U.S. FOOD & DRUG ADMINISTRATION

Food and Drug Administration
Center for Drug Evaluation and Research
Office of Pharmaceutical Quality
Office of Surveillance
Division of Quality Surveillance Assessment
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09/14/2017

Eurofins BioPharma Product Testing Spain SLU
Esplugues de Llobregat
Calle Josep Argemi 13-15
Esplugues de Llobregat, Barcelona , ES

Reference: Inspection Date(s): 06/27/2016 - 06/29/2016

Location: Eurofins BioPharma Product Testing Spain SLU
Calle Josep Argemi 13-15
Esplugues de Llobregat
Barcelona, 08950 , ES

Dear Mr. Salvi,

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 301-796-1287.

fda_phone

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

Sincerely,

Merideth K. Rose -S

Digitally signed by Merideth K. Rose -S
DN: c=US, o=U.S. Government, ou=HHS, ou=FDA, ou=People, 0.9.2342.19200300.100.1.1=1300419551, cn=Merideth K. Rose -S
Date: 2017.05.19 09:50:43 -04'00'

FEI: 3007914833

Enclosure: Establishment Inspection Report (EIR)