



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service  
Food and Drug Administration

CENTER FOR DRUG EVALUATION AND RESEARCH

REÇU 10 AVR. 2013

Office of Manufacturing and Product Quality  
Division International Drug Quality  
International Compliance Branch  
10903 New Hampshire Avenue  
Building #51, Room 4234  
Silver Spring, MD 20993

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March 28, 2013

Mr. Michel Butin  
Managing Director and CEO  
Albhades Provence  
ZA Les Roubines, B.P. 51  
Oraison, 04700, France

Reference: FEI 3006218344

Dear Mr. Butin:

We have completed our review of the Establishment Inspection Report (EIR) for the inspection conducted at your control testing laboratory in Oraison, France by Investigator Vlada Matusovsky and Chemist Temar Q. Williams during the period of December 17 - 18, 2012. An FDA-483, Notice of Inspectional Observations was issued at the conclusion of the inspection.

We have also reviewed your company's response dated January 4, 2013 with supportive documentation. Based on the profile class covered during the inspection, we are classifying your facility as acceptable. This letter is not intended as an endorsement or certification of the facility. It remains your responsibility to assure continued compliance with current good manufacturing practice (CGMP).

Please be advised that all manufacturers must register annually as required by 21 C.F.R. § 207.40. Information on how to register is available at [http://www.fda.gov/cder/drls/registration\\_listing.htm](http://www.fda.gov/cder/drls/registration_listing.htm)

Additionally, we enclose a copy of the establishment inspection report (EIR). Releasing this EIR to you is part of FDA's effort to make its regulatory process and activities more transparent to the regulated industry. It is being provided to you for information purposes only and may reflect some redactions made by the Agency in accordance with the Freedom of Information Act and 21 C.F.R. Part 20. Copies provided to other requestors may have additional redactions of trade secret and confidential commercial information.

If you have any questions regarding this letter, you may contact me at the above address or number.

Sincerely,

*Mary E. Farberman - Mary E. Farberman*

Alicia Mozzachio  
Branch Chief  
Division of International Drug Quality

Enclosure: EIR