



DEPARTMENT OF HEALTH & HUMAN SERVICES

Food and Drug Administration  
Rockville MD 20857

G-003299-K-0000

Norchim  
Attention: Jean Pierre Allaire  
Chief Executive Office/Site Director  
33 Quai D'amont  
Saint Leu D'Esserent,  
France

NOV 02 2006

Re: GMP Inspection that occurred July 10 - 13, 2006

Dear Mr. Allaire:

On July 10 - 13, 2006, an inspection was conducted at your facility located at Norchim, France by FDA Investigator David M. Beltran. The inspection revealed no serious cGMP deficiencies; therefore, no Form FDA-483 was issued at the conclusion of the inspection.

Based on the above inspection, your facility is classified as acceptable. We remind you that it is your responsibility to assure continued compliance to current good manufacturing practices.

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.

If you have questions regarding the enclosed EIR, please contact me at (301) 827-6963.

Sincerely,

Mai X. Huynh  
Team Leader  
Antimicrobial Team  
Division of Manufacturing Technologies  
Office of New Animal Drug Evaluation  
Center for Veterinary Medicine

Enclosure