

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration Center for Drug Evaluation and Research Office of Pharmaceutical Quality Office of Surveillance Division of Quality Surveillance Assessment 10903 New Hampshire Avenue Building 51, Room 4316 Silver Spring, MD 20993 TELEPHONE: (301) 796-3254 FAX: (301) 847-8742

08/17/2016

Wickham Laboratories Limited Hoeford Point, Barwell Lane Gosport, Hampshire, GB

Reference: Inspection Date(s): 02/15/2016 - 02/17/2016

Location: Wickham Laboratories Limited Hoeford Point, Barwell Lane Gosport, PO13 0AU, GB

Dear Mr. John Mckenzie,

We are enclosing a copy of the establishment inspection report (EIR) for the inspection that the U.S. Food and Drug Administration (FDA) conducted at your premises on the referenced locale and date(s). When the Agency concludes that an inspection is "closed" under 21 CFR 20.64(d)(3), it will release a copy of the EIR to the inspected establishment. This procedure is applicable to EIRs for inspections completed on or after April 1, 1997.

The Agency continually works to make its regulatory process and activities more transparent to the regulated industry. Releasing this EIR to you is part of this effort. The copy being provided to you comprises the narrative portion of the report; it may reflect redactions made by the Agency in accordance with the Freedom of Information Act (FOIA) and 21 CFR Part 20. This, however, does not preclude you from requesting additional information under FOIA.

If there is any question about the released information, feel free to contact me at 240-402-6594.

For more information on the U.S. FDA, please visit our website at www.fda.gov.

FEI: 3009798720

Enclosure: Establishment Inspection Report (EIR)