

THE UNIVERSITY OF KANSAS  
PALEONTOLOGICAL CONTRIBUTIONS

August 1999

Number 10

STRATIGRAPHIC AND GEOGRAPHIC BRYOZOAN ABUNDANCE GRADIENTS IN  
THE CALCAREOUS SHALES OF THE WREFORD MEGACYCLOTHEM  
(LOWER PERMIAN, KANSAS)

Joseph F. Pachut and Roger J. Cuffey

Department of Geology, Indiana University-Purdue University at Indianapolis, 723 West Michigan Street, Indianapolis, IN 46202, jpachut@iupui.edu; Department of Geosciences, 412 Deike Building, Pennsylvania State University, University Park, PA 16802, cuffey@ems.psu.edu

*Abstract.*—Studies of bryozoans from the Wreford Megacyclothem have provided an extensive species-level database with which to examine paleoecologic patterns using multivariate statistical methods. Detrended correspondence (DCA) or gradient analysis was applied to fractional abundances of 17 species across 85 sampling intervals primarily within calcareous lithologies of the Wreford strata of Kansas.

Gradient analysis segregated the single robust trepostome species and all the encrusting-upright cystoporates from mostly small or delicate fenestrates, cryptostomes, and a ctenostome. The distribution of colonial morphologies and sizes suggests that these patterns were ecologically controlled and not the result of hydrodynamic sorting or taphonomic processes.

With minor exceptions involving the uppermost Speiser Shale and Schroyer Limestone Member of the Wreford Limestone, detrended correspondence analysis did not segregate any distinctive groupings of beds or lithologies. Speiser assemblages contain a diverse suite of abundant fenestrates and cryptostomes, whereas assemblages from the Schroyer maintain high fenestrate and cryptostome diversity while adding cystoporates and a trepostome.

The average position of each sampled interval on the first three detrended correspondence analysis axes indicates a general stratigraphic trend within the Wreford. Samples are arrayed in approximate stratigraphic order along axis 1 with axis 3 isolating Schroyer from Havensville samples.

Similar patterns result from gradient analyses of individual stratigraphic sampling intervals. Bryozoan faunas are dominated by fenestrates and cryptostomes, while trepostomes and cystoporates are generally rare or absent.

Regression of detrended correspondence analysis scores on a south-to-north distance of a sample from the Oklahoma-Kansas border indicates onshore-offshore gradients in species abundances within Wreford stratigraphic intervals. Significant trends occur in five of eight stratigraphic intervals, suggesting that gradient strength varied temporally within the uniformly shallow Wreford sea.

## INTRODUCTION

The bryozoans of the Lower Permian (Wolfcampian) Wreford Megacyclothem have been examined extensively over the past thirty years (Cuffey, 1967, 1977; Newton, 1971; Warner & Cuffey, 1973; Fry & Cuffey, 1976; Simonsen & Cuffey, 1977, 1980; Lutz-Garrihan & Cuffey, 1979; Cuffey & Hall, 1985; Pachut & Cuffey, 1991; Pachut, Anstey, &

Cuffey, 1991). This information and stratigraphic conclusions were thoroughly documented in these previous publications; a detailed survey of all the data and the reasoning supporting these conclusions is far beyond the scope of this paper. Studies of systematics, morphology, patterns of development and heterochrony, and paleoautecology have provided an extensive quantitative database with which to examine possible biotic and spatiotemporal gradients in

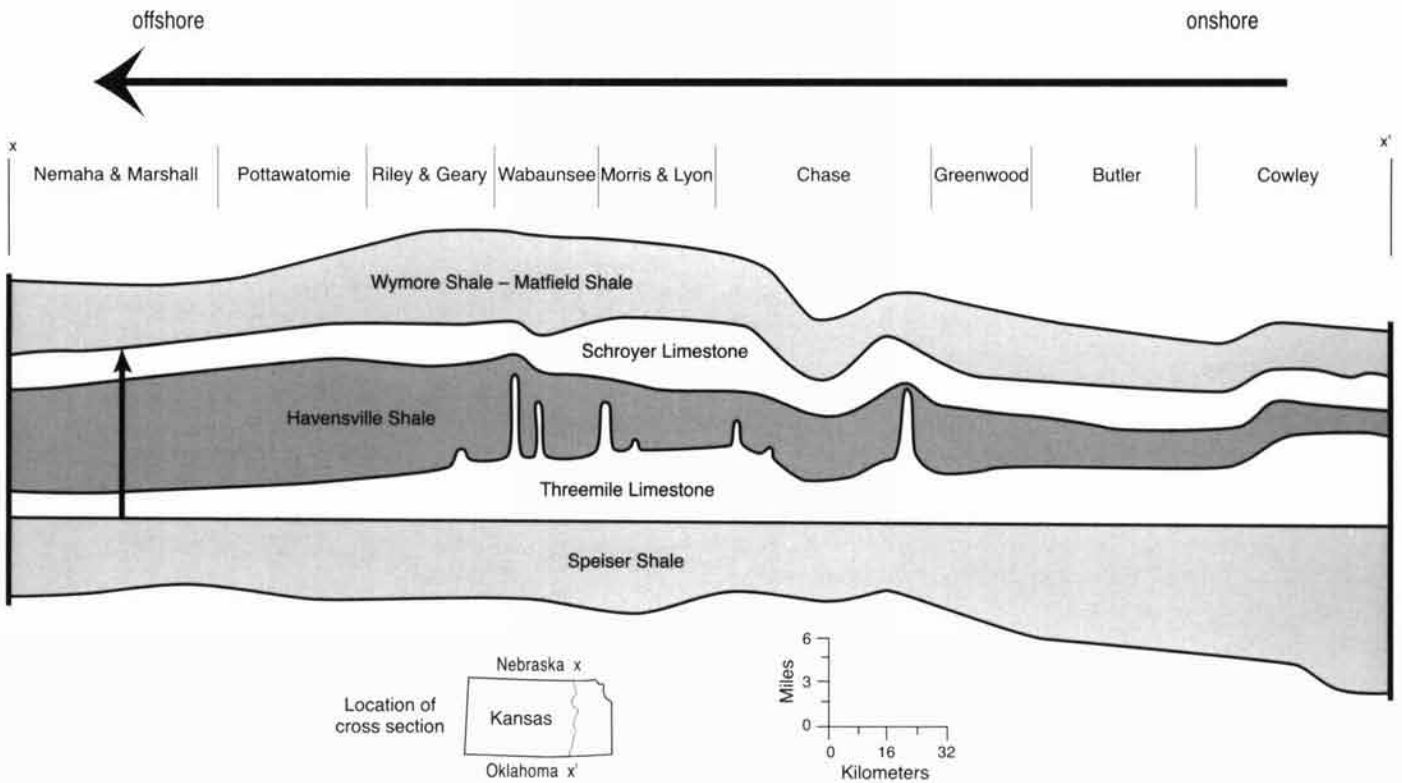


Figure 1. Schematic cross section of the Wreford Megacyclothem across Kansas (adapted from Cuffey, 1967, fig. 2, p. 14–15). County boundaries are indicated along the top, and Wreford stratigraphic units are indicated within the figure; Wreford Limestone indicated by vertical arrow (new).

these strata. The delineation of Wreford bryozoan species-level associations has paleosynecological significance in potentially permitting the delineation of paleoecological conditions, community structure, and biogeographic distributions that may ultimately be useful in establishing patterns of organism and community evolution.

*General stratigraphic setting.*—Rocks of the Wreford Megacyclothem (Fig. 1) are exposed along a south-to-north trending outcrop belt extending from northern Oklahoma across east-central Kansas into southernmost Nebraska. The Wreford Megacyclothem includes, in ascending order, the Speiser Shale of the Council Grove Group; the Threemile Limestone, Havensville Shale, and Schroyer Limestone Members of the Wreford Limestone; and the Wymore Shale Member of the Matfield Formation (Hattin, 1957). These units consist of a succession of 22 very thin beds that are roughly time parallel along the outcrop belt and within each of which may occur facies gradations (Hattin, 1957; Newton, 1971; Lutz-Garihan & Cuffey, 1979). These rocks were deposited in a shallow but extensive shelf sea and consist mostly of micritic limestones and mudstones indicative of paleoenvironments influenced partly by fluctuations in water depth and partly by variations in the supply of fine clastics (Cuffey & Hall, 1985). Some beds have since been recognized as diagenetically overprinted

by soil-forming (pedogenic) processes (Miller & West, 1993).

*Previous studies of Wreford bryozoan associations.*—Cuffey (1967) made an initial attempt to delineate Wreford paleo-environmental patterns prior to having a species-level taxonomic framework by examining the relative proportions of bryozoan growth forms present in sampled populations. He recognized that the same species sometimes constructed colonies that had different growth forms, whereas similar growth forms could be generated by different species. He also identified ten different colony growth forms among Wreford species (Table 1) but could not detect clear patterns of variation in percentages of growth forms within any of the lithologies across the Kansas outcrop belt. Several aspects of colony growth forms, however, became apparent within facies (Cuffey, 1967, fig. 6–10). 1. Delicate branching, small pinnate, and fenestrate growth forms numerically dominated faunas throughout the Wreford. 2. Bryozoans were most abundant in calcareous shales and cherty limestones presumably representing times of maximum marine submergences (based on the symmetry of the cyclothems around the cherty limestones). 3. Rarer growth forms occurred only in those same maximum-submergence rocks indicative of transgressions, whereas the three dominant colony forms were also found in units

indicative of more extreme ecological conditions associated with nearshore environments. 4. Delicate branching and fenestrate colonies became more abundant than small pinnate forms upward in the uppermost Speiser, probably paralleling environmental changes affecting sediment types, such as deposition of molluscan limestone followed by calcareous shale and then cherty limestone. 5. Bryozoans were far more abundant and widely distributed in more normal-marine, offshore (northern) areas of Kansas than in onshore (southern) areas where clastic input and possibly salinity varied.

Cuffey and Hall (1985) extended the earlier study of growth forms by completing the species-level taxonomy of Wreford bryozoans and examining their distributions qualitatively and quantitatively through Fourier analysis of species abundances. Only one species, the tiny, ramose cystoporate *Filiramoporina kretaphilia*, was environmentally restricted, occurring only in chalky limestones, particularly in carbonate mudbanks in the middle upper part of the upper Threemile Limestone. The other nineteen species were more widely distributed. Some were found only in calcareous shales and cherty limestones and others in varying abundances in most rock types. No mutually exclusive sets of species could be delineated that defined different communities that simultaneously occupied different habitats on the Wreford sea floor. Similarly, no consistent patterns were discernible in species abundances either from bed to bed or from nearshore habitats to offshore habitats (south-to-north) within the five principal calcareous shales of the Wreford: uppermost Speiser, middle Threemile, basal and topmost Havensville, and middle Schroyer. Cuffey and Hall (1985, p. 259) concluded that "no assemblages could be delineated by matching similar abundance variations among the Wreford bryozoans within particular rock types at several horizons."

The bryozoan assemblages they were able to define were based on an examination of overlapping ranges of species within lithologies (Cuffey & Hall, 1985, table 1, p. 258). 1. A full assemblage, containing all species except *Filiramoporina kretaphilia*, occurs in calcareous shale, brachiopod-molluscan limestone interbedded with calcareous shale, cherty limestone, and some algal limestones. 2. A partial assemblage, consisting of eight species, each with at least three zoaria, and single zoaria for each of two species, is found in molluscan limestone, most brachiopod-molluscan limestone, and some algal limestone. 3. A bank assemblage, comprising mostly (44 percent) *Filiramoporina kretaphilia* but with contributions from nine other species (four containing three or fewer zoarial fragments), is found in chalky limestone, especially in buildups. 4. A reduced assemblage containing seven species (three represented by three colony fragments each) occurs in green shale and gray-yellow mudstone. Assemblages 1 and 2 appear to have lived under normal-marine, offshore conditions. Reduced assemblage 4 lived in more nearshore, perhaps periodi-

Table 1. Current taxonomic assignments and colony growth forms of bryozoan species known to occur in the Wreford Megacyclothem of Kansas. The assignment of species of *Condranema* to the Ctenostomata follows Cuffey (1967). We recognize that they may be cyclostomes and require additional study (new).

Order (Suborder)	Species	Colony Growth Form
Ctenostomata	<i>Condranema magna</i>	encrusting threads
	<i>Condranema parvula</i>	encrusting threads
Fenestrata	<i>Acanthocladia guadalupensis</i>	small pinnate
	<i>Fenestella spinulosa</i>	delicate fenestrate
	<i>Fenestella tenax</i>	delicate fenestrate
	<i>Minilya binodata</i>	delicate fenestrate
	<i>Penniretepora auernigiana</i>	tiny pinnate
	<i>Penniretepora curvula</i>	tiny pinnate
	<i>Penniretepora flexistriata</i>	tiny pinnate
	<i>Polypora aestacella</i>	robust fenestrate
	<i>Polypora nodolinearis</i>	robust fenestrate
	<i>Protoretepora elliptica</i>	robust fenestrate
	<i>Septopora spinulosa</i>	delicate fenestrate
Cryptostomata (Rhabdomesina)	<i>Rhombopora lepidodendroides</i>	delicate ramose
	<i>Syringoclemis wrefordensis</i>	delicate ramose
Trepostomata	<i>Tabulipora carbonaria</i>	robust ramose
Cystoporata (Fistuliporina)	<i>Filiramoporina kretaphilia</i>	tiny ramose
	<i>Fistulipora carbonaria</i>	encrusting sheets
	<i>Fistulipora incrustans</i>	encrusting sheets
	<i>Meekopora prosseri</i>	upright sheet (bifoliate)

cally brackish and turbid waters, and assemblage 3 occurred only in chalky limestone mudbanks that formed, apparently, under conditions that were not conducive to the development of a normal, offshore, faunal association. Finally, 440 Fourier spectra (20 species from 22 samples) were calculated from species abundance profiles. They did not successfully delineate species assemblages, however. This could have resulted from the inability of spectra to delineate meaningful groupings or from random variations in species abundances across facies (Cuffey & Hall, 1985).

Holdener (1997) delineated geographic and clinal morphologic variation among populations of *Rectifenestella* sp. from the upper upper Speiser and middle Threemile of the Wreford. Patterns of variation were complex; morphologic characters that distinguished populations within each of these units generally were not the same, and those that were useful in distinguishing populations from both units did not always exhibit the same patterns of covariation. Populations of *Rectifenestella* sp. became more gracile with time, and patterns of morphologic variation in populations paralleled changes among populations of a Wreford trepostome bryozoan, *Tabulipora carbonaria* (Pachut & Cuffey, 1991). Among *T. carbonaria*, clinal geographic variation, different population growth trajectories, and heterochronic modifications of growth were demonstrated

Table 2. Ordination scores for 17 species calculated across 85 sampling intervals by detrended correspondence analysis (DCA) (DECORANA; Hill, 1979). Taxonomic assignments and growth habits of species are listed in Table 1 (new).

Species	Eigenvalue	Axis 1 0.42	Axis 2 0.22	Axis 3 0.14	Axis 4 0.12
<i>Fistulipora carbonaria</i>	386	173	-51	126	
<i>Tabulipora carbonaria</i>	345	133	132	195	
<i>Fistulipora incrustans</i>	321	150	244	254	
<i>Meekopora prosseri</i>	314	474	275	291	
<i>Protoretepora elliptica</i>	297	334	311	308	
<i>Fenestella tenax</i>	279	-43	1	73	
<i>Condranema magna</i>	194	153	232	125	
<i>Syringoclemis wrefordensis</i>	194	295	-105	246	
<i>Rhombopora lepidodendroides</i>	178	201	64	75	
<i>Penniretepora curvula</i>	175	27	144	20	
<i>Septopora spinulosa</i>	165	35	237	222	
<i>Polypora aestacella</i>	117	124	257	456	
<i>Minilya binodata</i>	96	87	236	74	
<i>Acanthocladia guadalupensis</i>	42	52	78	232	
<i>Penniretepora flexistriata</i>	0	42	172	136	
<i>Penniretepora auernigiana</i>	-1	97	217	12	
<i>Fenestella spinulosa</i>	-33	100	204	-23	

in three calcareous shales across an inferred onshore-offshore environmental gradient.

The purpose of this study is to expand on earlier quantitative analyses of colonial growth forms and qualitative attempts to delineate recurring assemblages of bryozoan species within the Wreford Megacyclothem of Kansas. Visual inspection of the relative frequencies of colony growth forms and of overlapping stratigraphic and geographic ranges of species has provided a general framework within which multivariate techniques may be applied to define more rigorously intergrading groups of species and to indicate the geographic and stratigraphic patterns of species occurrences within sampled units. Finally, the presence of clinal variation, previously delineated in *Tabulipora carbonaria* from three calcareous shales, can be evaluated in the entire bryozoan fauna across all calcareous shales of the Wreford.

## ANALYSES AND RESULTS

Wreford strata were extensively examined at about 250 localities by Cuffey (1967), Newton (1971), and Lutz-Garrihan (1976). We selected the eight most fossiliferous beds for analysis and obtained 509 samples. Limestone samples were mechanically broken; shales were disaggregated using standard kerosene treatment and sieved. The resulting approximately 16,000 bryozoan colony fragments were sorted into species. Table 1 lists the twenty species that occur in Wreford lithologies along with their colony growth forms.

Samples containing three or more species were retained for analysis; those containing fewer than three species provide little information on both the composition of species associations and on geographic groupings of samples.

Included were 85 fossiliferous samples representing collections from units at different localities within the Wreford and encompassing 11,015 colonies or fragments of colonies (hereinafter referred to as colonies). This screening of the data eliminated *Filiramoporina kretaphilia* and *Polypora nodolinaris*, leaving 18 species in the analysis. Numbers of colonies within each sample were transformed by calculating fractional abundances equal to the number of specimens of each species divided by the total number of specimens in the entire data set (Fishbein & Patterson, 1993).

An examination of sampled units across formations and members and across horizons at all localities permits a spatio-temporal analysis that attempts to delineate recurring associations of both species and sampling intervals occurring stratigraphically and geographically across Kansas. Calcareous shales were by far the most commonly sampled lithologic facies (68 samples), followed by calcareous shales interbedded with brachiopod-molluscan limestones (12 samples). Other facies contained locally sufficient numbers of species to justify analytical treatment, including gray-yellow mudstones (3 samples) and cherty limestones (2 samples). The overwhelming predominance of calcareous shales among our samples indicates both preserved abundances of fossils in that rock type and probably also ease of collection and extraction. As a byproduct, this lithologic dominance permits analyses to concentrate upon the detection of biotic variations related to environmental factors operating along onshore-offshore gradients across the shallow Wreford sea that are independent of sediment type.

Cuffey (1967) and Lutz-Garrihan and Cuffey (1979) interpreted the depositional environments of these rocks. Calcareous shale appears to have been deposited in normal salinity, low turbulence, shallow-marine waters, perhaps between 3 and 16 meters deep (see also Lane, 1964). Similar conditions prevailed during deposition of cherty limestone, possibly during times of deepest waters in the Wreford sea (but only slightly deeper than 16 m). Brachiopod-molluscan limestone also accumulated under similar conditions that at times were somewhat more turbulent and perhaps of slightly lower salinity than those of calcareous shales. In contrast, gray-yellow mudstone represents very shallow, nearshore, slightly brackish-marine habitats, perhaps shallower than 3 meters (Cuffey, 1967; Lutz-Garrihan & Cuffey, 1979). In places, these rocks have also been overprinted by soil-forming processes, as noted above. In general, each calcareous shale unit has been interpreted as representing an approximately time-parallel deposit that graded from nearshore environments in southern Kansas to progressively more offshore areas northward (Cuffey, 1967, p. 70-71).

*Detrended correspondence analysis (DCA).*—DCA, also referred to as gradient analysis, is an ordination technique related to principal component analysis. It is based on an eigenvalue algorithm that simultaneously calculates sample



Table 3. Ordination scores calculated for 85 sampling sites across 17 species using DCA (DECORANA; Hill, 1979). A generalized stratigraphic section (Simonsen & Cuffey, 1980, fig. 1) and a locality map (Pachut & Cuffey, 1991, fig. 1) have been published previously. County abbreviations are followed by a sample number and stratigraphic unit code. Abbreviations of Kansas counties are as follows: *BU*, Butler; *CH*, Chase; *CY*, Cowley; *GE*, Geary; *GR*, Greenwood; *LY*, Lyon; *ML*, Marshall; *MS*, Morris; *PT*, Pottawatomie; *RY*, Riley; *WA*, Wabaunsee. Unit codes are: *Sp*, upper upper Speiser Shale; *MTm*, middle Threemile Shale; *LLLH*, lower lower lower Havensville Shale; *ULLH*, upper lower lower Havensville Shale; *MH*, middle Havensville Shale; *UUH*, upper upper Havensville Shale; *LMSH*, lower middle Schroyer Limestone; *UMSH*, upper middle Schroyer Limestone (new).

Stratigraphic Unit Eigenvalue	Axis 1 0.42	Axis 2 0.22	Axis 3 0.14	Axis 4 0.12
ML03LLLH	322	141	113	178
GE01UMSh	270	122	161	177
PT09LLLH	267	67	26	118
CY21UUH	256	147	103	159
PT16LMSH	244	176	221	218
WA05LMSH	240	34	101	150
MS05UUH	238	33	61	126
GE18ULLH	234	235	149	167
CH24UUH	231	100	85	130
CH04MH	228	33	61	95
CH56UUH	223	193	64	124
PT04UMSh	221	200	91	151
GE18LMSH	221	161	163	174
GE04UMSh	218	61	165	186
CH22ULLH	216	7	64	102
MS06UUH	213	27	69	127
RY14ULLH	211	144	79	98
BU04UUH	209	103	77	148
GR01MH	203	109	71	121
GE30UMSh	203	162	104	119
CH56LMSH	198	145	52	106
GE30LMSH	197	112	133	141
CH44MH	194	63	70	83
GE18UMSh	193	125	70	74
GE18MTm	192	103	39	104
LY08LLLH	191	216	110	133
CY18Sp	190	231	0	162
GE01LMSH	190	67	193	145
ML03MTm	180	189	51	91
WA04MH	180	0	54	147
CH08UUH	180	58	51	135
CH44UUH	180	42	85	144
CH52LMSH	180	120	147	180
LY08ULLH	179	202	64	82
BU04ULLH	179	131	62	110
GR01LLLH	175	140	68	116
GR01ULLH	175	140	68	116
BU04Sp	170	188	79	81
CY09LMSH	168	123	199	227
RY06UUH	166	61	95	151
CH22MH	165	124	114	135
BU04UMSh	165	63	141	162
CH24ULLH	164	32	98	106
CH52MH	162	38	138	178
MS21LMSH	162	57	55	123
CY14Sp	161	145	95	120
WA04ULLH	159	38	91	162
CH18Sp	156	132	63	115
CH52LLLH	156	146	77	139
BU05Sp	153	181	73	86

Table 3. Continued.

Stratigraphic Unit Eigenvalue	Axis 1 0.42	Axis 2 0.22	Axis 3 0.14	Axis 4 0.12
RY14LLLH	143	51	100	168
BU04LMSH	142	103	144	237
CH39MTm	141	177	82	76
GE21Sp	140	152	101	112
BU04MH	140	132	106	111
RY04LMSH	140	124	178	105
RY04Sp	137	159	98	96
GE18UUH	137	39	120	169
GR01Sp	132	74	97	165
GR01UUH	130	152	52	162
ML01MTm	125	152	128	63
WA03Sp	112	130	78	141
ML01Sp	108	55	85	165
CH22UUH	108	74	140	104
PT04MH	105	42	91	205
CH53Sp	96	103	81	176
CH06MTm	91	135	129	34
CH39Sp	90	91	140	94
CH45UMSh	88	97	146	126
CH56Sp	87	102	105	148
PT09Sp	86	90	98	157
PT02Sp	78	77	90	203
CH32Sp	76	90	124	103
PT15Sp	75	83	133	203
CH04Sp	72	112	127	101
CH33Sp	72	111	127	92
LY01Sp	68	74	89	202
CH16Sp	66	76	141	122
ML03Sp	62	73	69	232
GE18Sp	55	75	135	136
LY13Sp	39	79	138	123
WA08Sp	37	81	140	115
MS22Sp	37	80	154	97
WA04LLLH	3	109	208	0
WA08MTm	0	93	161	51

scores (as the means of scores of taxa in the samples) and taxon scores (as the means of scores of the samples within which the taxa occur; Patzkowsky, 1995). DCA focuses on the abundances and distributions of taxa relative to physical environmental gradients that may be established by the interplay of multiple factors (Cisne & Rabe, 1978, p. 347). These factors may not be easily separable, but they parallel variation in some overriding environmental variable. In marine-benthic communities, including reef-dwelling bryozoans (Pachut & Cuffey, 1995), continuous change has been documented along depth-related or onshore-offshore gradients (Sanders & Hessler, 1969; Johnson, 1970, 1971). In fossil applications (Cisne & Rabe, 1978; Patzkowsky, 1995), ordination scores calculated by gradient analysis appear to represent the environment as experienced by the organisms and can be viewed as generalized measures of the environment. The technique has been used to delineate environmental and temporal gradients, to establish correlations, to evaluate sedimentation and sea level changes, and to examine clinal morphologic evolution (Cisne & Rabe, 1978; Cisne & others, 1980; Cisne, Molenock, & Rabe, 1980; Cisne, Chandlee, & others, 1982; Cisne, Karig, & others, 1982; Cisne & Chandlee, 1982;

Table 4. Data matrix illustrating the distribution of Wreford species across samples based upon first axis scores from detrended correspondence analysis (DCA). Values in the matrix represent original rather than fractional species abundances. Species and samples were rearranged to correspond to the scores generated by DCA; Kansas county abbreviations are listed in Table 3, and the eigenvalues for species and samples are listed in Tables 2 and 3. Two species groups and three sample groups are indicated along the top and right side of the figure, respectively, with average abundances also given for the latter (based on abundances per total number of cells in which species could have been found). Shading highlights differences discussed in the text. Average abundances within subsets, defined as combined species group-sample group intervals (number or letter, respectively, indicates shaded or unshaded rectangles), are as follows: 1A = 4.24, 2A = 7.41; 1B < 0.00, 2B = 4.52; 1C = 0.15, 2C = 22.21. Mean Brillouin diversity and evenness within each sample group = 1.23 and 0.72 for Group A, 1.05 and 0.69 for Group B, and 1.20 and 0.67 for Group C; calculations are discussed in the text (new).

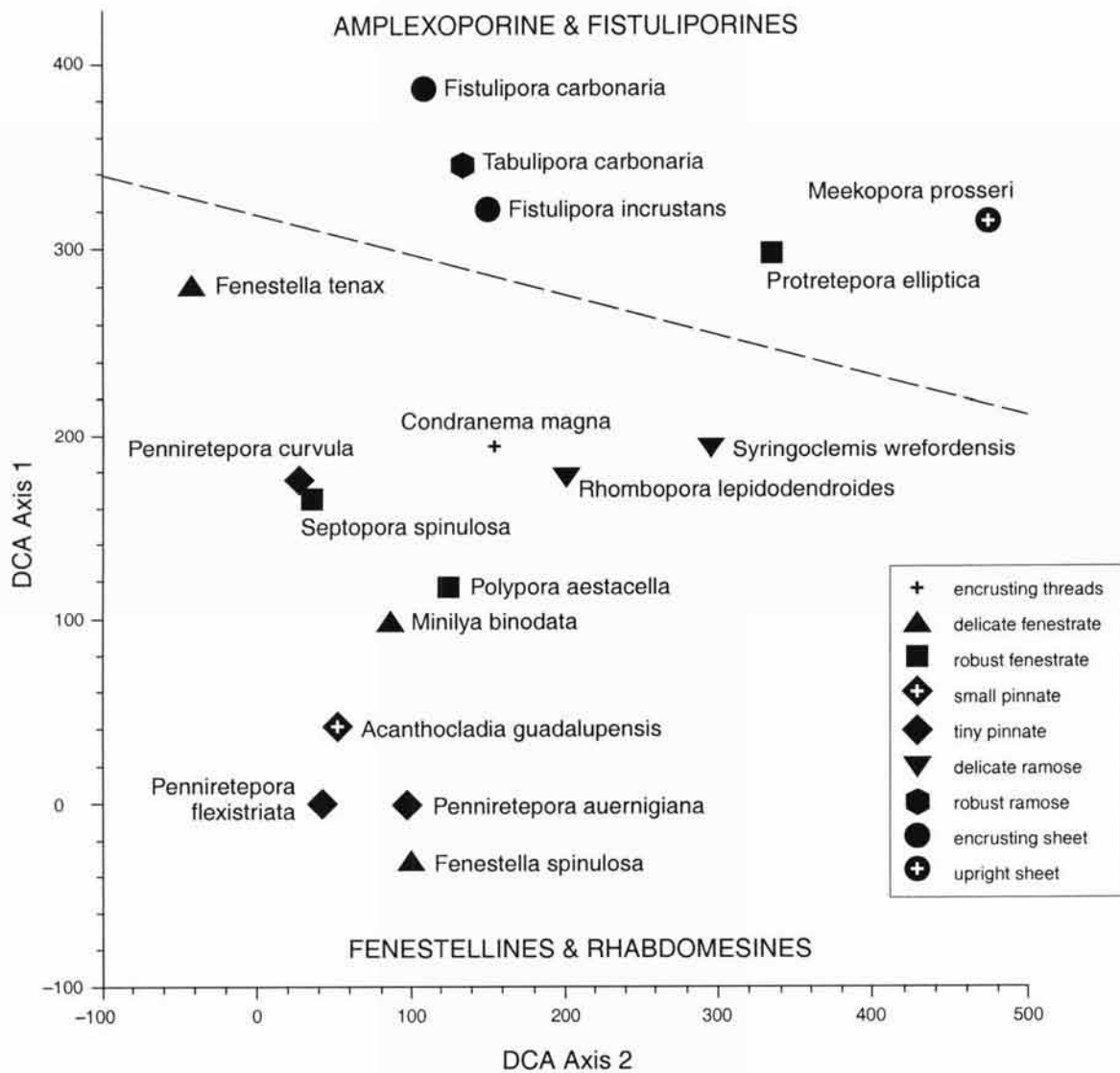


Figure 2. Plot of the positions of Wreford species on the first two DCA axes. High values on axis 1 separate a robust trepostome and cystoporates displaying encrusting-upright growth forms from generally small fenestrates, cryptostomes, and a single ctenostome. *Protretepora elliptica*, a robust fenestrate, is also allied with the trepostome-cystoporate group. Detrended correspondence analysis scores are listed in Table 2 (new).

Table 4. For explanation, see facing page.

SPECIES GROUP			1					2										SPECIES GROUP				
SPECIES SCORE			386	345	321	314	297	270	194	194	178	175	165	117	96	42	0	-1	-33	SPECIES SCORE		
SAMPLE	UNIT	SAMPLE SCORE	<i>Fusulipora carbonaria</i>	<i>Tabulipora carbonaria</i>	<i>Fusulipora incrassata</i>	<i>Moekepora prosseri</i>	<i>Protostopora elliptica</i>	<i>Fenestella imax</i>	<i>Conobanema magna</i>	<i>Syringolemus vorfontenai</i>	<i>Rhombopora leptodermoides</i>	<i>Penniretepora curvula</i>	<i>Fenestella spinulosa</i>	<i>Phyllopora anastella</i>	<i>Mitribya bimodata</i>	<i>Acanthoblastia guadalupensis</i>	<i>Penniretepora flexistriata</i>	<i>Penniretepora aenigmatica</i>	<i>Septopora spinulosa</i>	TOTAL	AVERAGE ABUNDANCE	SAMPLE GROUP
ML05	LLLH	322	7	39	5		4															
GE01	UMSh	270		65	8	2	4															
PT09	LLLH	267	12	5																		
CN21	UUh	256		5																		
PT10	UMSh	244		5																		
WA05	UMSh	240																				
MS05	UUh	238		20	7																	
GE18	ULLH	234		5	4	2																
CI24	UUh	231	2	50	8																	
CI104	MH	228		1																		
CI56	UUh	223		5																		
PT04	UMSh	221			2																	
GE18	UMSh	221	1	7	6	1	10															
GE01	UMSh	218		9	1																	
CI22	ULLH	216																				
MS06	UUh	213	4	46	6		1															
RY14	ULLH	211		2																		
BU04	UUh	209		23																		
GR01	MH	205																				
GE30	UMSh	203		1																		
CI56	UMSh	198																				
GE30	UMSh	197	1	18	15	8	33															
CI44	MH	194																				
GE18	UMSh	193																				
GE18	MIm	192																				
LY08	LLLH	191					8															
CN18	Sp	190		2																		
GE01	UMSh	190					1															
ML05	MIm	180																				
WA04	MH	180																				
CI108	UUh	180																				
CI44	UUh	180		2																		
CI52	UMSh	180			1																	
LY08	ULLH	179																				
BU04	ULLH	179																				
GR01	LLLH	175		5																		
GR01	ULLH	175		5																		
BU04	Sp	170																				
CN09	UMSh	168					2															
RY06	UUh	166																				
CI22	MH	165																				
BU04	UMSh	165		2	2																	
CI24	ULLH	164																				
CI52	MH	162			1																	
MS21	UMSh	162																				
CN14	Sp	161																				
WA04	ULLH	159																				
CI18	Sp	156																				
CI52	LLLH	156		2																		
BU05	Sp	155																				
RY14	LLLH	145																				
BU04	UMSh	142		3	2																	
CI39	MIm	141																				
GE21	Sp	140																				
BU04	MH	140																				
RY04	UMSh	140		1	1																	
RY04	Sp	137																				
GE18	UUh	137																				
GR01	Sp	132					1															
GR01	UUh	130		1	1																	
ML01	MIm	125																				
WA03	Sp	112																				
ML01	Sp	108					4															
CI22	UUh	108		1	1																	
PT04	MH	105																				
CI53	Sp	96																				
CI106	MIm	91			1																	
CI39	Sp	90																				
CI45	UMSh	88																				
CI56	Sp	87			1																	
QF09	Sp	86																				
PT02	Sp	78																				
CI52	Sp	76		1																		
PT15	Sp	75					2															
CI104	Sp	72																				
CI33	Sp	72																				
LY01	Sp	68																				
CI16	Sp	66																				
ML03	Sp	62																				
GE18	Sp	55		1			2															
LY13	Sp	39																				
WA08	Sp	37																				
MS22	Sp	37																				
WA04	LLLH	3																				
WA08	MIm	0																				
TOTAL			17	335	73	13	89	1423	60	208	1933	167	1491	26	554	3360	171	92	975	10996	TOTAL	
AVERAGE					1.24																	

Cisne, Gildner, & Rabe, 1984; Springer & Bambach, 1985; Anstey, Rabbio, & Tuckey, 1987; Anstey & Rabbio, 1990; Patzkowsky, 1995). Results of these studies support the view (Warme & others, 1976) that such taphonomic factors as time averaging may enhance the expression of paleocommunity gradients relative to similar gradients in living communities.

*Gradients across formations and members.*—Sample- and species-ordination scores for Wreford fractional abundance data were calculated using the program DECORANA (Hill, 1979). An initial DCA analysis indicated that *Condranema parvula* was an outlier distorting gradient structure. Its score on axis 2 was 892, while scores for all other species ranged between 0 and 366. It was essentially behaving like *Filiramoporina kretaphilia* and *Polypora nodolinearis* in forming a dense monotypic accumulation. *C. parvula* was therefore removed, and DCA was computed again using 17 species and 85 sampling intervals, reducing the sample size to 10,996 colonies. Both species and samples (Tables 2 and 3, respectively) were reordered in Table 4 to match the arrangements produced by DCA. This diagram illustrates resulting gradational patterns more clearly than the individual sets of DCA scores. Figure 2 plots scores of each species on the first two DCA axes to illustrate graphically the associations of species.

Two groupings of species are apparent in Table 4 and Figure 2 and form opposite poles of an abundance gradient (most easily seen in Table 4 in the upper left to lower right). Species Group 1 contains only robust and encrusting colonial morphologies that include the single trepostome and all of the cystoporate bryozoans sampled from the Wreford. These include the robust trepostome *Tabulipora carbonaria*; two encrusting cystoporates, *Fistulipora carbonaria* and *Fistulipora incrustans*; and the upright, sheet-forming cystoporate *Meekopora prosseri*. *Protoretopena elliptica*, a robust fenestrate, also displays an affinity for the trepostome-cystoporate grouping (Fig. 2). Abundances range between 1 and 66 specimens per sample with an average abundance of only 1.24 specimens per sample. Species are not uniformly distributed; most have high DCA scores and occur at the end of the sample gradient.

Species Group 2 contains all of the remaining fenestrate and cryptostome bryozoans, nearly all of which have small or delicate growth forms. *Fenestella spinulosa*, *F. tenax*, *Minilya binodata*, and *Septopora spinulosa* are delicate fenestrates, whereas *Acanthocladia guadalupensis* is a small pinnate fenestrate, and *Penniretepora auernigiana*, *P. curvula*, and *P. flexistriata* are tiny pinnate fenestrate morphotypes. Both of the sampled cryptostomes, *Rhombopora lepidodendroides* and *Syringoclemis wrefordensis*, also display delicate ramose growth habits. Finally, one questionable ctenostome, *Condranema magna*, exhibits an encrusting, threadlike growth form. Only the robust fenestrate *Polypora aestacella* deviates from this overall pattern of small and delicate morphologies, but it occurs sporadically in only 7 samples

and at low abundances (26 total specimens with 10 specimens in one sample). Species abundances in Group 2 range between 1 and 443 specimens per sample with a mean abundance of 10.26 specimens per sample. In contrast to Species Group 1, these species become more common as sample DCA scores decrease (Sample Group C in Table 4).

The ordination of samples (the sample gradient) was divided into three groupings (Table 4). The boundary of Sample Group A was placed between samples GE30LMSH and CH44MH, marking the last abundant occurrence of the trepostome and cystoporates of Species Group 1 (see Table 3 for explanation of sample codes). Sample Groups B and C were divided between samples ML01MTm and WA03Sp. This boundary is more arbitrary than the first but was located at a point where a trend of small decreases in sample scores was punctuated by a larger decrease in scores (from 125 to 112). Larger changes occur near the origin of DCA axis 1: a decrease of 16 units occurs between GE18Sp and LY13Sp; and a decrease of 34 units between MS22Sp and WA04LLLH. The first change, however, involves only three samples that contain the same species and have the same abundances as other samples in Sample Group C. The larger change of scores separates two samples, WA04LLLH and WA08MTm, that have only two and five species, respectively. They do not warrant designation as a fourth sample group and were also retained in Sample Group C.

Abundances of specimens vary between sample groups (not only across samples). Raw abundances for Group A range between 1 and 265 specimens with an average abundance of 6.47 specimens per sample (total of 374 possible collections of species within localities), between 1 and 77 specimens with a mean of 3.26 specimens per sample for Group B (663 samples), and between 1 and 443 specimens with an average of 15.72 specimens per sample for Group C (408 samples).

Combining the species and sample gradients illuminates several distributional and abundance patterns. First, representatives of Species Group 1, consisting of a trepostome, all Wreford cystoporates, and one robust fenestrate, occur abundantly only in Sample Group A where their average abundance is 4.24 specimens per sample. These species are virtually absent in the other two sample groups, where average abundances do not exceed 0.15 specimen per sample (in Species Group C). Fenestrates and cryptostomes of Species Group 2 occur even more abundantly within Sample Group A than do species of Group 1, having an average abundance of 7.41 specimens per sample. Of these, *Fenestella tenax*, *Rhombopora lepidodendroides*, *Acanthocladia guadalupensis*, and *Septopora spinulosa* are most abundant. These species continue to dominate Sample Group B, having an average abundance of 4.52 specimens per sample. Here, species from Group 1 are very sparsely distributed (average abundance < 0.01),



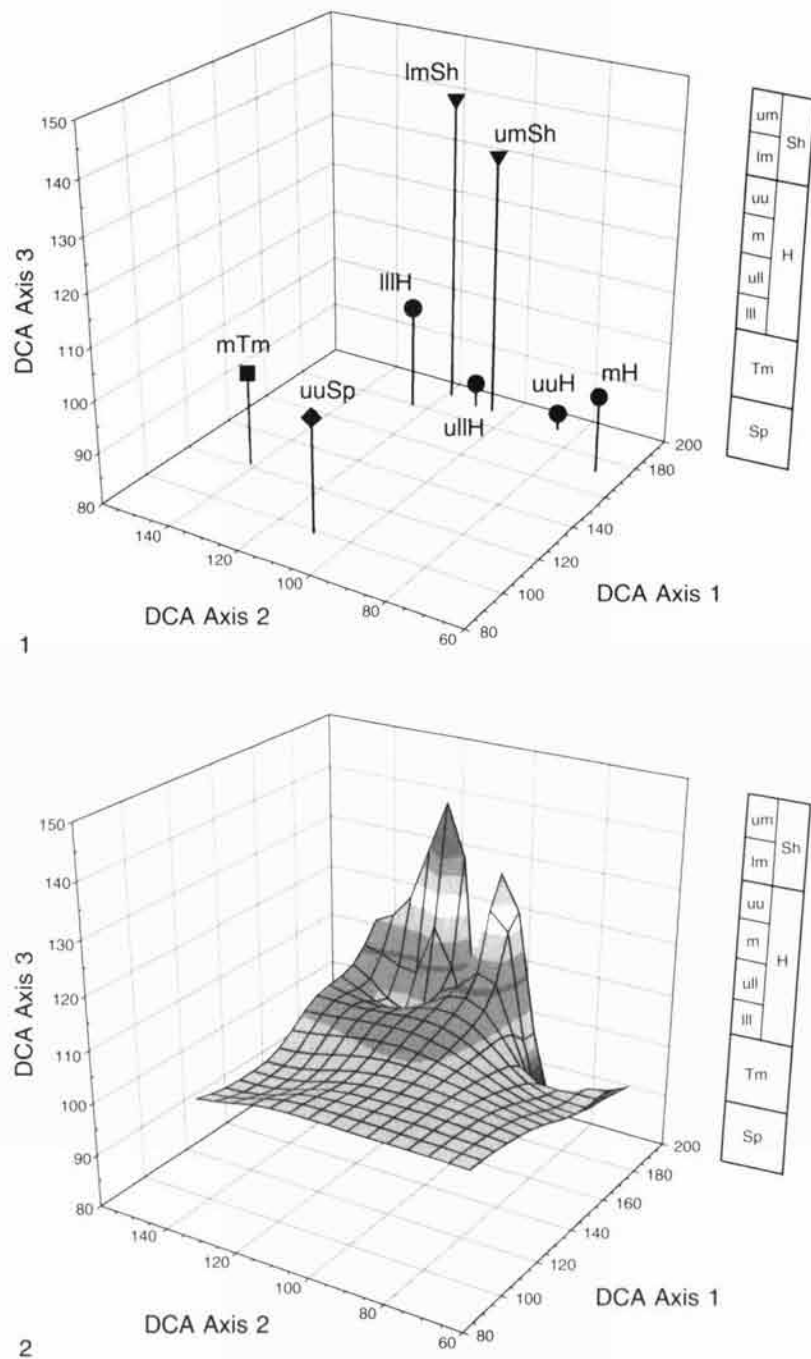


Figure 3. Three-dimensional plots of the mean positions of Wreford stratigraphic sampling intervals on the first three detrended correspondence analysis (DCA) axes. 1, Raw means and 2, 3-dimensional fitted surface illustrate a stratigraphic gradient that extends from the Speiser and Threemile (lower left) to Havensville (lower right) to Schroyer (upper right) sampling intervals. The schematic stratigraphic column indicates finer-scale subdivisions of the Havensville and Schroyer Members discussed in the text. DCA scores for each sample across axes are listed in Table 3; the distributions of samples and species on DCA axis 1 are listed in Table 4 (new).

which lowers the mean overall species abundance across samples to 3.26 specimens per sample. Sample Group C has the most abundant and diverse assemblage of fenestrates and cryptostomes. In addition to the dominant species of Groups A and B—*Fenestella tenax*, *Rhombopora lepidodendroides*, *Acanthocladia guadalupensis*, and *Septopora spinulosa*—

Group C has large numbers of *Fenestella spinulosa* and *Minilya binodata* along with three nearly ubiquitous species of *Penniretepora* that occurred more sporadically in the other two sample groups: *P. curvula*, *P. flexistriata*, and *P. auernigiana*. Considering only the portions of samples that include species of Group 2, mean abundance is 22.21

specimens per sample. If Species Group 1 is added (average = 0.15 specimen per sample), the average abundance across samples decreases to 15.72 specimens per sample.

In summary, these Wreford samples are dominated by fenestrates and cryptostomes. Cystoporates and a trepostome contribute, however, at one end of the gradient where DCA scores are high (on axis 1). More abundant and diverse fenestrates and cryptostomes occur at the other end of the gradient when DCA scores are low. A similar taxonomic dichotomy was noted in the Devonian Threeforks Formation (Prezbindowski & Anstey, 1978) and across Upper Ordovician bryozoan biomes or provinces (Anstey, 1986). In the latter, the Cincinnati biome was dominated by trepostomes and cystoporates and the Red River-Stony Mountain province by cryptostomes.

Species diversity shows this pattern well with some interesting caveats. Diversity indices have been interpreted as indicators of important aspects of community organization and structure including niche partitioning, trophic stability, and food chain length (see Pachut & Anstey, 1979, and references therein; Dodd & Stanton, 1990). The Brillouin diversity index (Pielou, 1974) was calculated for each sample and sample group. Unlike the more widely used Shannon-Weaver index, Brillouin's equation was developed to work with samples rather than with infinitely large populations. This and most other dominance diversity measures, however, confound species richness or number with evenness, a measure of the numerical distribution of species within a sample (Pielou, 1974); evenness is maximized (ranges between 0.0 and 1.0) when species have equal numbers of individuals (here, individuals = colonies or colony fragments).

For these Wreford samples, mean diversity and evenness are highest in Sample Group A (1.22 and 0.72, respectively) due to the presence of cystoporates and a trepostome and because the abundances of species are more uniform than in the other two groups. Sample Group B has the lowest diversity (1.05) but the second highest index of evenness (0.69). Although fewer species occur, abundances are similar across species. The mean diversity of samples from Sample Group C is only slightly lower (1.20) than that of Group A, but it is coupled with the lowest value of evenness (0.67). Diversity levels are maintained in spite of the rarity of species from Species Group 1 because species of fenestrates and cryptostomes of Species Group 2 reach their maximum diversity in Group C. Large numbers of approximately one-half of the species, however, create disparities in abundances within the assemblage that lower species evenness. Therefore, the single full assemblage of Cuffey and Hall (1985) has been subdivided based upon subtle, intergradational variations in species numbers and abundances, even within rocks of relatively uniform lithology.

Species assemblages occur across Kansas and do not appear to have been restricted to single time units or to

specific formations or members of the Wreford Megacyclothem. This is not surprising because these fossiliferous samples are primarily from calcareous shales, all very similar lithologies that were apparently deposited under relatively similar environmental conditions across a very gentle onshore-offshore (south-to-north) depth gradient. The various species appear to have been responding to spatially patchy paleoenvironmental conditions that did not affect preservable characteristics of the accumulating sediments in different regions of the Wreford sea.

*Stratigraphic gradients.*—The mean location of each sampled stratigraphic interval on the first three DCA axes is plotted in Figure 3.1 (data in Table 2). Axis 1 arrays samples approximately in ascending stratigraphic order: Speiser, Threemile, Havensville, and Schroyer, with some overlap among samples of the latter two members. Overlap is removed by axis 3, which separates Schroyer samples from all others. This suggests the presence of a complex stratigraphic gradient extending from the Speiser and Threemile toward the Havensville and curving upward to the Speiser, represented by the fitted surface in Figure 3.2. As discussed above, the scarcity of trepostomes and cystoporates sets Speiser, and to a lesser extent, Threemile samples apart from all others. Differing abundances and diversity of fenestrates and cryptostomes differentiate between Havensville and Schroyer samples. Holdener (1997) studied morphologic patterns in populations of *Rectifenestella* sp. from the Wreford. Populations from the Speiser and, to a lesser extent, the Threemile were also distinct morphologically from those of other members. The lack of independent physical evidence preserved in the rocks, however, leaves the sources of these differences unresolvable.

*Faunal composition of individual stratigraphic intervals.*—Sample and species ordination scores were calculated separately for each of eight Wreford sampling intervals to evaluate possible stratigraphic differences in faunal composition. For each analysis, we omitted species that only occurred within a single sample and sample locations containing fewer than two co-occurring species. As before, both species and samples were reordered in data matrices (Table 5–12) to match the arrangements produced by DCA.

It is not surprising that fenestrates and cryptostomes dominate assemblages across sampling intervals. In fact, after removing single-species occurrences, the middle Threemile and middle Havensville contain no trepostomes or cystoporates at all; and two other intervals, the upper upper Speiser and upper lower lower Havensville, include only one trepostome species (*Tabulipora carbonaria*). The lower lower lower Havensville and upper middle Schroyer include *T. carbonaria* and a single species of *Fistulipora* each (*F. carbonaria* and *F. incrustans*, respectively). Only the upper upper Havensville and the lower middle Schroyer have more than two trepostome and cystoporate species.

Table 5. Distribution of species and sample intervals for the upper upper Speiser. Original abundances have been rearranged to correspond to the placement of species and samples by detrended correspondence analysis. Species occurring in fewer than two samples and samples containing fewer than two species were removed prior to the analysis. Scores = ordination values calculated by detrended correspondence analysis; Kansas county abbreviations listed in Table 3 (new).

Species	upper upper Speiser																				Total			
	Score	ML03	ML01	GR01	GE18	CH18	LY01	PT02	PT09	WA03	CY14	PT15	CH56	GE21	RY04	CH32	CH16	LY13	CH39	CH04		WA08	CH33	MS22
<i>F. tenax</i>	272	113	106	93	88	88	85	84	65	65	57	52	50	45	34	33	22	18	14	14	12	11	0	0
<i>S. urefordensis</i>	253	22	8	1	14	1	1	3	7	1	24	2	1	1	1	26	17	4	26	6	6	20	334	
<i>P. elliptica</i>	181	4	1	1	2	1	1	2	3	2	2	2	2	2	2	5	1	1	1	1	3	1	1	46
<i>C. magna</i>	116	6	1	1	6	1	1	1	1	1	1	1	2	1	1	2	1	1	1	1	1	1	1	13
<i>A. guadalupensis</i>	106	223	443	56	294	11	179	24	94	24	6	161	32	7	10	97	155	242	74	68	202	48	269	2496
<i>R. lepidodendroides</i>	45	2	88	30	28	27	33	6	44	29	18	34	18	25	49	53	29	36	75	16	44	53	49	784
<i>S. spinulosa</i>	-9	11	55	20	49	14	3	3	12	1	4	28	3	2	14	14	14	27	20	10	32	6	63	377
<i>P. flexistriata</i>	-24	33	33	32	32	11	7	7	7	7	9	9	3	3	2	3	5	11	13	1	37	4	12	162
<i>P. curvula</i>	-35	11	11	11	11	1	1	1	1	1	1	1	1	1	1	5	5	5	13	1	3	8	2	67
<i>T. carbonaria</i>	-56	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	2
<i>P. auernigiana</i>	-68	11	11	11	17	17	4	4	4	4	32	32	7	3	8	18	87	29	65	17	13	6	31	83
<i>M. binodata</i>	-73	2	6	6	52	2	7	3	3	2	24	24	5	2	5	97	79	171	72	62	185	52	327	1999
<i>F. spinulosa</i>	-92	37	1	156	7	7	15	15	15	2	2	24	5	2	5	97	79	171	72	62	185	52	327	1999
Total	258	855	137	681	47	235	33	190	56	30	317	63	43	76	321	388	533	371	177	526	181	826	6086	

Table 6. Distribution of species and sample intervals for the Threemile. Original abundances have been rearranged to correspond to the placement of species and samples by detrended correspondence analysis. Species occurring in fewer than two samples and samples containing fewer than two species were removed prior to the analysis. Scores = ordination values calculated by detrended correspondence analysis; Kansas county abbreviations listed in Table 3 (new).

Species	middle Threemile										Total				
	Score	ML03	GE18	CH39	ML01	CH06	WA08	Score	ML03	GE18		CH39	ML01	CH06	WA08
<i>Syringoclemis urefordensis</i>	310	155	134	111	93	60	0	310	155	134	111	93	60	0	5
<i>Rhombopora lepidodendroides</i>	138	5	61	20	33	14	5	138	5	61	20	33	14	5	133
<i>Pennimtopora curvula</i>	88	1	1	2	2	4	6	88	1	1	2	2	4	6	
<i>Acanthocladia guadalupensis</i>	71	18	18	2	11	1	22	71	18	18	2	11	1	22	43
<i>Fenestella spinulosa</i>	-41	3	3	3	3	12	55	-41	3	3	3	3	12	55	81
Total	7	7	84	22	35	19	27	268	7	84	22	35	19	27	268

Table 7. Distribution of species and sample intervals for the lower lower lower Havensville. Original abundances have been rearranged to correspond to the placement of species and samples by detrended correspondence analysis (DCA). Species occurring in fewer than two samples and samples containing fewer than two species were removed prior to the analysis. Scores = ordination values calculated by DCA; county abbreviations listed in Table 3. Lower lower lower Havensville samples are generally arranged from south to north (ML03 to RY14, except for the last of these; new).

		lower lower lower Havensville						
Species	Score	ML03	PT09	LY08	CH52	GR01	RY14	Total
	Score	253	138	96	87	80	0	
<i>Fistulipora carbonaria</i>	286	7	2					9
<i>Tabulipora carbonaria</i>	276	39	3		2	5		49
<i>Syringoclemis wrefordensis</i>	138	1	1	1		3		6
<i>Condranema magna</i>	134	1		1		2	1	5
<i>Rhombopora lepidodendroides</i>	105	6	3	31	9	77	13	139
<i>Fenestella tenax</i>	82		8			15	18	41
<i>Acanthocladia guadalupensis</i>	-20		1	3	5	20	34	63
<i>Septopora spinulosa</i>	-114					6	20	26
Total		54	18	36	16	128	86	338

Table 8. Distribution of species and sample intervals for the upper lower lower Havensville. Original abundances have been rearranged to correspond to the placement of species and samples by detrended correspondence analysis (DCA). Species occurring in fewer than two samples and samples containing fewer than two species were removed prior to the analysis. Scores = ordination values calculated by DCA; county abbreviations listed in Table 3 (new).

		upper lower lower Havensville							
Species	Score	GE18	RY14	GR01	BU04	WA04	CH22	CH24	Total
	Score	192	160	150	130	63	1	0	
<i>Tabulipora carbonaria</i>	273	3	2	5					10
<i>Syringoclemis wrefordensis</i>	237	1		3	3				7
<i>Condranema magna</i>	231	2	1	2				1	6
<i>Rhombopora lepidodendroides</i>	192	9	12	77	21	6	19	6	150
<i>Acanthocladia guadalupensis</i>	97			20	4	16	24	18	82
<i>Septopora spinulosa</i>	72	3	1	6	3	11	25	6	55
<i>Pennirelepora curvula</i>	24				3		8		11
<i>Fenestella tenax</i>	-36		3	15	6	14	147	41	226
<i>Minilya binodata</i>	-99						23	21	44
Total		0	3	10	3	0	0	1	23

Nonetheless, trepostome-cystoporate subgroups are not found because many samples contain only one to three specimens per species for these two taxa.

Sampling localities are generally arrayed geographically from south-to-north in the lower lower lower Havensville and upper middle Schroyer and from north-to-south in the lower middle Schroyer. Otherwise, DCA scores vary from one location to another across Kansas.

If not simply the result of small sample sizes, these compositional and distributional patterns may represent a paleontological example of the Gleasonian perspective on community ecology (see Roughgarden, 1989). In this model, communities are viewed as mere samples of a regional species pool that co-occur simply because they are capable of tolerating the existing environmental conditions (Jackson, Budd, & Pandolfi, 1996). It is possible that

patchy habitats and Gleasonian sampling (i.e., random) of taxa combined to produce no clear-cut compositional distinctions and often no geographic patterns within individual sampling intervals of the Wreford.

*Ecological gradients within stratigraphic intervals.*—DCA scores of samples were used to evaluate the presence of geographic, onshore-offshore, bryozoan faunal gradients in the Wreford. Many ecological gradients are weakly developed and do not possess sharp discontinuities in species abundances (Cisne & Rabe, 1978). In such instances, results of gradient analysis will show these ecological continua more accurately than will divisive clustering algorithms that force samples into discrete cluster groups. To test for statistically significant ecological or geographic gradients, sample ordination scores were regressed (using SYSTAT; Wilkinson, 1990) on south-to-north (onshore-to-offshore)



Table 9. Distribution of species and sample intervals for the middle Havensville. Original abundances have been rearranged to correspond to the placement of species and samples by detrended correspondence analysis (DCA). Species occurring in fewer than two samples and samples containing fewer than two species were removed prior to the analysis. Scores = ordination values calculated by DCA; county abbreviations listed in Table 3 (new).

middle Havensville										
Species	Score	CH44	BU04	GR01	CH04	CH22	WA04	CH52	PT04	Total
Score		151	126	114	104	91	40	17	0	
<i>Penniretepora curvula</i>	232	12	5		1				2	20
<i>Fenestella spinulosa</i>	204		4		2					6
<i>Syringoclemis wrefordensis</i>	175	2	2	2	23			2		31
<i>Rhombopora lepidendroides</i>	173	8	22	10		11		10		61
<i>Fenestella tenax</i>	107	10		6	58	1	10	33		118
<i>Septopora spinulosa</i>	-18	1	7	5	15	6	2	71	7	114
<i>Acanthocladia guadalupensis</i>	-37	3	6			2	7	34	9	61
<i>Mimilya binodata</i>	-64				2			6		8
Total		36	46	23	101	20	19	156	18	419

Table 10. Distribution of species and sample intervals for the upper upper Havensville. Original abundances have been rearranged to correspond to the placement of species and samples by detrended correspondence analysis (DCA). Species occurring in fewer than two samples and samples containing fewer than two species were removed prior to the analysis. Scores = ordination values calculated by DCA; county abbreviations listed in Table 3 (new).

upper upper Havensville												
Species	Score	CH22	GE18	RY06	GR01	CH44	CH08	MS06	BU04	CH24	MS05	Total
Score		133	101	91	87	68	62	27	25	21	0	
<i>Fenestella spinulosa</i>	227	51	2				5	7		4	3	72
<i>Condranema magna</i>	204	2								2		4
<i>Septopora spinulosa</i>	177	29	28	3		14	11	62		15	6	168
<i>Acanthocladia guadalupensis</i>	144	26	35	3	32	18	21	91	19	16	22	283
<i>Rhombopora lepidodendroides</i>	68	17	6	3	37	12	13	40	20	68	8	224
<i>Syringoclemis wrefordensis</i>	18	1			10		10	6	2	4	4	37
<i>Fenestella tenax</i>	-23	22	18	3		26	30	265	12	56	84	516
<i>Mimilya binodata</i>	-39		3			2		7		4	1	17
<i>Tabulipora carbonaria</i>	-59	9			1	2		46	23	50	26	157
<i>Penniretepora curvula</i>	-109		5					23	4		7	39
<i>Fistulipora carbonaria</i>	-132							4		2		6
<i>Fistulipora incrustans</i>	-171				1			6		8	7	22
Total		157	97	12	81	74	90	557	80	229	168	1545

distances of sampling localities from the Oklahoma-Kansas border (Cuffey, 1967; Pachut & Cuffey, 1991). This approach is analogous to the ones used to study the effects of environmental gradients on the evolution of Ordovician trilobites (Cisne & others, 1980; Cisne, Molenock, & Rabe, 1980). Significance values in Table 13 suggest that gradients were developed in five of the eight sampled Wreford units on either axis 1 or 2: upper upper Speiser, lower lower Havensville, middle Havensville, upper upper Havensville, and the lower middle Schroyer. The geographic development of ecological gradients in the calcareous shales in the middle Threemile, the upper lower Havensville Shale, and upper middle Schroyer lacks statistical support; small sample sizes (total of 19 sampling sites across the 3 intervals) and concentrations of sampling sites in relatively small geographic areas may have contrib-

uted to this. In addition, patchy environmental conditions in this shallow epicontinental sea throughout Wreford deposition could also account for variations in the development and strength of gradients. Gradients delineated in virtually all previous studies of other sequences have been much steeper, particularly those studied in the Taconic trench (Cisne & others, 1980; Cisne, Molenock, & Rabe, 1980; Cisne, Chandler, & others, 1982; Cisne, Karig, & others, 1982), where depths may have reached 500 meters. In stark contrast, Wreford deposition has been estimated to have taken place in shallow waters having depths ranging between 10 or 15 m up to a maximum of only 35 m (Cuffey, 1967; Newton, 1971; Lutz-Garrihan & Cuffey, 1979). Another possible explanation of the variable development of bryozoan abundance gradients, operating independently or together with rather patchy environmental

Table 11. Distribution of species and sample intervals for the lower middle Schroyer. Original abundances have been rearranged to correspond to the placement of species and samples by detrended correspondence analysis (DCA). Species occurring in fewer than two samples and samples containing fewer than two species were removed prior to the analysis. Scores = ordination values calculated by DCA; county abbreviations listed in Table 3. Lower middle Schroyer samples are generally arranged from north to south (BU04 to RY04, but with some reversals; new).

lower middle Schroyer													
Species	Score	BU04	MS21	CH52	CY09	CH56	GE30	GE18	WA05	PT16	GE01	RY04	Total
Score		181	129	98	88	82	61	55	48	15	1	0	
<i>Polypora aestacella</i>	282	6					10						16
<i>Acanthocladia guadalupensis</i>	260	13	3		2		24	4	1				47
<i>Penniretepora flexistriata</i>	250		1				1						2
<i>Syringoclemis wrefordensis</i>	181	1		2		1		1				1	6
<i>Condranema magna</i>	157	1					5	1		1			8
<i>Fistulipora incrustans</i>	139	2		1			15	6				1	25
<i>Tabulipora carbonaria</i>	89	3					18	7		5		1	34
<i>Septopora spinulosa</i>	88	7		10	4	1	97	7	3	9	3	1	142
<i>Rhombopora lepidodendroides</i>	79	6	3	6	1	6	166	16		3		3	210
<i>Fenestella tenax</i>	38		4			2	97	3	8	2	1		117
<i>Meekopora prosseri</i>	14						8	1					9
<i>Fistulipora carbonaria</i>	-4						1	1					2
<i>Penniretepora curvula</i>	-66						10	1			2		13
<i>Protoretepora elliptica</i>	-78				2		33	10	2	13	1		61
<i>Minilya bimodata</i>	-81						44	6		2	1	9	62
Total		39	11	19	9	10	529	64	14	35	8	16	754

Table 12. Distribution of species and sample intervals for the upper middle Schroyer. Original abundances have been rearranged to correspond to the placement of species and samples by detrended correspondence analysis (DCA). Species occurring in fewer than two samples and samples containing fewer than two species were removed prior to the analysis. Scores = ordination values calculated by DCA; county abbreviations listed in Table 3; upper middle Schroyer samples are generally arranged from south to north (GE18 to GE04, except for the last of these; new).

upper middle Schroyer								
Species	Score	GE18	GE30	CH45	BU04	GE04	Total	
Score		139	98	81	52	0		
<i>Rhombopora lepidodendroides</i>	162	5	4	1	11	3	24	
<i>Acanthocladia guadalupensis</i>	103			1	14	1	16	
<i>Fenestella tenax</i>	83	2			7	11	20	
<i>Fistulipora incrustans</i>	80				2	1	3	
<i>Septopora spinulosa</i>	-20	1	1		25	32	59	
<i>Condranema magna</i>	-28				1	1	2	
<i>Tabulipora carbonaria</i>	-99		1		2	9	12	
Total		7	6	3	62	58	136	

conditions, is that the steepness of environmental gradients may have varied temporally. It is regrettable that gradient steepness may have varied in concert with biotic or abiotic environmental conditions that left no obvious signatures in these rocks. The geographic and stratigraphic patterns discussed above, however, mirror those studied by Pachut and Cuffey (1991) in *Tabulipora carbonaria* from three Wreford units: the lower and upper Havensville Shale and the middle Schroyer Member. In an analysis of clinal morphological variation, multivariate analyses delineated statistically significant differences in morphology between populations of *Tabulipora*, evidence for heterochronic modifications, and an increase in developmental canalization in

an onshore-offshore direction. Additionally, Holdener (1997) found that patterns of morphologic variation in populations of *Rectifenestella* sp. paralleled changes in equivalent characteristics of *Tabulipora carbonaria* examined by Pachut and Cuffey (1991). Nearly identical patterns occurred stratigraphically in this study of Wreford bryozoan species abundances after pooling data within each of the sampled units. These similarities among studies suggest that, in spite of difficulties in identifying specific environmental parameters responsible for establishing them, onshore-offshore ecologic gradients in the Wreford Megacyclothem existed and can be delineated by detrended correspondence analysis.

## CONCLUSIONS

1. Detrended correspondence analysis (DCA) of the fractional abundances of bryozoan species from the Wreford Megacyclothem of Kansas permit the recognition of two species and three sample subgroups in the sampled calcareous shales. Subgroupings lack sharp compositional discontinuities and intergrade with one another.

2. The single, full species assemblage of Cuffey and Hall (1985) was partitioned by DCA into two assemblages based upon intergradational differences in the relative abundances of taxa across sampling sites. One assemblage includes moderate abundances of a trepostome and several cystoporates plus a single robust fenestrate, whereas the other generally lacks those species and includes abundant, mainly small fenestrates, cryptostomes, and a ctenostome.

3. No distinctive groupings of beds or lithologies were identified other than a cluster of samples collected mostly from the upper upper Speiser Shale, where trepostomes and cystoporates are nearly absent, and the monospecific assemblages containing either *Filiramoporina kretaphilia*, *Polypora nodolinearis*, or *Condranema parvula*. The three sample subgroups vary in relative species abundances, diversity, and evenness. This suggests environmental patchiness during the deposition of Wreford calcareous lithologies that, otherwise, appear to have been deposited under rather similar environmental conditions.

4. A general stratigraphic trend is indicated by a plot of the average position of each Wreford stratigraphic sampling interval on the first three DCA axes. Factors responsible for the establishment of this trend remain obscure.

5. Analyses of individual stratigraphic sampling intervals indicate similar biotic responses by bryozoan faunas throughout the deposition of Wreford sediments. Fenestrates and cryptostomes dominate, with only the upper upper Havensville and lower middle Schroyer displaying more than two species of trepostomes or cystoporates.

6. Significant geographic trends occurred in five of eight sampled stratigraphic intervals in spite of the shallowness of the onshore-offshore depth gradient across Kansas.

Table 13. Statistical significance of linear regressions of DCA scores for samples on onshore-offshore distances (south-to-north) across Kansas. DCA scores were calculated separately for each sampling interval. Sampled intervals are arranged in stratigraphic order with values representing the calculated level of significance for each regression; *N*, number of samples (new).

Sampling Interval	N	Axis 1	Axis 2
upper middle Schroyer	6	0.25	0.31
lower middle Schroyer	11	0.02**	0.99
upper upper Havensville	10	0.70	0.02**
middle Havensville	8	0.07*	0.28
upper lower lower Havensville	7	0.42	0.34
lower lower lower Havensville	6	0.27	0.05**
middle Threemile	6	0.62	0.59
upper upper Speiser	23	0.09*	0.05**

Levels of significance noted by \* at  $p \leq 0.10$ , \*\* at  $p \leq 0.05$ .

7. These stratigraphic and geographic abundance gradients parallel onshore-offshore clinal morphologic variation studied in *Tabulipora carbonaria* from the lower and upper Havensville Shale and middle Schroyer Member (Pachut & Cuffey, 1991) and in *Rectifenestella* sp. (Holdener, 1997). This strengthens our interpretation of the existence of an overall onshore-offshore gradient in the Wreford sea.

## ACKNOWLEDGMENTS

We thank R. L. Anstey, A. S. Horowitz, and S. J. Hageman for their reviews of the manuscript. The study also benefited from discussions with R. L. A. regarding the application of detrended correspondence analysis and interpretations of results. Finally, we thank R. E. Hall for assistance in preparing Figure 1. Partial support for this project (to J.F.P.) was provided by the National Science Foundation (grant #EAR-9506003) and by the donors of the Petroleum Research Fund of the American Chemical Society (grant #20464-B8).

## REFERENCES

- Anstey, R. L. 1986. Bryozoan provinces and patterns of generic evolution and extinction in the Late Ordovician of North America. *Lethaia* 19:33–51.
- Anstey, R. L., & S. F. Rabbio. 1990. Regional bryozoan biostratigraphy and taphonomy of the Edenian stratotype (Kope Formation, Cincinnati area): graphic correlation and gradient analysis. *Palaios* 4:574–584.
- Anstey, R. L., S. F. Rabbio, & M. E. Tuckey. 1987. Bryozoan bathymetric gradients within a Late Ordovician epeiric sea. *Paleoceanography* 2:165–176.
- Cisne, J. L., & G. O. Chandlee. 1982. Taconic foreland basin graptolites: age zonation, depth zonation, and use in ecostratigraphic correlation. *Lethaia* 15:343–363.
- Cisne, J. L., G. O. Chandlee, B. D. Rabe, & J. A. Cohen. 1980. Geographic variation and episodic evolution in an Ordovician trilobite. *Science* 209:925–927.
- . 1982. Clinal variation, episodic evolution, and possible parapatric speciation: the trilobite *Flexicalymene senaria* along an Ordovician depth gradient. *Lethaia* 15:325–341.
- Cisne, J. L., R. F. Gildner, & B. D. Rabe. 1984. Epeiric sedimentation and sea level: synthetic ecostratigraphy. *Lethaia* 17:267–288.
- Cisne, J. L., D. E. Karig, B. D. Rabe, & B. J. Hay. 1982. Topography and tectonics of the Taconic outer trench slope as revealed through gradient analysis of fossil assemblages. *Lethaia* 15:229–246.
- Cisne, J. L., J. Molenock, & B. D. Rabe. 1980. Evolution in a cline: the trilobite *Triarthrus* along an Ordovician depth gradient. *Lethaia* 13:47–59.
- Cisne, J. L., & B. D. Rabe. 1978. Coenocorrelation: gradient analysis of fossil communities and its applications in stratigraphy. *Lethaia* 11:341–364.
- Cuffey, R. J. 1967. Bryozoan *Tabulipora carbonaria* in Wreford Megacyclothem (Lower Permian) of Kansas. The University of Kansas Paleontological Contributions, Article 43 (Bryozoa):1–96.
- . 1977. Ctenostome bryozoans and burrowing barnacles of the Wreford Megacyclothem (Lower Permian; Kansas-Oklahoma-Nebraska). Geological Society of America, Abstracts with Programs 9:587–588.
- Cuffey, R. J., & W. B. Hall. 1985. Species-level bryozoan assemblages within the Lower Permian Wreford Megacyclothem of Kansas, Oklahoma, and Nebraska. In J. T. Dutro, & H. W. Pfefferkorn, eds., *Neuvième Congrès International de Stratigraphie et de Géologie du Carbonifère*, *Compte Rendu Volume 5*, Paleontology, Paleogeology, Paleogeography. Southern Illinois University Press. Carbondale and Edwardsville. p. 256–260.
- Dodd, J. R., & R. J. Stanton, Jr. 1990. *Paleoecology, Concepts and Applications*. John Wiley and Sons. New York. 502 p.
- Fishbein, E., & R. T. Patterson. 1993. Error-weighted maximum likelihood (EWML): a new statistically based method to cluster quantitative micropaleontological data. *Journal of Paleontology* 67:475–485.
- Fry, H. C., & R. J. Cuffey. 1976. *Filiramoporina kretaphilia*—a new genus and species of bifoliate tubulobryozoan (Ectoprocta) from the Lower Permian Wreford of Kansas. University of Kansas Paleontological Contributions, Paper 84:1–9.
- Hattin, D. E. 1957. Depositional environment of the Wreford Megacyclothem (Lower Permian) of Kansas. *Kansas Geological Survey Bulletin* 124:1–150.
- Hill, M. O. 1979. DECORANA, a FORTRAN program for detrended correspondence analysis and reciprocal averaging. Ecology and Systematics Department, Cornell University. Ithaca, New York.
- Holdener, E. J. 1997. Morphometric approaches to analysis of microevolution: morphologic variation in Paleozoic Bryozoa. Unpublished Ph.D. dissertation. University of Illinois. Urbana. 262 p.
- Jackson, J. B. C., A. F. Budd, & J. M. Pandolfi. 1996. The shifting balance of natural communities? In D. Jablonski, D. H. Erwin, & J. H. Lipps, eds., *Evolutionary Paleobiology*. University of Chicago Press. Chicago. p. 89–122.
- Johnson, R. G. 1970. Variations in diversity within benthic marine communities. *American Naturalist* 104:135–147.
- . 1971. Animal-sediment relations in shallow water benthic communities. *Marine Geology* 11:93–104.
- Lane, N. G. 1964. Paleogeology of the Council Grove Group (Lower Permian) in Kansas, based upon microfossil assemblages. *Kansas Geological Survey, Bulletin* 170:1–23.
- Lutz-Garihan, A. B. 1976. *Composita subtilita* (Brachiopoda) in the Wreford Megacyclothem (Lower Permian) in Nebraska, Kansas, and northern Oklahoma. The University of Kansas Paleontological Contributions, Paper 81:1–19.
- Lutz-Garihan, A. B., & R. J. Cuffey. 1979. Stratigraphy of the Lower Permian Wreford Megacyclothem in southernmost Kansas and northern Oklahoma. *Kansas Geological Survey Bulletin* 216:1–19.
- Miller, K. B., & R. R. West. 1993. Reevaluation of Wolfcampian cyclothems in northeastern Kansas: significance of subaerial exposure and flooding surfaces. *Kansas Geological Survey Bulletin* 235:1–26.
- Newton, G. B. 1971. Rhabdomesid bryozoans of the Wreford Megacyclothem (Wolfcampian, Permian) of Nebraska, Kansas, and Oklahoma. The University of Kansas Paleontological Contributions, Article 56 (Bryozoa):1–71.
- Pachut, J. F., & R. L. Anstey. 1979. A developmental explanation of stability-diversity-variation hypotheses: morphogenetic regulation in Ordovician bryozoan colonies. *Paleobiology* 5:168–187.
- Pachut, J. F., R. L. Anstey, & R. J. Cuffey. 1991. The concepts of astogeny and ontogeny in stenolaemate bryozoans, and their illustration in colonies of *Tabulipora carbonaria* from the Lower Permian of Kansas. *Journal of Paleontology* 65:213–233.
- Pachut, J. F., & R. J. Cuffey. 1991. Clinal variation, intraspecific heterochrony, and microevolution in the Permian bryozoan *Tabulipora carbonaria*. *Lethaia* 24:165–185.
- . 1995. Depth-related associations of cryptic-habitat bryozoans from the leeward fringing reefs of Bonaire, Netherlands Antilles. *Palaios* 10:254–267.
- Patzkowsky, M. E. 1995. Gradient analysis of Middle Ordovician brachiopod biofacies: biostratigraphic, biogeographic, and macroevolutionary implications. *Palaios* 10:154–179.
- Pielou, E. C. 1974. *Population and Community Ecology*. Gordon and Breach Science Publishers. New York. 424 p.
- Prezbindowski, D. R., & R. L. Anstey. 1978. A Fourier-numerical study of a bryozoan fauna from the Threeforks Formation (Late Devonian) of Montana. *Journal of Paleontology* 52:353–369.
- Roughgarden, J. 1989. The structure and assembly of communities. In J. Roughgarden, R. M. May, and S. A. Levin, eds., *Perspectives in Ecological Theory*. Princeton University Press. Princeton. New Jersey. p. 203–226.
- Sanders, H. L., & R. R. Hessler. 1969. Ecology of the deep-sea



- benthos. *Science* 163:1419–1424.
- Simonsen, A. H., & R. J. Cuffey. 1977. Fenestrate and pinnate bryozoans in the Wreford Megacyclothem (Lower Permian; Kansas, Oklahoma, and Nebraska). *Geological Society of America, Abstracts with Programs* 9:652.
- . 1980. Fenestrate, pinnate, and ctenostome bryozoans and associated barnacle borings in the Wreford Megacyclothem (Lower Permian) of Kansas, Oklahoma, and Nebraska. *The University of Kansas Paleontological Contributions, Paper* 101:1–38.
- Springer, D. A., & R. K. Bambach. 1985. Gradient versus cluster analysis of fossil assemblages: a comparison from the Ordovician of southwestern Virginia. *Lethaia* 18:181–198.
- Warne, J. E., A. A. Ekdale, S. F. Ekdale, & C. H. Peterson. 1976. Raw material of the fossil record. In R. W. Scott, & R. R. West, eds., *Structure and Classification of Paleocommunities*. Dowden, Hutchinson and Ross, Inc. Stroudsburg, Pennsylvania. p. 143–169.
- Warner, D. J., & R. J. Cuffey. 1973. Fistuliporacean bryozoans of the Wreford Megacyclothem (Lower Permian) of Kansas. *The University of Kansas Paleontological Contributions, Paper* 65:1–24.
- Wilkinson, L. 1990. SYSTAT: The System for Statistics. SYSTAT, Inc. Evanston, Illinois.