

## NOTES AND CORRESPONDENCE

### Effects of bacteria-virus interaction on *Synechococcus* spp. growth in the coastal waters

An-Yi Tsai<sup>1,2,\*</sup>

<sup>1</sup>Institute of Marine Environment and Ecology, National Taiwan Ocean University, Keelung City, Taiwan

<sup>2</sup>Center of Excellence for the Oceans, National Taiwan Ocean University, Keelung City, Taiwan

#### Article history:

Received 18 May 2020

Revised 20 July 2020

Accepted 13 August 2020

#### Keywords:

Nanoflagellate grazing, Viral lysis, Heterotrophic bacteria, *Synechococcus* spp., Modified dilution approach, Recycled nutrients

#### Citation:

Tsai, A.-Y., 2020: Effects of bacteria-virus interaction on *Synechococcus* spp. growth in the coastal waters. Terr. Atmos. Ocean. Sci., 31, 691-696, doi: 10.3319/TAO.2020.08.13.01

#### ABSTRACT

This study aims to investigate the relative importance of nanoflagellate grazing and viral lysis for heterotrophic bacteria and *Synechococcus* spp. mortality in coastal waters of the north-eastern Taiwan. The results showed that viral lysis and nanoflagellate grazing are the main factors responsible for the mortality of heterotrophic bacteria, however, the presence of viruses had a positive effect on the *Synechococcus* spp. growth. Moreover, this study observed different growth responses between heterotrophic bacteria and *Synechococcus* spp. population by using the modified dilution approach. The results also showed that there was lower net growth rate of *Synechococcus* spp. in 30 kDa dilution series than that observed in 0.2  $\mu\text{m}$  series. Furthermore, based on the results obtained from modified dilution approach, we can estimate that about 12.8  $\text{ng N L}^{-1} \text{h}^{-1}$  and 2.8  $\text{ng P L}^{-1} \text{h}^{-1}$  was released by nanoflagellate grazing. In addition, this study estimated that, during the process of viral lysis, locally recycled nutrients released about 224  $\text{ng N L}^{-1} \text{h}^{-1}$  and 24.9  $\text{ng P L}^{-1} \text{h}^{-1}$  in this system. Our findings suggest that due to viral lysis, substantial portion of the recycled nutrients required for *Synechococcus* spp. growth was supplied sufficiently, emphasizing the importance of the viral shunt, especially in oligotrophic regions.

## 1. INTRODUCTION

Picophytoplankton (< 2  $\mu\text{m}$ ) contribute significantly to phytoplankton biomass and production in oligotrophic environments (Bell and Kalff 2001; Winder 2009; Grob et al. 2011). Among these small size picophytoplankton, *Synechococcus* spp. is virtually ubiquitous in marine environments and contributes to more than 50% of the primary production in oligotrophic oceans (Jardillier et al. 2010; Tsai et al. 2012). Compared to other larger group phytoplankton, *Synechococcus* spp. possesses high specific growth rates due to its small size, and has high specific affinity for nutrients (Agawin et al. 2000). Grazing and viral lysis are the two major causes of *Synechococcus* spp. mortality (Baudoux et al. 2008; Tsai et al. 2012). Grazing leads to the transfer of biomass to higher trophic levels, whereas viral

lysis recycles carbon and nutrients within the microbial loop (Ayukai 1996).

The dilution approach, proposed by Landry and Hassett (1982), has been accepted as a standard method for estimating phytoplankton growth and grazing mortality rates (Ayukai 1996; Redden et al. 2002; Juhl and Murrell 2005; Anderson et al. 2018). Following this technique, Evans et al. (2003) estimated viral lysis and grazing effect on picoeukaryote, *Micromonas* simultaneously using a modified dilution method. One of the main assumptions of the dilution approach is that prey growth rate is independent of the dilution (Landry and Hassett 1982). However, in some nutrient limited waters, growth of *Synechococcus* spp. is lower in incubations that are diluted with virus-free diluents than those diluted with 0.2  $\mu\text{m}$  diluents, containing viruses (Shelford et al. 2012; Tsai et al. 2018; Gong and Tsai 2019). Similarly, Tsai et al. (2018) found that *Synechococcus* spp. growth rate was clearly lower in treatments in which the

\* Corresponding author  
E-mail: anyitsai@mail.ntou.edu.tw

viral abundance was reduced by using the modified dilution approach. These observations may violate the assumption associated with the dilution approach that prey growth rate is independent of dilution effect, but highlights that the presence of viruses had a positive effect on the growth of *Synechococcus* spp. Furthermore, Weinbauer et al. (2011) studied the interactions among *Synechococcus* spp., heterotrophic bacteria and viruses. Based on the results, they suggested that nutrient cycles that are controlled by viral lysis of heterotrophic bacteria may also control *Synechococcus* spp. growth. This is persuasive evidence denoting that viruses are important agents in controlling nutrient cycle in various aquatic environments.

A potential problem associated with reduced *Synechococcus* spp. growth in diluted water has been recognized in a recent study (Tsai et al. 2018). The study mentioned that *Synechococcus* spp. present in oligotrophic waters may depend on nutrients regenerated through trophic interactions between heterotrophic bacteria and grazers or viruses. However, initial experiments in this study of Tsai et al. (2018) were not performed to assess viral-induced mortality in heterotrophic bacteria, and therefore, it may not provide valuable insights into the source of nutrients that *Synechococcus* spp. primarily depends on. This study hypothesizes that there are different growth responses present between heterotrophic bacteria and *Synechococcus* spp. population, and tests the hypothesis by using the modified dilution approach. It aims to study how viral lysis and nanoflagellate grazing affect the mortality of heterotrophic bacteria and *Synechococcus* spp.

## 2. MATERIALS AND METHODS

### 2.1 Sampling

Samples were collected on 14 and 18 July 2019, between 8 am and 9 am (local time) from the surface water of the coastal area of north-eastern Taiwan (25°09.4'N, 121°46.3'E). Previous long-term sampling at this study site has revealed significant seasonal variability in temperature, nutrients and picoplankton abundance (Tsai et al. 2005, 2008, 2012, 2013). Water temperature was measured immediately after the sampling bucket was cast. All samples were transported to the laboratory within 30 min.

### 2.2 Modified Dilution Approach

In order to measure the contribution of grazers and viruses in association with heterotrophic bacteria and *Synechococcus* spp. mortality, modified dilution method (Evans et al. 2003) was conducted. Sequential dilution of the natural community managed with two different types of seawater filters was used to reduce predator-prey (0.2  $\mu\text{m}$ ) and viral-host (30 kDa tangential flow filtration system) encounter rates. Filtered seawater sample (< 10  $\mu\text{m}$ ) was then diluted

with the 0.2  $\mu\text{m}$  and 30 kDa filtered seawater in a 4-point dilution series: 25, 50, 75, and 100% seawater (< 10  $\mu\text{m}$ ). The size fractionation used for grazers (< 10  $\mu\text{m}$ ) was chosen based on previous studies at this site to eliminate ciliates but not nanoflagellates (Tsai et al. 2012). Two modified dilution experiments were performed during the warm season (July), following the details of experimental set-up and incubation conditions described in Tsai et al. (2018). All treatments were incubated for 12 h in triplicate in 200 mL polycarbonate bottles under natural light in a water bath set at the *in situ* temperature of the seawater at the time of sampling. In the present study, nutrients were not added to the experimental bottles due to the following reasons: (1) to determine bacterial and *Synechococcus* spp. growth and the effect of grazing and viral lysis without involving the bias of nutrient addition; (2) to highlight the importance of the effects of recycling nutrients which are required for *Synechococcus* spp. growth.

Bacteria and *Synechococcus* spp. were counted using an epifluorescence microscope (Nikon Optiphot-2; 1000 $\times$ ) and abundance count was also conducted as described in Tsai et al. (2005). Significance of the regression lines was tested using analysis of variance (ANOVA). Moreover, the significance between the slopes of the 30 kDa and 0.2  $\mu\text{m}$  dilution series was determined using the *F*-test. If the regression slopes of 30 kDa and 0.2  $\mu\text{m}$  dilution series were significantly different, we then calculated the magnitude of viral lysis. STATISTICA 7.0 software was used to perform all statistical operations. A probability value of < 0.05 was considered significant.

## 3. RESULTS AND DISCUSSION

Both sampling procedures maintained the same temperature range (28.5 - 29°C). Bacterial abundance measured in the collected seawater (14 and 28 July) was  $6 \times 10^5$  and  $8 \times 10^5$  cells  $\text{mL}^{-1}$ , respectively. Furthermore, abundance of *Synechococcus* spp. was  $5 \times 10^4$  and  $7 \times 10^4$  cells  $\text{mL}^{-1}$  on 14 and 28 July, respectively. A good linearity with negative slope was found while measuring bacterial net growth rate in both dilution series (Figs. 1a and b). The regression coefficients (slopes) were 0.056 and 0.065  $\text{h}^{-1}$  for the 0.2  $\mu\text{m}$  fractionated series, and 0.10 and 0.172  $\text{h}^{-1}$  for the 30 kDa fractionated series in the two experiments (Figs. 1a and b). Additionally, there were significant differences noted between the two slopes of regression lines as seen in both experiments (*F*-test,  $p < 0.05$ ) (Figs. 1a and b). The difference between the two slope values were 0.044 and 0.107  $\text{h}^{-1}$ , indicating that bacterial mortality was caused mainly due to lysis of viruses. In the present study, we observed higher net growth rate of bacteria in the 30 kDa series than that in the 0.2  $\mu\text{m}$  series (Figs. 1a and b), which suggests that under these conditions lytic pressure is relatively high (Baudoux et al. 2008). Furthermore, the y-intercepts of these regression lines were 0.09

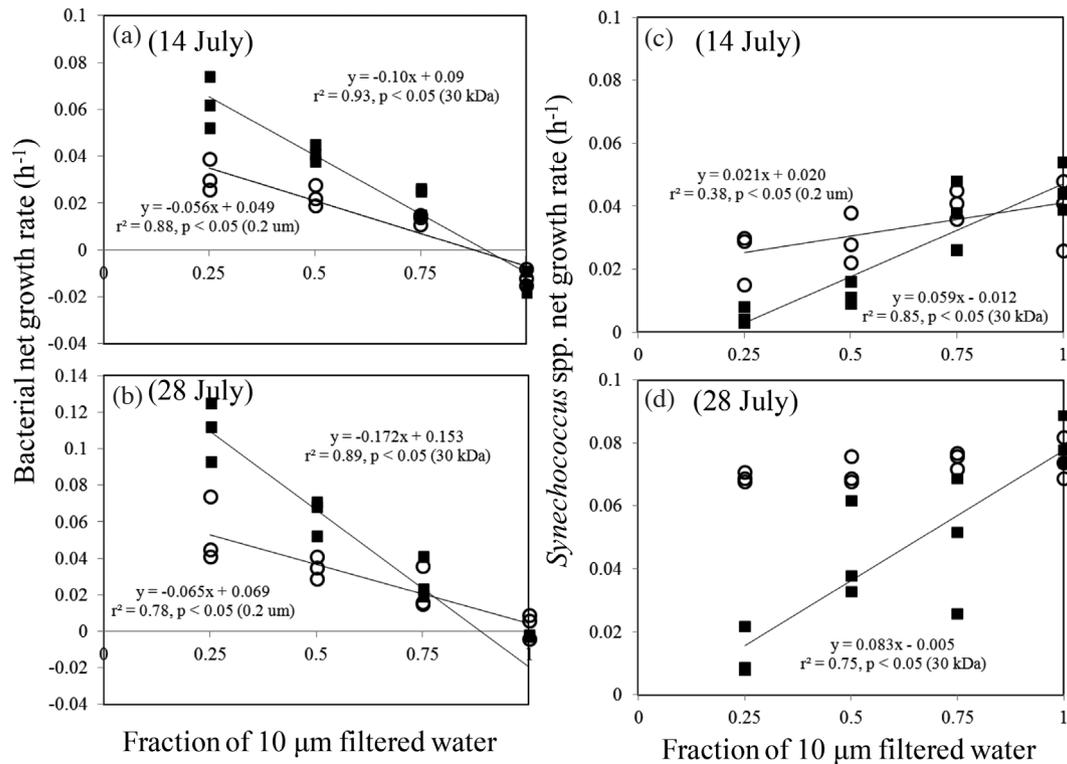


Fig. 1. Net growth rate vs. fraction of 10 µm filtered water for the parallel dilution experiments for bacteria (a) (b) and *Synechococcus* spp. were conducted on 14th and 28th July. Experiments were performed in 30 kDa ultra filtrate (■) and 0.2 µm filtered seawater (○).

and 0.153 h<sup>-1</sup>, respectively for 30 kDa fractionated series in both experiments (Figs. 1a and b). These values represent the bacterial growth rate in the absence of grazing pressure and viral lysis.

We investigated the role of *Synechococcus* spp. in association with growth incubation experiments, and the results confirmed a positive relationship between net growth rate of *Synechococcus* spp. and dilution factor, implying that the assumption of the dilution method was not met (Figs. 1c and d). Some modified dilution experiments have been tested to estimate viral lysis rates of phytoplankton (Evans et al. 2003; Tsai et al. 2018). This study did not estimate the effect of viral lysis on *Synechococcus* spp., as it was suspected that *Synechococcus* spp. might not be top-down controlled by viral lysis or nanoflagellate grazing. In this study, it was evident that *Synechococcus* spp. growth was bottom-up controlled with nutrient limitation being an important factor. This study also confirmed that different growth responses exist between heterotrophic bacteria and *Synechococcus* spp. population by using the modified dilution approach. Interestingly, our results reported that the net growth rate of *Synechococcus* spp. in 30 kDa dilution series was lower than that in 0.2 µm series (Figs. 1c and d). Similar patterns have been observed in other studies, namely reduced net growth rate detected at higher dilution levels; this suggests possible nutrient limitation during their

research (Baudoux et al. 2007; Shelford et al. 2012; Tsai et al. 2018). Concerning the different patterns of heterotrophic bacteria and *Synechococcus* spp. in response to dilution experiments, the explanation for reduced *Synechococcus* spp. growth rate in 30 kDa dilution series is still unclear. This finding is consistent with previous studies (Weinbauer et al. 2011; Shelford et al. 2012), which pointed out that viruses enhance the growth rate of *Synechococcus* spp. These results suggested an indirect effect of viral lysis of heterotrophic bacteria. Over the past decades, there has been an increased focus to study the importance of nutrients regeneration in marine environments, and the role of viruses in this process (Ayukai 1996; Weinbauer et al. 2011; Shelford et al. 2012; Gong and Tsai 2019).

This study found that nanoflagellate grazing and viral lysis both are the major factors for mortality of bacteria in the experiments, with an average of 0.06 and 0.08 h<sup>-1</sup>, respectively (Fig. 2). Many studies on the degree of viral lysis and grazing have focused on heterotrophic bacteria (Jacquet et al. 2005; Fischer et al. 2006; Wells and Deming 2006; Tsai et al. 2013), and reported that the impact of viral lysis and grazing varies according to divergent factors such as different seasons, environments and host organisms. Moreover, loss of bacterial cells either through grazing or due to viral lysis can be expected to have different biogeochemical and ecological consequences. It is observed that grazing leads

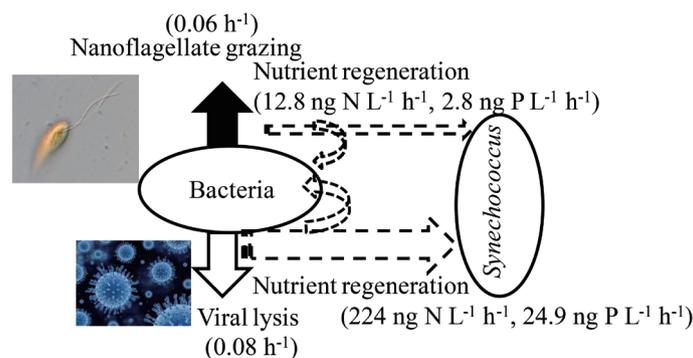


Fig. 2. Picture of nanoflagellate grazing and viral lysis of bacteria, their contribution in balancing carbon flux, and maintaining the release of nutrients for bacterial and *Synechococcus* spp. growth.

to transfer of energy and biomass partly to higher trophic levels, whereas viral lysis is considered to recycle carbon and nutrients within the microbial food web (Suttle 2007). Concerning the role of nanoflagellate grazing, besides their contribution in promoting carbon flux, they may also contribute to the release of nutrients and directly influence nitrogen (N) or phosphorus (P) cycle in water (Chase and Price 1997). Hence, rough calculation can be made from the experiments conducted in this study to assess nitrogen and phosphorus levels, as they are potentially released by the process of nanoflagellate grazing on bacteria. In this study, a carbon content of  $20 \text{ fg C cell}^{-1}$  (Lee and Fuhrman 1987) and C:N:P ratio of 45:9:1, respectively for bacteria (Goldman et al. 1987) was used to estimate the concentration of nitrogen and phosphorus released by grazing or viral lysis. Additionally, based on the average heterotrophic bacterial biomass ( $14 \mu\text{g C L}^{-1}$ ) and grazing rate ( $0.06 \text{ h}^{-1}$ ), we estimated that about  $12.8 \text{ ng N L}^{-1} \text{ h}^{-1}$  and  $2.8 \text{ ng P L}^{-1} \text{ h}^{-1}$  was released by nanoflagellate grazing, which highlights that nanoflagellate can release a significant fraction (10% of N and 20% of P) of ingested prey organic matter (Caron et al. 1990; Ferrier-Pagès et al. 1998) (Fig. 2).

In our study on the interactions between heterotrophic bacteria, viruses and *Synechococcus* spp., the results demonstrated that if viral lysis alone was taken into consideration, viruses were responsible for 57% of the heterotrophic bacterial mortality [ $(0.08)/(0.06 + 0.08) \times 100\%$ ] (Fig. 2). This study also found that reduced viral abundance at higher dilution factor decreased the growth of *Synechococcus* spp. (Figs. 1c and d), which was evidenced by the fact that viral lysis of heterotrophic bacteria potentially benefits *Synechococcus* spp. growth by supplying nutrients. Based on the estimations suggested above and the average rate of viral lysis ( $0.08 \text{ h}^{-1}$ ) (Fig. 2), viral process was estimated to release locally recycled nutrients of  $224 \text{ ng N L}^{-1} \text{ h}^{-1}$  and  $24.9 \text{ ng P L}^{-1} \text{ h}^{-1}$  in this system (Fig. 2). Due to viral lysis, substantial portion of the recycled nutrients required for *Synechococcus* spp. growth is properly supplied, emphasize-

ing the importance of the viral shunt, especially in oligotrophic regions. However, non-exclusive, heterotrophic bacteria can compete with *Synechococcus* spp. for these recycled nutrients (Kirchman and Wheeler 1998) thus resulting in reduced nutrients availability to *Synechococcus* spp. Interestingly, we observed that the growth rate of *Synechococcus* spp. in unfiltered water (100% fraction of whole water) was about  $0.03 - 0.09 \text{ h}^{-1}$  (Figs. 1c and d), suggesting that the processes of grazing and viral lysis releases enough nutrients required to maintain *Synechococcus* spp. growth. Similarly, previous studies have reported that *Synechococcus* spp. displayed higher proportion of dividing *Synechococcus* spp. cells (FDC) in virus existing treatment than that in the virus-reduced treatment (37% vs. 5%) (Tsai et al. 2014).

In conclusion, we confirmed the hypothesis that there are different growth responses present between heterotrophic bacteria and *Synechococcus* spp. population by using the modified dilution approach. In the present study, the results showed that nanoflagellate grazing and viral lysis were the major factors causing mortality of bacteria in the month of July. However, it was noticed that *Synechococcus* spp. may not have been top-down controlled by viral lysis or nanoflagellate grazing. During this study period, it was evident that *Synechococcus* spp. growth was bottom-up controlled with nutrient limitation being an important factor in the study, and viral lysis of heterotrophic bacteria potentially benefited *Synechococcus* spp. growth by supplying nutrients.

**Acknowledgements** This study was supported by a grant (NSC 108-2611-M-019-013) received from the Ministry of Science and Technology, ROC. We appreciate the language editing and helpful comments obtained from Choice Language Service related to this manuscript.

## REFERENCES

Agawin, N. S. R., C. M. Duarte, and S. Agustí, 2000: Nutrient and temperature control of the contribution of

- picoplankton to phytoplankton biomass and production. *Limnol. Oceanogr.*, **45**, 591-600, doi: 10.4319/lo.2000.45.3.0591. [[Link](#)]
- Anderson, S. R., Q. P. Diou-Cass, and E. L. Harvey, 2018: Short-term estimates of phytoplankton growth and mortality in a tidal estuary. *Limnol. Oceanogr.*, **63**, 2411-2422, doi: 10.1002/lno.10948. [[Link](#)]
- Ayukai, T., 1996: Possible limitation of the dilution technique for estimating growth and grazing mortality rates of picoplanktonic cyanobacteria in oligotrophic tropical waters. *J. Exp. Mar. Biol. Ecol.*, **198**, 101-111, doi: 10.1016/0022-0981(95)00208-1. [[Link](#)]
- Baudoux, A.-C., M. J. W. Veldhuis, H. J. Witte, and C. P. D. Brussaard, 2007: Viruses as mortality agents of picophytoplankton in the deep chlorophyll maximum layer during IRONAGES III. *Limnol. Oceanogr.*, **52**, 2519-2529, doi: 10.4319/lo.2007.52.6.2519. [[Link](#)]
- Baudoux, A.-C., M. J. W. Veldhuis, A. A. M. Noordeloos, G. van Noort, and C. P. D. Brussaard, 2008: Estimates of virus- vs. grazing induced mortality of picophytoplankton in the North Sea during summer. *Aquat. Microb. Ecol.*, **52**, 69-82, doi: 10.3354/ame01207. [[Link](#)]
- Bell, T. and J. Kalff, 2001: The contribution of picophytoplankton in marine and freshwater systems of different trophic status and depth. *Limnol. Oceanogr.*, **46**, 1243-1248, doi: 10.4319/lo.2001.46.5.1243. [[Link](#)]
- Caron, D. A., K. G. Porter, and R. W. Sanders, 1990: Carbon, nitrogen, and phosphorus budgets for the mixotrophic phytoflagellate *Poteroiochromonas malhamensis* (Chrysophyceae) during bacterial ingestion. *Limnol. Oceanogr.*, **35**, 433-443, doi: 10.4319/lo.1990.35.2.0433. [[Link](#)]
- Chase, Z. and N. M. Price, 1997: Metabolic consequences of iron deficiency in heterotrophic marine protozoa. *Limnol. Oceanogr.*, **42**, 1673-1684, doi: 10.4319/lo.1997.42.8.1673. [[Link](#)]
- Evans, C., S. D. Archer, S. Jacquet, and W. H. Wilson, 2003: Direct estimates of the contribution of viral lysis and microzooplankton grazing to the decline of a *Micromonas* spp. population. *Aquat. Microb. Ecol.*, **30**, 207-219, doi: 10.3354/ame030207. [[Link](#)]
- Ferrier-Pagès, C., M. Karner, and F. Rassoulzadegan, 1998: Release of dissolved amino acids by flagellates and ciliates grazing on bacteria. *Oceanol. Acta*, **21**, 485-494, doi: 10.1016/S0399-1784(98)80032-7. [[Link](#)]
- Fischer, U. R., C. Wieltchnig, A. K. T. Kirschner, and B. Velimirov, 2006: Contribution of virus-induced lysis and protozoan grazing to benthic bacterial mortality estimated simultaneously in microcosms. *Environ. Microbiol.*, **8**, 1394-1407, doi: 10.1111/j.1462-2920.2006.01032.x. [[Link](#)]
- Goldman, J. C., D. A. Caron, and M. R. Dennett, 1987: Regulation of gross growth efficiency and ammonium regeneration in bacteria by substrate C:N ratio. *Limnol. Oceanogr.*, **32**, 1239-1252, doi: 10.4319/lo.1987.32.6.1239. [[Link](#)]
- Gong, G.-C. and A.-Y. Tsai, 2019: Reduced daytime net growth rate of *Synechococcus* spp. in the East China Sea in summer estimated using a dilution approach. *Estuar. Coast. Shelf Sci.*, **219**, 90-96, doi: 10.1016/j.ecss.2019.01.029. [[Link](#)]
- Grob, C., M. Hartmann, M. V. Zubkov, and D. J. Scanlan, 2011: Invariable biomass-specific primary production of taxonomically discrete picoeukaryote groups across the Atlantic Ocean. *Environ. Microbiol.*, **13**, 3266-3274, doi: 10.1111/j.1462-2920.2011.02586.x. [[Link](#)]
- Jacquet, S., I. Domaizon, S. Personnic, A. S. Pradeep Ram, M. Hedal, S. Duhamel, and T. Sime-Ngando, 2005: Estimates of protozoan- and viral-mediated mortality of bacterioplankton in Lake Bourget (France). *Freshwat. Biol.*, **50**, 627-645, doi: 10.1111/j.1365-2427.2005.01349.x. [[Link](#)]
- Jardillier, L., M. V. Zubkov, J. Pearman, and D. J. Scanlan, 2010: Significant CO<sub>2</sub> fixation by small prymnesiophytes in the subtropical and tropical northeast Atlantic Ocean. *The ISME Journal*, **4**, 1180-1192, doi: 10.1038/ismej.2010.36. [[Link](#)]
- Juhl, A. R. and M. C. Murrell, 2005: Interactions between nutrients, phytoplankton growth, and microzooplankton grazing in a Gulf of Mexico estuary. *Aquat. Microb. Ecol.*, **38**, 147-156, doi: 10.3354/ame038147. [[Link](#)]
- Kirchman, D. L. and P. A. Wheeler, 1998: Uptake of ammonium and nitrate by heterotrophic bacteria and phytoplankton in the sub-Arctic Pacific. *Deep-Sea Res. Part I-Oceanogr. Res. Pap.*, **45**, 347-365, doi: 10.1016/S0967-0637(97)00075-7. [[Link](#)]
- Landry, M. R. and R. P. Hassett, 1982: Estimating the grazing impact of marine micro-zooplankton. *Mar. Biol.*, **67**, 283-288, doi: 10.1007/BF00397668. [[Link](#)]
- Lee, S. and J. A. Fuhrman, 1987: Relationships between biovolume and biomass of naturally derived marine bacterioplankton. *Appl. Environ. Microbiol.*, **53**, 1298-1303.
- Redden, A. M., B. G. Sanderson, and D. Rissik, 2002: Extending the analysis of the dilution method to obtain the phytoplankton concentration at which microzooplankton grazing becomes saturated. *Mar. Ecol. Prog. Ser.*, **226**, 27-33, doi: 10.3354/meps226027. [[Link](#)]
- Shelford, E. J., M. Middelboe, E. F. Møller, and C. A. Suttle, 2012: Virus-driven nitrogen cycling enhances phytoplankton growth. *Aquat. Microb. Ecol.*, **66**, 41-46, doi: 10.3354/ame01553. [[Link](#)]
- Suttle, C. A., 2007: Marine viruses — major players in the global ecosystem. *Nat. Rev. Microbiol.*, **5**, 801-812, doi: 10.1038/nrmicro1750. [[Link](#)]
- Tsai, A.-Y., K.-P. Chiang, J. Chang, and G.-C. Gong, 2005: Seasonal diel variations of picoplankton and

- nanoplankton in a subtropical Western Pacific coastal ecosystem. *Limnol. Oceanogr.*, **50**, 1221-1231, doi: 10.4319/lo.2005.50.4.1221. [[Link](#)]
- Tsai, A.-Y., K.-P. Chiang, J. Chang, and G.-C. Gong, 2008: Seasonal variations in trophic dynamics of nanoflagellates and picoplankton in coastal waters of the western subtropical Pacific Ocean. *Aquat. Microb. Ecol.*, **51**, 263-274, doi: 10.3354/ame01196. [[Link](#)]
- Tsai, A.-Y., G.-C. Gong, R. W. Sanders, K.-P. Chiang, J.-K. Huang, and Y.-F. Chan, 2012: Viral lysis and nanoflagellate grazing as factors controlling diel variations of *Synechococcus* spp. summer abundance in coastal waters of Taiwan. *Aquat. Microb. Ecol.*, **66**, 159-167, doi: 10.3354/ame01566. [[Link](#)]
- Tsai, A.-Y., G.-C. Gong, and J. Hung, 2013: Seasonal variations of virus- and nanoflagellate-mediated mortality of heterotrophic bacteria in the coastal ecosystem of subtropical western Pacific. *Biogeosciences*, **10**, 3055-3065, doi: 10.5194/bg-10-3055-2013. [[Link](#)]
- Tsai, A.-Y., G.-C. Gong, and Y.-W. Huang, 2014: Importance of the viral shunt in nitrogen cycling in *Synechococcus* spp. growth in subtropical western Pacific coastal waters. *Terr. Atmos. Ocean. Sci.*, **25**, 839-846, doi: 10.3319/TAO.2014.06.11.01(Oc). [[Link](#)]
- Tsai, A.-Y., G.-C. Gong, C.-C. Chung, and Y.-T. Huang, 2018: Different impact of nanoflagellate grazing and viral lysis on *Synechococcus* spp. and picoeukaryotic mortality in coastal waters. *Estuar. Coast. Shelf Sci.*, **209**, 1-6, doi: 10.1016/j.ecss.2018.05.012. [[Link](#)]
- Weinbauer, M. G., O. Bonilla-Findji, A. M. Chan, J. R. Dolan, S. M. Short, K. Šimek, S. W. Wilhelm, and C. A. Suttle, 2011: *Synechococcus* growth in the ocean may depend on the lysis of heterotrophic bacteria. *J. Plankton Res.*, **33**, 1465-1476, doi: 10.1093/plankt/fbr041. [[Link](#)]
- Wells, L. E. and J. W. Deming, 2006: Significance of bacterivory and viral lysis in bottom waters of Franklin Bay, Canadian Arctic, during winter. *Aquat. Microb. Ecol.*, **43**, 209-221, doi: 10.3354/ame043209. [[Link](#)]
- Winder, M., 2009: Photosynthetic picoplankton dynamics in Lake Tahoe: Temporal and spatial niche partitioning among prokaryotic and eukaryotic cells. *J. Plankton Res.*, **31**, 1307-1320, doi: 10.1093/plankt/fbp074. [[Link](#)]