



# Characterization of MODIS-derived euphotic zone depth: Results for the China Sea

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## ARTICLE INFO

### Article history:

Received 4 June 2010

Received in revised form 4 August 2010

Accepted 24 August 2010

### Keywords:

MODIS

Remote sensing

China Sea

Euphotic zone depth

Chlorophyll-a

## ABSTRACT

Euphotic zone depth ( $Z_{eu}$ ) products from ocean color measurements are now produced from MODIS ocean color measurements, one of which is based on inherent optical properties (IOP-approach) and the other is based on chlorophyll-a concentration (Chl-approach). For the first time, the quality of these satellite  $Z_{eu}$  products is assessed with extensive field-measured  $Z_{eu}$  (in the China Sea), where 78% of the measurements were made on the continental shelf ( $\leq 200$  m). For the data with matching location and time window, we have found that the overall average difference ( $\epsilon$ ) between satellite and *in situ*  $Z_{eu}$  is 21.8% ( $n = 218$ ,  $Z_{eu}$  ranges from 4 to 93 m) with a root mean square error in log scale (RMSE) of 0.118 by the IOP-approach, while it is 49.9% (RMSE = 0.205) by the Chl-approach. These results suggest that 1) MODIS  $Z_{eu}$  products for waters in the China Sea are robust, even in shelf waters; and 2)  $Z_{eu}$  produced with IOPs are more reliable than those produced with empirically derived Chl. Spatial and seasonal variations of  $Z_{eu}$  in the China Sea are briefly described with  $Z_{eu}$  products generated by the IOP-approach. These results will facilitate further research on carbon cycling and environmental changes on both local and global scales.

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## 1. Introduction

Most marine phytoplankton photosynthesize in the euphotic zone (Kirk, 1994). Although the euphotic zone has recently attracted increasing attention (Buesseler et al., 2007), the euphotic zone is nevertheless the most important zone in the context of both ecosystem dynamics (Platt & Sathyendranath, 1988) and in air–sea interaction through transfer of either heat (Sathyendranath et al., 1991) or gases, especially with regards to green house gases such as carbon dioxide ( $\text{CO}_2$ ) (Takahashi et al., 2002). In practice, the euphotic zone depth ( $Z_{eu}$ ) is defined as the depth at which the PAR (photosynthetic available radiation) value is 1% of the surface value (Kirk, 1994). The value of  $Z_{eu}$  is an important input parameter for many models that estimate basin-scale primary production (Behrenfeld & Falkowski, 1997a,b). Both Secchi disk depth (Preisendorfer, 1986; Tyler, 1968) and  $Z_{eu}$  measure water clarity, but  $Z_{eu}$  can be measured objectively with advanced electro-optical instruments. Because  $Z_{eu}$  is a cumulative measure of biogeochemical properties in the upper-water column, changes in the  $Z_{eu}$  depict environmental patterns that might be associated with climate. Thus it is not surprising to see that  $Z_{eu}$  is included as a standard ocean color product in the Global Change Observation Mission (Japan, [http://suzaku.eorc.jaxa.jp/GCOM\\_C](http://suzaku.eorc.jaxa.jp/GCOM_C)) and that  $Z_{eu}$  products for the global

ocean are now derived from MODIS (Moderate Resolution Imaging Spectroradiometer) measured water-leaving radiance. Numerous studies have been carried out to evaluate the quality of satellite-derived chlorophyll concentration (Chl) and the diffuse attenuation coefficient (Bailey & Werdell, 2006; Darecki & Stramski, 2004; Melin et al., 2007; Zhang et al., 2006). While it is important to know the quality of satellite-derived products before they can be further applied to the study of biogeochemistry and environmental changes, there hasn't been any published independent research (to our knowledge) to evaluate or characterize the quality of MODIS-derived  $Z_{eu}$ .

In general, there are two approaches to estimate  $Z_{eu}$  from ocean color remote sensing (Lee et al., 2007; Morel et al., 2007). The simple Chl-approach is an empirical method based on Case-1 assumptions (Morel et al., 2007), where  $Z_{eu}$  is calculated from concentration of chlorophyll-a (Chl). A different approach (IOP-approach) is centered on waters' inherent optical properties (IOPs) (Lee et al., 2007);  $Z_{eu}$  is computed after the absorption and backscattering coefficients at 490 nm are provided. The inputs, either Chl or IOPs, can be obtained from water samples or from the inversion of measured water color (Gordon & Morel, 1983; IOCCG, 2006). Results from both approaches have been tested or validated with ship borne measurements (Lee et al., 2007; Morel et al., 2007), but there hasn't been any test or evaluation of these products when they are derived from satellite data.

In this study, using  $Z_{eu}$  data measured in the China Sea over the past six years, we assessed the  $Z_{eu}$  products (from both Chl-approach and IOP-approach) derived from MODIS measured ocean color. These measurements were mainly made in the shelf water ( $\leq 200$  m) to

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gauge the quality of satellite products over coastal waters. Finally, we discuss spatial and seasonal variations of  $Z_{eu}$  in the China Sea.

## 2. Data and methods

### 2.1. In situ $Z_{eu}$ and its uncertainty

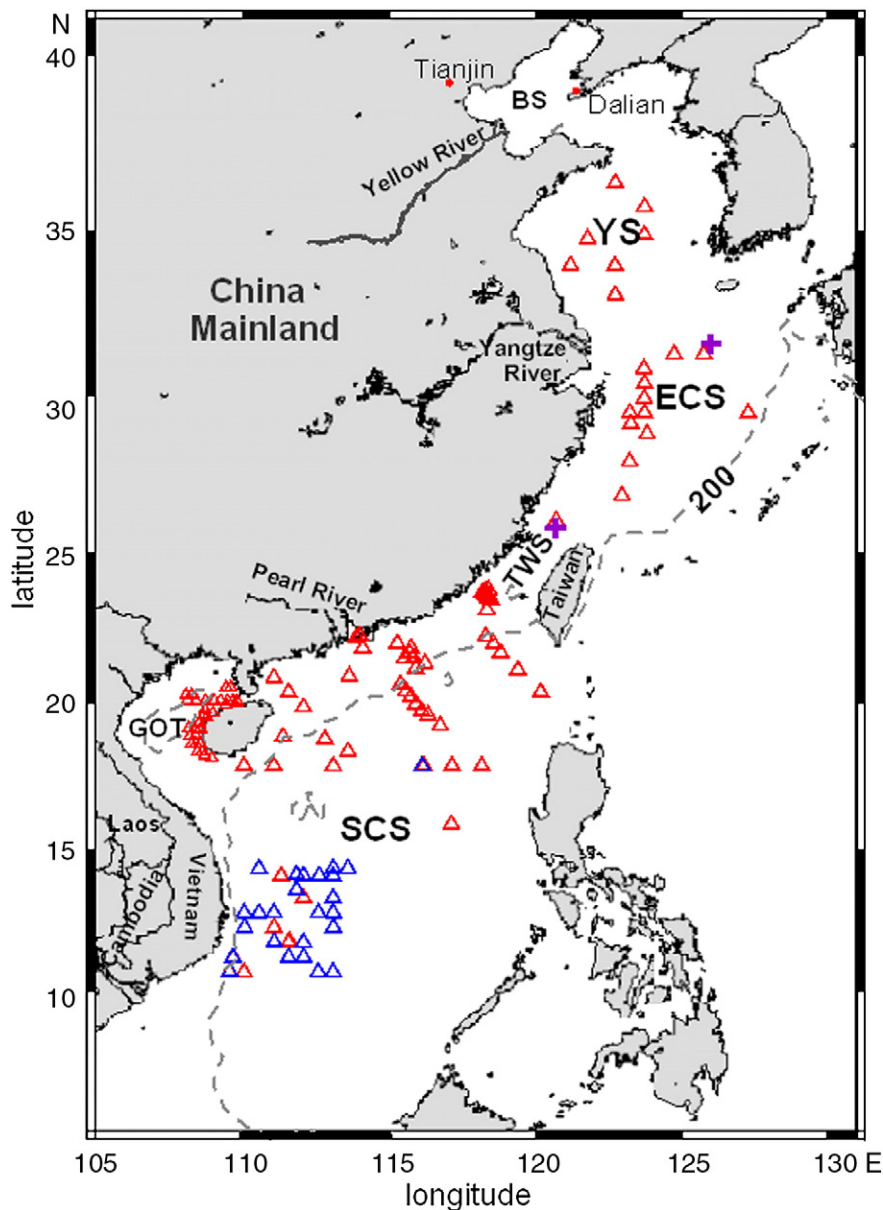
A series of field measurements of biological and optical properties were carried out between 2004 and 2009 in different seasons in the China Sea (Fig. 1). Based on equipment availability, the vertical profiles of PAR were measured with a hyperspectral irradiance meter or a broadband PAR sensor, and  $Z_{eu}$  values were calculated (detailed below) from the spectrally-integrated PAR profiles. Because  $Z_{eu}$  measures a relative difference between  $PAR(z)$  ( $z$  represents depth from sea surface) and  $PAR(0)$ , the absolute value that represents PAR has only minor impacts on the value of  $Z_{eu}$  (Morel & Gentili, 2004).

#### 2.1.1. $Z_{eu}$ from hyperspectral $E_d$

In 2007, hyperspectral (350–700 nm, ~3 nm spectral resolution) downwelling irradiance ( $E_d(\lambda)$ ) was measured with a Profiler II (Satlantic, Inc) in the South China Sea (blue symbols in Fig. 1). The Profiler II was lowered into water ~50 m away from an operational ship.  $E_d$  profiles were measured in the upper ~150 m and were processed with Satlantic software (Prosoft 7.7.10). Vertical profiles of the total downwelling irradiance in the visible domain  $E_{vis}(z)$  were calculated by summing  $E_d(\lambda, z)$  from 350 nm to 700 nm for each depth, and then the profiles of  $E_{vis}(z)/E_{vis}(0)$  were obtained.  $Z_{eu}$  was derived at the depth where  $E_{vis}(z)/E_{vis}(0) = 1\%$  (Lee et al., 2005, 2007).

#### 2.1.2. $Z_{eu}$ from PAR sensor

For the cruises during the period of 2004–2009, PAR (400–700 nm) was measured with a PAR sensor (PAR<sub>Bios</sub>, Biospherical



**Fig. 1.** Map of the study region and location of the field sampling stations. The blue symbols indicate those having both CTD and Profiler II measurements and the purple symbols indicate the two stations for analyzing small scale spatial-temporal variations in  $Z_{eu}$  (see text in 3.1); BS: Bohai Sea; YS: Yellow Sea; ECS: East China Sea; TWS: Taiwan Strait; SCS: South China Sea; and GOT: Gulf of Tonkin.

Instruments, Inc.) mounted on a CTD package; PAR profiles were processed with software provided by Biospherical Instruments Inc.

Because these CTD profiles were taken close to the operation ship, the PAR results at sea surface might have been contaminated by ship shadowing or reflected light from the hull. Consequently, the measured PAR(0) might be subject to larger uncertainty. To reduce this measurement-introduced uncertainty, a different approach was used for the derivation of  $Z_{eu}$  from the PAR profiles.

Based on Lee et al. (2005, 2007), the vertical variation of PAR(z) can be expressed as:

$$\frac{PAR(z)}{PAR(0)} = \exp(-K_{PAR}(z) \times z) \quad (1)$$

with

$$K_{PAR}(z) = K_1 + \frac{K_2}{(1+z)^{0.5}} \quad (2)$$

$K_{PAR}(z)$  is the vertical attenuation coefficient of PAR, and  $K_1$  and  $K_2$  are two model constants independent of depth (dependent on the sun angle and the inherent optical properties of the water). As  $Z_{eu}$  is the depth that satisfies  $K_{PAR}(Z_{eu}) \cdot Z_{eu} = 4.605$ , then the values of  $K_1$  and  $K_2$  are required for the derivation of  $Z_{eu}$ . Following this guidance, the vertical profile of each PAR measurement is expressed as:

$$\ln(PAR(z)) = \ln(PAR(0)) - \left( K_1 + \frac{K_2}{(1+z)^{0.5}} \right) \times z \quad (3)$$

and values of  $\ln(PAR(0))$ ,  $K_1$  and  $K_2$  for each station were derived by minimizing the difference between Eq. (3) – described profile and the PAR sensor measured profile. In the process, PAR(z) of depths generally deeper than 1 m were used. After  $K_1$  and  $K_2$  are known,  $Z_{eu}$  of a specific station can then be calculated algebraically (see Section 2.2).

### 2.1.3. Comparison between $Z_{eu}$ measured by PAR<sub>Bios</sub> and by Profiler

To obtain characterization and confidence of the  $Z_{eu}$  values derived from the two sensors and measurements,  $Z_{eu}$  derived from the PAR<sub>Bios</sub> (represented as  $Z_{eu}(PAR_{Bios})$ ) and the  $Z_{eu}$  derived from hyperspectral Profiler (represented as  $Z_{eu}(Profiler)$ ) for co-located stations (32 stations, see locations in Fig. 1, the blue symbols) were compared to each other and are presented in Fig. 2. For  $Z_{eu}$  in a range of 35–90 m, the average percentage difference between the two sets of  $Z_{eu}$  is 7.9%, with a correlation coefficient (R) of 0.942. Such results indicate self-consistency in  $Z_{eu}$  from two independent determinations in the field measurements, and provide a first order estimation of uncertainty of *in situ*  $Z_{eu}$ . In order to maintain the same standard for assessing MODIS  $Z_{eu}$ , and because  $Z_{eu}$  values from the Profiler are subject to less

uncertainty, for all stations where only  $Z_{eu}$  from PAR<sub>Bios</sub> were available their  $Z_{eu}(PAR_{Bios})$  were converted to equivalent  $Z_{eu}(Profiler)$  with:

$$Z_{eu}(Profiler) \approx 1.075 Z_{eu}(PAR_{Bios}). \quad (4)$$

Therefore, the final *in situ*  $Z_{eu}$  data used to assess MODIS  $Z_{eu}$  are  $Z_{eu}$  from the Profiler (where Profiler data were available) and  $Z_{eu}$  derived from Eq. (4) (where only PAR<sub>Bios</sub> data were available). For simplification, hereafter the  $Z_{eu}$  values derived from field measurements are annotated as  $Z_{eu}^{Mea}$ ; values from the Chl-approach are  $Z_{eu}^{Chl}$ ; and from the IOP-approach are  $Z_{eu}^{IOP}$ .

## 2.2. $Z_{eu}$ derived from satellite measurements

MODIS daily Level-2 normalized water leaving radiance (nLw) (MODIS/Aqua Reprocessing 1.1) were obtained from the NASA Distributed Active Archive Center and subsequently converted to remote-sensing reflectance (Rrs) via the ratio of nLw to extra-terrestrial solar irradiance (F0) (Gordon, 2005; also see <http://oceancolor.gsfc.nasa.gov/forum/oceancolor/>). The resolution of these data is approximately 1 km x 1 km at the nadir pixel.

### 2.2.1. $Z_{eu}$ from empirically derived Chl

This satellite Rrs dataset was used in the OC3M band ratio algorithm (O'Reilly et al., 2000) to calculate Chl.  $Z_{eu}$  was subsequently derived by the approach of Morel et al. (2007):

$$\log(Z_{eu}) = 1.524 - 0.436X - 0.0145X^2 + 0.0186X^3 \quad (5)$$

with  $X = \log(Chl)$ .

### 2.2.2. $Z_{eu}$ from semi-analytically derived IOPs

The absorption and backscattering coefficients at 490 nm ( $a(490)$  and  $b_b(490)$ ), were derived from the Rrs by applying the Quasi-Analytical Algorithm (Lee et al., 2002, 2007) (also see [http://www.iocccg.org/groups/Software\\_OCA/QAA\\_v5.pdf](http://www.iocccg.org/groups/Software_OCA/QAA_v5.pdf) for recent updates).  $K_{PAR}(z)$  was then calculated (Lee et al., 2005) as a function of  $a(490)$  and  $b_b(490)$ , and the sun angle ( $\theta_a$ ) matching relevant *in situ* measurements. The IOP-centered  $Z_{eu}$  (Lee et al., 2007) was thus derived based on the following equation:

$$K_{PAR}(Z_{eu}) \cdot Z_{eu} = 4.605 \quad (6)$$

The root mean square error in log scale (RMSE) and an averaged percentage error ( $\varepsilon$ ) were then used as measures to evaluate the consistency between the *in situ* and satellite data sets:

$$\varepsilon = \left( \frac{1}{n} \sum_{i=1}^n \left| \frac{(Z_{eu}^{IOP})_i - (Z_{eu}^{Mea})_i}{(Z_{eu}^{Mea})_i} \right| \right) \times 100\% \quad (7)$$

$$RMSE = \sqrt{\frac{1}{n} \sum_{i=1}^n (\log((Z_{eu}^{IOP})_i) - \log((Z_{eu}^{Mea})_i))^2} \quad (8)$$

RMSE and  $\varepsilon$  were calculated for  $Z_{eu}^{Chl}$  by substituting  $Z_{eu}^{IOP}$  with  $Z_{eu}^{Chl}$  in Eqs. (7) and (8).

## 3. Comparison between MODIS $Z_{eu}$ and *in situ* $Z_{eu}$

### 3.1. Matching between satellite and *in situ* measurements

Due to cloud coverage and cost, *in situ* measurements are extremely difficult to obtain with simultaneous measurements between ship surveys and satellite sensors. In order to gain large numbers of matching data points, Bailey et al. (2002) suggest a relaxation in the temporal difference to within  $\pm 2-3$  h of the satellite overpass. However, temporal mismatch is just one of the factors

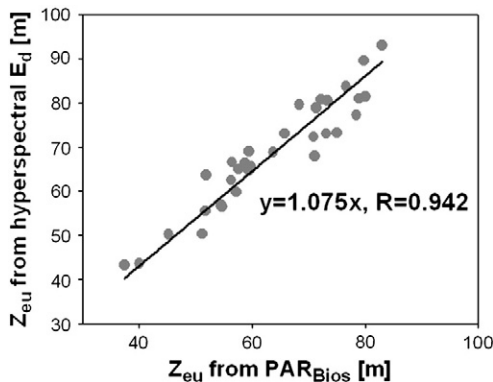


Fig. 2.  $Z_{eu}$  derived from PAR<sub>Bios</sub> versus  $Z_{eu}$  derived from hyperspectral  $E_d$ .

affecting the quality of matching between satellite and *in situ* measurements. Another important factor is the spatial mismatch. A satellite sensor (e.g., MODIS) provides a measurement of an averaged property within  $\sim 1$  km by 1 km, while *in situ* measurements represent an average area of 1 m by 1 m. Consequently, even if the two measurements are carried out simultaneously, data from the two measurements do not “match” each other, unless the observation area is completely homogeneous (i.e., independent of sampling location). This is a more serious source of uncertainty for coastal or near-shore waters where there are greater spatial variations. Because of such inherent mismatches, resulting from measurement strategies, matching-up data between satellite and *in situ* measurements are not always ideal.

In order to obtain an appropriate number of matching observations for meaningful statistical analysis, we relaxed the time-window of overpass to  $\pm 48$  h. This is based on an evaluation of temporal variations of the research regions and the spatial changes in the 1 km spatial resolution as detailed below.

Short-term variations of  $Z_{eu}$  within 13 days were calculated from MODIS Rrs by using the IOP-approach at two field sampling stations (the two purple symbols in Fig. 1) in summer 2004 and 2009 – during cruise surveys (Fig. 3a–b). The 2004 location was at 26.023°N, 120.486°E in the Taiwan Strait (TWS), and in 2009 it was at 32.250°N, 125.750°E in the East China Sea (ECS). For the Taiwan Strait, the value of  $Z_{eu}$  within 13 days was  $53 \pm 4$  m. The mean percentage variation of  $Z_{eu}$  within  $\pm 2$  days (range/mean) was  $\sim 13\%$ . For the East China Sea, these numbers were  $38 \pm 3$  m and  $\sim 15\%$ , respectively. These temporal variations are comparable to the uncertainty of *in situ* measured  $Z_{eu}$ . In addition, Fig. 3c–d presents spatial variations of  $Z_{eu}$  around the two stations (extending 1 km to 8 km away from the stations, based on satellite data availability) on

August 7, 2004 and August 24, 2009, respectively. The maximum spatial variations of these locations within 8 km distance were  $\sim 18\%$  and  $33\%$ , respectively. These analyses indicate that no significant extra uncertainties would be introduced from the relaxation of time windows for characterizing MODIS  $Z_{eu}$  products.

Following the above data processing method, along with the temporal window for MODIS, a total of 218 match-ups were compiled between *in situ* and satellite  $Z_{eu}$  data, which covers a range of 4–93 m. The following characterizes and discusses how satellite  $Z_{eu}$  compared with  $Z_{eu}$  determined from *in situ* measurements.

### 3.2. MODIS $Z_{eu}$ compared with *in situ* $Z_{eu}$

The comparisons between MODIS  $Z_{eu}$  and *in situ*  $Z_{eu}$  are presented in Fig. 4. For this dataset, while the non-shelf waters have a  $Z_{eu}$  range of 35–93 m, the shelf waters (bathymetry  $\leq 200$  m) show quite a wide range of  $Z_{eu}$ : 4–75 m. Such an observation indicates that shelf waters are dynamic and complex in biogeochemical contents, and signify different status of ecosystem health.

From satellite estimates, MODIS  $Z_{eu}^{chl}$  shows a high correlation with *in situ*  $Z_{eu}$  ( $R = 0.95$ ), along with a relative difference of 49.9% and an RMSE of 0.205 (Table 1). The bigger relative difference and RMSE values are because MODIS  $Z_{eu}^{chl}$  is apparently (and systematically) higher than *in situ*  $Z_{eu}$ ; this is also described in Lee et al. (2007) when comparing  $Z_{eu}$  with  $Z_{eu}^{chl}$  derived from ship-borne Rrs. In addition to the uncertainties introduced by the mismatches of spatial and temporal scales between *in situ* and satellite measurements, other sources of uncertainty for MODIS  $Z_{eu}^{chl}$  include the empirical estimation of surface Chl and the empirical relationship between Chl and  $Z_{eu}$ . Both relationships (Morel et al., 2007; O'Reilly et al., 2000) were developed with data that excluded the China Sea. Fig. 5

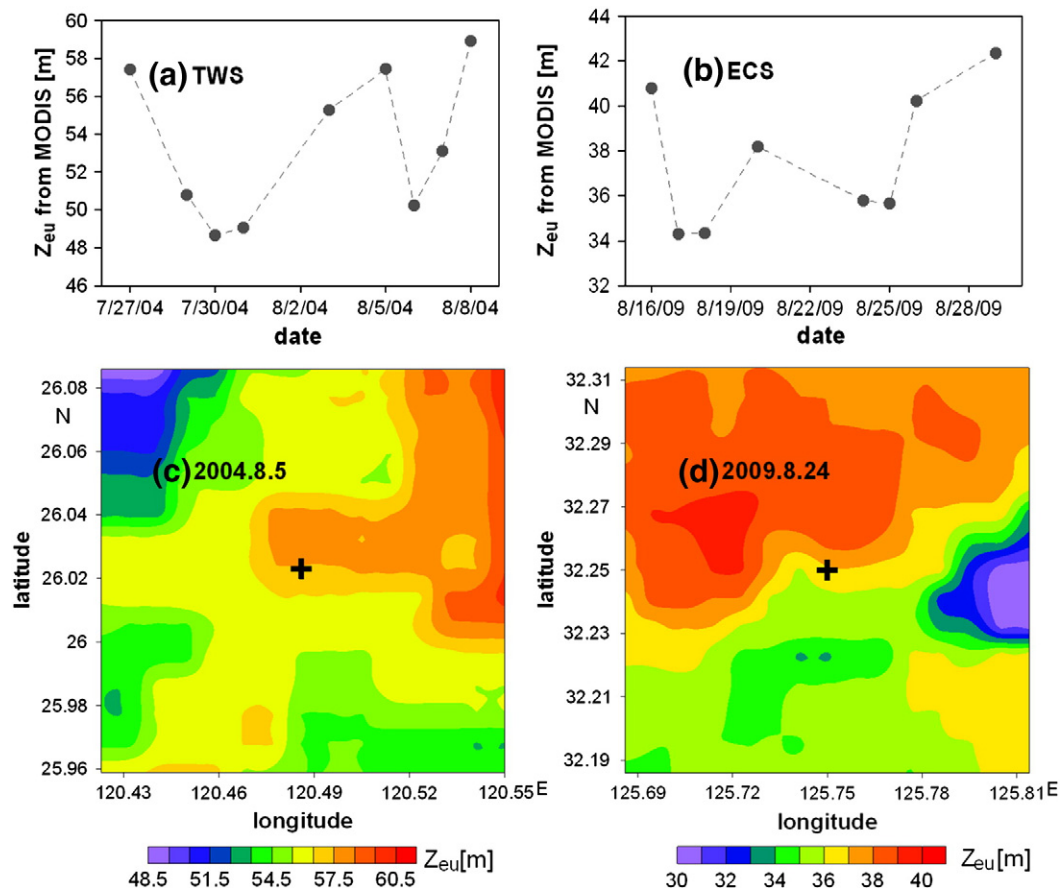


Fig. 3. Temporal and spatial variations of MODIS  $Z_{eu}$  at two stations. (a) and (c): TWS (26.023°N, 120.486°E) in July–August 2004; (b) and (d): ECS (32.250°N, 125.750°E) in August 2009.



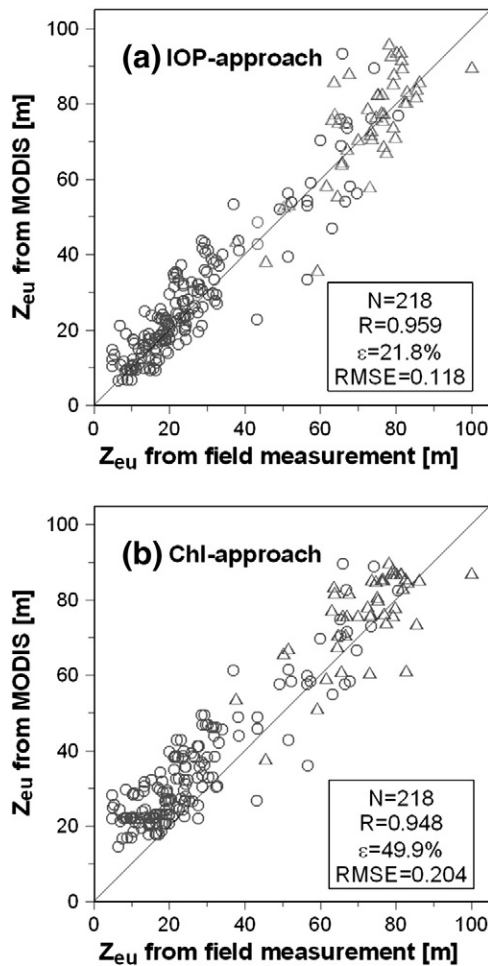


Fig. 4. Comparison between satellite  $Z_{eu}$  (by the IOP-approach (a) and Chl-approach (b)) and the  $Z_{eu}$  derived from *in situ* measurements. Circles are samples on the shelf (depth  $\leq 200$  m) and triangles are those in the basin (depth  $> 200$  m); the same applied to the subsequent two figures (Figs. 5–6).

compares Chl estimated by MODIS with Chl from the water samples (Huang et al., 2008; Zhai et al., 2009) (measured fluorometrically (Lalli & Parsons, 1993), also with relaxed time-window of satellite overpass to  $\pm 48$  h). Although the difference is large (RMSE = 0.363), there is no detectable systematic under- or over-estimation of Chl from

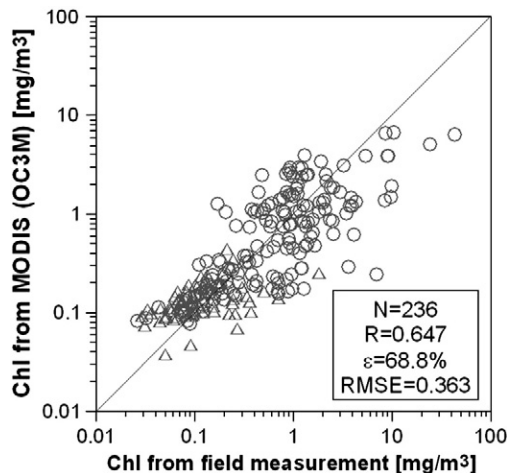


Fig. 5. Comparison between Chl derived from MODIS and Chl from surface water samples.

MODIS for this dataset. The over-estimation of MODIS  $Z_{eu}^{chl}$  apparently resulted from the empirical relationship between  $Z_{eu}$  and Chl, as indicated in Fig. 6 where *in situ*  $Z_{eu}$  is compared with *in situ* Chl. For the China Sea, at least for this dataset,  $Z_{eu}$  would be overestimated if the global empirical model (Morel et al., 2007) was applied (also see comparison of empirical  $Z_{eu}$ -Chl relationship in Siegel et al. (2001)). Because the waters studied here are subject to high runoff from major river systems, these results highlight that  $Z_{eu}$  of waters with a high suspended sediment load could not be explained well by Chl alone.

For the same dataset ( $Z_{eu}$  in a range of 4–93 m from *in situ* measurements), MODIS  $Z_{eu}^{IOP}$  also shows a high correlation with *in situ*  $Z_{eu}$  ( $R = 0.96$ ). Compared to  $Z_{eu}^{chl}$ , however, the average relative difference ( $\varepsilon$ ) between MODIS  $Z_{eu}^{IOP}$  and *in situ*  $Z_{eu}$  is 21.8% (RMSE is 0.118 (Table 1)). Because of extra sources of uncertainty (such as uncertainty in atmosphere correction and/or a mismatch of spatial and temporal scales), the value of  $\varepsilon$  (and RMSE) between *in situ*  $Z_{eu}$  and MODIS  $Z_{eu}^{IOP}$  is higher than that obtained between *in situ*  $Z_{eu}$  and *in situ*  $Z_{eu}^{IOP}$  (Lee et al., 2007). Nevertheless, the results indicate a significantly improved estimation of  $Z_{eu}$  from MODIS with the IOP-approach, and consequently improvements of estimation of primary production where the  $Z_{eu}$  product is required (Behrenfeld & Falkowski, 1997a; Shang et al., 2010). This result is even more exciting because most (78%) of the dataset were measured on the shelf, and the composition of shelf water constituents are generally complex and dynamic. Such an improvement on the satellite  $Z_{eu}$  estimation may be attributed to the fact that there is no dependence on Case-1 assumptions when applying the IOP-approach. Meanwhile, the discrepancy between MODIS  $Z_{eu}$  and *in situ*  $Z_{eu}$  shown above suggests an upper limit of  $Z_{eu}$  uncertainty from MODIS since there are extra sources of uncertainty in satellite remote sensing, while the results in Lee et al. (2007) indicated a lower limit of uncertainty, when the IOP-approach is applied.

Note that the RMSE value is 0.363 and the average relative difference is  $\sim 69\%$  when *in situ* Chl was compared with the match-up satellite Chl (see Fig. 5). Our results indicate that, at least for waters in the China Sea, MODIS-derived  $Z_{eu}^{chl}$  (Fig. 4, right) has better accuracy than MODIS-derived Chl. This is in part because euphotic zone depth is an optical property, while chlorophyll-a concentration is a biological property. When they are derived from a remote sensing reflectance, the former has no dependence on the concentration-normalized optical properties (e.g., chlorophyll-specific absorption coefficient), while the latter does (explicitly or implicitly). MODIS-derived  $Z_{eu}^{IOP}$ , however, has much higher accuracy (a factor of 3 smaller RMSE and relative difference) than MODIS-derived Chl.

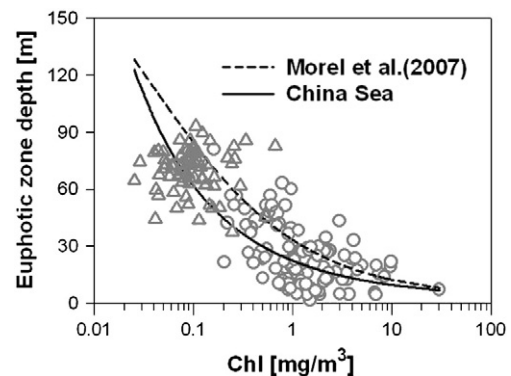


Fig. 6. Relationship between *in situ*  $Z_{eu}$  and *in situ* Chl, with all data from the China Sea. The empirical, global, relationship developed by Morel et al. (2007) is superimposed on the figure (dashed line). The local empirical relationship (solid line) for this dataset is:  $\log_{10}(Z_{eu}) = 1.35 - 0.4026 X + 0.0375 X^2$ ,  $X = \log_{10}(\text{Chl})$ ,  $R = 0.767$ ,  $N = 190$ .

**Table 1**

Error statistics for the comparison between the  $Z_{eu}$  derived from *in situ* measurements and from MODIS Rrs.

	IOP-approach		Chl-approach		N
	$\epsilon(\%)$	RMSE	$\epsilon(\%)$	RMSE	
TWS	18.5	0.091	37.3	0.157	56
GOT	37.6	0.171	107.9	0.331	43
SCS	15.9	0.103	34.5	0.165	80
YS and ECS	21.3	0.107	35.7	0.148	39
Shelf water	25.1	0.130	60.6	0.229	170
Basin water	10.2	0.060	12.2	0.062	48
Total	21.8	0.118	49.9	0.205	218

Especially, the IOP-approach is *not* a regression analysis to develop empirical relationships (and coefficients), but rather an *independent* test/evaluation of algorithms that were developed earlier. Such efforts provide users more insights of performances and characterizations of independently developed algorithms in a broad range of environments. Furthermore, a product with low RMSE adds confidence in applying MODIS products for monitoring water clarity and/or ecosystem health.

#### 4. Distribution and variation of MODIS $Z_{eu}$ in the China Sea

We then used the IOP-approach to derive a climatological seasonal mean  $Z_{eu}$  for the China Sea (see Fig. 7) from the monthly mean MODIS Rrs for the time frame of December 2002 through November 2009. The results presented hereafter are limited to the region west of the dashed black line in Fig. 7.

In general,  $Z_{eu}$  is deep (maximum ~110 m) in the oligotrophic South China Sea (SCS, ~5000 m at maximum depth (Liu et al., 2002)) when compared to  $Z_{eu}$  in the complex and relatively eutrophic East China Sea (ECS) (Gong et al., 2003). This is because the ECS has a wide continental shelf and is associated with freshwater discharge from the Yangtze River (the third largest river in the world). There is a distinct tongue of low  $Z_{eu}$  water out of the Yangtze River estuary. The value of  $Z_{eu}$  within this tongue is extremely low in winter (~<10 m overlaid on the ECS background level of ~40 m). The tongue of low  $Z_{eu}$  reaches further offshore in winter, a dry season when the river discharge is low (minimum ~30 km<sup>3</sup>/month, mean value for 1950–1990, (Xu & Milliman, 2009)), while it shrinks in summer. This suggests that sediment resuspension (driven by strong winds in winter) (Song et al., 2006) has a higher impact on the shallowing of  $Z_{eu}$  in this region – when compared to the river outflow (Xu & Milliman, 2009).

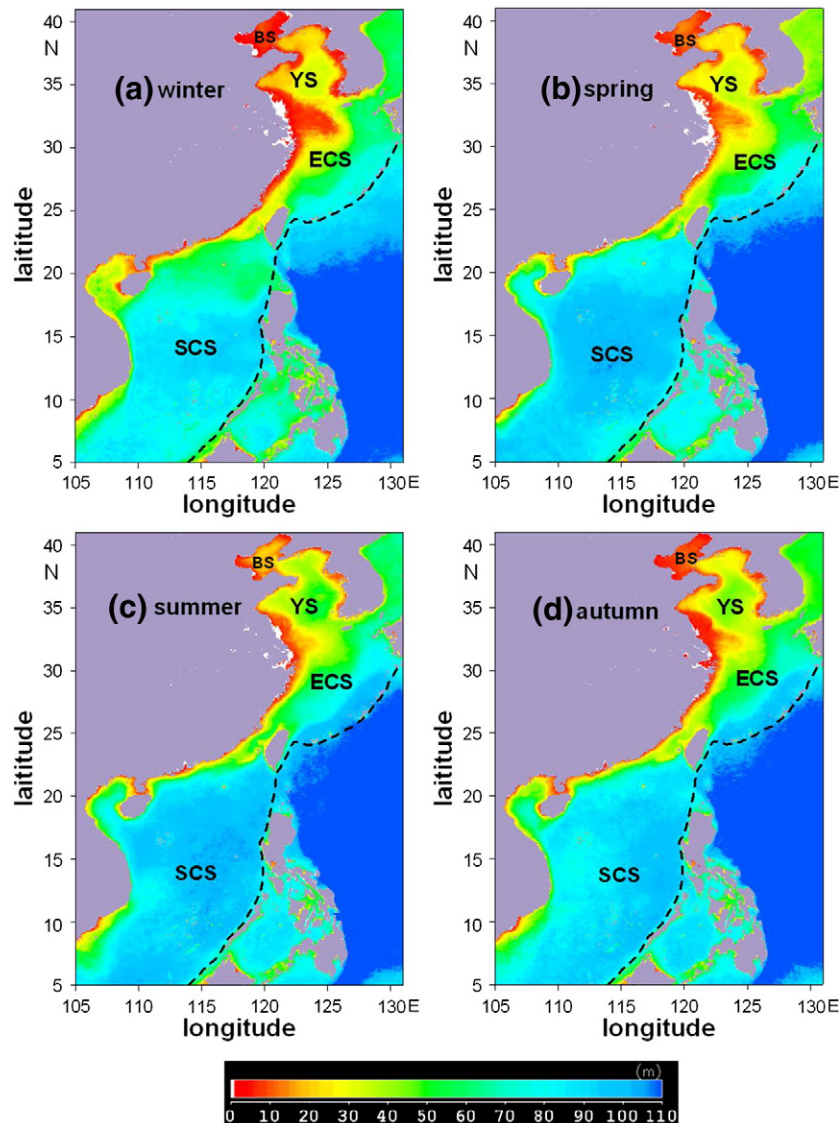


Fig. 7. Climatological seasonal mean  $Z_{eu}$  in the China Sea (IOP-approach, 2002–2009).

In the SCS, winter is the season with the highest primary production in the surface layer, which is attributed to nutrient pumping from deep water, which is in turn induced by strong wind mixing (Chen, 2005). Consequently,  $Z_{eu}$  is shallower in winter (~85 m) when compared to other seasons (~110 m) for the SCS. On the other hand, because of the intrusion of the oligotrophic Kuroshio and Taiwan Warm Current to the ECS (driven by southerly winds) (Lee & Chao, 2003),  $Z_{eu}$  is generally deeper in summer in the ECS, although primary productivity in some areas is three times higher in summer than that in other seasons (Gong et al., 2003).

In addition, the Bohai Sea is an area of notably shallow  $Z_{eu}$  (~6 m in winter, ~9 m in spring, ~8 m in autumn, and ~14 m in summer). Considering that the Bohai Sea is a semi-enclosed shallow sea (average water depth is ~18 m) surrounded by several big cities (e.g., Tianjin and Dalian), and that the Yellow River (the sixth longest river in the world) empties into this marine system, this character is quite reasonable and expected.

## 5. Conclusions

For the first time, the quality of the two MODIS  $Z_{eu}$  products is characterized with extensive field-measured  $Z_{eu}$  in the China Sea, a marginal sea that features one of the widest continental shelves in the world and has an input from the two large rivers. Three main results are found from this study. First, MODIS  $Z_{eu}$  products in the China Sea are robust, even in shelf waters ( $\leq 200$  m). Second,  $Z_{eu}$  produced with the IOP-approach (which has no dependence on Case-1 assumptions) is more reliable than those produced with empirically derived Chl (which is dependent on Case-1 assumptions). And third,  $Z_{eu}$  is generally shallower in winter in the China Sea (when compared to summer), and is shallower in the East China Sea than in the South China Sea.

Results shown here provide the research and monitoring communities an increased confidence when applying the satellite  $Z_{eu}$  products (especially those produced with the IOP-approach) to the primary production estimates and environmental change research – both at the local and global scales.

## Acknowledgements

This work was supported jointly by the High-Tech R&D Program of China (#2006AA09A302 & 2008AA09Z108), the National Basic Research Program of China (#2009CB421200 & 2009CB421201), NSF of China (#40821063) and the China Scholarship Council. Supports provided by NASA's Water- and Energy Cycle Program (Lee) are greatly appreciated. We thank Drs. H. Hong, M. Dai, H. Gao, Y. Li, J. Hu, B. Huang, J. Sun, W. Zhai, Z. Liu and L. Huang for providing *in situ* measured PAR and Chl data. We thank the Ocean Biology Processing Group (OBPG) at the Goddard Space Flight Center for generation and distribution of the MODIS products.

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