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Macrobenthos of the subtidal Wadden Sea: revisited after 55 years

W. Riesen & K. Reise

II. Zoologisches Institut der Universität Göttingen; D-3400 Göttingen,
Federal Republic of Germany
and
Biologische Anstalt Helgoland (Litoralstation); D-2282 List/Sylt,
Federal Republic of Germany

ABSTRACT: During the years 1923-1926 Hagmeier & Kändler (1927) sampled the macrofauna of subtidal shallows and channels of the Wadden Sea close to the Island of Sylt (German Bight, North Sea). Reinvestigating this study area in 1980, a substantially altered faunal composition was recorded. An approach is made to quantify the comparison in terms of abundance, species richness and diversity of invertebrate taxa. Human interference is assumed to be responsible for the major changes. Natural oyster beds have been overexploited and the local population of Ostrea edulis has been driven to extinction. Subsequently, mussels (Mytilus edulis) spread in the entire region, promoted by shell fishery. Particularly barnacles and many polychaetes took advantage of the expansion of mussel banks which is substantiated by correlation analysis. Reefs of the colonial polychaete Sabellaria spinulosa stood in the way of shrimp trawling and became destroyed together with the associated fauna. A subtidal Zostera marina bed was wiped out in 1934 by a natural epidemic disease but never succeeded in reestablishing itself. The associated fauna disappeared. Large epibenthic predators and scavengers (crabs, snails and starfish) survived all these changes. The total number of species remained approximately at the same level but molluscs experienced losses and polychaetes diversified. Overall abundance increased with a disproportionately large share of a few species (Mytilus edulis, Balanus crenatus, Cerastoderma edule, Scoloplos armiger). The subtidal fauna of the Wadden Sea proved to be vulnerable to human disturbance; thus, the present community can no longer be viewed as the outcome of entirely natural processes.

INTRODUCTION

An understanding of the benthic ecology of the Wadden Sea necessitates an exploration of its past. Do we have a resilient community in natural balance, a continuously changing assemblage in a hazardous environment, or a pile of organisms disturbed because of repeated human interference? Here we make an attempt to present a comparison of the two quantitative surveys, 55 years apart, on the macrofauna of subtidal shallows and channels to the east of the Island of Sylt. Inspired by C. G. J. Petersen and urged by the declining oyster fishery, Hagmeier started in 1923 a sampling program with dredge and Petersen grab in the North Frisian Wadden Sea (Hagmeier, 1925; Hagmeier & Kändler, 1927). His sampling scheme and data presentation does not allow a sound, statistically-backed comparison with the present, however, the magnitude of the differences suggests considerable change in species composition and abundance of macrofauna.

AREA AND METHODS

The area of investigation is part of the North Frisian Wadden Sea, east of the Island of Sylt. The subtidal benthic fauna is subject to holomarin conditions, soft sediments, and strong tidal currents while the island provides some shelter from the rough North Sea.

Grab samples were taken at 9 sites down the slope of the channel "Lister Ley" and on 2 subtidal shallows once occupied by oyster beds (Fig. 1). Sites on the transect refer to those described by Hagmeier & Kändler (1927, p. 32) as I, II, III, IV, VII, VIII, IX, X, XII with a depth range from 0.5 m to 18.5 m. Sites in the shallows (depth 3 m) correspond to Nos. 36 and 42 on p. 41, hereafter called "Ellenbogen" and "Huntje".

We used a 0.1 m² V^an Veen grab. Quantity and quality of the sediment was recorded. Further treatment of samples was done in the lab. The samples were sieved through a 1 mm meshed screen in a huge tub. The sieve content was washed into white plastic dishes to pick up the macrofauna. Numbers of more than 100 *Balanus crenatus* individuals were estimated, colonies of Porifera, Hydrozoa and Bryozoa were noted as present.

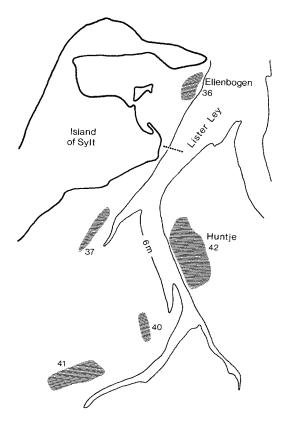


Fig. 1. Wadden Sea of northern Sylt (German Bight), the 6m-isoline marks the channel system, Former oyster beds are shaded, numbers of stations are from Hagmeier & Kändler (1927). The dotted line at the western slope of Lister Ley shows the transect with 9 sampling stations investigated in 1923–1926 and in 1980

In addition to grab samples, dredge hauls were conducted in 5 subtidal shallows including Ellenbogen and Huntje (Fig. 1). The sites correspond to Hagmeier's & Kändler's Nos. 36, 37, 40, 41 and 42 (p. 44). For the purpose of comparison, the dredge of both authors was rebuilt, with a blade length of 1 m and a mesh size of 6 cm. Dredge samples were treated qualitatively on board.

RESULTS

Habitat structure

In 1980, the sites revisited showed partially altered habitat conditions. In the subtidal shallows, where there were once oyster beds on a sandy bottom, half of the samples contained silty sediment due to silt-assembling mussel banks. Along the transect down the slope of Lister Ley, the same phenomenon occurred at some upper sites, while the sites in the steep part showed signs of erosion. Stones, shell gravel and coarse grained sand occurred. A *Zostera marina* bed, once in the shallow upper part of the transect, was absent in 1980 and so were reefs of the tube-building polychaete *Sabellaria spinulosa* on the lower slope (Fig. 2).

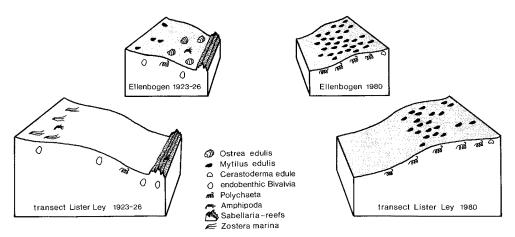


Fig. 2. Cartoon of benthic assemblages in Lister Ley, a comparison between 1923–1926 and 1980. Block-diagrams above correspond to site 36 and the two below to the dotted transect in Fig. 1

Overall shifts in abundance and species composition

Most striking is an increase in individuals per area (Fig. 3a). Only two of the sites investigated – both occupied by *Sabellaria* reefs in 1923–26 – show a decline. *Balanus crenatus* and *Mytilus edulis* contributed most to high abundances in 1980. However, even without these two, abundance was generally higher in 1980 because of more polychaetes.

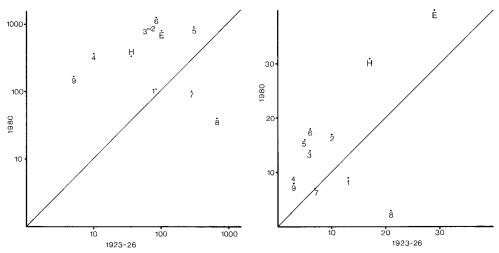


Fig. 3. Comparison between 1923–1926 and 1980 in macrofaunal abundance (logarithmic scale) (left) and in species density (right). Nos 1–9: transect Lister Ley, E=Ellenbogen, H=Huntje. The line at 45° marks equality between surveys; points above represent higher values in 1980, points below higher values in 1923–1926. All ratios refer to areas of 0.1 m², except in (b) where E=0.9 m² and H=0.4 m²

Ranking of the 8 most abundant species reveals notable changes in dominance since the 1920s (Table 1). Only two species remained in the top ranks until 1980. *B. crenatus* was able to take advantage of the increased offer of secondary, hard substrate provided by mussel shells. All amphipods abundant during the years 1923–26 were associated with *Sabellaria* reefs and the *Zostera* bed. Most of the sites investigated show higher species density in 1980, mainly caused by polychaetes (Fig. 3b). The only sites with a reverse trend are those of former *Sabellaria* reefs and *Zostera* beds, where a formerly rich epifauna has gone.

While abundance and species density increased at most sites, the total number of species seems to have remained stable. Hagmeier & Kändler (1927) recorded 54 species gathered from 22 grab samples and 5 dredge hauls. We found 89 species in 123 grabs

Table 1. Most abundant macrobenthic species in the subtidal Wadden Sea at Sylt during the period 1923–1926 and in 1980. Three sites, Ellenbogen, Huntje and the transect have been pooled in equal proportions and individuals \cdot m⁻² (N) are presented

1923–1926	N	1980	N
Balanus crenatus	520	Balanus crenatus	2300
Sabellaria spinulosa	250	Mytilus edulis	1988
Corophium bonelli	105	Scoloplos armiger	226
Scoloplos armiger	29	Cerastoderma edule	164
Abra alba	17	Heteromastus filiformis	116
Eumida sanguinea	17	Nereis virens	62
Ericthonius difformis	14	Pholoe minuta	58
Caprella linearis	10	Capitella capitata	30

and 50 dredge hauls. Converted to their sampling effort, we obtained 49 species. This is close to the 54 recorded by Hagmeier & Kändler. Thus, the increase in species density is caused by a more even distribution of the species in 1980.

Transect along the slope of Lister Ley

Since the 1920s, the benthos on the slope of Lister Ley has changed considerably (Fig. 2, Table 2). In the *Zostera* bed on the upper part of the slope, Hagmeier & Kändler found dense populations of amphipods, snails and young starfish. Endobenthic bivalves settled all along the slope. *Sabellaria* reefs on the lower slope were populated by polychaetes, amphipods and cnidarians. In 1980, *Mytilus edulis* occupied the upper half of the slope, while *Balanus crenatus* and many polychaetes were abundant along the entire range. Dense patches of *Cerastoderma edule* were observed at the three uppermost sites.

While Mollusca and Crustacea have decreased in species number and diversity since the 1920s, Polychaeta have increased (Table 3). Each site on its own shows little similarity in 1980 to the faunal composition recorded by Hagmeier & Kändler (Fig. 4). Few species common to both investigations were found. Sites with a high number of species in 1923–26 (sites 1, 2, 7, 8) were the grass bed and the reefs. Sites with a high number of species in 1980 are dominated by *M. edulis* (sites 2 and 6). In Figure 4, dendrograms at sites 1, 7 and 8 separate the sample obtained by Hagmeier & Kändler from all recent ones. At the other sites, some degree of similarity is brought about mainly by spatial heterogeneity. Rather than having species in common, samples within subgroups lack many species otherwise present.

Past oyster beds - mussel banks today

The constancy of large epibenthic species was estimated from dredge hauls (Table 4). Of the 5 species abundant in 1923–26 but absent in 1980, Lepidochitona cinerea and Psammechinus miliaris are known to be still present in the area, while the other three are probably gone. A definite increase is substantiated for Mytilus edulis and Conopeum reticulum. The latter was mentioned by Hagmeier & Kändler for some areas south of Sylt only. Large sized predators and scavengers do not seem to be affected in their constancy by the changed composition in the benthic community.

Most of the former oyster beds are covered by clusters of mussels today. As a consequence, grab samples taken during 1924–26 and those in 1980 contain quite different sets of species or at least different abundances where the same species did persist (Table 5). Ostrea edulis has vanished from the area. It was abundant prior to 1874 when yields of the oyster fishery dropped dramatically (Möbius, 1893; Hagmeier & Kändler, 1927), remaining at a low level over a period of 50 years; and since 1925, oyster fishing became so unprofitable that it was finally abandoned (Reise, 1980). To our knowledge, no live specimen has been dredged during the last 30 years.

While mussels are a permanent component of the subtidal benthos, the population of *Cerastoderma edule* at Ellenbogen did not persist. Many were already tied together by byssus threads of *M. edulis* which settled on top of the cockles. One year later, in 1981, *C. edule* was rare at the site but empty shells were abundant within clusters of mussels.

Table 2. Individuals · 0.1 m⁻² of macrofauna at 9 sites along a transect down the slope of Lister Ley during the period 1923–26 and in 1980. As

Hagn	ıeier &	Hagmeier & Kändler (1 3 sampl	ndler (1 3 sampl		took o appro	ne sa ximat	mple te the	at eacl mean	927) took one sample at each site and we took 7 samples, we included in the 1980-data all species occurring in at least es to approximate the mean number of species in one sample. \times = present but individuals not counted	led in presen	the 19 t but	980-de indiv	ita all	speci s not c	es occ	ourring d	y in at	leas
			19	1923–1926	326									1980				
18,	5 14.0	18.5 14.0 12.0	6.5	4.5	3.0	1.5	1.0	0.5	Depth (m) below low tide level	0.5	1.0	1.5	3.0	4.5	6.5 1	12.0 1	12.0 14.0 18.5	3.5
									Mollusca									
	4		4	က	8		က	H	Abra alba (S. Wood)									
	1						~		Angulus tenuis (Da Costa)									
	15								Venerupis pullastra (Montagu)									
	2							8	Mya spec. juv.									
	5								Petricola pholadiformis Lamarck									
						7			Spisula subtruncata (Da Costa)									
						1		က	Retusa truncatula (Bruguière)									
							₩.	4	Nassarius reticulatus (L.)									
						7	က		Littorina littorea (L.)									
								 i	Hydrobia ulvae (Pennant)									
	7		1				_		Macoma balthica (L.)			7			1			
							25		Cerastoderma edule (L.)	42		309						
	19			\vdash				9	Mytilus edulis L.	13	395	134	148	501	540			
									Polychaeta									
	317	300	-						Sabellaria spinulosa Leuckart									
	₩	2	က	7		+	2		Nephtys caeca (F.) + hombergii S.	က		7				₩.	7	2
m	8	က	က	4		4	4		Scoloplos armiger (O. F. Müller)	15	9	11	Ţ	က	7	15	4	8
									Nereis virens Sars	5	5	E	12	9	10	9		7
									Eulalia viridis (L.)		-			3	9			
									Anaitides mucosa (Oersted)	₩	က	5						
									Kefersteinia cirrata (Keferst.)					7				
~									Lepidonotus squamatus (L.)					1	က			
									Gattyana cirrosa (Pallas)		5							
									Harmothoe imbricata (L.)		3	4		7	6			

		Macrob	enthos of the Wadden Sea
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Harmothoe impar (Johnston) Pholoe minuta (Fabricius) Capitella capitata (Fabricius) Heteromastus filiformis (Clap.)	Crustacea Corophium bonelli Milne Edwards Cheirocratus sundevallii (Rath.) Phoxocephalus holboelli (Krøyer) Parapleustes bicuspis (Krøyer) Bathyporeia guilliamsoniana (Ba.) Caprella linearis (L.) Ericthonius difformis Milne Edw. Apherusa bispinosa (Sp. Bate) Dexamine spinosa (Montagu)	fuotea batunca (ranas) Eupagurus bernhardus (L.) Carcinus maenas (L.) Balanus crenatus Bruguière Gammarus locusta (L.)	Other species Sagartiogeton viduata (O. F. M.) Metridium senile (L.) Asterias rubens L. Ophiura albida Forbes Amphipholis squamata (D. Chiaje) Hydractinia echinata (Fleming) Laomedea longissima (Pallas) Alcyonidium polyoum (Hässall) Conopeum reticulum (L.) Halichondria panicea (Pallas) Taelia felina (L.)
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Table 3. Numerical characteristics of major taxa occurring along the transect down the slope of Lister Ley during the period 1923–26 and in 1980. All 9 stations have been combined to 0.9 m² samples, and the data given refer to this sample size. $H' = -\sum p_i \cdot \ln p_{ii}$ $D = (\sum p_i^2)^{-1}$; $F = (D-1)/(\exp H' - 1)$

Major taxa		usca	•	haeta		tacea
Survey	1923–26	1980	1923–26	1980	1923–26	1980
Number of individuals	101	2110	653	294	826	2028
Number of species	12	3	3	13	14	4
Shannon-Weaver diversity H'	2.05	0.48	0.24	2.13	1.10	0.04
Simpson's diversity D	6.04	1.42	1.11	6.82	2.33	1.01
Hill's evenness F	0.74	0.68	0.41	0.78	0.66	0.27

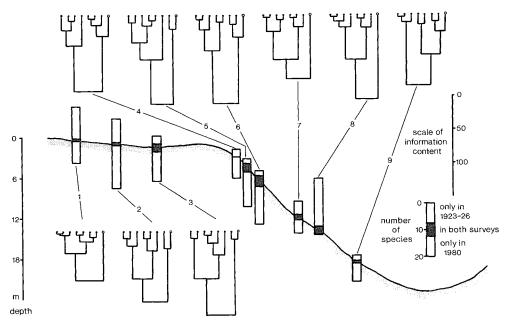


Fig. 4. Similarity of macrofauna between 1923–1926 and 1980 for 9 sites at the slope of Lister Ley. Dendrograms depict similarity between our sets of 7 samples (·) and the single ones taken by Hagmeier & Kändler (o), calculated from presence-absence data according to Williams et al. (1966). The scale is given to the right. Vertical bars along the transect present numbers of species in both surveys and those occurring only once (see explanations to the right). For comparison with the single samples of the first survey, we included all species occurring in at least 3 out of 7 samples

With regard to polychaetes, only few species were more abundant in the early survey. In 1980, numbers of small as well as large sized species (i. e. Polynoidae, *Nereis virens*) were considerably higher. Many of them being abundant on the adjacent tidal flats too. In Crustacea, increased numbers of barnacles represent the major difference between the surveys. In large-sized decapods no changes are apparent. High numbers of

Table 4. Frequencies (%) of large macrobenthos and epigrowth caught with the dredge during the period 1923–1926 (5 "representative" hauls) and in 1980 (50 hauls). Because of ambiguities in Hagmeier & Kändler's data, *Hydractinia echinata, Crangon crangon, Balanus crenatus* and fishes are excluded from the comparisons made

Species	1923–26	1980
Sertularia cupressina L.	100	0
Sagartiogeton viduata (O. F. Müller)	100	0
Ostrea edulis L.	100	0
Lepidochitona cinerea (L.)	100	0
Psammechinus miliaris (Gmelin)	100	0
Hyas araneus (L.)	100	38
Macropipus holsatus (Fabricius)	80	46
Lepidonotus squamatus (L.)	100	74
Alcyonidium 2 spp.	100	74
Taelia felina (L.)	80	62
Asterias rubens L.	100	82
Buccinum undatum L.	80	72
Eupagurus bernhardus (L.)	100	90
Carcinus maenas (L.)	80	94
Mytilus edulis L.	40	100
Conopeum reticulum (L.)	0	100

Table 5. Individuals \cdot m⁻² of macrofauna at Ellenbogen (E) and Huntje (H) during 1924–1926 (n = 9 + 4 grabs) and in 1980 (n = 30 + 30 grabs). In the 1980-data all species are included which occurred in at least 4 or 3 samples to approximate the mean number of species in 9 or 4 samples respectively. Hagmeier & Kändler (1927) did not differentiate between species of *Nephtys* thus we lumped them together too. x = present but individuals not counted

Survey	1924	-26	1	.980
Sites	Е	Н	Е	Н
Mollusca				
Lepidochitona cinerea (L.)	3	3		
Abra alba (S. Wood)	1	25		
Spisula subtruncata (Da Costa)		3		
Ostrea edulis L.	1			
Petricola pholadiformis Lamarck	1			
Venerupis pullastra (Montagu)	0.2			
Mya spec. juv. (truncata?)	1			
Buccinum undatum L.	1	3	1	
Cerastoderma edule (L.)			117	
Macoma balthica (L.)			14	8
Mytilus edulis L.			2804	1425
Polychaeta				
Gattyana cirrosa (Pallas)	15	3		
Sabellaria spinulosa Leuckart		5		
Ampharete acutifrons (Grube)		5		1
Eumida sanguinea (Oersted)	50	3	2	2
Lanice conchilega (Pallas)	31		4	

Table 5 (continued)

Survey	192	4–26	1	.980
Sites	E	Н	Е	Н
Pectinaria koreni Malmgren	3		2	
Nephtys caeca + N. hombergii	J	17	9	21
Scoloplos armiger (O. F. Müller)	32	27	118	483
Lepidonotus squamatus (L.)	1	_,	24	10
Harmothoe imbricata (L.)	_		39	11
Harmothoe impar (Johnston)			24	13
Pholoe minuta (Fabricius)			90	78
Heteromastus filiformis (Clap.)			198	110
Capitella capitata (Fabricius)			56	6
Nereis virens Sars			103	34
Eulalia viridis (L.)			26	4
Eteone longa (Fabricius)			3	4
Anaitides mucosa (Oersted)			3	3
Kefersteinia cirrata (Keferst.)			6	
Malacoceros tetracerus (Schmarda)			3	11
Malacoceros fuliginosus (Clap.)			2	
Tharyx marioni (Saint-Joseph)			2	3
Spio filicornis (O. F. Müller)			5	
Pygospio elegans (Claparède)			7	3
Crustacea				
Cheirocratus sundevallii (Rath.)	14	12		
Caprella linearis (L.)		7		
Apherusa bispinosa (Sp. Bate)	1			
Dexamine spinosa (Montagu)	1			
Bodotria scorpioides (Montagu)	1			
Macropipus holsatus (Fabricius)	18		1	
Eupagurus bernhardus (L.)	15	3	5	2
Crangon crangon (L.)	1	7	1	7
Carcinus maenas (L.)	11		48	9
Balanus crenatus Bruguière	800	260	4000	1000
Hyas araneus (L.)			7	
Corophium bonelli Milne Edwards			4	
Gammarus locusta (L.)				6
Bathyporeia sarsi Watkin				24
Other species				
Sagartiogeton viduata (O. F. Müller)	6			
Metridium senile (L.)	1			
Psammechinus miliaris (Gmelin)	6			
Asterias rubens L.	1			
Ophiura albida Forbes	1			
Sertularia cupressina L.		×		
Hydractinia echinata (Fleming)	×	×	×	×
Taelia felina (L.)			5	5
Laomedea longissima (Pallas)			×	
Laomedea flexuosa Alder			×	
Alcyonidium polyoum (Hässall)			×	×
Conopeum reticulum (L.)			×	×
Tubificoides benedeni (Udekem)			3	1

these species include juveniles in both sets of samples. The herbivor Hyas araneus was absent from the grab samples during 1924–26 but occurred in almost all dredges at that time.

Because of all these differences in species composition and species abundance, numerical parameters of the benthic assemblage also changed from the early survey to the recent one (Table 6). In all major taxa reported the total number of individuals increased, and in polychaetes species richness is much higher in 1980. Diversity and evenness decreased in Mollusca and Crustacea, while in Polychaeta there was an increase in diversity at Ellenbogen.

Table 6. Numerical characteristics of major taxa occurring at Ellenbogen (E) and Huntje (H) during the period 1924–1926 and in 1980. Number of species refer to 9 grabs at E and 4 grabs at H. See also Table 3

Major taxa Survey		Mol	lusca 5 1980	Polyc 1924–26	haeta 1980	Crus 1924–26	tacea 5 1980
Individuals · m ⁻²	E	8	2936	133	726	862	4066
	H	34	1433	60	628	289	1048
Number of species	E	6	4	7	22	9	7
	H	4	2	6	18	5	6
Shannon-Weaver diversity H'	E	1.67	0.20	1.46	2.23	0.37	0.10
	H	0.87	0.03	1.43	1.46	0.45	0.25
Simpson's diversity D	E	4.57	1.09	3.75	6.71	1.16	1.03
	H	1.77	1.01	3.31	2.45	1.23	1.10
Hill's evenness F	E	0.83	0.41	0.83	0.69	0.36	0.29
	H	0.56	0.37	0.73	0.44	0.41	0.34

Species associated with mussels

Mytilus edulis gave the past oyster beds their new structure. Clusters of mussels accumulate silt, provide shelter for other species and surfaces for epigrowth. At the sites Ellenbogen and Huntje, 60 grab samples were taken in 1980 containing variable amounts of mussels. Out of the 12 most frequently sampled species, 7 are positively correlated with the number of mussels present (Table 7). B. crenatus is an abundant epigrowth on mussels, while polychaetes will profit from shelter and accumulated silt. There is a significant positive correlation of the total number of polychaetes, the number of polychaete species and their diversity with mussel density. Thus, the considerable increase in polychaetes since the survey of 1923–26 may be a consequence of the spread in M. edulis. However, not all polychaete species abundant in 1980 are correlated with mussel density and the genus Nephtys shows a negative correlation.

DISCUSSION

On comparing the results of the two surveys, a trend towards a massive long-term increase of the mussel banks and associated species, particularly polychaetes and

Table 7. Abundance of macrofaunal species at Ellenbogen and Huntje as functions of mussel abundance (Mytilus edulis). Linear regressions calculated from n = 60 grab samples taken in July 1980. Individuals \cdot 0.1 m⁻² = a + b (Mytilus \cdot 0.1 m⁻²). a, b = coefficients, * indicates significance (P < 0.05) of the correlation coefficient r. All species present in at least 20 samples are listed

Species	a	b	r
Lepidonotus squamatus	0.033	+ 0.008	+ 0.732*
Eulalia viridis	0.190	+ 0.006	+ 0.714*
Nereis virens	2.426	+ 0.021	+ 0.705*
Balanus crenatus	62.967	+ 0.835	+ 0.702*
Harmothoe imbricata	0.682	+ 0.009	+ 0.686*
Pholoe minuta	4.237	+ 0.020	+ 0.541*
Heteromastus filiformis	7.004	+ 0.040	+ 0.441*
Carcinus maenas	1.743	+ 0.005	+ 0.245
Capitella capitata	1.681	+ 0.004	+ 0.239
Harmothoe impar	1.318	+ 0.003	+ 0.198
Scoloplos armiger	22.347	-0.006	-0.118
Nephtys 2 spp.	1.976	-0.002	- 0.303*

barnacles becomes evident. Species richness in molluscs, as well as the abundance of some amphipods associated with seagrass and *Sabellaria* reefs decreased. In contrast, large epibenthic predators and scavengers could hold their position.

To what extent may the claimed long-term differences result from artifacts or unequal sampling efficiency? For dredging we used exactly the same device as Hagmeier & Kändler (1927) did, trawling for 10 min and about 1000 m. Although Hagmeier & Kändler present numbers of individuals encountered in a trawl assumed to be "typical" for a given site, we confined our comparison to presence-absence data. Thus, efficiency in catching large epibenthic invertebrates is not considered to be different in the two surveys.

Instead of using a Petersen grab as Hagmeier & Kändler did, we had to use a Van Veen grab of the same size. Particularly on sandy bottoms the latter collects more sediment and the biting profiles are slightly different (Gallardo, 1965; Ziegelmeier, 1968). However, currents, waves, and experience in handling the grab are variables of equal importance. On average, 5 l of sediment were obtained with both grabs, and where a sample contained less it was rejected. The mesh of sieves was equal but the sorting procedures differed. Hagmeier & Kändler picked up the organisms directly out of the sieves. Because this takes some time, slender worms have a good chance to escape. In 1980 we did the sorting in dishes at the lab and thus probably worked with higher efficiency regarding some polychaetes such as Spionidae, Capitellidae and Orbiniidae. On the other hand, differences in sorting efficiency cannot account for an increase in such large sized polychaetes as Lepidonotus squamatus, Harmothoe spp., Eulalia viridis and Nereis virens. While improved sorting affects our estimates on polychaete abundance, it will hardly alter the number of species encountered, as we do not report on species smaller than those listed by Hagmeier & Kändler.

Severe winters are known to have lasting effects on benthic macrofauna in the North Sea (Smidt, 1944; Ziegelmeier, 1964, 1970; Dörjes, 1980; Reichert & Dörjes, 1980; Buhr, 1981). Ziegelmeier (1978) noted a general shift from dominance of molluscs to domi-

nance of polychaetes in the years following a very cold winter. As such a shift is claimed to be one of the major long-term trends in the present study, the sequence of severe winters throughout the period of concern is worth recording: 1923/24, 1928/29, 1939/40, 1940/41, 1941/42, 1946/47, 1955/56, 1962/63, 1969/70 and 1978/79. There is no trend in frequency or amplitude, and both surveys are conducted subsequent to a particularly cold winter. Thus, winter effects cannot account for the differences observed but the possibility of population fluctuations triggered off by other environmental factors in the course of decades cannot be ruled out. Unfortunately, we have no reliable long-term data on physical factors in the study area. More frequent surveys in the future are needed to test our conclusions.

Are all the proposed long-term changes ultimately caused by human interference? Oyster beds, once common in the shallow parts of the North Sea are absent from the German Bight today. There has been some debate as to the actual cause of the decline (Möbius, 1877; Hagmeier & Kändler, 1927) but more recent accounts on the subject leave little doubt that overexploitation by oyster fishery is responsible for the extinction of *Ostrea edulis* in the area considered here (Linke, 1937; Korringa, 1980; Reise, 1980, 1982).

In the Wadden Sea, oyster beds were intermediate in the vertical distribution between mussel banks above and *Sabellaria* reefs below, and both tended to invade the oyster beds (Hagmeier & Kändler, 1927; Linke, 1937). At Sylt, the entire range of the former oyster beds is now occupied by mussel banks. It is not known whether this occurred with or without the aid of the mussel fishery. In any case, since the 1950s subtidal banks have been stocked with spat, dredged from tidal flats. Today, there are more extensive banks even in the intertidal zone at Sylt than were recorded in 1934 by Wohlenberg (1937). The intertidal banks are of no interest to sea fisheries. There may very well be a natural component in the spread of *Mytilus edulis* but fisheries certainly promoted this trend.

Sabellaria reefs are completely lost to the area. Local fishermen claim to have ground them with heavy gear because the reefs ripped apart the nets when fishing for shrimp. There is no other evidence. The loss of an extensive subtidal seagrass bed of the large growth form of Zostera marina L. during the general epidemic seagrass disease (1933–34) is regarded as a natural event (Wohlenberg, 1935; Den Hartog, 1970). As in the Dutch western Wadden Sea, Z. marina failed to reestablish itself in the subtidal zone of our study area. Van den Hoek et al. (1979) suppose that increased turbidity following the closure of the Zuiderzee in 1932 prevented recolonization in the Dutch western Wadden Sea. For the Sylt area, we have no data on turbidity but dams connecting the Island of Sylt (since 1927) and the Island of Rømø (since 1948) with the mainland are likely to have similar effects as the closure of the Zuiderzee. Taken together, there is little doubt that all major habitat shifts were caused, promoted or made permanent by human interference.

In recent years, a number of species immigrated into the tidal flats of the Wadden Sea (reviewed by Michaelis, 1978, 1981) without any apparent environmental cause, i.e. at Sylt *Tharyx marioni* (Saint-Joseph), *Spio filicornis* (O. F. Müller) and *Chelon lambrosus* (Risso). Recent gains and losses of species in the Wadden Sea give no evidence of any directional change in climate.

Long-term comparisons similar to the present study have been conducted by Ceder-

wall & Elmgren (1980) and Persson (1981) in the Baltic Sea. They noted a considerable increase in the biomass of benthic macrofauna which the former attribute to recent eutrophication while the latter assumes decreased predation pressure following exploitation of flatfish. Also at Sylt, an increase in biomass must be assumed as a consequence of increased abundance in such large species as *Mytilus edulis* and *Nereis virens*. In this area, however, we assume the spread of mussel banks promoted by fisheries to be the principle cause of increased biomass. Predators on bivalves (Carcinus maenas, Asterias rubens) do not seem to be less abundant today than 55 years ago.

It is well known that most terrestrial communities have been severely altered by man's impact. Here we point out that the same applies to the coastal sea. The present assemblage of organisms is not the pure outcome of a natural chain of events. Thus, there is no reason to expect a stable balance between trophic levels, brought about by coevolution. The present state is apt to continuous change not only because of man's continued interference but because the relative abundances are to a high degree artificial.

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