. y las 5:00 p.m., llamando al (661) 632-1590 en Bakersfield o al 1-800-391-2000 fuera de Bakersfield o si no puede oír o hablar bien, por favor llame al 711.
- **Por escrito:** llene un formulario de apelación o escriba una carta y envíela a:

Kern Family Health Care
2900 Buck Owens Boulevard
Bakersfield, CA 93308

La oficina de su doctor va a tener formularios de apelación disponibles. Su plan de salud puede enviarle también un formulario.

- Electrónicamente: visite la página web de su plan de salud. Vaya a <http://www.kernfamilyhealthcare.com/>.

Usted mismo puede presentar una queja o puede tener a un pariente, amigo, defensor, doctor o abogado que presente la apelación por usted. Usted puede enviar cualquier tipo de información que desee que su plan de salud revise. Un doctor que es diferente que el doctor quien tomó la primera decisión, va a ver su apelación.

Su plan de salud tiene 30 días para darle una respuesta. En ese momento, usted recibirá una carta de “Aviso de Resolución de la Apelación”. Esta carta le dirá lo que el plan de salud ha decidido. **Si usted no recibe una carta dentro de los 30 días, usted puede:**

- Solicitar por una **“Revisión Médica Independiente” (IMR, por sus siglas en inglés)** y una persona de fuera, que no está relacionada con el plan de salud, va a revisar su caso.
- Solicitar por una **“Audiencia Estatal”** y un juez va a revisar su caso.

Por favor lea la sección siguiente para las instrucciones de cómo solicitar una IMR o una Audiencia Estatal.

APELACIONES DE URGENCIA

Si usted cree que esperar 30 días va a dañar su salud, usted puede tener una respuesta dentro de las 72 horas. Cuando presente su apelación, diga por qué el esperar va a dañar su salud. Asegúrese de solicitar por una **“apelación de urgencia”**.

SI NO ESTÁ DE ACUERDO CON LA DECISIÓN DE LA APELACIÓN

Si usted presenta una apelación y recibe una carta de “Aviso de Resolución de Apelación” informándole que su plan de salud todavía no le proporcionará los servicios o que **nunca recibió una carta informándole de la decisión y ya han pasado 30 días**, usted puede:

- Solicitar una **“Revisión Médica Independiente” (IMR, por sus siglas en inglés)** y una persona de fuera, que no está relacionada con el plan de salud, va a revisar su caso.
- Solicitar por una **“Audiencia Estatal”** y un juez va a revisar su caso.

Usted puede solicitar ambos, una IMR y una Audiencia Estatal al mismo tiempo.

También puede solicitar una antes que la otra, para ver si se resuelve su problema primero. Por ejemplo, si usted solicita por una IMR primero, pero no está de acuerdo con la decisión, todavía puede solicitar una Audiencia Estatal más tarde. Sin embargo, si solicita por una Audiencia Estatal primero, pero la audiencia ya ha tenido lugar, ya no puede solicitar por una IMR. En este caso, La Audiencia Estatal tiene la última palabra.

Usted no tiene que pagar por una IMR o Audiencia Estatal.

REVISIÓN MÉDICA INDEPENDIENTE (IMR, POR SUS SIGLAS EN INGLÉS)

Si usted quiere una IMR, tiene que presentar primero una apelación con su plan de salud. Si usted no tiene noticias de su plan de salud dentro de los 30 días o si está descontento con la decisión de su plan de salud, entonces puede solicitar una IMR. Tiene que solicitar por una IMR dentro de los **180 días** de la fecha de la carta del “Aviso de la Resolución de Apelación”.

Usted puede obtener una IMR de inmediato, sin presentar primero una apelación. Esto es en los casos que su salud está en peligro inmediato o que la solicitud fue denegada, porque el tratamiento es considerado experimental o de investigación.

El párrafo siguiente va a proporcionarle con la información de cómo solicitar una IMR. Tenga en cuenta que el término “quejas” está hablando de “apelaciones”.

El Departamento de Atención Médica Administrada es responsable de regular los planes de servicio de cuidado de salud. Si usted tiene una queja en contra de su plan de salud, debería llamarlos primero al **1-800-391-2000** y usar su proceso de quejas del plan de salud antes de contactar al Departamento. Utilizando este procedimiento de quejas no prohíbe ningún derecho legal potencial o remedios que pueden estar disponibles para usted. Si necesita ayuda con esta queja involucrando una emergencia, una queja que no ha sido resuelta satisfactoriamente por su plan de salud o una queja que se ha mantenido sin resolver por más de 30 días, usted puede llamar al Departamento para asistencia. También puede ser elegible para una Revisión médica Independiente (IMR, por sus siglas en inglés). Si usted es elegible para una IMR, el proceso de la IMR va a proporcionar una revisión imparcial de las decisiones médicas tomadas por el plan de salud, relacionadas a la necesidad médica del servicio propuesto o el tratamiento, las decisiones cubiertas para los tratamientos que son experimentales o por naturaleza de investigación y las disputas de pago por emergencia o los servicios médicos urgentes. El Departamento también tiene un número de teléfono gratuito (**1-888-HMO-2219**) y una línea TDD (**1-877-688-9891**) para la discapacidad auditiva y del habla. El sitio web del Departamento (<http://www.hmohelp.ca.gov>) tiene formularios de quejas, formularios para aplicaciones de la IMR e instrucciones en línea.

AUDIENCIA ESTATAL

Si desea una Audiencia Estatal, tiene que solicitarla dentro de los **120 días** de la fecha de la carta "Aviso de Resolución de la Apelación". Puede solicitar una Audiencia Estatal por teléfono o por escrito:

- Por teléfono: llame al **1-800-952-5253**. Este número puede estar bien ocupado. Usted puede recibir un mensaje para que llame más tarde. Si no puede hablar o escuchar bien, por favor llame **TTY/TDD 1-800-952-8349**.
- Por escrito: llene un formulario de Audiencia Estatal o envía una carta a:

**California Department of Social Services
State Hearing Division
P.O. Box 944243, Mail Station 9-17-37
Sacramento, CA 94244-2430**

Asegúrese de incluir su nombre, dirección, número de seguro social y la razón por la que quiere una Audiencia Estatal. Si alguien le está ayudando a solicitar una Audiencia Estatal, añada su nombre, dirección y número de teléfono en el formulario o carta. Si necesita un intérprete, díganos qué idioma habla. No tiene que pagar por un intérprete. Le daremos uno.

Después de solicitar la Audiencia Estatal, podría tomar hasta 90 días para decidir su caso y se le envíe una respuesta. Si usted cree que esperar tanto tiempo puede dañar su salud, puede obtener una respuesta dentro de los 3 días hábiles. Solicite a su doctor

o al plan de salud que le escriban una carta. La carta tiene que explicar en detalle cómo esperar hasta 90 días para que su caso sea decidido va a dañar seriamente su vida, su salud o su habilidad de lograr, mantener o recuperar la función máxima. Entonces, asegúrese de solicitar por una “**audiencia de emergencia**” y proporcionar la carta con su solicitud para una audiencia.

Usted mismo puede hablar con la Audiencia Estatal o puede tener a un pariente, amigo, defensor, doctor o abogado que hable por usted. Si usted quiere otra persona que hable por usted, entonces tiene que decirle a la oficina de la Audiencia Estatal que la persona tiene permiso de hablar en su nombre. Esta persona se llama “representante autorizado”.

AYUDA LEGAL

Usted puede obtener ayuda legal gratuita. Llame a Greater Bakersfield Legal Assistance al (661) 325-5943. También puede llamar al Legal Aid Society en su condado al 1-888-804-3536.

HOW TO FILE A GRIEVANCE

If you believe that Kern Family Health Care has failed to provide these services or discriminated in another way on the basis of race, color, national origin, age, disability, or sex, you can file a grievance with Kern Family Health Care. You can file a grievance by phone, in writing, in person, or electronically:

- **By phone:** Contact Kern Family Health Care between 8:00 a.m. to 5:00 p.m. by calling (661) 632-1590 inside Bakersfield, or 1-800-391-2000 outside Bakersfield. Or, if you cannot hear or speak well, please call 711.
 -
 - **In writing:** Fill out a complaint form or write a letter and send it to:

Kern Family Health Care
2900 Buck Owens Boulevard
Bakersfield, CA 93308
 - **In person:** Visit your doctor's office or Kern Family Health Care and say you want to file a grievance.
 - **Electronically:** Visit Kern Family Health Care's website at <http://www.kernfamilyhealthcare.com/>.
-

OFFICE OF CIVIL RIGHTS

You can also file a civil rights complaint with the U.S. Department of Health and Human Services, Office for Civil Rights by phone, in writing, or electronically:

- **By phone:** Call **1-800-368-1019**. If you cannot speak or hear well, please call **TTY/TDD 1-800-537-7697**.
- **In writing:** Fill out a complaint form or send a letter to:

**U.S. Department of Health and Human Services
200 Independence Avenue, SW
Room 509F, HHH Building
Washington, D.C. 20201**

Complaint forms are available at <http://www.hhs.gov/ocr/office/file/index.html>.

- **Electronically:** Visit the Office for Civil Rights Complaint Portal at <https://ocrportal.hhs.gov/ocr/portal/lobby.jsf>.

CÓMO PRESENTAR UNA QUEJA

Si usted cree que Kern Family Health Care ha fallado en proporcionar estos servicios o ha discriminado de alguna otra manera en base a raza, color, origen nacional, edad, discapacidad o sexo, usted puede presentar una queja con Kern Family Health Care. Puede presentar una queja por teléfono, por escrito, en persona o electrónicamente:

- Por teléfono: contacte a Kern Family Health Care entre las 8:00 a.m. a las 5:00 p.m. llamando al (661) 632-1590 en Bakersfield o al 1-800-391-2000 fuera de Bakersfield o si no puede oír o hablar bien, por favor llame al 711.
 - Por escrito: llene un formulario de quejas o escriba una carta y envíela a:

Kern Family Health Care
2900 Buck Owens Boulevard
Bakersfield, CA 93308
 - En persona: visite la oficina de su doctor o de Kern Family Health Care y diga que desea presentar una queja.
 - Electrónicamente: visite el sitio web de Kern Family Health Care en <http://www.kernfamilyhealthcare.com/>.
-

OFICINA DE LOS DERECHOS CIVILES

Usted también puede presentar una queja de los derechos civiles, con el Departamento de Salud y Servicios Humanos, Oficina para los Derechos Civiles, por teléfono, por escrito o electrónicamente:

- Por teléfono: llame al **1-800-368-1019**. Si no puede hablar o no puede oír bien, por favor llame al: TTY/TDD 1-800-537-7697.
- Por escrito: llene un formulario de quejas o envíe una carta a:

**U.S. Department of Health and Human Services
200 Independence Avenue, SW
Room 509F, HHH Building
Washington, D.C. 20201**

Los formularios de quejas están disponibles en:
<http://www.hhs.gov/ocr/office/file/index.html>.

- Electrónicamente: visite el portal de quejas de la Oficina para los Derechos Civiles en <https://ocrportal.hhs.gov/ocr/portal/lobby.jsf>.

MEDWATCHThe FDA Safety Information and
Adverse Event Reporting ProgramFor VOLUNTARY reporting of
adverse events and product problems

Page ____ of ____

FOCUS ONLYTriage unit
sequence #**A. PATIENT INFORMATION**

1. Patient Identifier	2. Age at Time of Event: or Date of Birth:	3. Sex <input type="checkbox"/> Female <input type="checkbox"/> Male	4. Weight ____ lbs or ____ kgs
-----------------------	--------------------------------------------------	----------------------------------------------------------------------------	-----------------------------------------

In confidence

B. ADVERSE EVENT OR PRODUCT PROBLEM1. ☐ Adverse Event and/or ☐ Product Problem (e.g., defects/malfunctions)2. Outcomes Attributed to Adverse Event
(Check all that apply)

- ☐ Death: _____ (mo/day/yr)
☐ Life-threatening
☐ Hospitalization - initial or prolonged
☐ Disability
☐ Congenital Anomaly
☐ Required Intervention to Prevent Permanent Impairment/Damage
☐ Other: _____

3. Date of Event (mo/day/yr)

4. Date of This Report (mo/day/yr)

5. Describe Event or Problem:

6. Relevant Tests/Laboratory Data, including Dates

7. Other Relevant History, including Preexisting Medical Conditions (e.g., allergies, race, pregnancy, smoking and alcohol use, hepato/biliary dysfunction, etc.)

C. SUSPECT MEDICATION(S)

1. Name (Give labeled strength & manufacturer, if known)

#1

#2

2. Dose, Frequency & Route Used

#1

#2

3. Therapy Dates (If unknown, give duration)
from/to (or best estimate)

#1

#2

4. Diagnosis for Use (Indication)

#1

#2

6. Event Abated After Use
Stopped or Dose Reduced?#1 ☐ Yes ☐ No ☐ Doesn't Apply#2 ☐ Yes ☐ No ☐ Doesn't Apply

6. Lot # (If known)

#1

#2

7. Exp. Date (If known)

#1

#2

8. Event Reappeared After
Reintroduction?#1 ☐ Yes ☐ No ☐ Doesn't Apply#2 ☐ Yes ☐ No ☐ Doesn't Apply

9. NDC# (For product problems only)

10. Concomitant Medical Products and Therapy Dates (Exclude treatment of event)

D. SUSPECT MEDICAL DEVICE

1. Brand Name

2. Type of Device

3. Manufacturer Name, City and State

4. Model #

Lot #

Catalog #

Expiration Date (mo/day/yr)

Serial #

Other #

5. Operator of Device

- ☐ Health Professional
☐ Lay User/Patient
☐ Other: _____

6. If Implanted, Give Date (mo/day/yr)

7. If Explanted, Give Date (mo/day/yr)

8. Is this a Single-use Device that was Reprocessed and Reused on a Patient?

☐ Yes ☐ No

9. If Yes to Item No. 8, Enter Name and Address of Reprocessor

10. Device Available for Evaluation? (Do not send to FDA)

☐ Yes ☐ No ☐ Returned to Manufacturer only

(mo/day/yr)

11. Concomitant Medical Products and Therapy Dates (Exclude treatment of event)

E. REPORTER (See confidentiality section on back)

1. Name and Address

Phone #

2. Health Professional?

☐ Yes ☐ No

3. Occupation

4. Also Reported to:

- ☐ Manufacturer
☐ User Facility
☐ Distributor/Importer

5. If you do NOT want your identity disclosed
to the manufacturer, place an "X" in this box: ☐

PLEASE TYPE OR USE BLACK INK



Mail to: **MEDWATCH**
 6500 Fishers Lane
 Rockville, MD 20852-9787

-or- FAX to:
 1-800-FDA-0178

ADVICE ABOUT VOLUNTARY REPORTING

Report adverse experiences with:

- Medications (drugs or biologics)
- Medical devices (including in-vitro diagnostics)
- Special nutritional products (dietary supplements, medical foods, infant formulas)
- Cosmetics
- Medication errors

Report product problems - quality, performance or safety concerns such as:

- Suspected contamination
- Questionable stability
- Defective components
- Poor packaging or labeling
- Therapeutic failures

Report SERIOUS adverse events. An event is serious when the patient outcome is:

- Death
- Life-threatening (real risk of dying)
- Hospitalization (initial or prolonged)
- Disability (significant, persistent or permanent)
- Congenital anomaly
- Required intervention to prevent permanent impairment or damage

Report even if:

- You're not certain the product caused the event
- You don't have all the details

How to report:

- Just fill in the sections that apply to your report
- Use section C for all products except medical devices
- Attach additional blank pages if needed
- Use a separate form for each patient
- Report either to FDA or the manufacturer (or both)

Confidentiality: The patient's identity is held in strict confidence by FDA and protected to the fullest extent of the law. FDA will not disclose the reporter's identity in response to a request from the public, pursuant to the Freedom of Information Act. The reporter's identity, including the identity of a self-reporter, may be shared with the manufacturer unless requested otherwise.

If your report involves a serious adverse event with a device and it occurred in a facility outside a doctor's office, that facility may be legally required to report to FDA and/or the manufacturer. Please notify the person in that facility who would handle such reporting.

Important numbers:

- 1-800-FDA-0178 -- To FAX report
- 1-800-FDA-1088 -- To report by phone or for more information
- 1-800-822-7967 -- For a VAERS form for vaccines

To Report via the Internet:

<http://www.fda.gov/medwatch/report.htm>

The public reporting burden for this collection of information has been estimated to average 30 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to:

Department of Health and Human Services
Food and Drug Administration
MedWatch; HFD-410
5600 Fishers Lane
Rockville, MD 20857

Please DO NOT
RETURN this form
to this address.

OMB statement:

"An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number."

U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES
Food and Drug Administration

FORM FDA 3500 (9/03) (Back)

Please Use Address Provided Below -- Fold In Thirds, Tape and Mail

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service
Food and Drug Administration
Rockville, MD 20857

Official Business
Penalty for Private Use \$300

BUSINESS REPLY MAIL

FIRST CLASS MAIL PERMIT NO. 946 ROCKVILLE MD

POSTAGE WILL BE PAID BY FOOD AND DRUG ADMINISTRATION

MEDWATCH

The FDA Safety Information and Adverse Event Reporting Program
Food and Drug Administration
5600 Fishers Lane
Rockville, MD 20852-9787

NO POSTAGE
NECESSARY
IF MAILED
IN THE
UNITED STATES
OR APO/FPO

