Observed on the measurement and interpretation of clearance rate variations in suspension-feeding bivalve shellfish

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ABSTRACT: Mussels *Mytilus edulis* (35 ± 2 mm shell length) were fed on cultured *Isochrysis galbana* in a flow-through system, and measures were undertaken to quantify separate effects of inflow cell concentration, outflow cell concentration, percentage reduction (i.e. recycling), and flow rate and/or feeding history on clearance rate measures (CR; l h⁻¹). Findings identify the following features of physiological regulation: (1) a lag phase in the feeding response to changes in food concentration, of around 30 min; (2) a trigger level in algal cell (chlorophyll [chl] a) volume/concentration (4000 cells ml⁻¹, 0.5 µg l⁻¹ chl a) below which filtering in most mussels ceases; (3) saturation reduction, or satiation, resulting in reduced CR and valve closure after feeding for over 2 h at 30 000 or more cells ml⁻¹ (~6 µg l⁻¹ chl a); and (4) at food levels between Features 2 & 3, CR increases to maximal rates (ca. 2 to 2.5 l h⁻¹ ind⁻¹ or 6 to 7.5 l⁻¹ h⁻¹ g⁻¹ dry weight). Our findings also help resolve uncertainties associated with the measurement and interpretation of clearance rate variations. These uncertainties are associated with: differences that result from analyses of feeding responses to different components of the available seston; whether variations in seston availability are physiologically relevant; analyses of individual shellfish as compared with average responses computed for >1 ind.; and comparisons between short-term and integrated measures. We think it is important to consider whether measures of feeding are really required per individual in real time, and we stress that population averaging and temporal integration are required for the scaling up of results when simulating ecological interrelations between shellfish populations and their environments.

KEY WORDS: Filter feeding · Suspension feeding · Clearance rate · *Mytilus edulis* · Bivalve shellfish · Flow-through · Physiological regulation · Supply rate

INTRODUCTION

The current understanding of filter feeding and its measurement in bivalve shellfish is beset by a number of controversies, and several reviews have summarised the current knowledge with a view to resolving these controversies (Jorgensen 1996, Bayne 1998, Riisgård & Larsen 2000, 2001, Riisgård 2001a,b). Riisgård (2001b) highlighted methodological shortcomings and possible misinterpretations of data, suggesting guidelines to help obtain reliable data for measures of the rate with which suspension-feeding bivalves remove particles from seawater (clearance rate, CR). Conclusions on the through-flow chamber method were collated in an appendix by Larsen (2001). These guidelines stimulated further work on the comparison of 3 different methods of measuring CR (Petersen et al. 2004), resulting in more comment and debate (Bayne 2004, Petersen 2004, Riisgård 2004). More recently Filgueira et al. (2006) again addressed the design and validation of the flow-through chamber method for CR measurements.

There is general recognition of the need to better understand the intriguing complexity of feeding processes in bivalve shellfish, which shows remarkable regulation and morphological adaptation of particle processing mechanisms within and between species, respectively (e.g. Beninger & St-Jean 1997, Beninger et al. 1997, 2008). Here, we address some key environ-
mental factors and behavioural issues affecting variations in CR that are observed within species. Using mussels fed on cultured algae in a flow-through system, experiments were undertaken to quantify separate effects of inflow cell concentration, outflow cell concentration, percentage reduction (i.e. recycling) and flow rate and/or feeding history on CR measurements. Notable reductions or cessation of feeding were observed at both low (below trigger level) and high (through satiation) food concentrations. The interpretation of results is intended to help researchers ensure that the measures they use are ‘fit for purpose’, including whether averaging and integration may be required to simulate ecological interrelations between shellfish populations and their environments.

MATERIALS AND METHODS

Mussels Mytilus edulis L., at shell lengths of 35 ± 2 mm, were collected at Great Bull Point (50°36.8’N, 3°26.0’W), near the mouth of the Exe estuary, SW England (Hilbish et al. 2002). After removal of any epibiotic growth, individual animals were numbered and then maintained in a 7 l vessel linked to a 3000 l system of circulating seawater at 14 ± 1°C and 34 ± 3°C. 2 mm, were collected at Great Bull Point (50° 36.8’ N, 3° 26.0 W), near the mouth of the Exe estuary, SW England (Hilbish et al. 2002). After removal of any epibiotic growth, individual animals were numbered and then maintained in a 7 l vessel linked to a 3000 l system of circulating seawater at 14 ± 1°C and 34 ± 1°C for a period of 10 d before the experiments. During this time and between experiments, mussels were fed on a maintenance ration (approximately 2000 cells ml⁻¹) of the microalga Isochrysis galbana. In the feeding experiments, 14 ind. were each placed in a chamber within a flow-through raceway system as described by Hawkins et al. (1996) to measure CR (litres of water cleared of particles h⁻¹) over separate ranges of algal cell concentration, water flow rate and percentage reduction in food particles, as water passed from inflow to outflow. Mussels were orientated with their inhalant apertures towards the incoming flow. The volume of the chambers was approximately 200 ml, and experimental flow rates through each chamber ranged from 30 to 400 ml min⁻¹, which for comparative purposes equate to maximal current speeds of <0.003 m s⁻¹ through those chambers, measured with a Valeport 800-175 electromagnetic current meter. As particle sedimentation within the mussel chambers was found to be negligible, 2 ‘control’ chambers were maintained without mussels. At prescribed intervals, 20 ml aliquots from the outflows of the 2 control chambers, and from chambers containing an individual mussel, were analysed using a Coulter Multisizer II (100 µm diameter orifice) with Coulter Accucomp software. Both the total number and volume of particles between 3 and 6 µm equivalent spherical diameter, which spanned the size range of I. galbana, were measured and recorded in replicate 0.5 ml samples. We concede that using an algal monoculture as food may produce different responses from those using natural seston, but it was chosen here as a more consistent and comparable food source to study the experimental variables.

Clearance rate was determined by 2 equations that are in common use:

\[ CR = \frac{Fl \times [(C_i - C_o)/C_i]}{H_{20862}} \]  
\[ CR = \frac{Fl \times ([C_i - C_o]/C_o)}{H_{20862}} \]

where Fl is the flow through the chamber, and C_i and C_o are the number of particles per millilitre within the inflow and outflow, respectively, and

where the outflow concentration is used as the denominator, and is normally used in steady-state conditions (Hildreth & Crisp 1976, Riisgård 2001b). Concentrations measured in the outflows from control chambers containing experimental mussels. To avoid errors associated with potential recycling, we have normally restricted calculations of CR to mussels effecting a reduction in particles between inflow and outflow of 5 to 30% (Hawkins et al. 1999). It should be noted that, for analytical purposes, greater reductions have in some cases deliberately been induced here.

Experiments were designed to resolve any separate effects of inflow cell concentration, outflow cell concentration, percentage reduction (recycling) and/or flow rate (current speed) on CR. To achieve this, 3 experimental phases were undertaken:

1) Expt 1. To confirm the time course in response of CR to stepwise increases in food supply (cell concentration), CR was measured simultaneously in 14 mussels after 5, 30, 60 and 180 min following a change from the maintenance ration to between 4000 and 20 000 cells ml⁻¹. At each cell concentration, flow rates were adjusted to give between 5 and 30% reduction in algal cell concentration between inflow and outflow.

2) Expt 2. To resolve the effects of flow rate, percentage reduction (recycling) and outflow cell concentration on CR, a series of experiments was undertaken using constant inflow concentrations (5, 10, 20, 30 and 50 000 cells ml⁻¹), and flow rates were varied to give mean percentage reductions in cell concentration between inflow and outflow of between 5 and 65%. On the basis of findings from our first experimental phase above, and as explained in the results, CR measures were standardised after 30 min under each condition.

3) Expt 3. To assess the relative importance of inflow concentration versus outflow concentration, mussel chamber outflow concentrations were maintained as close as possible to 10 000 cells ml⁻¹, applying the same flow rates as in Experiment 2 above, and varying the algal dosing rate (inflow concentration) to give the
same percentage reductions in cell number as had been recorded at each flow rate during Experiment 2. Because feeding history may affect the feeding responses of bivalve shellfish, care was taken to limit the time that each experimental mussel was exposed to high food rations, thus minimising any satiation that may have led to reduced CRs. On a day to day basis, experiments administering high rations were alternated with those at low rations. At each sampling point, individual mussels that were closed and not feeding, or which were producing pseudofaeces, were noted.

Chlorophyll $a$ ($chl\ a$) was measured in filtered aliquots from the outflows of each control tray. Samples were collected on separate 47 mm Whatman GF/F filters that were frozen at $-20^\circ$C before using standard procedures for acetone extraction and fluorometric analysis (Holm-Hansen et al. 1965).

Following feeding experiments, all soft tissues were excised from each mussel and dried at $60^\circ$C before weighing to a constant total soft tissue dry weight (mean ± 2 SE: 0.168 ± 0.019 g). Clearance rates presented here were not weight standardised. Instead, the stated mean total soft tissue dry weight affords the opportunity for others to do so at their discretion.

To resolve any effects of inflow or outflow cell concentrations, percentage reduction (recycling) and/or flow rates, the differences between mean CR measures were first tested for statistical significance with 1-way analysis of variance (ANOVA), applying a Bonferroni correction (Sokal & Rohlf 1995). Upon any suggested difference, specific pair-wise comparisons were undertaken using a Tukey’s honestly significant difference test. These analyses were carried out using SYSTAT 11 (Systat Software Inc.).

RESULTS

Experiment 1

The effects on CR of stepwise increases in *Mytilus edulis* cell concentrations after 5, 30, 60 and 180 min following change from the maintenance ration to between 4000 and 200 000 cells ml$^{-1}$ are summarised in Fig. 1. These CR values were measured under conditions when cell reductions ranged from 5 to 30%, and were computed using Eq. (1), with inflow concentration as the denominator. Findings illustrate that reduced average clearance rates were often associated with lower numbers of actively feeding animals. At low cell concentrations averaging <4000 cells ml$^{-1}$ (less than ~0.5 µg chl $a$ l$^{-1}$), CR remained at <1 l h$^{-1}$ throughout the time course of up to 3 h, when low proportions of the mussels were actively feeding. At cell concentrations of 5000 to 200 000 cells ml$^{-1}$ (~0.6 to 25 µg chl $a$ l$^{-1}$), CR increased to values of between 1.93 and 2.38 l h$^{-1}$ after 30 min, compared to <1 l h$^{-1}$ below 4000 cells ml$^{-1}$ ($p < 0.01$). ‘Saturation reduction’ occurred after 180 min at high concentrations as evidenced by a higher CR of >2 l h$^{-1}$ at 30 000 cells ml$^{-1}$, compared with <1.5 l h$^{-1}$ at >100 000 cells ml$^{-1}$ ($p < 0.05$).

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**Fig. 1.** *Mytilus edulis*. Relationships between clearance rate (CR; mean ± 2 SE) and algal cell concentration after different time intervals at each food ration. (A,B,C,D) Responses after 5, 30, 60 and 180 min at each ration, respectively. Data are for all animals, including those not feeding. (●) number of animals feeding
**Experiment 2**

On the basis of the above results, further CR measures were standardised to 30 min after any experimental change in feeding conditions. Fig. 2 illustrates all CR measures for which the percentage reduction in cell concentration between inflow and outflow was >5%, and shows that average CR increases with cell concentration from 5000 (mean = 1.58) to around 20 000 cells ml⁻¹ (mean = 2.14) and then decreases to around 1.5 at higher concentrations of about 50 000 cells ml⁻¹. CR values at the lowest 3 concentrations were all significantly different from each other (p < 0.01), and that at 20 000 cells ml⁻¹ was significantly different from all others (p < 0.01). Fig. 3 shows how, for given flow rates, CR calculated using Eq (1) increased with percentage reduction in cell concentration between inflow and outflow. Flow rate appeared to be the dominant parameter, as slopes of relations between CR and percentage reduction increased with increasing flow. Also, at low flow rates, percentage reductions were high, whilst CR remained low. The same data plotted according to Riisgård (2001b) show CR values approaching a plateau as flow rates rise above 150 ml min⁻¹, at which point virtually all values fall between the lines representing 5 and 30% reductions in particles (Fig. 4). At these higher flows of >150 ml min⁻¹, the maximal recorded average CR of around 38 ml min⁻¹ (or 2.28 l h⁻¹) is similar to that predicted by Møhlenberg & Riisgård (1979) for *Mytilus edulis* of this size.

**Fig. 2.** *Mytilus edulis*. Relationships between clearance rate (CR) and algal cell concentration showing mean values (±2 SE) at each nominal concentration for all animals (with a >5% reduction in particles between inflow and outflow) and at all flow rates from Experiment 2.

**Fig. 3.** *Mytilus edulis*. Clearance rate (CR) versus percentage particle reduction, plotted for all animals at each nominal flow rate (shown in the legend, ml min⁻¹). The box shows values lying between 5 and 30% reduction in particle numbers.

**Fig. 4.** *Mytilus edulis*. Relationships between clearance rate (CR) and flow rate following Riisgård (2001b), to validate the flow-through chamber design. Lines are plotted to show CR (i.e. flow rate), and the limits of 5 and 30% reduction in particles. Horizontal dashed line: theoretical CR (2.28 l h⁻¹) derived for the size of *M. edulis* used here, from the equation CR = 7.45W^{0.66}, where W is g dry soft tissue (Møhlenberg & Riisgård 1979).

**Experiment 3**

Fig. 5 illustrates relationships between CR and flow rate or percentage reduction according to inflow and outflow concentrations, including any differences resulting from the use of Eqs. (1) versus (2). Results are compared for feeding conditions with constant inflow (Fig. 5A & B) and outflow (Fig. 5D & C) concentrations, all of about 10 000 cells ml⁻¹. Fig. 5 Panels A & B show that at low flow rates, differences between cell concentrations in inflows and outflows (percentage reductions) were high due to likely recycling of
the water, which results in CR calculated using Eq. (2) being much greater than CR calculated using Eq. (1). This may highlight the lack of ‘steady state’ conditions in this case. Also, the latter may be depressed, as the concentration of algal cells falls towards the trigger level. At high flow rates, percentage reductions were very low, and, consequently, there is little difference between CR calculated using Eqs. (1) & (2). In this case, the animals are filtering at maximal rates, but they are only clearing a small percentage of the total supply, as most of the flow is bypassing them (see Larsen 2001). As the formulae would suggest, differences in CR values resulting from the 2 equations are clearly proportional to the percentage reduction. For example, given a reduction of 20%, CR calculated using Eq. (2) will be 20% higher than CR derived from Eq. (1).

Results from feeding conditions with constant outflows at about 10 000 cells ml\(^{-1}\) (Fig. 5D & C) are similar to those with constant inflow concentrations (Fig. 5A & B), since, although inflow concentrations at low flow rates are markedly different, the percentage reductions, and hence CR values calculated using either Eq. (1) or (2), are similar whether measured with constant outflows or inflows. The same data plotted against elapsed time show that CR calculated using Eq. (1) was inversely related to inflow concentration, and again illustrates a dependency related to flow rates <150 ml min\(^{-1}\) (Fig. 6). The effect of feeding history is also illustrated here, as CR values are lower and more variable after feeding at higher inflow concentrations and for a longer time.

![Fig. 5. Mytilus edulis. Relationships between clearance rate (CR; mean ± 2 SE) and both flow rate and percentage reduction in particles, for 2 separate conditions: (A,B) maintaining a constant inflow concentration of 10 000 cells ml\(^{-1}\) and (C,D) maintaining a constant outflow concentration of 10 000 cells ml\(^{-1}\). CR values were derived from both Eq. (1) (■) and Eq. (2) (○), and cell concentrations are shown for both inflows (●) and outflows (○).](image)

![Fig. 6. Mytilus edulis. Interrelationships between mean clearance rates (CR; mean ± 2 SE) derived from Eq. (1) (●) and algal cell concentrations against time elapsed from food change (i.e. Expt 3). A constant outflow concentration of 10 000 cells ml\(^{-1}\) was maintained. Cell concentrations are shown for both inflows (●) and outflows (○), and flow rates for each measure are shown in bold italics.](image)
DISCUSSION

Our findings are consistent with ‘physiological regulation’ of bivalve filter feeding, according to indicators of such regulation suggested by Riisgård (2001a,b): (1) a lag phase was evident in the feeding response to changes in food concentration, that lag being between 5 and 30 min in the present study, when mussels were acclimated to a low food ration of approximately 2000 cells ml⁻¹ giving 0.25 µg l⁻¹ chl a; (2) a trigger level was reached in algal cell (chl a) concentration, below which filtering in most mussels ceases, and which, in this study, appeared to be around 0.5 µg l⁻¹ chl a; (3) saturation, or satiation, was reduced, resulting in reduced clearance rates and valve closure, after feeding for some time at high food rations, evidenced here after feeding for >2 h at ≥30 000 cells ml⁻¹ (~6 µg l⁻¹ chl a); and (4) at suitable flow rates and food levels between Points 2 & 3, CR increased to maximal rates, unless affected by other environmental conditions. There has been uncertainty over whether the transitions from no feeding to maximal CR, both at low and very high cell concentrations, entail abrupt stepwise events or a more gradual change resulting in a bell-shaped response of CR versus cell concentration. We suggest that different observations will result from measures based on instantaneous particle depletion by individuals, as compared with measures that average the responses of >1 ind., thus disguising significant inter-individual variability. Alternative methods may temporally integrate CR and other responses over longer time periods when measured as particle depletion within static seawater volumes (Widdows 1985, Petersen et al. 2004) or through the collection of faecal biodeposits (Cranford & Hargrave 1994, Hawkins et al. 1996, Cranford & Hill 1999, Navarro & Velasco 2003, Velasco & Navarro 2005). To illustrate this point, Fig. 7 compares differences between individual responses to food concentration in 3 mussels (Fig. 7A) with the averaged values for those same individuals (Fig. 7B).

It is clear that considerable variation in CR values can be produced by different methods of measurement. Problems have been recognised and recommendations advanced for the optimization and standardization of those methods, addressing chamber geometry, flow rate and the relative definitions of filtration rate and clearance rate (Riisgård 1977, Møhlenberg & Riisgård 1979, Widdows 1985, Riisgård 2001b). In the present study, one aim has been to re-assess our flow-through system as used by Hawkins et al. (1996, 1999, 2002) according to Riisgård’s (2001b) suggested protocol. Figs. 3, 4 & 5 illustrate how—providing the normal limits of flow (>150 ml min⁻¹) and percentage reductions (5 to 30%) are met—measures made with our system are entirely consistent with the guidelines proposed. Indeed, maximal CR values of around 2.5 l h⁻¹ described here for mussels of 0.168 g mean dry weight and 35 ± 2 mm shell length compare favourably with related measures of 2.29 l h⁻¹ from CR = 7.45W⁰.⁶⁶ (Møhlenberg & Riisgård 1979) and 2.42 from CR = 0.0012L^{2.14} (Kiørboe & Møhlenberg 1981), where W and L are dry soft tissue weight and shell length, respectively.

Since Riisgård’s (2001b) review, further methodological investigations have included inter-calibration by Petersen et al. (2004), comparing the flow-through method, the biodeposition method and the indirect (or clearance) method, which further stimulated criticism and debate (Bayne 2004, Petersen 2004, Riisgård 2004). Although this inter-calibration (Petersen et al. 2004) still left some uncertainty over the correct methodology and measurements, their suggested maximal CR of 3.6 l h⁻¹ is, in fact, close to estimates from early published equations, e.g. CR = 7.45W⁰.⁶⁶ (Møhlenberg & Riisgård 1979). Widdows (1985) showed that the different methods (static, flow-through and steady-state) and their relevant equations should produce similar CR values. Also, as we have highlighted above, the dif-

![Fig. 7](image-url)
ference in CR values using Eqs. (1) & (2) for a flow-through system is proportional to the percentage reduction in particles. A clear validation of 2 different flow-through systems was made by Filgueira et al. (2006). They investigated the fluid dynamics in individual cylindrical experimental (flow-through) chambers (ICEC) and then, with *Mytilus galloprovincialis*, established that the system complied with the requirements for accurate CR measurements as suggested by Riisgård (2001b). Their proposed protocol for validation—identifying phases where CR is dependent on flow rate, where it is independent of flow, and the transitional phases—allows simple analysis of any flow-through chamber, even those used in previous studies. An average of 20% (range 13 to 25) reduction in particles is recommended for valid measurements, and our Eq. (1) is deemed correct for the flow-through chamber method. Accuracy and validity of the method relies to some extent on a suitable ratio of animal size to chamber size, and the suggested scaling of the chamber by reference to the allometric length–volume relationship of mussels is both simple and practical. Filgueira et al. (2006) also validate a mesocosm system by comparison with results from the ICEC, so, overall, their work represents a major step forward in the debate on measurement of clearance rates in bivalves.

Figs. 5 & 6 show the difference between CR values using the 2 equations, and confirm that Eq. (1) is the better representation of true CR. However, care needs to be taken over the limits of flow rate and percentage reduction, and also feeding history. Results presented here validate our methods for measurement of clearance rates, and confirm that guidelines for the limits of percentage particle reduction suggested by some previous authors were correct (Hawkins et al. 1999, 2002, Larsen 2001, Filgueira et al. 2006).

Our findings for *Mytilus edulis* complement those illustrating regulatory feeding behaviour in the New Zealand green-lipped mussel *Perna canaliculus* (Hawkins et al. 1999) and the Chinese scallop *Chlamys farreri* (Hawkins et al. 2002). CR has been observed to increase with the availability of natural suspended sediments in turbid environments, up to experimental maxima of as much as about 80 mg total particulate matter l⁻¹ (e.g. Hawkins et al. 1996). However, particles suspended in turbid environments are dominated by silt, when chl a may only be present at up to about 3 µg l⁻¹ (e.g. Iglesias et al. 1992, Newell & Shumway 1993, Hawkins et al. 1996). As reported here for *M. edulis*, it is notable that maximum CRs recorded in species from temperate latitudes typically average up to about 8 l h⁻¹ g⁻¹ dry soft tissue and are most commonly observed in mussels feeding on cultured unicellular algae at concentrations affording up to about 5 µg chl a l⁻¹ (e.g. Møhlenberg & Riisgård 1979, Riisgård 1988, Clausen & Riisgård 1996, Jørgensen 1996, Hawkins et al. 1997). Alternatively, at higher concentrations of cultured unicellular algae, CR has frequently been observed to decrease in a negative exponential relation to increasing food availability (Winter 1978, Griffiths 1980, Iglesias et al. 1992, Arief & Bell-Dell-Young 1997, Hawkins et al. 1997, 1998, Yukihiro et al. 1998, Denis et al. 1999). This is again consistent with observations for *M. edulis* here, when saturation reduction, or satiation, resulted in reduced CRs and valve closure after feeding for >2 h at 30 000 or more cells ml⁻¹ (~6 µg chl a l⁻¹) (Fig. 1).
Three main points may be emphasised. Firstly, CR in suspension-feeding shellfish is highly flexible. Reductions from maximal rates occur, when costs of feeding (e.g. Widdows & Hawkins 1989) may outweigh potential returns, both at low particle availabilities and upon satiation at high food availabilities, when coincident rates of ingestion and absorption are nevertheless maintained, that is, until such elevated concentrations are reached to incur physical inhibition (e.g. clogging) of the feeding mechanism. Secondly, adjustments in CR are best understood in relation to the utilisable organic component of available seston, rather than the total suspended load; whether that utilisable component is measured as chlorophyll, as a marker for living organic matter, or as the remaining detrital organic matter, which is increasingly understood to play an important role in the nutrition of bivalve shellfish (Navarro et al. 2009, this Theme Section). Thirdly, responsive adjustments in CR are most evident at low availabilities of that utilisable component. Chl a may reach as much as 150 µg l⁻¹ in dense blooms of algae within unmixed eutrophic waters. Yet, maximal CR in Mytilus edulis in the present study was observed at <6 µg chl a l⁻¹. Similarly, maximal CR in Chlamys farrelli occurred at about 5 µg chl a l⁻¹, this species experiencing between about 1 and 20 µg chl a l⁻¹ during culture in China (Sun et al. 1996). More striking still, maximal CR in Perna canaliculus occurred at <2 µg chl a l⁻¹ (Hawkins et al. 1999), normally experiencing between about 0.5 and 5 µg chl a l⁻¹ during culture in New Zealand (Gibbs & Vant 1998). On this basis, variations in availability of suspended organic matter are of greatest physiological consequence in suspension-feeding bivalves whilst over the lower end of observed natural ranges, when responses are higher per unit change in food abundance and have the highest proportional impacts upon net energy balance (Hawkins et al. 1999).

A further advance in this field might be to formulate the relationship between the effects of both flow and food concentration on CR, by using the product of the two as units of food supply, provision, or availability, expressed as grams of chl a or carbon per litre per hour. Fig. 8 shows these relationships, both separately and combined as the supply rate, using data derived in the present study for chl a content in Isochrysis galbana. Over a wide range of supply rates, all the CR measures that we have judged to be valid in this study, i.e. those measured at flow rates of >150 ml min⁻¹, lie within a limited range of from 1.5 to 2.5 l⁻¹ h⁻¹, averaging about 2 l⁻¹ h⁻¹ or 6 l h⁻¹ g⁻¹ when standardised according to dry soft tissue weight, and which are similar to the near-maximal or ‘autonomous filtration’ condition suggested by Riisgård (2001a) for moderate food levels.

In any continued quest for accuracy, it is important to consider what measures of feeding are intended for, and whether they are ‘fit for purpose’. For example, we often measure feeding in individual bivalves and ignore the variation between individuals by taking mean values for given conditions. Extrapolating these data for use at the population or ecosystem level, for example, in modelling growth rates and impacts of bivalve feeding, is a significant step requiring some understanding of interactions between neighbouring individuals, the hydrodynamics of the system and, of course, temporal variations in food quantity and quality. In this context, there is a wealth of literature concerning the effects of flow or current velocity on bivalve feeding in nature (Wildish & Kristmanson 1997, Dolmer 2000, Newell et al. 2001, Widdows et al. 2002). It is worth emphasising that most or all of this research relates to relatively high current velocities (flows), including associated effects of external and internal pressure fields on the ciliary pump, neither of which are normally relevant to the effects of flow rate measured using a flow-through chamber system. These studies, whether using field measurements or attempting realistic simulations in a flume, usually involve flows well above 0.05 m s⁻¹, whereas maximal flow through our chambers was measured at <0.003 m s⁻¹.

Recently, Saurel et al. (2007) examined the feeding behaviour of mussel beds in the Menai Strait in relation to multiple environmental factors. Based on in situ video measurements of valve gape area (VA) over 2 tidal cycles, they concluded that mussel feeding was regulated solely by chl a concentration measured at 1 m above the mussel bed (0.6 to 2.5 µg l⁻¹), and not by flow velocity, changes in suspended particulate matter, or the presence of predators. However, at times of low flow during slack water, concentrations of chl a decreased to levels below the threshold for maximal CR and mussels consequently reduced their feeding activity. In fact, Saurel et al. (2007) estimated that maximal CR (VA of 80 to 100%) occurred for only ca. 45 min d⁻¹, suggesting that optimal feeding conditions are rarely experienced in the Menai Strait. Suggestions are made, with reference to the review by McKindsey et al. (2006), as to how these and other suitable in situ measurements may facilitate assessment and modelling of carrying capacity. Other work on in situ measurements has assessed the use of valve gape or exhalent siphon area as proxies for pumping rate in bivalves and propose these as useful methods for following the dynamic changes and physiological adaptations during natural bivalve feeding (Riisgård et al. 2003, 2006, Maire et al. 2007). Many of these field studies emphasise that CR varies greatly under natural conditions, such that dynamic modelling is required to assess ecological interrelations between bivalve sus-
pension feeders and their environment. Dynamic feeding responses have increasingly been established using the biodeposition method, which, through separate coincident collections of both true faeces and pseudofaeces and in contrast to measures of CR based solely upon particle depletion, not only integrates over time, but also enables calculations of ingestion rate and absorption efficiency (Hawkins et al. 1996, 2002, Cranford et al. 1998, Cranford & Hill 1999). Measured responses may then be used to simulate each physiological component of growth (Hawkins et al. 2002), for use in integrated ecological modelling (Ferreira et al. 2007, 2008).

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