



# 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  
10903 New Hampshire Avenue  
Building 51, Room 4316  
Silver Spring, MD 20993  
TELEPHONE: (301) 796-3254  
FAX: (301) 847-8742

02/02/2017

AmatsiAvogadro  
Parc De Genibrat  
Fontenilles, , FR

Reference: Inspection Date(s): 09/26/2016 - 09/28/2016

Location: AmatsiAvogadro  
Parc De Genibrat  
Fontenilles, , FR

Dear Mr. Arnaud,

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 301-796-3254.

For more information on the U.S. FDA, please visit our website at [www.fda.gov](http://www.fda.gov).

Sincerely,

*Monique Hagler*

FEI: 3006118993

Enclosure: Establishment Inspection Report (EIR)