LACHMAN CONSULTANT SERVICES, INC.
CONSULTANTS TO THE PHARMACEUTICAL AND ALLIED INDUSTRIES

1600 STEWART AVENUE, WESTBURY, NY 11590
(516) 222-6222 • FAX (516) 683-1887

2012 JAN -4 A 9:52

January 3, 2012

OVERNIGHT COURIER 1/3/2012

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

2006-P-0078 (Legacy No. 2006P-0122)
Cetirizine Hydrochloride Orally Disintegrating Tablets, 5 mg and 10 mg)
Request to Withdraw Petition

Dear Sir or Madam:

Reference is made to the above cited petition which was filed on March 17, 2006, under Section 505(j)(2)(C) of the Federal Food, Drug and Cosmetic Act, and in accordance with 21 CFR 10.30 on behalf of a client requesting the Commissioner of the Food and Drug Administration to declare that the drug product Cetirizine Hydrochloride Orally Disintegrating Tablets, 5 mg and 10 mg, is suitable for consideration in an abbreviated new drug application (ANDA).

We are hereby requesting FDA take action to withdraw this petition.

Respectfully submitted,

Robert Pollock

Robert W. Pollock PK
Executive Vice President

RWP/pk

cc: Johnny Young (OGD)

T05_Petition_Withdrawal_Cetirizine_010312

FDA-2006-P-0078 WDL

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