

Research



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What limits photosynthetic energy conversion efficiency in nature? Lessons from the oceans

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Constraining photosynthetic energy conversion efficiency in nature is challenging. In principle, two yield measurements must be made simultaneously: photochemistry, fluorescence and/or thermal dissipation. We constructed two different, extremely sensitive and precise active fluorometers: one measures the quantum yield of photochemistry from changes in variable fluorescence, the other measures fluorescence lifetimes in the picosecond time domain. By deploying the pair of instruments on eight transoceanic cruises over six years, we obtained over 200 000 measurements of fluorescence yields and lifetimes from surface waters in five ocean basins. Our results revealed that the average quantum yield of photochemistry was approximately 0.35 while the average quantum yield of fluorescence was approximately 0.07. Thus, closure on the energy budget suggests that, on average, approximately 58% of the photons absorbed by phytoplankton in the world oceans are dissipated as heat. This extraordinary inefficiency is associated with the paucity of nutrients in the upper ocean, especially dissolved inorganic nitrogen and iron. Our results strongly suggest that, in nature, most of the time, most of the phytoplankton community operates at approximately half of its maximal photosynthetic energy conversion efficiency because nutrients limit the synthesis or function of essential components in the photosynthetic apparatus.

This article is part of the themed issue 'Enhancing photosynthesis in crop plants: targets for improvement'.

1. Introduction

The conversion of one form of energy to another inevitably comes at a thermodynamic cost of heat. The energy conversion efficiency in photosynthesis has been inferred from several different experimental protocols, including photoacoustics [1–3], amplitude-based variable fluorescence [4–6] and fluorescence lifetimes [7], yet seldom has there been closure on the energy budget in model organisms, let alone in nature. Fundamentally, closure can be achieved by measuring any two of the three rate constants that ultimately determine the overall quantum yield. The basic argument is that photons absorbed by a photosynthetic reaction centre have only three possible fates: they can be used to drive an electrochemical reaction, reradiated as fluorescence, or dissipated as heat [8,9]. In unicellular algae, the first two reactions can be precisely measured using two active fluorescence techniques, namely amplitude and lifetime analyses, respectively (figure 1). Moreover, by using very sensitive detectors, these two signals can be obtained from natural phytoplankton communities in the global oceans [12].

Phytoplankton are a highly diverse group of unicellular photosynthetic algae, including cyanobacteria, capable of oxygenic photosynthesis [14]. One clade of these organisms gave rise to higher plants, but phytoplankton are much more diverse: there are one prokaryotic and thirteen eukaryotic phyla containing

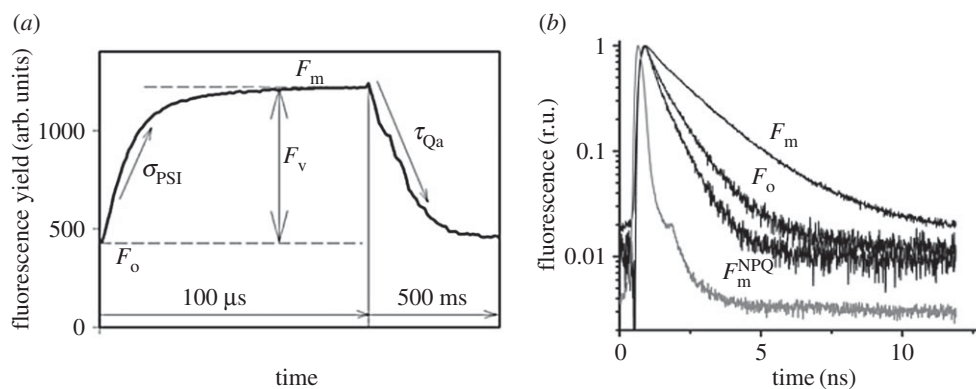


Figure 1. The concept of amplitude-based variable fluorescence (a) and picosecond lifetime (b) measurements. (a) Amplitude-based variable fluorescence techniques, such as fast repetition rate (FRR) or fluorescence induction and relaxation (FIRE), rely on recording the changes in fluorescence intensities induced by saturating flashes of excitation light on micro- and milli-second time scales [10,11]. The quantum yield of photochemistry in photosystem II (PSII) (i.e. photosynthetic efficiency) is deduced from the relative change in fluorescence yield (F_v/F_m). The effective absorption cross section of PSII (σ_{PSII}) is derived from the rate of fluorescence rise during the first 100 μs of the induction curve. The subsequent relaxation in fluorescence yield on millisecond time scale reflects the rates of photosynthetic electron transport delivered on the acceptor side of PSII. Most of those electrons are used in carbon fixation. All amplitude-based fluorometers record fluorescence yields in arbitrary units and do not measure absolute quantum yields of fluorescence. (b) Picosecond-resolved measurements of fluorescence lifetimes—fluorescence is induced by an ultra-short laser pulse (grey line) and the decay kinetics is recorded on pico- and nano-second time scales. The rate of fluorescence decay (i.e. lifetime) is directly proportional to the absolute quantum yield of fluorescence. The three profiles show fluorescence kinetics of phytoplankton at different physiological states of photosystem II reaction centres. F_0 was recorded in dark-adapted cells with open reaction centres (when the quantum yield of photochemistry is maximum), F_m in dark-adapted cells with closed centres (when photochemistry is nil), F_m^{NPQ} under high light exposure ($1500 \mu\text{mol quanta m}^{-2} \text{s}^{-1}$) with closed centres. The reduction in the fluorescence lifetime under high light is due to non-photochemical quenching (NPQ) [12,13]. The grey trace is the instrument response function. Both variable fluorescence and lifetime instruments (namely a mini-FIRE and picosecond lifetime fluorometer (PicoLiF), respectively [12]) have extremely high sensitivity; they can accurately measure fluorescence signals in samples as low as $0.01 \text{ mg chlorophyll/m}^3$ —effectively the lowest concentration of chlorophyll recorded anywhere in the upper ocean.

oxygenic photosynthetic species [15]. All of these have the same two photosynthetic reaction centres as found in plants, but are highly diverse in the pigmentation in their light harvesting antenna systems. Regardless of antenna pigmentation however, the maximum photosynthetic energy conversion efficiency in photosystem II (PSII), obtained when grown under optimal conditions in cultures, is remarkably constant. When measured with a single turnover saturating flash (e.g. approx. 100 μs duration), variable fluorescence normalized to the maximal fluorescence (F_v/F_m) is about 0.65 [16]. Assuming that the first order loss processes to fluorescence have the quantum yield of ca. 0.05, this suggests that about 30% of the absorbed excitation energy is dissipated as heat in PSII. The efficiency of photosystem I (PSI), which does not fluoresce at room temperature, is much more difficult to assess in phytoplankton [3] because their concentration is usually so low in natural waters that absorption spectra cannot be easily obtained [17]. Regardless, as PSI operates in series with PSII, its efficiency can only reduce the overall calculation of photosynthetic energy conversion efficiency.

2. Variability and controls of photosynthetic energy conversion efficiency in the ocean

Based on numerous experiments with cultured phytoplankton, if allowed sufficient time to photoacclimate, F_v/F_m is insensitive to growth irradiance, but not to nutrient insufficiency [16,18]. Nitrate and iron, in particular, are two nutrients that can lead to marked reductions in F_v/F_m . Based on laboratory experiments, analyses of natural variations in F_v/F_m strongly indicate that in every major ocean gyre there is significant nutrient limitation—but it is highly patchy [14]. That is, in some regions of the ocean, injection of nutrients from deep waters due to (e.g.) storm events or eddy pumping, can

temporarily lead to elevated F_v/F_m values, approaching those obtained under optimal growth conditions in the laboratory or observed in nutrient-rich coastal waters. This interpretation was further supported by deliberate iron fertilization experiments in high-nutrient-low-chlorophyll (HNLC) regions of the open ocean [19–22]. To date, thirteen iron fertilization experiments have been conducted in the three major iron limited regions of the world's oceans: the Subarctic Pacific, the eastern Equatorial Pacific and the Southern Ocean [23,24]. An increase in F_v/F_m was the first detectable signal of release from iron deficiency in every iron enrichment experiment.

We obtained more than 200 000 discrete measurements of variable fluorescence from the North and South Pacific, North Atlantic, Arctic and Southern Oceans. These measurements comprise the global map of photosynthetic energy conversion efficiencies in phytoplankton in the upper ocean (figure 2). The nighttime *in situ* measurements of F_v/F_m ranged from 0.12 to 0.60 with a mean of 0.35 ± 0.11 . These values span virtually the entire range of published values of photosynthetic conversion efficiencies obtained from cultured phytoplankton by single turnover flashes, and reflect extraordinary variability in phytoplankton photophysiology in the global ocean. F_v/F_m values in ocean phytoplankton communities are two- to three-fold lower than those observed in higher plants, suggesting the fundamental difference in nutrient availability between terrestrial and aquatic ecosystems.

The highest F_v/F_m values (more than 0.4) were observed along continental margins, in the Antarctic convergence, the Subtropical Atlantic and Pacific oceans. In contrast the lowest photosynthetic efficiencies were observed in HNLC regions of the equatorial Pacific Ocean and the Southern Ocean, where primary production is limited by a paucity of iron [25,26], a micronutrient that is critical for the function of photosystem II [18]. The exceptionally low F_v/F_m values

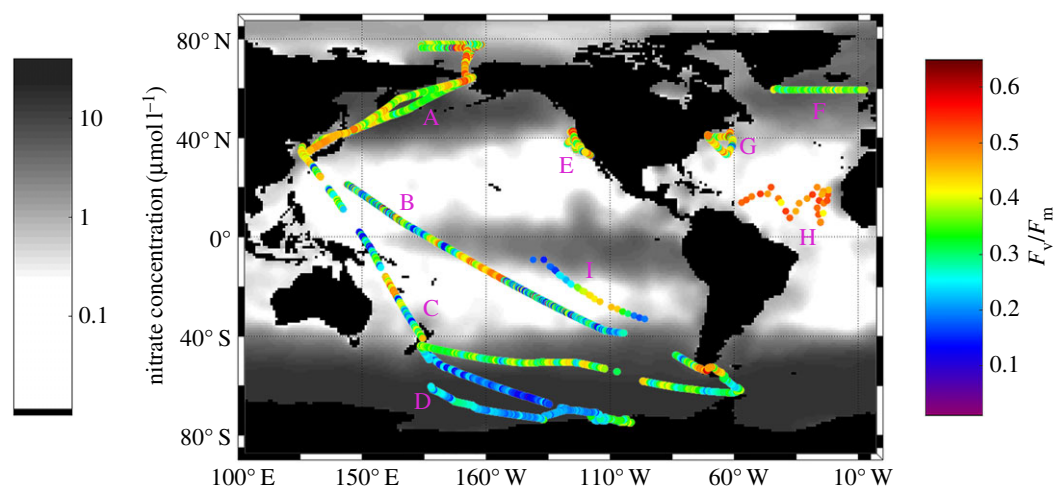


Figure 2. Global distribution of ship-based measurements of the photosynthetic energy conversion efficiencies (F_v/F_m) in the upper ocean superimposed on the climatological map of surface nitrate concentrations in the world ocean. Periods of *in situ* sampling were (A) July–Aug 2011; (B) Oct–Nov 2011; (C) Oct–Nov 2010; (D) Jan 2012; (E) July 2014; (F) Sept 2010; (G) May 2014; (H) Aug 2008; (I) Nov. 2004.

(less than 0.20) in these regions are consistent with extremely long fluorescence lifetimes and suggest that there is a significant fraction of non-functional PSII reaction centres and/or energetically uncoupled antenna pigment–protein complexes [18,27,28].

With literally hundreds of thousands of measurements of F_v/F_m , and other parameters that can be derived from variable fluorescence analysis of single and multiple turnover flashes, including the effective absorption cross section of PSII and the electron transport rate on the acceptor side of the reaction centre, we turned to fluorescence lifetime analyses as a possible approach to retrieving the absolute quantum yields of fluorescence and to closing the energy budget for phytoplankton in nature.

3. Theoretical bases of fluorescence yields and lifetimes

The quantum yield of fluorescence (ϕ_f) is defined as the ratio of the photons reradiated to those absorbed. The biophysical basis of fluorescence measurements derives from the three possible fates of solar energy absorbed by any photosynthetic organism [9]. Absorbed photons can (i) generate photochemical reactions (with the rate k_p), (ii) be dissipated as heat (k_t), or (iii) be emitted back to the environment as fluorescence (k_f). The rate k_p is at first order proportional to the fraction of open or active reaction centres. The rate k_t is the sum of dark component (k_D) and light-dependent component (k_{NPQ}), driven by non-photochemical quenching.

In a dark-adapted state or under low irradiance (when k_{NPQ} is nil and k_t is constant), the quantum yield of chlorophyll fluorescence, $\phi_f (=k_f/(k_p + k_t + k_f))$, is inversely related to the quantum yield of photochemistry in PSII, $\phi_p = k_p/(k_p + k_t + k_f) = F_v/F_m$:

$$\phi_f = \phi_{fm} \left(1 - \frac{F_v}{F_m} \right), \quad (3.1)$$

where $\phi_{fm} (=k_f/(k_t + k_f))$ is the maximum fluorescence yield obtained when the quantum yield of photochemistry is nil (e.g. at saturating background light).

This biophysical model predicts an inverse linear relationship between the quantum yield of photochemistry and that

of chlorophyll fluorescence. However, by the early 1980s it was realized that exposure to high irradiance can generate a suite of thermal dissipation mechanisms, collectively called non-photochemical quenching (NPQ) [29,30]. This photoprotective response markedly decreases the quantum yield of chlorophyll fluorescence at high background light. Hence, the relationship between fluorescence yield and photochemistry becomes highly nonlinear as NPQ phenomena play an increasingly larger role in energy dissipation.

Fluorescence is a delayed light emission process, which is described by one or more exponential decay functions that can be parameterized by the lifetime, which is the *e*-folding time of the decay function. The fluorescence lifetime can be quantitatively related to the absolute quantum yield of fluorescence [31]:

$$\phi_f = \frac{\tau}{\tau_n}, \quad (3.2)$$

where τ is the observed lifetime and τ_n is the intrinsic (or ‘natural’) lifetime constant for the molecule. Thus, the longer the lifetime, the higher the quantum yield of fluorescence.

The ‘natural’ lifetime (τ_n) is that which would be observed if fluorescence emission were the only path of dissipation of excited state energy; this number cannot be measured directly. The calculated value of τ_n (which is a constant for a specific molecule) for chlorophyll *a* is 15 ns [32]. In a population of molecules, the actual measured lifetimes are inevitably shorter than the ‘natural’ lifetime due to intra-molecular conversion (i.e. energy dissipation as heat) and triplet state formation. The actual measured lifetimes of isolated chlorophyll *a* molecules range from ca. 3.0 to 5.1 ns, depending on the solvent polarity. These measured lifetimes correspond to quantum yields ranging from 20 to 32%. Fluorescence lifetimes in living cells are even shorter (ca. 0.3 to approx. 1.5 ns), as a significant fraction of the absorbed energy is used in photochemical reactions [7], and reflect the physiological state of the cells.

4. Lifetimes and quantum yields of chlorophyll fluorescence in the ocean

Accurate, optical measurements of the quantum yield of fluorescence are among the most complicated measurements to

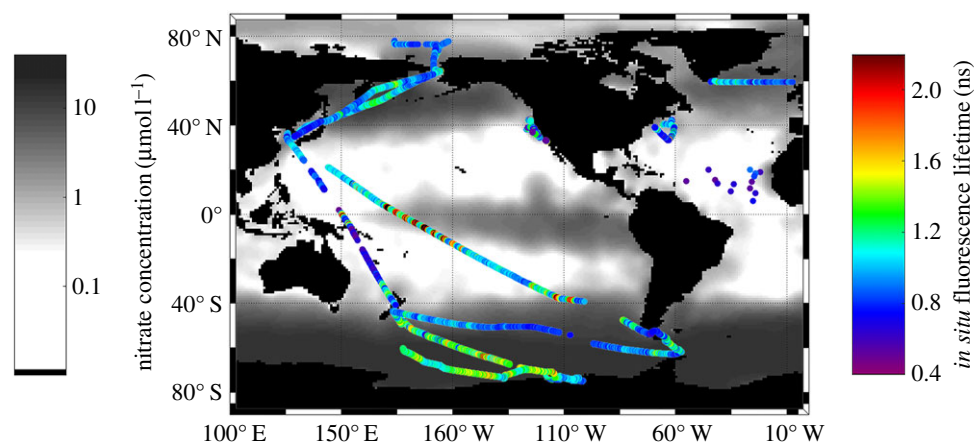


Figure 3. Global distribution of ship-based measurements of chlorophyll fluorescence lifetimes in the upper ocean superimposed on the climatological map of surface nitrate concentrations in the world ocean. (Data from [12]).

obtain in applied physics. By definition, such measurements require precise determination of the number of quanta absorbed by the sample and the number of quanta emitted over the entire fluorescence spectrum integrated over 4π steradians. In living phytoplankton cells, the measurements are further complicated by the ‘packaging’ effect (i.e. self-shading) of photosynthetic pigments embedded within layers of thylakoid membranes, as well as scattering and refraction of light by cell walls and membranes [32]. The measurement of fluorescence lifetimes conveniently circumvents all these potential obstacles.

To that end, we constructed a sea-going picosecond lifetime instrument that employs a time-correlated single photon counting detector [12]. The instrument has extremely high sensitivity; it can measure lifetimes in samples as low as $0.01 \text{ mg chlorophyll/m}^3$ —effectively the lowest concentration of chlorophyll recorded anywhere in the upper ocean. Furthermore, this instrument employs a multi-component kinetic analysis, which is crucial for interpretation of fluorescence kinetics from a living cell [7]. We obtained more than 200 000 discrete chlorophyll fluorescence lifetime measurements from the Pacific, Atlantic, Arctic and Southern (Antarctic) Oceans. These comprise the first map of *in situ* measurements of quantum yields of chlorophyll fluorescence from phytoplankton in the upper ocean (figure 3). The nighttime *in situ* lifetimes ranged from 0.5 to 2.7 ns with a mean of $1.13 \pm 0.33 \text{ ns}$ [12]. These values span the entire range of published lifetimes of *in vivo* chlorophyll fluorescence obtained from cultured phytoplankton and reflect extraordinary variability in phytoplankton photophysiology and nutrient status in the global ocean. The general pattern of the fluorescence lifetimes in the central gyres of the global ocean is rather featureless [12], although phytoplankton growth is subject to both macro- and micronutrient limitation [15]. The shortest fluorescence lifetimes (less than 1 ns) were observed along continental margins, in the Antarctic convergence, the Subtropical Atlantic and Pacific oceans. These lifetime distributions support the hypothesis that phytoplankton in the central gyres are acclimated to broad scale and persistent nutrient limitation [25,33]. In contrast, the longest fluorescence lifetimes were observed in HNLC regions of the equatorial Pacific Ocean and the Southern Ocean where primary production is limited by a paucity of iron [25,26]. The exceptionally long fluorescence lifetimes (greater than 2.0 ns) in these areas of the global ocean exceed the typical

maximum lifetime values observed in algal cultures with fully closed reaction centres [34] and thus cannot, entirely, be explained by the presence of inactive PSII reaction centres alone. Therefore, our lifetime measurements suggest the presence of both a large fraction of non-functional PSII reaction centres and energetically uncoupled antenna pigment–protein complexes in Fe-limited phytoplankton communities. The combination of variable fluorescence and lifetime measurements allows us to quantify the contributions of these two modifications in the photosynthetic apparatus under nutrient limitation. This analysis revealed that up to ca. 50% of reaction centres are inactive and up to 30% of light-harvesting antennae become uncoupled in iron-limited phytoplankton in the Southern Ocean [35].

Simultaneous measurements of amplitude-based fluorescence yields and lifetimes were startling. While there was significant spatial and temporal variability, F_v/F_m in the upper ocean averaged ca. 0.35 at night, when non-photochemical quenching is nil. These results indicate that most of the contemporary open surface waters are extremely nutrient deficient. Moreover, the average lifetime obtained was about 1 ns, and only varied by ca. 10% between day, when NPQ is maximal, and night, when NPQ is nil but photochemical quenching is maximal. Assuming a natural lifetime of 15 ns, these results suggest that approximately 7% of the absorbed photosynthetic energy is reradiated as fluorescence (figure 3). Combining the two measurements suggests that approximately 60% of the absorbed excitation energy in natural phytoplankton is converted to heat (figure 4). Thus, photosynthetic energy conversion efficiency in the open ocean is almost the exact opposite of that obtained under optimal growth conditions for unicellular algae in culture.

5. Phytoplankton physiology from space

Variable fluorescence signals can be readily measured remotely in both phytoplankton and terrestrial plants using shipboard or airborne Light Detection And Ranging (LIDAR) instruments, such as pump-and-probe LIDARs [36,37]. Chlorophyll fluorescence lifetimes can potentially be recorded in terrestrial plants, but not from phytoplankton in the water column, using Picosecond LIDARs [38,39]. In terrestrial ecosystems however, the operational use of such instruments is

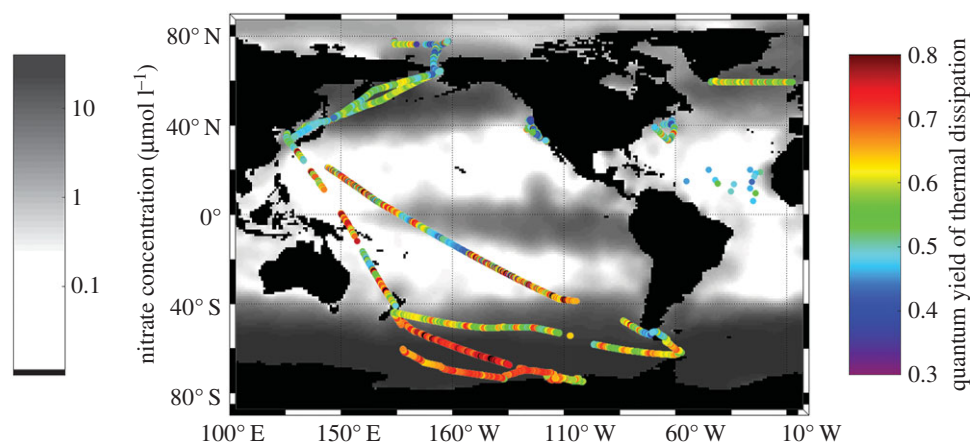


Figure 4. Global distribution of the quantum yields of thermal dissipation in photosynthetic reactions in the upper ocean superimposed on the climatological map of surface nitrate concentrations in the world ocean. These values were reconstructed from *in situ* variable fluorescence and lifetime measurements.

restrained by heterogeneity of the plant canopy, which makes it extremely difficult to deconvolute kinetic profiles generated by multiple layers of leaves within the canopy [38].

Variable fluorescence signals cannot be recorded from space without high-power lasers or some other source of light—which is not practical, let alone potentially dangerous. An alternative approach to infer phytoplankton physiology and photosynthetic rates is based on measurements of the absolute quantum yields of chlorophyll fluorescence. With the launch of the MODerate Resolution Imaging Spectroradiometer (MODIS) and MEdium Resolution Imaging Spectrometer (MERIS) satellites, which possess the capability of detecting solar induced chlorophyll fluorescence signals from the global ocean, it became theoretically possible to calculate the quantum yield of chlorophyll fluorescence from space [40–42]. The MODIS/MERIS analytical algorithms retrieve the quantum yields of chlorophyll fluorescence from the ratio of two independent variables, namely the magnitude of solar-induced fluorescence and the number of quanta absorbed by phytoplankton. Solar induced fluorescence (SIF, also called passive fluorescence) from chlorophyll a is detected as a red peak (centred at ca. 683 nm) in spectra of water-leaving radiance [43–45]. Although the presence of phytoplankton in natural waters alters the entire visible spectrum of water-leaving radiance [44–48], SIF is the only signal emitted from the ocean and detectable from space that can be unambiguously ascribed to phytoplankton.

The natural variations of fluorescence yields are the sources of both errors and useful information. SIF yield is highly variable in nature [49–54]. While the apparently huge variability of chlorophyll fluorescence yield in the ocean (ca. 10 fold) is often correlated with environmental forcing [12,41,50,54], the mechanisms and interpretation of this relationship remain to be elucidated.

The development of remote sensing algorithms for interpretation of solar-induced fluorescence yields crucially depends on comparison with accurate *in situ* measurements. Our sea-going lifetime fluorometer provides a unique operational tool for ground-truthing of satellite-based retrievals of the quantum yields of solar-induced chlorophyll fluorescence [12]. Our analysis of more than 20 000 matchups revealed a weak, but significant correlation between satellite-derived estimates of the quantum yields of chlorophyll fluorescence with corresponding *in situ* measurements [12].

6. Applications and limitations to terrestrial plants

Unlike marine unicellular organisms, which are optically thin, terrestrial plants concentrate their photosynthetic pigments in leaves, which are optically thick and spatially very heterogeneous. This leads to three fundamental problems in applying active fluorescence techniques in terrestrial ecosystems. First, the pigment packaging effect in leaves leads to reabsorption of fluorescent photons, thereby significantly reducing the observed fluorescence yields and red-shifting the emission spectrum of chlorophyll. The application of active fluorescence techniques to terrestrial plants potentially can resolve variable fluorescence parameters on leaf surfaces [55]. However, the packaging effect greatly complicates the interpretation of fluorescence signals, and therefore the conversion of lifetimes to absolute quantum yields [38]. Second, under natural irradiance conditions, the spatial heterogeneity in terrestrial canopies presents a significant problem in sampling fluorescence parameters under controlled irradiance. Hence, perhaps paradoxically, although chlorophyll concentrations in the upper ocean are 3 to 5 orders of magnitude smaller on an areal basis compared to vegetated terrestrial ecosystems, the biophysical signals obtained from active fluorescence techniques, while harder to measure, are easier to interpret within the context of current models of photosynthetic energy conversion efficiency.

7. Conclusion

The results of our extensive field measurements of variable fluorescence and lifetimes strongly imply that most of the time the phytoplankton in the world oceans operate at a much lower photosynthetic efficiency than they are potentially capable of achieving.

Indeed, it would appear that approximately 60% of the absorbed solar energy in the photosynthetically useable portion of the spectrum is dissipated as heat. Regardless, these organisms, which account for less than 1% of the global photosynthetic biomass, contribute about 45% of the global carbon fixation on Earth [34]. One of the fundamental limitations to an even higher contribution to carbon fixation is nutrients, especially fixed inorganic nitrogen and, in HNLC

regions, iron. Deliberate iron fertilization experiments in the latter regions clearly show that photosynthetic energy conversion efficiency can be rapidly increased to its maximum potential. In the former, mesoscale (e.g. tens to hundreds of kilometres) measurements reveal spatial variations in photosynthetic energy conversion efficiency associated with eddy turbulence, storms and wind driven processes that advect nutrients from deep waters into the euphotic zone. In effect, phytoplankton ‘body surf’, waiting for a wave of nutrients to allow the synthesis of key components of their photosynthetic apparatus, ultimately stimulating bursts of growth.

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