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**Colourful coexistence : a new solution to the plankton paradox**

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*Citation for published version (APA):*

Stomp, M. (2008). Colourful coexistence : a new solution to the plankton paradox. Amsterdam.

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## Chapter 5

# **Colourful niches of phototrophic microorganisms shaped by vibrations of the water molecule**

### Abstract

The photosynthetic pigments of phototrophic microorganisms cover different regions of the solar light spectrum. Utilisation of the light spectrum can be interpreted in terms of classical niche theory, as the light spectrum offers opportunities for niche differentiation and allows coexistence of species absorbing different colours of light. However, which spectral niches are available for phototrophic microorganisms? Here, we show that the answer is hidden in the vibrations of the water molecule. Water molecules absorb light at specific wavebands that match the energy required for their stretching and bending vibrations. Although light absorption at these specific wavelengths appears only as subtle shoulders in the absorption spectrum of pure water, these subtle shoulders create large gaps in the underwater light spectrum due to the exponential nature of light attenuation. Model calculations show that the wavebands between these gaps define a series of distinct niches in the underwater light spectrum. Strikingly, these distinct spectral niches match the light absorption spectra of the major photosynthetic pigments. This suggests that vibrations of the water molecule have played a major role in the ecology and evolution of phototrophic microorganisms on our planet.

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This chapter is based on the paper: Stomp M, J Huisman, LJ Stal, HCP Matthijs (2007b) Colorful niches of phototrophic microorganisms shaped by vibrations of the water molecule. *ISME Journal* 1: 271-282.

## Introduction

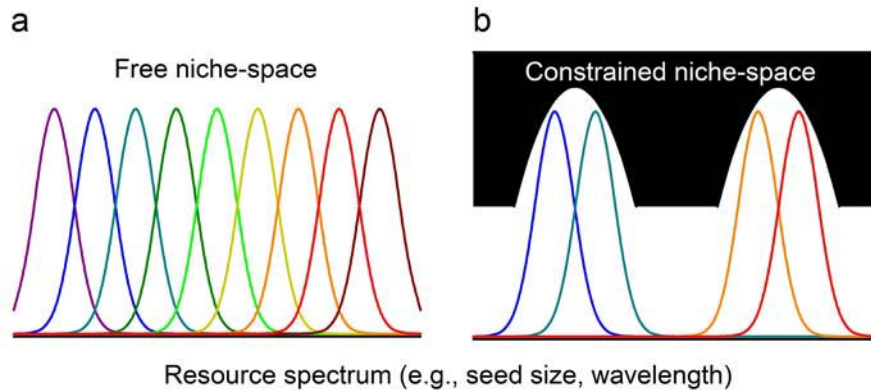
In the late 19<sup>th</sup> century, Professor Theodor W. Engelmann was the first to demonstrate that phototrophic organisms utilise specific parts of the light spectrum. He produced a “living action spectrum”, by illuminating filaments of the green alga *Spirogyra* with a light spectrum created by a prism glass. This revealed that oxygen-dependent bacteria accumulated near those parts of the algal filaments illuminated with red and blue light, thus demonstrating that the pigment chlorophyll absorbs red and blue light for photosynthesis (Engelmann 1882). One year later, in 1883, Engelmann discovered the utilisation of infrared wavelengths by purple bacteria (Engelmann 1883). Since then, many photosynthetic pigments have been identified, each with its own characteristic absorption spectrum (Pfennig 1967; Falkowski & Raven 1997; Des Marais 2000; Xiong *et al.* 2000; Bèjà *et al.* 2001; Falkowski *et al.* 2004). How can we explain the specific set of pigments that have evolved on planet Earth? Why is there not a single black pigment that absorbs all wavelengths?

Utilisation of the light spectrum can be interpreted in terms of classical ecological theory. Light offers a spectrum of resources. According to ecological theory, niche differentiation along a resource spectrum reduces competition among species, and thereby promotes their coexistence (Gause 1934; MacArthur & Levins 1967; May & MacArthur 1972; Rueffler *et al.* 2006). Darwin’s finches on the Galápagos Islands provide a famous example. Niche differentiation along a spectrum of different seed sizes allows a variety of finch species to coexist (Darwin 1859; Lack 1974; Grant & Grant 2002). Likewise, differences in light absorption spectra of species result in niche differentiation along the light spectrum. Indeed, competition models and experiments have shown that red and green picocyanobacteria can coexist by absorbing different parts of the light spectrum (Stomp *et al.* 2004).

Niche differentiation along the light spectrum is probably a common phenomenon in aquatic ecosystems. For instance, a recent field survey confirmed that the relative abundances of red and green picocyanobacteria in lakes and seas are related to the underwater light colour (Stomp *et al.* 2007). Red picocyanobacteria dominate in relatively clear waters where green light penetrates the deepest, while green picocyanobacteria dominate in turbid waters where red light penetrates the deepest. Coexistence of red and green picocyanobacteria is widespread in waters of intermediate colouration. Likewise, many other studies have revealed a close correspondence between the absorption spectra of phototrophic microorganisms and the prevailing underwater light spectrum (e.g., Pierson *et al.* 1990; Wood *et al.* 1998; Bèjà *et al.* 2001; Vila & Abella 2001; Rocap *et al.* 2003; Kühl *et al.* 2005; Bouman *et al.* 2006; Sabehi *et al.* 2007).

Which spectral niches are available for phototrophic microorganisms? If our planet would offer a continuum of spectral niches, then one would expect a free distribution of light absorption spectra along this continuum (as in Figure 5.1a). Alternatively, it might be that environmental conditions constrain part of the resource spectrum, such that only a few distinct niches are available (Figure 5.1b). In this paper, we show that vibrations of the water molecule create gaps in the underwater light spectrum. As a result, not all wavebands are

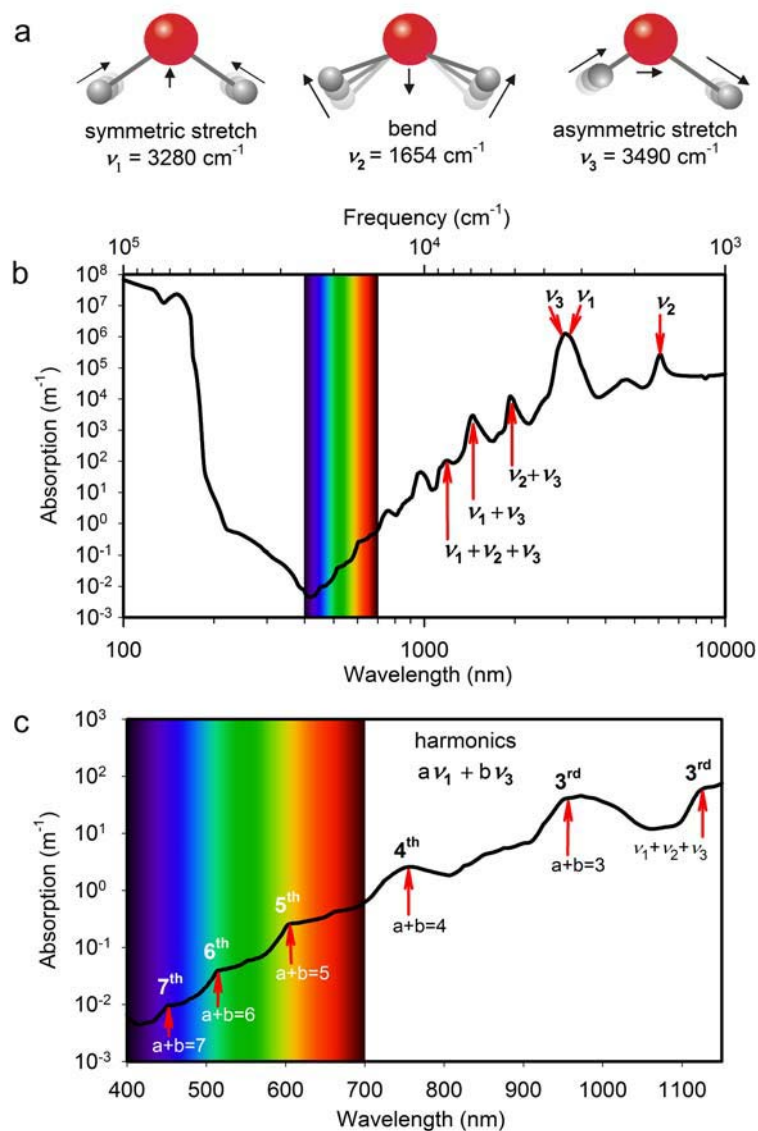
equally available for photosynthesis. This yields a series of distinct spectral niches for phototrophic microorganisms.



**Figure 5.1** Resource utilisation curves of competing species along a spectrum of resources. (a) Resources are freely distributed along the resource spectrum. (b) Environmental conditions impose constraints on the distribution of resources along the resource spectrum.

## Vibrations of water molecules

Water molecules are never at rest. They rotate and vibrate. Vibrations of water molecules occur in three modes, including symmetric stretching ( $\nu_1$ ), asymmetric stretching ( $\nu_3$ ), and bending ( $\nu_2$ ) of the water molecule (Figure 5.2a; see also, e.g., Braun & Smirnov 1993; Pegau *et al.* 1997; Sogandares & Fry 1997). The energy for these vibrations is obtained by absorption of radiation. The vibrations are most intense at wavelengths matching the specific energy requirements of these motions. These wavelengths can be recognised as peaks in the absorption spectrum of pure water (Figure 5.2b). Because the energy requirements for symmetric and asymmetric stretching are rather similar, their absorption peaks coalesce into a large absorption peak at around 3000 nm. The bending vibrations occur at a lower energy level, resulting in an absorption peak at around 6000 nm. Harmonics of these vibrations occur at higher energy levels (i.e., shorter wavelengths) that double or triple the required energy. As a result, harmonics of the bending and stretching vibrations can be recognised as shoulders in the visible and near-infrared range of the absorption spectrum of water. For instance, the distinct shoulders in Figure 5.2c, at 449 nm, 514 nm, 605 nm, 742 nm, and 972 nm have been identified as the seventh, sixth, fifth, fourth, and third harmonics, respectively, of the symmetrical and asymmetrical stretch vibration (Braun & Smirnov 1993; Pegau *et al.* 1997; Sogandares & Fry 1997). The shoulder at 1130 nm has also been identified as a third harmonics, composed of the combination of a symmetrical, asymmetrical, and bending vibration ( $\nu_1 + \nu_2 + \nu_3$ ). We will argue, below, that these subtle shoulders in the absorption spectrum of pure water have a major effect on the underwater light spectrum.



**Figure 5.2** (a) The three vibrational modes of the water molecule and their fundamental frequencies in liquid water: symmetric stretching ( $\nu_1$ ), bending ( $\nu_2$ ), and asymmetric stretching ( $\nu_3$ ). The atoms move in the directions indicated by arrows. (b) Absorption spectrum of pure water (Hale & Querry 1973; Segelstein 1981; Pope & Fry 1997). Peaks in the absorption spectrum correspond to the fundamental frequencies and higher harmonics of the vibrations of the water molecules. (c) Absorption spectrum of pure water in the visible and infrared region. Shoulders in the absorption spectrum correspond to the third, fourth, fifth, sixth and seventh harmonics of the symmetric and asymmetric stretch vibrations, as indicated.

## The underwater light colour

The underwater light spectrum of aquatic ecosystems depends on light absorption by pure water as well as by other components, including dissolved organic matter (known as ‘gilvin’ in the optics literature), inanimate particulate organic matter (known as ‘tripton’), and phytoplankton. More specifically, according to Lambert-Beer’s law the underwater light spectrum can be calculated as (Sathyendranath & Platt 1989; Kirk 1994; Stomp *et al.* 2007):

$$I(\lambda, z) = I_{in}(\lambda) \text{EXP}\left(-[K_w(\lambda) + K_{GT}(\lambda) + K_{PH}(\lambda)]z\right) \quad (5.1)$$

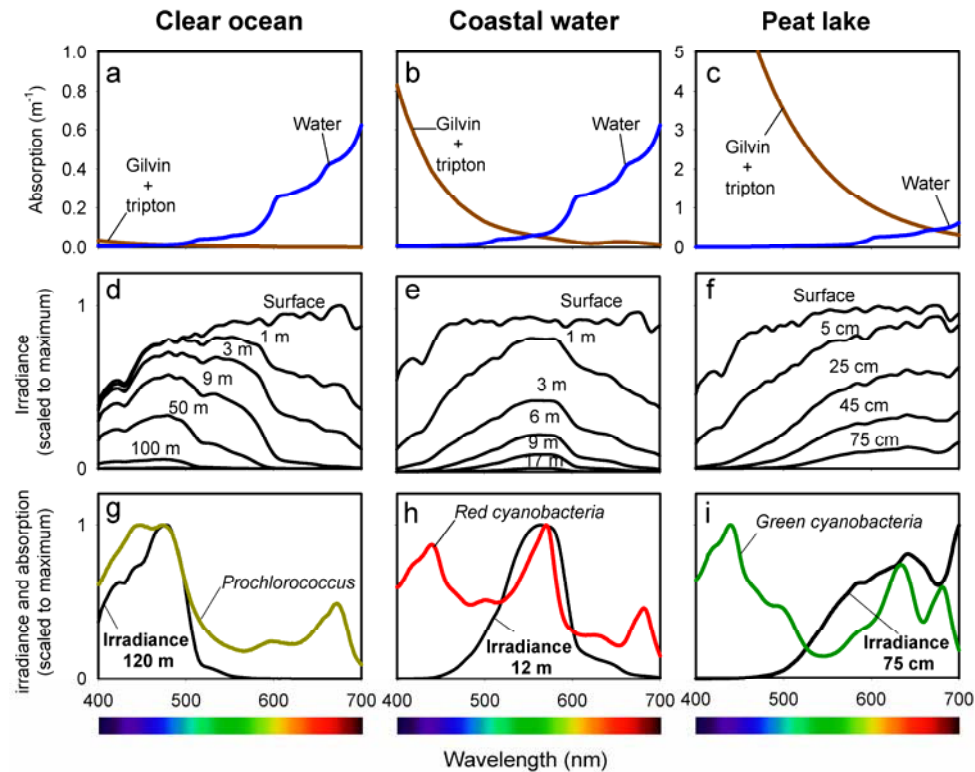
where  $I(\lambda, z)$  is the light intensity of wavelength  $\lambda$  at depth  $z$ ,  $I_{in}(\lambda)$  is the spectrum of the incident solar irradiance,  $K_w(\lambda)$  is the absorption spectrum of pure water (Figure 5.2b and c),  $K_{GT}(\lambda)$  is the absorption spectrum of gilvin and tripton, and  $K_{PH}(\lambda)$  is the absorption spectrum of the phytoplankton community.

The incident solar irradiance has essentially a white spectrum with a few small dips from 450 nm to 900 nm, and two large gaps in the infrared at 950 nm and 1120 nm. These dips in the incident solar spectrum are due to light absorption by oxygen and water molecules in the atmosphere (Kirk 1994). Pure water mainly absorbs red and infrared light, with several distinct shoulders (Figure 5.2c). In contrast, gilvin and tripton absorb strongly in the blue part of the spectrum (Figure 5.3a-c). More specifically, the absorption spectrum of gilvin and tripton is a decreasing function of wavelength, which can be described by a smoothly declining exponential curve (Bricaud *et al.* 1981; Kirk 1994):

$$K_{GT}(\lambda) = K_{GT}(440) \text{EXP}(-S(\lambda - 440)) \quad (5.2)$$

where  $K_{GT}(440)$  is the attenuation coefficient of gilvin and tripton at a reference wavelength of 440 nm, and  $S$  is the slope of the exponential decline. The attenuation coefficient  $K_{GT}(440)$  is proportional to the concentration of gilvin and tripton.

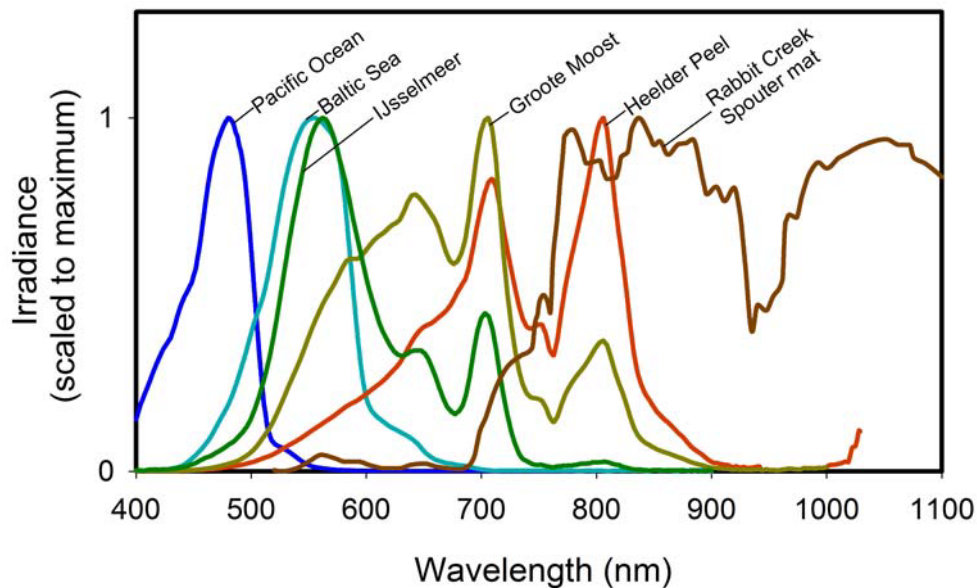
In waters with low concentrations of gilvin and tripton and low phytoplankton concentrations, light absorption by pure water dominates (e.g., Kirk 1994; Morel *et al.* 2007). This applies, for instance, to the oligotrophic waters of the subtropical Pacific Ocean (Figure 5.3a). Here, red light is absorbed by water within the upper 10 m, whereas blue light penetrates much deeper (Figure 5.3d). Indeed, selective absorption of red light is responsible for the blue colour of the oceans of our planet. In coastal waters, like the Baltic Sea, gilvin and tripton concentrations are higher, and their light absorption is often of a similar magnitude as light absorption by water itself (Figure 5.3b). As a result, green light penetrates the deepest (Figure 5.3e). In peat lakes, gilvin and tripton concentrations are extremely high, such that blue and green light are rapidly absorbed (Figure 5.3c). As a result, red light penetrates the deepest (Figure 5.3f). Hence, with increasing gilvin and tripton concentrations, the underwater light colour is shifted from the blue part towards the green and red part of the spectrum.



**Figure 5.3** Underwater light spectra measured in the subtropical Pacific Ocean (station ALOHA), the coastal waters of the Baltic Sea (stations near the Gulf of Finland), and a peat lake in The Netherlands (Lake Groote Moost). (a, b, c) Light absorption spectra of pure water (blue line) and gilvin plus tripton (brown line). Note that light absorption by gilvin plus tripton is low in the Pacific Ocean, intermediate in the Baltic Sea, and extremely high in Lake Groote Moost. (d, e, f) Underwater light spectra show that blue light penetrates very deep into the subtropical Pacific Ocean, green light penetrates deep into the Baltic Sea, while red light prevails in Lake Groote Moost. (g) The phytoplankton community sampled at 120 m depth in the subtropical Pacific Ocean was dominated by low-light adapted *Prochlorococcus*, which strongly absorb the available blue light using the pigments divinyl-chlorophyll *a* and *b* (absorption band at 450-500 nm). (h) The phytoplankton community sampled at 12 m depth in the Baltic Sea was dominated by red-coloured *Synechococcus* strains, which strongly absorb the available green light using the pigment phycoerythrin. (i) The phytoplankton community sampled at 75 cm depth in Lake Groote Moost was dominated by green cyanobacteria and green algae, which strongly absorb the available red light (absorption peaks of phycocyanin at 635 nm and chlorophyll *a* at 680 nm). Materials and methods for these measurements are described in Appendix B.

To investigate in further detail how gilvin concentrations affect the underwater light spectrum, we measured the underwater light spectrum in a variety of different aquatic ecosystems, ranging from blue waters of the Pacific Ocean to brown waters of very humic lakes. In addition, we searched the literature for light spectra measured in microbial mats, and found a beautiful spectrum from the murky microbial mats surrounding Rabbit Creek Spouter in Yellowstone National Park (Boomer *et al.* 2000). Figure 5.4 plots the underwater light spectra measured at the euphotic depth. The euphotic depth is here defined as the depth at which the irradiance over the entire photosynthetically available spectrum for aquatic microorganisms

(400-1100 nm) equals 1% of the irradiance at the water surface. With increasing gilvin and tripton concentration, the underwater light colour is shifted towards the red and even the infrared region of the spectrum in Lake Groote Moost, Lake Heelder Peel and the Rabbit Creek Spouter mat. Surprisingly, the data do not show a smooth shift in the underwater light spectrum, but reveal a striking landscape of peaks and valleys (Figure 5.4). Some underwater light spectra consist of a single peak, like the spectra of the Pacific Ocean and the Baltic Sea. Other underwater light spectra display several peaks and valleys. Moreover, the same peaks and valleys reoccur in different aquatic ecosystems. For instance, Lake IJsselmeer shows a similar peak at 560 nm as the Baltic Sea, and shares two peaks at 640 nm and 700 nm with Lake Groote Moost. Lake Groote Moost and Lake Heelder Peel share conspicuous peaks at both 700 nm and 800 nm, separated by a deep valley at 740-760 nm (Figure 5.4). The spectrum of the Rabbit Creek Spouter mat partly overlaps with Lake Heelder Peel, and extends further into the infrared, with a large dip at around 935 nm. If the absorption spectrum of gilvin and tripton is a smoothly decreasing function of wavelength (Equation 5.2), then why do underwater light spectra produce such a striking landscape of peaks and valleys?



**Figure 5.4** Underwater light spectra measured at the euphotic depth of 6 aquatic ecosystems. These ecosystems cover the entire range from very clear waters with low gilvin and tripton concentrations in the subtropical Pacific Ocean to extremely turbid conditions in the microbial mat of Rabbit Creek Spouter (a hot spring in Yellowstone National Park). Euphotic depths vary accordingly, from 120 m in the clear blue waters of the subtropical Pacific Ocean to less than a few millimeters in the microbial mat. Materials and methods for these measurements are described in Appendix B. The irradiance spectrum of the Rabbit Creek Spouter mat is from Boomer *et al.* (2000).



## Small shoulders, large gaps

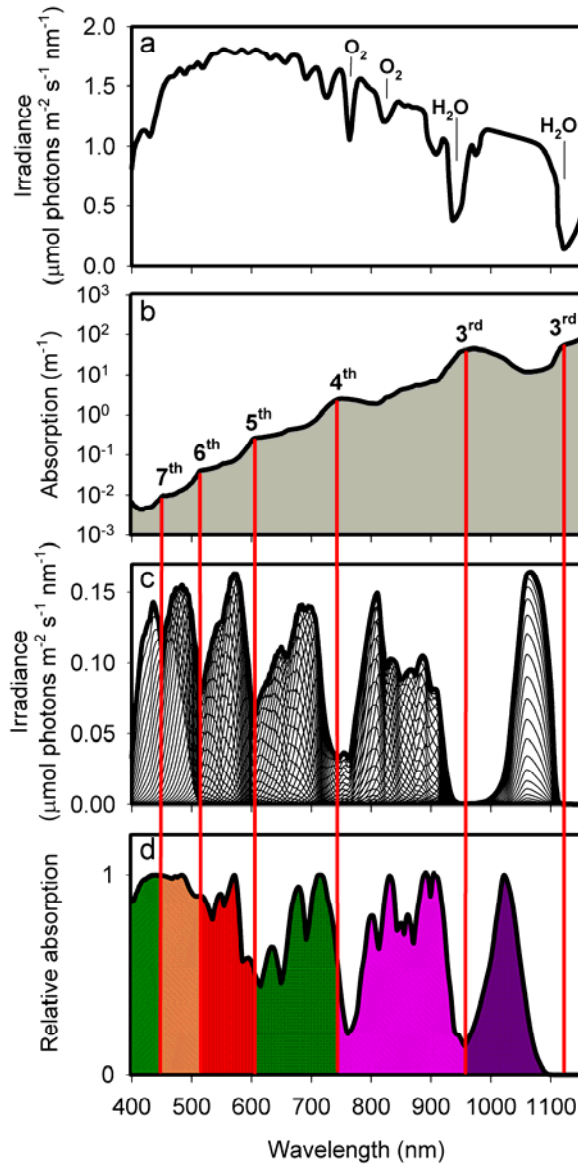
The spectrum of the incident solar irradiance (Figure 5.5a) might offer one possible explanation for the striking landscape of peaks and valleys in the underwater light spectra of Figure 5.4. However, according to Equation 5.1, peaks and valleys in the solar spectrum are transferred linearly in the underwater light spectrum. That is, the peaks and valleys in the solar spectrum are not amplified with depth. Therefore, only major gaps in the solar spectrum can be recognised in underwater light spectra. For example, the sharp dip in the solar light spectrum at 765 nm, due to atmospheric oxygen (Figure 5.5a), can be recognised in the underwater light spectra of Lake Groote Moost and Lake Heelder Peel. Similarly, the major gap in the solar light spectrum at 935 nm, due to atmospheric water, can be recognised in the irradiance spectrum of the Rabbit Creek Sprouter mat. The remainder of the solar spectrum in the range of 450-1100 nm is rather flat, however, and does not bear any resemblance with the measured underwater light spectra (compare Figure 5.4 and Figure 5.5a).

The subtle shoulders in the light absorption spectrum of pure water (Figure 5.5b) may offer an alternative explanation for the striking landscape of peaks and valleys in Figure 5.4. According to Equation 5.1, subtle shoulders in the light absorption spectrum of water are amplified with depth. That is, consider two wavelengths,  $\lambda_1$  and  $\lambda_2$ . For simplicity, assume that there is a subtle difference in light absorption by water at these two wavelengths, while the incident irradiance and light attenuation by other components would be equal for  $\lambda_1$  and  $\lambda_2$ . Now, according to Equation 5.1, the light intensities at these two wavelengths will diverge exponentially with depth:

$$\frac{I(\lambda_1, z)}{I(\lambda_2, z)} = \text{EXP}([K_w(\lambda_2) - K_w(\lambda_1)]z) \quad (5.3)$$

This shows that, due to the exponential nature of light absorption, subtle shoulders in the absorption spectrum of water create large gaps in the underwater light spectrum.

To investigate the latter hypothesis in further detail, we calculated the underwater light spectrum in the absence of phytoplankton. In this way, we obtained the available niches in the underwater light spectrum that can be exploited as a potential playfield for the ecology and evolution of phototrophic microorganisms. More specifically, we used Equations 5.1 and 5.2 to calculate 100 different underwater light spectra at the euphotic depth. The absorption spectrum of pure water was taken from the literature (Hale & Querry 1973; Segelstein 1981; Pope & Fry 1997). The slope  $S$  in Equation 5.2 typically varies between 0.010 and 0.020  $\text{nm}^{-1}$ , and we here assumed a typical value of  $S = 0.017 \text{ nm}^{-1}$  (Kirk 1994). The calculations were made for a wide range of different chlorophyll and tripton concentrations, with  $K_{CT}(440)$  values from 0.003  $\text{m}^{-1}$  in very clear ocean waters (Morel *et al.* 2007) to more than 5000  $\text{m}^{-1}$  in extremely turbid systems representative for microbial mats in sediments (Kühl & Jørgensen 1994). As a consequence, the euphotic depths ranged from more than 200 m in clear ocean water to only a few mm in turbid sediments and microbial mats.



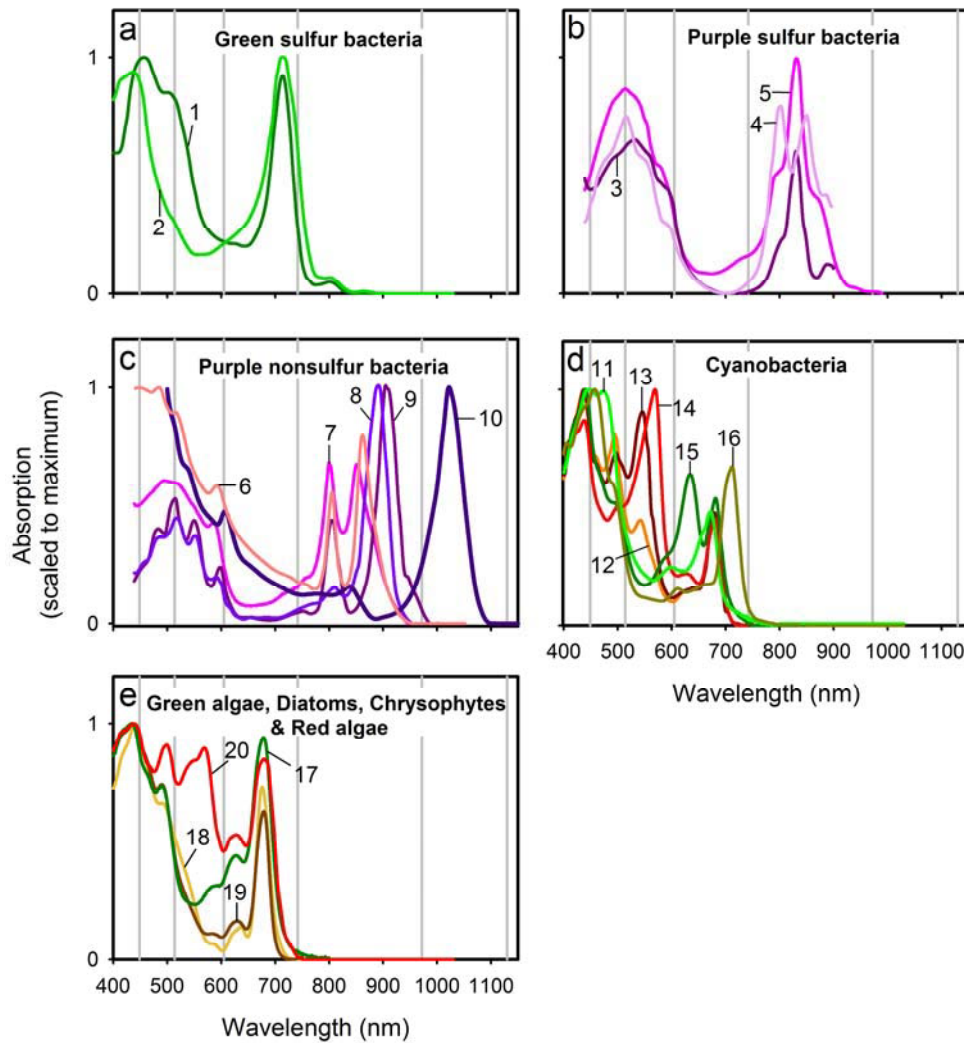
**Figure 5.5** The absorption spectrum of water creates a series of distinct niches in the underwater light spectrum. (a) Light spectrum of the incident solar irradiance at the water surface. Dips in the incident irradiance are caused by absorption of photons by oxygen and water molecules in the atmosphere. (b) Absorption spectrum of pure water, plotted at a log scale. The different harmonics of the stretching and bending vibrations of the water molecule are indicated. (c) Overlay of 100 underwater light spectra at the euphotic depth. The light spectra are calculated from Equations 5.1 and 5.2, using a wide range of different gilvin and tripton concentrations, from the clearest ocean waters to very turbid systems such as microbial mats. (d) Overlay of measured light absorption spectra of 20 phototrophic species, including purple sulfur bacteria, green sulfur bacteria, purple nonsulfur bacteria, cyanobacteria, green algae, red algae, diatoms and chrysophytes. Light absorption spectra of each individual species are shown in Figure 5.6.

Figure 5.5c shows an overlay of all 100 underwater light spectra thus calculated, which reveals a landscape of peaks and valleys quite similar to the measured underwater light spectra in Figure 5.4. Comparison of Figure 5.5a and Figure 5.5c show that the dips in solar irradiance have some effect on the underwater light spectra, but this effect is relatively small. For instance, the dip caused by atmospheric oxygen at 765 nm (Figure 5.5a) creates a small secondary valley in the calculated underwater light spectra (Figure 5.5c). This small secondary valley was also visible in the measured light spectra of Lake Groote Moost and Lake Heelder Peel (Figure 5.4). In contrast, our calculations show that, consistent with Equation 5.3, the subtle shoulders in the absorption spectrum of water create large gaps in the underwater light spectrum. The shoulder of the sixth harmonics in the absorption spectrum of water (Figure 5.5b) creates a large gap in the underwater light spectrum at ~514 nm (Figure 5.5c). This gap separates the measured underwater light spectra of the Pacific Ocean and the Baltic Sea (Figure 5.4). Likewise, the shoulder at the fifth harmonics in the absorption spectrum of pure water (Figure 5.5b) creates a gap in the underwater light spectrum at ~600 nm (Figure 5.5c), which separates the measured underwater light spectra of the Baltic Sea and Lake IJsselmeer from the peaks of Lake Groote Moost and Lake Heelderpeel (Figure 5.4). The next shoulder in the absorption spectrum of pure water, at the fourth harmonics (Figure 5.5b), creates a large gap in the underwater light spectrum at 740-760 nm (Figure 5.5c). This corresponds to the gap within the measured underwater light spectra of Lake Groote Moost and Lake Heelder Peel (Figure 5.4). Finally, at ~950 nm, the combination of a deep trough in the incident solar irradiance caused by water molecules in the atmosphere (Figure 5.5a) and a large shoulder at the third harmonics of liquid water (Figure 5.5b) create a deep gap in the underwater light spectrum (Figure 5.5c). This gap can be clearly recognised in the irradiance spectra of microbial mats (Pierson *et al.* 1990; Boomer *et al.* 2000), as exemplified by the Rabbit Creek Spouter mat (Figure 5.4). In other words, this exercise shows that the underwater light spectrum does not present a homogeneous playfield for the ecology and evolution of phototrophic microorganisms. Instead, the underwater light spectrum offers a number of distinct niches at specific wavebands, separated by deep gaps created by the shoulders in the light absorption spectrum of the water molecule.

## Filling the niches

Have phototrophic microorganisms adapted the absorption spectra of their pigments to these distinct niches in the underwater light spectrum? Three examples are presented in Figure 5.3. The phytoplankton community in the subtropical Pacific Ocean is dominated by picocyanobacteria of the genus *Prochlorococcus* (Chisholm *et al.* 1988; Letelier *et al.* 1993). *Prochlorococcus* effectively absorbs the available blue light with its pigments divinyl-chlorophyll *a* and *b* (Figure 5.3g). In the Baltic Sea, the phytoplankton community at the euphotic depth is dominated by red-coloured picocyanobacteria of the *Synechococcus* group (Stomp *et al.* 2007), which effectively absorb the available green light with their pigment phycoerythrin (Figure 5.3h). In peat lakes, the phytoplankton community is often dominated by green-coloured

phytoplankton species, like green cyanobacteria and green algae, which absorb the available red light with pigments such as phycocyanin and chlorophylls *a* and *b* (Figure 5.3i). This is a first indication that the light absorption spectra of phytoplankton communities are often well tuned to their underwater light environment.



**Figure 5.6.** In vivo absorption spectra of intact cells of 20 phototrophic species, including purple sulfur bacteria, green sulfur bacteria, purple non-sulfur bacteria, cyanobacteria, green algae, red algae, diatoms and chrysophytes. The names of the species and their main photosynthetic pigments are listed in Table 5.1. Vertical lines indicate the location of the harmonics of the stretching and bending vibrations of the water molecule.

To extend our analysis to the full light spectrum available for photosynthesis, from 400 to 1100 nm, we gathered absorption spectra of 20 phototrophic species containing a wide variety of different pigments. The species belong to the green sulfur bacteria, purple sulfur bacteria, purple nonsulfur bacteria, cyanobacteria, green algae, diatoms, chrysophytes and red algae (Table 5.1). We measured the light absorption spectra of seven species with an Aminco DW-2000 double-beam spectrophotometer. Light absorption spectra of the other 13 species were taken from the literature (Table 5.1). The absorption spectra of the individual species are shown in Figure 5.6. An overlay of all 20 absorption spectra is presented in Figure 5.5d. The match between the peaks and valleys in the underwater light spectrum and the peaks and valleys in the light absorption spectra of this rich variety of photosynthetic pigments is striking. That is, the harmonics in the light absorption spectrum of pure water (Figure 5.5b) create a series of distinct niches in the underwater light spectrum (Figure 5.5c), which are effectively captured by the light absorption spectra of the major groups of phototrophic microorganisms inhabiting our planet (Figure 5.5d). Peaks in the underwater light spectrum at wavelengths below the sixth harmonics ( $< 514$  nm) are captured by chlorophyll *a* and *b*, divinylchlorophylls *a* and *b*, and accessory carotenoids. The peak in the underwater light spectrum between the sixth and fifth harmonics (514-604 nm) is captured by the phycoerythrins of red algae and cyanobacteria. Peaks in the underwater light spectrum between the fifth and fourth harmonics (604-760 nm) are captured by phycocyanin, chlorophylls *a*, *b* and *d*, and bacteriochlorophyll *e*. Bacteriochlorophyll *a* in various phototrophic bacteria captures the peaks in the underwater light spectrum between the fourth and third harmonics (760-960 nm). Finally, bacteriochlorophyll *b* in the purple bacterium *Blastochloris viridis* (formerly known as *Rhodospseudomonas viridis*) harvests the light energy available at wavelengths between both third harmonics (960-1130 nm). Interestingly, the absorption peak of *B. viridis* seems a bit shifted towards shorter wavelengths compared to its spectral niche (compare Figure 5.5c and d). Perhaps our model calculations do not provide a very accurate description of spectral niches in microbial mats, where scattering of light can play an important role (Pierson *et al.* 1990). Furthermore, the infrared part of the light absorption spectrum of water is sensitive to temperature (Collins 1925; Braun & Smirnov 1993), which may shift this spectral niche to slightly shorter wavelengths at high temperatures. Also, the exact location of the light absorption peak of *B. viridis* is probably sensitive to temperature, and the range of measurements on lab cultures may not be exactly the same as in the bacterium's native environment (Kiang *et al.* 2007). It would be interesting to investigate these issues further. Perhaps other species containing bacteriochlorophyll *b* are capable to harvest light at even longer wavelengths, and therefore match this infrared niche more closely.

## Discussion

In this paper, we have developed the hypothesis that vibrations of the water molecule generate a series of distinct niches in the underwater light spectrum, which are effectively utilised by the different phototrophic microorganisms inhabiting our planet. Our hypothesis implicitly

assumes that the underwater light spectrum is an important selective factor for the ecology and evolution of phototrophic microorganisms. This hypothesis is supported by several lines of evidence. Many physiological studies have shown that light colour affects photosynthesis and growth rates of phototrophic microorganisms, as has been demonstrated for, e.g., green sulfur bacteria (Montesinos *et al.* 1983; Vila & Abella 1994), cyanobacteria (Wyman & Fay 1986; Hauschild *et al.* 1991; Callieri *et al.* 1996), and eukaryotic phytoplankton (Holdsworth 1985; Glover *et al.* 1987). Furthermore, laboratory competition experiments have shown that light colour can act as a selective factor. For instance, Parkin & Brock (1980) studied competition between green and purple sulfur bacteria isolated from the sulfide containing waters of three stratified lakes with different underwater light spectra. They observed that green sulfur bacteria became dominant in flasks exposed to red light, whereas purple sulfur bacteria became dominant in flasks exposed to green light. Likewise, Stomp *et al.* (2004) studied competition between red and green picocyanobacteria isolated from the Baltic Sea. They developed a competition model that predicts that red picocyanobacteria should become dominant in green light, green picocyanobacteria should become dominant in red light, whereas red and green picocyanobacteria can coexist in the full spectrum provided by white light. The results of their competition experiments were consistent with these predictions. These studies demonstrated that light colour plays a decisive role in the species composition of phototrophic communities, at least in controlled laboratory experiments.

Numerous field studies have confirmed that the species composition of phototrophic microorganisms is related to the underwater light spectrum. For instance, Figure 5.3 shows that *Prochlorococcus* in the subtropical Pacific Ocean is well tuned to the prevailing blue light, red picocyanobacteria in the Baltic Sea are well tuned to the prevailing green light, and green cyanobacteria and green algae in turbid waters are well tuned to the prevailing red light. Recent work has shown that relative abundances of red and green picocyanobacteria show a clear link with the underwater light spectrum across many ecosystems (Stomp *et al.* 2007). Likewise, the relative abundances of proteorhodopsin containing bacteria absorbing either blue or green light have been explained by prevailing spectral light conditions (Béjà *et al.* 2001; Sabeji *et al.* 2007). Similar observations have shown a good correspondence between the absorption spectra of phototrophic microorganisms and the underwater light spectrum in clear oceans (Ting *et al.* 2002; Rocap *et al.* 2003; Bouman *et al.* 2006), coastal waters (Olson *et al.* 1990; Wood *et al.* 1998; Katano *et al.* 2007), lakes (Pick 1991; Vörös *et al.* 1998; Vila & Abella 2001) and microbial mats (Pfennig 1967; Pierson *et al.* 1990; Kühl *et al.* 2005). Thus, there is overwhelming evidence from theory, laboratory experiments and field data that the underwater light spectrum is a major determinant of the composition of phototrophic communities.

The novel part of our hypothesis is that the underwater light spectrum does not offer a continuum of niches (as in Figure 5.1a), but consists of a series of distinct niches (as in Figure 5.1b) created by vibrations of the water molecule. This rather unexpected prediction is supported by the striking similarity between the calculated peaks and valleys in the underwater light spectra (Figure 5.5c) and the observed peaks and valleys in the absorption spectra of phototrophic microorganisms (Figure 5.5d). That is, the absorption spectra of the major

photosynthetic pigments fit the available niches in the underwater light spectrum. Moreover, each spectral niche can be occupied by several pigments, with slightly different absorption peaks. For example, the niche between the sixth and fifth harmonics is occupied by two types of phycoerythrin, known as PUB (peak at 494 nm) and PEB (peak at 545 nm) (Toledo 1999). Likewise, several variants of bacteriochlorophyll *a* cover the niche between the fourth and third harmonics. This indicates that, within each spectral niche, absorption peaks of phototrophic organisms may diverge, possibly driven by the evolutionary process of adaptive radiation (Schluter 2000; Rueffler 2006).

Yet, this part of our hypothesis is clearly open for further testing. For instance, laboratory competition experiments could simulate light environments that deviate from the underwater light niches predicted by our model. As a first test, mixtures of phototrophic microorganisms could be exposed to wavebands that are less available in underwater light spectra. For example, wavelengths around 600 nm are relatively less available due to strong absorption by the fifth harmonics of water (Figure 5.5c) and currently few microorganisms have their absorption peak at 600 nm (Figure 5.6). If this waveband becomes the prevailing light colour in a long-term laboratory selection experiment, will selection favour new mutants that shift their absorption peak to 600 nm? Another interesting test could be based on selection experiments in artificial water that lacks the characteristic absorption peaks of normal water (H<sub>2</sub>O). In so-called heavy water (D<sub>2</sub>O), the hydrogen atoms are replaced by heavier deuterium atoms. As a consequence, molecular vibrations of D<sub>2</sub>O occur at other frequencies, and the harmonics of heavy water molecules are shifted to the far-red compared to H<sub>2</sub>O. Hence, the absorption spectrum of D<sub>2</sub>O is completely different from H<sub>2</sub>O (Tam & Patel 1979; Braun & Smirnov 1993). Our hypothesis therefore predicts that, in light-limited systems, selection experiments in D<sub>2</sub>O will lead to phototrophic communities with other light absorption spectra than selection experiments in H<sub>2</sub>O.

In conclusion, our findings point at a striking causal relationship between the stretching and bending vibrations of the water molecule, the underwater light spectra of aquatic ecosystems, and the ecology and evolution of phototrophic microorganisms.

## Acknowledgements

We thank M. Laamanen and D.M. Karl for the opportunity to join cruise Cyano-04 on the Baltic Sea and HOT cruise 174 on the Pacific Ocean, and the crew of research vessels Aranda and Kilo Moana for their great help during sampling. We also thank B. Pex, H. van Overzee, R. Poutsma and students of the MSc program Limnology & Oceanography 2005 for their help in the Dutch lakes, and H.J. Gons and J.C. Kromkamp for the underwater light spectrum of Lake IJsselmeer, and their help with the filterpad method. Special thanks to O. Béjà and the anonymous referee for their helpful comments on the manuscript. The research of M.S. and J.H. was supported by the Earth and Life Sciences Foundation (ALW), which is subsidised by the Netherlands Organization for Scientific Research (NWO). L.J.S. acknowledges support from the European Commission through the project MIRACLE (EVK3-CT-2002-00087).

**Table 5.1** Phototrophic organisms presented in Figure 5.5d and Figure 5.6.

No	Species	Main Pigments	Reference
<b>Green sulfur bacteria</b>			
1	<i>Pelodictyon phaeoclathratiforme</i>	BChl <i>e</i>	Overmann & Pfennig 1989
2	<i>Prosthecochloris aestuarii</i>	BChl <i>a, c</i>	Overmann <i>et al.</i> 1991
<b>Purple sulfur bacteria</b>			
3	<i>Thiocapsa marina</i>	BChl <i>a</i>	Caumette <i>et al.</i> 2004
4	<i>Thiocapsa roseopersicina</i>	BChl <i>a</i>	Caumette <i>et al.</i> 2004
5	<i>Chromatium okenii</i>	BChl <i>a</i>	Pfennig 1967
<b>Purple nonsulfur bacteria</b>			
6	<i>Rhodobacter capsulatus</i>	BChl <i>a</i>	Zubova <i>et al.</i> 2005
7	<i>Rhodobacter sphaeroides</i>	BChl <i>a</i>	This study
8	<i>Rhodospirillum rubrum</i>	BChl <i>a</i>	Pfennig 1967
9	<i>Roseospirillum parvum</i>	BChl <i>a</i>	Glaeser & Overmann 1999
10	<i>Blastochloris viridis</i>	BChl <i>b</i>	Pfennig 1967
<b>Cyanobacteria</b>			
11	<i>Prochlorococcus sp.</i>	Divinyl Chl <i>a, b</i>	This study
12	<i>Synechococcus</i> WH7803	Chl <i>a</i> , PUB/PEB	Toledo <i>et al.</i> 1999
13	<i>Synechococcus</i> WH8103	Chl <i>a</i> , PUB/PEB	Toledo <i>et al.</i> 1999
14	<i>Synechococcus</i> BS5	Chl <i>a</i> , PC, PE	This study
15	<i>Synechococcus</i> BS4	Chl <i>a</i> , PC	This study
16	<i>Acaryochloris marina</i>	Chl <i>d</i>	Kühl <i>et al.</i> 2005
<b>Green algae</b>			
17	<i>Chlamydomonas sp.</i>	Chl <i>a</i> , Chl <i>b</i>	This study
<b>Diatoms</b>			
18	<i>Phaeodactylum tricornutum</i>	Chl <i>a</i>	This study
<b>Chrysophytes</b>			
19	<i>Isochrysis sp.</i>	Chl <i>a</i>	This study
<b>Red algae</b>			
20	<i>Palmaria palmata</i>	Chl <i>a</i> , PE	Cordi <i>et al.</i> 1997

BChl = bacterio-chlorophyll, Chl = chlorophyll, PC = phycocyanin, PE = phycoerythrin, PEB = phycoerythrobilin, PUB = phycourobilin



*Chapter 5*