United Research Laboratories, Inc.
Mutual Pharmaceutical Company, Inc.

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Philadelphia, PA 19124

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July 1, 2003

Dockets Management Branch
Food and Drug Administration (HFA-305)
Department of Health and Human Services
5630 Fishers Lane, Room 1061
Rockville, MD 20852

Re: **Docket Number 2003P-0214**

Dear Sir or Madam:

On May 18, 2003 I submitted a Citizens Petition requesting the Food and Drug Administration to correct the listing of Mutual Pharmaceutical's quinidine gluconate extended-release tablets in the Orange Book. That Citizens Petition was assigned Docket Number 2003P-0214.

It has come to my attention that the FDA has made the requested change in the Orange Book and Mutual's quinidine gluconate product is now identified as the reference listed drug.

Consequently, I hereby withdraw my Citizens Petition regarding this matter.

Sincerely

Robert Dettery

Robert Dettery
Vice President, Regulatory Affairs
Mutual Pharmaceutical Company

Cc: G. Davis (Office of Generic Drugs)
C. Parise (Office of Generic Drugs)
D. Hare (Office of Generic Drugs)

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