

NOAA Technical Memorandum NMFS



SEPTEMBER 1992

THE KEWALO RESEARCH FACILITY, 1958-92: OVER 30 YEARS OF PROGRESS

Richard W. Brill
(compiler and editor)

NOAA-TM-NMFS-SWFSC-171

U.S. DEPARTMENT OF COMMERCE
National Oceanic and Atmospheric Administration
National Marine Fisheries Service
Southwest Fisheries Science Center



NOAA Technical Memorandum NMFS

This TM series is used for documentation and timely communication of preliminary results, interim reports, or special purpose information; and have not received complete formal review, editorial control, or detailed editing.

SEPTEMBER 1992

THE KEWALO RESEARCH FACILITY, 1958-92: OVER 30 YEARS OF PROGRESS

Richard W. Brill
(compiler and editor)

Honolulu Laboratory, SWFSC
National Marine Fisheries Service
2570 Dole Street
Honolulu, Hawaii 96822-2396

NOAA-TM-NMFS-SWFSC-171

U.S. DEPARTMENT OF COMMERCE

Barbara H. Franklin, Secretary

National Oceanic and Atmospheric Administration

John A. Knauss, Under Secretary for Oceans and Atmosphere

National Marine Fisheries Service

William W. Fox, Jr., Assistant Administrator for Fisheries

CONTENTS

	Page
The Kewalo Research Facility	1
Research Activities	2
The Early Years	2
The Later Years	10
Tunas	10
Thermoregulation	10
Energetics	10
Spawning and rearing	13
Geomagnetic sensitivity	13
Lobsters	15
The Kewalo Research Facility Today	16
Tuna Research	16
Olfaction	16
Research to Determine Energy and Habitat Requirements of Tunas	19
Tracking Tuna Movements	20
Burnt Tuna	21
Spawning and Rearing	21
Protected Species	25
Hawaiian Monk Seals	25
Hawaiian Sea Turtles	27
Relationship with the University of Hawaii	28
List of Visiting Investigators	29
Partial List of Scientific Publications Resulting from Research at the Kewalo Research Facility	32

THE KEWALO RESEARCH FACILITY

Kewalo can be translated from the Hawaiian as "the place of wailing." Historical descriptions of the area called Kewalo on the island of Oahu give meaning to the translation. In ancient times, this section of land contained a spring which was used as a place for human sacrifices. Here *kauwa* (outcasts) were first drowned before being taken to the *Heiau of Kanelaau* (temple) on the slopes of Punchbowl Crater for burning in the *imu ahi* (fire oven). Kewalo Basin, as part of the modern city of Honolulu, is of course no longer used for such purposes. Today it is the home of many commercial and recreational fishing boats, tour boats, a fresh fish auction house, and other marine-related enterprises. Kewalo Basin is also the site of the Honolulu Laboratory's **Kewalo Research Facility**.

The area occupied by the Kewalo Research Facility was once a shallow, submerged coral reef. In 1945, the U.S. Navy dredged a small harbor, which became known as Kewalo Basin and was later turned over to the Territory of Hawaii. The harbor was subsequently enlarged, and artificial and sanitary fill was dumped on the adjacent coral reef to create protective land areas. In July 1958, the Honolulu Laboratory of the National Marine Fisheries Service, then a part of the U.S. Fish and Wildlife Service, negotiated a lease to the grounds and building on the spit of artificial land created at the southeast entrance of Kewalo Basin and established the Kewalo Research Facility.



An aerial view of the Kewalo Research Facility in Honolulu, Hawaii.

The facility has a low profile and goes unnoticed by the many tourists, surfers, and fishermen that frequent the area. But within the 0.4 hectare (0.98 acre) area is a unique research laboratory. The main building houses offices, laboratories tailored for various research activities, a machine shop, and storage areas. A saltwater well on the adjoining grounds has the capacity to produce high-quality, coral-filtered seawater at a rate of over 3,785 liters (1,000 gallons) per hour. The seawater is pumped to aerators to be oxygenated and then distributed to various tanks, including a series of five 75,706-liter (20,000-gallon) circular pools, a 757,060-liter (200,000-gallon) oceanarium, and specially designed experimental tanks of various sizes.

The Kewalo Research Facility is today, as it has been since its inception, the only research center in the world capable of maintaining live tunas in captivity throughout the year for use in behavioral and physiological research. The uniqueness of this facility and the past 30 years of quality research have engendered it an enviable international reputation. Indeed, its reputation continues to attract established scientists of diverse backgrounds and expertise to this unique laboratory where experiments that require live tunas and other marine animals can be conducted.

In recent years, the role of the Kewalo Research Facility has expanded. It has served the research needs of scientists charged with the responsibility of enhancing the survival of threatened and endangered species, such as the Hawaiian monk seal (*Monachus schauinslandi*) and green sea turtle (*Chelonia mydas*). The Facility now has a fully functional larval culture laboratory which has successfully reared, from eggs, species as diverse as mahimahi (*Coryphaena hippurus*), spiny lobster (*Panulirus marginatus*), and deepwater shrimp (*Heterocarpus* spp.).

RESEARCH ACTIVITIES

The Early Years

Tuna stocks are distributed throughout the world's oceans and form an important economic resource for many countries. The value of worldwide tuna catches is currently estimated at close to \$4 billion each year. The United States alone consumes over 600 million pounds of canned tuna per annum, valued at approximately \$1 billion. Despite the high economic value of tuna stocks, very little research had been done with live specimens before 1958, because no facility existed to maintain tunas in captivity. Analyses of fisheries data showed correlations between the apparent abundance of tunas and various oceanographic and meteorological conditions, but the mechanisms that determine the horizontal and vertical distributions, migrations, and vulnerability of tunas to specific types of fishing gear were completely unknown. Temperature, oxygen, and salinity all seemed to exert an influence. Predicting the changing abundance and gear vulnerability of the various tuna species that comprise the resource was obviously a major biological problem and impediment to effective management.

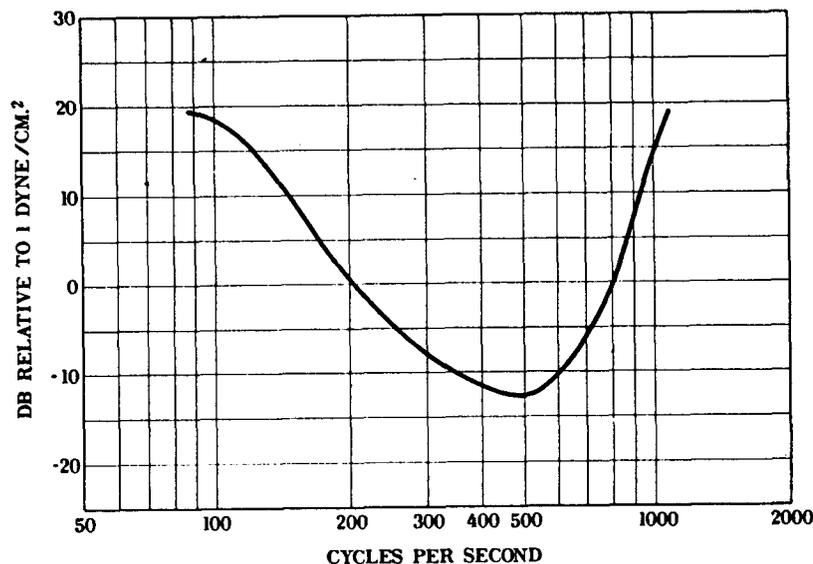
The initial goal of the Kewalo Research Facility was the development of procedures for keeping live tunas in captivity for experimentation. Because this was truly a pioneering effort, early research was aimed at collecting data that would serve as the foundation for future investigations. This early work uncovered several interesting facts about tunas:

- Tunas are heavier than water and must continuously swim to keep from sinking.
- Tunas breathe by simply opening their mouths so that water is forced over their gills as they swim; they sink and suffocate if they stop swimming.
- Basal swimming speeds of tunas are dependent upon the lifting area of fins and the density of the fish and are not a function of either respiratory requirements or the search for food.
- All tunas have the following adaptations for continuous swimming: (1) a high hemoglobin level in the blood to carry sufficient oxygen to maintain continuous muscle activity; (2) a large proportion of the muscle composed of red muscle fibers specialized for continuous activity, as are the muscles of the heart; and (3) a streamlined body shape to reduce hydrodynamic drag.
- Larger species of tunas have evolved two morphological features to reduce the energy required to keep from sinking: (1) pectoral fins became larger to produce more lift; and (2) gas bladders developed to decrease density. [Although gas bladders are very effective in reducing fish density, they limit the vertical movements of tunas. A fast, vertical ascent to the surface can cause large changes in volume and, in extreme cases, burst the gas bladder. Small species of tunas, such as skipjack tuna (*Katsuwonus pelamis*), do not have gas bladders.]

Other early experiments were designed to determine the sensory abilities of tunas--how well they smell, taste, hear, see, and sense changes in water temperature. The rationale for these studies was that a basic understanding of the sensory capabilities of tunas would be useful in designing fishing gear, developing new fishing methods, and locating tunas.

To determine how well tunas can see, studies were conducted on their visual acuity (the ability to see the fine details of objects). Of the three species tested, it was determined that yellowfin tuna (*Thunnus albacares*) can see better than skipjack tuna which, in turn, see better than kawakawa (*Euthynnus affinis*). Further experiments on the optical system of restrained tuna showed that they are color-blind and are most sensitive to blue light.

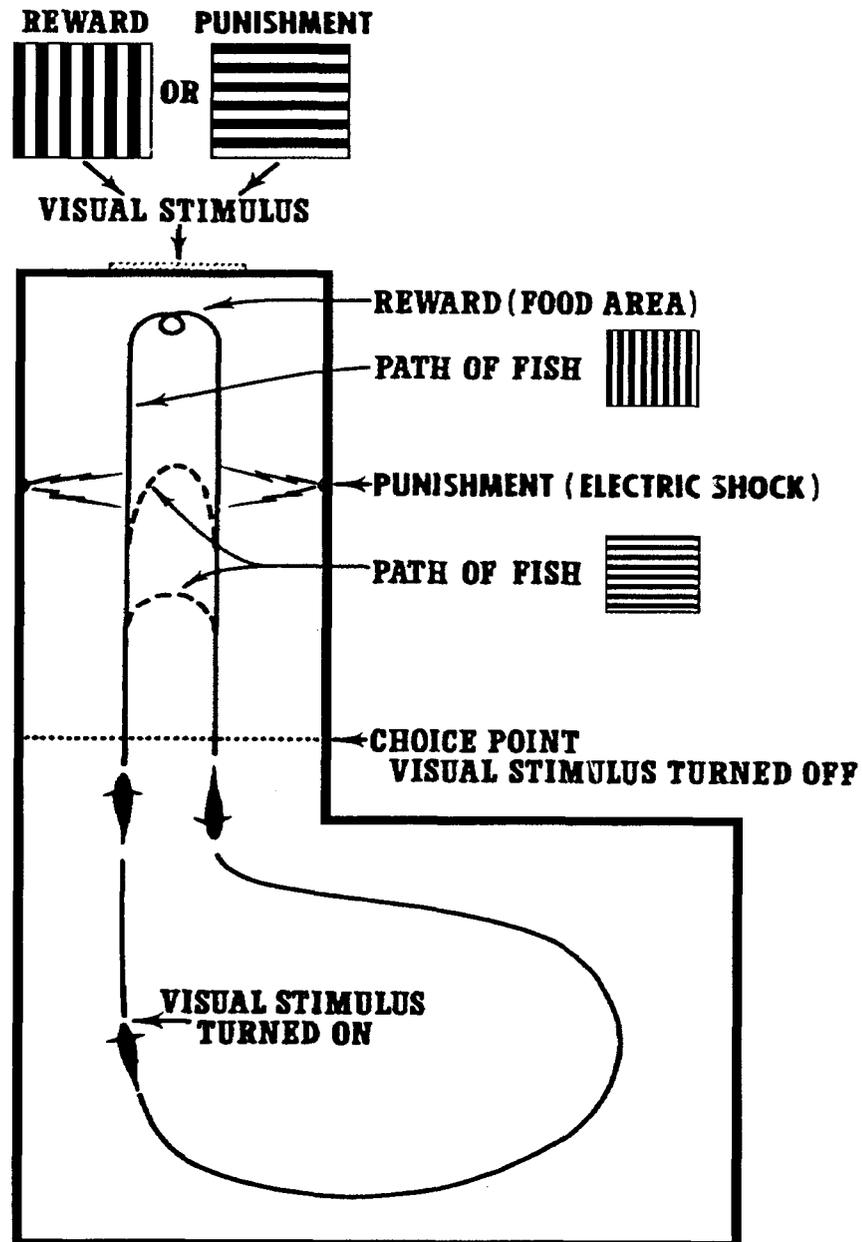
Experiments to define the hearing ability of tunas made it possible to construct a tuna hearing curve--the first ever for a scombrid--and to determine their auditory thresholds (the lowest level of sound that can be heard at a specific frequency). The hearing range of yellowfin tuna is about 200-2,000 hertz (cycles per second), and their hearing is most acute at 500 hertz.



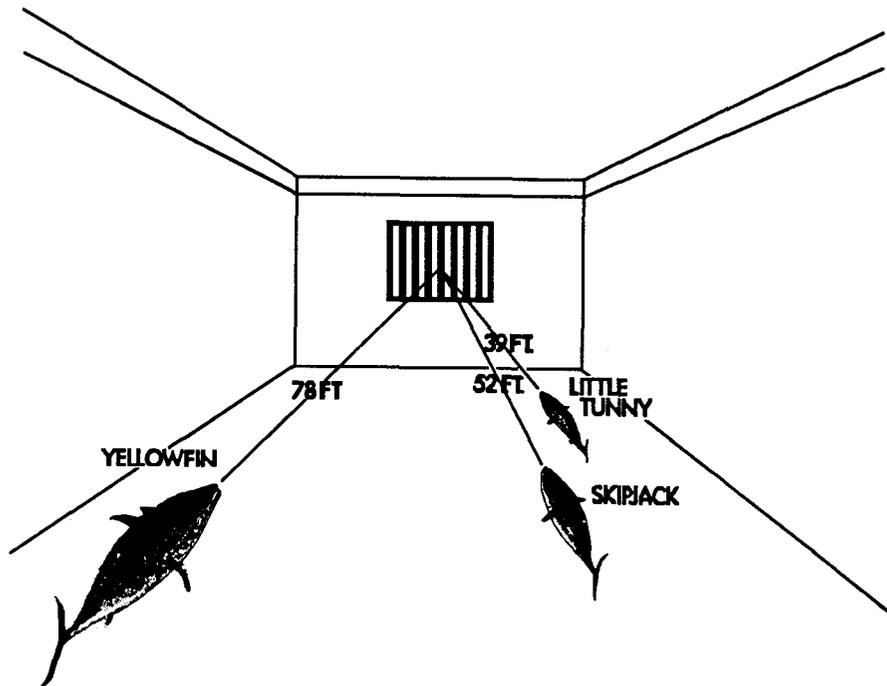
Experiments to determine the hearing ability of tunas were conducted in a pool specially constructed to insulate the fish from outside sounds and electronic interference. The test fish were first trained to recognize a pure "white" sound and then to react to the sound stimulus by swimming through a maze for a reward. The yellowfin tuna best hears sounds that are near 500 Hz, as shown by the dip at that frequency in the hearing curve. Sounds near this frequency (e.g., the sound produced by the swimming of a school of small fish) are common in the ocean.

Experiments at the Kewalo Research Facility also showed that tunas have a highly developed sense of smell. A strong response was elicited from a school of kawakawa when a liter (1.06 quarts) of water in which a small fish (a smelt weighing 10 grams or 0.4 ounce) had been dipped for 10 seconds was introduced to their holding tank. The response was elicited even though the water had been further diluted by its introduction through the inflow seawater system! A study of the morphological structure of the nares (nose) of the tunas revealed that they can "sniff" the water. Each jaw movement of a tuna produces a pumping action that forces water past its nasal rosettes (odor receptors). Observations of fish in captivity showed this pumping action to be continuous.

Two other research projects were designed to determine the ability of tunas to perceive changes in water temperature. One experiment made use of the observation that the heart rate of a restrained tuna slows when the fish is presented with an external stimulus, such as a change in water temperature. In the second experiment, a free-swimming fish was rewarded with food each time it was able to recognize a temperature difference when cooler or warmer water was added to the tank. In restrained fish, a temperature change of 1°C elicited a response. Free-swimming fish were able to do even better; they perceived a temperature difference of as little as 0.1°C.



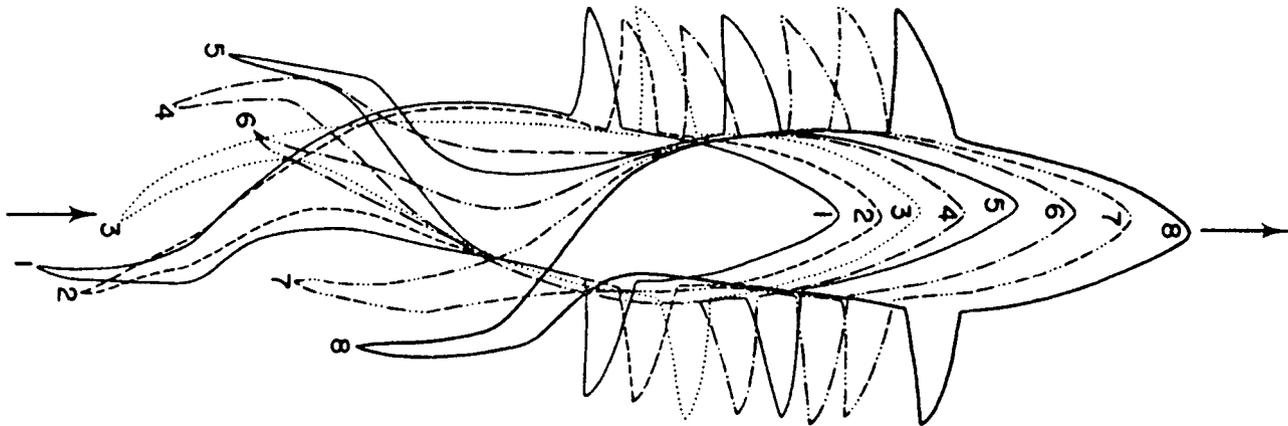
This diagram illustrates the experimental method used to determine the visual acuity of tunas. The method involves training a fish to respond to a visual stimulus (horizontal or vertical stripes), projected on an opal glass plate placed against a tank window. When the stripes are vertical, the fish is trained to swim down the tank to a food-drop area where it is rewarded; when the stripes are horizontal, the fish is trained to turn before reaching the food-drop area and to return to the far end of the tank. If the fish fails to turn, it receives a mild electrical shock.



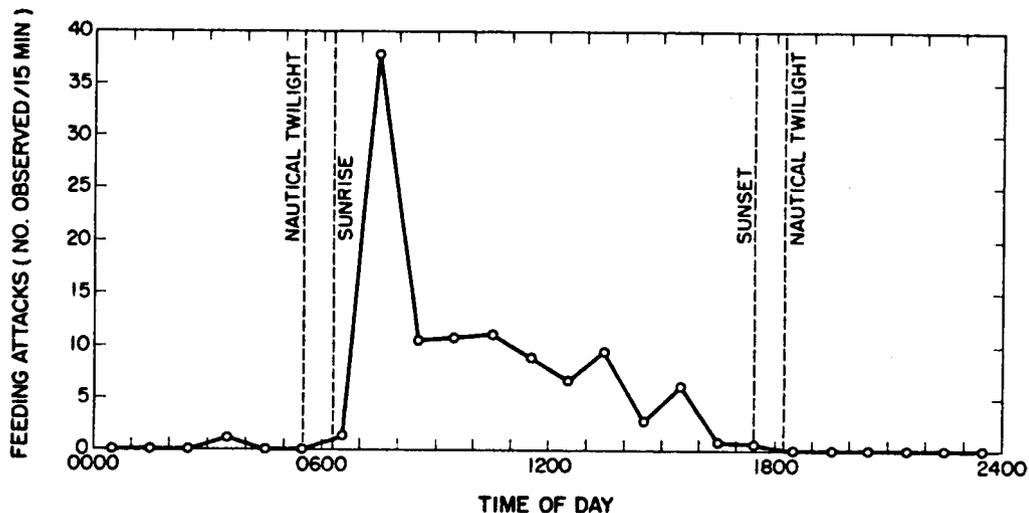
To measure tunas' reactions to various sensory stimuli, an observer must be able to detect the fishes' responses to these stimuli. Tunas can be trained to perform a specific act in response to stimuli if they are rewarded. To measure how well they can see, tunas were trained to respond differently to vertical and horizontal bars, which were projected onto an underwater screen by giving rewards (food) or punishment (electric shock). These experiments showed that at a constant brightness, a yellowfin tuna sees details of an object better than a skipjack tuna which, in turn sees better than a kawakawor a little tunny.

Early work on the feeding and digestion rates showed that tunas can digest a meal several times faster than other fish species. Prey organisms are not homogeneously distributed in the open ocean, but are found in patchy concentrations in space and time. Tunas therefore exist in a "feast or famine" situation and must eat whenever they find food. Knowledge of digestion and feeding rates of fishes adapted to such environments is important for understanding growth and worldwide distribution of tunas and can be of practical value to commercial fishermen.

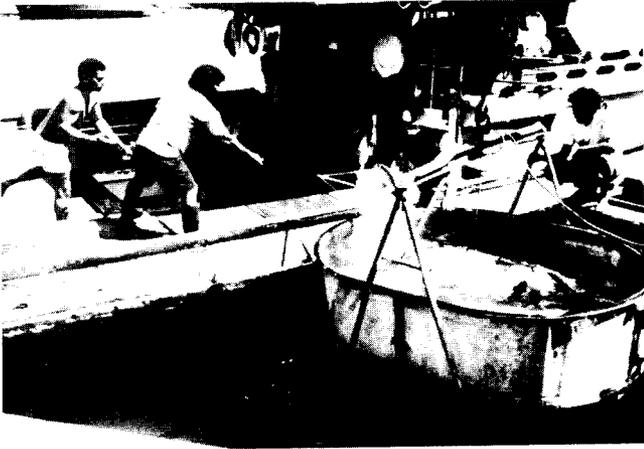
As techniques for capture, transport, and maintenance of tunas improved, the number of live tunas available for experimental purposes increased proportionately. This made it possible to increase the variety of behavioral and physiological studies conducted at the Kewalo Research Facility.



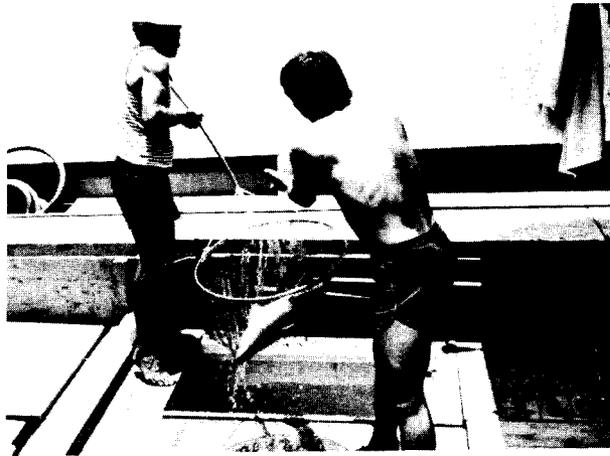
Specialized tanks at the Kewalo Research Facility made it possible to closely observe captive fish and produced the first high-speed movies of swimming kawakawa. The analysis of the film provided intimate details of swimming speed, tail beat rates, body postures and flexures, and how the changing positions of fins and finlets provide drag reduction features. It was determined that the tail provides nearly 100% of the forward thrust and the fish attains burst speeds of twice that of nonscombrid fish. The line drawing shown here was traced from successive cine frames (camera speed, 100 frames per second) for one complete caudal fin beat cycle of a kawakawa. The swimming speed of this fish was 8.2 body lengths per second, which was produced by a tail beat frequency of 14.3 tail beats per second.



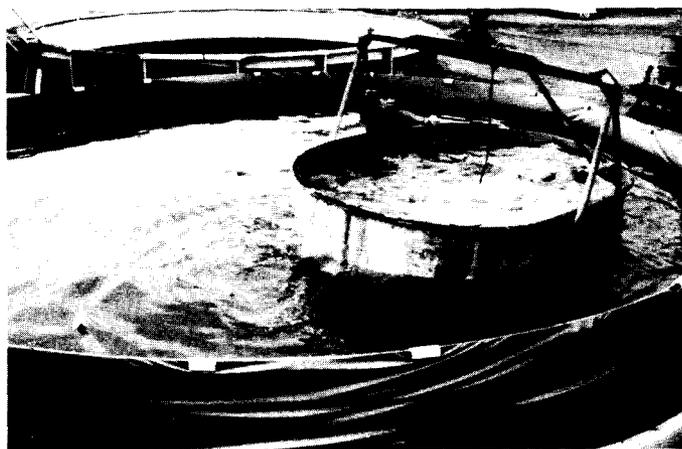
The changes in the feeding activity of kawakawa during a 24-hour period as shown in this graph is typical for all tunas. When tunas in captivity were provided with a constant supply of food, feeding motivation was highest at early morning, followed by a rapid decrease through noon and two smaller peaks at mid-afternoon. They showed no attempts to feed at night. This behavior is consistent with the high consumption and digestion rates which are from two to five times faster than those of other fish. When tunas are fed continuously, an equivalent of 15% of body weight or two times their stomach volume is eaten. The drive to attack prey is dependent on the amount of food in the stomach. Intense feeding always occurs in the morning when the stomach is empty. However, the stomach is not filled to capacity at the first feeding frenzy, and the fish feeds throughout the day. Feeding slows when the stomach is 80% filled.



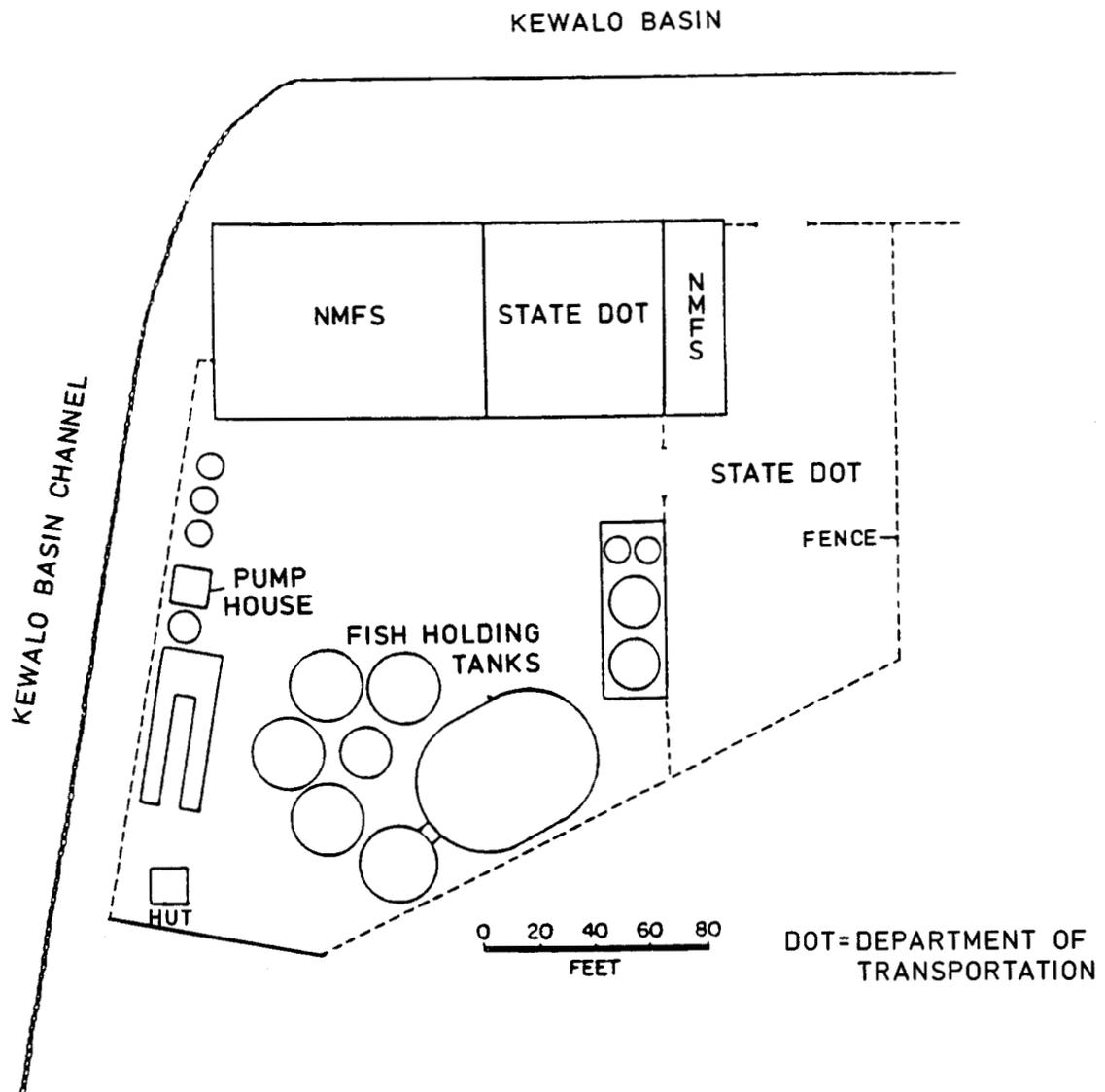
Live tunas are caught with barbless hooks by cooperating commercial pole-and-line fishermen and then dropped into the vessel's bait-holding tanks.



When the vessel returns to Kewalo Basin, the fish are moved to a transfer tank with a chamois-lined dip net.



The transfer tank is then transported by a crane and lowered into a holding tank where the fish can swim freely. This handling technique, which has evolved over the years, minimizes injury to the fish.



The Kewalo Research Facility is able to obtain and maintain tunas in captivity because of several conditions unique to its site. Commercial live-bait tuna fishing boats dock in front of the Facility. Also, because the Facility was built over a filled-in coral reef and Hawaii has a mild climate, the saltwater well is able to provide clean seawater at the appropriate temperature for holding tunas year-round.

The Later Years

Tunas

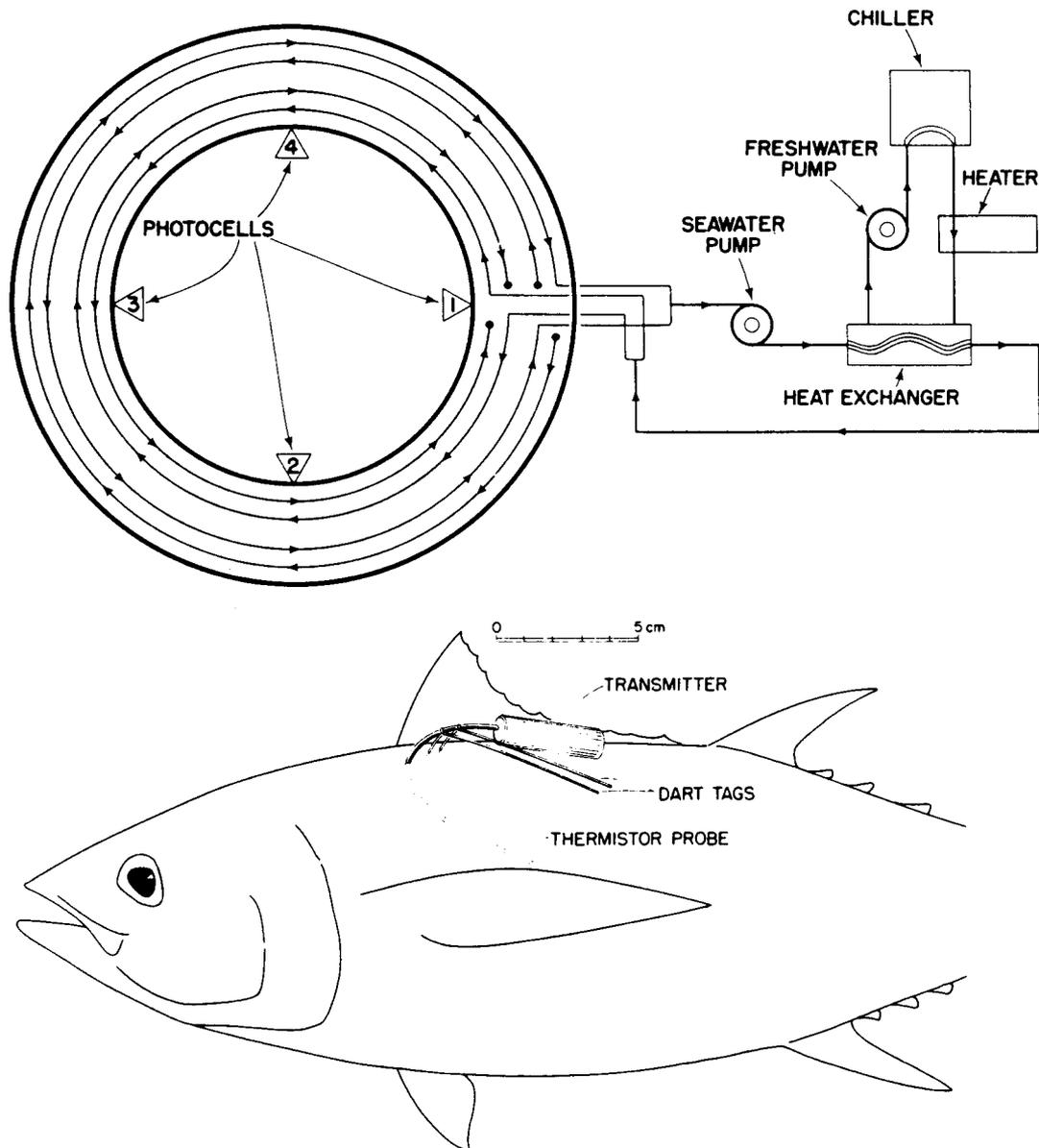
Thermoregulation--Additional studies at the Kewalo Research Facility confirmed that tunas have a remarkable ability to maintain body temperatures higher than the water in which they swim. This ability is attributable to vascular countercurrent heat exchangers that retain the heat produced by metabolic activity within the muscles. In other fishes, metabolic heat is lost to the surrounding water via the gills and body surface. As a predator, the ability to maintain an elevated body temperature probably gives tunas an advantage over other fishes, because it allows them to operate at higher activity levels. Depending on the activity and size of the fish, muscle temperatures of tunas can range from 2° to 21°C above ambient.

The accumulation of knowledge on the effects of temperature on tuna physiology allowed work on more sophisticated experiments, such as those designed to determine whether tunas can physiologically or behaviorally thermoregulate. The first evidence of physiological thermoregulation in tunas was obtained in experiments with yellowfin tuna. Fish, placed in a doughnut-shaped tank, were able to alter their rates of heat loss independently of swimming speed (that is, physiologically thermoregulate) as the water temperature was changed at 12-hour intervals. This ability to physiologically thermoregulate, however, has not yet been demonstrated in all species of tuna.

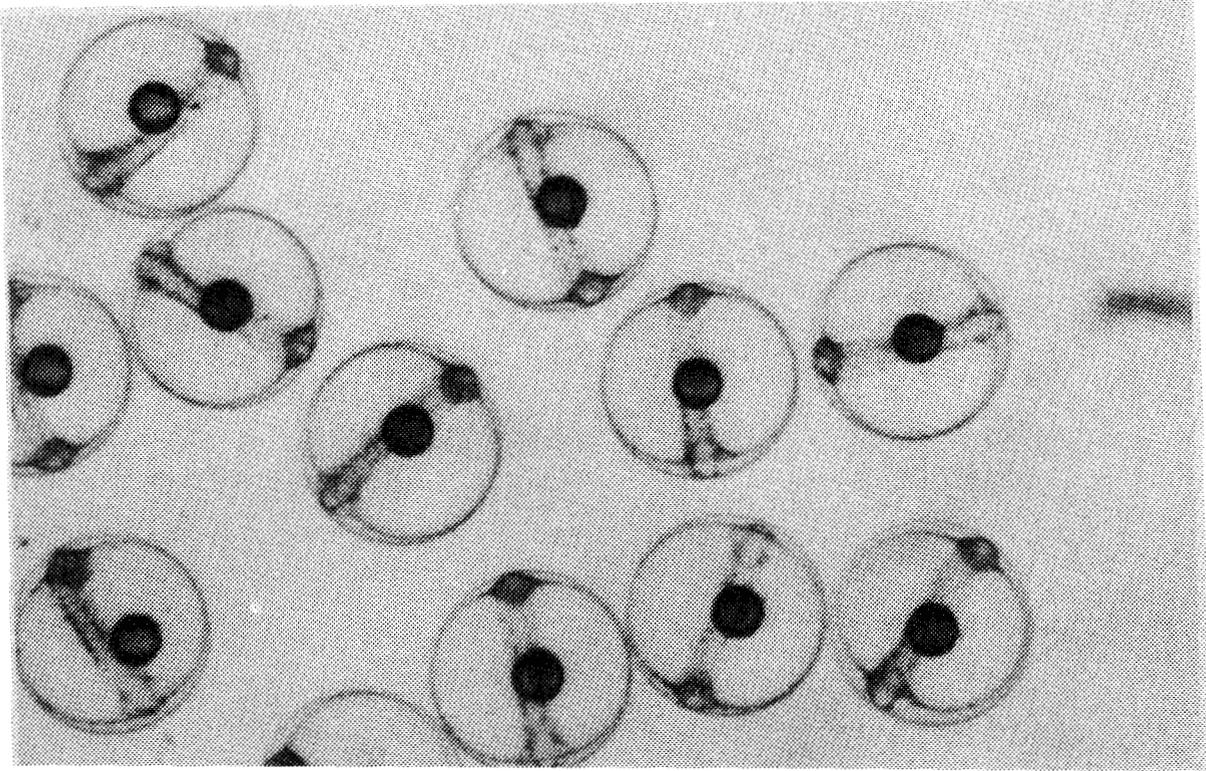
Energetics--Tuna metabolic rates present interesting paradoxes. Tunas have higher energy demands than other fishes, yet they inhabit a very food-poor environment: the tropical oceans. How do tunas obtain energy when they live in a virtual desert? Surprisingly, tunas require more energy to swim at their cruising speed than do other fishes swimming at the same speeds, despite tunas' having a streamlined shape and five swimming fins which can each be withdrawn into a slot or recess leaving the body surface perfectly smooth. Shouldn't tunas be more energy efficient?

The measurement of tuna metabolic rates has a long history at the Kewalo Research Facility. Past projects included measuring standard metabolic rate (metabolic rate at zero activity) and studying the effects of size, temperature, and speed on active metabolic rate. More recent work was designed to reexamine earlier results, which were based on oxygen consumption (respirometry), by directly measuring changes in the energy content (calorimetry) of individual fish.

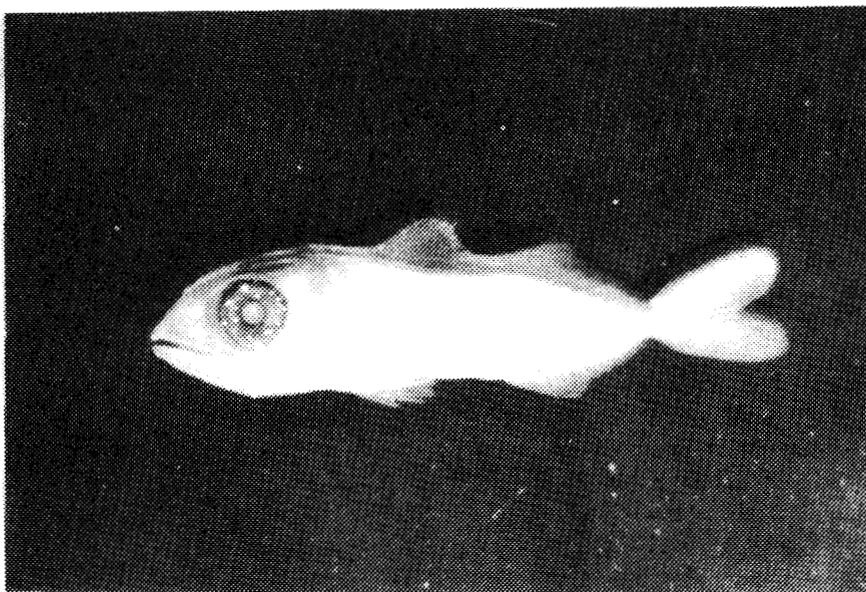
The answer to the paradox of high metabolic rates may come from the advantages they provide with respect to agility and mobility in hunting and capturing prey. Studies indicated tunas become more efficient than other fishes at higher swimming speeds. For tunas, high metabolic rates during low activity appear to be a physiological necessity for greater efficiency at high swimming speeds during feeding or when escaping from predators. The unique ability of tunas to conserve metabolic heat may also make high metabolic rates advantageous by keeping the tunas' swimming muscles warm when they penetrate cold, deep water in pursuit of prey.



Schematic diagram of the annular test tank (doughnut tank) system used to measure the physiological thermoregulatory ability of skipjack and yellowfin tunas. Seawater is delivered to and removed from the swimming channel through countercurrent perforated pipes, so that longitudinal temperature gradients do not develop. A computer continually calculates fish swimming speed based on data coming from the four photocells which monitor fish position in the swimming channel. The fish's deep red muscle temperature is measured by a thermistor probe connected to an ultrasonic transmitter. In this way, muscle temperature could be measured in free-swimming fish. Water temperature is controlled within 0.05°C by a heater and chiller system.



These skipjack tuna eggs are within an hour or two of hatching. The dark spot visible in each egg is an oil globule that provides energy and ensures flotation for the tiny egg, whose actual size is smaller than 1 millimeter in diameter.



This skipjack tuna larva is 26 days old (actual size: 16.72 millimeters long).

Spawning and rearing--The first successful attempts to artificially induce spawning in captive tunas were accomplished at the Kewalo Research Facility. The technique involved a periodic biopsy of kawakawa to determine the developmental stage of the eggs in the ovaries. After the eggs attained a critical size, hormone treatments were administered to induce spawning.

Eventually, because of advances in techniques and knowledge, hormone treatments were no longer used, and during the summer months, skipjack tuna were routinely spawned in the shoreside tanks of the Facility. This enabled researchers to investigate many of the techniques needed to rear larval pelagic fishes.

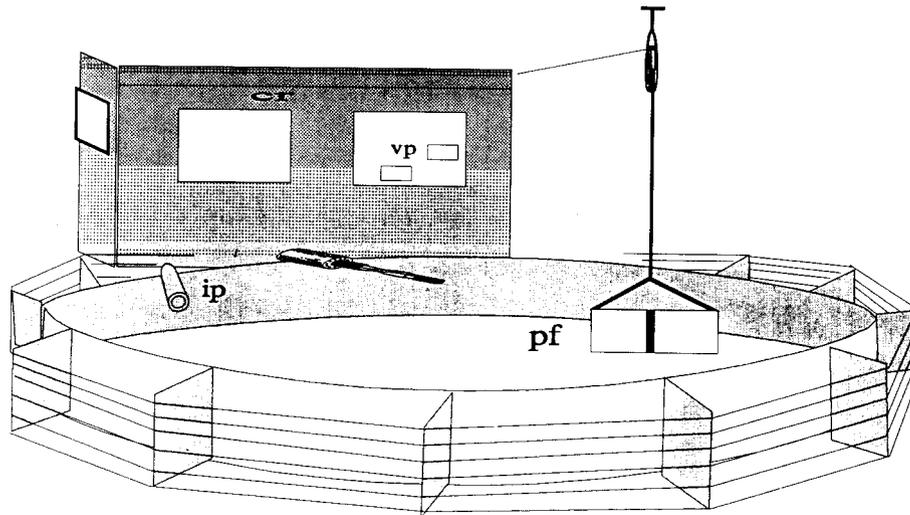
It was discovered that tuna eggs hatch about 24 hours after fertilization, and the yolk sacs of larvae are absorbed in about 2 days. At this critical stage, larval tuna must forage for food. To meet their nutritional requirements, a culture system for phytoplankton, rotifers, and copepods was begun and is continued today. The technology developed to rear larval tuna has opened new fields of research that focus on the previously unobservable, day-to-day development and early life history of pelagic fish.

Building upon the experience gained with tunas, subsequent studies involved spawning and rearing mahimahi (*Coryphaena hippurus*). Besides basic studies on the nutritional requirements, energetics, and growth of larval mahimahi, a series of studies on the species' tolerance to cold shock was also undertaken. These later experiments were designed to help assess the potential impacts of large-scale ocean thermal energy conversion projects, which move massive amounts of deep, cold ocean water to the surface in a process (roughly analogous to a steam turbine) that generates electricity. Although this process produces no air pollutants, unlike fossil-fuel based electricity production, the potential biological impact of the cold water brought to the surface needed to be evaluated. Again, the Kewalo Research Facility with its unique combination of animal holding facilities and laboratories proved an ideal place for the work.

Geomagnetic sensitivity--Tunas are among the most highly migratory fishes. They routinely make transoceanic migrations but also can precisely navigate on a daily basis. An understanding of the mechanism guiding the daily movements and long distance migrations of tunas is therefore central to understanding the biology of these species.

Since migration represents a substantial investment of energy, there has probably been intense evolutionary pressure to develop accurate sensory systems capable of guiding these movements. However, no special abilities useful in navigation had been detected among the tunas' common previously recognized sense systems (vision, smell, taste, etc.). Yet one other possibility existed: Tunas could possess a magnetic compass sense.

After the discovery that yellowfin tuna have up to 10 million crystals of magnetite (a biologically generated magnetic crystal) in the ethmoid bones of the skull and that the fish produce the magnetite under very closely controlled conditions of size, shape, and chemical composition, studies were undertaken to test the ability of yellowfin tuna to discriminate between different magnetic fields. The fish were trained to perform a



This system is for measuring the ability of yellowfin tuna to detect local changes in the Earth's magnetic field. The vertical component of the Earth's magnetic field is altered by passing an electrical current through the coil of wire encircling the tank. The tuna is trained to swim through a rectangular pipe frame (pf) lowered into the tank. The fish's ability to detect the changes is measured by the number of passes through the pipe frame per minute. Correct responses are rewarded by food delivered from an automatic dispenser; incorrect responses are punished by the food reward being withheld. To avoid any possible cues being given by the researcher to the fish, all observations must be made through a one-way glass window in the small hut situated near the tank.

conditioned response (swimming through a hoop) at a consistent rate. They were then tested by being rewarded with food when one magnetic field was present in the tank and punished (food withheld) when the second field was present. If the fish were able to detect the difference between the two magnetic fields, maintaining a high rate of response during positively reinforced trials would maximize food rewards, whereas a low rate of response during negatively reinforced trials would minimize the cost of responding. Thus, discrimination would be measured as a difference in the rates at which the fish swim through the hoop in anticipation of positive or negative reinforcement. These experiments were clearly able to show that yellowfin tuna can learn to use magnetic field information to make appropriate decisions--the first proof tunas possess a magnetic sense which is probably used for navigation.

Related studies showed that a large branch of the anterior, lateral-line nerve ramifies in the area of the ethmoid bones which contain the magnetite crystals. It is therefore possible that a branch of this nerve may be associated with the magnetite crystals and form the magnetoreceptor organ, although this still remains to be determined.

Lobsters

Starting in the late 1970s, the commercial fishery targeting spiny lobster (*Panulirus marginatus*) and slipper lobster (*Scyllarides* spp.) in the Northwestern Hawaiian Islands rapidly expanded. Field research conducted by the Honolulu Laboratory showed that most small (i.e., sublegal) lobsters released after being captured in traps were eaten by fish before reaching the ocean bottom. Escape vents installed in commercial lobster traps were obviously needed to prevent this problem and to retain a viable commercial fishery. But how big and what shape should these escape vents be? Where on the traps should they be located?

Studies were begun at the Kewalo Research Facility where populations of known sizes of lobster were set up in the laboratory's shoreside tanks. Lobster traps with various sizes and shapes of escape vents were added to the tanks, and the sizes of lobsters retained were carefully monitored. Optimal results were obtained using two escape-vented panels with two circular openings of 67 millimeters in diameter. Field trials conducted in the Northwestern Hawaiian Islands using similarly equipped traps confirmed the efficacy of the vents.

As a result of this important laboratory and field research, all commercial lobster traps used in the Northwestern Hawaiian Islands since January 1, 1988, must contain these escape-vented panels.

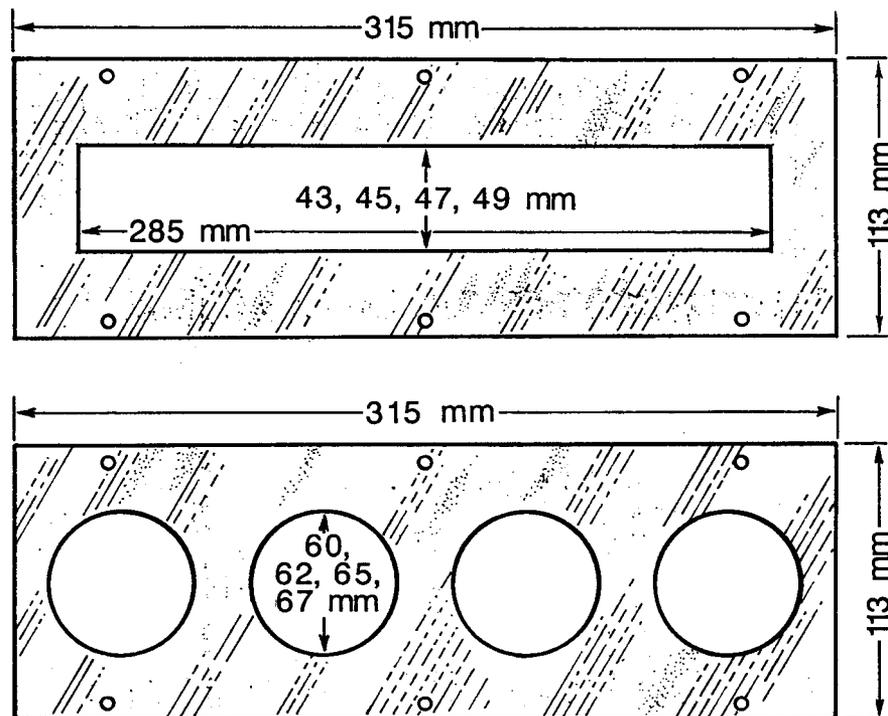
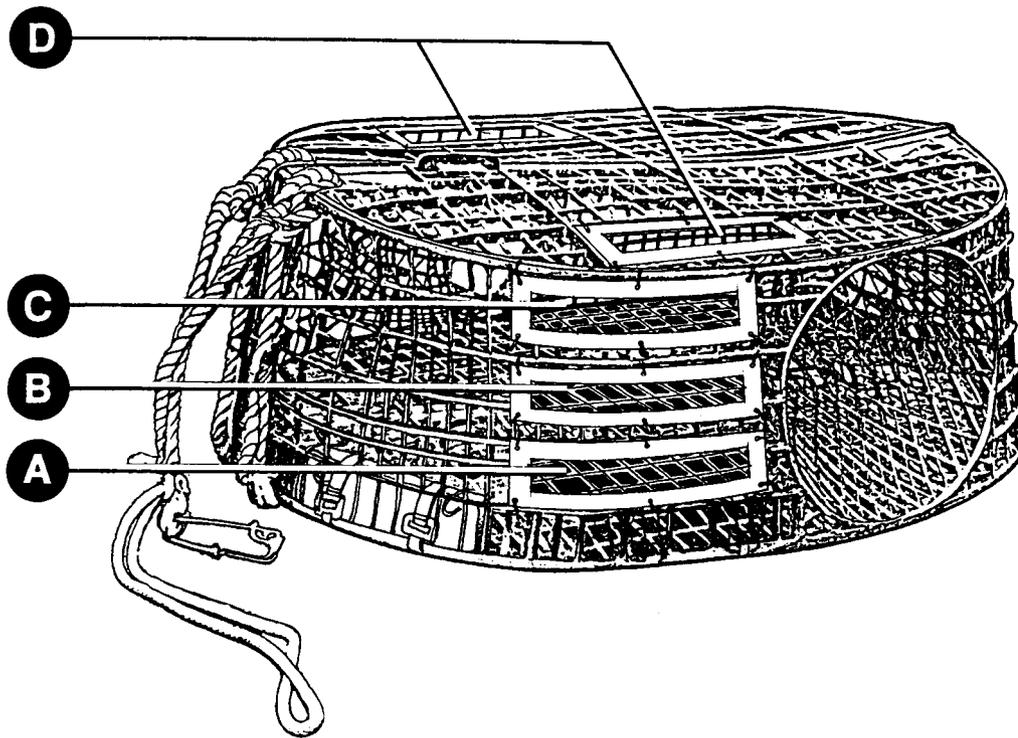


Diagram of escape vents, varying in size and shape, tested at the Kewalo Research Facility.



This diagram depicts a Fathoms Plus shellfish trap, which by 1984 had almost completely replaced the wire mesh trap used by lobster fishermen. This trap has been fitted with several escape vents that Honolulu Laboratory scientists tested to determine the locations which best allow the release of undersized lobsters. This diagram shows the vertical placement of the escape vents tested: (A) 45 millimeters from the bottom, (B) 115 millimeters from the bottom, (C) 195 millimeters from the bottom, and (D) top of trap.

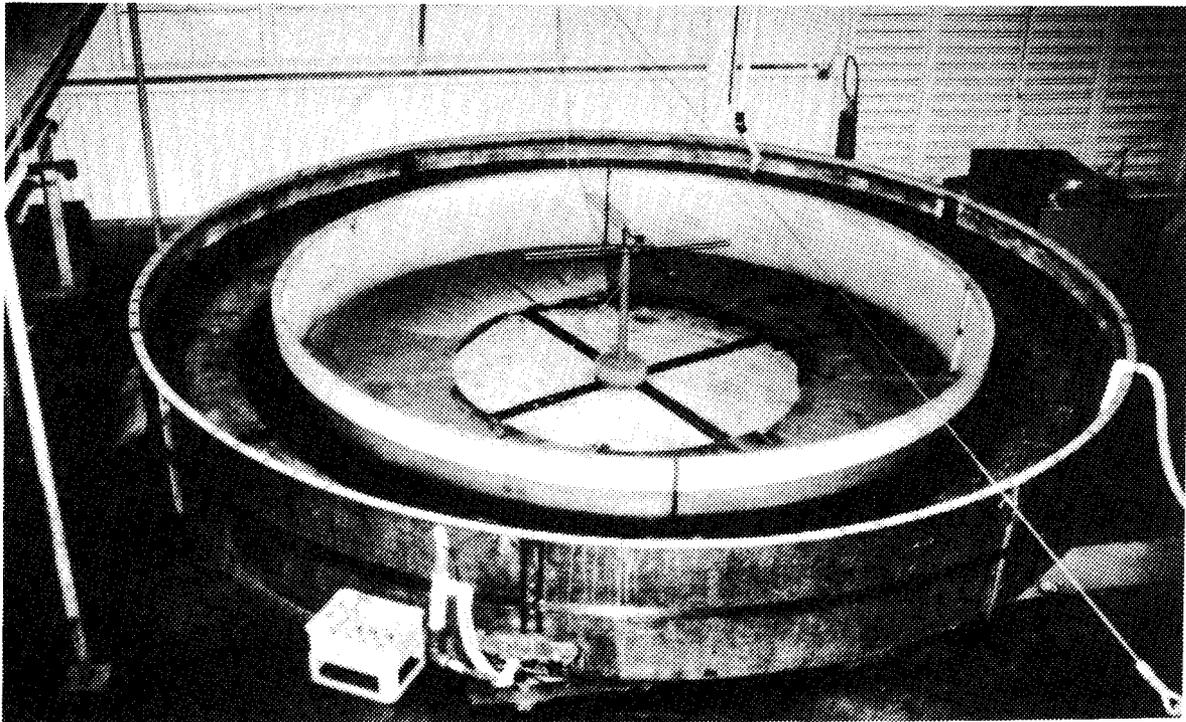
THE KEWALO RESEARCH FACILITY TODAY

Tuna Research

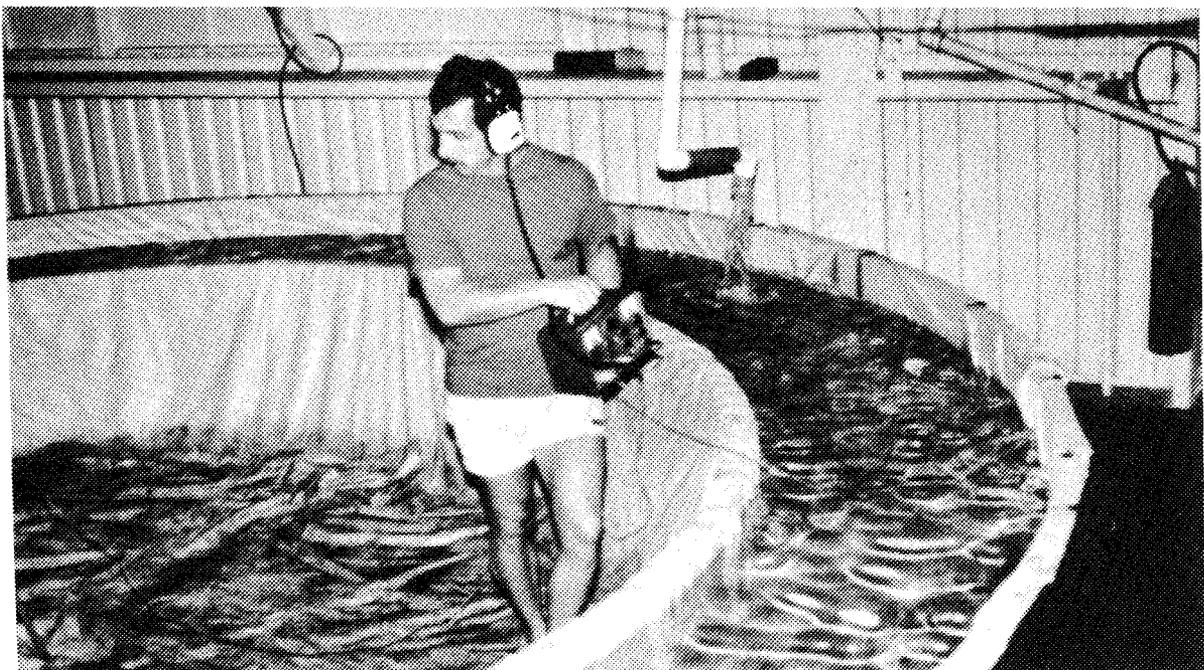
Olfaction

Work conducted 20 years ago at the Kewalo Research Facility established that tunas have an excellent sense of smell, capable of detecting the very dilute odor of their prey. Recent and ongoing research with captive tunas has shown that they can distinguish between odors of different types of prey, and that some prey odors cause stronger search behavior than others, indicating tunas probably use their sense of smell to detect prey before they come within visual range.

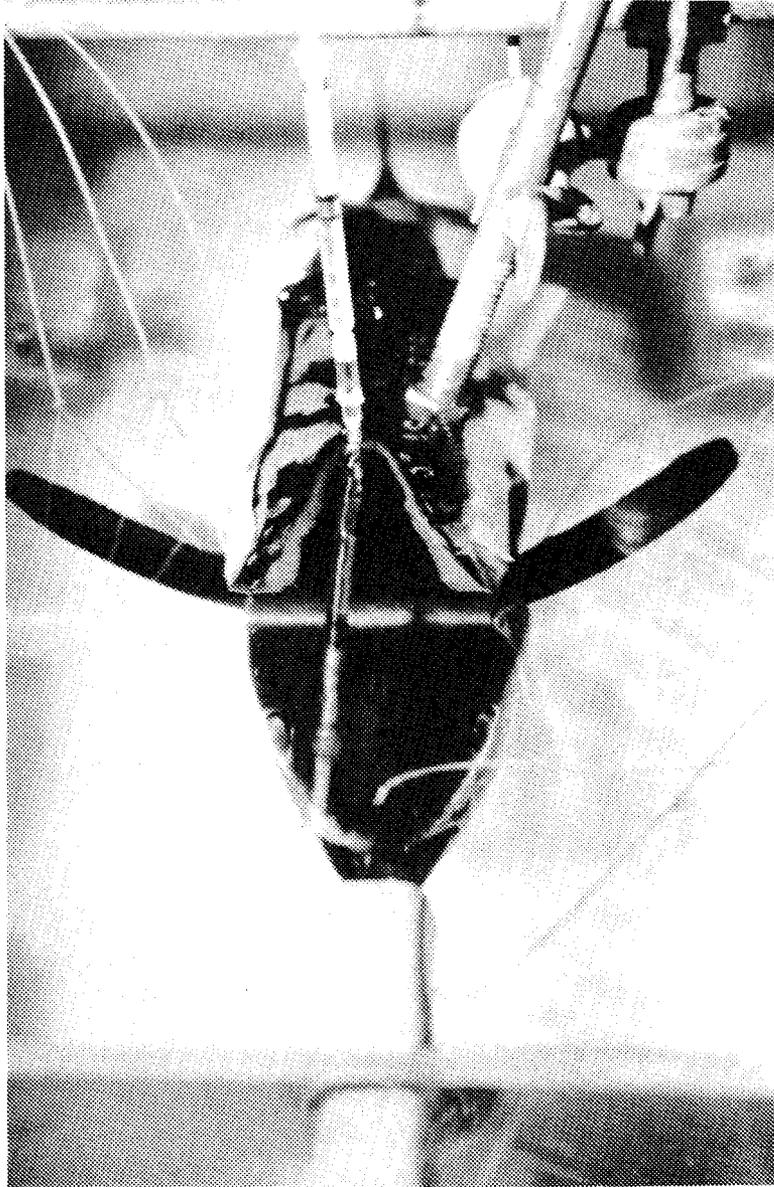
Recent research has been aimed at analyzing the chemical structure of natural prey odors, developing synthetic prey odors, and testing the efficacy of these synthetic prey odors for eliciting a feeding response. Eventually, it may be possible to use natural or synthetic odors to enhance the effectiveness of traditional fishing techniques. If an inexpensive synthetic odor can be formulated, it could be used in the live-bait and the



This "doughnut" tank is used for physiological experiments on swimming tunas. The center section of the tank is dry which allows investigators to follow a fish while carrying physiological (for example, heart rate or cardiac output) monitoring equipment. Data gathered in this manner are used to confirm similar observations made on non-swimming fish in the laboratory.



Much effort at the Kewalo Research Facility has been put forth to determine the habitat requirements of skipjack and yellowfin tunas, especially their temperature and minimum ambient oxygen requirements. Here an investigator is monitoring the cardiac output of a yellowfin tuna by using a Doppler ultrasonic blood flowmeter. The blood flow probe, mounted over the main vessel carrying blood from the heart, is carried by the fish, while the electronic instrumentation needed to quantify blood flow is carried by the investigator. The output of the blood flowmeter is sent to a chart recorder (not shown). The head phones are used to monitor the functioning of the flowmeter.



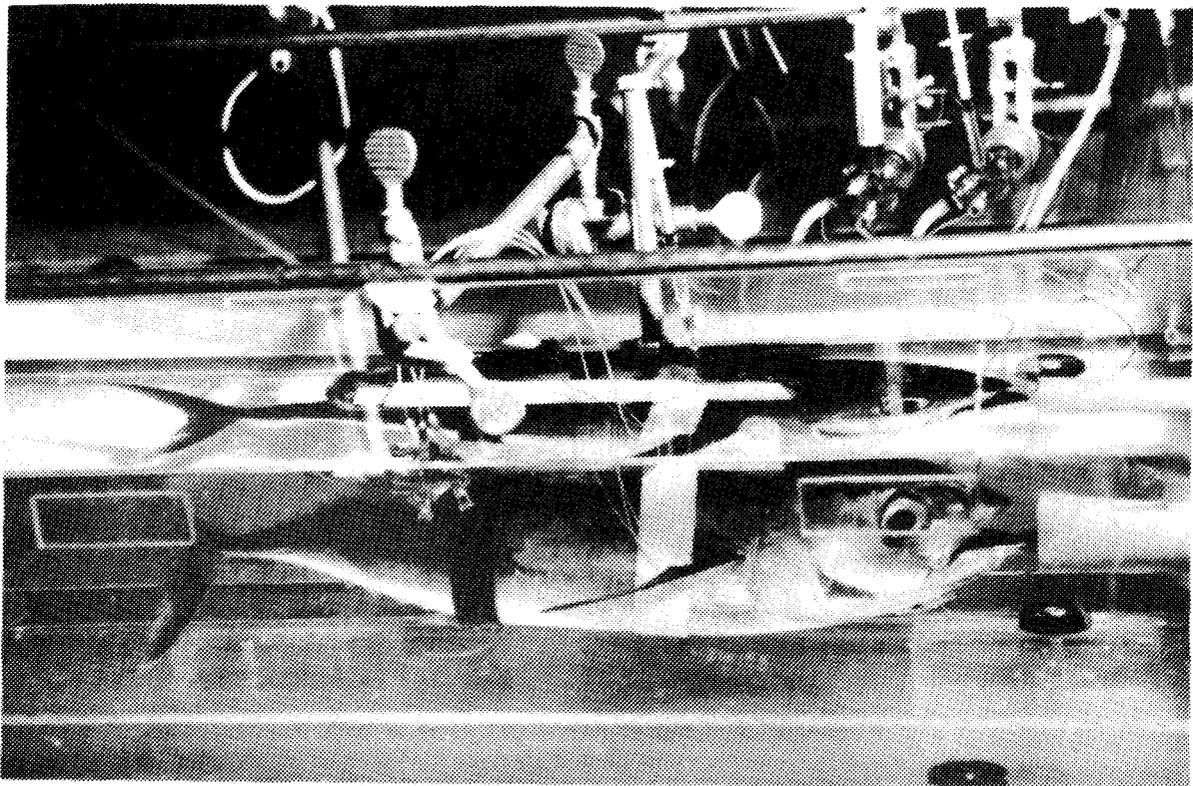
Overhead view (above photo) and side view (right photo) of a live yellowfin tuna being held on the operating table designed specifically to test the physiological responses and tolerances of tunas to changes in environmental conditions. Tunas normally must swim to ventilate their gills; however, in this situation, water is supplied to the fish by the pipe immediately in front of the mouth. With this system, blood pressure, ventilation volume, cardiac output, and other physiological parameters can be simultaneously monitored.

handline tuna fisheries to increase catch success and decrease dependency on expensive natural bait.

Research to Determine Energy and Habitat Requirements of Tunas

Mathematical models of the energetics of tunas enable scientists to better explain and predict abundance and maximum sustainable yields. A large portion of the data collected over the past quarter century at the Kewalo Research Facility has been directed toward acquiring the data necessary for these models. Results to date indicate that the growth rate of skipjack tuna less than 11.8 kilograms (26 pounds) is governed by food consumption, whereas the growth of fish larger than 15.0 kilograms (33 pounds) is limited by the energetic demands of activity. Other models, integrating data from laboratory experiments on tunas with oceanographic information, indicated the distribution of small tunas is most likely dependent on the availability of food, whereas the distribution of larger fish is dependent on environmental conditions, such as temperature and oxygen levels.

To further investigate the physiological abilities and tolerances of tunas to temperature and oxygen conditions, a laboratory specifically designed to conduct physiological experiments on tunas has been developed at the Kewalo Research Facility. The laboratory contains a vibration-free operating table with running seawater and extensive physiological monitoring equipment. The temperature and oxygen levels of the



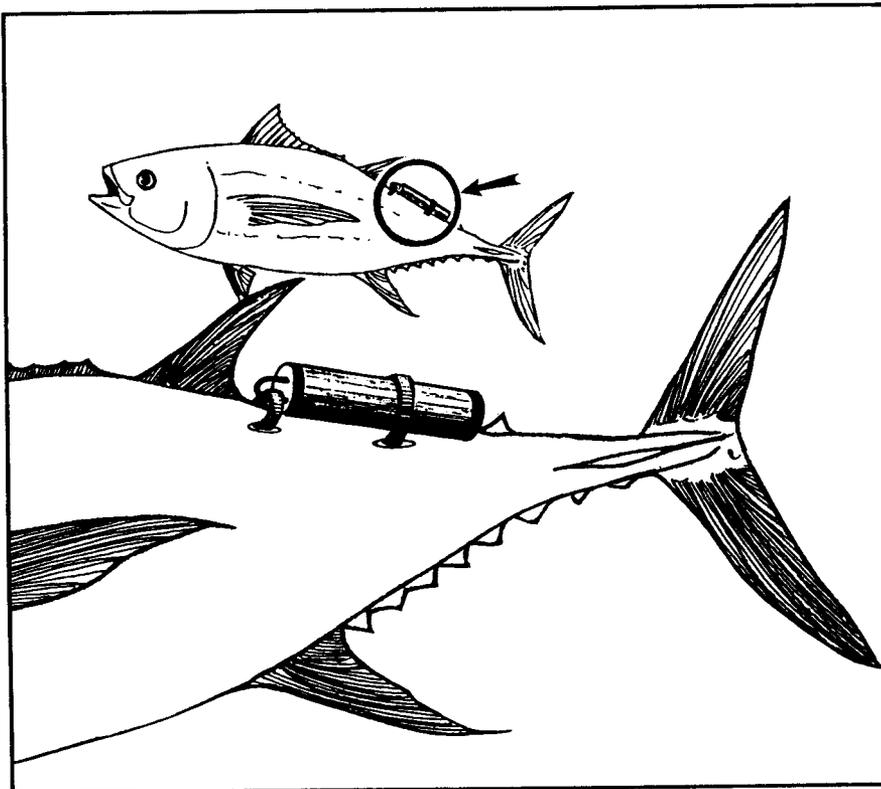
water supplied to the operating table can be closely controlled and monitored. Tunas gently restrained on the table respond normally when subjected to changes in environmental conditions.

Results of recent experiments using this system have shown that tunas are sensitive to even minute reductions in ambient oxygen; they begin making physiological adjustments to reductions in ambient oxygen far smaller than those needed to elicit swimming speed changes. Data have also been obtained on the effects of rapid temperature change on the metabolic rates and blood acid base chemistry of tunas and their truly remarkable ability to recover from strenuous exercise.

Models of the function of the tuna cardiorespiratory system (which removes oxygen from the water passing over the gills) have been recently developed using data obtained in the physiology laboratory. Surprisingly, the results generated by these models imply that the unique anatomy/physiology/biochemistry of tunas has evolved to permit rapid repayment of oxygen debts (i.e., rapid lactate metabolism) rather than high sustained cruising speeds.

Tracking Tuna Movements

The Kewalo Research Facility has at its disposal the *Kaahale 'ale*, a 33-foot research vessel equipped with sophisticated electronic and navigational equipment to track the vertical and horizontal movements of tunas and billfishes carrying ultrasonic transmitters.



This drawing depicts the placement of an ultrasonic transmitter which can be used to track tuna and other animals in the ocean. The transmitter is attached to the fish by two nylon straps that are inserted through the dorsal musculature and pterygiophores associated with the second dorsal fin.

The vessel is an integral part of the facility and is in constant demand to test the results of theoretical and experimental investigations.

Recent tracking work using the *Kaahale 'ale* has determined the short-term horizontal and vertical movements of tunas associated with fish aggregating devices (FADs). Much of the success of the tracking studies is due to the ability to hold tunas in captivity and to test various ultrasonic transmitter attachment methodologies on captive fish. The project is now moving into its second phase, that of simultaneously monitoring tuna movements and physiological parameters such as body temperature and tail beat frequency (i.e., swimming activity). The data collected in these experiments will be incorporated into more sophisticated computer models capable of predicting tuna movements, distribution, and vulnerability to specific types of fishing gear. Again, the ability to test new telemetry devices on fish held at the Kewalo Research Facility is helping to keep the tracking project at the forefront of this area of science.

Burnt Tuna

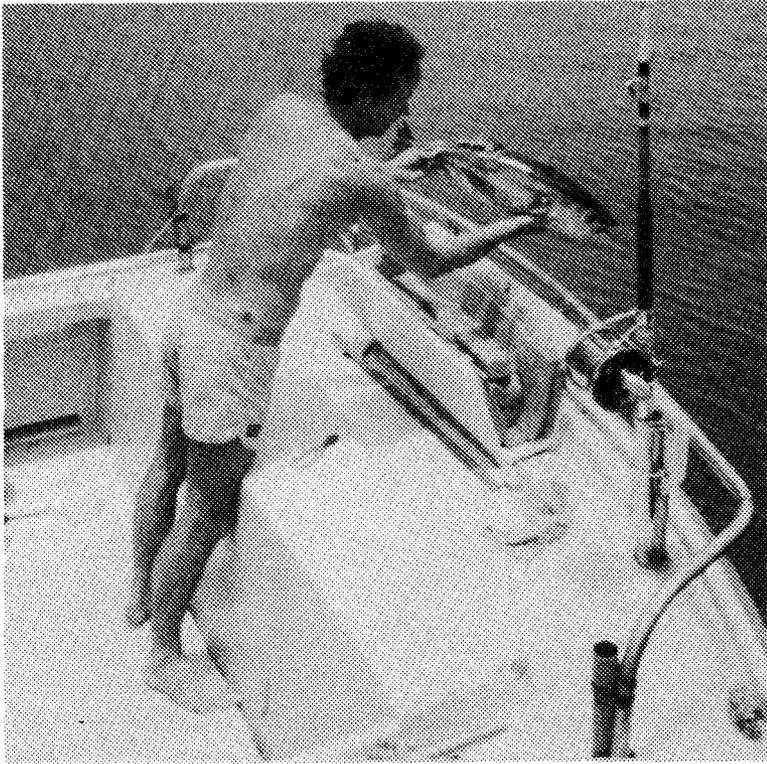
In Hawaii, the handline fishery for large yellowfin tuna and bigeye tuna (*Thunnus obesus*) land fish intended primarily for raw consumption as sashimi. The current value of the fishery is estimated at over \$5 million annually. International interest in this type of fishing is growing because of its low initial capital investment, low operating and fixed expenses, strong export markets, and high profitability. Unfortunately, the tuna handline and, primarily, recreational troll fisheries are plagued by a product quality problem known as "burnt tuna" or, in Japanese, as *yake niku* (literally translated as "cooked meat"). When fish are intended for raw consumption, product quality is obviously of utmost importance.

For years, the high muscle temperatures and high muscle acidity created during the landing of large tunas on handlines were hypothesized to be the underlying cause of burnt tuna. However, when samples of burnt tuna muscle were examined at the histological and biochemical level, the observations did not fit the hypothesis. Based on the work of scientists at the Kewalo Research Facility, a new concept was developed: burnt tuna is caused by activation of the proteolytic (protein-destroying) enzyme known as "calcium activated neutral protease" or more commonly as "calpain." Current efforts are directed at proving this hypothesis, developing a thorough understanding of the etiology of burnt tuna, and, more importantly, developing strategies that could be used by fishermen to successfully mitigate this problem.

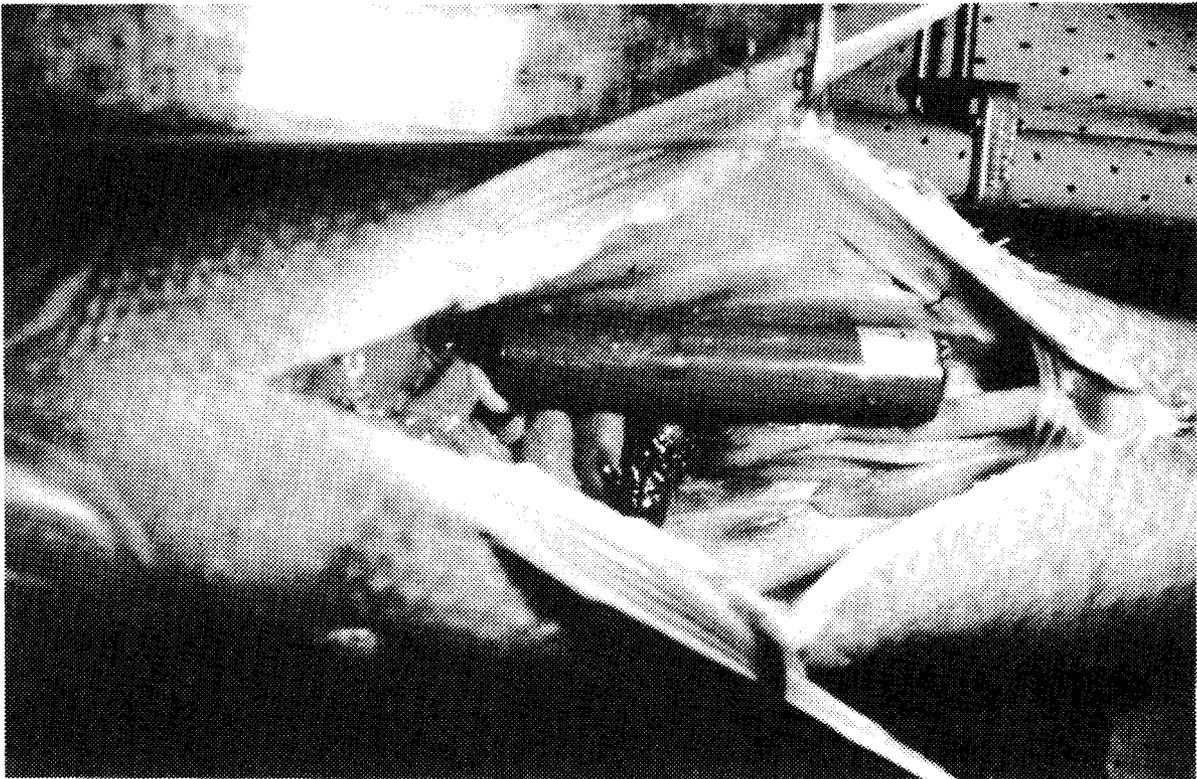
Surprisingly, it also now appears that burnt tuna is not an isolated phenomenon, but rather is biochemically identical to processes occurring in heart muscle during a heart attack and to some forms of human muscular dystrophy. Research begun at the Kewalo Research Facility to answer a specific fishery's product problem may prove to have medical importance!

Spawning and Rearing

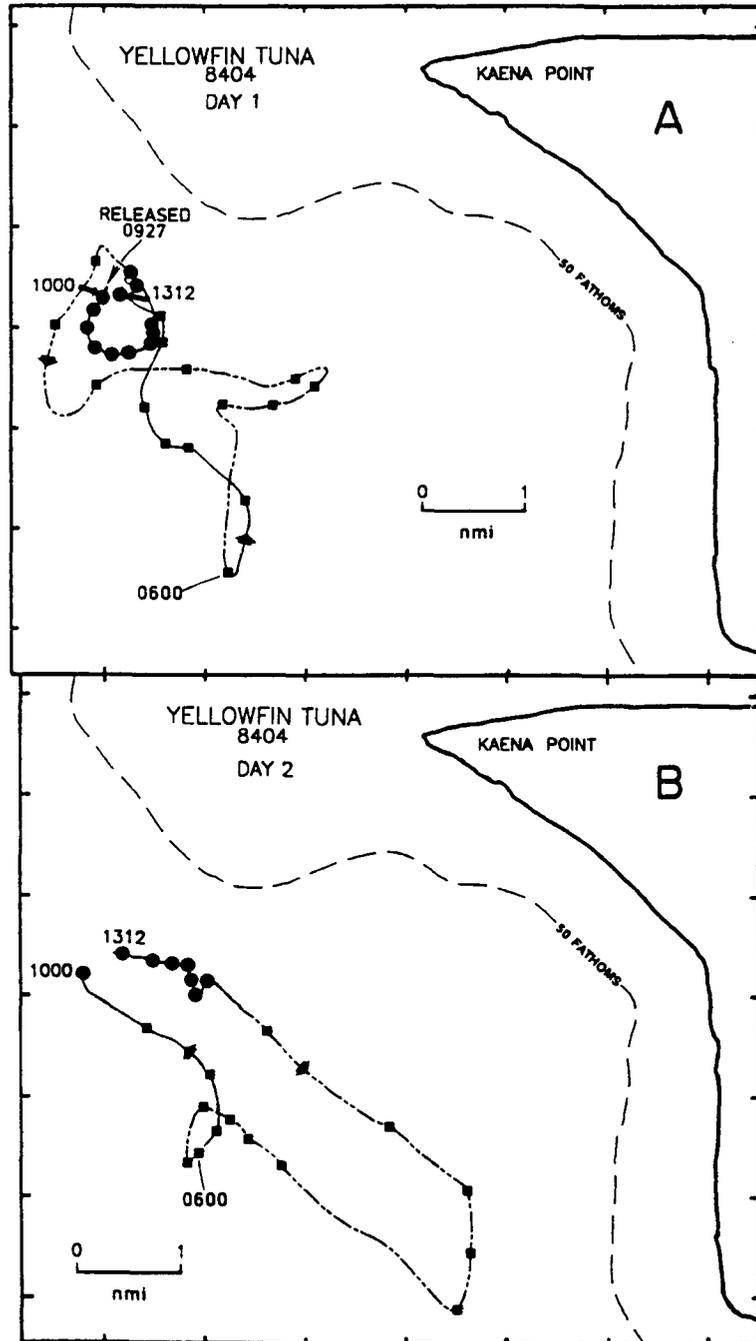
The ability to spawn and rear fishes at the Kewalo Research Facility provides known-age specimens which can be used to validate methods used for estimating the age



A sonic transmitter is attached to a freshly caught tuna, which is then released so that the fish's swimming depth and direction can be tracked to better understand its horizontal and vertical movement patterns.

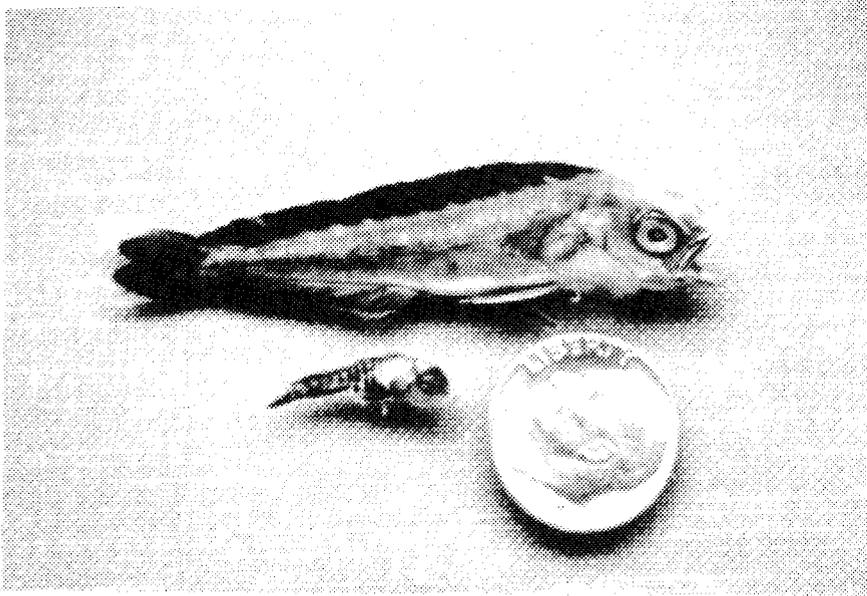


This ultrasonic transmitter is being surgically implanted in the body cavity of an anesthetized opakapaka (*Pristipomoides filamentosus*), so that scientists will eventually be able to track the movements of individuals of this species. The Kewalo Research Facility affords the opportunity to test various transmitter attachment schemes before they are attempted in the field.



The Honolulu Laboratory was one of the pioneers in the use of ultrasonic tags to track tuna in the open ocean. With the use of ultrasonic tags, research has shown that skipjack and yellowfin tunas are temporarily territorial and remain in a given area for some time in Hawaiian waters. They repeatedly return to the same area each morning; this behavior implies that tuna can navigate and have a sense of time.

These figures show a 48-hour track of a yellowfin tuna (No. YF8404) tagged and released at 0927 at a fish aggregating device (FAD) off the island of Oahu. Circles show each hour the fish was near the FAD; squares show each hour the fish was away from the FAD. The solid line equals the daytime movements, the broken line equals nighttime movements. On the first day of tracking (A), the fish remained very close to the FAD which moved in a circular path in the current. The fish departed the FAD after nightfall, and returned at 1312 the following day. On the second day (B), the fish behaved similarly, despite a different pattern of movement by the FAD.

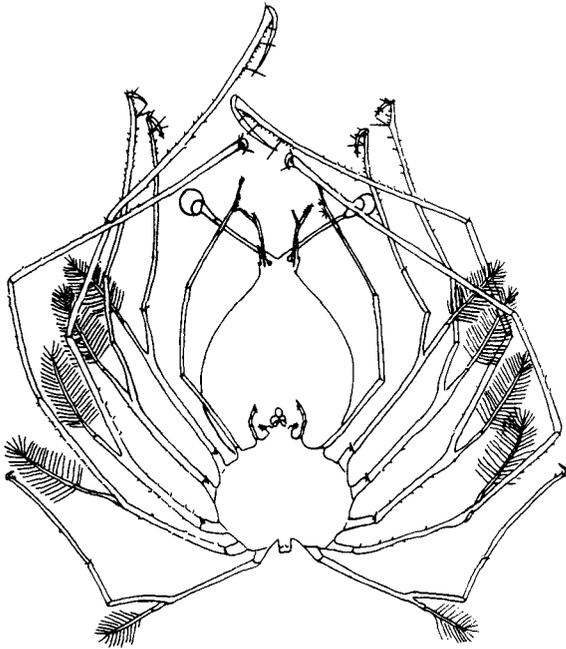


These mahimahi (*Coryphaena hippurus*) were reared from fertilized eggs. The lower individual is about 40 days old, and the upper individual is about 50 days old. When they first hatched, these mahimahi were less than 1 millimeter long (less than the thickness of the dime pictured).

of tropical fishes, especially short-lived (less than one year) species. Such work has been completed for nehu (*Encrasicholina purpurea*), skipjack tuna (*Katsuwonus pelamis*), and mahimahi (*Coryphaena hippurus*).

Most tropical fishes appear to be partial spawners. Only recently, with the discovery of postovulatory follicles, has it become possible to estimate the frequency of spawning. Skipjack tuna and mahimahi observed spawning in tanks have been sampled to determine the deterioration rate of postovulatory follicles which appeared to exist for about only 24 hours after spawning. Thus it is now possible to estimate the spawning frequency of skipjack tuna and mahimahi in the wild.

Recent spawning and rearing studies at the Kewalo Research Facility centered on rearing larval slipper lobster (*Scyllarides haanii*) and deepwater shrimp (*Heterocarpus laevigatus*). Deepwater shrimp, collected at sea, were successfully hatched and the larvae reared for up to 139 days in the laboratory, during which time they went through 37 molts! Similar techniques applied to larval lobster have enabled scientists to rear them in captivity for up to 123 days. The objective of these studies is to provide information useful for identifying the larval lobster and shrimp that often are a large part of specimens caught during plankton sampling. These surprisingly long larval stages also help explain how widespread apparently isolated adult populations can be genetically related. Not the adults, but rather their long-lived planktonic larvae are migrating over long distances. This information also has obvious fishery management implications.



This drawing is of a lobster larva (phyllosome) such as those hatched and reared at the Kewalo Research Facility. Little is known about the early life history of the commercially important lobster species caught near the Hawaiian Islands. Rearing lobster larvae from eggs in captivity allows scientists to (1) identify the larval stages caught in plankton nets in the open ocean, (2) determine how long the larvae of various lobster species remain in the plankton, and therefore (3) calculate how far the larvae could possibly be transported by oceanic currents.

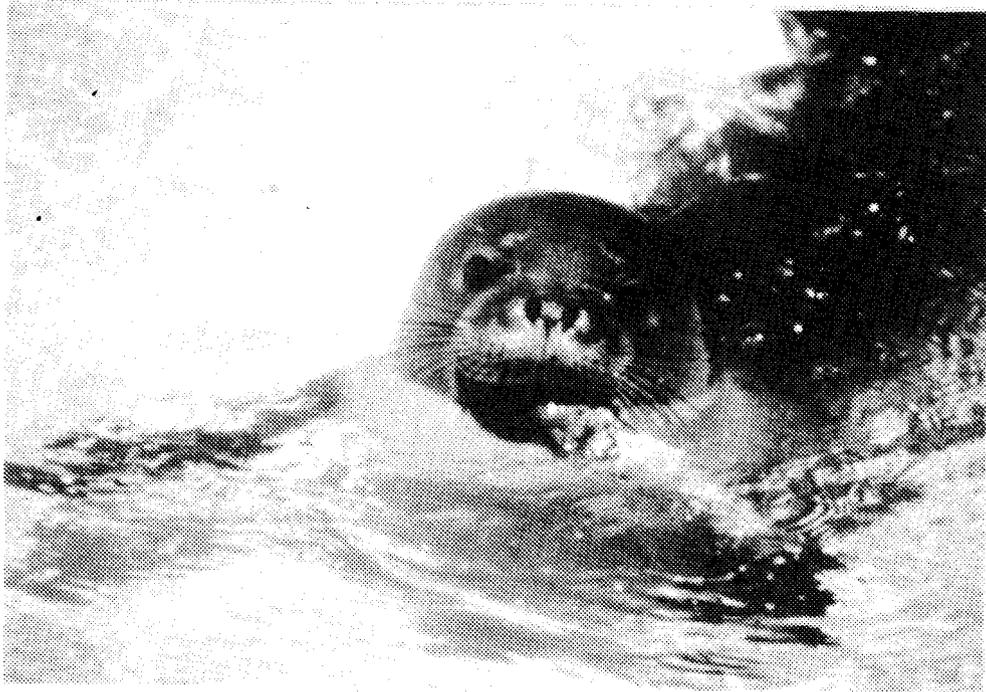
Protected Species

Hawaiian Monk Seals

The isolated atolls and beaches of the Northwestern Hawaiian Islands are the primary terrestrial habitat of the Hawaiian monk seal (*Monachus schauinslandi*). This species and its Mediterranean cousin (*M. monachus*) have remained virtually unchanged for 15 million years and are sometimes referred to as living fossils. The Caribbean monk seal (*M. tropicalis*) became extinct in the mid-1950s, the Mediterranean population has fewer than 500 animals, and the Hawaiian monk seal has around 1,500 animals. These are highly endangered animals.

Monk seals, for the most part, lead a pelagic existence and may be at sea for a month or more. They are also able to dive for food up to 400 feet. The seals haul out on deserted beaches and atolls to rest, pup, and nurse their pups. Although docile, monk seals are extremely sensitive to any human disturbance and will leave preferred haul-out areas, sometimes even deserting their pups. Since monk seals are in constant danger from predation by sharks (most adult monk seals carry large shark-caused scars), more time spent in shark-infested waters due to human disturbance means higher mortality rates.

Since the early 1980s, the NMFS Honolulu Laboratory has been sending researchers to the Northwestern Hawaiian Islands to observe, tag, and count monk seals. One project (the Rehabilitation Project) has been transporting sick or abandoned pups to the Kewalo Research Facility and, more recently, to Sea Life Park to be "fattened up" overwinter and then transported to Kure Atoll where the population has been seriously depleted. This



A Hawaiian monk seal catching a bite to eat.



This ocean-beach enclosure at Kure Atoll is used by the Honolulu Laboratory's Head Start and Rehabilitation Projects to protect female Hawaiian monk seal pups and yearlings from attacks by aggressive male seals and sharks until the females are ready to be released into the wild.

project has been successful in bolstering the female seal population at Kure Atoll; already some of the project's seals have become successful breeding members of the expanding Kure population.

Research on monk seals has also been conducted at the Kewalo Research Facility. Studies have included testing techniques to lessen the aggressive behavior of adult male monk seals to reduce the problem of "mob mating" that occurs in certain areas where the male to female sex ratio of adult seals has become abnormally skewed over the years due to high female seal mortality. Studies have also been conducted at the Kewalo Research Facility on the metabolic rate of seals, information important for determining how much undisturbed feeding area adult seals may need.

Hawaiian Sea Turtles

The Kewalo Research Facility provides a badly needed laboratory where research on and rehabilitation of several species of threatened and endangered Hawaiian sea turtles can be conducted. Several successful recoveries of turtles either injured intentionally by spears or unintentionally by boat propellers have been made.

The most common sea turtle around Hawaii is the green turtle (*Chelonia mydas*) or *honu* in Hawaiian. The turtles are primarily vegetarian, eating algae growing on coral reefs.



Hawaiian green sea turtles basking in the sun at French Frigate Shoals in the Northwestern Hawaiian Islands. French Frigate Shoals is a major feeding and nesting area of this turtle species.

The smaller and rare hawksbill turtle (*Eretmochelys imbricata*) or *ea* in Hawaiian is found around the islands of Molokai and Hawaii. Large (up to 1,500-pound) leatherback turtles (*Dermochelys coriaca*) are never found near the islands, but are regularly seen in the open sea where they feed primarily on jellyfish.

Sea turtle populations have been decimated worldwide because of habitat destruction, direct exploitation of adults and eggs for food and other items, illegal poaching, ingestion of plastic debris, and unintended entanglement in fishing gear. In Hawaii, Florida, and elsewhere, another problem has recently cropped up--debilitating, life-threatening tumors which may grow up to 12 inches in diameter on green turtles. The prevalence of this disease is increasing at an alarming rate in a number of Hawaii's green turtle populations. Research on this problem is continuing and involves maintaining turtles in captivity and working cooperatively with scientists from Hawaii and the U.S. mainland, and conducting ultrasonic tracking of healthy and tumor-afflicted turtles.

RELATIONSHIPS WITH THE UNIVERSITY OF HAWAII

The Honolulu Laboratory (through its Kewalo Research Facility and in other ways) maintains a special relationship with the University of Hawaii. There is free dialogue and exchange of information among researchers working at the Kewalo Research Facility and scientists in the University's Departments of Zoology, Physiology, Oceanography, Biochemistry, Nutrition, and Animal Sciences. The Honolulu Laboratory provides part-time employment for University of Hawaii undergraduates and support for master's and doctoral degree candidates by providing laboratory space, experimental tuna, and monetary grants. Laboratory scientists also serve as advisors on graduate student thesis committees.

Numerous graduate students from the University of Hawaii and other universities have earned advanced degrees that involved work at the Kewalo Research Facility:

San Diego State University

Robert Olson

Scripps Institution of Oceanography

Heidi Dewar
Peter Fields

Shizuoka Prefectural Fisheries Experimental Station

Minato Yasui

Simon Fraser University

John Keen

University of Hawaii

Andrew Ayers
C. Scott Baker
Michael Barry
Gordon Bauer
Robert Bourke
Richard Brill
Peter Bushnell
Ron Dunn
Sharon D. Hendrix
Walter N. Ikehara
Barbara A. Kuljis
Christopher Moyes
Elizabeth A. Monckton
Linda M. B. Paul
Anjanette S. Perry
Anthony Sudekum
Michael M. Walker
Cheryl Watson

University of St. Andrews

John Salamonski

University of Miami

Daniel Benetti

University of British Columbia

Les Buck
 Michael Guppy
 Mark Heieis
 Michael Herrick
 Tom Petersen
 Manabu Shimazu
 Jean-Michele Weber
 Tim West

University of Wisconsin

Christofer Boggs
 Sherry Steffel

LIST OF VISITING INVESTIGATORS

The Kewalo Research Facility has, since its inception, hosted visiting scientists from around the world. Many have come because it is one of the few laboratories in the world that routinely maintains live tunas specifically for behavioral and physiological research. Scientists who have worked at the Kewalo Research Facility include the following:

Aquatic Farms

Dr. Edward D. Scura

Bedford Institute of Oceanography

Dr. Barry S. Muir

Boston University

Dr. Jelle Atema

Canada Center for Inland Waters

Dr. Arthur J. Niimi

**Centre National pour l'Exploitation
des Oceans**

Dr. F. Havard-Duclois

Huntsman Marine Laboratory

Dr. Thomas W. Moon

**Indiana University School of
Medicine**

Dr. Kenneth R. Olson

**Institute of Oceanology, Academy of
Sciences, Moscow**

Dr. Sergei M. Kashin

**Inter-American Tropical Tuna
Commission**

Dr. Gary D. Sharp

John G. Shedd Aquarium

Dr. William P. Braker

Kinki University

Dr. Teruo Harada
 Dr. Shigeru Miyashita

Kyoritsu Women's University

Dr. Hiroki Abe

**Laboratory of Comparative
Biochemistry**

Dr. Grant R. Bartlett

Long Island University

Dr. Phyllis H. Cahn

Massey UniversityDr. Peter Davie
Dr. Craig Franklin**McMaster University**

Dr. Steve Perry

Micronesia Maritime Authority

Michael A. McCoy

Montana State University

Dr. Calvin M. Kaya

Nagoya UniversityDr. Hiroshi Niwa
Dr. Tamotsu Tamura**Naval Undersea Center**

Dr. A. Earl Murchison

Oceanarium, Inc.

Dr. John H. Prescott

Pacific Gamefish Foundation

Dr. Charles Daxboeck

**Resources Systems Institute
East-West Center**

Dr. John E. Bardach

Rutgers University

Dr. Bryant T. Sather

Scripps Institution of Oceanography

Dr. Jeffrey Graham

Simon Fraser UniversityDr. Anthony Farrell
Jeff Johansen
Haruyo Kashihara
Dr. Glen Tibbits**South Pacific Commission**

Dr. Robert E. Kearney

State University of New York

Dr. E. E. Suckling

**Technion-Israel Institute of
Technology**

Dr. Daniel Weihs

Texas A&M University

Dr. William H. Neill

University of British ColumbiaDr. Peter Arthur
Dr. Robert Blake
Dr. Brian Emmett
Dr. Christopher French
Dr. Peter W. Hochachka
Dr. C. S. Holling
Dr. William C. Hulbert
Dr. David Jones
Dr. William Milsom**University of Tokyo**

Dr. Isao Hanyu

University of HawaiiDr. Jean L. Cramer
Dr. E. Gordon Grau
Dr. Kim Holland
Hank Marrow
Dr. Martin D. Rayner
Dr. Terence A. Rogers**University of Guelph**

Dr. E. Don Stevens

University of San Diego

Dr. Alan R. Hargens

University of Birmingham, England

Dr. Pat Butler

University of Notre Dame

Dr. F. W. Goetz, Jr.

University of New York

Dr. J. A. Suckling

University of California, Los AngelesDr. Malcolm S. Gordon
Dr. Vladimir Walters

University of Toronto

Dr. F. E. J. Fry

University of California, Davis

Dr. Claude M. Nagamine

University of North Carolina

Dr. John M. Miller

University of California, San Diego

Dr. Ted Bullock

Dr. Odile Mathieu-Costello

University of Chicago

Dr. Barbara Block

University of Wisconsin

Dr. James F. Kitchell

Dr. John J. Magnuson

Dr. Warren P. Porter

ACKNOWLEDGMENTS

Many people over the years have contributed to the ongoing success of the Kewalo Research Facility. I would like to take this opportunity to thank them all and especially recognize the pioneering efforts of Richard Barkley, Andrew Dizon, John Magnuson, John Marr, Eugene Nakamura, Bill Neill, Richard Shomura, Don Stevens, and Albert Tester. If those of us currently associated with the Kewalo Research Facility can reach new heights, it is only because we stand on the accomplishments of the people who went before us.

I also acknowledge the people who contributed to the preparation of this booklet: George Balazs, George Boehlert, Peter Bushnell, Randolph Chang, William Gilmartin, Kim Holland, Tom Kazama, Michael Walker, Cheryl Watson, Leslie Williams, and Howard Yoshida.

**PARTIAL LIST OF SCIENTIFIC PUBLICATIONS
RESULTING FROM RESEARCH AT THE
KEWALO RESEARCH FACILITY**

- Abe, H., R. W. Brill, and P. W. Hochachka.
1986. Metabolism of L-histidine, carnosine, and anserine in skipjack tuna. *Physiol. Zool.* 59(4):439-450.
- Arthur, P.G., T.G. West, R.W. Brill, P.M. Shulte and P.W. Hochachka.
1992. Recovery metabolism in skipjack tuna (*Katsuwonus pelamis*) white muscle; rapid and parallel changes of lactate and phosphocreatine after exercise. *Can. J. Zool.* 70:1230-1239.
- Atema, J., K. Holland, and W. Ikehara.
1971. The sense of taste in fishes. *In*: Beidler, L. M. (editor), Handbook of sensory physiology IV. Chemical senses 2. Taste, p. 293-336. Springer-Verlag, Berlin, Heidelberg, and NY.

1980. Olfactory responses of yellowfin tuna (*Thunnus albacares*) to prey odors: Chemical search image. *J. Chem. Ecol.* 6:457-465. Bardach, J. E., and J. Atema.
- Barkley, R. A., W. H. Neill, and R. M. Gooding.
1978. Skipjack tuna, *Katsuwonus pelamis*, habitat based on temperature and oxygen requirements. *Fish. Bull., U.S.* 76:653-662.
- Barry, M.
1978. Behavioral response of yellowfin tuna, *Thunnus albacares*, and kawakawa, *Euthynnus affinis*, to turbidity. M.S. thesis, Univ. Hawaii, Honolulu, 31 p. + tables.
- Bone, Q.
1978. Myotomal muscle fiber types in *Scomber* and *Katsuwonus*. *In*: Sharp, G. D., and A. E. Dizon (editors), The physiological ecology of tunas, vol. 3, p. 183-205. Academic Press, NY.
- Bourke, R. E., J. Brock, and R. M. Nakamura.
1987. A study of delayed capture mortality syndrome in skipjack tuna, *Katsuwonus pelamis* (L). *J. Fish Dis.* 10:275-287.
- Brill, R. W.
1978. Temperature effects on speeds of muscle contraction and stasis metabolic rate. *In*: Sharp, G. D., and A. E. Dizon (editors), The physiological ecology of tunas, vol. 5, p. 277- 283. Academic Press, NY.

1979. The effect of body size on the standard metabolic rate of skipjack tuna, *Katsuwonus pelamis*. *Fish. Bull., U.S.* 77:494-498.

1979. The thermal physiology of tuna. Ph.D. dissertation, Univ. Hawaii, Honolulu, 215 p.

1987. On the standard metabolic rates of tropical tunas, including the effect of body size and acute temperature change. *Fish. Bull., U.S.* 85:25-35.
- Brill, R. W., and P. G. Bushnell.
1989. CARDIO--A Lotus 1-2-3 based computer program for rapid calculation of cardiac output from dye dilution or thermal dilution curves. *Comput. Biol. Med.* 19:361-366.

1991. Effects of open and closed system temperature changes on blood-oxygen dissociation curves of yellowfin tuna (*Thunnus albacares*) and skipjack tuna (*Katsuwonus pelamis*). *Can. J. Zool.* 69:1814:1821.

1991. Metabolic and cardiac scope of high energy demand teleosts--the tunas. *Can. J. Zool.* 69:2002-2009.
- Brill, R. W., and A. E. Dizon.
1979. Effect of temperature on isotonic twitch of white muscle and predicted maximum swimming speeds of skipjack tuna, *Katsuwonus pelamis*. *Environ. Biol. Fishes* 4:199-205.
1979. Red and white-muscle fibre activity in swimming skipjack tuna, *Katsuwonus pelamis* (Linnaeus). *J. Fish Biol.* 15:679-685.
- Brill, R. W., D. L. Guernsey, and E. D. Stevens.
1978. Body surface and gill heat loss rates in restrained skipjack tuna. *In: Sharp, G. D., and A. E. Dizon* (editors), *The physiological ecology of tunas*, vol. 4, p. 261-276. Academic Press, NY.
- Brill, R. W., K. N. Holland, and J. S. Ferguson.
1984. Use of ultrasonic telemetry to determine the short-term movements and residence times of tunas around fish aggregating devices. *Proceedings of the Pacific Congress on Marine Technology, PACON84*, p. MRM 1/1-1/7.
- Brill, R. W., R. Bourke, J. A. Brock, and M. D. Dailey.
1987. Prevalence and effects of infection of the dorsal aorta in yellowfin tuna, *Thunnus albacares*, by the larval cestode, *Dasyrhynchus talismani*. *Fish. Bull., U.S.* 85:767-776.
- Brill, R. W., P. G. Bushnell, D. R. Jones, and M. Shimazu.
1992. Effects of temperature change, *in vivo* and *in vitro*, the on acid-base status of blood from yellowfin tuna (*Thunnus albacares*). *Can. J. Zool.* 70:645-662.
- Brown, C. E., and B. S. Muir.
1970. Analysis of ram ventilation of fish gills with application to skipjack tuna (*Katsuwonus pelamis*). *J. Fish. Res. Board Can.* 27:1637-1652.
- Buck, L. T., R. W. Brill, and P. W. Hochachka.
1992. Gluconeogenesis in hepatocytes isolated from skipjack tuna (*Katsuwonus pelamis*). *Can. J. Zool.* 70:1254-1257.
- Bushnell, P. G., and R. W. Brill.
1991. Responses of swimming skipjack (*Katsuwonus pelamis*) and yellowfin (*Thunnus albacares*) tunas and exposed to acute hypoxia, and a model of their cardio-respiratory function. *Physiol. Zool.* 64:787-811.
1992. Oxygen transport and cardiovascular responses in skipjack tuna (*Katsuwonus pelamis*) and yellowfin tuna (*Thunnus albacares*) exposed to acute hypoxia. *J. Comp. Physiol. B.* 162:131-142.
- Bushnell, P. G., R. W. Brill, and R. E. Bourke.
1990. The cardiorespiratory responses of skipjack tuna, *Katsuwonus pelamis*; yellowfin tuna, *Thunnus albacares*; and bigeye tuna, *T. obesus*, to acute reductions of ambient oxygen. *Can. J. Zool.* 68:1857-1865.
- Cahn, P. H.
1972. Sensory factors in the side-to-side spacing and positional orientation of the tuna, *Euthynnus affinis*, during schooling. *Fish. Bull., U.S.* 70:197-204.
- Chang, R. K. C., and J. J. Magnuson.
1968. A radiographic method for determining gas bladder volume of fish. *Copeia* 1968:187-189.

- Dizon, A. E.
1977. Effect of dissolved oxygen concentration and salinity on swimming speed of two species of tunas. *Fish. Bull., U.S.* 75: 649-653.
- Dizon, A. E., and R. W. Brill.
1979. Thermoregulation in tunas. *Am. Zool.* 19:249-265.
1979. Thermoregulation in yellowfin tuna, *Thunnus albacares*. *Physiol. Zool.* 52:581-593.
- Dizon, A. E., R. W. Brill, and H. S. H. Yuen.
1978. Correlations between environment, physiology, and activity and the effects on thermoregulation in skipjack tuna. *In: Sharp, G. D., and A. E. Dizon (editors), The physiological ecology of tunas, vol. 3, p. 233-259. Academic Press, NY.*
- Dizon, A. E., T. C. Byles, and E. D. Stevens.
1976. Perception of abrupt temperature decrease by restrained skipjack tuna, *Katsuwonus pelamis*. *J. Therm. Biol.* 1:185-187.
- Dizon, A. E., W. H. Neill, and J. J. Magnuson.
1977. Rapid temperature compensation of volitional swimming speeds and lethal temperature in tropical tunas (Scombridae). *Environ. Biol. Fishes* 2:83-92.
- Dizon, A. E., and G. D. Sharp.
1978. Perspectives: The past, present, and future of tuna physiology. *In: Sharp, G. D., and A. E. Dizon (editors), The physiological ecology of tunas, vol. 2, p. 451-458. Academic Press, NY.*
- Dizon, A. E., E. D. Stevens, W. H. Neill, and J. J. Magnuson.
1974. Sensitivity of restrained skipjack tuna (*Katsuwonus pelamis*) to abrupt increases in temperature. *Comp. Biochem. Physiol.* 49A:291-299.
- Farrell, A. P., P. S. Davie, C. E. Franklin, J. Johansen, and R. W. Brill.
1992. Cardiac physiology in tunas: I. In vitro perfused heart preparations in yellowfin and skipjack tunas. *Can. J. Zool.* 70:1200-1210.
- Fierstine, H. L., and V. Walters.
1968. Studies in locomotion and anatomy of scombrid fishes. *Mem. South. Calif. Acad. Sci.* 6:1-31.
- Gooding, R. M.
1963. The olfactory organ of the skipjack *Katsuwonus pelamis*. *Proceedings of the World Scientific Meeting on the Biology of Tunas and Related Species, La Jolla, Calif., 2-14 July 1962. FAO Fish. Rep.* 6, 3:1621-1631.
- Gooding, R. M., W. H. Neill, and A. E. Dizon.
1981. Respiration rate and low-oxygen tolerance limits in skipjack tuna, *Katsuwonus pelamis*. *Fish. Bull., U.S.* 79:31-48.
- Gordon, M. S.
1968. Oxygen consumption of red and white muscles from tuna fishes. *Science (Wash., D.C.)* 159:87-90.
1972. Comparative studies on the metabolism of shallow-water and deep-sea marine fishes. I. White-muscle metabolism in shallow-water fishes. *Mar. Biol. (Berl.)* 13:222-237.
1972. Comparative studies on the metabolism of shallow-water and deep-sea marine fishes. II. Red-muscle metabolism in shallow-water fishes. *Mar. Biol. (Berl.)* 15:246-250.

- Graham, J. B., H. Dewar, N. Chin Lai, P. A. Fields, R. Shabetai, and R. W. Brill.
In press. Swimming physiology of pelagic fishes. Proceedings of Society of Experimental Biology Symposium.
- Guppy, M.
1978. Skipjack tuna white muscle: A blueprint for the integration of aerobic and anaerobic carbohydrate metabolism. *In: Sharp, G. D., and A. E. Dizon (editors), The physiological ecology of tunas, vol. 2, p. 175-181. Academic Press, NY.*
- Guppy, M., and P. W. Hochachka.
1978. Controlling the highest lactate dehydrogenase activity known in nature. *Am. J. Physiol.* 234:R136-R140.

1979. Pyruvate kinase functions in hot and cold organs of tuna. *J. Comp. Physiol.* 129:185-191.
- Hanyu, I., T. Tamura, and H. Niwa.
1973. Electroretinograms and retinal ganglion cell responses in the skipjack tuna. *Bull. Jpn. Soc. Sci. Fish.* 39:265-273.
- Hochachka, P. W., and R. W. Brill.
1987. Autocatalytic pathways to cell death: a new analysis of the tuna burn problem. *Fish Physiol. Biochem.* 4(2):81-87.
- Hochachka, P. W., W. C. Hulbert, and M. Guppy.
1978. The tuna power plant and furnace. *In: Sharp, G. D., and A. E. Dizon (editors), The physiological ecology of tunas, vol. 1, p. 153-174. Academic Press, NY.*
- Holland, K.N., R.W. Brill, J. S. Ferguson, and R. Yost.
1986. A small vessel technique for tracking pelagic fish. *Marine Fisheries Review.* 47(4):26-32.
- Holland, K. N., R. W. Brill, and R. K. C. Chang.
1990. Horizontal and vertical movements of tunas (*Thunnus* spp.) associated with fish aggregating devices. *Fish. Bull., U.S.* 88:493-507.
- Holland, K. N., R. W. Brill, R. K. C. Chang, and R. Harman.
1990. Horizontal and vertical movements of Pacific blue marlin captured using sport fishing techniques. *Fish. Bull., U.S.* 88:397-402.
- Holland, K. N., R. Brill, R. Chang, J. Sibert, and D. Fournien.
In Press. Physiology and behavioral thermoregulation in bigeye tuna (*Thunnus obesus*). *Nature*.
- Hunter, J. R., B. J. Macewics, and J. R. Sibert.
1986. The spawning frequency of skipjack tuna, *Katsuwonus pelamis*, from the south Pacific. *Fish. Bull., U.S.* 84:895-903.
- Ikehara, W., J. Atema, A. Brittain, J. Bardach, A. Dizon, and K. Holland.
1978. Reactions of yellowfin tuna to prey scents. [Abstract] *Pac. Sci.* 32:97.
- Ikehara, W. N., and J. E. Bardach.
1981. Chemosensory attracting and guiding of yellowfin tuna, *Thunnus albacares*. Southwest Fish. Cent. La Jolla Lab, Natl. Mar. Fish. Serv., NOAA, La Jolla, CA 92038, Southwest Fish. Cent. Admin. Rep. LJ-81-07C.

Iversen, R. T. B.

1966. Hearing in tunas with special reference to *Euthynnus yaito* Kishinouye. [Abstract] Abstracts of Papers Related With Fisheries, Marine and Freshwater Science. Divisional Meeting on Fisheries Sciences, p. 25. Proc. 11 Pac. Sci. Congr., Tokyo, 1966, vol. 7. Iversen, R. T. B., and J. O. Puffinburger.

1967. Response of yellowfin tuna (*Thunnus albacares*) to underwater sound. In: Tavalga, W. N. (editor), Marine bio-acoustics 2:105-119, discussion, p. 119-121. Proceedings of the Second Symposium on Marine Bio-Acoustics held at the American Museum of Natural History, New York, April 13-15, 1966. Pergamon Press, Oxford and NY.

1969. Auditory thresholds of the scombrid fish *Euthynnus affinis*. with comments on the use of sound in tuna fishing. In: Ben-Tuvia, A., and W. Dickson (editors), Proceedings of the FAO Conference on Fish Behaviour in Relation to Fishing Techniques and Tactics, Bergen, Norway, 19-27 October 1967. FAO Fish. Rep. 62, 3:849-859.

1977. Capture, transportation, and pumping of threadfin shad, *Dorosoma petenense*. In: Shomura, R. S. (editor), Collection of tuna baitfish papers, NOAA Tech. Rep. NMFS Circ. 408., Natl. Oceanic Atmos. Adm., Natl. Mar. Fish. Serv., p. 127-136. Jemison, H. A., III, A. E. Dizon, and M. M. Walker.

Jamison, H. A., A. E. Dizon, and M. M. Walker

1982. An automatic feeder for liquids, or wet or dry solids. Behav. Res. Methods Instrum. 14:54-55.

Johnston, I. A., and R. W. Brill.

1984. Thermal dependence of contractile properties of single skinned muscle fibres from antarctic and various warm water marine fishes including skipjack tuna (*Katsuwonus pelamis*) and kawakawa (*Euthynnus affinis*). J. Comp. Physiol. B Biochem. Syst. Environ. Physiol. [Metab. Transp. Funct.] 155:63-70.

Jokiel, P. L.

1989. Effects of marine mining dredge spoils on eggs and larvae of a commercially important species of fish, the mahimahi, *Coryphaena hippurus*. Mar. Min. 8:303-315.

Jokiel, P. L., R. Y. Ito, and T. K. Kazama.

1986. A preliminary study on the effects of suspended sediment on the survival of early life stages of the mahimahi, *Coryphaena hippurus*. Southwest Fish. Cent. Honolulu Laboratory, Natl. Mar. Fish. Serv., NOAA, 2570 Dole St., Honolulu, HI 96822-2396. Southwest Fish. Cent. Admin. Rep. H-86-18, 13 p.

Jones, D. R., R. W. Brill, P. J. Butler, P. G. Bushnell, and M. R. A. Heieis.

1989. Measurement of ventilation volume in swimming tunas. J. Exp. Biol. 149:491-498

Jones, D. R., R. W. Brill, and D. C. Mense.

1986. The influence of blood gas properties on gas tensions and pH of ventral and dorsal aortic blood in free-swimming tuna, *Euthynnus affinis*. J. Exp. Biol. 120:201-213.

Jones, D. R., R. W. Brill, and P. G. Bushnell.

In press. Ventricular and arterial dynamics of anesthetized and swimming tunas. J. Exp. Biol.

Kashin, S. M., R. W. Brill, W. N. Ikehara, and A. E. Dizon.

1981. Induced locomotion by midbrain stimulation in restrained skipjack tuna, *Katsuwonus pelamis*. J. Exp. Zool. 216:327-329.

Kaya, C. M., A. E. Dizon, and S. D. Hendrix.

1981. Induced spawning of tuna, *Euthynnus affinis*. Fish. Bull., U.S. 79:185-187.

- Kaya, C. M., A. E. Dizon, S. D. Hendrix, T. K. Kazama, and M. K. K. Queenth.
1982. Rapid and spontaneous maturation, ovulation, and spawning of ova by newly captured skipjack tuna, *Katsuwonus pelamis*. Fish. Bull., U.S. 80:393-396.
- Kaya, C. M., M. K. K. Queenth, and A. E. Dizon.
1984. Capture and restraining technique for experimental work on small tuna in large laboratory holding tanks. Prog. Fish-Cult. 46(4):288-290.
- Keen, J. E., A. P. Farrell, G. F. Tibbits, and R. W. Brill.
1992. Cardiac dynamics in tunas. II. Effect of ryanodine, calcium, and adrenaline on force-frequency relationships in atrial strips from skipjack tuna., *Katsuwonus pelamis*. Can. J. Zool. 70:1211-1217.
- Kitchell, J. F., J. J. Magnuson, and W. H. Neill.
1977. Estimation of caloric content for fish biomass. Environ. Biol. Fishes 2:185-188.
- Kitchell, J. F., W. H. Neill, A. E. Dizon, and J. J. Magnuson.
1978. Bioenergetic spectra of skipjack and yellowfin tunas. In: Sharp, G. D., and A. E. Dizon (editors), The physiological ecology of tunas, vol. 3, p. 357-368. Academic Press, NY.
- Lamadrid-Rose, Y., and G. W. Boehlert.
1988. Effects of cold shock on egg, larval, and juvenile stages of tropical fishes: Potential impacts of ocean thermal energy conversion. Mar. Environ. Res. 25:175-193.
- Magnuson, J. J.
1963. Tuna behavior and physiology, a review. Proceedings of the World Scientific Meeting on the Biology of Tunas and Related Species, La Jolla, Calif., 2-14 July 1962. FAO Fish. Rep. 6, 3:1057-1066.
1964. Activity patterns of scombrids. [Abstracts] Proc. Hawaiian Acad. Sci., 39 Annu. Meet., 1963-1964, p. 26.
1964. Salt well water facilities at the Bureau of Commercial Fisheries Biological Laboratory, Honolulu. In: Clark, J. R., and R. L. Clark (editors), A collection of papers on sea-water systems for experimental aquariums. U.S. Fish Wildl. Serv., Res. Rep. 63:169-172.
1964. Tuna behavior program at Honolulu. In: Modern fishing gear of the world 2:560-562. Fish. News (Books) Ltd., Lond.
1965. Tank facilities for tuna behavior studies. Prog. Fish-Cult. 27:230-233.
1966. A comparative study of the function of continuous swimming by scombrid fishes. [Abstract] Abstracts of Papers Related With Fisheries, Marine and Freshwater Science. Symposium on Biological Studies of Tunas and Sharks in the Pacific Ocean, p. 15. Proc. 11 Pac. Sci. Congr., Tokyo, 1966, vol. 7.
1969. Digestion and food consumption by skipjack tuna (*Katsuwonus pelamis*). Trans. Am. Fish. Soc. 98:379-392.
1969. Swimming activity of the scombrid fish *Euthynnus affinis* as related to search for food. In: Ben-Tuvia, A., and W. Dickson (editors), Proceedings of the FAO Conference on Fish Behaviour in Relation to Fishing. Techniques and Tactics, Bergen, Norway, 19-27 October 1967. FAO Fish. Rep. 62, 2:439-451.
1970. Hydrostatic equilibrium *Euthynnus affinis*, a pelagic teleost without a gas bladder. Copeia 1970:56-85.

1978. Locomotion by scombrid fishes: Hydromechanics, morphology, and behavior. *In*: Hoar, W. S., and D. J. Randall (editors), *Fish physiology*, vol. 7, p. 240-315. Academic Press, NY. @LIT/L2 = 1973. Comparative study of adaptations for continuous swimming and hydrostatic equilibrium of scombroid and xiphoid fishes. *Fish. Bull.*, U.S. 71:337-356.
- Manar, T. A.
1965. Tuna behavior. A growing field for research. *Pac. Fisherm.* 63(11):9-11.
- Marr, J. C.
1967. Research programme of the U.S. Bureau of Commercial Fisheries Biological Laboratory, Honolulu, Hawaii. *Proceedings of the Symposium on Scombroid Fishes, Part 3*, p. 1154-1157. *Mar. Biol. Assoc. India. Symp. Ser.* 1.
- Mathieu-Costello, O., P. J. Agey, R. B. Logermann, R. W. Brill, and P. W. Hochachka.
1992. Capillary-to-fiber geometrical relationships in tuna red muscle. *Can. J. Zool.* 70:1218-1229.
- Milsom, W.K. and R.W. Brill.
1986. Oxygen sensitive afferent information arising from the first gill arch of yellowfin tuna. *Resp. Physiol.* 66:193-203.
- Monckton, E. A.
1978. Stress in captive skipjack tuna, *Katsuwonus pelamis*. M.S. thesis, Univ. Hawaii, Honolulu, 13 p. + tables.
- Moon, T. W., R. W. Brill, P. W. Hochachka, and J.- M. Weber.
1987. L- (+)lactate transportation into the red blood cells of the skipjack tuna (*Katsuwonus pelamis*). *Can. J. Zool.* 65:2570-2573.
- Moyes, C. D., O. A. Mathieu-Costello, R. W. Brill, and P. W. Hochachka.
1992. Mitochondrial differences in muscles from a high performance species, skipjack tuna, (*Katsuwonus pelamis*) and a low performance species, common carp, (*Cyprinus carpio*). *Can. J. Zool.* 70:1246-1253.
- Muir, B. S., and C. E. Brown.
1971. Effects of blood pathway on the blood pressure drop in fish gills, with special reference to tunas. *J. Fish. Res. Board Can.* 28:947-955.
- Murchison, A. E., and J. J. Magnuson.
1966. Notes on the coloration and behavior of the common dolphin, *Coryphaena hippurus*. *Pac. Sci.* 20:515-517.
- Nakamura, E. L.
1960. Confinement of skipjack in a pond. [Abstract] *Proc. Hawaiian Acad. Sci.*, 35 Annu. Meet. 1959-1960, p. 24-25.
1962. Observations on the behavior of skipjack tuna, *Euthynnus pelamis*, in captivity. *Copeia* 1962:499-505.
1964. A method of measuring visual acuity of scombrids. [Abstract] *Proc. Hawaiian Acad. Sci.*, 39 Annu. Meet., 1963-1964, p. 26-27.
1966. Fiberglass tanks for transferring of pelagic fishes. *Prog. Fish-Cult.* 28:60-62.
1968. Visual acuity of two tunas, *Katsuwonus pelamis* and *Euthynnus affinis*. *Copeia* 1968: 41-49.

1969. A review of field observations on tuna behavior. *In*: Ben-Tuvia, A., and W. Dickson (editors), Proceedings of the FAO Conference on Fish Behaviour in Relation to Fishing Techniques and Tactics, Bergen, Norway, 19-27 October 1967. FAO Fish. Rep. 62, 2:59-68.
1969. Visual acuity of yellowfin tuna, *Thunnus albacares*. *In*: Ben-Tuvia, A., and W. Dickson (editors), Proceedings of the FAO Conference on Fish Behaviour in Relation to Fishing Techniques and Tactics, Bergen, Norway, 19-27 October 1967. FAO Fish. Rep. 62, 3:463-468.
1972. Development and uses of facilities for studying tuna behavior. *In*: Winn, H. E., and B. L. Olla (editors), Behavior of marine animals. Current perspectives in research, vol. 2, s, p. Vertebrates, p. 245-277. Plenum Publ., NY.
- Nakamura, E. L., and J. J. Magnuson.
1965. Coloration of the scombrid fish *Euthynnus affinis* (Cantor). *Copeia* 1965:234-235.
- Neill, W. H., and T. C. Byles.
1972. Automatic pellet dispenser for experimental feeding of fishes. *Prog. Fish-Cult.* 34:170.
- Neill, W. H., R. K. C. Chang, and A. E. Dizon.
1976. Magnitude and ecological implications of thermal inertia in skipjack tuna, *Katsuwonus pelamis* (Linnaeus). *Environ. Biol. Fishes* 1:61-80.
- Neill, W. H., J. J. Magnuson, and G. D. Chipman.
1972. Behavioral thermoregulation by fishes: A new experimental approach. *Science* (Wash., D.C.) 176:1433-1445.
- Neill, W. H., and E. D. Stevens.
1974. Thermal inertia versus thermoregulation in "warm" turtles and tunas. *Science* (Wash., D.C.) 184:1008-1010.
- Perry, S. F., C. Daxboeck, B. Emmett, P. W. Hochachka, and R. W. Brill.
1985. Effects of exhausting exercise on acid-base regulation in skipjack tuna (*Katsuwonus pelamis*) blood. *Physiol. Zool.* 58:421-429.
1985. Effects of temperature change on regulation in skipjack (*Katsuwonus pelamis*) blood. *Comp. Biochem. Physiol. A Comp. Physiol.* 81A:49-53.
- Radtke, R. C.
Otolith formation and increment deposition in laboratory-reared skipjack tuna, *Euthynnus pelamis*, larvae. *In*: Prince, E. D., and L. K. Pulos (editors), Proceedings of the International Workshop on Age Determination of Oceanic Pelagic Fishes: Tunas, Billfishes, and Sharks, p. 99-103. U.S. Dep. Commer., NOAA Tech. Rep. NMFS 8.
- Rayner, M. D., and M. J. Keenan.
1967. Role of red and white muscles in the swimming of the skipjack tuna. *Nature* (Lond.) 214:392-393.
- Sather, B. T., and T. A. Rogers.
1967. Some inorganic constituents of the muscles and blood of the oceanic skipjack, *Katsuwonus pelamis*. *Pac. Sci.* 21:404-413.
- Shaklee, J.B., R.W. Brill, and R. Acerra.
1983. Biochemical genetics of Pacific blue marlin, *Makaira nigricans*, from Hawaiian waters. *Fish. Bull.*, U.S. 81:85-90.

- Steffel, S.
1973. Temperature discrimination thresholds in a tuna, the kawakawa (*Euthynnus affinis*), as determined by operant conditioning. M.S. thesis, Univ. Wisconsin, Madison, 52 p.
- Steffel, S., A. E. Dizon, J. J. Magnuson, and W. H. Neill.
1976. Temperature discrimination by captive free-swimming tuna, *Euthynnus affinis*. *Trans. Am. Fish. Soc.* 105:588-591.
- Stevens, E. D.
1972. The effect of changes in ambient temperature on spontaneous activity in skipjack tuna. *Comp. Biochem. Physiol.* 42A:803-805.
1972. Some aspects of gas exchange in tuna. *J. Exp. Biol.* 56:809-823.
- Stevens, E. D., and F. E. J. Fry.
1971. Brain and muscle temperatures in ocean caught and captive skipjack tuna. *Comp. Biochem. Physiol.* 38A:203-211.
- Struhsaker, P., and J. Uchiyama.
1976. Age and growth of the nehu, *Stolephorus purpureus* (Pices:Engraulidae, from the Hawaiian Islands as indicated by dialy growth increments of sagittae. *Fish. Bull. U.S.* 74-9-17.
- Suckling, E. E.
1967. Electrophysiological studies on the trunk lateral line system of various marine and freshwater teleosts. *In: Cahn, P. H. (editor), Lateral line detectors*, p. 97-103. Indiana Univ. Press, Bloomington.
- Suckling, J. A.
1967. Trunk lateral line nerves: Some anatomical aspects. *In: Cahn, P. H. (editor), Lateral line detectors*, p. 45-52. Indiana Univ. Press, Bloomington.
- Tamura, T., I. Hanyu, and H. Niwa.
1972. Spectral sensitivity and color vision in skipjack tuna and related species. *Bull. Jpn. Soc. Sci. Fish.* 38:799-802.
- Tester, A. L.
1959. Summary of experiments on the response of tuna to stimuli. *In: Kristjonnsson, H. (editor), Modern fishing gear of the world*, p. 538-542. Fish. News (Books) Ltd., Lond.
- Tibbits, G. F., H. Kashihara, and R. W. Brill.
1992. Properties of myocardial sarcolemma isolated from skipjack tuna, *Katsuwonus pelamis*. *Can. J. Zool.* 70:1240-1245.
- Uchiyama, J. H., and P. Struhsaker.
1981. Age and growth of skipjack tuna, *Katsuwonus pelamis* and yellowfin tuna, *Thunnus albacares*, as indicated by daily growth increments of sagittae. *Fish. Bull., U.S.* 79:151-162.
- Walker, M. M.
1982. Conditioned response for use in magnetic sensory discrimination studies in yellowfin tuna, *Thunnus albacares*. (Abstr.) Tenth Annual Conference of the International Marine Animal Trainers Association (IMATA), Honolulu, Hawaii, October 25-29, 1982. (Mimeogr.)
1984. Learned magnetic field discrimination in yellowfin tuna, *Thunnus albacares*. *J. Comp. Physiol.* A155:673-679.

1984. Magnetic sensitivity and its possible physical basis in the yellowfin tuna, *Thunnus albacares*. In: McCleave, J. D., G. P. Arnold, J. J. Dodson, and W. H. Neill (editors), Mechanisms of migration in fishes, p. 123-141. Plenum Press, NY (NATO Conference Ser. IV. Marine Science 14.)
- Walker, M. M., A. E. Dizon, and J. L. Kirschvink.
1982. Geomagnetic field detection by yellowfin tuna. *Oceans* 82, p. 755-758. Conference sponsored by Marine Technology Society, IEEE Council on Oceanic Engineering, Wash., D.C., September 20-22, 1982.
- Walker, M. M., J. L. Kirschvink, R. S.-B. Chang, and A. E. Dizon.
1984. A candidate magnetic sense organ in the yellowfin tuna, *Thunnus albacares*. *Science* (Wash., D.C.) 224:751-753.
- Walker, M. M., J. L. Kirschvink, and A. E. Dizon.
1985. Magnetoreception and biomineralization of magnetite, fish. In: Kirschvink, J. L., D. S. Jones, and B. J. MacFadden (editors), Magnetite biomineralization and magnetoreception in organisms, a new magnetism, p. 417-437. Plenum Publ., NY.
- Walker, M. M., J. L. Kirschvink, A. Perry, and A. E. Dizon.
1985. Detection, extraction, and characterization of biogenic magnetite. In: Kirschvink, J. L., D. S. Jones, and B. J. MacFadden (editors), Magnetite biomineralization and magnetoreception in organisms, a new biomagnetism, p. 155-166. Plenum Press, NY.
- Walters, V.
1966. On the dynamics of filter feeding by the wavyback skipjack (*Euthynnus affinis*). *Bull. Mar. Sci.* 16:209-221.
- Watson, C., R. Bourke, and R. W. Brill.
1988. A comprehensive theory on the etiology of burnt tuna. *Fish. Bull.*, U.S. 86:376-3.
- Watson, C. L., H. Morrow, and R. W. Brill.
In press. Proteolysis of skeletal muscle in yellowfin tuna: Evidence of calpain activation. *Comp. Biochem. Physiol.*
- Weber, J.-M., R. W. Brill, and P. W. Hochachka.
1986. Mammalian metabolic flux rates in a teleost: lactate and glucose turnover tuna. *Am. J. Physiol.* 250: R452-R458.

RECENT TECHNICAL MEMORANDUMS

Copies of this and other NOAA Technical Memorandums are available from the National Technical Information Service, 5285 Port Royal Road, Springfield, VA 22167. Paper copies vary in price. Microfiche copies cost \$4.50. Recent issues of NOAA Technical Memorandums from the NMFS Southwest Fisheries Science Center are listed below:

- NOAA-TM-NMFS-SWFSC- 161 Report of ecosystem studies conducted during the 1990 eastern tropical Pacific dolphin survey on the research vessel *McArthur*.
V.A. PHILBRICK, P.C. FIEDLER, S.B. REILLY, R.L. PITMAN, L.T. BALLANCE and D.W. BEHRINGER
(May, 1991)
- 162 Predicting sablefish age using otolith characteristics.
A. McBRIDE and J.E. HIGHTOWER
(August, 1991)
- 163 CHARTOPS: simulating short-term use of the tuna purse-seine fleet to survey dolphin schools in the eastern tropical Pacific Ocean.
E.F. EDWARDS and P. KLEIBER
(August, 1991)
- 164 Results of the southern California sportfish economic survey.
C.J. THOMSON and S.J. CROOKE
(August, 1991)
- 165 Status of Pacific oceanic fishery resources of interest to the USA for 1991.
STAFF OF THE SOUTHWEST FISHERIES SCIENCE CENTER
(September, 1991)
- 166 Methods used to identify pelagic juvenile rockfish (Genus *Sebastes*) occurring along the coast of central California.
EDITED BY T.E. LAIDIG and P.B. ADAMS
(November, 1991)
- 167 Spatial and temporal variability in growth of widow rockfish (*Sebastes entomelas*)
D.E. PEARSON and J.E. HIGHTOWER
(December, 1991)
- 168 Documentation of three computer programs used to assess daily age from the hard structures of animals.
T.E. LAIDIG and D.E. PEARSON
(June, 1992)
- 169 Report of a marine mammal survey of the California coast aboard the research vessel *McArthur*, July 28-November 5, 1991.
P.S. HILL and J. BARLOW
(July, 1992)
- 170 A program for the Microsoft Windows environment to collect analog-to-digital and serial communication data on a personal computer based system.
R.C. HOLLAND
(July 1992)