Growth and population structure of *Paracentrotus lividus* (Echinodermata: Echinoidea) in two contrasting habitats

X. Turon, G. Giribet, S. López, C. Palacín

Department of Animal Biology (Invertebrates), Faculty of Biology, University of Barcelona, Avinguda Diagonal, 645, E-08028 Barcelona, Spain

**ABSTRACT.** Two populations of *Paracentrotus lividus* (Lamarck) (Echinodermata: Echinoidea) from the Northwestern Mediterranean were compared to ascertain the plasticity and adaptive value of certain biological parameters. These populations were found in habitats which differed in terms of stability of environmental parameters and availability of food. The density was higher and more fluctuating in the unstable community. Sigmoidal growth functions were fitted from plate ring analysis, and reflected a higher growth rate in the stable community, resulting in higher mean diameters at equal age. The maximal growth rate was attained in the unstable community at an earlier age than in the stable one. Size-frequency analyses reflected drastic month-to-month changes in population structure in the unstable community, in which the smaller size classes were almost absent. In contrast, at the stable site the distribution was much more uniform through time, and featured a polymodal shape consistent with the development of several annual cohorts. The smaller size classes were the most abundant. It is suggested that different dynamics underlie both populations, the one at the unstable site being driven primarily by episodic storms, which cause high mortalities but carry new individuals to the site. The stable community relies instead on an annual settlement and features a lower and more predictable mortality which allows for the development of a well-structured population.

**KEY WORDS:** Population dynamics · Growth · Gompertz function Echinoidea · *Paracentrotus*

**INTRODUCTION**

The sea urchin *Paracentrotus lividus* (Lamarck) (Echinodermata: Echinoidea) is an Atlanto-Mediterranean form found in great numbers in the littoral zone, especially in the sublittoral level down to 20 m. Many aspects of its biology and population parameters are poorly known, although it is an edible species of commercial interest.

In this study, the main life-history parameters of 2 populations of this species were investigated. These 2 populations were found in ecological situations which differed in terms of stability of environmental parameters and food availability. The aim of this study was to describe the biology and population dynamics of this species and to ascertain the plasticity and adaptive value of different biological parameters. In another paper, Lozano et al. (1995, this issue) reported the main results concerning reproductive and feeding cycles, as well as timing and intensity of recruitment, of the populations studied. The present paper focuses on the density fluctuations, growth and structure of the populations in these communities. Both populations seemed to be well established in the sense that a fair number of sea urchins at both sites have been observed year round, not only during the study period, but also in the many years of marine biology research in diverse fields conducted at these localities. Our hypothesis in this paper was that these dense, apparently well-established populations which belong to such diverse habitats, must develop following differing processes and, therefore, feature different structure and dynamics. Knowledge of these dynamics is, on the other hand, a necessary prerequisite for a management of this resource. It has long been recognized that echinoid growth rates are highly sensitive to factors such as food availability and hydrodynamism (Ebert 1968, Himmelman 1986). Fewer data, however, are available on habitat influences in population structure and other biological parameters (Himmelman 1986, Byrne 1990, King et al. 1994).
Several techniques have been employed for the study of growth and population structure in echinoids: the study of size-frequency distributions, the analysis of growth rings in the test, mark-recapture techniques (usually with tetracycline labelling, first used by Kobayashi & Taki 1969), and monitoring of animals in enclosures. Each has its own limitations, and a combination of several of them is more likely to produce reliable results. In this study, we have combined data from growth marks in the plates with results from the changes in size-frequency distributions through time to obtain data on the growth and dynamics of the populations analyzed.

**MATERIAL AND METHODS**

**Sampling sites.** The study was performed on specimens from 2 localities, Tossa de Mar (41° 43.2' N, 2° 56.4' E) and Cubelles (41° 12.0' N, 1° 40.8' E), both on the northeast coast of Spain (Northwestern Mediterranean). A map of the area and descriptions of each zone are presented in Lozano et al. (1995). Essentially, the site at Tossa de Mar is representative of a well-established, stable (with respect to most physical parameters) community in rocky vertical walls between 3 and 10 m in depth, with high coverage of diverse fleshy algal species. This will be referred to hereafter as the stable community. The site at Cubelles was an open, shallow (0.5 m in depth) beach with small-to-medium sized boulders on which a poorly developed community of small coralline algae was found. This site is referred to as the unstable community.

Unless otherwise stated, the samplings for the different parameters were carried out approximately every 2 wk from June 1992 to July 1993 at Tossa, and from July 1992 to August 1993 at Cubelles. Results are presented as monthly means to reduce stochastic variation.

**Density fluctuations.** Fixed transects were censused at both localities to assess density fluctuations of the populations. At Tossa de Mar, 3 transects 25 m long × 1 m wide were placed horizontally at a depth of 3, 6 and 9 m in a nearly vertical wall. At Cubelles, a 25 × 1 m transect was deployed, perpendicular to the coast line, running from 0.3 to 0.7 m in depth. The specimens present in these transects were counted at approximately the same hour (13:00 h) during daylight on each sampling date.

**Growth ring measurements.** Samples of 20 individuals of *Paracentrotus lividus* of all sizes were taken every 2 wk at both localities. The samples were fixed and preserved in formalin. Growth rings were studied in 509 individuals from Tossa de Mar and 537 from Cubelles (these were the same specimens as those used in Lozano et al. 1995) for the study of biological cycles. The maximum diameter of the individuals was measured to the nearest 0.1 mm in the laboratory. The body was then oven dried at 120°C for 48 h and weighed. The test was cleaned of spines and an interambulacral series of plates was dissected and used to count growth rings (Jensen 1969, Allain 1978, Azzolina 1988, Gage 1991). Within one coronal series, the larger plates corresponding to the zone of the ambitus were the most suitable for ring counting. Since in the older ones (near the peristomial membrane) the distal bands were hardly distinct and resorption phenomena were present (Smith 1980). Interambulacral series were used because they were free of podia orifices. The observation of the rings was easy after drying, and the preparation procedure of Jensen (1969) was not necessary. Rings were visualized simply by immersing the plates in xylene and observing them under a stereomicroscope. The number of translucent rings (dark in reflected light) was counted. They were narrower than the opaque ones and easier to count.

Statistical analyses were performed on the size data using 2-way ANOVA for unbalanced designs. The Tukey test (Zar 1984) was used for multiple comparisons. Prior to performing parametric tests, the assumptions of normality and homoscedasticity were tested by Kolmogorov-Smirnov and Bartlett tests, respectively. The SYSTAT package (v. 5.0) was used for the analyses.

The data on size and number of growth rings were transformed to size-at-age under the assumptions of an annual formation of the rings (see 'Discussion') and of 1 main settlement episode per year (Lozano et al. 1995). After trying different models (see 'Discussion'), the Gompertz equation was used to fit a growth function to these measurements. It has the form (we used the same formulation as Cellario & Fénaux 1990):

\[ L_t = L_w e^{-b t} \]

where \( L_t \) is the test diameter at time \( t \), \( L_w \) is the maximum test diameter, the scale parameter \( b = \ln L_w / L_0 \), where \( L_0 \) is the diameter at \( t = 0 \) (settlement on bottom), and \( k \) is a constant of the model.

The means of the sizes of each age-class were used to fit the growth function. Parameter estimation was performed using a nonlinear procedure (NONLIN module of the SYSTAT program with the default Quasi-Newton minimization algorithm).

In order to ascertain the timing of ring formation, a labelling experiment was started in March 1994 in the stable community. A total of 272 individuals of all sizes were collected, taken to the shore in 20 l jars, and a 1% aqueous solution of tetracycline hydrochloride (Boehringer Mannheim) was injected through the peristome. The sea urchins were then returned to their
natural habitat. The dose was set at 0.1 ml per 10 g wet body weight, following Gage (1991). We measured the diameters of the sea urchins to estimate body weight, after the corresponding regression figures calculated with data of the specimens collected for biological cycles. The time of labelling was visible in the test of the sea urchins as a fluorescent tag of tetracycline incorporated into the structure of the test and visible under ultraviolet light. This experiment is planned to continue for 2 yr, but we report here the results concerning some individuals (47 were collected; 29 featured the tetracycline tag) that were recovered from the same place at the end of July 1994. The nature of the outermost band in the plates was also recorded at Tossa throughout the year as indirect evidence of the pattern of ring formation.

Size-frequency distributions. For size-frequency analyses, 1 fixed transect was set at each locality at the same site and was perpendicular to the transects used for density estimations. At Tossa the transect was 6 m long × 1 m wide and was placed between 4 and 8 m in depth following a nearly vertical crevice in the wall. At Cubelles, a 15 × 1 m line was placed parallel to the coast at 0.5 m depth.

All the specimens in these transects were measured (maximum diameter without spines) once a month. Care was taken to examine all cryptic spaces such as undersurfaces of boulders or small crevices. Most of the specimens were satisfactorily measured using sharpened-point calipers without causing harm to the urchins. A slight overestimation of the diameter was unavoidable, due to the underwater working conditions and the presence of spines, which made the use of calipers difficult. The measurements were made at approximately the same hour (13:00 h) during daylight.

This part of the study was performed from January 1993 to January 1994 at Tossa de Mar, and from December 1992 to August 1993 at Cubelles. After this date, the construction of a nearby jetty altered the zone of the transect and we thought it advisable to discontinue the study at Cubelles.

Size-frequency histograms were constructed, and a moving average 3 points wide was applied to filter the noise from the data. Modal analyses were performed to identify cohorts in the size distributions using Battacharya's method (Battacharya 1967), with the aid of the ELEFAN package (ICLARM software, Pauly & David 1981).

RESULTS

Density fluctuations

Fig. 1 shows the temporal trend of the number of ind. m⁻² censused in the 3 fixed transects of 25 m² at Tossa and in the single transect at Cubelles. At Tossa the densities fluctuated between 2 and 11 ind. m⁻², with a trend of decreasing numbers with depth. The global minima were found in September and May, although no sharp variations were detected from one month to the next.

At Cubelles, the single transect surveyed featured mean densities higher than those at Tossa (from 3 to 31 ind. m⁻²), and some abrupt variations were detected, especially at the end of summer in 1992 (from 14 August to 2 September, the population dropped from 765 to 92 individuals in the 25 m²), coinciding with strong storms and easterly winds in the area.

Growth ring measurements

Growth rings were successfully counted in 463 individuals of Tossa (90.96% of the collected specimens) and 492 individuals at Cubelles (91.62%). Most of the specimens in which the reading was not possible, due to merging of rings, were large-sized forms from both localities.

![Graphs showing density fluctuations at Tossa and Cubelles](image-url)
Fig. 2. *Paracentrotus lividus*. Box plots of the diameters of individuals from both localities as a function of the number of translucent rings in the plates of the interambulacral series. Horizontal lines within the boxes are the medians. The upper and lower limits of the boxes indicate the first and third quartiles, respectively. The vertical lines reach the highest (upper line) and lowest (lower line) values within 1.5 times the interquartile distance from the limits of the boxes. Values outside these limits are plotted with asterisks. Numbers of specimens measured in each ring class are indicated. The rectangles in the abscissa indicate ring classes whose mean diameters proved not significantly different in a Tukey test (see 'Results: growth ring measurements').

Fig. 2 displays the median and dispersion of the measures of test diameter against number of growth rings at both localities in the form of box plots (McGill et al. 1978). A 2-way ANOVA with locality and number of rings as factors was performed on the diameter values (ring classes 0 and 11 at Tossa excluded). Both factors proved highly significant (locality: *F*-ratio = 266.67, df = 1, 931, *p* < 0.001; ring number: *F*-ratio = 681.617, df = 9, 931, *p* < 0.001), as was also the interaction term (*F*-ratio = 22.038, df = 9, 931, *p* < 0.001). In the presence of a significant interaction term, we analysed each factor at fixed levels of the other (Underwood 1981). Tukey tests between localities (for a fixed number of rings) showed that the sea urchins from the stable community had significantly larger diameters for all ring classes, except for the forms with a single growth ring. Tukey tests among ring classes (locality factor fixed) revealed that the smaller (up to 2 rings) and larger (from 7 onwards) ring classes were not significantly different within each locality (Fig. 2). This provided evidence of a sigmoid shape of the growth function, with asymptotic left and right legs.

The fitting of a growth function to ring data implies, besides accepting an annual pattern of formation of these marks (see below), the estimation of the time between settlement, formation of the first ring, and time of capture (Sime & Cranmer 1985). We assume here that the main settlement of this species takes place in the beginning of summer (Lozano et al. 1995), and that the formation of the first opaque growth band (the nucleus region, formed during the first months, was not considered) takes place during the period of active growth in the next spring (see below), followed by the formation of the first translucent ring counted here. Therefore, an urchin with 1 translucent ring was between 1 year old (if captured at the beginning the formation of this translucent ring after the spring following the year of settlement) and 2 years old (at the end of the formation of the second opaque ring). In general, as we pooled measures from sea urchins collected year round, we expected that, on average, the sea urchins had an age (in years) = \( \frac{N_{\text{rings}}}{2} + 0.5 \). This value has been adopted to translate data on number of translucent rings to age in years. Fig. 3 represents the fitted Gompertz function against age. The means of the diameters of the different ring classes are also depicted. The parameters of the function at Tossa were \( L_{\infty} = 77.451, b = 2.645, k = 0.252 \), and the residual sum of squares (RSS) of the corresponding regression analysis was 55.59. At Cubelles the figures were \( L_{\infty} = 53.67, b = 2.524, k = 0.287 \) and RSS = 21.444.

Fig. 4 shows the trend of the growth rates with age, obtained from the derivative of the Gompertz function. The maximal growth rate was higher in the stable community (maximum of 7.7 mm yr\(^{-1}\)), and was reached at 4.78 yr. The highest rate at Cubelles (5.8 mm yr\(^{-1}\)) was found at 4.17 yr.

The results of the examination of the marginal ring in sea urchins from Tossa are presented in Fig. 5. The percentage of specimens in which the marginal band was an opaque ring was higher from March to June 1993, with maxima near 80% in May and June. The minimum was found in July 1993 (21%). This indicated that the growth period was concentrated in spring, although actively growing individuals were found year round. The period of low growth leading to the formation of a translucent ring could be somewhat
variable between individuals, but seemed to be concentrated during the gonad building phase (Lozano et al. 1995).

As for the individuals labelled with tetracycline in March 1994 and recaptured at the end of July 1994, in 2 of the 29 specimens the tetracycline tag formed a poorly defined band at the very margin of the plates and these were discarded (the injection could have damaged some organs and the sea urchins featured little or no growth during the intervening period). The distance between the tetracycline mark and the margin of the coronal plates in the other 27 specimens was 0.359 ± 0.179 mm (mean ± standard deviation, measured on the largest plates of the coronal series, which, although they do not feature the highest growth, were the ones used for ring counting in this study). In all except 5 sea urchins the tetracycline mark was found within a narrow translucent ring. Of the 5 remaining specimens, in 2 of them the mark was placed slightly before (i.e. closer to the nucleus) this ring, and in the other 3 slightly after (closer to the margin) it. An opaque band always appeared between the translucent ring and the margin of the plate, which featured the beginning of another translucent ring in most (28 out of 29) individuals. There were, however, fine intervening translucent lines within the opaque band formed after the tetracycline injection. The tetracycline may have caused some disturbances in the growth during this period that were responsible for these poorly defined subsidiary marks. This preliminary result indicated that the period of growth (opaque band formation) was mostly found between March and July, although the observation of marked specimens at later times will be more informative to confirm the pattern of band deposition and avoid the influence of short-term effects of the marking procedure (reported also by Gage 1991).
Size-frequency distributions

Figs. 6 & 7 present, in size classes of 1 mm, the distributions obtained at Tossa (1099 individuals measured) and Cubelles (1002 individuals) during the months studied. A moving average 3 points wide was used to smooth the series. Information on the sizes of the different ring classes in each month, as well as for all months pooled (top of figures), is also included (see figure legends).

No attempt was made to decompose the complete size distributions in modal components, given the low numbers of individuals in some months and the evidence of overlap of the larger cohorts. Instead, the Battacharya method was used to ascertain the normal components which showed up more distinctly in the data, i.e., those with a correlation coefficient higher than 0.6 and a separation index higher than 2 (Pauly & David 1981). Usually, no components were identified above the sizes at which overlap occurs. The normal
Turon et al. Growth and population structure of *Paracentrotus lividus*

Growth and population structure of *Paracentrotus lividus*

The most abundant classes always corresponded to the smallest specimens (less than 10 mm). These tiny individuals could be confused visually with specimens of *Genocidaris maculata* Agassiz, an echinoid which hardly reaches 10 mm in diameter. However, complementary samplings confirmed that the population of sea urchins less than 10 mm in diameter at this site was composed almost exclusively (about 98%) of *Paracentrotus lividus* at all seasons.

The picture for the unstable community was quite different (Fig. 7). The size distribution was variable from one month to the next, and it shifted from a multimodal to a bi- (e.g. June) or uni-modal (e.g. May) shape without apparent regularity. The smallest size classes were always poorly represented (in fact, few individuals less than 20 mm in diameter were recorded), and most specimens were between 30 and 50 mm, a range of sizes in which a superposition of cohorts was suggested from the growth ring measurements. Although the study ended in August, the data for the 8 mo studied seem enough to assign a fluctuating pattern to this community. Some normal components were identified (Fig. 7), but most of the distribution lay in the zone above 40 mm, where no attempt was made to distinguish cohorts, which would have no meaning due to the overlap of the true annual classes.

The instability of the modal components from one month to the next prevented further studies of the histograms at Cubelles. In the stable community, however, the modes below the zone of overlap were reasonably identified and followed with the aid of the data from the ring measurements. Assuming that there was a major recruitment event in the year, and that the translucent ring is primarily formed in summer, we present in Fig. 6 the hypothesized growth of 4 annual classes, corresponding to the sea urchins settled in 1992, 1991, 1990 and 1989. From the sequential increases in size of the frequency modes, we calculated the means of the monthly gains in size of these 4 cohorts (Fig. 8). Important growth variations during the year can be seen, with more active growth in winter-spring and lower rates in August and autumn. Note that the graph in Fig. 8 is quite coincident with the one in Fig. 5 on the formation of the opaque band.
population, although the fact that the yearly recruitment (and, possibly, the yearly mortality) were variable in intensity prevented a smooth decrease in the right part of the size distribution (the 'bumps' corresponding to the more successful year-classes), and made mortality estimates only indicative. To obtain such a gross estimate of the mortality, we fitted a simple exponential function of abundance ($N_t$) as a function of time $N_t = N_0e^{-Zt}$, or $\ln(N_t) = \ln(N_0) - Zt$ to the data on number of individuals per age (by transforming the size values in mm to age in years using the corresponding growth equation estimated above). The value of $Z$ (the instantaneous rate of mortality on a yearly basis) calculated for the above regression at rates (in mm mo$^{-1}$) of the 4 cohorts identified at Tossa was $0.349 (r^2$ of the regression $= 0.716)$, or an annual mortality rate of $1 - e^{-Z} = 0.295$ for specimens larger than 5 mm. It is clear from Fig. 9, however, that the mortality rates were not constant throughout life, and the decrease in number of individuals was steeper where a whole year has been monitored, this graph at sizes below 20 mm (ca 2.5 yr). The instantaneous mortality rate for specimens below this size was $Z = 0.600 (r^2 = 0.937)$ or 0.451 annual rate. The corresponding figures for specimens larger than 20 mm were $Z = 0.258, r^2 = 0.610$, and $1 - e^{-Z} = 0.227$. At the unstable community, the smaller classes were under-represented, and the graph indicated a dominance of forms larger than 30 mm. There was a much steeper decrease in the number of individuals above 40 mm, indicating a higher mortality at this site. However, estimating mortality parameters only in this part of the size distribution is not advisable, since we would miss most of the size classes present in the zone and because information on a complete annual period was not available.

**DISCUSSION**

The use of skeletal growth zones seemed to be the best method for estimating growth in the population studied, due to the difficulty in correctly identifying and interpreting the sequence of size-frequency modes, and the overlap in size of the older age-classes. Alternating phases of high (in which opaque bands are deposited) and low (in which translucized bands are formed) growth have been attributed to the annual reproductive cycle found in many echinoids and, therefore, an annual pattern of band formation has been assumed in studies of growth.
Paracentrotus lividus. However, care must be taken until evidence is obtained that the bands are formed seasonally in any particular species under study (Pearse & Pearse 1975). A further difficulty with the growth bands is the formation of supplementary translucent rings of low growth due to episodic stresses. This can produce an overestimation of the age, especially in old specimens that are more likely to have undergone such stresses.

Validation of the annual pattern of band formation can be achieved indirectly, from the agreement between data obtained from growth bands and from the study of size-frequency distributions (Crapp & Willis 1975, Lumingas & Guillou 1994), or more directly by following the timing of formation of the bands by considering the outermost ring in sequential samples (Taki 1972a, b, Crapp & Willis 1975). Tetracycline labelling has also been used to ascertain the consistency of the bands formed after the tetracycline mark with an annual pattern (Taki 1972a, b, Gage 1991, 1992a, b). In the case of Paracentrotus lividus, Smith (1980) remarked upon the distinctiveness of the growth banding in this species as compared to other echinoids, and Crapp & Willis (1975) concluded from several sources of evidence that the growth structures can be used to age individuals of this species from Ireland on the basis of an annual period of high growth.

Our results on the type of marginal band found in the plates throughout the year, as well as the preliminary information from the tetracycline-labelled individuals, support the annual nature of the pattern of band deposition. The good agreement between the population structure derived from size-frequency data and from ring class data also supports the validity of this method. However, the possibility of overestimation of age in the larger specimens due to the above mentioned problem with the supplementary bands should be taken into account. This is more likely to occur in the unstable than in the stable community.

Selection of a suitable growth function is also a crucial step (Ebert & Russell 1993). The inaccuracy of functions that do not allow a sigmoid shape of the growth curve (such as the Von Bertalanffy model), especially when juveniles are included in the analyses, has been recognized (Gage & Tyler 1985, Sime & Cranmer 1985). Among the functions describing a sigmoid curve, we tried the Richards, Gompertz, and single logistic functions (Richards 1959, Schnute 1981), and finally adopted the Gompertz model on the basis of its lower residual sum of squares. Gage et al. (1986) and Gage & Tyler (1985) also adopted the Gompertz model after comparing different functions. On the other hand, the Richards model was found to be most appropriate by Kenner (1992) and Ebert & Russell (1992). A compromise should be sought between identifying the best fitting function for each set of data and the adoption of a common function allowing for comparison among studies. At present it is not clear which function will provide the best compromise between these needs.

The growth equations fitted to the data showed noticeably lower growth rates (the maximal one was 25% lower), as well as a smaller asymptotic size (about 30% less), in the unstable community with respect to the stable one. This result was as expected, since the scarcity and low quality of the food available in the unstable habitat (Lozano et al. 1995) was combined with more direct exposure to the surf effects in this shallow water zone, and with a higher density of the population. Food and hydrodynamism are key factors in determining the growth rate in echinoids (Ebert 1968, Himmelman 1986), which has also proved to be density dependent (Levitan 1988). The higher density and smaller sizes at the unstable site found here confirm the hypotheses that food limitation has more influence on the size than on the numbers of sea urchins (Ebert 1968) and that density-dependent mortality is not as prevalent in this group as in other invertebrates (Andrew 1989). Besides, a higher investment in gonad production has been observed at the unstable site (Lozano et al. 1995), contributing to the shortage of resources available for somatic growth. A greater longevity at the stable site is also suggested by the fact that individuals with up to 10 (and even 11) rings have been encountered, while only 2 specimens out of 492 featured 10 rings at Cubelles, in spite of the fact that we expected more supernumerary bands in this community, as mentioned above.

Few data on growth of this species are available for comparison with our results. Allain (1978) found different sizes in 2 populations from the French Atlantic coasts, and estimated a longevity of 10 to 11 yr. In one of the populations, a Von Bertalanffy curve was fitted, obtaining an asymptotic size of 61.8 mm, half-way between the 2 values found in our study. Azzolina (1987), in populations from French Mediterranean shores, obtained from the Von Bertalanffy equation a maximal size of 59.29 mm from data on growth rings (which indicated longevities up to 7.5 yr), and of 46.3 mm from growth measures in situ in enclosures. These values are similar to those observed in the unstable community and much lower than those reached at the stable site. Azzolina studied populations in a shallow water bay occupied by a Posidonia oceanica bed whose conditions may be closer to our unstable than to the stable community.

It is also worth noting that the highest growth rate is attained about 6 mo earlier in the unstable community. Although this advance is moderate, this result agrees...
with the general advance of other features, such as gonadal development and percentage of maturity (Lozano et al. 1995): the percentage of mature individuals (all year pooled) plotted against size reached the asymptotic part of the curve at sizes between 20 and 30 mm at Cubelles, while the asymptote was reached at sizes between 40 and 50 mm at Tossa. This finding cannot be accounted for by only the smaller sizes reached at equal age at the former site (a sea urchin of 25 mm at the unstable site is, according to the growth function, 4.164 yr old, while a 45 mm specimen in the stable population is 6.283 yr old).

The use of size-frequency data is useful in describing the general structure of the populations, but ascertaining growth parameters from them is difficult unless the different components or cohorts are well defined and constant through time. This is rarely the case with echinoids, and changing or poorly defined distributions are commonly encountered (Gage et al. 1986, Fénaux et al. 1987, Rowley 1990, Lumingas & Guillou 1994). The overlap of the older age classes in the same size range is a further difficulty in this group (Allain 1978, Azzolina & Boudouresque 1984). Our analyses are thus restricted to the more distinct components that could be determined using the Battacharyya method, and the data on ring classes are incorporated to support the results. Only in the stable community were the modes consistent enough in successive sampling times to suggest a sequence of growth for some cohorts. This complemented the data from the growth function fitted to the size-at-age data by adding a seasonal element. The results showed a seasonally variable growth rate, in which the highest growth is obtained during spring and beginning of summer. This is fully consistent with the data on reproductive periodicities and on deposition of the large opaque band. The period of high growth corresponds to the phase in which mature ova are accumulated in the gonads, and the slow growth phase commenced just after spawning, when gonadal development restarts and the nutritive layer is restored (Lozano et al. 1995). Azzolina (1987) similarly found the highest growth rate of *Paracentrotus lividus* in the French Mediterranean in spring.

The evidence obtained from the structure of the population, together with larval settlement data (presented in Lozano et al. 1995) indicated that markedly different dynamics underlie the arrival of new members to the 2 populations. The recruitment at Tossa, although variable in intensity between years, appeared more constant and predictable, resulting in a population in which the younger size classes predominated (Figs. 6 & 9). At Cubelles, no recruitment was observed in the 2 yr of study, and the evidence from the size distributions (Figs. 7 & 9) strongly suggested that this population underwent a strong limitation in larval settlement.

Echinoids can feature a high interannual variation in recruitment, resulting in exceptionally good years intermingled with periods without apparent recruitment (Ebert 1983). Recruitment limitation has been described in open populations of *Diadema antillarum* (Karlson & Levitan 1990), and this also seems to be the case at Cubelles. Either the population is maintained by episodic recruitment events, such as those described by Peerce & Hines (1987) for *Strongylocentrotus purpuratus* (and the last one must have taken place several years before this study, judging from the absence of small individuals), or it is dependent on migration of larger forms. The latter hypothesis seems the most likely, since after storms it is common to find high numbers of dead sea urchins on the beach at this locality. It is likely that, during storms, individuals coming from a deeper zone (possibly from a nearby *Posidonia oceanica* bed) reached the zone of study. Although many of them, together with sea urchins already instilled in the sampled zone, would be cast onto the beach, some could remain in this shallow area. Besides, the movement of the boulders due to wave action would provoke high mortalities in the population. All this can explain the changing size structure and fluctuating densities found at Cubelles.

In a broad sense, recruitment means addition of new individuals to a population and thus refers not only to larval settlement, but also to immigration (Ebert 1983). We suggest that the reproductive component of recruitment predominates in the stable community, while the migration component explains the dynamics of recruitment more fully in the unstable one.

The population structure found in our 2-site comparison is markedly different from that reported by Himmelman (1986) on *Strongylocentrotus droebachiensis* at 3 sites along a gradient of exposure in Newfoundland. There, the mean size-frequency distributions showed the lack of juveniles at the more protected site, while specimens of the smaller size classes were the most abundant in the exposed zone. The lack of recruitment (or, alternatively, the high juvenile mortality) in the protected area was explained by Himmelman by the high temperatures of the water in this zone, which may not be tolerated by the larvae, and by factors associated with water movement. These contrasting results clearly indicated that the distinction between exposed and sheltered habitats, although important, does not in itself determine the dynamics of the inhabiting echinoid populations, and other factors must be taken into account in each particular case.

Taken together, the results obtained here and those reported in Lozano et al. (1995) indicated a high plas-
ticity in most of the biological parameters studied, which conferred a wide range of adaptive responses to environmental conditions to the sea urchins of this species. In particular, the comparison of a stable and an unstable community suggested that the population inhabiting the former was more settlement-dependent, while the latter population was more migration-dependent, with higher fluctuations of abundance and a changing population structure. Longevity was greater and mortality lower in the stable community. There was an advancement of the age at maturity, and the maximal growth rate was attained at younger ages in the unstable community. The investment in reproduction was higher at the unstable site in spite of the poorer food available and the higher density, and as a result the growth rates were markedly lower and the sizes smaller than in the stable habitat.

Acknowledgements. The authors are grateful to Dr M. Balles-teros, I.I. Dantart, J. Lozano, J. Galera, G. Morera and S. Carner, from the University of Barcelona, for field and laboratory assistance. This work was supported by the Fisheries Department of the Catalan Government.


This article was submitted to the editor

Manuscript first received: September 13, 1994
Revised version accepted: March 16, 1995
Sulfide stress and tolerance in the lugworm *Arenicola marina* during low tide

Susanne Völkel, Kerstin Hauschild, Manfred K. Grieshaber

Institut für Zoologie, Lehrstuhl für Tierphysiologie, Heinrich-Heine-Universität, Universitätsstr. 1, D-40225 Düsseldorf, Germany

ABSTRACT: In the present study environmental sulfide concentrations in the vicinity of and within burrows of the lugworm *Arenicola marina* during tidal exposure are presented. Sulfide concentrations in the pore water of the sediment ranged from 0.4 to 252 pM. During 4 h of tidal exposure no significant changes of pore water sulfide concentrations were observed. Up to 32 pM sulfide were measured in the water of lugworm burrows. During 4 h of low tide the percentage of burrows containing sulfide increased from 20 to 50% in July and from 36 to 77% in October. A significant increase of median sulfide concentrations from 0 to 14.5 pM was observed after 5 h of emersion. Sulfide and thiosulfate concentrations in the coelomic fluid and succinate, alanopine and strombine levels in the body wall musculature of freshly caught *A. marina* were measured. During 4 h of tidal exposure in July the percentage of lugworms containing sulfide and maximal sulfide concentrations increased from 17% and 5.4 pM to 62% and 150 pM, respectively. A significant increase of median sulfide concentrations was observed after 2 and 3 h of emersion. In October, changes of sulfide concentrations were less pronounced. Median thiosulfate concentrations were 18 to 32 pM in July and 7 to 12 pM in October. No significant changes were observed during tidal exposure. Succinate accumulated in the tissue of *A. marina*, indicating the onset of an environmental and sulfide dependent anaerobiosis. Recovery experiments after 4 h of tidal exposure showed that sulfide is completely removed from the coelomic fluid after 30 min. Succinate levels began to decrease immediately after the disappearance of sulfide, reaching control levels after 60 min of recovery. Thiosulfate concentration showed a significant increase after 30 min, indicating that internal sulfide is removed by its oxidation to thiosulfate. The present study shows that *A. marina* exhibits the same mechanisms of sulfide tolerance under habitat conditions as under experimental conditions. In addition, it is demonstrated that the lugworm is able to recover from tidal sulfide stress within 1 h.

KEY WORDS: *Arenicola marina* · Sulfide tolerance · Tidal exposure · Sediment · Recovery

INTRODUCTION

Animals inhabiting the sediment of the intertidal zone can be exposed to pronounced fluctuations of abiotic conditions. The tidal rise and fall of the sea causes periodical changes in temperature, salinity and oxygen supply. The lugworm *Arenicola marina* lives about 10 to 30 cm deep in the sediment of intertidal flats. During high tide its U-shaped burrow is irrigated by peristaltic movements of its body wall thus providing the animal with oxygen (Krüger 1971). At low tide when the burrow is emersed, ventilation becomes impossible and the lugworm is exposed to increasing hypoxia. Jones (1955) demonstrated that the P<sub>O₂</sub> in the remaining water of the lugworm burrow decreases from about 33 to 13 torr during 2 h of tidal emersion. Correspondingly, blood oxygen content drops nearly to zero within the first hour of tidal emersion (Toulmond 1973). In the hypoxic period *A. marina* reduces its ventilatory movements and oxygen consumption and switches from aerobic to anaerobic metabolism (Schöttler et al. 1984a, Toulmond & Tchernigovtzeff 1984, Toulmond 1987, Grieshaber et al. 1992). Thus, the accumulation of typical anaerobic metabolites such as succinate, acetate and propionate in the body wall tissue and blood of *A. marina* was observed during tidal exposure (Pionetti & Toulmond 1980, Schöttler et al. 1984b). When the tide comes back in, the burrow is ventilated again and the lugworm can return to an aerobic metabolism. Pörtner et al. (1997) showed that...
A. marina recovered from anoxia with most of the tissue metabolites reach control levels during 1 to 2 h of normoxic incubation.

Apart from irrigated burrows of the tubebuilding infauna, oxygen is not measurable in the pore water of the deeper layers of the marine sediment (Brafield 1964, Watling 1991). These reduced layers are commonly characterized by the presence of sulfide, which is mainly produced by sulfate-reducing bacteria (Jørgensen & Fenchel 1974, Kröger et al. 1988). Sulfide concentrations in the sediment pore water of marine habitats can range from a few μM up to several mM (see Bagarinao 1992) depending on the structure of the sediment and organic matter production (Fenchel & Riedl 1970). The rate of sulfate reduction shows seasonal fluctuations which are mainly caused by changes of temperature and organic matter supply (Nedwell & Floodgate 1972, Jørgensen 1977). Sulfide concentrations in sediment pore water are usually high in summer and low during the cold months (Jørgensen 1977, Vökel & Grieshaber 1992). Diel fluctuations of sulfide can occur as a consequence of changing light conditions (De Wit et al. 1989).

In the vicinity of lugworm burrows sulfide concentrations up to 340 μM have been observed (Groenendaal 1979, Vökel & Grieshaber 1992). Sulfide is assumed to diffuse from the pore water along the chemical gradient into the lugworm burrow (Aller 1980, Wastenchuk et al. 1983). During high tide sulfide may be delivered to the overlying water as the lugworm irrigates its burrow. Sulfide is, moreover, likely to be oxidized rapidly by the oxygen-rich seawater (Cline & Richards 1989, Millero 1986). Sulfide will therefore only be rarely found in burrows which are covered by tide. During tidal exposure, however, the burrow water is stagnant and becomes hypoxic (see above). Sulfide diffusing into the burrow cannot be flushed out or oxidized and Arenicola marina may be exposed to increasing sulfide concentrations as long as the burrow is emersed. Arp et al. (1992), for example, measured sulfide concentrations up to 66 μM in the burrow water of Urechis caupo after 2 h of tidal exposure. As soon as the tide comes back in, the worm is able to ventilate again and sulfide will be removed from the burrow. A. marina is thus exposed not only to a repetitive lack of oxygen but also to periodically changing sulfide concentrations during tidal cycles.

Sulfide is a strong inhibitor of the cytochrome c oxidase (National Research Council 1979). Despite sulfide’s toxicity, numerous organisms possess several mechanisms of sulfide tolerance and can live in sulfide-rich habitats (for reviews see Vetter et al. 1991, Vismann 1991). Arenicola marina is highly insensitive to sulfide (Groenendaal 1980). Its sulfide tolerance has been investigated in the laboratory and is based upon 3 main strategies: (1) in the presence of oxygen, sulfide entering the body is rapidly oxidized to thiosulfate which accumulates in the coelomic fluid of A. marina. The oxidation of sulfide could be localized in the mitochondria of its body wall tissue (Vökel & Grieshaber 1992, 1994). (2) During hypoxia, the sulfide concentration in the body of A. marina is reduced by the acidification of the coelomic fluid (Groenendaal 1981, Vökel & Grieshaber 1992). (3) The increase of sulfide in the body can be tolerated by switching to an anaerobic metabolism as indicated by the accumulation of anaerobic metabolites (Vökel & Grieshaber 1992, 1994, see also Grieshaber et al. 1992). Although these mechanisms have been demonstrated under extreme experimental conditions, up to now it is not known whether they also play a role in the lugworm’s habitat. The aim of this study, therefore, was (1) to determine the environmental sulfide concentrations to which A. marina is exposed during prolonged low tide, (2) to investigate whether A. marina exhibits the same mechanisms of sulfide tolerance in its natural habitat as under experimental conditions, and (3) to investigate the time course of recovery of A. marina from sulfide exposure during tidal emersion when reimmersed.

**MATERIAL AND METHODS**

**Study site and sampling conditions.** The studies took place at a mudflat located at the harbor of St. Pol de Léon, Brittany, France (English Channel). The site is characterized by a high population density of Arenicola marina. Sampling was performed during July 1991, July and October 1992 and October 1993 and 1994 when tidal heights were 0.9 to 1.5 m and peaks of low tide were around noon. All environmental and animal data were obtained within an area of about 10 × 10 m which was exposed for 4 to 5 h during low tide. Sampling started as soon as the water had receded (t = 0) and was repeated after 1, 2, 3 and 4 h of emersion. During collecting trips air temperatures ranged from 12.5 to 24.9°C in July and 4.8 to 12.4°C in October (minimal and maximal temperatures, respectively; data obtained from the Station météorologique, Brest, France). The weather was dry during all collecting trips with the exception of 15 July 1991 and 28 October 1992 when it started to rain after 1 h and 3 h, respectively.

**Sulfide concentration of the pore water.** Sulfide concentration of the pore water of the sediment was determined during 3 collecting trips in July 1991. Each time pore water was sampled from 2 different sites at a depth of 10, 20 and 30 cm. Pore water samplers as described by Vökel & Grieshaber (1992) were inserted into the sediment immediately after emersion of the
area and remained there for 4 h. Every hour, 0.5 to 1 ml pore water was collected, 3 samples from each depth and site. The samples were fixed in zinc acetate and NaOH and analyzed for sulfide within 1 wk with the methylene blue method (Gilboa-Garber 1971) as described by Völkel & Grieshaber (1992).

Sulfide concentration of the burrow water. The sulfide concentration of the burrow water was determined in July 1992 and in October 1992, 1993 and 1994. Burrow water was sampled using a polyethylene tube (length 30 cm, inner diameter 1 mm) which was attached to a 1 ml tuberculin syringe. After the removal of the lugworm's cast the tube was carefully inserted into the burrow until it was hindered by the worm. Samples of 0.5 to 1 ml were taken from each burrow. At each time point, 2 to 3 burrows were analyzed at the same time. The first sample was taken immediately after the emersion of the burrows and additional samples were taken hourly for 4 h. In October 1993 the period of emersion was long enough to obtain samples after 5 h of emersion during 1 collecting trip. Each burrow was used only once to prevent artifacts arising from diffusion from the environment. Water samples were fixed and analyzed for sulfide as described above.

Determination of sulfur compounds and anaerobic end products in freshly caught Arenicola marina. Specimens of Arenicola marina were collected in July 1991, July and October 1992 and in October 1993. During each collecting trip, 2 to 3 lugworms h⁻¹ were collected from unanalyzed burrows during the 4 h of emersion. The freshly caught lugworms were quickly washed in seawater. Coelomic fluid was collected by dorsally cutting the body wall. After that, the worms were dissected and the body wall tissue was freeze-clamped (Wollenberger et al. 1960) and stored in liquid nitrogen.

For determination of sulfide and thiosulfate the coelomic fluid was immediately mixed with monobromobimane (3 mmol l⁻¹ final concentration; Calbiochem, Giessen, Germany) and HEPES/EDTA (50/5 mmol l⁻¹, pH 8.0) according to a modified method described by Vetter et al. (1989). As a control, parallel samples were prepared using 2-pyridyl disulfide (Vetter et al. 1989). After a reaction time of 30 min methansulfonic acid (25 mmol l⁻¹) was added and the samples were frozen and stored in liquid nitrogen for 2 mo at most. In the laboratory, the samples were thawed and spun for 10 min (14 550 X g, Biofuge A, Heraeus Christ, Osterode, Germany). The supernatant was immediately analyzed for sulfur compounds by high-performance liquid chromatography (HPLC) as described by Völkel & Grieshaber (1994).

For determination of succinate, alanopine and strombine the tissue was extracted according to Beis & Newsholm (1975). Succinate was measured spectrophotometrically as described by Beutler (1985). Alanopine and strombine were determined by HPLC using a DX–100 Ion Chromatograph (DIONEX, Idstein, Germany) for pumping and conductivity detection. The opines were isocratically separated at 45°C with a PolyspherAR AC cation exchange column (100-6.5, Merck, Darmstadt, Germany) using 7.5 × 10⁻⁵ N H₂SO₄ as a solvent (0.6 ml min⁻¹).

Recovery experiments. During 2 collecting trips in October 1993, 12 specimens of Arenicola marina were collected after 4 h of emersion. The lugworms were immediately placed into 10 l aerated seawater from their habitat and were allowed to recover in darkness. After 15, 30, 60 and 120 min, respectively, 3 worms were removed from the tank. Coelomic fluid was collected and prepared for analysis of sulfur compounds and the body wall musculature was stored for determination of anaerobic end products as described above.

Data treatment. Results are reported as single data and medians respectively. Since most of the data were not normally distributed, significance of differences between medians were evaluated using the non-parametric Mann-Whitney U-test (2-tailed test) at the p = 0.05 level (Beyer 1988).

RESULTS

Sulfide in the pore water of the sediment

Sulfide concentrations in the pore water ranged from 0.4 to 252.2 µM (Fig. 1). Variations were high both between different sampling sites and between different sampling days. For instance, at a depth of 10 cm and at t = 0 h, sulfide concentrations from 13.7 to 177.2 µM could be measured. For the first 2 collecting trips (a, 12 July; b, 13 July 1991) sulfide concentrations in the pore water tended to increase during low tide. For the third collecting trip (c, 15 July 1991) sulfide concentrations increased in the first hour of emersion but decreased during the remaining 3 h of emersion. In almost all cases sulfide concentrations decreased with depth, e.g. from 77.8 µM (a, site I, t = 4 h) at a depth of 10 cm to 7.8 µM at 30 cm. Due to the high variability of the data changes in sulfide concentrations during low tide were not significant.

Sulfide in the burrow water

Sulfide was present in 23% of the lugworm burrows analyzed during low tide in July 1992 (n = 26) with concentrations ranging from 0.2 to 32 µM (Fig. 2). Immedi-
Sulfur compounds in the coelomic fluid of *Arenicola marina*

**Sulfide**

During collecting trips in July 1991 and 1992 sulfide was measured in the coelomic fluid of 46% of 37 freshly caught specimens of *Arenicola marina*, with concentrations ranging from 5.4 to 150 \( \mu M \) (Fig. 3A). During tidal exposure, the percentage of lugworms containing sulfide, the medians of the sulfide concentrations and the maximal sulfide concentrations increased from 17%, 0 \( \mu M \) and 5.4 \( \mu M \) at the beginning to 62%, 35.1 \( \mu M \) and 150.0 \( \mu M \), respectively, after 4 h of emersion. Due to the high variations between lugworms the increase of sulfide concentration of the coelomic fluid was only significant after 2 and 3 h but not after 4 h of emersion when compared to \( t = 0 \) h (Fig. 3A). In October 1992 and 1993, 49 specimens of *A. marina* were analyzed, 57% of which contained sulfide ranging from 0.7 to 13 \( \mu M \) (Fig. 3A). As in July, the percentage of lugworms containing sulfide, the median and the maximal sulfide concentrations increased with duration of low tide (33%, 0 \( \mu M \) and
centrations of October specimens after 3 and 4 h were much (significantly at \( t = 3 \) h) lower than the corresponding July values (0 and 6.4 \( \mu M \) as opposed to 46.0 and 35.1 \( \mu M \) respectively; Fig. 3A). Likewise, maximal sulfide concentrations were only 9 to 13 \( \mu M \) in lugworms collected in October as compared to 66 to 150 \( \mu M \) in those collected in July after 2 to 4 h of emersion.

8.4 \( \mu M \) at the beginning to 78%, 6.4 \( \mu M \) and 12.8 \( \mu M \), respectively, after 4 h of tidal exposure). A significant increase of median sulfide concentrations could only be measured after 4 h of emersion. Median sulfide concentrations of October specimens after 3 and 4 h were much (significantly at \( t = 3 \) h) lower than the corresponding July values (0 and 6.4 \( \mu M \) as opposed to 46.0 and 35.1 \( \mu M \) respectively; Fig. 3A). Likewise, maximal sulfide concentrations were only 9 to 13 \( \mu M \) in lugworms collected in October as compared to 66 to 150 \( \mu M \) in those collected in July after 2 to 4 h of emersion.

**Thiosulfate**

In July 1991 and 1992 thiosulfate could be measured in the coelomic fluid of 76% of the lugworms (\( n = 50 \)). The concentrations ranged from 10.2 to 73.0 \( \mu M \), median concentrations were 17.7 to 32.2 \( \mu M \) (Fig. 3B). During 4 h of tidal emersion no significant changes of median or maximal thiosulfate concentrations or of the percentage of lugworms containing thiosulfate could be observed. In October 1992 and 1993, 58 specimens of *Arenicola marina* were analyzed, 95% of which contained thiosulfate at concentrations of 0.4 to 34.7 \( \mu M \). As in July, no changes of median concentrations (6.9 to 11.6 \( \mu M \)), maximal concentrations or percentage of thiosulfate-containing lugworms were found. Medians were significantly lower in October than in July (at \( t = 2 \) and 3 h). Maximal thiosulfate concentrations in the coelomic fluid of lugworms collected in October were 1.5- to 4-fold lower than of those collected in July (Fig. 3B).

**Anaerobic end products in the tissue of *Arenicola marina***

The concentrations of succinate (July 1991 and 1992 and October 1992 and 1993), alanopine and strombine (July 1991 and 1992) in the body wall musculature of freshly caught *Arenicola marina* were determined in order to follow the onset of an anaerobic metabolism. In July median succinate concentrations increased significantly from 0.33 \( \mu mol \cdot g^{-1} \) wet wt at the beginning to 0.81 \( \mu mol \cdot g^{-1} \) wet wt after 2 h and 0.92 \( \mu mol \cdot g^{-1} \) wet wt after 4 h of emersion (Fig. 3C).
The concentrations ranged from 0.07 to 1.24 \( \mu \text{mol g}^{-1} \) wet wt at \( t = 0 \) h, from 0.39 to 1.43 \( \mu \text{mol g}^{-1} \) wet wt at \( t = 2 \) h and from 0.23 to 2.75 \( \mu \text{mol g}^{-1} \) wet wt at \( t = 4 \) h. Similarly, median and maximal succinate concentrations increased in the body wall tissue of lugworms collected in October with the length of tidal emersion (Fig. 3C). The median concentrations were 0.21 \( \mu \text{mol g}^{-1} \) wet wt in the beginning and increased significantly to 0.62 \( \mu \text{mol g}^{-1} \) wet wt after 2 h and to 1.01 \( \mu \text{mol g}^{-1} \) wet wt after 4 h of emersion. The respective concentrations ranged from 0.06 to 0.31 \( \mu \text{mol g}^{-1} \) wet wt at \( t = 0 \) h, from 0.34 to 1.21 \( \mu \text{mol g}^{-1} \) wet wt at \( t = 2 \) h and from 0.72 to 2.02 \( \mu \text{mol g}^{-1} \) wet wt at \( t = 4 \) h. No significant differences were observed between lugworms collected in July and those collected in October.

Alanopine in the tissue remained constant (at a level of about 1 \( \mu \text{mol g}^{-1} \) wet wt) during 4 h of emersion although maximal alanopine concentrations tended to increase in the first 3 h of emersion (Table 1). The median strombine level increased significantly from 0.66 \( \mu \text{mol g}^{-1} \) wet wt at \( t = 0 \) h to 1.53 \( \mu \text{mol g}^{-1} \) wet wt after 3 h of emersion. Similarly, maximal strombine concentrations increased from 1.10 to 6.35 \( \mu \text{mol g}^{-1} \) wet wt in the same period of time (Table 1). However, median and maximal strombines concentrations decreased to 0.88 and 3.95 \( \mu \text{mol g}^{-1} \) wet wt after 4 h of emersion.

<table>
<thead>
<tr>
<th>Time of emersion (h)</th>
<th>n</th>
<th>Min. value</th>
<th>Max. value</th>
<th>Median</th>
</tr>
</thead>
<tbody>
<tr>
<td>Alanopine</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>0</td>
<td>8</td>
<td>0.49</td>
<td>1.40</td>
<td>0.86</td>
</tr>
<tr>
<td>1</td>
<td>11</td>
<td>0.25</td>
<td>2.70</td>
<td>0.88</td>
</tr>
<tr>
<td>2</td>
<td>12</td>
<td>0.06</td>
<td>3.36</td>
<td>0.47</td>
</tr>
<tr>
<td>3</td>
<td>10</td>
<td>0.25</td>
<td>4.23</td>
<td>1.27</td>
</tr>
<tr>
<td>4</td>
<td>10</td>
<td>0.14</td>
<td>2.25</td>
<td>1.08</td>
</tr>
<tr>
<td>Strombine</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>0</td>
<td>8</td>
<td>0.22</td>
<td>4.10</td>
<td>0.96</td>
</tr>
<tr>
<td>1</td>
<td>11</td>
<td>0.13</td>
<td>3.74</td>
<td>0.93</td>
</tr>
<tr>
<td>2</td>
<td>12</td>
<td>0.09</td>
<td>4.13</td>
<td>0.91</td>
</tr>
<tr>
<td>3</td>
<td>10</td>
<td>0.30</td>
<td>6.35</td>
<td>1.53</td>
</tr>
<tr>
<td>4</td>
<td>10</td>
<td>0.10</td>
<td>3.95</td>
<td>0.88</td>
</tr>
</tbody>
</table>

**Recovery**

Specimens of Arenicola marina were collected after 4 h of emersion and were incubated in normoxic seawater in order to investigate recovery from tidal exposure. The median sulfide concentration in the coelomic fluid (\( \mu \text{M} \), , ) and succinate in the body wall tissue (\( \mu \text{mol g}^{-1} \) wet wt, , ) and (B) thiosulfate in the coelomic fluid (\( \mu \text{M} \), , ) of lugworms collected in October. Lugworms were collected after 4 h of tidal emersion (\( t = 0 \) min) and were allowed to recover in normoxic seawater. Open symbols represent data from individual lugworms; closed symbols are median concentrations. (*) Median concentration is significantly different from the median at \( t = 0 \) min. The data for \( t = 0 \) min are identical with \( t = 4 \) h (October) in Fig. 3. For clarity, sulfide data in (A) are shifted to the right. The number of lugworms at \( t = 0 \), 15, 30, 60 and 120 min of recovery was (with the number of lugworms containing no sulfide in parentheses): 9 (2), 5 (2), 6 (6), 5 (4) and 6 (3). Thiosulfate and succinate were found in all lugworms (\( n = 5 \) or 6).
fluid began to decrease from 6.4 μM (see above) to 4.2 μM after 15 min of recovery (Fig. 4A). Sulfide was completely removed from the coelomic fluid after 30 min of recovery. Succinate concentrations were still high after 15 min of recovery (median = 1.21 μmol g⁻¹ wet wt), began to decrease after 30 min (0.7 μmol g⁻¹ wet wt) and reached control values after 60 and 120 min of recovery (medians = 0.33 and 0.17 μmol g⁻¹ wet wt, respectively) (Fig. 4A). In contrast, the median thiosulfate concentration in the coelomic fluid of A. marina showed a sudden significant increase from 14.4 to 32.4 μM after 30 min of recovery with concentrations ranging from 10.6 to 103.5 μM (Fig. 4B). After 60 min of recovery, the median thiosulfate concentration had returned to the control value of 12.3 μM, concentrations ranging from 1.6 to 30.9 μM.

DISCUSSION

Environmental sulfide concentrations

Sulfide was measured in the pore water of the sediment in the vicinity of lugworm burrows in order to see whether sulfide concentrations vary during tidal exposure. Like many other marine sediments (Ott & Novak 1989) the area exhibited a pronounced heterogeneity (Fig. 1). Maximal sulfide concentrations were comparable to earlier investigations [250 μM in this study as compared to 340 μM in Völkel & Grieshaber (1992) with both values obtained from July measurements]. Sulfide concentrations decreased with depth (Fig. 1): the maximal concentration was only 44 μM at a depth of 30 cm, which may be due to a decreasing rate of sulfate reduction parallel to a decreasing sulfate gradient in the sediment (Jørgensen 1977).

Although in some cases pore water sulfide concentrations increased during 4 h of tidal emersion, variations between different sampling sites were high and no significant changes of sulfide concentrations could be observed.

Sulfide concentrations in the water of lugworm burrows were generally lower than pore water concentrations although sulfide was detected in 42% of the analyzed burrows. Most of them did not contain any sulfide just after the water had receded. During 4 h of tidal emersion the percentage of burrows containing sulfide doubled whereas median sulfide concentrations in the burrow rose slightly but not significantly. A significant increase to 14.5 μM was only observed after 5 h of emersion in October (Fig. 2). In July, however, sulfide concentrations up to 30 μM could be measured in single burrows over the whole period of emersion. These sporadic high values may be due to sediment heterogeneity as mentioned above. On the other hand, sulfide concentrations in burrow water is frequently higher in summer than in winter and spring (Arp et al. 1989).

Arp et al. (1992) found sulfide in 50 to 70% of Urechis caupo burrows with maximal concentrations of 25 to 65 μM (samples from 2 different sites, respectively). In their study samples were taken about 2 h after emersion and unfortunately these authors give no sulfide concentrations at the beginning of low tide. Waslenchuk et al. (1983) analyzed burrows of Callianassa spp. which were covered with water over the whole sampling period. They found sulfide concentrations of 2.0 to 26 μM in the burrow water as opposed to less than 0.1 μM in the overlying water and 100 to 1000 μM in the pore water. Although the shrimps vigorously flush their burrows, irrigation is not continuous and therefore seems to be insufficient to maintain sulfide-free burrows (Waslenchuk et al. 1983). Arenicola marina also exhibits an intermittent irrigation pattern (Davey et al. 1990). It is, therefore, possible that A. marina is occasionally exposed to short-term increases of sulfide concentrations in its burrows during high tide. This may explain why sulfide was already found in some burrows at the beginning of low tide (Fig. 2).

Sulfur compounds in freshly caught Arenicola marina

Sulfide concentrations in the coelomic fluid of Arenicola marina were measured every hour during 4 h of tidal emersion in order to see how much sulfide is taken up by the worms under habitat conditions. When the flat fell dry, the percentage of lugworms containing sulfide increased with exposure time, along with both the median and the maximal sulfide concentrations in the coelomic fluid. Due to the high variability between lugworms, the increase of median sulfide concentration was only significant after 2 and 3 h in July and after 4 h in October (Fig. 3A). Immediately after the water had receded only low sulfide concentrations (<10 μM) were found in a few lugworms whereas no sulfide could be measured in about 70% of the lugworms. These data indicate that either no sulfide was present in the burrow water of these lugworms or that some sulfide was present but that entering sulfide could be oxidized due to a sufficiently high oxygen supply during high tide. In earlier investigations (Völkel & Grieshaber 1994) we observed that under normoxic conditions A. marina is able to prevent an increase of sulfide in the coelomic fluid up to an external sulfide concentration of at least 330 μM, since sulfide entering the body is rapidly oxidized to thiosulfate. Fig. 3B shows that thiosulfate concentrations up to 40 μM are present immediately after the water has receded, indicating that some sulfide must have been present in the body of these lugworms and that this sul-
Sulfide had been oxidized during high tide. Oeschger & Vetter (1992) measured sulfide and thiosulfate in the hemolymph of freshly caught priapulid Halicryptus spinulosus living in anoxic sediments of the Western Baltic. In their study sulfide concentrations of 86 to 445 μM were measured whereas thiosulfate amounted to 21 to 84 μM. These high internal sulfide levels may be a consequence of limited oxygen conditions together with pore water sulfide concentrations up to 665 μM (Oeschger & Vetter 1992).

In the coelomic fluid of Arenicola marina collected in July, sulfide concentrations rose significantly to about 35 μM after 3 h of emersion and maximal concentrations were even higher (Fig. 3A). In the lugworm burrows sulfide levels were 20 to 30 μM at most (Fig. 2). Sulfide concentrations within lugworms, therefore, can be higher than burrow water concentrations. In laboratory experiments sulfide concentrations in the coelomic fluid were 60 μM after 8 h of hypoxic (P0, = 3 torr) sediment-free incubation at an external sulfide level of 200 μM (Völkel & Grieshaber 1992). Internal sulfide concentrations reported by Völkel & Grieshaber were comparable to those which were found in the present study. Under laboratory conditions, however, sulfide concentrations in the coelomic fluid were much lower than external sulfide concentrations. This was due to a pH-dependent limited influx of sulfide into the body of A. marina (Groenendaal 1981, Völkel & Grieshaber 1992). Under field conditions sulfide may have entered the lugworm by it feeding on sulfide-rich sediment.

The increase of sulfide within lugworms during low tide indicates that sulfide oxidation was not possible due to the lack of oxygen. In an earlier investigation (Völkel & Grieshaber 1994) sulfide oxidation proved to be oxygen dependent. Thiosulfate concentrations in the coelomic fluid of Arenicola marina were only 2 μM after hypoxic sulfide incubations at a P0, of 0.7 torr as opposed to 4.2 mM at a P0, of 130 torr (8 h, external sulfide 550 μM; Völkel & Grieshaber 1994). Although thiosulfate concentrations were slightly higher in July than in October, no changes of thiosulfate concentrations were observed during tidal exposure (Fig. 3b). Völkel (1992) demonstrated that during hypoxia thiosulfate disappears very slowly from the coelomic fluid of A. marina, which may explain the constant thiosulfate concentrations during tidal exposure.

Anaerobic metabolites in freshly caught Arenicola marina

During tidal exposure the stagnant water conditions in lugworm burrows can cause anoxia as well as enhanced sulfide load. Both factors prevent aerobiosis, leading to anaerobic metabolism. In an earlier study we demonstrated the accumulation of the anaerobic metabolites succinate, alanopine and strombine in the body wall tissue of Arenicola marina during sulfide incubations under hypoxic conditions (Völkel & Grieshaber 1992, 1994). In the present investigation tissue concentrations of anaerobic metabolites were measured in order to follow the onset of anaerobic metabolism during tidal exposure. Succinate in the body wall tissue of Arenicola marina collected in October increased significantly from 0.21 μmol g⁻¹ wet wt to 1.01 μmol g⁻¹ wet wt after 4 h of emersion (Fig. 3C). These data show that anaerobiosis commenced at least after 2 h of tidal exposure. In laboratory experiments succinate amounted to about 1 μmol g⁻¹ wet wt after 8 h of hypoxia (P0, 1 to 2 torr) and to about 2 μmol g⁻¹ wet wt after 8 h of hypoxic sulfide incubations (external sulfide 200 and 1000 μM, respectively) (Völkel & Grieshaber 1992). The above data correspond well to succinate levels in the body wall tissue of freshly caught A. marina found in the present study.

Unfortunately, it is impossible to distinguish between environmental and sulfide-dependent anaerobiosis since we do not know the oxygen and sulfide concentrations in the burrow water of the individual lugworms. Fig. 5 shows that most of the lugworms which contain sulfide exhibit high succinate levels (0.9 to 1.2 μmol g⁻¹ wet wt), as aerobiosis cannot be maintained due to inhibition of the cytochrome c oxidase. In contrast, less succinate (0 to 0.6 μmol g⁻¹ wet wt) was found in most lugworms without sulfide. In many other specimens, however, succinate levels were high although no sulfide could be found in their coelomic fluid. In that case, anaerobiosis was probably caused by anoxia and not by sulfide.

Schöttler et al. (1984b) measured 0.14 μmol g⁻¹ wet wt succinate in the tissue of Arenicola marina at the beginning of low tide, increasing to 0.25 after 2 h and to 0.28 μmol g⁻¹ wet wt after 4 h of emersion. In the above study the authors give no sulfide concentrations in the environment or in the lugworms. Possibly, the lower succinate level, as compared to the present study, was a consequence of a lower sulfide stress. Schöttler (1989) demonstrated that the extent of anaerobic metabolism also depends on the season, the locality of the intertidal burrow and the development of gametes. Correspondingly, summer specimens of A. marina accumulated significantly more succinate during normoxic sulfide incubations than winter specimens (Völkel & Grieshaber 1994). In the present study, however, no differences of succinate levels between lugworms collected in July and those collected in October were observed (Fig. 3C) although sulfide exposure was significantly higher in July (Fig. 3A). Sulfide entering the body of A. marina may quickly reach a critical level; the cytochrome c oxidase is blocked, also
leading to maximal anaerobiosis which does not depend on the sulfide concentration. During hypoxic sulfide incubations there was no difference between succinate accumulation at external sulfide concentrations of 200 or 1000 μM (Volkel & Grieshaber 1992).

In addition to succinate, we measured opine levels in the body wall tissue of lugworms collected in July. Alanopine did not change significantly although maximal concentrations tended to increase in the first 3 h of emersion (Table 1). Strombine concentrations increased significantly during the first 3 h of tidal exposure. Alanopine and strombine both are glycolytic end products, with alanopine being accumulated preferentially during functional and strombine during environmental anaerobiosis (Siegmund et al. 1985, see also Grieshaber et al. 1992). Strombine therefore accumulates predominantly in Arenicola marina during tidal hypoxia with maximal concentrations being higher than maximal alanopine concentrations. Volkel & Grieshaber (1992) demonstrated that much more strombine is accumulated during 8 h of hypoxia, hypoxia plus sulfide, and normoxia plus sulfide as compared to alanopine. Tissue levels of alanopine were about 1 to 1.5 μmol g⁻¹ wet wt whereas strombine amounted to 4 μmol g⁻¹ wet wt (Volkel & Grieshaber 1992).

**Recovery**

During tidal exposure, sulfide and succinate accumulate in the body of Arenicola marina whereas thiosulfate concentrations remain constant (Fig. 3). As soon as the tide comes back in, the lugworm can irrigate its burrow again. Sulfide which has accumulated in the burrow water is flushed out and the presence of oxygen enables the worm to oxidize sulfide which may have entered its body. When internal sulfide has disappeared, the cytochrome c oxidase is no longer inhibited and A. marina is able to switch back to aerobic metabolism. In the present study we mimicked the incoming tide by placing the lugworms into aerated seawater tanks after 4 h of tidal exposure in the sediment. Sulfide concentrations in the coelomic fluid began to decrease immediately and reached control levels after 30 min of recovery (Fig. 4A). Two mechanisms may be involved in the disappearance of sulfide: the diffusion into the external medium and the oxidation of sulfide to thiosulfate. The diffusion of sulfide may certainly play a role because biological membranes are permeable to sulfide (Beerman 1924, Julian & Arp 1992, Volkel & Grieshaber 1992). On the other hand, Fig. 4B shows that thiosulfate in the coelomic fluid of A. marina increases significantly after 30 min of recovery. A rough estimation reveals that sulfide disappears from the coelomic fluid with a rate of 0.12 nmol min⁻¹ g⁻¹ whereas thiosulfate production is 0.26 nmol min⁻¹ g⁻¹ (assuming a lugworm weight of about 10 g and a relative amount of coelomic fluid of 40%). One mole of produced thiosulfate is equivalent to 2 moles of oxidized sulfide. Therefore, thiosulfate production during the first 30 min of recovery must be the consequence of the oxidation of 0.52 nmol min⁻¹ g⁻¹ sulfide which is 4-fold higher than the measured rate of sulfide removal. On the other hand, sulfide concentrations in the body wall tissue are much higher as compared to the coelomic fluid. For example, after 7 h of hypoxic incubations at an external sulfide concentration of 25 μM, sulfide concentrations increased by about 2 to 3 μM in the coelomic fluid and about 30 μM in the body wall tissue of A. marina (K. Hauschild unpubl. results). Unfortunately, it was not possible to determine tissue sulfide levels because we were not able to determine the fresh weight of the tissue under field conditions. If we assume a similar increase of sulfide in the body wall musculature as seen under experimental conditions, the total removal of sulfide would require a rate of approximately 0.49 nmol min⁻¹ g⁻¹. This rate is com-
rapidly return to an aerobic metabolism. Although the mechanisms of sulfide removal during recovery from sulfide exposure remain to be investigated in detail, it can be assumed that the oxidation of sulfide to thiosulfate plays a major role in this process. After 60 min of recovery thiosulfate in the coelomic fluid reached the same level as during low tide (Fig. 4B). Very little is presently known about degradation or excretion of thiosulfate. It may be further oxidized to sulfate or it may be excreted via the nephridia. Thiosulfate is not completely removed from the coelomic fluid during 120 min of recovery. Possibly, any thiosulfate oxidizing or transporting system is only activated by thiosulfate concentrations above a critical threshold.

Parallel to the disappearance of sulfide, the succinate level in the body wall tissue decreases during recovery. Although sulfide had reached control levels after 30 min of recovery, loss of succinate appeared to be slower (Fig. 4A). Pörtner et al. (1979) showed that succinate began to decrease immediately after the onset of aerobic conditions and reached control levels after 1 to 2 h of recovery from anaerobiosis. From the change of energy status and the immediate cessation of fumarate reduction they concluded that Arenicola marina is able to utilize oxygen as soon as it is available. In the presence of sulfide, however, a rapid onset of aerobicosis is not possible. Fig. 4A shows that considerably high sulfide concentrations are still present after 15 min of recovery. Correspondingly, the succinate level is still high at this moment. Only when sulfide has disappeared can A. marina switch back to an aerobic metabolism, as indicated by decreasing succinate concentrations after 30 min of recovery (Fig. 4A).

During recovery, thiosulfate concentration in the coelomic fluid of Arenicola marina was maximal after 30 min when sulfide had reached control levels and succinate has decreased (Fig. 4B). Thiosulfate production, however, started earlier when sulfide and succinate levels may still have been high. In an earlier investigation (Volkel & Grieshaber 1994) we observed an accumulation of thiosulfate in the coelomic fluid of A. marina during normoxic sulfide incubations (PO2 = 130 torr, external sulfide 1 mM). In the latter study sulfide was oxidized although internal levels of sulfide and succinate were high. These data indicate that sulfide oxidation in the presence of oxygen is possible even when aerobic metabolism is inhibited by sulfide. Previously we demonstrated that mitochondrial sulfide oxidation in A. marina is sulfide insensitive and assumed the existence of an alternative terminal oxidase (Volkel & Grieshaber 1994). This sulfide detoxification system enables the worm to remove sulfide quickly from its body and to rapidly return to an aerobic metabolism.

Conclusions

The lugworm Arenicola marina can be exposed to considerably high sulfide concentrations during tidal exposure. Sulfide concentrations can also increase sporadically in some burrows during high tide. Under habitat conditions the lugworm seems to exhibit the same mechanisms of sulfide tolerance as under experimental conditions. In the presence of oxygen, sulfide entering the body of A. marina is oxidized to thiosulfate. During high tide, therefore, an increase of sulfide in the body is prevented. Under hypoxic conditions during tidal exposure sulfide cannot be oxidized and sulfide concentrations in the coelomic fluid increase. The accumulation of sulfide together with hypoxia prevent aerobic metabolism and the lugworm switches to anaerobiosis. The lower coelomic pH as compared to burrow water (B. Giebels unpubl. results) may keep internal sulfide concentrations in some lugworms below the external level. However, coelomic sulfide concentrations increase above external concentrations in many other specimens. When the tide returns and the burrows are immersed sulfide in the body of A. marina is rapidly oxidized, which enables the worm to return to aerobic metabolism. After 1 h of recovery in normoxic seawater sulfide, succinate and thiosulfate have reached the same level as at the beginning of low tide, indicating that 1 high tide is more than sufficient for recovery from sulfide stress during tidal exposure.

Acknowledgements. We thank Silke Jakob for skilful technical assistance. Thanks are also due to our Düsseldorfer colleagues for their help in collecting the lugworms. K.H. is a fellow of the Konrad-Adenauer Foundation. Financial support from the Bundesminister für Forschung und Technologie (DYSMON 03P0123B) is also acknowledged.

LITERATURE CITED


Völkel S, Grieshaber MK (1992) Mechanisms of sulfide tolerance in the peanut worm Sipunculus nudus (Sipunculida) and in the lugworm Arenicola marina (Polychaeta). J comp Physiol 162B:469–477

This article was submitted to the editor.