

Aquaflor[®] Veterinary Feed Directive (Florfenicol)



Client: _____ **Veterinarian:** _____
Address: _____ **Address:** _____

Phone: _____ **Phone:** _____
Fax: _____ **Fax:** _____
E-mail: _____ **E-mail:** _____

Catfish and/or Freshwater-reared Salmonids to be Treated: Number, Total Weight (Biomass): _____
Farm Location: Farm Address, Raceway/Pond Identification (Raceway/Pond Number, etc.): _____

Indications: Catfish: For the control of mortality in catfish due to enteric septicemia of catfish associated with *Edwardsiella ictaluri*.
 Freshwater-reared Salmonids: For the control of mortality in freshwater-reared salmonids due to coldwater disease associated with *Flavobacterium psychrophilum* and furunculosis associated with *Aeromonas salmonicida*.

Mix into Type C Medicated Feed to Provide: Check one: 182 g/ton 300 g/ton 454 g/ton 908 g/ton 1816 g/ton

Amount of Final (Type C) Feed: _____ (Pounds or Tons) **Feeding Rate:** _____ % Biomass

Feeding Directions: Feed as the sole ration for 10 consecutive days. Feeding at this rate will deliver 10 mg florfenicol per kg of fish.

Feeding Rate	Florfenicol Concentration in Feed	Amount of Aquaflor [®] (florfenicol) per Ton	Biomass of Fish Medicated per Ton of Feed per 10-day Treatment Period
% Biomass	Grams/ton	lbs	lbs
0.5	1,816	8.00	40,000
1.0	908	4.00	20,000
2.0	454	2.00	10,000
3.0	300	1.32	6,666
5.0	182	0.80	4,000

Special Instructions

Date of Treatment: _____ (Month/Day/Year)
Expiration Date of VFD: _____ Month/Day/Year (Not to exceed 15 days from date of issuance.)
Veterinarian's Signature: _____ **Date:** _____
License Number and State: _____

Caution: Federal law limits this drug to use under the professional supervision of a licensed veterinarian. Animal feed bearing or containing this veterinary feed directive drug shall be fed to animals only by or upon a lawful veterinary feed directive (VFD) issued by a licensed veterinarian in the course of the veterinarian's professional practice. Extra-label use (i.e., use of this VFD feed in a manner other than as provided by the VFD drug approval) is strictly prohibited.

Feed containing Aquaflor[®] (florfenicol) shall not be fed to catfish or freshwater-reared salmonids for more than 10 days. Following 10 days administration, fish should be re-evaluated by a licensed veterinarian before initiating a further course of therapy. The expiration date for VFD for Aquaflor (florfenicol) must not exceed 15 days from the date of issuance. VFD for Aquaflor (florfenicol) shall not be refilled. For catfish, a dose related decrease in hematopoietic/lymphopoietic tissue may occur. The time required for the hematopoietic/lymphopoietic tissues to regenerate was not evaluated. The effects of florfenicol on reproductive performance have not been determined.

RESIDUE WARNING: Catfish: Feeds containing Aquaflor[®] (florfenicol) must be withdrawn 12 days prior to slaughter.

RESIDUE WARNING: Freshwater-reared Salmonids: Feeds containing Aquaflor[®] (florfenicol) must be withdrawn 15 days prior to slaughter.

Warning: Avoid inhalation, oral exposure, and direct contact with skin or eyes. Operators mixing and handling Aquaflor[®] should use protective clothing, gloves, goggles and NIOSH-approved dust mask. Wash thoroughly with soap and water after handling. If accidental eye contact occurs, immediately rinse thoroughly with water. If irritation persists, seek medical attention. Not for human consumption. Keep out of reach of children. The Material Safety Data Sheet (MSDS) contains more detailed occupational safety information. For a copy of the MSDS, call 1-800-770-8878. To obtain more information or to report adverse effects, call 1-800-211-3573.

NADA 141-246, Approved by FDA.
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