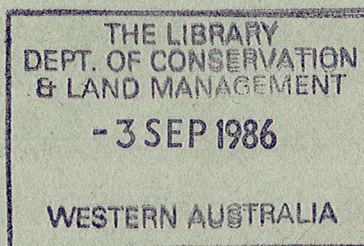


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**ANALYSIS OF CORAL COMMUNITY
DATA USING MULTIVARIATE
TECHNIQUES, AND THEIR
APPLICATION TO OTHER
COMMUNITY DATA**



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ANALYSIS OF CORAL COMMUNITY DATA USING
MULTIVARIATE TECHNIQUES, AND THEIR APPLICATION
TO OTHER COMMUNITY DATA

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DAMPIER ARCHIPELAGO MARINE STUDY

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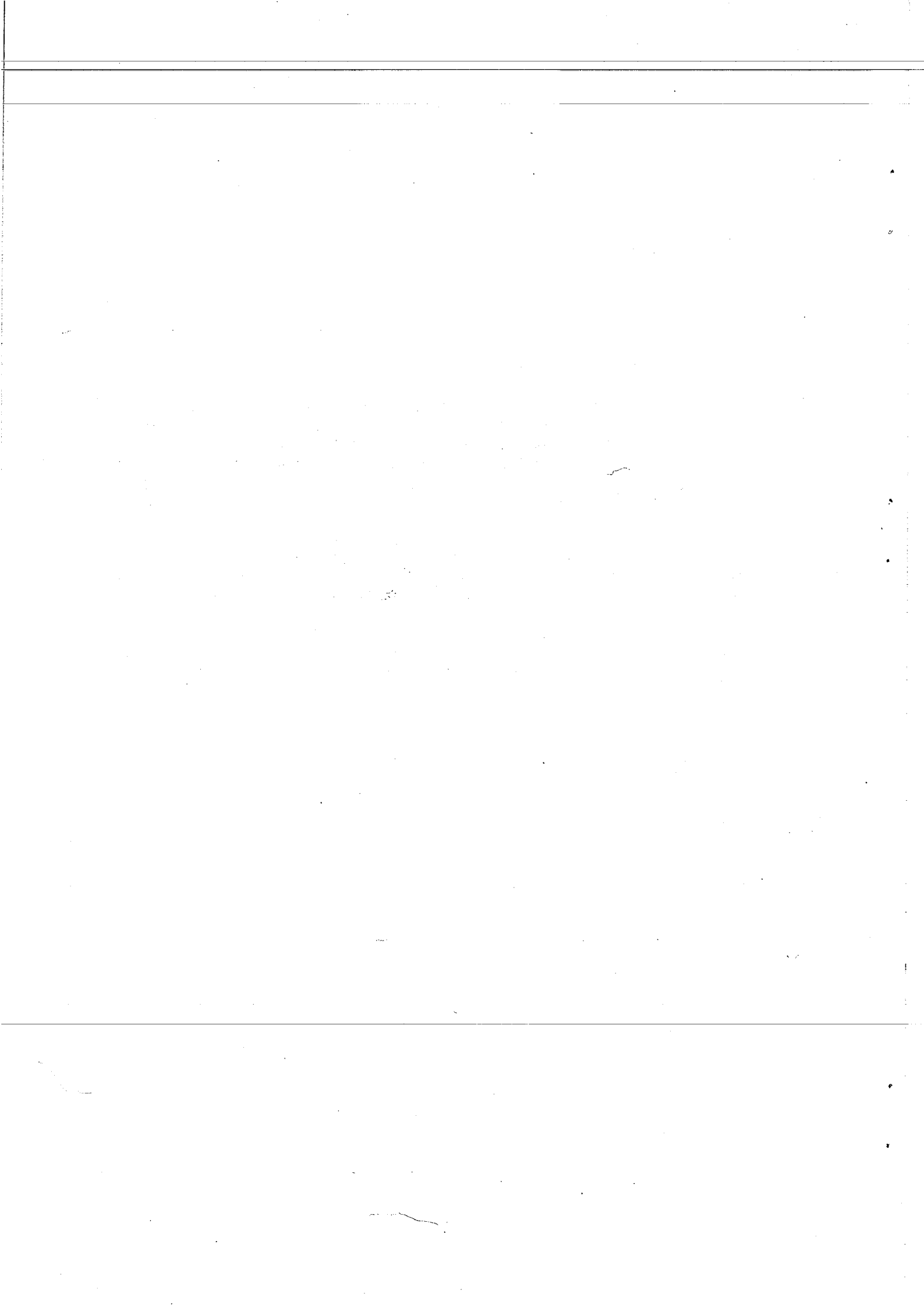
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ABSTRACT

Coral data from the Dampier Archipelago, Western Australia are used as an example to illustrate the procedures necessary to convert raw field data, in the form of species lists, into a form that can be analysed by the Cornell Ecology Program Series. Classification of the data was carried out using composite clustering and two way indicator species analysis. Ordination was performed by detrended correspondence analysis. A brief description of each program is given. Data coding and input, program execution and output interpretation are also discussed.

Sites were classified into three main groups that correlated with the distribution of offshore, exposed and protected habitats within the Archipelago. Coral species diversity tended to be lower at protected and offshore sites than at exposed sites. There was a notable trend of species classification into their preferred habitats. Information derived from these analyses agreed well with data from the literature.

The successful application of these multivariate techniques in analysing coral data indicates that they may have a more widespread use; suggestions are made for their use in other studies.



GENERAL INTRODUCTION

Multivariate techniques (ordination and classification) are used extensively in terrestrial ecology (Kershaw, 1973; Sneath and Sokal, 1973; Whittaker, 1978a, 1978b), and have also been applied to data from marine environments (Veron and Done, 1979; Wiegleb, 1980; Moran, 1981; Done, 1983).

The availability of computers, and the large range of programs for use in community studies has resulted in the use of these techniques on data not originally collected for ordination and classification analyses. While there is generally a wide choice of programs for data analyses (Gauch and Whittaker, 1981), the final decision usually rests on what information is being sought and which programs are available at respective research institutes. As a result a number of methods have been used and quoted in the literature (eg. Sneath and Sokal, 1973; Green and Vascotto, 1978; Veron and Done, 1979).

This report is divided into two sections. Section A briefly describes four Cornell Ecology Programs; this includes the methods of data coding and input as well as program execution and interpretation of output. Section B applies these techniques to data, not originally collected for this type of analysis, on corals from the Dampier Archipelago, Western Australia (Marsh, 1978, Fig. 1). The value of these techniques in analysing other descriptive community data is also discussed.

A MULTIVARIATE TECHNIQUES

1. INTRODUCTION

The mechanics of computer program execution are often a deterrent to researchers. To assist in understanding the Cornell Ecology Programs applied in this study, a brief description is given below. In addition, data coding and input, program execution and output interpretation are discussed, using the coral data described in Section B.

These are brief descriptions only and the Cornell Programs' capabilities surpass the overview discussed here. The reader is directed to the appropriate program manuals (Gauch, 1982) for further information on their full capabilities and use.

2. PROGRAM DESCRIPTION

All programs described here (CONDENSE, COMPLUS, TWINSpan, DECORANA) were from the Cornell Ecology Program Series (Gauch, 1982) maintained at Murdoch University, Western Australia. These programs at the time of writing were also available for use at the University of Western Australia.

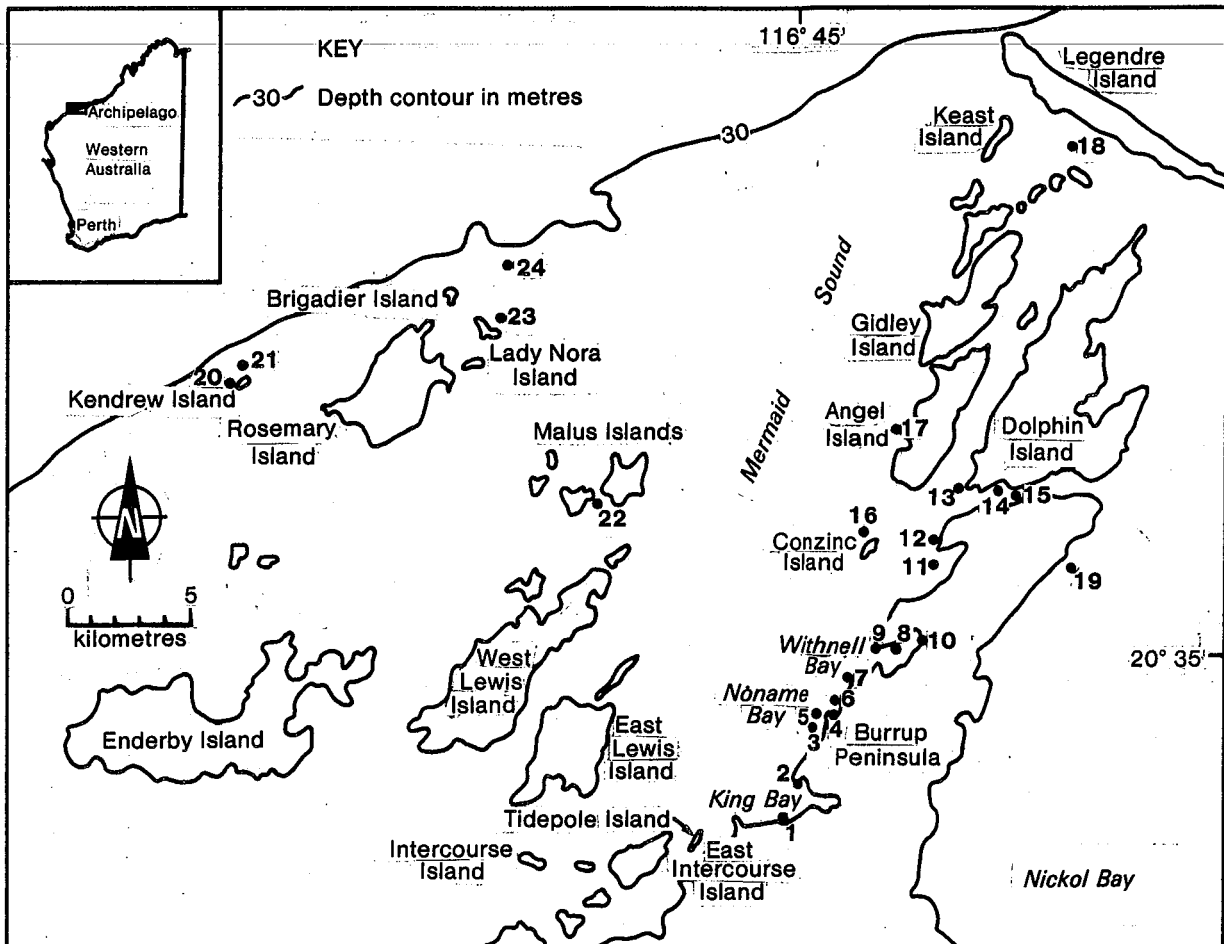


Figure 1; Sample sites in the Dampier Archipelago where coral species were listed by Marsh (1978).

To use the programs, COMPCLUS, TWINSpan and DECORANA, input data had to be in a 'restricted condensed format' (Gauch, 1982) in samples (or sites) order. To achieve this restricted format, data was read into the computer in a matrix of samples by species and copied into a form acceptable to the above programs using a program called CONDENSE (Singer and Gauch, 1979). Once data had been condensed it could be used in the COMPCLUS, TWINSpan and DECORANA programs.

COMPCLUS (Composite Clustering)

Written by Gauch (1979), this is a FORTRAN program designed to enable large data sets (species and sites) to be clustered initially. This is useful where there are large numbers of species and samples. It is used to identify outlying samples and major groupings of samples (clusters). Each cluster produces a composite sample, which is an average of the samples contained, and these can be used for subsequent analysis by ordination or classification.

Clustering involves two operations; a sample is selected at random and samples clustered within a user-specified radius (several distance measures are available (Gauch, 1982)). This is repeated until all samples are allocated to a cluster, ignoring any sample that has been assigned to a cluster already. The second operation reassigns samples from small clusters (having fewer than a specified number of members) into the nearest large cluster, provided that the sample is within a user-specified radius (which is larger than the radius specified for the first operation).

The value of COMPCLUS is largely dependent on the initial size of the data set. A large array of figures can be reduced by grouping similar samples into classes. Relationships can then be studied among a smaller, comprehensible number of groups.

COMPCLUS was used in this study, but the relatively small number of sites (24) and species (99) did not require further reduction before analysis.

TWINSpan (Two Way Indicator Species Analysis)

This is a package designed for use on data collected on the occurrence of a set of species in a set of samples (Hill, 1979a). This program uses a polythetic, divisive technique (ie. whereby the initial data set is divided sequentially into smaller groups ('divisive'), using a number of different characters ('polythetic')) to first construct a classification of the samples. The program then uses this to obtain a classification of the species according to their ecological preferences. The two classifications are then used together to obtain an ordered two-way table that expresses the species' relations as closely as possible.

TWINSpan and other classification programs are compared by Gauch and Whittaker (1981) and the general theory of indicator species analysis (of which TWINSpan is a further development) is discussed by Hill *et al.* (1975).

TWINSpan was used in this study to classify coral communities based on site and species data.

DECORANA (Detrended Correspondence Analysis)

This is a FORTRAN program for detrended correspondence analysis (DCA) and reciprocal averaging (Hill, 1979b). Its main purpose is to make ordinations using DCA (Gauch, 1982).

Ecological ordination refers to the arrangement of samples (or species) either in relation to environmental gradients, or in relation to axes that may correspond to environmental gradients (Kessel, 1979). A major purpose of such an arrangement is to enable recognition of joint variation in community composition and environmental factors (Gauch *et al.*, 1977).

If ordination produces axes that are ecologically meaningless or species and site patterns that cannot be interpreted, the process is not useful.

DECORANA, by default, produces four axes of variation, each with a calculated eigenvalue (see Sneath and Sokal, 1973 for further definition). An eigenvalue is equal to the variance along its corresponding axis. Thus the principal axis corresponding to the largest eigenvalue is the dimension that accounts for the greatest amount of variance from the sample (Sneath and Sokal, 1973). Likewise the next highest eigenvalue is that of the second most important axis and accounts for the second largest variance, and so on.

DECORANA also produces scatter diagrams of samples and species. These usually need to be replotted, using the values given for each axis, and then examined.

The present study applied DECORANA to determine whether the distribution of coral communities could be related to some combination of environmental factors.

3. DATA CODING AND INPUT

The allocation of a unique number to each coral species recorded in the Dampier Archipelago generated, for each coral site, a species presence/absence list. This enabled data to be coded into a site by species matrix (a proforma of which is shown in Fig. 2) for input into the Cornell Ecology Programs. Relative abundance figures given by Marsh (1978) were modified to allow them to be used in the programs as follows; 0: absent, 1: rare, 2: common, 3: abundant.

The data file was then placed on computer disk with the appropriate FORTRAN statements necessary to read the data into the CONDENSE program (Fig. 3). The generated output (Fig. 4) was then in the restricted format required for COMPCLUS, TWINSpan and DECORANA.

CONVERT 10(I3), 10(I3, F2.0))									
99 24DAMPIER CORAL COMMUNITY DATA 1978 - 99 SPECIES 24 SAMPLES T*DAMP**									
(3(2X, 30F2.0), 2X, 9F2.0)									
30									
1 000000000000001000000000000000									
1 001010000100000001000000000000									
1 000000200000000001000000000000									
1 000011010									
2 000000000100101000000010000101									
2 011010010100100101100111001111									
2 000000011001000110000000000010									
2 000022101									
3 000000000000100000000010001000									
3 000020010000000001210010012001									
3 000101010001000210000000000000									
3 000001001									
4 000300200102102000000010002010									
4 101010020001000002000200000011									
4 101010020001000002000200000011									
4 000011110									
- Additional data here -									
20012010011010000000000001000000									
20001020000000010021001100002011									
20001000000000000001110000000000									
20000000000									
21021110002011000000000020000000									
210001100000000000021200100002210									
21001001000101000011100000000000									
21010201000									
22000000001022001000010030002200									
22000030000100000002200221102010									
22000001000000010120010000020000									
22000001000									
23000000000001010000011000000000									
23100000000100000001000100000000									
230000000000000000000000000000									
23000011000									
24000000001011000000001000000000									
24000101000000000000010001100000									
240000001000000000000000000000									
24000000000									
MILI	MILP	PSAD	PSAC	PSAS	STYG	POCD	POCE	POCV	STPP
ACRH	ACRH	ACR2	ACR3	ACR4	ACR5	ACR6	ACR7	ASTM	MON1
MONR	MONE	PACD	PAVM	PACR	CUSS	FUNF	HERL	POLT	PODC
ALVS	GON1	GON2	GONT	POR1	POR2	POR3	CAUT	CYPM	CYP1
CYPS	FAVA	FAVF	FAVT	FAVX	FAVP	FAVS	FAV1	FATA	FATC
FATP	FAT1	GOSA	GOSU	GOSE	GOSP	GOSR	ECHL	HYDE	HYDM
HYDR	LEPP	LETP	MOTC	MOTC	PLAD	PLAS	PLEV	MOSL	DIPH
TRAG	GALF	GALC	MERA	SCA1	ACAE	LOBC	LOBH	SYMR	ECNA
OXYL	MYCE	MYCT	MYC1	PECP	PEC1	CATJ	EUPG	EUPF	PLR1
DEN1	DEN2	DUNA	TUBA	TURB	TURF	TURM	TURP	TURR	
	1	2	3	4	5	6	7	8	9
	11	12	13	14	15	16	17	18	19
	21	22	23	24					20

Figure 3; The input file to CONDENSE which consists of FORTRAN statements, the data, species codes and site numbers.

COMPCLUS

L CORNCLUS	[Loads the COMPCLUS program]
AS 3.CON:	[Input and output directed to the console]
AS 5.DAMPCOND.DAT	[Assigns the input file to be 'DAMPCOND.DAT' - the product of the CONDENSE program]
AS 6.PR:	[Assigns output to the lineprinter]
AS 7.NULL:	[Indicates no assignation]
ST	[Starts the program]

TWINSpan

L CORNTWIN	[Loads the TWINSpan program]
AS 1.CON:	[Input from the console]
AS 3.PR:	[Assigns output to the lineprinter]
AS 4.CON:	[Output to the console]
AS 5.DAMPCOND.DAT	[Assigns the input file to be 'DAMPCOND.DAT']
AS 7.NULL:	[Indicates no assignation]
ST	[Starts the program]

DECORANA

L CORNDECO	[Loads the DECORANA program]
AS 1.DAMPDECO.COMD	[Assigns a command file which specifies program parameters]
AS 3.PR:	[Assigns output to the lineprinter]
XAL DAMPDECO.OUT.IN132	[Creates an ouput file called 'DAMPDECO.OUT']
AS 4.DAMPDECO.OUT	[Assigns output to the file 'DAMPDECO.OUT']
AS 5.DAMPCOND.DAT	[Assigns the input file 'DAMPCOND.DAT']
AS 7.NULL:	[Indicates no assignation]
ST	[Starts the program]

Within each program there exists the capability of changing certain parameters; for example, sites and species may be omitted or data transformed. These manipulations are detailed in the appropriate program manuals and the reader is directed to these for further information.

5. OUTPUT INTERPRETATION

The output derived from the Cornell Programs, is described by the appropriate manuals. The main features of output for the present study are discussed briefly below.

COMPCLUS

The output of COMPCLUS consisted of a description of the clusters and which sites were included in each. Also detailed were the most abundant species present in each cluster.

99 24DAMPIER CORAL COMMUNITY DATA 1978 - 99 SPECIES 24 SAMPLES TCDAMP**S									
(13, 10(13, F2.0))									
10									
1 151. 331. 351. 401. 481. 672. 771. 951. 961. 981.									
2 101. 131. 151. 231. 281. 301. 321. 331. 351. 381.									
2 401. 431. 461. 481. 491. 521. 531. 541. 571. 581.									
2 591. 601. 681. 691. 721. 761. 771. 891. 952. 962.									
2 971. 991.									
3 131. 231. 271. 352. 381. 481. 492. 501. 531. 561.									
4 43. 72. 101. 122. 131. 152. 231. 272. 291. 311.									
4 331. 351. 382. 421. 482. 522. 591. 601. 661. 691.									
4 712. 721. 741. 772. 791. 891. 951. 961. 971. 981.									
- Additional data here -									
20 21. 32. 51. 81. 91. 111. 241. 331. 352. 441.									
20 472. 481. 511. 521. 572. 591. 601. 631. 771. 781.									
20 791.									
21 22. 31. 41. 51. 92. 111. 121. 232. 341. 351.									
21 472. 481. 492. 521. 572. 582. 591. 631. 661. 701.									
21 721. 771. 781. 791. 921. 942. 961.									
22 91. 112. 122. 151. 201. 233. 272. 282. 353. 401.									
22 482. 492. 522. 532. 541. 551. 572. 591. 661. 741.									
22 761. 772. 801. 862. 961.									
23 111. 131. 191. 201. 311. 401. 481. 521. 951. 961.									
24 91. 111. 121. 211. 341. 361. 491. 531. 541. 671.									
0									
MILI	MILP	PSAD	PSAC	PSAS	STYG	POCD	POCE	POCV	STPP
ACRH	ACRH	ACR2	ACR3	ACR4	ACR5	ACR6	ACR7	ASTM	MON1
MONR	MONE	PACD	PAVM	PACR	CUSS	FUNF	HERL	POLT	PODC
ALVS	GON1	GON2	GONT	POR1	POR2	POR3	CAUT	CYPM	CYP1
CYPS	FAVA	FAVF	FAVT	FAVX	FAVP	FAVS	FAV1	FATA	FATC
FATP	FAT1	GOSA	GOSU	GOSE	GOSP	GOSR	ECHL	HYDE	HYDM
HYDR	LEPP	LETP	MOTC	MOTC	PLAD	PLAS	PLEV	MOSL	DIPH
TRAG	GALF	GALC	MERA	SCA1	ACAE	LOBC	LOBH	SYMR	ECNA
OXYL	MYCE	MYCT	MYC1	PECP	PEC1	CATJ	EUPG	EUPF	PLR1
DEN1	DEN2	DUNA	TUBA	TURB	TURF	TURM	TURP	TURR	
	1	2	3	4	5	6	7	8	9
	11	12	13	14	15	16	17	18	19
	21	22	23	24					10
									20

Figure 4; The output file generated by CONDENSE from the file shown in Fig. 3. This file is now ready for use as input to other Cornell Ecology Programs.

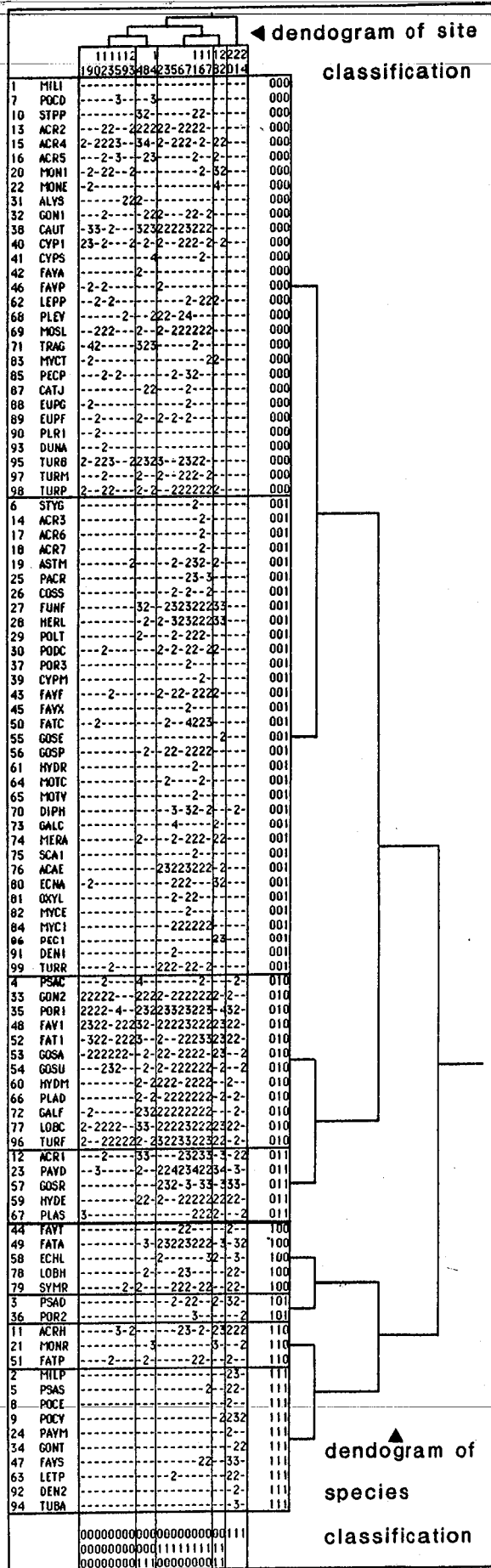


Figure 5; An example of a site by species table produced by TWINSpan.

TWINSPAN

Output gained from TWINSPAN allowed dendograms to be drawn of sites and species (as in Fig. 8). Indicator species for each division were also produced. The tabular arrangement of species and sites (Fig. 5) indicated the arrangement of species with regard to their site preferences. For example, those species shown at the bottom of Fig. 5 tended to occur in the sites numbered 20, 21 and 24.

DECORANA

Ordinations produced by DECORANA appeared as four axes for both sites and species, in which sites were ranked from highest to lowest. Values for each axis (Fig. 6) have been multiplied by 100. Hence a value of 310 (Fig. 6, *) can be interpreted as 3.1 sd (standard deviations), and so on. These axes were also plotted in scatter diagrams (Fig. 7), although for clarification it was necessary to replot the axes using the printed values, as the scatter diagrams did not show species and site numbers. The main function of the scatter diagram output by DECORANA (Fig. 7) was to illustrate the presence of obvious gradients. The replotting of data from Fig. 6 can be seen in Fig. 11.

99 24DAMPIER CORAL COMMUNITY DATA 1978 - 99 SPECIES 24 SAMPLES TCDAMP
DECORANA OPTIONS - DOWNWEIGHTING 0 RESCALING 4 ANALYSIS 0 SEGMENTS 26 THRESHOLD 0.00
TRANSFORMATION 0.00 0.00

SAMPLE SCORES - WHICH ARE WEIGHTED MEAN SPECIES SCORES

N	NAME	AX1	AX2	AX3	AX4	RANKED 1 EIG=0.362		RANKED 2 EIG=0.194		RANKED 3 EIG=0.135		RANKED 4 EIG=0.088					
1	1	112	102	108	3	21	21*	310	18	18	225	20	20	206	10	10	168
2	2	121	59	71	114	20	20	305	5	5	139	19	19	168	9	9	124
3	3	177	34	60	92	24	24	218	24	24	129	23	23	138	19	19	121
4	4	119	0	54	59	22	22	194	22	22	120	15	15	116	2	2	114
5	5	177	139	48	98	17	17	183	15	15	119	1	1	108	7	7	112
6	6	152	78	98	86	3	3	177	1	1	102	6	6	98	5	5	98
7	7	158	81	83	112	5	5	177	20	20	99	9	9	95	11	11	93
8	8	120	41	91	42	7	7	158	11	11	96	22	22	95	3	3	92
9	9	58	95	95	124	16	16	154	9	9	95	8	8	91	6	6	86
10	10	98	49	52	168	6	6	152	17	17	88	13	13	90	22	22	85
11	11	137	96	12	93	11	11	137	21	21	85	7	7	83	17	17	83
12	12	82	64	68	68	19	19	135	13	13	84	18	18	79	20	20	71
13	13	86	84	90	39	18	18	124	7	7	81	2	2	71	12	12	68
14	14	0	67	55	51	2	2	121	6	6	78	12	12	68	21	21	66
15	15	99	119	116	45	8	8	120	23	23	76	3	3	60	18	18	62
16	16	154	56	59	0	4	4	119	14	14	67	16	16	59	4	4	59
17	17	183	88	45	83	1	1	112	12	12	64	14	14	55	14	14	51
18	18	124	225	79	62	15	15	99	2	2	59	4	4	54	23	23	50
19	19	135	11	168	121	10	10	98	16	16	56	10	10	52	15	15	45
20	20	305	99	206	71	23	23	93	10	10	49	5	5	48	8	8	42
21	21	310	85	25	66	13	13	86	8	8	41	17	17	45	13	13	39
22	22	194	120	95	85	12	12	82	3	3	34	21	21	25	24	24	25
23	23	93	76	138	50	9	9	58	19	19	11	11	11	12	1	1	3
24	24	218	129	0	25	14	14	0	4	4	0	24	24	0	16	16	0

Figure 6; Axes produced by DECORANA for all species and all sites. *Values are multiplied by 100 and must be reduced when replotting for scatter diagrams.

the printed values, as the scatter diagrams did not show species and site numbers. The main function of the scatter diagram output by DECORANA (Fig. 7) was to illustrate the presence of obvious gradients. The replotting of data from Fig. 6 can be seen in Fig. 11.

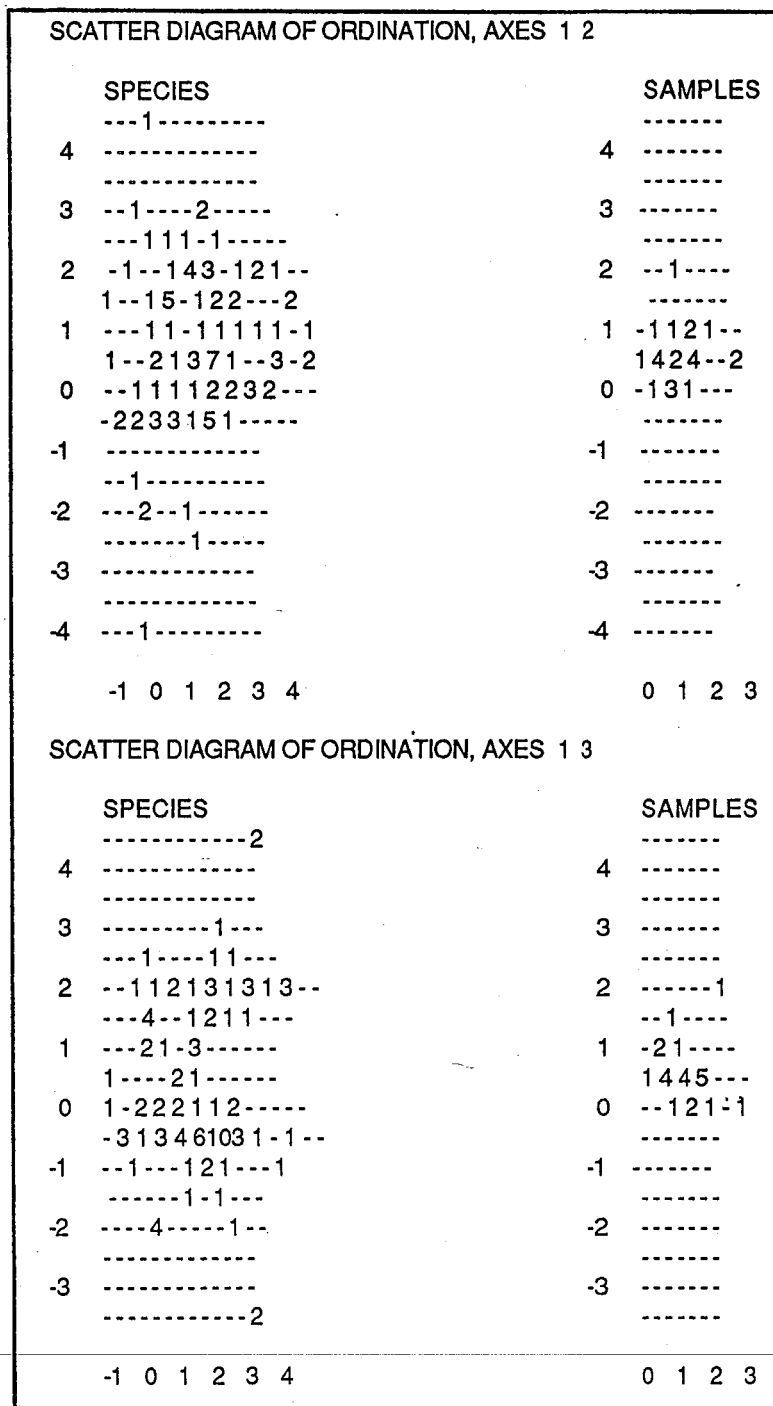


Figure 7; Examples of scatter diagrams produced by DECORANA for all species and all sites. Numbers within each diagram refer to the frequency of occurrence of species or sites. For example in the scatter diagram of samples from axes 1 and 2, the number '4' signifies that four sites are included in that 'area' (ie. within the resolution of the diagram).

The eigenvalue is given at the head of each axis (Fig. 6). According to Hill (1979b) axes which have eigenvalues much lower than the highest value are probably not significant. For example, in Fig. 6, where values are 0.362, 0.194, 0.135 and 0.088 for axes 1 to 4 respectively, only the first three axes are likely to be significant.

B. ANALYSIS OF DAMPIER ARCHIPELAGO CORAL COMMUNITIES

1. INTRODUCTION

In the past the structure and zonation of coral communities have been explained on the basis of prevailing environmental conditions (Braithwaite, 1971; Dana, 1976; Glynn, 1976; Done, 1977; Veron and Done, 1979). Recent information, however, has suggested that aperiodic, episodic events may also determine coral zonation and community structure (Connel, 1978; Done, 1983). Stated simply, the question is to what degree the species present in a community are there as a result of the surrounding physical and biotic environment.

The interpretation of community composition is complicated by the presence of widespread species that may overlap communities so much that their presence is not an indication of the long term physical conditions. This observation has led to the suggestion that the growth forms of ubiquitous coral species may be more distinctly zoned in their distribution. Thus alternatively, the form of a species may be a better indicator of the environment than the presence or absence of species (Done, 1983).

Light, and water movement (waves, currents and tides), are environmental factors recognised as major determinants of coral distribution (Done, 1983). Water movement also determines sediment dynamics (sediment depth, grain size, suspended load and deposition), which is another factor determining coral distribution patterns (Loya, 1976).

The aim of this study was to use the multivariate techniques described in Section A to determine broad patterns in coral community composition and to find their physical determinants in different areas of the Dampier Archipelago. The approach taken was to group sites on the basis of species composition and to determine if physical factors could be used to explain the grouping of these communities.

2. STUDY AREA

The Dampier Archipelago is a marine area characterised by groups of islands, rocky reefs, coral reefs and shoals that rise from a general deep water plain (Semenuik *et al.*, 1982). Numerous

geomorphic units are recognised within the Archipelago based mainly on substrate type, substrate heterogeneity and tidal level. These units have been developed by a combination of factors which include: erosion versus sedimentation, underlying stratigraphic sequence, tidal level, wave action and ancestral (pre-Holocene) physiography (Semenuik *et al.*, 1982).

Virtually every shoreline in this region has exposed and protected settings. In addition there is a wide range of habitats throughout the Archipelago that vary from those exposed to wave action on outer islands with clear waters (eg. Sites 20, 21, 22 and 24, Table 1, Fig. 1), to sheltered areas within bays that can become extremely turbid (eg. Sites 8, 9 and 10, Table 1, Fig. 1). Intermediate conditions exist for shores of a different aspect, topography, slope and varying current regimes (Marsh, 1978).

3. METHODS

Coral data

Data used for clustering (classification) and ordination were obtained from coral species lists reported in Marsh (1978). These data were collected from a number of sites in the Dampier Archipelago in 1978 (Fig. 1). At each site transects were made from the intertidal zone to a depth of 7 to 9 m where coral reefs gave way to a sandy bottom (Semenuik *et al.*, 1982). Sites were surveyed by divers using a boat-mounted hookah apparatus, SCUBA and snorkelling. The depth, substrate type, an indication of species abundance at each site (in terms of absent, rare common and abundant), and the percent cover of living coral were also noted (Marsh, 1978).

To facilitate analysis, sample sites were renumbered (Table 1), and a complete species list of all corals was recorded. All species were given an individual number to generate presence and absence data for each site, this information was then stored on computer file and subsequently transformed to a 'restricted condensed format' using the CONDENSE program (Section A).

Site and species classification

The program TWINSpan (Hill, 1979a) was used to classify sites and species using four divisive levels. Initially all species (99) and sites (24) were used in the classification. A second classification, however, was used after removing rare species (leaving a total of 79) and one site (Site 9; Withnell Bay barge). This was performed because the distribution of rare species (those occurring at one site only) was not considered a good indicator of environmental conditions. Furthermore the barge in Withnell Bay was an artificial and unique substrate. General patterns were being sought, which could be obscured by rare species and outlying samples (Gauch, 1977).

Groups of sites and species derived from TWINSpan were then examined on the basis of predominant substrate type and other physical characteristics at each site. Species preferences for water conditions (for example; exposed, protected, turbid or clear) were determined for the majority of species present using taxonomic information described by Braithwaite (1971), Marsh (1978), Mather and Bennet (1978), Veron and Pichon (1976, 1979, 1982) and Veron *et al.* (1977).

Table 1; Code numbers allocated for this study to sample sites, coral genera and species. Data and original site codes from Marsh (1978), (see Fig. 1).

Sample site	Site code (Marsh, 1978)	Site code (This study)	Number of genera	Number of species	Range of % living cover
King Bay/Dampier	ML16	1	8	10	<1%
North King Bay	ML15	2	21	32	5-50%
South Noname Bay	ML14	3	16	20	5%
Noname Bay	ML2	4	23	30	1-50%
Noname Rocks	ML3	5	32	38	<1-30%
Near Noname Bay	ML1	6	22	33	0-50%
Withnell/Noname Bay	ML5	7	35	55	<1-60%
Withnell Bay	ML7	8	17	25	10-20%
Withnell Bay barge	ML27	9	13	15	-
Withnell east	ML28	10	15	16	1-5%
Conzinc Bay	ML9a	11	34	51	<1-80%
North Conzinc Bay	ML10b	12	16	25	<1-20%
Southwest Dolphin Is.	ML11a	13	11	16	5-10%
West end of Boat Passage	ML11b	14	16	20	5-20%
Central Boat Passage	ML11c	15	9	12	-
North Conzinc Is.	ML12	16	29	48	5-10%
West Angel Is.	ML13	17	27	40	5-50%
North Gidley/Legendre Is.	-	18	23	29	50-100%
Nickol Bay	-	19	7	7	<1%
Kendrew Is.	WAM1	20	14	21	1-20%
Kendrew Is.	WAM2	21	21	27	5-50%
Malus Is. (Whalers Bay)	ML30	22	19	25	60-80%
North Lady Nora Is.	ML17a	23	8	10	<1%
Miller Rocks	ML17b	24	9	10	1-5%

Site ordination

Ordination of sites was performed using the technique of detrended correspondence analysis incorporated into the computer package DECORANA (Hill, 1979b). The axes of variation derived from this process were then examined using known site characteristics. A second ordination process was carried out after removing rare species and Site 9, for the reasons given above.

4. RESULTS

Site classification

TWINSpan, for both classifications, separated the offshore from the nearshore sites (20, 21 and 24) at the first division (Fig. 8). An additional outer site (Lady Nora Island - Site 23), in both classifications, was grouped with sites occurring on or near the Burrup Peninsula (Fig. 9). This occurred because there were species present both at Site 23 and on the Burrup Peninsula that did not occur at the other offshore sites. These species were *Alveopora* sp. (Poritidae); *Acropora* sp., *Astreopora myriophthalma* (Acroporidae); *Favites* sp. (Faviidae) and *Turbinaria bifrons* (Dendrophylliidae).

The main subgroups occurring on the Burrup Peninsula were the relatively sheltered sites (Group 1b: Sites 1, 8, 10, 12-15) which tended to have a lower number of species present (Fig. 8(b)); and the more exposed sites (Group 2a: Sites 2-7, 11, 16, 17) which had more species.

In general there was less living coral (<1% - 20% cover, Marsh, 1978) and a low species diversity in protected sites on the Burrup Peninsula (Group 1b, Fig. 8(b), Table 2) compared with more

Table 2; The number of genera and species present in coral families occurring in protected and exposed sites on the Burrup Peninsula and at offshore sites, (see Fig. 8).

Family	Protected sites on the Burrup Peninsula (1b) (Fig. 8b)															Exposed sites on the Burrup Peninsula (2a) (Fig. 8b)										Offshore sites (3) (Fig. 8b)																					
	1	8	10	12	13	14	15	2	3	4	5	6	7	11	16	17	20	21	22	24																											
	G	S	G	S	G	S	G	S	G	S	G	S	G	S	G	S	G	S	G	S	G	S	G	S	G	S																					
Milleporidae				1	1																					1	1	1	1																		
Thamnaestriidae					1	1																					1	2	1	3																	
Astrocoeniidae											1	1	1	1			1	1	1	1	1	1	1	1																							
Pocilloporidae			1	1						2	2						2	2	2	2						1	2	1	1	1	1	2															
Acroporidae	1	1	1	4	1	1	2	5	2	3	2	3	1	3	1	2	1	1	1	3	2	2	1	4	2	5	2	4	3	9	1	1	1	1	1	2	2	4	2	2							
Agericiidae					1	1											1	1	1	1	1	1	1	1	2	2	2	2	1	1	2	2															
Siderastreidae																	1	1																													
Fungiidae			2	2			1	1									2	2	1	1	2	2	4	4	2	2	4	4	4	4	4	3	3	3	3			3				2	2				
Poritidae	2	2	2	3	2	2	7	10	1	1	2	3	1	1	2	3	1	1	3	3	2	2	2	2	2	2	4	2	4	2	2	2	3	2	2	2	2	1	1	2	2						
Faviidae	3	3	8	1	7	8		6	7	7	7	3	4	10	16	9	12	8	10	14	10	14	10	14	22	20	12	21	11	12	21	22	0	5	9	9	11	6	10	4	4						
Merulinidae																																															
Mussaidae	1	1	1	2	1	1	1	1	1								2	2	2	2	2	3	3	3	4	3	4	2	2	3	3	3	3	2	3	2	3	2	2								
Pectiniidae						1	1																			4	4	2	2	4	5	3	3	1	1	1	2								2	2	
Caryophylliidae			1	1	2	2											1	1			1	1	1	1			2	3																			
Dendrophylliidae	1	3	1	1	1	1	2	5	1	4	1	3	1	1	4	1	2	1	4	2	4	1	4	1	4	1	5	1	5	1	3	1	4														
Total: Genera (G)	8	17	15	16	11	16	9										21	16	23	32	22	35	34	29	27																						
Species (S)	10	25	16	25	16	20	12										32	20	30	38	33	55	51	48	40																						

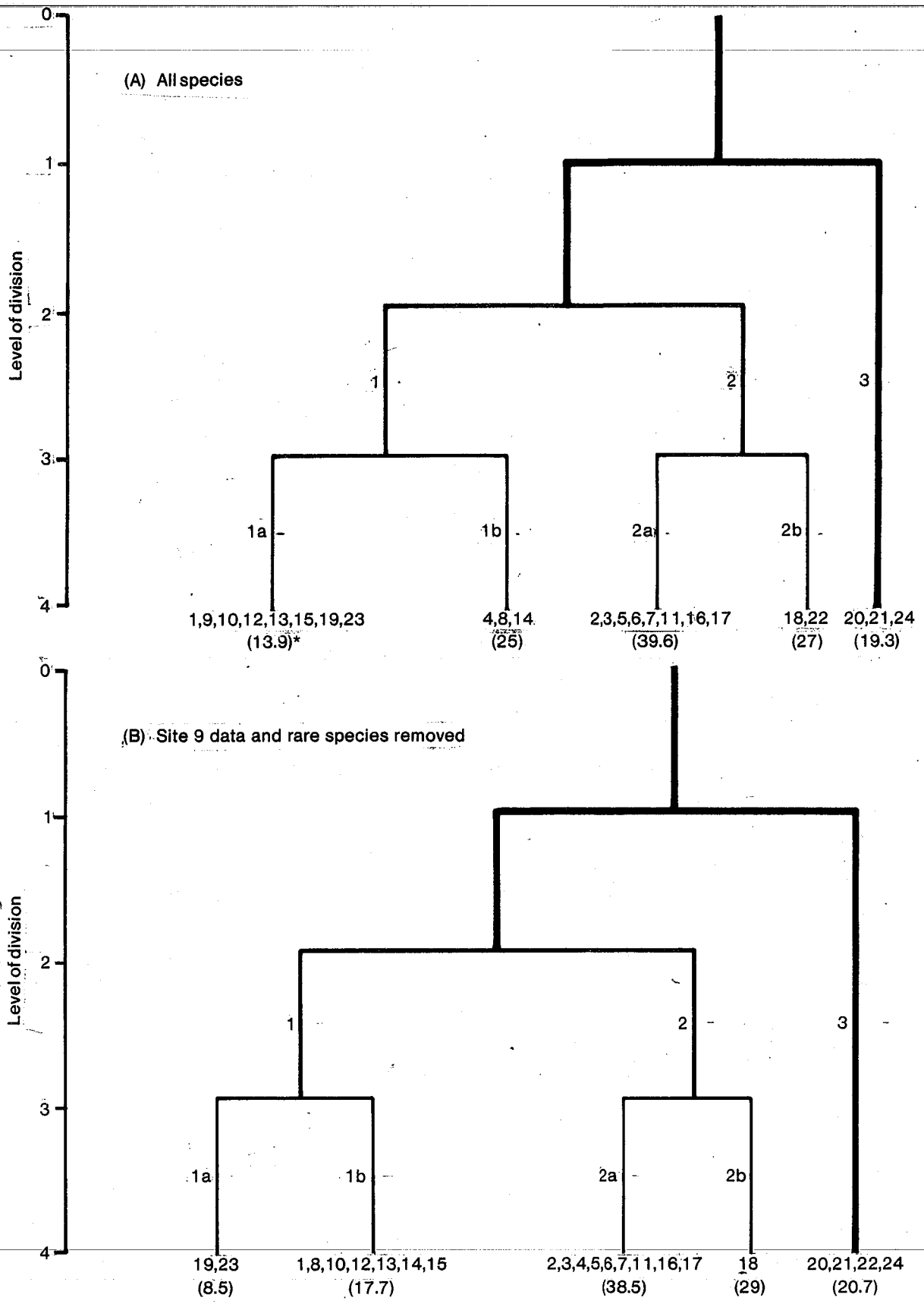


Figure 8; Two dendrograms of sample sites produced by TWINSpan. *Refers to the average number of species per site.

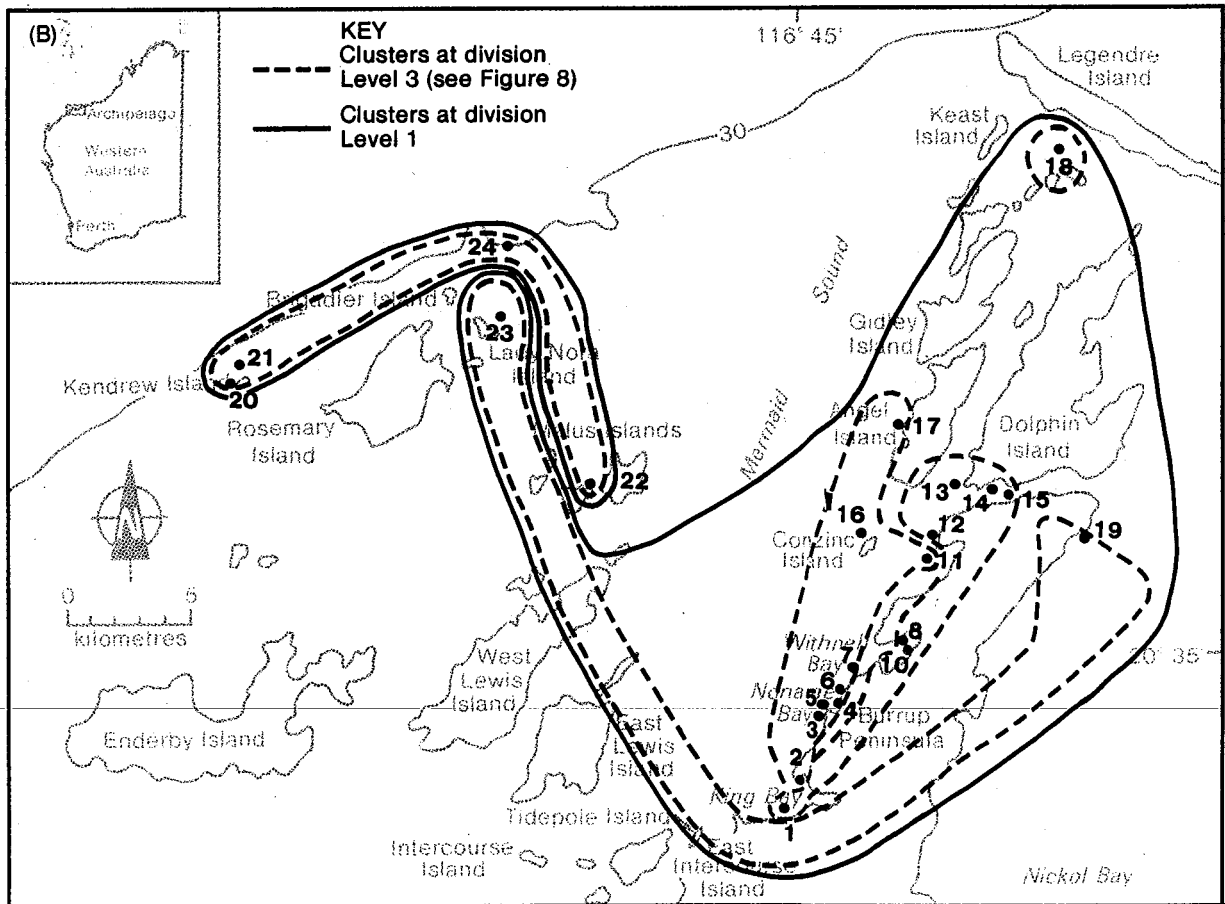
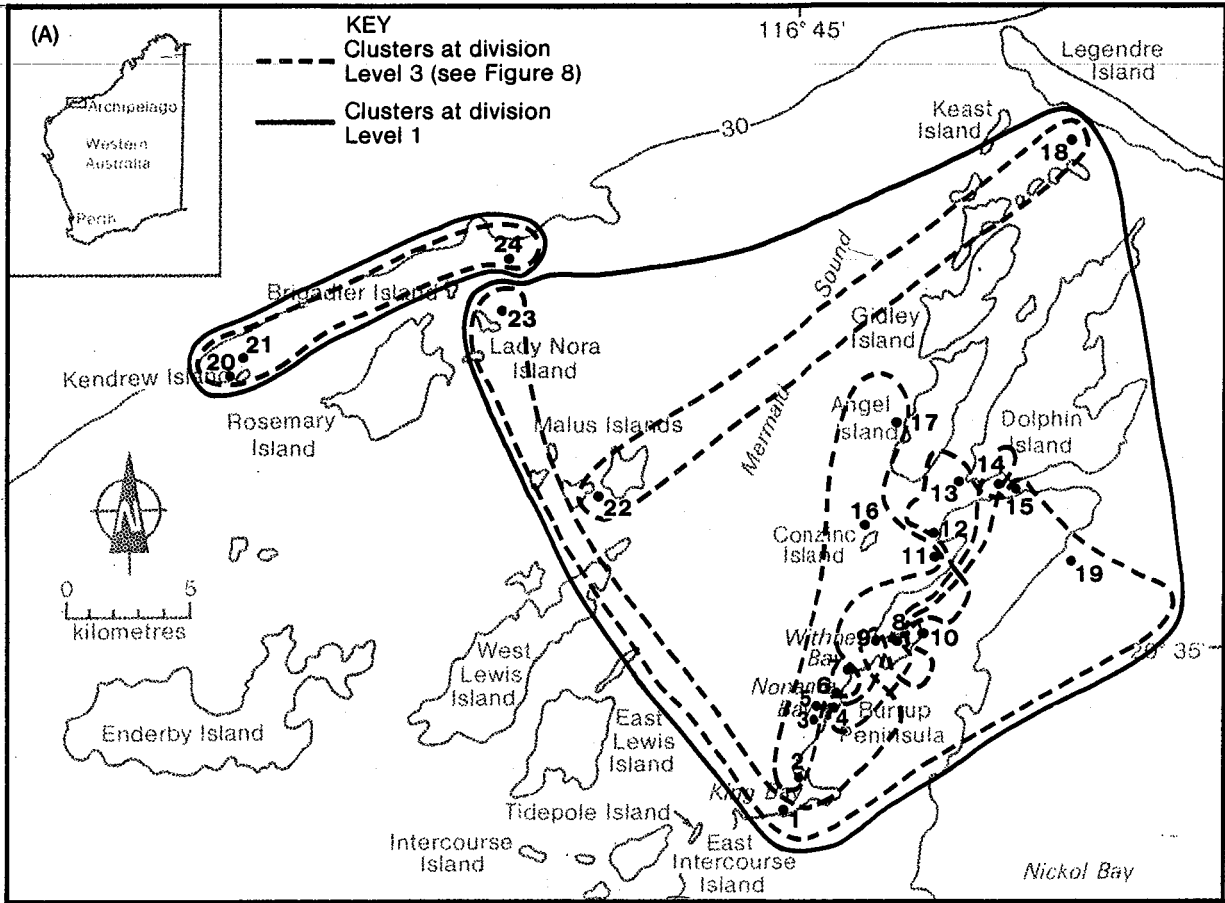


Figure 9; Site groupings produced by TWINSPLAN. Map A for all species; Map B has Site 9 and rare species removed.

exposed sites (Group 2a, Fig 8(b) that had more living coral (up to 80% cover) and a greater diversity (Table 2). At offshore sites (20, 21, 22 and 24) on average there was less living coral, though cover up to 80% was noted (Marsh, 1978), and species numbers were also low when compared with the exposed sites on the Burrup Peninsula.

Species classification

Species assemblages in general reflected the site classifications (Fig. 10). The most notable trend was the classification of some species into their preferred habitats. Species known to be rarely found in turbid waters, such as *Pocillopora eydouxi*, *P. verrucosa*, *Millepora platyphylla* and *Favia stelligera* were exclusively found in the offshore habitats (20, 21, 24). On the other hand, species preferring or tolerating silty conditions or turbid waters, *Moseleya latistellata*, *Trachyphyllia geoffroyi*, *Euphyllia glabrescens* and *E. (Fimbriaphyllia) sp.* (Fig. 10), were confined to inshore sites (Groups 1 & 2: Sites 1-17 Fig. 8). Species ubiquitous at most sites included *Acropora hyacinthus*, *Favites* species, *Porites* species and *Turbinaria frondens*.

Site ordination

The greatest separation (variance) of sites occurred on the higher axes of ordination (axis 1: eigenvalue = 0.362, and axis 2: eigenvalue = 0.194). Axes 3 and 4 had eigenvalues of 0.135 and 0.088 respectively.

Substrate characteristics (the proportion of live coral found on rock, sand or coral rubble) appeared most able to explain the distribution of sites on the axes (Fig. 11). Ordination in general confirmed those groups classified by TWINSpan. The grouping of Site 23 with sites on the Burrup Peninsula, just as TWINSpan grouped them, was also evident using the DECORANA ordination program. Both Site 19 and Site 23, which were grouped together by TWINSpan, were exposed to waves only from an easterly direction (Fig. 11).

GENERAL DISCUSSION

The groups of sites identified here by computer analysis are similar to those described by Marsh (1978). The Dampier Archipelago has a rich coral fauna compared with other areas in Western Australia (Wilson and Marsh, 1979), chiefly because there is a wide range of coastal habitats varying from exposed to very sheltered (Marsh, 1978; Semenuik *et al.*, 1982).

While there were no distinctly different coral assemblages in each habitat, 'indicator species' were present which, by their tolerance of (or preference for) certain environmental conditions, gave an indication of some site characteristics. The distribution of families in the sites classified by

DENDROGRAM	SPECIES	CODE	HABITAT TYPE			
			S	P	O	V
	Millepora intricata	MILI				•
	Pocillopora demicornis	POCD				•
	Stylophora pistillata	STPP				•
	Acropora sp. (small clumps)	ACR2				•
	Acropora sp. (coarse bushy)	ACR4				•
	Acropora sp. (fine staghorn)	ACR5				•
	Montipora sp. (encrusting)	MOM1				•
	Montipora sequituberculata	MONE				•
	Alveopora sp.	ALYS				•
	Goniopora sp. (encrusting)	GONI				•
	Caulastrea tumida	CAUT	•			
	Cyphastrea spp.	CYP1				•
	Cyphastrea serotilla	CYPS				•
	Berberetia mirabilis	FAYA				•
	Favia pallida	FAYP				•
	Leptastrea pruinosa	LEPP				•
	Plesiosireia versicolor*	PLEV				•
	Moseleya latistellata	MOSL	•			•
	Trachyphyllia geoffroyi	TRAG	•			•
	Mucedium tenuicostatum	MYCT				•
	Pectinia paeonia	PECP				•
	Catalaphyllia jardnei	CATJ	•			•
	Euphyllia glabrescens	EUPG	•			•
	Euphyllia (Fimbriaphyllia) sp.	EUPF	•			•
	Pterogyra sp.	PLR1	•			•
	Duncanopodmmia axifuga	DUNA	•			•
	Turbinaria bifrons	TURB	•			•
	Turbinaria mesenterina	TURM	•			•
	Turbinaria pellata	TURP	•			•
	Stylocoenello guentheri	STYG				•
	Acropora sp. (fine bushy)	ACR3				•
	Acropora sp. (Brown staghorn)	ACR6				•
	Acropora sp. (knobby staghorn)	ACR7				•
	Astreaopora myriophthalma	ASTM				•
	Pachyseris rugosa	PACR	•			•
	Coccolinia sp.	COSS				•
	Fungi fungus	FUNF				•
	Herpolitha limax	HERL	•			•
	Polyphyllia talpina	POLT				•
	Podabectia crustacea	PODC				•
	Porites sp. (columnar)	POR3				•
	Cyphastrea microphthalma	CYPM				•
	Favia favaus	FAYF				•
	Favia maxima	FAYX				•
	Favites chinensis	FATC				•
	Goniastrea edwardsi	GOSE				•
	Goniastrea pectinosa	GO3P				•
	Hydnophora pilosa	HYDR	•			•
	Montastrea curta	MOTC				•
	Montastrea valenciennesi	MOTV				•
	Diploastrea heliopora	DIPH				•
	Galaxea fascicularis	GALC	•			•
	Merulina ampliata	MERA				•
	Scaphophyllia sp.	SCA1				•
	Acanthastrea echinata	ACAE				•
	Echinopullia aspera	ECHA				•
	Ocyropsis lacera	OCYL	•			•
	Mucedium elephantotus	MYCE				•
	Mucedium sp.	MYC1				•
	Pectinia lectuca	PEC1	•			•
	Dendrophyllia sp. 1	DEN1				•
	Turbinaria reniformis	TURR				•
	Psammocora superficialis	PSAS				•
	Goniopora sp. (massive)	CON2				•
	Porites sp.	POR1				•
	Favia spp.	FAY1				•
	Favites spp.	FAT1				•
	Goniastrea aspera	GOSA	•			•
	Goniastrea australensis	GOSU	•			•
	Hydnophora microconos	HYDM				•
	Platygyra deedelea	PLAD				•
	Galaxea fascicularis	GALF	•			•
	Lobophyllia corymbosa	LOBC				•
	Turbinaria frondens	TURF				•
	Acropora sp. (coarse tabular)	ACR1				•
	Pavona decussata	PAYD				•
	Goniastrea retiformis	GOSR	•			•
	Hydnophora exesa	HYDE				•
	Platygyra sinensis	PLAS				•
	Favia mathai	FAYT				•
	Favites abdita	FATA				•
	Echinopora lamellosa	ECHL				•
	Lobophyllia hemprichii	LOBH				•
	Symphylia radians	SYMR				•
	Psammocora digitata	PSAD				•
	Porites sp. (branched)	POR2				•
	Acropora hyacinthus	ACRH				•
	Montipora ramosa	MOMR				•
	Favites pentagona	FATP				•
	Millepora platyphylla	MILP				•
	Psammocora superficialis	PSAS	•			•
	Pocillopora eydouxi	POCE				•
	Pocillopora verrucosa	POCV				•
	Pavona minuta	PAYM				•
	Goniopora tenuidens	GONT	•			•
	Favia stelligera	FAYS				•
	Leptoria phrygia	LETP				•
	Dendrophyllia sp. 2	DEN2				•
	Tubastrea aurea	TUBA				•

KEY

Habitat type - known preferences of species:
 S - silty or turbid
 P - protected
 O - clear waters
 V - variable conditions

Figure 10; Species classification (for all sites and species) produced by TWINSpan.

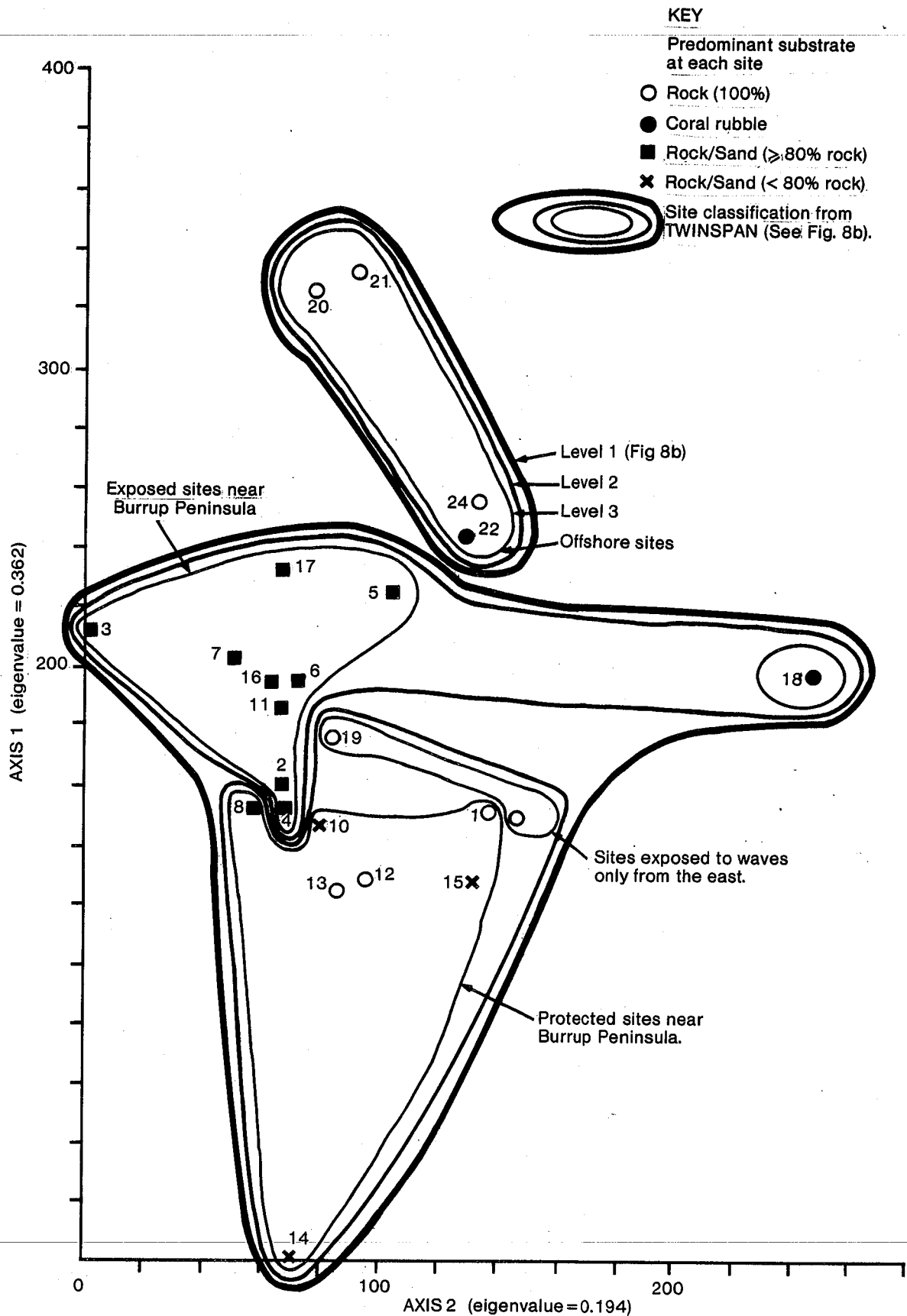


Figure 11; Ordination produced by DECORANA. These axes, 1 and 2, were determined from data with rare species and Site 9 removed. Axes are measured in units of standard deviation + 100.

TWINSpan conformed to what is known about the genera within them. Families tending to occur in exposed and offshore groups (Table 2) contained those species described as occurring in open water (Marsh, 1978; Veron and Pichon, 1976, 1979); for example, Thamnasteriidae (*Psammocora superficialis*), Pocilloporidae (*Pocillopora eydouxi*, *P. verrucosa*) and Agariciidae (*Pavona minuta*). Caryophylliidae, a family more common at sites on the Burrup Peninsula (Groups 1b, 2a, Figs. 8(b) and 9, Table 2), contained species tolerant of silty conditions; *Euphyllia glabrescens*, *E. (Fimbriaphyllia) sp.* and *Cataphyllia jardnei* (Fig. 10). Ubiquitous families were the Poritidae, Acroporidae, Faviidae, Mussidae and the Dendrophylliidae. These were indicated by Done (1983) to be tolerant of a wide range of physical and topographical conditions.

An offshore site, Lady Nora Island (Site 23, Fig. 1), was consistently classified with sites occurring on the Burrup Peninsula (Figs. 8 and 9). This was in contrast to its habitat; sheltered with relatively clear water. TWINSpan, on the basis of commonality of species grouped Site 23 with the inner sites. Site ordination (Fig. 11), also failed to separate this site with the other outer sites. One explanation is that the site had a high proportion of algal cover with corals occurring only on exposed rock. It was also more sheltered from wave action than the other outer sites; in effect a semi-lagoonal environment. Thus, both the Burrup sites and Site 23 were sheltered, the former being turbid and the latter non-turbid (L. Marsh pers. comm.).

The site classification (Fig. 8) appears to relate highly to the predominant substrate on which species were found (Fig. 11). To determine which group of variables corresponded to each axis, however, was difficult without detailed physical data for each site. The positioning of Site 14 (Boat Passage) away from other points on the ordination axes may reflect that it was a rather specialised habitat. This site was shallow, partly exposed to air at low tide and subject to strong currents (Marsh, 1978).

The higher species numbers at the more exposed sites may have resulted from the greater availability of habitats. Diversity was low in the protected sites because conditions were extreme; high turbidity and a wide temperature range occurred at these sites. The low species numbers at the outer sites may have resulted from a number of factors. These sites were more exposed than the Burrup sites to wave action and environmental disturbances such as cyclones (Connell, 1978). Marsh (1978) attributed a reduction of the number of genera at Kendrew Island (Sites 20 and 21) between 1974 (36 genera) and 1978 (21 genera) to *Acanthaster planci*, the Crown of Thorns starfish, predation and cyclone damage in the intervening period.

Furthermore, the percentage living coral cover at Kendrew Island was much higher in 1974 (50-80% coral cover) before predation and cyclone damage than in 1978 (1-50% coral cover).

While the techniques described here are useful for data already collected, further information can be obtained by collecting data with these types of analyses in mind. On this basis, two recommendations are made for future data collection. Firstly the need for pilot studies and secondly the measurement of physical data.

Before the main data collection begins, a pilot study to allow for initial analysis of data is useful for the following reasons:

- A suitable, efficient sampling method can be designed, so that the number of samples needed and the sample area size can be determined to minimise time, equipment and costs.
- Major environmental gradients may be recognised and the sampling modified accordingly.
- Field data sheets may be constructed, along with the appropriate codes for species, abundance and environmental factors etc. A standardised field sheet is particularly valuable when a number of workers are collecting data.

The measurement of environmental parameters when species are noted will help the final interpretation of biological data. The amount and type of physical data could include (see also Scheer, 1978):

- Water temperatures
- Water depths (species groups and average site depth)
- Salinity
- Turbidity
- Light
- Wind
- Currents
- Tidal amplitude and period
- Wave action

- Site aspect (sheltered, exposed)
- Habitat type (slope, reef flat, bottom topography)
- Sediment (composition, type, influx, resuspension)
- Substrate composition (sand, silt, rubble, rock)
- Biotic factors (competition, predation)

GENERAL CONCLUSION

TWINSpan classified sites into groups that correlated with the distribution of offshore, exposed and protected habitats within the Archipelago. TWINSpan could not determine distinct coral assemblages in terms of species groups, probably because of the variety of habitats available for colonisation. It is arguable therefore that this conclusion can be reached without using computer analysis. The treatment of data in this way however, is rapid, simple and would be a valuable aid to community classification for other environments where little physical data are known. DECORANA ordination produced similar results to TWINSpan; offshore groups being separated from inner sites. The failure of ordination, however, to produce any distinct separation of sites into groups was likely, once again, to be caused by the wide range of habitats. Gradients, if they occur are usually the result of many environmental variables such as current, sediment and wave exposure. For this study there was little environmental data available.

The successful application of the Cornell Ecology Programs to these coral data (as described in Section A) indicate that they could have more widespread use. TWINSpan, for example, may be used without accompanying physical site data and is a useful technique for site and species classification on the basis of presence/absence data alone. Hence, there is no reason why other data should not be used for analysis in this way. Examples of habitats or studies where TWINSpan may be useful include: island fauna and flora; algal communities; taxonomic studies (presence/absence of characters); bottom faunal communities; bird and mammal populations; gut floras; fish communities; and plant species stands.

Ordination using DECORANA, confirmed the results derived from TWINSpan, and its usefulness would increase with the availability of more environmental data. Though, if clearly defined environmental gradients are present, the use of ordination when little site data is available, may provide valuable information on the causes of site species composition.

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