



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service
Food and Drug Administration

CENTER FOR DRUG EVALUATION AND RESEARCH

Office of Manufacturing and Product Quality
Division International Drug Quality
International Compliance Branch
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June 26, 2014

Dr. Christian Poinot
CEO & Scientific Director
Icare
B.P. 60006
St. Beauzire, France 63360

Reference: FEI 3008220949

Dear Dr. Poinot:

We have completed our review of the Establishment Inspection Report (EIR) for the inspection conducted at your contract control testing laboratory for pharmaceutical and medical devices in St. Beauzire, France by Investigator Kenneth Klobus during the period of March 17, 2014 to March 18, 2014.

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 practices (CGMPs).

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.

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,

Alicia Mozzachio
Branch Chief
Division of International Drug Quality

Enclosure: EIR