

Public Health Service

Food and Drug Administration Rockville, MD 20857

NDA 20-125/S-007

Pfizer Inc.

Attention: Kathleen Collins Manager, Worldwide Regulatory Strategy 235 East 42nd Street (685/18/62) New York, NY 10017

SUPPLEMENT APPROVAL

Dear Ms. Collins:

Please refer to your supplemental new drug application (NDA) dated April 24, 2009 submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act for Accuretic (quinapril hydrochloride/hydrochlorothiazide) tablets.

This changes being effected supplemental new drug application provides for the following addition under the **PRECAUTIONS**, **Drug Interactions** section of the package insert:

Gold: Nitritoid reactions (symptoms include facial flushing, nausea, vomiting and hypotension) have been reported rarely in patients on therapy with injectable gold (sodium aurothiomalate) and concomitant ACE inhibitor therapy.

We have completed our review of this application, and it is approved, effective on the date of this letter, for use as recommended in the electronic draft labeling text.

If you issue a letter communicating important information about this drug product (i.e., a "Dear Health Care Professional" letter), we request that you submit a copy of the letter to this NDA and a copy to the following address:

MEDWATCH Food and Drug Administration 5515 Security Lane HFD-001, Suite 5100 Rockville, MD 20852

Marketing the product with labeling that is not identical to the approved labeling text may render the product misbranded and an unapproved new drug.

We remind you that you must comply with reporting requirements for an approved NDA (21 CFR 314.80 and 314.81).

If you have any questions, please call Michael Monteleone, M.S., Regulatory Project Manager,

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at (301)796-1952.

Sincerely,

{See appended electronic signature page}

Mary Ross Southworth, Pharm.D.
Deputy Director for Safety
Division of Cardiovascular and Renal Products
Office of Drug Evaluation I
Center for Drug Evaluation and Research

cc: Enclosed Labeling Text

Application Type/Number	Submission Type/Number	Submitter Name	Product Name
 NDA-20125	SUPPL-7	PFIZER PHARMACEUTICA LS LTD	ACCURETIC (QUINAPRIL/HCTZ) TABLETS
		electronic record s the manifestation	
/s/			
MARY R SOUTH	WORTH		

09/18/2009