

Journal of Applied Remote Sensing

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Caroline Andrade, Nariane Bernardo, Alisson Carmo, Enner Alcântara, Milton Kampel, "Assessment of quasi-analytical algorithm for estimating the inherent optical properties in a complex cascade system," *J. Appl. Remote Sens.* **12**(3), 036014 (2018), doi: 10.1117/1.JRS.12.036014.

Assessment of quasi-analytical algorithm for estimating the inherent optical properties in a complex cascade system

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Abstract. Several quasi-analytical algorithm (QAA) versions were developed to make it suitable for different inland water systems. QAA_{BBHR} and QAA_{OMW} were reparameterized based on two reservoirs from the Tietê River cascading system (São Paulo State, Brazil), which present widely differing compositions. Considering the purpose of monitoring the entire cascade through a unique QAA version, we aimed to assess the suitability of these two QAA versions and, in addition, another two QAA native forms (versions 5 and 6), for retrieving inherent optical properties (IOPs) in Ibatinga hydroelectric reservoir (IHR), situated in the same cascading system. In addition to that, we addressed bio-optical characterization of IHR, using spectral and water quality data collected in a field campaign conducted in July 2016. Wide spatial variability of optically significant constituent (OSC) in IHR and colored dissolved organic matter predominance in its absorption budget was observed. None of the tested QAA versions were completely suitable in retrieving absorption coefficients for IHR in all wavelengths. However, results for wavelengths commonly used as proxy for OSC concentration retrieval were satisfactory in some of the models. Therefore, the results obtained in this study shows that QAAs versions can be used for specific purposes (e.g., chlorophyll-*a* mapping), by employing the best model for IOPs retrieval at a specific wavelength. This highlights the challenge of copying with high optical variability in cascading systems. In this sense, further research is necessary, for either achieving a QAA reparameterized version appropriate for aquatic systems with widely differing optical properties or another analytical scheme. © 2018 Society of Photo-Optical Instrumentation Engineers (SPIE) [DOI: [10.1117/1.JRS.12.036014](https://doi.org/10.1117/1.JRS.12.036014)]

Keywords: bio-optical models; water quality; water resources monitoring; remotely sensed data.

Paper 180369L received May 3, 2018; accepted for publication Aug. 17, 2018; published online Sep. 4, 2018.

1 Introduction

Remotely sensed data have been used to retrieve the optically significant compounds (OSC) concentrations in inland waters, using bio-optical models. These models are often empirical or semianalytical models. Empirical models directly relate remotely sensed measurements to the OSC of interest, usually through statistical regression, whereas semianalytical is based on radiative transfer inverse modeling. Empirical models can be time and site limited with no physical meaning.

Semianalytical and quasianalytical algorithms (QAA)¹ are based on radiative transfer theory and often include empirical steps. A semianalytical model estimates the total absorption coefficient $a_t(\lambda)$ by sum of phytoplankton absorption coefficient, $a_p(\lambda)$, colored dissolved organic matter (CDOM) absorption coefficient, $a_{CDOM}(\lambda)$, and nonalgal particle absorption coefficient, $a_{NAP}(\lambda)$ ²⁻⁴ while QAA estimates $a_t(\lambda)$ using exclusively remote sensing reflectance (R_{rs}).

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The backscattering coefficient, $b_b(\lambda)$, is also estimated via semianalytical models by summing b_b of each in-water constituent, except CDOM, whereas QAA also retrieves backscattering coefficient of suspended particles $b_{bp}(\lambda)$ using R_{rs} , though analytical and empirical steps.

QAA is a special case because it includes empirical and analytical steps for retrieving inherent optical properties (IOPs) and was developed to retrieve IOPs in the open ocean and coastal waters, and later for turbid waters. There are mainly three limitations of QAA applicability in inland waters: (1) the model used to estimate the $a_t(\lambda)$ is empirically derived from synthetic data, (2) the model used to estimate η —spectral power for $b_{bp}(\lambda)$ derivation, and (3) the equations used to estimate the phytoplankton absorption coefficient.^{5,6}

The applicability of QAA in optically complex inland waters is certainly more challenging than when working on case 1 waters.² To make QAA suitable for more complex waters, calibration and reparameterization of the original versions are required as demonstrated by several authors.^{5–8} Considering that at shorter wavelengths, a_t is highly influenced by a_ϕ and CDOM plus detritus absorption coefficient, a_{CDM} , in inland waters, reparameterization includes reference wavelength (λ_0) shifting to red-edge and near-infrared regions.^{6–10} In addition to that, other usual enhancements in inland water QAA versions are a calibration of intermediary empirical and semianalytical steps, such as $a_t(\lambda_0)$ and $b_{bp}(\lambda)$ estimation, and reparameterization of equations or coefficients for a_ϕ and a_{CDM} estimation, according to the IOPs presented in the specific inland water body.

When it comes to cascading reservoirs systems, variability trends are also influenced by cascading effects,¹¹ as well as by the different contributions arriving from the drainage basin, tributary rivers, and floodgates mechanisms.¹² In such aquatic systems, there are widely differing optical properties and the efficiency of parameterized versions is uncertain.

Tietê River cascading system is located in São Paulo state, a densely populated region of Brazil where water quality is an urgent matter. Recently, two QAA versions were parameterized for two reservoirs from this system: QAA_{BBHR}¹³ and QAA_{OMW}.¹⁴ The QAA_{BBHR} was parameterized for the eutrophic Barra Bonita hydroelectric reservoir (BBHR),¹⁵ which is the first reservoir of the cascading system, receiving large amounts of pollutants from the metropolis of São Paulo and also from agriculture and cattle raising.¹¹ Suspended particulate matter (SPM) in this reservoir is dominated by the organic fraction.^{12,13} The QAA_{OMW} version was parameterized for the oligo-to-mesotrophic Nova Avanhandava reservoir (NHR), located further downstream, presenting relatively low chlorophyll-*a* (Chl-*a*) and SPM concentrations, and inorganic predominance in its SPM.¹²

Ibitinga hydroelectric reservoir (IHR) is also situated in the Tietê River, downstream from BBHR and upstream from NHR. Chl-*a* and SPM concentration values in IHR are typically between the ranges found in BBHR and NHR.¹⁶ However, due to some particularities, as organic matter predominance and slightly higher Chl-*a* concentrations, IHR is possibly more similar to BBHR bio-optical characteristics. Considering that IHR bio-optical status is comparable with BBHR ones, the hypothesis here tested is that QAA_{BBHR} could be able to consistently retrieve a_ϕ in IHR but may be not capable of accurately retrieve a_{CDM} , as QAA_{BBHR} was parameterized considering phytoplankton features of the BBHR reservoir.¹³

To test the aforementioned hypothesis, we aim to assess the performance of QAA_{BBHR} and QAA_{OMW}, in addition two latest native forms of QAA – QAA_{V5}¹⁷ and QAA_{V6}¹⁸—in retrieving a_ϕ and a_{CDM} for IHR; we also will retrieve a_t for each QAA version. We decided to also test the QAA_{V5} and QAA_{V6} to compare the results from the original and reparameterized versions. This investigation relates to the purpose of monitoring all the cascading system in an integrated manner, through a unique QAA version properly adjusted for capturing the bio-optical variability occurring along the system. We also addressed water quality and bio-optical characterization of IHR, intending to associate it with algorithms efficiency, as this was not previously reported in the literature.

2 Materials and Methods

2.1 Study Area

IHR is located in the middle course of Tietê River, central area of São Paulo State, Brazil (21°45' S and 48°59' W). It is the third of six cascading reservoirs [Fig. 1(a)]. Its flooded area is around

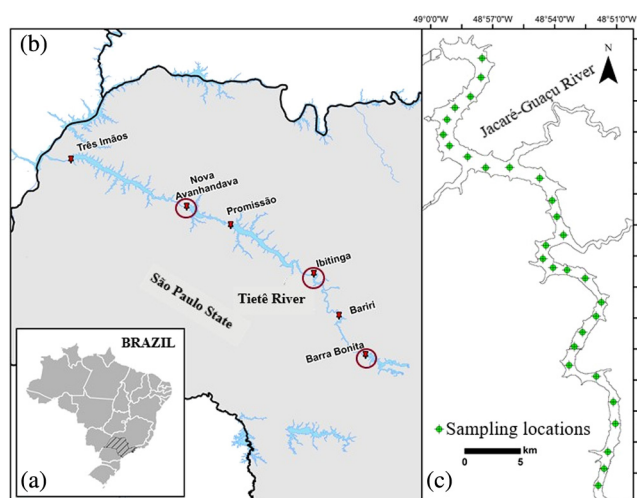


Fig. 1 (a) São Paulo State location in Brazil, (b) cascading reservoirs in Tietê River. Please note the IHR upstream from NHR and downstream from Barra Bonita reservoir (see text for more details), and (c) distribution of 29 sampling stations in IHR.

114 km², with ~9 m of average depth and an average flow of 525 m³ s⁻¹. IHR is situated in a temperate humid climate, characterized by well-defined dry and wet seasons. The surrounding area presents grazing land, sugarcane crops, and minor reforestation areas.¹⁹ The river receives wastewater discharges from domestic and industrial sources along its course, besides diffuse sources of pollution arising from agriculture and cattle breeding activities. The trophic state of the reservoir is highly spatially and temporally variable, and some areas are classified as eutrophic, whereas other ones are considered mesotrophic.^{16,20} A field campaign was conducted during austral winter, from July 19 to 23, 2016, in which water quality and spectral data were obtained in 29 sampling stations [Fig. 1(c)], as well as water samples collected just below air–water interface.

2.2 Water Quality Data

Secchi disk depth (m), turbidity (NTU), and electric conductivity ($\mu\text{S}/\text{cm}$) data were obtained in all 29 sampling stations [Fig. 1(c)]. Water samples were collected to estimate Chl-*a* concentration, through acetone extraction method.^{21,22} The SPM concentrations, as well as organic (OSM) and inorganic suspended matter (ISM) fractions were estimated according to the American Public Health Association protocol.²³

2.3 Radiometric Data

Total upwelling radiance [$L_t(\lambda)$; W m⁻² sr⁻¹ nm⁻¹] and atmospheric diffuse radiance [$L_s(\lambda)$; W m⁻² sr⁻¹ nm⁻¹] were measured using two RAMSES-ARC hyperspectral radiometers. Downwelling irradiance [$E_d(\lambda)$; W m⁻² nm⁻¹] was measured using a RAMSES-ACC sensor (TriOS, Oldenburg, Germany). All radiometric sensors used were operated in the spectral range between 350 and 900 nm, with a spectral resolution of 3.3 nm. The acquisition geometry followed Ref. 24, with $L_t(\lambda)$ measured at a zenith angle (θ) of 140 deg, $L_s(\lambda)$ at $\theta = 40$ deg, and $E_d(\lambda)$ measured with the sensor aligned to the zenith, i.e., $\theta = 0$ deg; all measurements were taken at an azimuth angle of 90 deg.

These radiometric quantities were then applied to estimate R_{rs} (sr⁻¹) spectra, using the spectral optimization approach proposed by Ref. 25, to remove surface-reflected radiance [$L_r(\lambda)$].²⁶ R_{rs} spectra, which were estimated through *in situ* hyperspectral measurements, were then resampled to simulate satellite data. The spectral response functions of each ocean and land colour instrument—Sentinel 3 band were used to derive band-weighted data.

2.4 Phytoplankton Absorption Coefficient (a_p), Detritus Plus CDOM Absorption Coefficient (a_{CDM}), and Total Absorption Coefficient (a_t)

To estimate CDOM absorption coefficient (a_{CDM} , m^{-1}), water samples were filtered through Whatman nylon membrane—0.22- μm porosity and 47-mm diameter. The absorbance of the filtrates was read using a 2600 UV-VIS spectrophotometer (Shimadzu, Japan), and the results were applied to calculate a_{CDM} as proposed by Ref. 27 [Eq. (1)]:

$$a_{CDM} = 2.3 \frac{OD_{CDM}(\lambda)}{r}, \quad (1)$$

where $OD_{CDM}(\lambda)$ is the optical density of CDOM and r is the cuvette path length (0 and 1 m).

To determine the total particulate (algal and detritus) absorption coefficient (a_p , m^{-1}), water samples were filtered through fiberglass GF/F Whatman—0.7- μm porosity and 47-mm diameter. Then, transmittance–reflectance (T–R) method described by Refs. 28 and 29 was employed using a double-beam 2600 UV-VIS spectrophotometer equipped with an integrating sphere. The spectral sampling ranged from 280 to 800 nm, with a spectral resolution of 1 nm. To bleach pigments from the filters, a 10% sodium hypochlorite (NaClO) solution was used and T–R measurements were taken again. By eliminating pigments influence, it was possible to obtain detritus absorption coefficient (a_d , m^{-1}); a_p and a_d are then obtained according to Eq. (2):

$$a_{p,d}(\lambda) = \frac{2.303 \times OD_{p,d}(\lambda)}{V \times A}, \quad (2)$$

where $OD_p(\lambda)$ is the optical density of total particulate, $OD_d(\lambda)$ is the optical density of detritus, V is the filtered volume (m^3), and A is the filter clearance area (m^2). The phytoplankton absorption coefficients (a_p , m^{-1}) were calculated by subtracting a_d from a_p . Finally, a_{CDM} is calculated by adding a_{CDM} to a_d .

The a_t is calculated as the sum of the absorption coefficients of pure water (a_w), phytoplankton (a_p), nonalgal particles (a_{NAP}), and CDOM (a_{CDM}).

2.5 Quasianalytical Algorithm

QAA_{V5} and QAA_{V6} performances were evaluated for IHR. It is important to highlight that QAA_{V6} was developed to improve QAA accuracy in water systems in which R_{rs} at 670 nm is $>0.0015 \text{ sr}^{-1}$, i.e., coastal regions and areas with high sediments concentration. QAA_{BBHR} and QAA_{OMW}, the two reparameterized versions for Tietê River cascading system, were also tested. The dataset considered by Ref. 13 for QAABBHR parameterization presented high values and a broad range of Chl-*a* concentration (from 17.7 to 797.8 $mg \text{ m}^{-3}$; average 274.5 $mg \text{ m}^{-3}$) and SPM concentration ranging from 3.6 to 44.0 $g \text{ m}^{-3}$ (average 14.6 $g \text{ m}^{-3}$), with its organic fraction corresponding to $\sim 90\%$ of total SPM in the samples. Differently, the dataset used by Ref. 14 for QAA_{OMW} parameterization showed low average Chl-*a* concentration (16.15 $mg \text{ m}^{-3}$, ranging from 2.46 to 38.59 $mg \text{ m}^{-3}$), as well as low average SPM concentration (1.85 $g \text{ m}^{-3}$, varying from 0.10 to 5.30 $g \text{ m}^{-3}$); inorganic fraction was predominant in total SPM. Statistic metrics used to assess the QAAs performances for IHR were normalized root mean square error (NRMSE) and mean absolute percentage error (MAPE), considering a_p , a_{CDM} , and a_t estimated from water samples obtained in the field campaign as reference data.

The main improvements applied by the authors to adequate QAA for BBHR were changing λ_0 to 709 nm; calibration of χ and of the coefficients of the polynomial fit for empirical derivation of $a_t(\lambda_0)$; parameterization of ξ based on *in situ* a_{CDM} spectral slope and empirical calibration of ζ . These alterations aimed to adjust the model for the highly eutrophic environment of BBHR. Even presenting an oligo-to-mesotrophic status, the adjustment of λ_0 toward NIR region also showed to be satisfactory in NHR. To parameterize QAA for NHR, in addition to shifting λ_0 , the combination of wavelengths in the equation for χ estimation was modified, as well as the wavelengths for η calculation, which were calibrated based on *in situ* a_t measurements. Considering local data, band ratio and the offset coefficient for ζ estimation were changed in QAA_{OMW}, along with S optimization for ξ derivation; final steps for obtaining $a_{CDM}(\lambda)$

followed Ref. 8. For a_φ derivation, QAA_{OMW} follows an approach combining Refs. 30 and 31, using *in situ* dataset to reproduce the spectral shape for $a_\varphi(\lambda)$.

3 Results and Discussion

3.1 Water Quality Characterization

Chl-*a* and SPM concentrations varied widely (1.37 to 119.04 mg m⁻³ and 1.00 to 8.10 g m⁻³, respectively), showing significant spatial variability (Table 1). OSM represented 63% of the SPM, which means organic matter is slightly predominant. Turbidity values also showed some fluctuation, although it was lower than the variation presented by Chl-*a* and SPM (coefficient of variation 23.28%); in general, turbidity presented a relatively low average value (4.24 NTU). Secchi disk presented an average of 2.23 m and the lowest coefficient of variation (15.50%).

Regarding the absorption coefficients, $a_d(443)$ and $a_\varphi(443)$ presented similar average values, whereas $a_{CDOM}(443)$ presented the highest values— $a_d(443)$ and $a_\varphi(443)$. Detritus can be constituted of mineral matter, humus, or organic remains; considering that organic fraction of SPM showed to be slightly higher in this dataset and a_φ is not dominant, detritus in the reservoir is probably mainly constituted of humus and organic remains that are not resulting from phytoplankton degradation.³²

3.2 Bio-Optical Characterization

In situ measured R_{rs} spectra [Fig. 2(a)] show absorption features around 675 nm and a reflectance peak around 700 nm that can be associated with Chl-*a*. However, these R_{rs} features are not

Table 1 Descriptive statistics of water quality parameters and optical data in IHR; SD, standard deviation; CV, coefficient of variation; $n = 29$.

	Minimum	Maximum	Mean	SD	CV (%)
Chl- <i>a</i> (mg m ⁻³)	1.37	119.04	19.34	24.71	127.79
SPM (g m ⁻³)	1.00	8.10	2.45	1.40	57.19
OSM/SPM	0.29	0.88	0.63	0.15	23.28
ISM/SPM	0.12	0.71	0.37	0.15	40.11
Depth (m)	9.50	21.60	14.90	4.29	28.77
Turbidity (NTU)	2.82	8.87	4.24	1.19	28.01
Secchi depth (m)	1.60	3.20	2.23	0.35	15.50
$a_\varphi(443)$ (m ⁻¹)	0.06	1.88	0.30	0.36	120.44
$a_{CDOM}(443)$ (m ⁻¹)	1.06	2.78	1.61	0.30	18.64
$a_d(443)$ (m ⁻¹)	0.14	0.62	0.37	0.13	33.54
$a_{CDOM}(443)$ (m ⁻¹)	0.69	2.17	1.24	0.30	24.18
$a_t(443)$ (m ⁻¹)	1.13	4.68	2.06	0.80	38.95

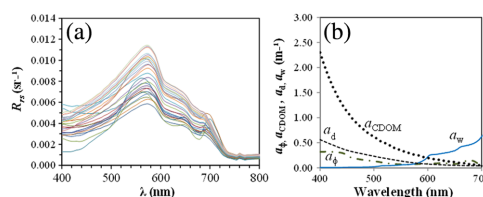


Fig. 2 (a) *In situ* R_{rs} spectra and (b) average absorption coefficients by phytoplankton (a_φ), CDOM (a_{CDOM}), detritus (a_d) and pure water (a_w); $n = 29$.

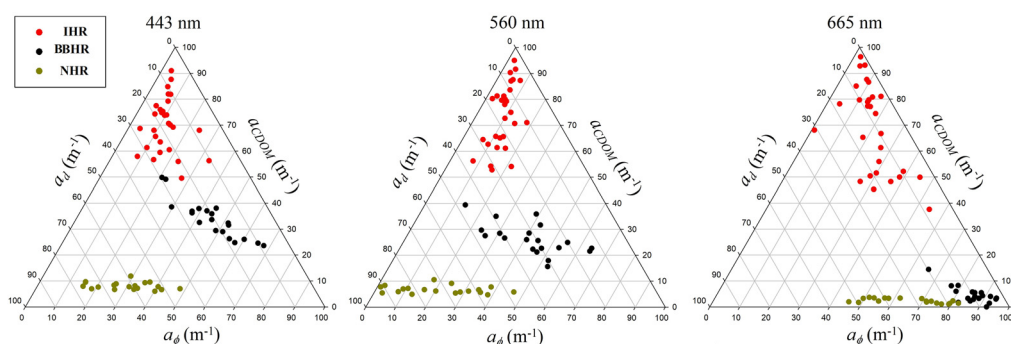


Fig. 3 Ternary plots presenting the relative contribution of detritus, phytoplankton and CDOM to the total absorption (without water fraction) at three different wavelengths for IHR (red dots), BBHR (black dots), and NHR (brown dots): for 443, 560, and 665 nm.

observed in all stations, indicating spatial variability in terms of optically significant constituents (OSC) at IHR. A prominent feature of reflectance in the green spectral region, around 550 nm, can be observed with different magnitudes [Fig. 2(a)]. This reflectance peak is usually related with scattering from algal cells,³³ and also with ISM, which shifts higher reflectance values toward longer wavelengths.³⁴ Average a_{CDOM} was the greatest contributor for a_t until around 660 nm [Fig. 2(b)]. Average a_d presented higher values than average a_ϕ through blue and green spectral regions; approximately at 600 nm average a_ϕ shows slight increase, and it is considerably greater than average a_d from 660 to 700 nm [Fig. 2(b)].

Ternary plots (Fig. 3) display the relative contribution of a_{CDOM} , a_ϕ , and a_d to the total absorption at IHR. Similar datasets from BBHR¹³ and NHR,¹⁴ collected in May, 2014, and May, 2016, respectively, are also plotted for comparison purpose. It is relevant to highlight that all three dataset considered here were collected during the dry season.

Absorption in IHR dataset is dominated by CDOM in all three wavelengths here considered, corresponding to $70.11\% \pm 3.94\%$ at 443 nm, $73.63\% \pm 4.67\%$ at 560 nm, and $68.94\% \pm 6.43\%$ at 665 nm. Detritus contributed to the total absorption budget with $18.13\% \pm 2.67\%$ at 443 nm and $17.18\% \pm 3.40\%$ at 560 nm. At 665 nm, phytoplankton contributed with $20.08\% \pm 5.27\%$ of the total while detritus contributed with only $10.97\% \pm 2.86\%$. In BBHR, total absorption was predominately due to phytoplankton in all three wavelengths, especially at 665 nm, with a_ϕ contributing with $85.72\% \pm 3.18\%$. However, at 443 and 560 nm the total absorption is more balanced among all OSC as can be observed in the scatter of dots around the central area of the plots [Figs. 3(a) and 3(b)]. In NHR dataset, otherwise, a_d was the major contributor at 443 and 560 nm, corresponding to $62.16\% \pm 4.42\%$ and $72.02\% \pm 6.52\%$, respectively. At 665 nm, a_ϕ represented $65.15\% \pm 5.97\%$ of total absorption, whereas a_d was the second greatest contributor with $32.51\% \pm 5.72\%$.

3.3 QAA Performances

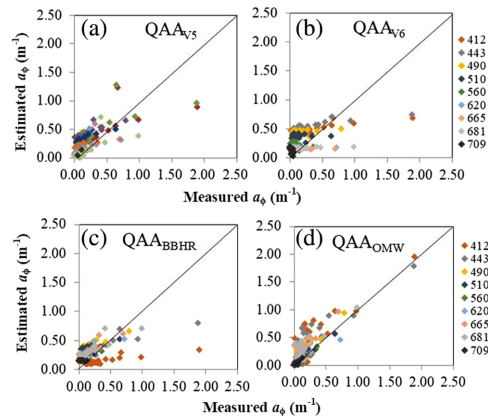
3.3.1 Absorption coefficient of phytoplankton (a_ϕ)

QAA_{OMW} presented the lowest average NRMSE (%) and MAPE (m^{-1}) for a_ϕ retrieval (Table 2). However, if we consider only 665-nm band, which is a diagnosis wavelength for Chl-*a*, this same version shows the highest errors (NRMSE = 31.74%, MAPE = 6.03 m^{-1}).

QAA_{OMW} tuning was based on NHR dataset, comprising Chl-*a* concentration values lower than the ones measured in IHR (NHR average 7.94 mg m^{-3} ; IHR average 19.34 mg m^{-3}), and this difference can be related to its poor performance at 665-nm band. QAA_{BBHR}, differently, was developed for a highly productive reservoir, and IHR bio-optical status did not showed to be similar to BBHR one, justifying why it was not completely suitable for retrieving a_ϕ in IHR. QAA_{V6} showed higher values of NRMSE and MAPE than QAA_{V5} in all wavelengths, although this performance was not initially expected as QAA_{V6} was proposed for R_{rs} (670) $> 0.0015 \text{ sr}^{-1}$, which fits IHR dataset [$R_{rs}(670) = 0.00427 \text{ sr}^{-1}$]. Comparatively,

Table 2 QAAs performances for a_φ retrieval, according to each band, based on NRMSE (%) and MAPE (m^{-1}).

Bands (nm)	QAA _{V5}		QAA _{V6}		QAA _{BBHR}		QAA _{OMW}	
	NRMSE (%)	MAPE (m^{-1})	NRMSE (%)	MAPE (m^{-1})	NRMSE (%)	MAPE (m^{-1})	NRMSE (%)	MAPE (m^{-1})
412	14.04	1.36	17.14	1.93	19.46	0.70	21.94	1.84
443	15.86	1.35	19.82	1.91	13.80	0.86	21.02	1.15
490	30.77	2.44	40.53	3.29	30.86	2.46	26.07	0.98
510	32.46	2.61	44.12	3.58	35.94	2.86	19.92	0.67
560	38.14	3.21	54.41	4.55	49.70	4.03	20.49	0.60
620	14.19	1.38	21.25	3.49	20.01	3.48	16.07	0.84
665	15.35	1.19	20.23	4.09	18.06	5.46	31.74	6.03
681	15.49	0.78	19.17	2.32	14.03	2.56	23.79	1.94
709	112.00	3.78	118.04	10.99	89.48	10.70	30.64	0.45
Average	32.03	2.01	39.41	4.02	32.37	3.68	23.52	1.61

**Fig. 4** Comparison between estimated and measured a_φ using: (a) QAA_{V5}, (b) QAA_{V6}, (c) QAA_{BBHR}, and (d) QAA_{OMW}.

QAA_{V5} showed to be the most accurate version at 665 nm (NRMSE = 15.35%, MAPE = 1.19 m^{-1}).

Scatterplots of a_φ estimated by the QAAs and a_φ measured in the laboratory are shown in Fig. 4. In general, QAA_{V5} overestimated a_φ in great part of wavelengths [Fig. 4(a)], except for the longer ones—specifically 681 and 709 nm—for which the version underestimated the values. The other versions overestimated most of the a_φ values at all wavelengths, excluding QAA_{BBHR} at 412 nm. It can be clearly observed that a sampling spot disagrees with the other ones in almost all wavelengths for all four QAA tested; this spot corresponds to the highest Chl-*a* and SPM concentrations of the dataset.

3.3.2 Absorption coefficient of CDOM plus detritus (a_{CDM})

For a_{CDM} , all QAAs variants demonstrated similar average NRMSE and MAPE, presenting reasonable accuracy (Table 3). An improvement in the NRMSE values can be observed with increasing wavelengths. QAA_{OMW} showed slight better results (NRMSE = 21.74%, MAPE = 0.44 m^{-1}); NHR is a nonproductive reservoir and it is dominated by a_{CDM} in the blue–green

Table 3 QAAs performances for a_{CDM} retrieval, according to each band, based on NRMSE (%) and MAPE (m^{-1}).

Bands (nm)	QAA _{V5}		QAA _{V6}		QAA _{BBHR}		QAA _{OMW}	
	NRMSE (%)	MAPE (m^{-1})	NRMSE (%)	MAPE (m^{-1})	NRMSE (%)	MAPE (m^{-1})	NRMSE (%)	MAPE (m^{-1})
412	50.41	0.85	49.60	0.84	41.17	0.67	29.49	0.35
443	40.71	0.87	40.26	0.86	34.54	0.70	24.64	0.31
490	32.75	0.91	32.56	0.90	29.54	0.78	21.43	0.33
510	30.46	0.92	30.33	0.91	28.07	0.81	20.69	0.34
560	25.94	0.95	25.89	0.94	24.83	0.86	19.71	0.39
620	23.52	0.97	23.50	0.97	23.07	0.92	19.82	0.49
665	22.42	0.98	22.41	0.98	22.20	0.95	19.98	0.57
681	21.91	0.98	21.91	0.98	21.74	0.96	19.85	0.58
709	21.65	0.99	21.65	0.99	21.54	0.97	20.07	0.64
Average	29.97	0.94	29.79	0.93	27.41	0.85	21.74	0.44

region, just as IHR, and this can be associated with the lower errors presented by QAA_{OMW}. As generally the estimation of a_{CDM} from remotely sensed data uses the wavelength of 443, the lowest NRMSE was observed for QAA_{OMW} version.

Figure 5 shows that QAA_{V5}, QAA_{V6}, and QAA_{BBHR} consistently underestimated a_{CDM} in all wavelengths. It probably occurs due to high CDOM and detritus concentrations and its relative predominance in the absorption budget of IHR (see Fig. 3), which is not commonly found in ocean and coastal waters, and also differs from BBHR optical characteristics. In these scatter-plots, two sampling spots evidently differ from the others: one of them is discrepant in all wavelengths and also corresponds to the spot with the highest Chl-*a* and SPM concentrations (as in Fig. 4), whereas the second one disagrees mostly in the shortest wavelengths—except for QAA_{OMW}, which overestimated a_{CDM} values for all wavelengths—and it corresponds to the second-highest SPM concentration in the dataset.

On the other hand, QAA_{OMW} was able to retrieve, in general, more consistent values of a_{CDM} [Fig. 5(d)]. It is important here to highlight that QAA approach unifies absorption by detritus and CDOM as a_{CDM} . Taking this into account, and also considering that NHR absorption occurs mostly due to detritus, while in IHR the CDOM is predominant, it corroborates the capability

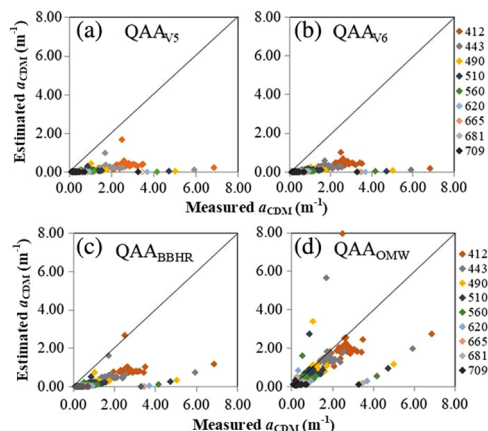
**Fig. 5** Comparison between estimated and measured a_{CDM} using: (a) QAA_{V5}, b) QAA_{V6}, (c) QAA_{BBHR}, and (d) QAA_{OMW}.

Table 4 QAAs performances for a_t retrieval, according to each band, based on NRMSE (%) and MAPE (m^{-1}).

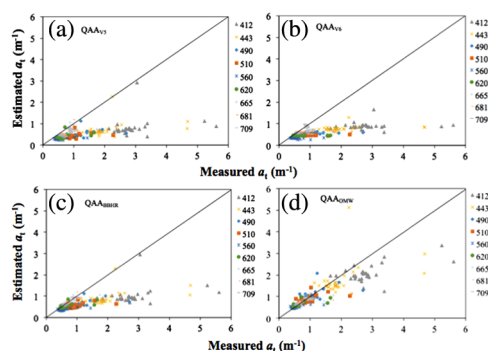
Bands (nm)	QAA _{V5}		QAA _{V6}		QAA _{BBHR}		QAA _{OMW}	
	NRMSE (%)	MAPE (m^{-1})	NRMSE (%)	MAPE (m^{-1})	NRMSE (%)	MAPE (m^{-1})	NRMSE (%)	MAPE (m^{-1})
412	55.76	2.88	53.59	2.26	50.41	1.93	32.97	0.41
443	42.51	2.06	40.90	1.60	37.10	1.30	24.41	0.27
490	36.00	1.36	33.54	1.01	29.19	0.78	17.26	0.18
510	34.56	1.27	32.15	0.94	27.78	0.71	16.37	0.18
560	31.67	0.97	29.35	0.71	24.56	0.49	17.23	0.18
620	28.52	0.58	27.42	0.37	20.78	0.20	18.73	0.16
665	25.43	0.43	24.49	0.24	16.02	0.11	15.89	0.13
681	21.84	0.40	22.16	0.27	15.72	0.13	16.15	0.16
709	78.17	0.27	66.89	0.17	28.39	0.07	33.05	0.08
Average	39.38	1.14	36.72	0.84	27.77	0.63	21.34	0.19

of QAA_{OMW} in retrieving a_{CDM} in nonphytoplankton-dominated waters and, consequently, its potential in determining carbon content in aquatic systems.

3.3.3 Total absorption coefficient (a_t)

Regarding the a_t retrieval, QAA_{OMW} performed better with NRMSE ranging from 15.89% to 32.97% (average of 21.34%) and QAA_{V5} was the one that presented the highest errors with 55.76% at 412 nm and average NRMSE of 39.38% (Table 4). The second best performance was observed for QAA_{BBHR} with NRMSE ranging from 15.72% to 50.51% (average of 27.77%). QAA_{OMW} great performance for a_t correlates with its good average results for a_φ and a_{CDM} , demonstrating that indeed, both initial steps for a_t estimation and the last ones—partitioning of a_t into a_φ and a_{CDM} —in QAA_{OMW} parameterization showed to be fairly adequate for IHR characteristics.

According to Tables 2 and 3, the highest errors for lower wavelengths observed in Table 4 were due to the estimation of a_{CDM} . The QAAs performed better for a_φ in lower wavelengths than in the longer ones. Scatterplots of a_t estimated by the QAAs and a_t measured in laboratory are shown in Fig. 6. These graphs show that the performance of QAA_{OMW} outperformed the other QAA versions.

**Fig. 6** Comparison between estimated and measured a_t using: (a) QAA_{V5}, (b) QAA_{V6}, (c) QAA_{BBHR}, and (d) QAA_{OMW}.

3.3.4 Implications for OSC monitoring

Neither originals nor reparameterized QAA versions tested here were capable of estimating accurately the absorption coefficients in all wavelengths. This shows a challenge in copying with high optical variability in cascading system. Although QAA_{BBHR} was parameterized for a high productive aquatic system, the model could not accurately retrieve the a_φ at 665 nm [$a_\varphi(665)$], which is used as a proxy to estimate Chl-*a* concentration. This possibly happened because Chl-*a* concentration in BBHR is very high ($\sim 797 \text{ mg m}^{-3}$), and due to this the R_{rs} spectra are masked by the package effect, as reported by Ref. 35. For $a_\varphi(665)$, the QAA_{V5} presented the lowest error. On the other hand, to estimate the a_{CDM} at 443 nm [$a_{\text{CDM}}(443)$], which is a proxy for carbon content³⁶ the QAA_{OMW} presented the lowest error. These results highlighted the limitation of such quasianalytical scheme to monitor the spatial-temporal OSC in the cascading system. However, the obtained results also demonstrate an opportunity to better understand the complexity of these aquatic systems and try to figure out how to improve the QAA to use only one version to estimate the OSC in the entire cascade. Up to now, the monitoring of OSCs from space operationally is still a challenge.

4 Conclusion

As QAA native versions, QAA_{V5} and QAA_{V6}, were designed for ocean and coastal waters, a poor performance for optically complex inland waters was already expected and it was confirmed. However, QAA_{BBHR} did not performed satisfactorily as supposed, demonstrating that even presenting some similarity regarding the OSCs, the bio-optical status of BBHR and IHR are substantially different and, thus, IHR IOPs cannot be derived through this reparameterized QAA. Therefore, our hypothesis can be rejected as QAA_{BBHR} was not suitable for retrieving neither a_φ nor a_{CDM} . Although QAA_{OMW} was able to retrieve relatively accurate average values for a_φ and a_{CDM} values in IHR, in addition performing relatively well for $a_{\text{CDM}}(443)$, it presented an unsuitable performance for $a_\varphi(665)$, indicating that this version also has its limitations for deriving IHR IOPs at wavelengths used as proxy for OSCs concentrations estimation. Variability of bio-optical characteristics along the cascading system was confirmed as none reparameterized version for reservoirs in the same system was completely suitable for IHR in all wavelengths. Even though the algorithms did not performed satisfactorily in all wavelengths, results for specific wavelengths show potential when it comes to OSCs concentration retrieval. We can highlight the good, it is possible to point out the good QAA_{V5} performance for $a_\varphi(665)$ estimation and QAA_{OMW} performance for $a_{\text{CDM}}(443)$. These results demonstrate the possibility of using QAA versions that showed to be suitable for a particular absorption coefficient at a certain wavelength for an application that relates to it, using them as a proxy and then applying it to estimate the OSC in inland waters from space. In any case, further research is necessary, either aiming to achieve a QAA reparameterization appropriate for waters with widely differing optical properties or analyzing the suitability of other analytical methods.

Disclosures

No potential conflict of interest was reported by the authors.

Acknowledgments

The authors thank FAPESP Projects (Process Nos. 2012/19821-1 and 2015/21586-9) and Professor Edivaldo Velini and staffs from FCA/UNESP for allowing the use of their laboratory facilities.

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