

## FORM LA200-1. Report on Receipt of Drug - Guide for Reporting Investigational New Animal Drug Shipments for Poikilothermic Food Animals

### INSTRUCTIONS

1. Investigator must fill out Form LA200-1 **immediately** upon receipt of LIQUAMYCIN<sup>®</sup> LA-200<sup>®</sup>.
2. Investigator should keep the original on file, and send one copy to the Study Monitor for review.
3. Within 10 days of receipt, the Study Monitor should send a copy to the AADAP Office.
4. **Note:** Both Investigator and Study Monitor should sign and date Form LA200-1.

*The sponsor, U.S. Fish and Wildlife Service, submits a notice of claimed investigational exemption for the shipment or delivery of a new animal drug under the provisions of Section 512 of the Federal Food, Drug, and Cosmetics Act.*

|  |   |  |             |
|--|---|--|-------------|
| Name of Drug   | <b>LIQUAMYCIN<sup>®</sup><br/>LA-200<sup>®</sup></b>                | INAD Number                                  | <b>9027</b> |
| Proposed Use of Drug                                 | To control certain bacterial diseases in a variety of fish species. |  |             |
| Date of CVM Authorization Letter                     | January 11, 2008  |  |             |
| <b>Date of Drug Receipt</b>                          |   | <b>Amount of Drug Received</b>               |             |
| <b>Drug Lot Number</b>                               |   | <b>Study Worksheet Number</b>                |             |
| <b>Name of Investigator</b>                          |   |  |             |
| <b>Address of Investigator</b>                       |   |  |             |
| <b>Location of Trial</b>                             |   |  |             |
| Pivotal Study (yes/no)                               |   | Non-pivotal Study (yes/no)                   |             |
| <b>Approximate Number of Treated Animals</b>         |   | <b>Approximate Number of Control Animals</b> |             |
| <b>Number of Animals Used Previously<sup>1</sup></b> |   |  |             |
| Study Protocol Number                                | 9027  |  |             |
| <b>Approximate dates of trial (start/end)</b>        |   |  |             |
| <b>Species, Size, and Type of Animals</b>            |   |  |             |
| Maximum daily dose and duration                      | 20 mg OTC/Kg body weight  |  |             |
| Methods(s) of Administration                         | Single injection  |  |             |
| Withdrawal Period                                    | 30 days   |  |             |

<sup>1</sup> To be filled out by the NIO

**Date Prepared:** \_\_\_\_\_ **Investigator:** \_\_\_\_\_

**Date Reviewed:** \_\_\_\_\_ **Study Monitor:** \_\_\_\_\_

**Date Reviewed:** \_\_\_\_\_ **Sponsor:** \_\_\_\_\_