

Form OTIMM-3: Results Report Form for use of Terramycin 343[®] under INAD 9033**INSTRUCTIONS**

- Investigator must fill out Form OTIMM-3 no later than 10 days after completion of the 30-day post-treatment observation period. Study Number must be recorded on all pages of Form OTIMM-3. Attach lab reports and other information.
- If Terramycin 343[®] was not used under the assigned Study Number, fill out only the Site Information portion on this page, and skip to the end of page 4 and fill out only the "Negative Report" section.
- Investigator should keep the original on file, and send a copy to the Study Monitor. Within 10 days of receipt, the Study Monitor should send a copy to the AADAP Office for inclusion in the permanent file.
- Note:** Both Investigator and Study Monitor should sign and date Form OTIMM-3.

SITE INFORMATION

Facility	Fish Hatchery A
Reporting Individual	John Doe

TREATMENT INFORMATION AND SCHEDULE

Drug lot number	111222	Total amount drug used (gm)	1,600
Fish species treated	White sturgeon	OTIMM dosage used (mg/L)	20
Duration of drug treatment (hours)	1	Number of treatments	4
Disease treated	BGD	Disease diagnosed by	Josey Jo
Average fish weight (gm)	80.0	Average fish length (in)	10.0
Number of fish per unit (e.g. 10,000 fish/raceway)			7,800
Number of treated units	1	Total number of treated fish	7,800
Number of control units	0	Total number of control fish	0
Treatment date(s)		3/3 – 3/6/08	

WATER QUALITY PARAMETERS

Ave pre-treatment temp (°F)	70	Dissolved Oxygen (mg/L)	9.0
Ave treatment temp (°F)	70	pH	6.8
Ave post-treatment temp (°F)	70	Hardness - CaCO ₃ (mg/L)	75

Daily Mortality Record

INSTRUCTIONS

1. Investigator should fill out the Daily Mortality Record as completely as possible.
2. Prior to initiation of the trial, fill out Rearing Unit ID, whether a rearing unit is Treated or Control, and the number of fish in each rearing unit.
3. Water temperature and individual tank mortality should be recorded on a daily basis.
4. Please mark all treatment days with an asterisk.
5. Use additional copies of this form if more than 6 rearing units are involved in the trial.

FACILITY		Fish Hatchery A							
Pre-treatment period	Rearing Unit ID		1						
	Treated or Control		T						
	Number of Fish		7,800						
	Day	Date	Water Temp (F°)	Mortality	Mortality	Mortality	Mortality	Mortality	Mortality
	1	2/27/08	70	0					
	2	2/28	70	1					
	3	2/29	70	2					
	4	3/1	70	8					
	5	3/2	70	10					
Treatment period	1	3/3	70	12					
	2	3/4	70	5					
	3	3/5	70	3					
	4	3/6	70	0					
Post-treatment period	1	3/7	70	0					
	2	3/8	70	0					
	3	3/9	70	0					
	4	3/10	70	0					
	5	3/11	70	1					
	6	3/12	70	0					
	7	3/13	70	0					
	8	3/14	70	1					
	9	3/15	70	0					
	10	3/16	70	0					

RESULTS: Describe in detail treatment results. Was treatment successful? If treatment did not appear to be successful, explain why not? Were there any mitigating environmental conditions that may have impacted treatment results? Were there any deviations from the Study Protocol?

Treatment appeared to be successful. Mortality returned to normal levels following the treatment period.

Pathology Report: Attach pathology report to this form. Report should include: 1) a description of how the pathogen(s) was identified; 2) disease identification records that confirm the presence of the pathogen; and 3) the name and title of the individual performing the diagnosis.

Pathology Report included: pre-treatment post-treatment

Toxicity observations: Report any apparent drug toxicity including a description of unusual fish behavior.

none

DRUG DISCHARGE RESULTING FROM THIS TREATMENT: Use Addendum 2: Discharge Worksheet for calculations and attach completed Discharge Worksheet to this form. Enter the value from Addendum 2 step 3 in this space.

0.033 ppm

OBSERVED WITHDRAWAL PERIOD: (Investigator should initial the appropriate box below)

Observed withdrawal period: _____ 21 days; Objectives A & B

Observed withdrawal period: x _____ 60 days; Objectives C & D

Estimated number of days between last treatment and first availability of fish for human consumption (ensure this time period meets the withdrawal period). >1000 days

NEGATIVE REPORT Oxytetracycline for Immersion was not used at this facility under this Study Number during the reporting period. (Investigator should initial for negative reports as soon as the Study Number is known to be no longer needed or valid.)

Date Prepared: 3/20/08

Investigator: Sign here

Date Reviewed: 3/20/08

Study Monitor: Sign here