

LOAN COPY ONLY

THIS REPRINT RESULTS FROM RESEARCH
SPONSORED BY CALIFORNIA SEA GRANT

HISTOLOGY AND ULTRASTRUCTURE
OF THE JELLIED CONDITION OF DOVER SOLE,
Microstomus pacificus (Lockington)

by

Robert A. Fisher

CIRCULATING COPY
Sea Grant Depository

A Thesis

Presented to

The Faculty of Humboldt State University

In Partial Fulfillment

of the Requirements for the Degree

Master of Science

December, 1986

ABSTRACT

Muscle samples of "normal" and "jellied" Dover sole, Microstomus pacificus, were taken from fish on the fillet line at Eureka Fisheries, Inc., Eureka, California and from whole, live fish as they were brought on board a commercial fishing vessel out of Eureka, California. Muscle tissue samples were processed for percent moisture content and for histological and ultrastructural analyses. Percent moisture content for fish samples from the fillet-line ranged from 82.8 to 92.1 percent. Samples from the commercial dragger ranged from 84.3 to 89.2 percent moisture. Histologically, normal Dover sole muscle is typical vertebrate muscle. Jellied muscle was characterized by much intercellular space with a proliferation of connective tissue, loosely packed muscle cells resulting in spherically shaped cells in cross-section and severe waviness of cells in longitudinal-section, hypertrophic nuclei, and degraded muscle contractile elements. Occassionally, dystrophic-like characteristics such as internally positioned nuclei, myofibril fragmentation, and variations in fiber size were also observed. Ultrastructural examination of the jellied condition showed loosely packed, randomly spaced myofibrils with few myofibrils occupying the periphery of the cell. In the majority of samples observed, myofilaments were densely packed within myofibrils. Torn myofilaments and pyknotic appearing nuclei with oversized nuclear envelopes were observed. Three out of 100 fillets contained Kudoa

clupeidae cysts, suggesting that the jellied muscle condition is not likely a result of myxozoan infection.

ACKNOWLEDGEMENTS

I wish to acknowledge the California Sea Grant College Program for funding which made this study possible and the cooperative effort of Humboldt State University's Fisheries and Biology Departments.

Invaluable guidance was afforded to me by my committee members Dr. Ron Fritzsche, for fisheries systematics, statistical procedures, and editing my drafts, Dr. Gary Hendrickson, for histological and pathological procedures, and editing and reorganizing my original draft, and Dr. Dennis Walker, for electron microscopy procedures and editing of my thesis drafts.

I wish to thank Eureka Fisheries, Inc., for providing fish samples, the crew of the F/V ANNA W (Tom Swisher, Pat Williamson, and Captain Rex Coffman) for permitting me to obtain my live fish samples during their fishing trips at sea, Max Puckett for laboratory assistance, Chris Toole, the Humboldt County Marine Extension Agent for his time and effort in procuring cooperation between Humboldt State University and the local fishing industry, and Dr. John Baird and the Biology Department at Long Beach State University for the use of their Olympus photomicrographic system.

I must also acknowledge my parents, Joe and Evelyn Fisher, for their continued interest and support, my wife Nancy, for her support and tolerance, and also my daughter Dylan 'Lee who has made me realize the importance of completing this thesis and starting my career.

I wish to dedicate this thesis to the memory of my brother William Patrick Fisher with whom I shared a love for the sea.

This work is a result of research sponsored in part by NOAA, National Sea Grant College Program, Department of Commerce, under grant number NA80AA-D-00120, project number R/F 89, through the California Sea Grant College Program, and in part by the California State Resources Agency. The U. S. Government is authorized to reproduce and distribute for governmental purposes.