



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service
Food and Drug Administration

CENTER FOR DRUG EVALUATION AND RESEARCH

Division of Manufacturing and Product Quality
Foreign Inspection Team, HFD-325
11919 Rockville Pike, MM2
Rockville, MD 20852

TELEPHONE: (301) 827-9021
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November 19, 2007

Graham Pettipher
General Manager
Reading Scientific Services LTD
Lord Zuckerman Research Center,
Whiteknights Campus, Pepper Lane
Reading
United Kingdom

Dear Mr. Pettipher:

We have completed our review of the Establishment Inspection Report (EIR) for the inspection conducted at your control testing laboratory in Reading, United Kingdom by FDA Investigator Regina T. Brown. A FDA 483, Notice of Inspectional Observations was issued at the conclusion of that inspection.

We have also reviewed the October 4, 2007 written response to the FDA 483 observation. Based on the corrections described in the response, we are classifying your facility as acceptable. The corrective actions will be further evaluated during the next routine inspection of this facility. It remains your responsibility to assure continued compliance with good manufacturing practices. This letter is not intended as an endorsement or certification of the facility.

Since the Agency is working to make its regulatory process and activities more transparent to the regulated industry, enclosed is a copy of the Establishment Inspection Report (EIR) for the above inspection. The enclosed copy contains only the narrative portion of the report. However, you may request additional information under the Freedom of Information Act.

You may contact me at Montrose Metro II, 11919 Rockville Pike, Rockville, MD 20852. You may also contact my office by telephone at (301)827-9073 or by fax at (301)827-8909.

Sincerely,

Anthony A. Charity,
Compliance Officer
Foreign Inspection Team, HFD-325

Enclosure: