Sampling Collection on Board



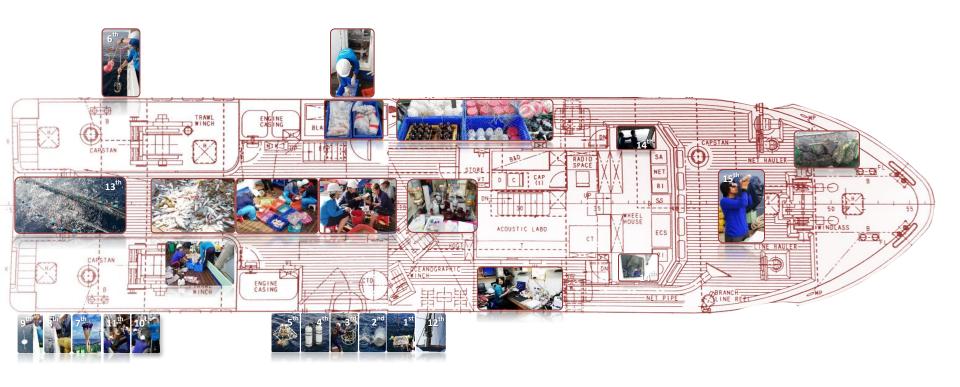
Supapong Pattarapongpan Fishery Oceanographer

M.V. SEAFDEC 2



M.V. SEAFDEC 2

Working space on M.V. SEAFDEC 2



Sampling collection onboard

Introduction

- Sampling collection are important for research
 - Objectives
 - Species composition
 - Juvenile rearing areas or general distribution
 - Spawning areas or general distribution
 - Sizes (length) of adult and juvenile
 - Relative abundance in selected areas
 - Method
 - Budget / Time / Effort

Oceanography

Soon after ship arrive station

- Secchi disc and Forel-Ule
 Scale
- CTD
- Water sampling
- Sampling for sediment and benthos
- Neuston net
- Bongo net



OCEANOGRAPHIC LOGSHEET



Oceanographic logsheet

- Cruise detail
- Station number
- Date and time
- Water depth
- Latitude, Longitude
- Vessel data
- Air
- Water, current
- **Bottom**
- Equipment

Record by	Sukchai A.								
Project name:		Name of vessel				Water			
Cruise no:	18-2/2006		M.V. SEAFDEC 2			Temperature:	29	° C	
Station no:	08(30)	Start (28-Feb-06) Finish(28-Feb-06)		Color:	7				
Date:	28-Feb-06	Time	13:55	Time	14:48	Transparency:	23co	os50 m	
Sounding depth	Sounding depth: 68 m.		05_29.95 N	Lat.	05_29.66 N	Current			
Air		Long.	099_29.05 E	Long.	099_28.70 E	Depth	Speed(Knts)	Direction	
Wind						Бери	Speed(Rins)	Direction	
Speed(Knts)	Direction	Bottom			surface	0.8	128		
8	50	Type: muddy		35 m	0.7	90			
Air temp: 29.06 °C		Color: black			m	N/A	N/A		
Air press: 1012.0 hpa		Smell: -			Memorandum:	:			
Humidity:	Humidity: 78 %		Temperature: - ° C						
Weather condit	Weather condition: partly cloudy		pH: -						
Stage of sea:	smoot	1			ŀ				

NR: not be recorder

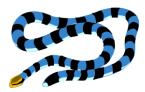
			Oceanographic ed	minmont				
CTD system		Doto filo no	0 1	Juihment	211000	2 10000		
		Data file name		s2d18008,s2u18008				
TSG system		Data file name		20060228(2),20060301(1):FronSt.07-St.08				
PRR system		Data file name						
BT data acquisi	ition system	Data file name						
Temp & Depth Recorder		Data file name						
Vandorn water sampler		List of sampling depth						
Sediment Piston core		No. of operation		8				
1	Smith McIntyre grab	No. of operation		8				
		Towing Towing speed		Start		Finish		
		depth (m.)	(Knots)	Time	13:55	Time	14:04	
		35	1.5-3	Lat.	05 29.95 N	Lat.	05_30.11 N	
				Long.		Long.	099 28.94 E	
Bongo	o net(oblique)	No. at flow meter		<u> </u>	Fish Larvae		olankton	
				21,056		10,833		
					21,030	1,	0,833	
1								
			neter Calibration					
		((cycle/m ³)					
		Towing	(cycle/m³) Towing speed		Start		inish	
		((cycle/m ³)	Time	Start 14:11	Time	Tinish 14:21	
		Towing depth (m.)	(cycle/m³) Towing speed (Knots)	Time Lat.			T	
		Towing	(cycle/m³) Towing speed		14:11	Time Lat.	14:21	
Bongo i	net(horizontal)	Towing depth (m.)	(cycle/m³) Towing speed (Knots)	Lat. Long.	14:11 05_29.95 N	Time Lat. Long.	14:21 05_29.99 N	
Bongo i	net(horizontal)	Towing depth (m.)	(cycle/m³) Towing speed (Knots)	Lat. Long.	14:11 05_29.95 N 099_29.00 E	Time Lat. Long. Zoop	14:21 05_29.99 N 099_28.93 E blankton	
Bongo i	net(horizontal)	Towing depth (m.) surface	(cycle/m³) Towing speed (Knots) 2 at flow meter	Lat. Long.	14:11 05_29.95 N 099_29.00 E	Time Lat. Long. Zoop	14:21 05_29.99 N 099_28.93 E	
Bongo i	net(horizontal)	Towing depth (m.) surface No.	(cycle/m³) Towing speed (Knots) 2 at flow meter neter Calibration	Lat. Long.	14:11 05_29.95 N 099_29.00 E	Time Lat. Long. Zoop	14:21 05_29.99 N 099_28.93 E blankton	
Bongo i	net(horizontal)	Towing depth (m.) surface No.	(cycle/m³) Towing speed (Knots) 2 at flow meter	Lat. Long.	14:11 05_29.95 N 099_29.00 E	Time Lat. Long. Zoop	14:21 05_29.99 N 099_28.93 E blankton	



Catch from trawl

- The basic feature of any good sampling systems is " a random sample"
- Deck sampling and recording procedure(FAO)

 Step 1: Remove all sea snakes and other venomous or otherwise dangerous animals.











- Step 2: Remove inorganic debris and plant material.
 Record type of material removed.
- Step 3: Remove the larger fish that are readily visible and place them in a box

Catch from trawl

 Step 4: Wash the remainder of catch (Small fish) if necessary, and mixes with shovel.



- Step 5: Put the mix catch in box while continuing to remove larger fish and putting them into the box mentioned in step3. The box should be filled simultaneously, not one after the other, and it should be made certain that all boxes contain approximately the same weight of fish.
- Step 6: Count the number of boxes with small fish and record.

- Step 7: A rule of thumb, is to take one box out of every five at random for sub-sampling. Record number of boxes taken to sub-sampling as B1, B2, B3,.....
- Step 8: The boxes taken for sub-sampling is (are) then treated as follow:
 - Weight total catch B1 and record.
 - Place fish of B1 on sorting table and sort to species as far as food fishes and valuable crustacean (e.g. shrimp) are concern and taxonomic grouping as well- defined as possible (e.g. genus, family) for other group (the non-edible fish and miscellaneous).
 Repeat procedure if appropriate for the other boxes B2, B3,...

- Step 9: If more than one box was sorted, compute, for each species (or higher taxonomy group) the total weight and number in all sorted boxes
- Step 10: Multiply the number and weight of fish and invertebrate by species (or higher taxonomic group) by the ratio of the number of unsorted to sorted boxes.
- Step 11: Weight and count the larger fish mention in step 3 and 5 by species, very large fish should be weighed individually and measure.



- Step 12: Add, when there is an overlap (when the fish of a certain species occurred both in the sorted boxes of small fish in the large fish box) the weights and number obtained in step 11 to weight and number in step 10
- Step 13: step 12 (as well as step 11 when there is no overlap) provided estimates of total catch, both in weight and number, by species and higher taxonomic groups. Record the total, both in weight and number in to the appropriate the fishing log sheet and convert to catch per unit if fishing time is less or morn than an hour. During surveys, this step must be complete after each haul every evening at the latest to preclude loss of information.

- Step 14: In addition to catch sampling, identifying and recording, the work of the fishery scientist general include among other things:
 - Collecting length-frequency data
 - Collecting miscellaneous biological information
 - Collecting and preserving specimens for further study onshore
 - Collecting oceanographic data

Pre-fixation

- Equipment
 - Pins
 - Plastic foam boards and food trays
 - Paintbrush
 - Formalin
- Preparation for fin spreading
 - Before starting fin spreading, the fishes are rinsed in water to remove the mucilaginous solution and dirt adhering

Pre-fixation

- Fin spreading
 - Positioning of the fish
 - The fish body is laid with the side facing up for fin spreading.
 - Fin spreading
 - Thin pins should be used during fin spreading to prevent damage to the fish membrane.

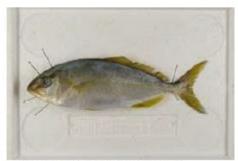
Pre-fixation



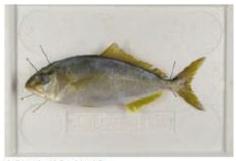
1. Fixed body axis.



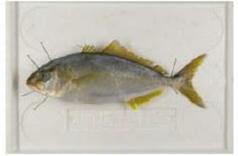
2. Spread and fixed caudal fin.



3. Spread and fixed dorsal fin ray from the rear.



4. Spread and fixed anal fin.



5. Spread and fixed dorsal spiny ray.



6. Spread and fixed pelvic fin.

Motomura and Ishikawa (2013)

Pre-fixation

- Application of formalin
 - A concentrated formalin solution is applied around the after spreading all fin.
 - Formalin fixation normally takes 5-10 min for small fish.
 (Smaller than 10 cm.)
 - Large fish specimens can be fixed in approximately 15 min.
 - To prevent the fish body from drying after application of formalin, it should be covered with a paper towel and sprayed with plenty water
 - Rinsing
 - After formalin fixation, the fish body is re-washed and prepared for obtaining photographs

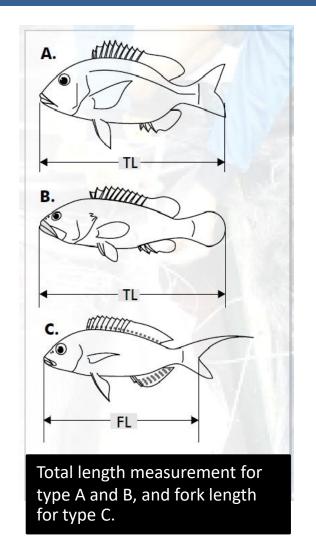
Pre-fixation

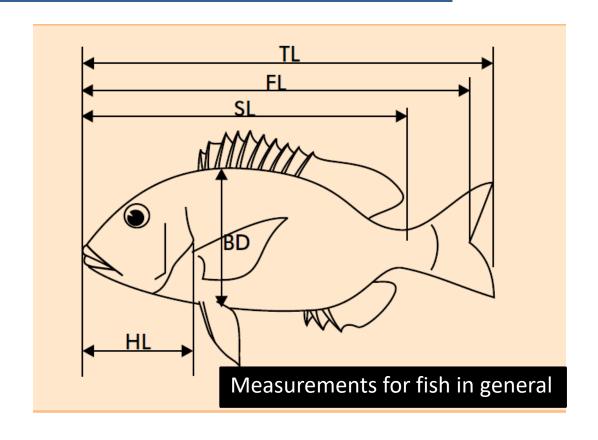


Application of formalin solution.

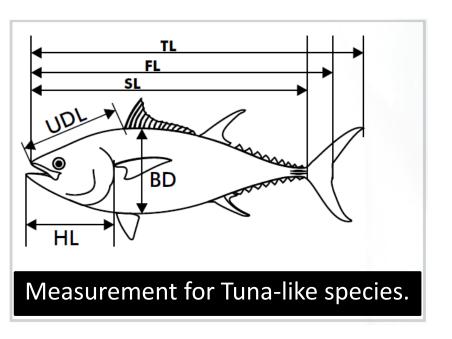


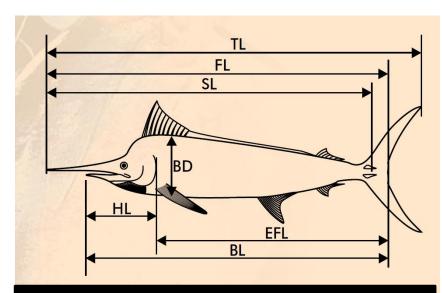
Fish body covered with a wet paper towel, sprayed with water to prevent drying of the fish.



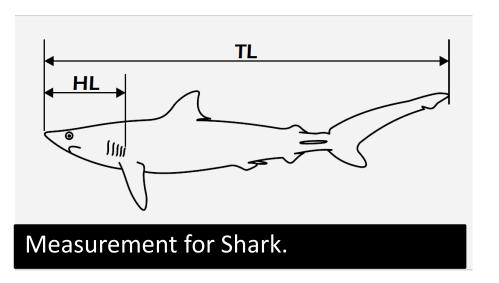


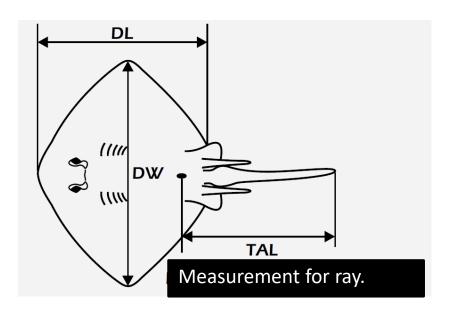
Length Measurement

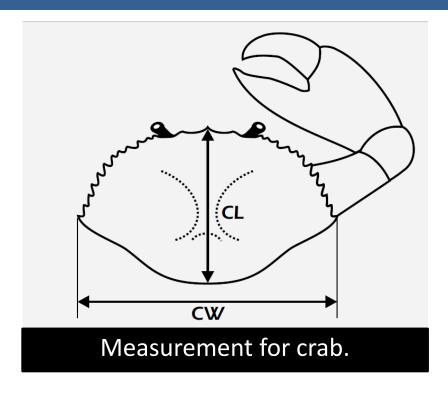


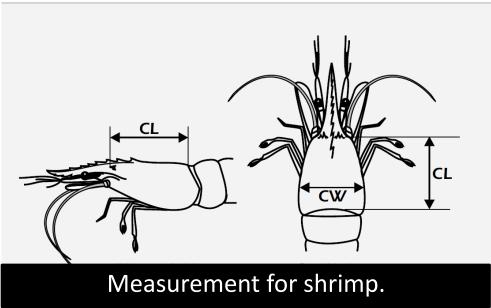


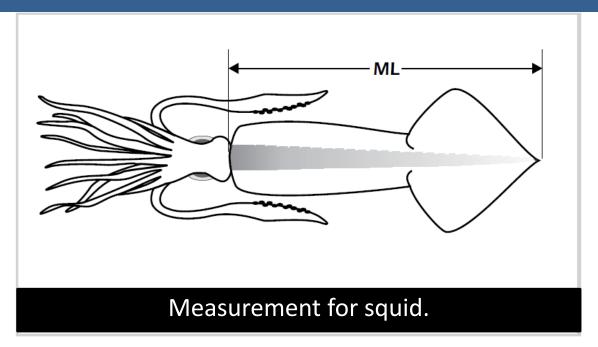
Measurement for marlin and sailfish species.



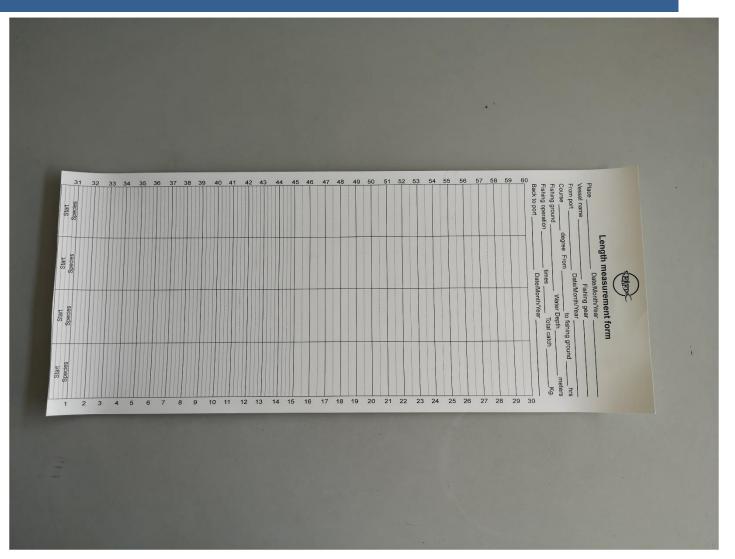








Punching Sheet



Specimen Preservation

- Specimens preservation means "long term preservation of organism either plant or animal is the best possible condition."
- Step for specimens preservation
 - Killed and relaxing of animal
 - Fixation
 - Storage

Specimen Preservation

Formalin

- It is use for vertebrates only.
- It is avoided for the long term storage
- Mostly formalin is used where colour is important since alcohol dissolves most colour immediately.
- Dilution
 - Concentrate formalin 100% = water saturated 40% with formaldehyde
 - 10% formalin = 4% formaldehyde (used for preservation)
 - Mix one part concentrated formalin to nine part water.
 - Fill about two-thirds the bottle's volume with 10% formalin

Specimen Preservation

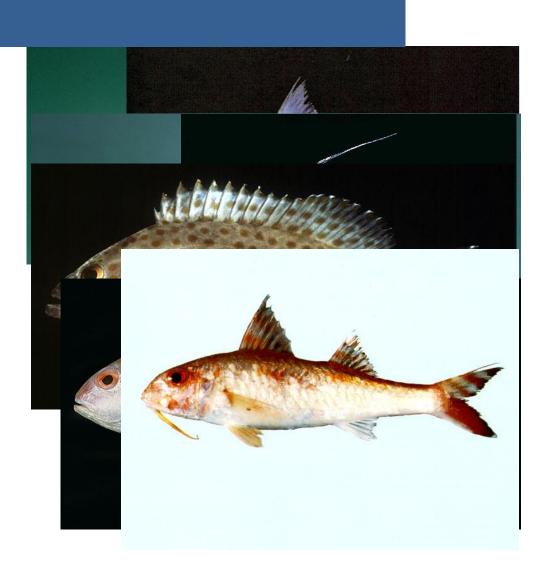
Alcohol

- is usually not used for killing and fixing vertebrates. But of course used for long term storage.
- Colour specimen is lost immediately.
 - A teaspoonful of glycerine in quart of alcohol helps to preserve natural colour and to keep integument flexible
- Alcohol usually comes in the 95% concentrates form.
- For long-term preservation, 70-75% strength in used
 - i. Formalin 1 week (fix soft tissue)
 - ii. Water 1 day (leach out the formalin)
 - iii. Alcohol long term storage

http://preserve.sivasothi.com

Species Identification

- Family Priacanthidae
- Family Engraulidae
- Family Synodontidae
- Family Platycephalidae
- Family Carangidae
- Family Leiognatidae
- Family Lutjanidae
- Family Serranidae
- Family Nemipteridae
- Family Mullidae



Species Identification

- Family Squillidae
- Family Penaeidae
- Family Portunidae
- Family Loliginidae
- Family Octopodidae





