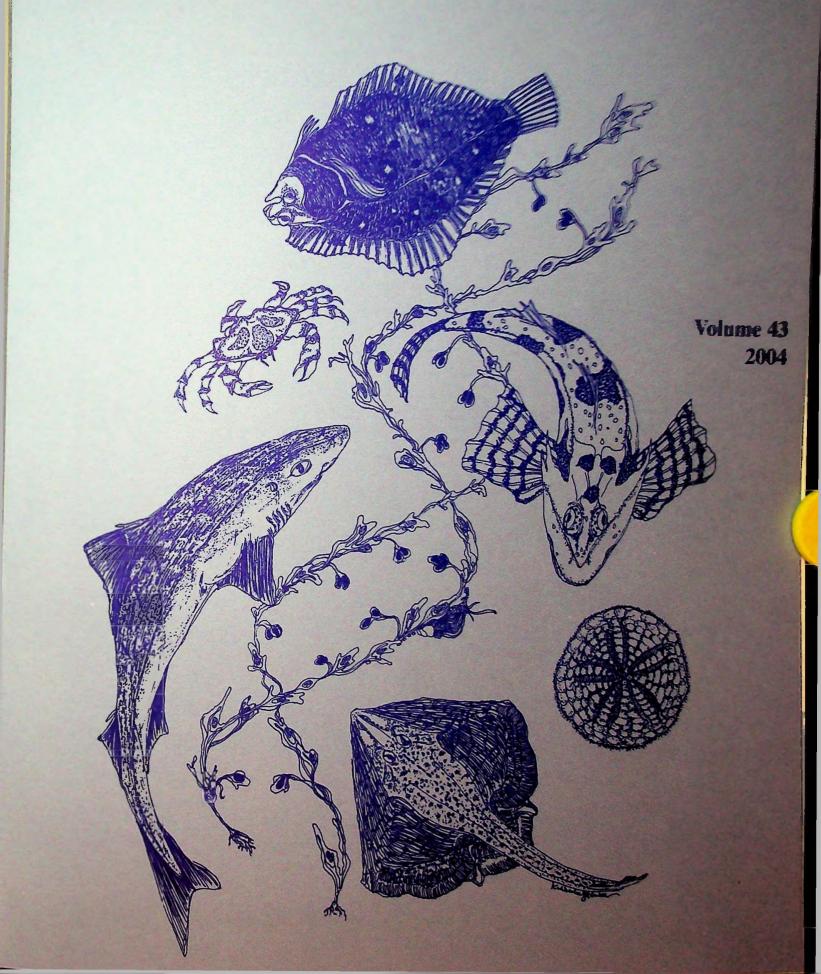
# THE BULLETIN Mount Desert Island Biological Laboratory



## THE 2004 BULLETIN EDITORIAL COMMITTEE

Editor Managing Editor Dr. J.B. Claiborne Michael P. McKernan

Dr. J.B. Claiborne, Chair

Dr. David Barnes

Dr. Raymond Frizzell

Dr. John Henson

Dr. Harmut Hentschel

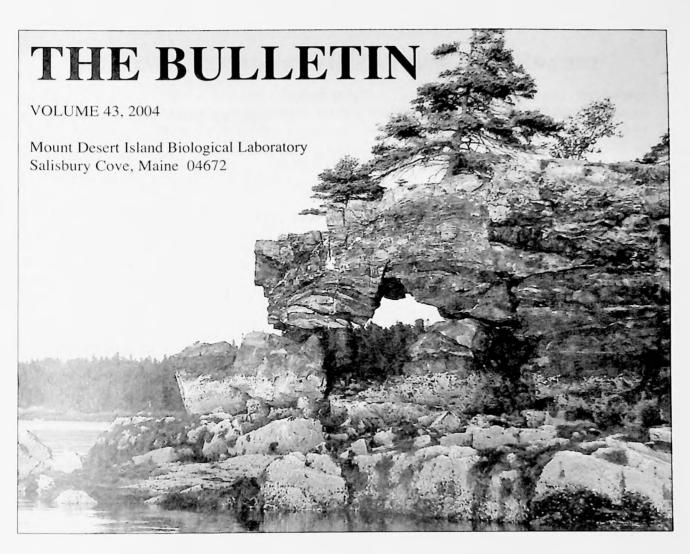
Dr. Karl Karnaky

Dr. David Miller

Dr. Robert L. Preston

Dr. Alice Villalobos

Published by the Mount Desert Island Biological Laboratory July 2004 \$10.00



## TABLE OF CONTENTS

Introduction	ii
Report Titles	vi
Reports	1-144
Officers and Trustees	145
Scientific Personnel	148
Summer Fellowship Recipients	155
Seminars, Workshops, Conferences, Courses	159
Publications	167
Author Index	171
Species Index	174
Keyword Index	175

## THE MOUNT DESERT ISLAND BIOLOGICAL LABORATORY

#### **RESEARCH AND EDUCATION IN THE BIOLOGY OF MARINE ANIMALS**

#### **INTRODUCTION**

The Mount Desert Island Biological Laboratory (MDIBL) is an independent, non-profit marine and biomedical research facility and international center for comparative physiology, toxicology and marine functional genomic studies. The Laboratory is located on the north shore of Mount Desert Island, overlooking the gulf of Maine about 120 miles northeast of the Portland near the mouth of the Bay of Fundy. The island, well known for Acadia National Park, provides a variety of habitats including shallow and deep saltwater, a broad intertidal zone, saltwater and freshwater marshes, freshwater lakes and streams, forests and meadows.

The Laboratory is among the oldest cold water research facilities in the Eastern United States, and its unique site provides an outstanding environment for studying the physiology of marine and freshwater flora and fauna. During 2003, the scientific personnel included 82 doctoral level scientists (including 55 Investigators), plus 137 students, and technical staff, representing 81 institutions in 27 states, Australia, Europe, and South America.

#### HISTORY AND ORGANIZATION

MDIBL was founded in 1898 at South Harpswell, Maine by J.S. Kingsley of Tufts University. Its present site at Salisbury Cove was donated by the Wild Gardens of Acadia, and relocation was completed in 1921. The Wild Gardens of Acadia, a land-holding group headed by George B. Dorr and John D. Rockefeller, Jr., who was instrumental in the founding of Acadia National Park.

In 1914, the Laboratory was incorporated under the laws of the State of Maine as a non-profit scientific and educational institution. Founded as a teaching laboratory, MDIBL is now a center for marine research and education that attracts investigators and students from across the U.S. and around the world. Since the pioneering work of H.W. Smith, E.K. Marshall and Roy P. Forster on various aspects of renal and osmoregulatory physiology of local fauna, the Laboratory has become known worldwide as a center for investigations in electrolyte and transport physiology, developmental biology, electrophysiology and marine molecular biology.

The Mount Desert Island Biological Laboratory is owned and operated by the Board of Trustees and Members of the Corporation; at present, there are 378 members. Officers of the Corporation - Chair, Vice-Chair, Director, Secretary, Treasurer, Clerk - and an Executive Committee are elected from among the Trustees. The Chair and Executive Committee oversee and promote long range goals of the Laboratory. The Director, with the aid of a full-time Administrative Director, staff and a Scientific Advisory Committee is responsible for implementing the scientific, educational and public service activities of the Laboratory.

#### NIEHS CENTER FOR MEMBRANE TOXICITY STUDIES

The Center for Membrane Toxicity Studies (CMTS), an NIEHS Marine and Freshwater Biomedical Sciences Center was established at the Mount Desert Island Biological Laboratory (MDIBL) in 1985. The purpose of this Center has been to involve a group of internationally recognized investigators, who are primarily experts in mechanisms of epithelial transport, to study the biological effects of environmental pollutants on cell and membrane transport functions. The primary emphasis of this research effort has been to elucidate the mechanisms of toxicity of environmental pollutants at the cellular and molecular level, using novel aquatic models developed at this laboratory.

The focus of the research programs of the Center has broadened in the last several years from the more narrow objective of identifying the molecular targets for the effects of heavy metals (or metal compounds) on cell functions, to include the effects of a broader range of environmental toxicants (including marine toxins) and the mechanisms by which the organism takes up and eliminates a wide range of xenobiotics and environmental pollutants. However, the concept that a "membrane lesion" accounts for the cellular toxicity of many environmental toxins still remains as a paradigm.

**Research Cores:** The Center consists of two highly integrated research cores or themes consisting of:

- Signal Transduction and Ion Transport
- Xenobiotic Transport and Excretion

Investigators in the Signal Transduction and Ion Transport Core are examining the basic mechanisms concerning the cell's signaling response to changes in its external environment, particularly as related to environmental stress, heavy metal exposure, marine toxins and environmental estrogens. These signaling pathways often involve mechanisms of homeostasis of ion transport, pH and cell volume regulation. Investigators in the Core are interested in determining the fundamental mechanisms by which cells regulate their cell volume, maintain internal pH and secretory functions and how these processes are disturbed by environmental influences. Investigators in the Xenobiotic Transport and Excretion Core are examining the processes that are used by various epithelial tissues such as the liver and kidney to take up and excrete drugs and xenobiotics and other toxic compounds that enter from the environment and to study the effects of toxicants on this process. Investigators in the Signal Transduction and Ion Transport Core.

Facilities Cores: The Center provides for five facility cores for Center investigators. These include:

• an Animal Core that is responsible for the acquisition, and maintenance of the many marine species available to investigators at this Center;

• an Instrumentation and Facilities Core that maintains the basic laboratory equipment that investigators would not otherwise be able to easily bring to the laboratory (a fully equipped cell

culture and molecular biology facility, Marine DNA Sequencing Center, and an electrophysiology facility);

• a Cell Isolation, Culture and Organ Perfusion Core that provides isolated cells and tissues from marine species to Center investigators;

• an Imaging Core that maintains and operates a confocal fluorescent microscope as well as providing other imaging technology including epifluorescence and video-enhanced microscopy;

• a Bioinformatics Core that is the site of development of a national Comparative Toxicogenomics Database and webpage design. This core incorporates molecular data on marine sequences with a highly annotated database and provides comparative information with human genes of toxicologic interest.

All Center members and pilot recipients have free access to these core facilities. Non-Center members who utilize these facilities are charged appropriate fees.

Community Outreach and Education Program: The Center's outreach program involves community education on water monitoring programs. This is directed primarily at high school and college students in the immediate area of the laboratory. However, an extensive summer research educational program includes high school students from both regional and national sites, the latter emphasizing minority student education as well as college and postdoctoral fellowship training.

Pilot Projects: The Pilot Project Program provides support for investigators who are interested in pursuing a new project related to environmental toxicology in one or more of the Center's Research Cores. The purpose of these Pilot grants is to obtain preliminary data to facilitate new grant submissions. Grants are awarded competitively and successful applicants receive up to \$10,000/season.

#### **APPLICATIONS AND FELLOWSHIPS**

Research space is available for the entire summer season (June 1 - September 30) or a halfseason (June 1 - July 31 or August 1 - September 30). Applications for the coming summer must be submitted by February 1st each year. Investigators are invited to use the year-round facilities at other times of the year, but such plans should include prior consultation with the *MDIBL* office concerning available facilities and specimen supply.

A number of fellowships and scholarships are available to research scientists, undergraduate faculty and students, and high school students. These funds may be used to cover the cost of laboratory rent, housing and supplies. Stipends are granted with many of the student awards. Applicants for fellowships for the coming summer research period are generally due in January.

For further information on research fellowships, please contact:

Dr. Patricia H. Hand Administrative Director Mount Desert Island Biological Laboratory P.O. Box 35 Salisbury Cove, Maine 04672 Tel. (207) 288-3605 Fax. (207) 288-2130 phand@mdibl.org

Students should contact:

Michael McKernan Director of Education and Conferences mmckernan@mdibl.org

#### ACKNOWLEDGEMENTS

The Mount Desert Island Biological Laboratory is indebted to the National Institutes of Health and National Science Foundation and for substantial support. Funds for building renovations and new construction continue to permit the Laboratory to expand and upgrade its research and teaching facilities. Individual research projects served by the Laboratory are funded by private and government agencies, and all of these projects have benefited from the NSF and NIH grants to the Laboratory. For supporting our educational initiative, MDIBL acknowledges the National Science Foundation Research Experience for Undergraduates, Maine Biomedical Research Infrastructure Network (NCRR/NIH), Cserr/Grass Foundation, Milbury Fellowship Fund, Northeast Affiliate of the American Heart Association, Cystic Fibrosis Foundation, Blum/Halsey Fellowship, Stanley Bradley Fund, Stan and Judy Fund, Adrian Hogben Fund, Bodil Schmidt-Nielsen Fellowship Fund, Maine Community Foundation, the Hearst Foundation, the Betterment Fund and many local businesses and individuals.

#### **REPORT TITLES**

Reports preceded by an asterisk were prepared by investigators funded by the NIEHS Center for Membrane Toxicity Studies at the Mount Desert Island Biological Laboratory

### **Invited Review**

## **Ionic Regulation**

*Spanings-Pierrot, C., & D.W. Towle. Salinity-related mRNA expression of Na <sup>+</sup> /K <sup>+</sup> /2Cl <sup>-</sup> cotransporter and V-type H <sup>+</sup> -ATPase in gills of the euryhaline crab <i>Pachygrapsus marmoratus</i>
Pedersen, S.F., King, S.A., Holt, M. & P.M. Cala. Signal transduction mechanisms involved in the regulation of NHE1 from <i>Pseudopleuronectes americanus</i> red blood cells by osmotic shrinkage, b-adrenergic stimuli, and calyculin A
*Pelis, R., Edwards, S., Claiborne, J. & L. Renfro. pH-dependent sulfate secretion by the renal proximal tubule of <i>Pleuronectes americanus</i>
Epstein, F.H., Sighinolfi, C., Hessler, K., Verkman, A.S., Riordan, J.R. & P. Silva. A synthetic CFTR inhibitor blocks stimulation of rectal gland secretion by CNP and VIP
Silva, P., Sighinolfi, C., Hessler, K., Spokes, K., Hays, R.M. & F.H. Epstein. Sildenafil citrate enhances the stimulation of the secretion of chloride by CNP in <i>Squalus acanthias</i> rectal gland 15
Silva, P., Sighinolfi, C., Hessler, K., Spokes, K., Hays, R.M. & F.H. Epstein. TMAO has no effect on the secretion of chloride or its stimulation by VIP or CNP in <i>Squalus acanthias</i> rectal gland 17
*Scharlau, D., Althoff, T., Hentschel, H. & R.K.H. Kinne. Immunohistochemical studies of Na <sup>+</sup> /D-glucose cotransporters in the intestine and kidney of <i>Squalus acanthias</i> and <i>LeucoRaja erinacea</i>
Catches, J.S. & J.B. Claiborne. NHE2 and Na <sup>+</sup> /K <sup>+</sup> -ATPase immunoreactivity in <i>Myoxocephalus</i> octodecimspinosus
Catches, J.S., Burns, J.M. & J.B. Claiborne. Na <sup>+</sup> /K <sup>+</sup> /2Cl <sup>-</sup> -cotransporter immunoreactivity in <i>Myoxocephalus octodecimspinosus</i>
*Preston, R., Clifford, R., Thompson, J., Slager, D., Petersen, C. & G. Kidder. CFTR mRNA expression in developing <i>Fundulus heteroclitus</i> embryos
*Bewley, M., Decker, S., Klein, C., Ratner, M., Kelley, C., Burks, K., Motley, W., Peters, A. & J.N. Forrest. The CFTR inhibitor-172 has minimal effects on shark CFTR as compared to human CFTR
*Decker, S., Klein, C., Ratner, M., Kelley, C., Epstein, M., Burks, K., Motley, W., Peters, A. & J.N. Forrest. Effects of quinidine and other $K^+$ channel inhibitors on chloride secretion in the rectal gland of the spiny dogfish, <i>Squalus acanthias</i>
Rigor, R.R., Zhuang, Z. & P.M. Cala. Regulatory volume decrease by <i>Pseudopleuronectes</i> americanus red blood cells: the nature of the potassium flux pathway

## **Comparative Biochemistry & Molecular Biology**

Fellner, S. & L. Parker. Endothelin B Ca2 <sup>+</sup> signaling in <i>Squalus acanthias</i> vascular smooth muscle: participation of IP3 and ryanodine receptors
*Roer, R. & D. Towle. Partial nucleotide sequence of a putative cuticular hexosaminidase from the blue crab, <i>Callinectes sapidus</i>
*Campbell, J.D., Jensen, T.J., Scarlet, C., Borchers, C., Rosenberg, M.F., Ford, R.C. & J.R. Riordan. Immunoisolation of CFTR from shark rectal gland for proteomic and structural studies43
Rose, R., Choe, K.P. & D.H. Evans. Sequencing of a putative COX-2 from the killifish ( <i>Fundulus heteroclitus</i> ) gill
*Sato, J.D., Chapline, M.C., Alestorm, P., Fan, L. & P. Collodi. Receptor kinase expression in germ-line competent zebrafish ( <i>D. rerio</i> ) ES cells
*Henson, J.H., Davis, J.E., Huang, K. & R.F. Murphy. Computer-based recognition of drug- induced changes in the distribution of fluorescently labled microtubules in NIH 3T3 cells
*Straub, P.F., Higham, M.L., Phoel, W.C. & B. J. Landau. Functional genomics and quantitative PCR for the study of environmentally related gene expression in <i>Pseudopleuronectes americanus</i> 51
*Sato, J.D., Sun, L., Yu, Y., Herley, M.T. & C.M. Chapline. Variable region characterization of monoclonal antibodies to the VEGF receptor Flt-1
Crockett, E.L., Hassett, R.P., Funk, K.R. & C.M. Doering. Activities of Na <sup>+</sup> /K <sup>+</sup> -ATPase and cholesterol contents are not altered by dietary supplementation with cholesterol in <i>Acartia</i> hudsonica and Calanus finmarchicus
*Fan, L., Crodian, J., Parton, A., Collodi, P. & P. Alestrom. Use of marine cell lines as a source of factors that maintain pluripotency of zebrafish ( <i>Danio rerio</i> ) ES cell cultures
*Bayne, C. & A. Parton. Derivation of cell lines from <i>Strongylocentrotus droebachiensis</i> , the Northern sea urchin
*Elmore, L.W., Parton, A., Barnes, D.A. & S.E. Holt. A novel role for telomerase in fish
*Gaskins, H.R., McCray, N.R., Collier, C.T., King, D.E., Thurmond, J.E. & R.I. Mackie. Molecular ecological and phylogenetic analyses of the intestinal microbiota of the ascidian tunicates <i>Boltenia echinata</i> , <i>B. ovifera</i> , <i>Halocynthia pyriformis</i> and <i>Ciona intestinalis</i>
Thomason, K., Towle, D. & R. Henry. Quantitative expression of carbonic anhydrase mRNA and protein-specific activity in the gills of the euryhaline green crab, <i>Carcinus maenas</i> , during low salinity acclimation
*Crane, M., Wong, S., Tilden, A. & D. Towle. Identification of Na <sup>+</sup> /Cl <sup>-</sup> -dependent neurotransmitter transporters in the lobster, <i>Homarus americanus</i>
*Stoller, J., Ryder, P., Gitler, A., Vosburgh, B. & J. Epstein. Gene knockdown using morpholino nucleotides in zebrafish ( <i>Danio rerio</i> ) to investigate the function of Nf1 and FoxP4 in cardiovascular development

Blakaj, A., Plattus, R.B., Berliner, N. & E.J. Benz, Jr.	Cloning of the 5' end of Na, K-ATPase
cDNA in the rectal gland of the Squalus acanthias	
Day, R.M., Nagase, H., Lee, Y.H., Kraev, A., Cleemann	n, L. & M. Morad. Sequencing the cardiac

## **Comparative Physiology**

*Pelis, R. & L. Renfro. Feeding and PKC activation inhibit sulfate secretion by the intestine of <i>Pleuronectes americanus</i>
*Knickelbein, R., Boyer, J.L. & N. Ballatori. Glutathione export from skate ( <i>Raja erinacea</i> ) liver is unaffected by taurocholic acid
Hyndman, K.A. & D.H. Evans. Immunolocalization of the endothelin receptors in the gill of the longhorn sculpin, <i>Myoxocephalus octodecimspinosus</i>
Giesbrandt, K., Choe, K.P. & D.H. Evans. The cardiovascular and branchial perfusion effects of endothelin in the longhorn sculpin ( <i>Myoxocephalus octodecimspinous</i> )
Crane, M., Hand, E., LeBlanc, J., Pfeiffer, A., Shanahan, K. & A. Tilden. Influence of tides, exercise, and melatonin on hemolymph glucose and lactate levels in the fiddler crab <i>Uca pugilator</i> 90
Litteral, J., Kirsch, T., Wortmann, J., Beese, M., Borley, K., Hentschel, H., Haller, H. & M. Elger. Evaluating mechanisms of nephrogenesis on tissue cultures from adult Little Skate, <i>Leucoraja</i> erinacea kidney
Swenson, K.E., Eveland, R.L, Reiter, C., Gladwin, M.T. & E.R. Swenson. Nitric oxide (NO) in vascular regulation of the spiny dogfish, <i>Squalus acanthias</i>
Long, J., Koob-Emunds, M. & T. Koob. The mechanical consequences of vertebral centra
Koob, T.J., Dean, M.N. & M.M. Koob-Emunds. Extracellular matrix compisition in the rostral gel of <i>Squalus acanthias</i> : A non-cartilaginous skeletal element
Freiji, A. & J.B. Claiborne. mRNA for the NHE2 exchanger is expressed in a variety of epithelial tissues in the spiny dogfish, <i>Squalus acanthias</i>
Lanier, C. & J.B. Claiborne. Analysis of branchial longhorn sculpin ( <i>Myoxocephalus octodecimspinosus</i> ) Na <sup>+</sup> /H <sup>+</sup> exchanger isoform 3 using Northern hybridization
Smith, K. & R.P. Henry. A carbonic anhydrase repressor is found in the hemolymph of the euryhaline green crab, <i>Carcinus maenas</i>
Baldwin, J., Goldsmith, C., Petersen, C., Preston, R. & G. Kidder. Synchronous hatching in <i>Fundulus heteroclitus</i> embryos: Production and properties
Mohebbi, N., Wellner, M., Herold, D. & M. Gollasch. Structure and function of the dicarboxylate transporter INDY
*Salinas, S., Brandvain, Y.J., Anderson, R., Marty, J., R.L. Preston, G.W. Kidder & C.W. Petersen. Reproductive ecology of <i>Fundulus heteroclitus</i> and <i>Fundulus diaphanus</i> in a New England watershed
Burdick, D., Hartline, D.K. & P.H. Lenz. Quantitative differences in escape responses to hydrodynamic stimuli by calanoid copepods

Henry, R. Initial characterization of a carbonic anhydrase repressor from the eyestalks of the euryhaline green crab, <i>Carcinus maenas</i>
*Althoff, T., Scharlau, D., Luig, J., Schuetz, H. & R.K.H. Kinne. Sodium-D-glucose transport in <i>Squalus acanthias</i> and <i>Leucoraja erinacea</i> : an update and new perspectives
*Hill, W.G., Mathai, J.C., Gensure, R.H., Zeidel, J.D., Saenz, J.P., Kinne-Saffran, E., Kinne, R. & M.L. Zeidel. Barrier function of teleost and elasmobranch gill apical membranes

## Molecular Toxicology & Xenobiotic Transport

- 2

*Sato, J.D., Coutermarsh, B., Chapline, M.C. & B.A. Stanton. Effects of arsenic, an inhibitor of CFTR function, on intracellular signaling in <i>Fundulus</i>
*Park, G., Knickelbein, R., Boyer, J.L. & N. Ballatori. The liver X receptor-alpha, LXRa, is expressed in the liver of the little skate, <i>Raja erinacea</i>
*Gordon, A., Renfro, J. L., Miller, D.S. & A.R.V. Villalobos. Effects of thermal stress on xenobiotic transport by <i>Squalus acanthias</i> choroid plexus
*Chen, C. & B. Mayes. Trophic transfer of metals in estuarine food webs
*Shaw, J.R., Curtis-Burnes, J., Stanton, B.A. & J.W. Hamilton. The toxicity of arsenic to the killifish, <i>Fundulus heteroclitus</i> : Effects of salinity
*Notenboom, S., Masereeuw, R., Russel, F. & D. Miller. Upregulation of multidrug resistance- associated protein (Mrp2) in renal proximal tubules from killifish, <i>Fundulus heteroclitus</i>
*Baehr, C., DiPasquale, K., Fricker, G. & D.S. Miller. Fluorescein-methotrexate (FL-MTX) transport in dogfish shark, <i>Squalus acanthias</i> , choroid plexus
*Decker, S., Klein, C., Ratner, M., Kelley, C., Epstein, M., Burks, K., Motley, W., Peters, A., Stanton, B.A. & J.N. Forrest. Arsenic inhibits chloride secretion in the perfused rectal gland and cultured tubularepithelial cells of the dogfish shark, <i>Squalus acanthias</i>
*Parton, A., Dowell L., Rafferty, J., Forrest, J.N., Boyer, J.L. & D. Barnes. Culture of marine elasmobranch cells in vitro
*Mayer, G.D., Berry, J.P., Patenaude, C.A. & P.J. Walsh. Effect of lipopolysaccharides from <i>Microcystis</i> and <i>Lyngbya</i> on metal toxicity in <i>Fundulus heteroclitus</i>