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Winter and summer metabolic rates of Arctic amphipods. Preliminary results

INTRODUCTION

The metabolic rates of terrestrial and marine Arctic invertebrates is fairly well known (Remmert, 1980; Opaliński, 1982; Klekowski & Opaliński, 1986, 1989, 1990; Opaliński & Klekowski, 1989; Opaliński & Węśławski, 1989a, b; Piepenburg *et al.*, 1995). However, the above cited literature deals with the summer - Polar Day period, and data on the winter metabolic rates are rather rare. Scarce literature on the planktonic and littoral Antarctic crustaceans (Rakusa-Suszczewski & Klekowski, 1973; Rakusa-Suszczewski, 1990) indicate that winter metabolic rates are lower than summer ones, which is consistent with the generally accepted rule on of the temperature influence on animal metabolism (Prosser, 1973; Duncan & Klekowski, 1975). The aim of the present paper is to answer the question: How does the Arctic marine invertebrate metabolism changes in the course of winter to summer?

METHODS

Animals

Four species of marine littoral and shallow sublittoral amphipod crustaceans were the object of the study (*Gammarus oceanicus*, *G.setosus*, *Anonyx sarsi*, *A. nugax*). All four are common and abundant in the coastal waters of Spitsbergen (Węśławski, 1993a, 1994).

G. oceanicus is an opportunistic omnivorous, Atlantic - boreal species. It occurs in the littoral from the Baltic Sea on the south to Spitsbergen on the north. In Horsund Fjord it is most common between 0 and 2 m depth among stones. Natural temperatures in its environment range from -1.8°C in the winter and spring to $+8^{\circ}\text{C}$ in August in the peak of summer. Salinity ranges from 5 to 34 ppt (Węśławski, 1994). *G.setosus* is an Arctic - boreal species of circumpolar distribution, common throughout the Svalbard

archipelago. This species is more cold water compared to the previous one, and is more often found in the inner fjord basins and eastern coast of archipelago, but generally its environmental conditions are the same as for the *G. oceanicus* (Węśławski, 1994).

A. sari is necrophagic and carnivorous, Arctic - boreal species. It occurs on soft bottom, between 2 and 30 m depth, where temperature is not exceeding +5° C, and salinity is above 15 ppt.

A. nugax is necrophagic and carnivorous, also an Arctic - boreal species. It occurs on the soft bottom, below 20 m depth, down to 200 m in fjords. Temperature and salinity in its environment stays below +3.5° C and over 33 ppt (Węśławski, 1994). More on the ecology of these species can be found in Opaliński & Węśławski (1989a, b), and Klekowski & Węśławski (1991).

Experimental animals were collected in the intertidal and shallow sublittoral zone in Isbjornhamna, Hornsund Fjord, near the Polish Arctic Station in Hornsund (Southern Spitsbergen, Svalbard, 77°N, 15°E) - Figure 1. A detailed description of the local environment might be found in Węśławski (1993b). Animals were collected with the use of a light dredge between February 1985 and July 1985 (Table 1).

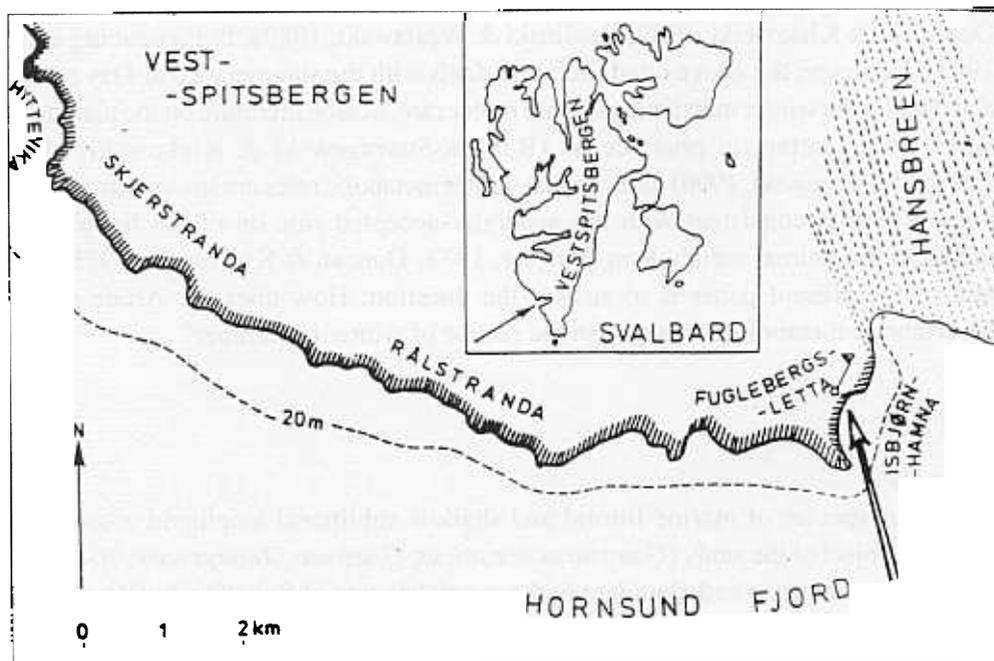


Fig. 1. Hornsund Fjord, Spitsbergen. The arrow indicate place of animal sampling.

Measurement methods

Animals were transported to the laboratory in the Polar Station and placed in aquaria for 5 to 12 hours to adapt for measurement conditions. Oxygen consumption

Table 1. Oxygen consumption and metabolic rate in Spitsbergen amphipods. n - number of measurements, T - temperature of measurements (°C), LC - animal body length (mm), W - animal wet weight (mg), R - oxygen consumption (cubic mm per individual per hour), MR - metabolic rate (cubic mm per mg wet wt per hour). Mean values ± Standard Error.

| Date | n | T | LC | W | R | MR |
|---------------------------|----|------|------------|--------------|-------------|---------------|
| <i>Gammarus oceanicus</i> | | | | | | |
| Feb. 24 | 8 | -1.0 | 17.6 ± 2.6 | 128.5 ± 56.3 | 20.6 ± 8.8 | 0.189 ± 0.026 |
| March 28 | 8 | -1.0 | 26.1 ± 1.3 | 284.6 ± 38.2 | 40.6 ± 4.0 | 0.164 ± 0.029 |
| May 16 | 9 | -1.0 | 23.2 ± 1.5 | 227.0 ± 35.7 | 39.6 ± 3.9 | 0.177 ± 0.035 |
| May 20 | 9 | 0.0 | 27.9 ± 1.4 | 360.3 ± 53.6 | 46.8 ± 3.7 | 0.146 ± 0.018 |
| May 24 | 9 | -1.6 | 18.5 ± 1.6 | 132.6 ± 32.0 | 20.2 ± 3.3 | 0.175 ± 0.016 |
| June 2 | 4 | -1.0 | no data | 268.0 ± 40.0 | 38.0 ± 6.4 | 0.149 ± 0.022 |
| June 6 | 6 | +1.0 | no data | 231.2 ± 59.3 | 32.1 ± 7.7 | 0.140 ± 0.010 |
| June 12 | 5 | +1.0 | 26.0 ± 1.6 | 244.0 ± 30.2 | 22.4 ± 3.3 | 0.147 ± 0.014 |
| July 4 | 25 | +2.0 | no data | 249.4 ± 28.1 | 37.4 ± 4.6 | 0.150 ± 0.010 |
| <i>Gammarus setosus</i> | | | | | | |
| Feb. 24 | 22 | -1.6 | 19.5 ± 1.3 | 149.7 ± 28.5 | 20.5 ± 4.4 | 0.126 ± 0.012 |
| Feb. 24 | 21 | 0.0 | 23.1 ± 1.2 | 245.7 ± 27.8 | 26.1 ± 2.7 | 0.120 ± 0.011 |
| March 26 | 11 | -1.0 | 29.9 ± 0.9 | 409.6 ± 39.1 | 52.7 ± 4.2 | 0.134 ± 0.013 |
| May 28 | 8 | -1.0 | 28.7 ± 2.3 | 405.7 ± 84.4 | 42.3 ± 7.8 | 0.120 ± 0.016 |
| July 8 | 26 | +2.0 | no data | 61.0 ± 8.0 | 10.9 ± 0.9 | 0.187 ± 0.013 |
| July 15 | 5 | +3.0 | no data | 89.0 ± 35.0 | 18.7 ± 6.9 | 0.212 ± 0.055 |
| <i>Anonyx sarsi</i> | | | | | | |
| Feb. 24 | 33 | -1.6 | 19.4 ± 0.7 | 250.0 ± 22.0 | 48.2 ± 13.3 | 0.156 ± 0.013 |
| Feb. 27 | 5 | -1.0 | 17.6 ± 1.7 | 180.0 ± 35.7 | 38.2 ± 11.2 | 0.220 ± 0.043 |
| April 2 | 5 | -1.6 | 18.8 ± 0.9 | 266.4 ± 24.3 | 46.2 ± 4.1 | 0.177 ± 0.017 |
| May 16 | 5 | -1.6 | 19.6 ± 0.8 | 292.0 ± 21.8 | 47.0 ± 5.2 | 0.159 ± 0.010 |
| May 20 | 4 | 0.0 | 15.8 ± 1.6 | 155.0 ± 42.5 | 23.8 ± 7.4 | 0.149 ± 0.010 |
| May 20 | 10 | +2.0 | 19.2 ± 1.6 | 219.2 ± 31.9 | 39.5 ± 6.9 | 0.176 ± 0.025 |
| May 24 | 12 | -1.6 | 17.8 ± 1.4 | 216.1 ± 40.9 | 21.4 ± 3.5 | 0.113 ± 0.019 |
| <i>Anonyx nugax</i> | | | | | | |
| April 12 | 9 | +3.0 | 29.2 ± 1.6 | 699 ± 114 | 46.7 ± 5.2 | 0.078 ± 0.011 |
| June 5 | 23 | +1.0 | no data | 652 ± 72 | 56.6 ± 6.0 | 0.085 ± 0.005 |
| July 3 | 21 | +2.0 | no data | 549 ± 70 | 44.6 ± 3.9 | 0.095 ± 0.008 |
| July 15 | 28 | +3.0 | no data | 658 ± 51 | 56.8 ± 5.7 | 0.086 ± 0.006 |

was measured using closed vessels method. Oxygen concentration in vessels was measured with the oxygen sensor OXI 57 (WTW, Germany).

Single animals were examined at natural environmental temperatures, controlled within an accuracy of 0.5° C. Erlenmeyer flasks of 200 cm were used as the respirometric vessels. The present paper is based on the data from 338 measurements. Details on sampling dates and the number of measurements for each species are given in Table 1.

As the criterion for the seasonal change from winter to summer, we took the rise of temperature in surface coastal waters from -1.88°C (winter) to -1.0°C (Węśławski & Adamski, 1987; Węśławski *et al.*, 1988; Węśławski, 1993b). In 1985 this took place in the turn of May to June, when the maximum of chlorophyll 'a' abundance was observed (Węśławski *et al.*, 1988).

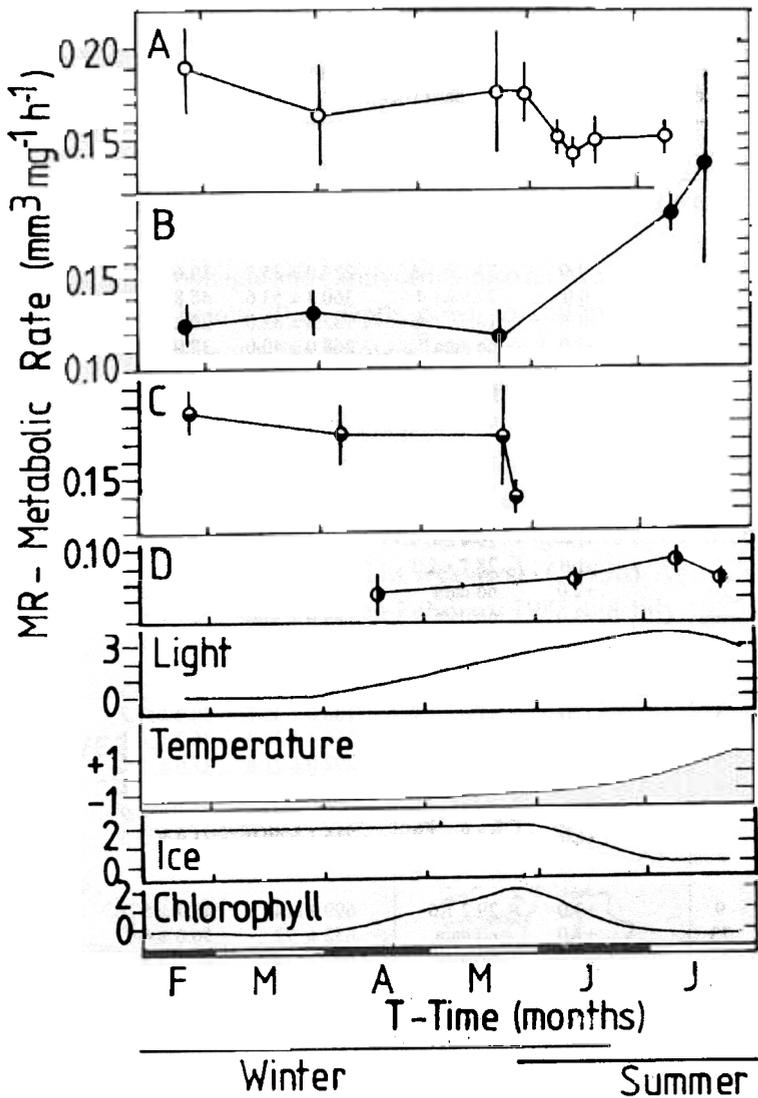


Fig. 2. Seasonal changes of metabolic rate in Spitsbergen coastal water amphipods and seasonal changes of some environmental factors in Hornsund Fjord coastal waters. A - *Gammarus oceanicus*, B - *Gammarus setosus*, C - *Anonyx sarsi*, D - *Anonyx nugax*. Light - in arbitrary units (0 - polar night, 1 - night longer than day, 3 - day longer than night, 4 - polar day), Temperature in °C, Ice - in arbitrary units (0 - fast ice, 1 - pack ice, 2 - open water), Chlorophyll - in mg m. Environmental data after Węśławski *et al.* (1988).

RESULTS

Gammarus oceanicus

The metabolic rates were lowest at the turn of the seasons (June 6th) and reached 0.140 mm/mg/h, while highest values were observed in the peak of the winter (February 25th) and reached 0.189 mm/mg/h, which makes a statistically important difference (Table 1). The *G. oceanicus* metabolic rates decrease steadily from winter to summer,

Table 2. Comparison of metabolic rate (MR - cubic mm per mg wet wt per hour) of Spitsbergen amphipods in winter and in summer. T - temperature (°C), n - number of measurements, W - animal wet weight (mg), R - animal oxygen consumption (cubic mm per individual per hour), MR - metabolic rate (cubic mm per mg wet wt per hour). Mean values \pm Standard Error.

| Season (date) | T | N | W | R | MR |
|----------------------------|--------------|----|------------------|----------------|-------------------|
| <i>Gammarus oceanicus</i> | | | | | |
| Winter (Feb.24 - June 2) | -1.6 to 0.0 | 58 | 220.3 \pm 18.6 | 32.3 \pm 2.2 | 0.171 \pm 0.009 |
| Summer (June 2 - July 4) | -1.0 to +2.0 | 31 | 254.3 \pm 24.2 | 36.9 \pm 3.8 | 0.148 \pm 0.014 |
| Winter + Summer | -1.6 to +2.0 | 83 | 233.2 \pm 15.5 | 34.5 \pm 2.1 | 0.165 \pm 0.007 |
| <i>Gammarus setosus</i> | | | | | |
| Winter (Feb.24 - May 20) | -1.6 to 0.0 | 62 | 260.7 \pm 22.7 | 30.9 \pm 2.6 | 0.124 \pm 0.006 |
| Summer (July 8 - July 15) | +2.0 to +3.0 | 39 | 130.1 \pm 28.6 | 17.7 \pm 2.6 | 0.206 \pm 0.015 |
| Winter + Summer | -1.6 to +3.0 | 93 | 193.5 \pm 18.2 | 24.4 \pm 2.0 | 0.159 \pm 0.009 |
| <i>Anonyx sarsi</i> | | | | | |
| Winter (Feb. 24 - May 20) | -1.6 to +2.0 | 81 | 225.4 \pm 13.1 | 34.7 \pm 2.3 | 0.165 \pm 0.008 |
| <i>Anonyx nugax</i> | | | | | |
| Winter (April 12 - June 5) | +1.0 to +3.0 | 32 | 663.5 \pm 82.3 | 49.2 \pm 5.3 | 0.082 \pm 0.007 |
| Summer (June 5 - July 15) | +1.0 to +3.0 | 72 | 580.3 \pm 41.9 | 47.8 \pm 3.5 | 0.089 \pm 0.003 |
| Winter + Summer | +1.0 to +3.0 | 82 | 568.2 \pm 65.9 | 43.6 \pm 4.7 | 0.087 \pm 0.004 |

with a clear drop in the beginning of June (Fig. 2a). The metabolic rates of *G. oceanicus* calculated for the winter period (Feb. 24th - June 2nd) were higher than for summer (June 6th - July 4th), and changed respectively from 0.171 to 0.148 mm/mg/h (Table 2). This clear picture gets worse when we compare not the mean values, but regression lines calculated for the oxygen consumption dependence on body mass (Fig. 3). It was found that large animals of weight over 200 mg showed the same level of metabolic rates in winter and in summer, while small animals of body mass below 100 mg showed higher winter metabolic rates compared to summer ones. The regression lines show the same pattern: intercept 'a' is high in winter animals (0.73), and is low in summer (0.14), with a very high regression coefficient (1.00) (Table 3).

Gammarus setosus

The reverse situation was observed in *Gammarus setosus*. The lowest metabolic rates were observed in the middle of the winter (Feb. 24th), and the highest in late summer (July 15th): respectively 0.120 and 0.212 mm/mg/h (Table 1). After the relatively stable winter metabolic rates, the summer values increase sharply (Fig. 2b). The mean metabolic rate of *G. setosus* calculated for the summer period is 0.206 mm/mg/hr (July 8th - July 15th), almost twice as high as the winter one: 0.124 mm/mg/hr (Feb. 24th - May 20th) (Table 2). Relation of oxygen uptake to body weight in *G. setosus* was similar to that in *G. oceanicus*, and large animals (100-200 mg weight) expressed almost the same metabolic rates in winter and in summer (Fig. 4). In small animals (below 100 mg) metabolic rates were higher in summer than in winter (contrary to the results with *G. oceanicus*). The intercept „a” in regression equations shows

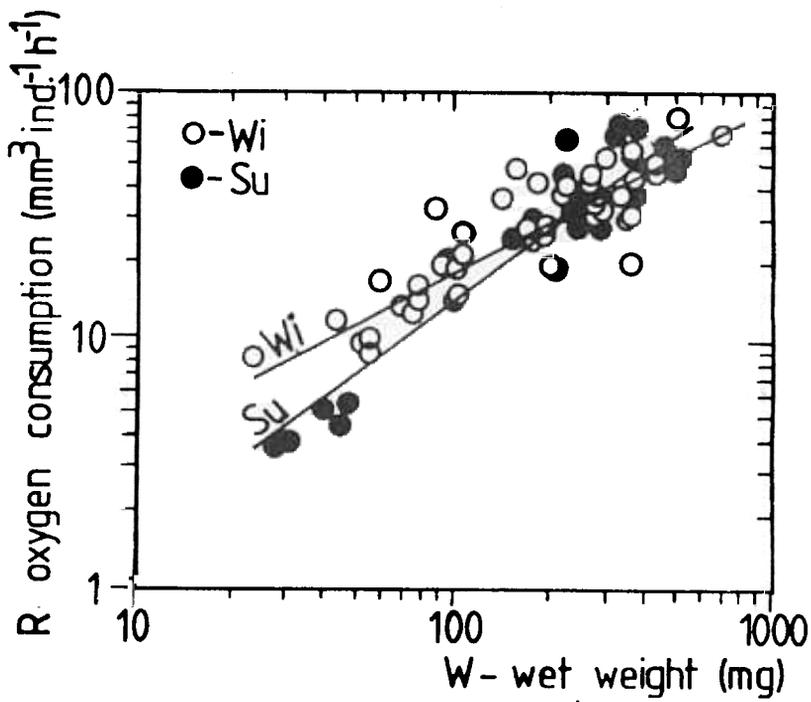


Fig. 3. The dependence between animal wet weight (W) and oxygen consumption (R) in *Gammarus oceanicus* from Hornsund Fjord in winter (Wi) and in summer (Su). See also Table 3.

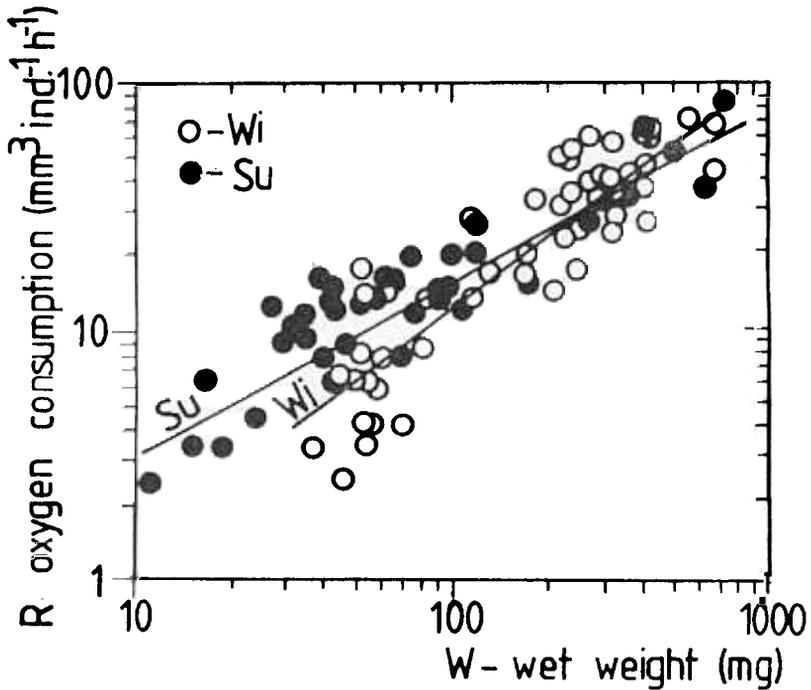


Fig. 4. The dependence between animal wet weight (W) and oxygen consumption (R) in *Gammarus setosus* from Hornund Fjord in winter (Wi) and summer (Su). See also Table 3.

Table 3. The dependence between oxygen consumption (R - cubic mm per individual per hour) and bod wet weight (W - mg) in Spitsbergen amphipods in winter and in summer. General formula $R = a W^b$. n - number of measurements.

| Season | n | W | R | Regression intercept a | Regression coefficient b | Regression r^2 |
|---------------------------|----|-----|------|------------------------|--------------------------|------------------|
| <i>Gammarus oceanicus</i> | | | | | | |
| Winter | 58 | 220 | 32.6 | 0.73 | 0.71 | 0.7904 |
| Summer | 31 | 254 | 36.9 | 0.14 | 1.00 | 0.8702 |
| Winter + Summer | 82 | 233 | 24.5 | 0.42 | 0.81 | 0.8110 |
| <i>Gammarus setosus</i> | | | | | | |
| Winter | 62 | 261 | 30.9 | 0.13 | 0.98 | 0.8400 |
| Summer | 39 | 130 | 17.1 | 0.80 | 0.66 | 0.8290 |
| Winter + Summer | 93 | 193 | 34.5 | 0.43 | 0.77 | 0.7904 |
| <i>Anonyx sarsi</i> | | | | | | |
| Winter | 81 | 225 | 34.7 | 0.36 | 0.83 | 0.5375 |
| <i>Anonyx nugax</i> | | | | | | |
| Winter | 32 | 663 | 49.2 | 0.57 | 0.69 | 0.6708 |
| Summer | 72 | 580 | 47.8 | 0.16 | 0.90 | 0.9109 |
| Winter + Summer | 82 | 568 | 43.6 | 0.16 | 0.89 | 0.9435 |

low values in winter (0.13) and in summer (0.80), with regression coefficients $b=0.98$ for winter and $b=0.66$ in summer (Table 3).

Anonyx sarsi

The metabolic rates of this species were measured in winter only (Feb. 24th - May 24th). They ranged between 113 to 220 mm/mg/h (Table 1). From the middle of winter to the late winter a slight decrease of *A. sarsi* metabolism was observed (Fig. 2c).

Table 4. Metabolic rate (MR - cubic mm per mg wet wt per hour), the dependence between oxygen consumption (R - cubic mm per individual per hour) and body wet weight (W - mg) (general formula $R = aW^b$) in some polar invertebrates in the winter and in the summer. The difference between winter and summer metabolic rate (W/S) is indicated in percent.

| Species | Season | Temperature °C | MR | W/S % | Regression | | Author |
|---------------------------|--------|----------------|-------|-------|------------|------|------------------------------------|
| | | | | | a | b | |
| <i>Gammarus oceanicus</i> | winter | -1.6 to 0.0 | 0.171 | - | 0.73 | 0.71 | present paper |
| | summer | -1.0 to +2.0 | 0.148 | -13 | 0.14 | 1.00 | present paper |
| <i>Gammarus setosus</i> | winter | -1.6 to 0.0 | 0.124 | - | 0.13 | 0.98 | present paper |
| | summer | +2.0 to +3.0 | 0.206 | +66 | 0.80 | 0.66 | present paper |
| <i>Anonyx nugax</i> | winter | +1.0 to +3.0 | 0.082 | - | 0.57 | 0.69 | present paper |
| | summer | +1.0 to +3.0 | 0.089 | +8 | 0.16 | 0.90 | present paper |
| <i>Paramoera walkeri</i> | winter | -1.9 | 0.050 | - | 0.09 | 0.71 | Klekowski et al 1973 |
| | summer | -1.2 | 0.090 | +80 | 0.10 | 0.65 | Rakusa-Suszczewski, Klekowski 1973 |
| <i>Euphausia superba</i> | winter | -1.8 | 0.060 | - | 0.08 | 0.95 | Rakusa-Suszczewski 1990 |
| | summer | 0.0 | 0.074 | +23 | 0.10 | 0.95 | Rakusa-Suszczewski 1990 |

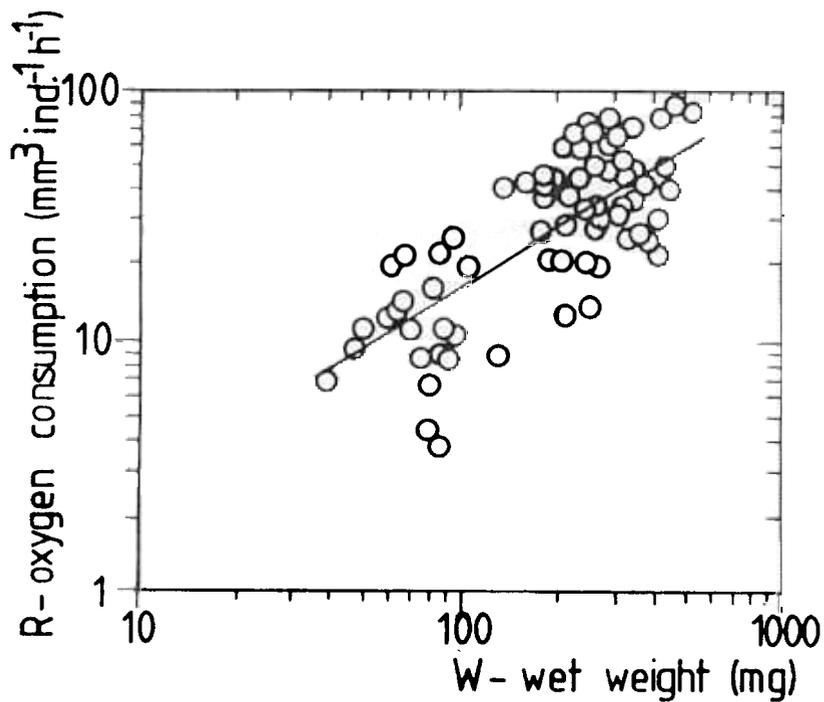


Fig. 5. The dependence between animal wet weight (W) and oxygen consumption (R) in *Anonyx sarsi* from Hornsund Fjord in winter. See also Table 3.

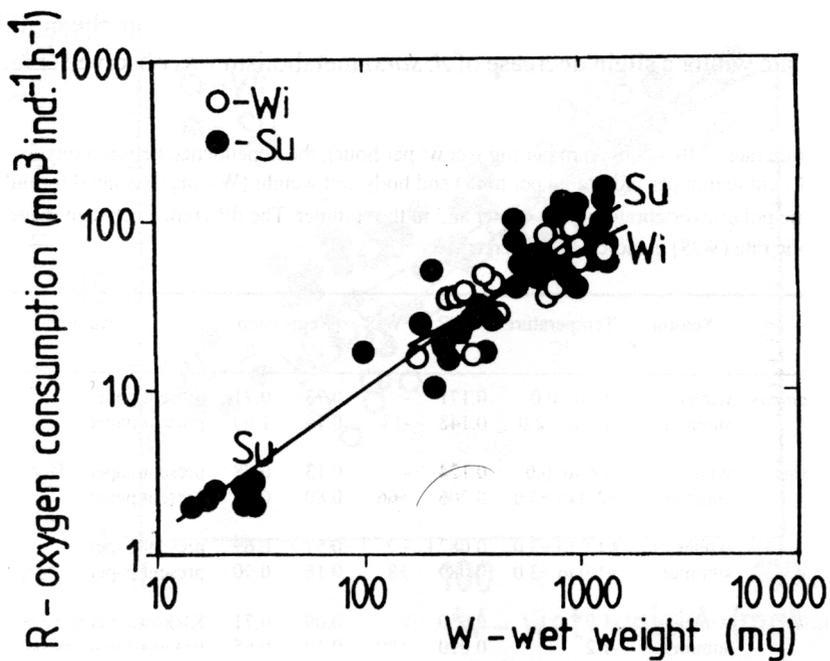


Fig. 6. The dependence between animal wet weight (W) and oxygen consumption (R) in *Anonyx nugax* from Hornsund Fjord in winter (Wi) and summer (Su). See also Table 3.

The mean metabolic rate of *A. sarsi* in winter was 0.165 mm /mg/h (Table 2). The correlation of oxygen consumption and body mass in *A. sarsi* is illustrated in Figure 5 and Table 3.

Anonyx nugax

The lowest metabolic rates of this species were found in winter (April 12th) and the highest in summer (July 5th), 0.078 and 0.095mm mg/h respectively (Table 1). The mean summer metabolic rates of *A. nugax* (June 5th and July 15th) were similar to the mean winter values (April 12th - June 5th) and ranged respectively from 0.089 to 0.082 mm /mg/h (Table 2, Fig. 2d). The relation of oxygen consumption to body weight in *A. nugax* is shown by the differences in winter and summer values of intercepts 'a' (0.57 to 0.16) and coefficients 'b' : 0.69 and 0.9 respectively (Table 3), however the measurements of adult animals showed almost identical regression lines (Fig. 6). The differences of regression parameters come from the low metabolic rates of juvenile specimens, observed in summer.

Presented results show all possible metabolic reactions connected with the change of seasons: the metabolism may drop from winter to summer (*G. oceanicus*), it may grow (*G. setosus*) or remain stable (*A. nugax*). The observed differences of metabolic rates in winter and summer are the effect of juvenile animals metabolism, the adults expressed an even level of metabolism from winter to summer.

DISCUSSION

Seasonal change from winter to summer is connected with the change of important environmental parameters like temperature, salinity, light, and food resources. The High Arctic region shows especially drastic changes of these parameters (Węśławski *et al.*, 1988; Węśławski, 1994).

How may poikilothermic animals react to those changes?

The increase of poikilothermic animals metabolic rates from winter towards summer seems to be a natural reaction caused by an increase in water temperature (Klekowski *et al.*, 1973; Opaliński, 1979a,b; Opaliński & Klekowski, 1992), increased sun radiation (Opaliński & Klekowski, 1991), food resources (Opaliński, 1991; Percy, 1993; Vernberg & Vernberg, 1993; Chappelle *et al.*, 1994), or the salinity drop caused by the ice melt (Klekowski & Opaliński, 1993). Such metabolic rates, which increase from winter to summer, have been observed among Antarctic animals. *Paramoera walkeri* (Amphipoda) metabolism expressed a summer increase of metabolic rates of 80% compared to winter values, and antarctic krill (*Euphausia superba*) showed a 23% increase (Table 4).

A rate of 8% increase of the metabolic rate from winter towards summer in *A. nugax* was statistically unimportant, while *G. oceanicus* expressed a decrease of metabolic rate of 13% (Table 4).

Further analysis is needed for answering the question of what are the specific external and internal factors, changeable from winter to summer, that might be

responsible for the metabolic rates seen in Arctic littoral amphipods. This will be the subject of future studies.

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