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# Assessing water quality in the northern Adriatic Sea from HICO<sup>™</sup> data

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### Assessing water quality in the northern Adriatic Sea from HICO<sup>TM</sup> data

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This letter focuses on water-quality estimation in the northern Adriatic Sea using physically-based methods applied to image obtained with the Hyperspectral Imager for the Coastal Ocean (HICO<sup>TM</sup>). Optical properties of atmosphere and water were synchronously measured to parameterise such methods. HICO<sup>TM</sup>-derived maps of chlorophyll-*a* (chl-*a*) and suspended particulate matter (SPM) indicated low values, in the range of 0–3 mg m<sup>-3</sup> and 0–4 g m<sup>-3</sup>, respectively, correlating significantly with field data ( $R^2 = 0.71$  for chl-*a* and  $R^2 = 0.85$  for SPM). The results, on analysis, identify clear waters in the open sea and moderately turbid waters near the coast due to river sediment discharge and organic matter from coastal lagoons. These findings support the use of HICO<sup>TM</sup> data to assess water-quality parameters in coastal zones and suggest the feasibility of integrating them with future-generation space-borne hyperspectral images.

#### 1. Introduction

Coastal zones are important interfaces between land, sea and atmosphere, acting as a boundary for oceanic circulation and currents, as well as receiving inputs from the land. River discharge, wind energy, tidal mixing and long-shore currents influence the distribution of suspended and dissolved matter in coastal zones, varying their concentration, size and physical-chemical composition. The organic component, be it dissolved or particulate, is also generated and converted within the coastal system (by photosynthesis and degradation) and its distribution may vary considerably. The impact of human activities (e.g. changes in land use, sewage discharge, aquaculture industry and port activities) on coastal ecosystems further increases their variability.

The Hyperspectral Imager for the Coastal Ocean (HICO<sup>TM</sup>) is the first space-borne hyperspectral sensor designed specifically for the coastal ocean and for estuarine,

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riverine, lacustrine and other shallow-water areas. The HICO<sup>TM</sup> acquires images from different angles in selected areas, at various times of day. Each pixel covers  $90 \times 90$  m, with 87 nominal spectral channels in the range of 400–900 nm providing information on water properties and bottom reflectance (see Lucke *et al.* 2011).

HICO<sup>TM</sup> images have been proved to be useful for estimating chlorophyll-*a* (chl-*a*) (Gitelson *et al.* 2011), as well as for monitoring suspended sediments in river plumes (Tufillaro *et al.* 2010). Most of the related studies focused on very turbid and productive waters, applying semi-empirical algorithms optimized for HICO<sup>TM</sup> spectral bands. In the present study, the aim is instead to use the HICO<sup>TM</sup> data for assessing water quality in moderately productive waters. This work concerns an area in the northern Adriatic Sea defined by Berthon and Zibordi (2004) as a frontal area between case 1 and case 2 conditions according to the Morel and Prieur (1977) optical water type classification.

Water quality was assessed by (1) defining a two-component bio-optical model parameterised with specific inherent optical properties for the study area, (2) correcting the HICO<sup>TM</sup> at-sensor radiances for both atmospheric and adjacent effects and (3) adopting a bio-optical model inversion technique. The study used data collected by the Aerosol Robotic Network-Ocean Colour (AERONET-OC) (Holben *et al.* 1998) at the Acqua Alta Oceanographic Tower (AAOT). These data were used both to run the atmospheric correction code and to validate HICO<sup>TM</sup>-derived reflectance. Water-quality parameters (e.g. concentrations of chl-*a*, suspended particulate matter (SPM) and coloured dissolved organic matter (CDOM)) and water reflectances collected during the satellite overpass were also considered for validation purposes.

#### 2. Materials and methods

#### 2.1 Study area and in situ measurements

The northern Adriatic Sea is in the northernmost part of the Mediterranean and is characterized by a shallow depth (averaging 35 m). It receives approximately 20% of the total Mediterranean river run-off, mainly from the Po (the longest river in Italy), and its hydrodynamics are seasonally influenced by meteorological physical forcing (winds and tides). The northern Adriatic is also considered to be one of the most productive regions of the generally oligotrophic Mediterranean Sea. The largest phytoplankton blooms occur in its surface layers in late winter and in summer, they occur in the western part of the region. According to Hooker *et al.* (2004), the area alternates between case 1 and case 2 water types. Case 2 conditions are mainly due to the effects of local winds, which resuspend bottom sediments, and due to the discharge from the Po River and other rivers to the north (Adige, Piave, Tagliamento and Isonzo), which are sources of terrestrial particulate and dissolved matter. The Venice Lagoon and the Grado–Marano Lagoon lie along the northern Adriatic coastline, and the water exchange between the lagoons and the sea is governed by tide and contributes to the input of organic matter in the coastal waters (Ferrarin *et al.* 2013).

A 7-day cruise was undertaken in the study area (figure 1) in the second week of May 2012 to gather data for the purpose of calibrating a bio-optical model and assessing the HICO<sup>TM</sup>-derived products, including water reflectance and water-quality parameters. At 18 stations (figure 1), discrete water samples were collected to measure absorption and concentration at two different depths (at the surface, by sampling the first integrated meter, and at the Secchi disk depth); then, the samples were filtered immediately *in situ* and were stored for subsequent laboratory analysis. The concentration



Figure 1. Study area with location of fieldwork activities carried out in the second week of May 2012 (green and red dots) and portion of Adriatic Sea imaged by HICO<sup>TM</sup>. The stations visited on 10, 11 and 12 May 2012 are indicated with red dots; the data from these stations (which include also AERONET-OC AAOT site) allow us to validate the proposed methodology.

of chl-*a* was measured using the trichromatic method (APHA 1981), the concentration of SPM was measured with the gravimetric method (Van der Linde 1998) and the concentration of CDOM was assumed from the  $a_{\text{CDOM}}(\lambda)$  at 440 nm ( $\lambda$  indicates the wavelength). The absorption spectra of particles retained on the filters,  $a_p(\lambda)$ , were obtained using the filter pad technique (Strömbeck and Pierson 2001) and were calculated according to Babin *et al.* (2003). The spectrophotometric determination and processing of the CDOM absorption spectra,  $a_{\text{CDOM}}(\lambda)$ , were derived from Kirk (1994). At the same stations where the water was sampled, remotesensing reflectance ( $R_{rs}$ ) values (omitting the wavelength dependence for the sake of clarity, apart from a few cases) were measured with a WISP-3 spectroradiometer (Water Insight, Wageningen, The Netherlands; Hommersom *et al.* 2012) in the optical range of 380–800 nm. A HydroScat-6 backscattering sensor (HOBILabs, Tucson, AZ, USA) was also used to estimate the backscattering coefficient of the particles ( $b_{b_p}(\lambda)$ ) (Maffione and Dana 1997).

Some of the *in situ* data used in this study were provided by the AERONET-OC AAOT station (Zibordi *et al.* 2009), located in the northern Adriatic approximately 15 km south-east of the Venice lagoon (12.51° E, 45.31° N); these data enabled us to identify the atmospheric parameters useful for the atmospheric correction of the HICO<sup>TM</sup> image, as well as providing reference water reflectance spectra to validate the HICO<sup>TM</sup>-derived reflectances.

#### 2.2 Bio-optical modelling

The bio-optical model used in this study was based on works by Lee *et al.* (1998, 1999), where the  $R_{\rm rs}(\lambda)$  above water was a function of the subsurface radiance reflectance subsequently calculated as a function of the absorption  $a(\lambda)$  and backscattering  $b_{\rm b}(\lambda)$ 

coefficients. To express the dependence of  $a(\lambda)$  and  $b_b(\lambda)$  on constituent concentrations (chl-*a*, CDOM and SPM), the spectral absorption coefficient,  $a(\lambda)$ , was modelled as

$$a(\lambda) = a_{\rm w}(\lambda) + a_{\rm CDOM}(440) \times e^{-S_{\rm CDOM} \times (\lambda - 440)} + [\text{chl} - a] \times a_{\rm p}^*(\lambda) \tag{1}$$

where,  $a_w(\lambda)$  is the pure-water absorption (Smith and Baker 1981, Pope and Fry 1997),  $S_{\text{CDOM}}$  is the slope factor commonly used in modelling the absorption spectra of CDOM, [chl-*a*] is the concentration of chl-*a* and  $a^*_{p}(\lambda)$  is the specific absorption of the particles, including phytoplankton and detritus. In this study,  $S_{\text{CDOM}}$  was established from each CDOM absorption spectrum measured, and the average slope (0.018) was used as the model parameter. The specific absorption for all particles,  $a^*_{p}(\lambda)$ , was the average value of the chlorophyll-specific particle absorption coefficients. In both cases, the average values were used because the data sets showed a low range of variation, with a coefficient of variation below 5% for both the CDOM slope and the chlorophyll-specific particle absorption coefficient.

The spectral backscattering coefficient,  $b_b(\lambda)$ , was modelled as

$$b_{\rm b}(\lambda) = b_{\rm b_w}(\lambda) + [{\rm SPM}] \times b^*_{\rm b_p}(\lambda)$$
<sup>(2)</sup>

where the backscattering coefficient of pure water,  $b_{b_w}(\lambda)$ , was taken from Dall'Olmo and Gitelson (2006), [SPM] is the SPM concentration and  $b^*_{b_p}(\lambda)$  is the particlespecific backscattering coefficient. The backscattering coefficient of the SPM was modelled as an inverse power function of wavelength.

The scalars enabling the total  $a(\lambda)$  and  $b_b(\lambda)$  to be correlated with the subsurface remote-sensing reflectance and  $R_{rs}(\lambda)$  were ascertained by fitting the forward run of the bio-optical model to *in situ* measurements of  $R_{rs}(\lambda)$  taken at the 18 stations. The bio-optical model needs to be inverted to retrieve in-water constituent concentrations. This was done with the BOMBER software (Giardino *et al.* 2012) that inverts the biooptical model using optimization techniques, while simultaneously retrieving waterquality parameters from remotely-sensed image atmospherically corrected to  $R_{rs}(\lambda)$ values.

#### 2.3 Image processing

A hyperspectral image was acquired by the HICO<sup>TM</sup> sensor on 12 May 2012. The signal-to-noise ratio (SNR) of the at-the-sensor radiance in the HICO<sup>TM</sup> data was investigated according to Gao (1993): for each sensor channel, the local standard deviation (LSD) was calculated for each block of pixels in the scene, identified using a moving window technique; the maximum value on the LSD histogram (representing the mean noise of the image) was then used to calculate the SNR, defined as the ratio between the mean value signal and the mean noise of the image.

The at-the-sensor radiance was converted into  $R_{rs}$  values above the water surface by adapting the algorithm presented in Bassani *et al.* (2010) to the HICO<sup>TM</sup> image. The algorithm, called the HICO@CRI (HICO<sup>TM</sup> atmospherically corrected reflectance image), is based on the atmospheric correction method presented in Vermote *et al.* (1997), using the 6SV radiative transfer code (Vermote *et al.* 1997, Kotchenova *et al.* 2008). The HICO@CRI also implements a correction for the adjacency effect according to Vermote *et al.* (1997) using the empirical formula

$$\rho_{i} = \rho_{i}^{s} + \frac{t_{d(\mu_{v})_{i}}}{e^{-\tau/\mu_{v}}} [\rho_{i}^{s} - \langle \rho_{i}^{s} \rangle]$$
(3)

where  $\rho_i$  is the at-ground reflectance of the *i*th channel,  $\rho_i^s$  is the reflectance with the environmental contribution and  $e^{-\tau/\mu_v}$  and  $t_d(\mu_v)_i$  are the direct and diffuse components of the atmospheric transmittance along the target-sensor direction, respectively. These radiative quantities depend on the cosine of the viewing zenith angle,  $\mu_v = \cos \theta_v$ . The direct component is also defined by the total optical thickness,  $\tau$ . The  $< \rho_i^s >$  is the average of the  $\rho_i^s$  for the *i*th channel, calculated on the whole image to ensure that the adjacency effect is removed from each pixel in the image.

The HICO@CRI inputs on aerosol optical thickness at 550 nm and precipitable water vapour were obtained at the AERONET-OC AAOT station, as retrieved from the direct sun algorithm at level 2.0 (Eck *et al.* 1999). The size distribution of the aerosol was obtained from the diffuse component corresponding to cloud-screened level 1.5 (Smirnov *et al.* 2000). The aerosol optical thickness at 550 nm (equal to 0.25) showed a low aerosol loading at the time of the HICO<sup>TM</sup> image acquisition, while the size distribution indicated an aerosol model dominated by fine particles, typical situation of the Adriatic Sea, as reported in Mélin *et al.* (2006).

#### 3. Results and discussion

The SNR for the HICO<sup>TM</sup> data showed an average value of about 80 for all the 87 sensor channels for an *albedo* value of about 1%. In particular, the analysis showed a decreasing SNR for increasing wavelengths, with an average value of about 100 in the first 30 bands from 400 to 570 nm, the minimum value of 40 only being reached at bands 74 and 75, at around 825 nm. These results are consistent with the findings of Moses *et al.* (2012).

The methods used to assess water quality from the HICO<sup>TM</sup> image were assessed using *in situ* data collected at the 9 stations (of the 18) visited synchronously with the HICO<sup>TM</sup> overpass on 12 May 2012 (figure 1, red dots) and 2 days previously, since a 2-day mismatch between *in situ* data and acquired image is acceptable for the purpose of validating water-quality products based on the assumption that environmental variables (e.g. wind and river discharge) are fairly stable (Odermatt *et al.* 2010).

Figure 2 shows the convergence of the  $R_{\rm rs}$  values obtained from atmospherically corrected image data, *in situ* spectroradiometric data and forward bio-optical modelling. The plot is given for the wavelengths measured by the radiometer at the AERONET-OC AAOT station (412, 442, 490, 530, 551, 667 and 868 nm), which are common to the HICO<sup>TM</sup> data and WISP-3 spectroradiometer (except for the band at 868 nm). The plotted spectra are the average values of the 9 measurements obtained at the stations measured in the footprint of the HICO<sup>TM</sup> during the satellite's overpass and on the 2 days beforehand. The HICO<sup>TM</sup>-derived  $R_{\rm rs}$  values were extracted from 3 × 3 pixel areas centred on the position of the *in situ* measurements, as suggested by Patt (2002) and Bailey and Werdell (2006). The *in situ*  $R_{\rm rs}$  are the average values of WISP-3 measurements combined with the data gathered at the AERONET-OC AAOT station. The modelled  $R_{\rm rs}$  values were obtained from the forward run of the bio-optical model using *in situ* data for the chl-*a*, SPM and CDOM concentrations.

Overall, a good consistency was obtained across the spectrum, with a relative root mean square error (RMSE) of 8% and 18%, respectively, when the modelled and HICO<sup>TM</sup> data were compared with *in situ* measurements. The HICO<sup>TM</sup>-derived  $R_{rs}$ 



Figure 2. In situ, forward-modelled and HICO<sup>TM</sup>-derived  $R_{rs}$  spectra. 'In situ' spectrum is the average value measured with a WISP-3 spectroradiometer of the stations visited on 10-11-12 May 2012 plus the AERONET-OC AAOT site data (cf. figure 1, red dots).

values diverged at the shorter wavelengths (412 and 442 nm), however, showing an inverse trend between 412 and 490 nm. This mismatching behaviour might make estimations of CDOM from image data unpredictable. In fact, the  $R_{rs}$  values in the blue region are often used to retrieve CDOM concentrations because they are sensitive to changes in CDOM concentrations (Kutser *et al.* 2005). As in previous works (e.g. Giardino *et al.* 2007), the bio-optical model for mapping water quality was inverted by keeping a fixed CDOM concentration; the average value of 0.06 m<sup>-1</sup> obtained from *in situ* measurements was used in the inversion performed with BOMBER.

Figure 3 presents the two BOMBER-retrieved HICO<sup>TM</sup> maps with ranges of variation of  $0-3 \text{ mg m}^{-3}$  for chl-*a* and  $0-4 \text{ g m}^{-3}$  for SPM. The chl-*a* map shows quite low concentrations, with homogenous patterns except in the south-west, where there was a residual phytoplankton bloom induced by the nutrients discharged by the River Po and then dispersed by coastal currents. The SPM map likewise depicts fairly



Figure 3. The BOMBER-retrieved products ((*a*) the SPM and (*b*) the chl-*a* concentrations) obtained from HICO<sup>TM</sup> data acquired on 12 May 2013. The boxes indicate the areas where there were marked differences in the SPM and chl-*a* concentrations (these areas were also used to label the spectra in figure 4).



Figure 4. HICO<sup>TM</sup>-derived  $R_{rs}$  spectra in waters affected by rivers plumes (1 and 3) and a clearer area (2), where were marked differences in the SPM and chl-*a* concentrations were found (cf. figure 3).

homogenous patterns with SPM concentrations around 1 g m<sup>-3</sup>. The patchiest areas are in the coastal zones, particularly near the Grado and Marano lagoon, where the northern tributaries (figure 1) carry significant amounts of inorganic detritus. There is little correlation between the maps, save in the south-west, where the correspondence between SPM and chl-*a* is due to the load of nutrients discharged into the Adriatic Sea by the Po river causing a rapid increase in phytoplankton bloom, so that most of the suspended matter in this area is due to phytoplankton (Giani *et al.* 2001).

Figure 4 shows the HICO<sup>TM</sup>-derived  $R_{rs}$  spectra coinciding with three areas where there were marked differences in the SPM and chl-*a* concentrations. These areas include waters affected by rivers plumes (figure 3, boxes 1 and 3) and a clearer area (figure 3, box 2) further away from the coast. The degree of variation in the three HICO<sup>TM</sup>-derived spectra explains the chl-*a* and SPM patterns shown in figure 3. The highest spectra (#3 in figure 4) reveal the contribution of SPM due to discharge from northern tributaries; the spectra decreasing with wavelengths (#2 in figure 4) are typical of clear marine waters (Morel and Prieur 1977) and explain the low concentrations of both SPM and chl-*a* (figure 3, box 1) and the third spectra (#1 in figure 4) peaking at 570 nm explain the patterns of moderately high chl-*a* concentrations found around the mouth of the Po river. Finally, all the spectra show a trend with peaks and dips at shorter wavelengths, where the HICO<sup>TM</sup> data appear to be less reliable (figure 4).

Figure 5 shows two scatter plots depicting the HICO<sup>TM</sup>-derived estimations of the chl-*a* and SPM concentrations versus the *in situ* measurements obtained at the 9 stations sampled at the time of the HICO<sup>TM</sup> overpass (figure 1, red dots). The HICO<sup>TM</sup> image was again averaged on a  $3 \times 3$  pixel area centred on the location of the sampling stations. The HICO<sup>TM</sup>-derived SPM concentration was consistent with the *in situ* data, with a correlation coefficient *r* of 0.92, a coefficient of determination  $R^2$  of 0.85 and a relative RMSE of 25.1%, and coming close to the 1:1 line with a slope of 0.985 (figure 5(*a*)). The chl-*a* data validation was positive too, showing an *r* of 0.84, an  $R^2$  of 0.71, a relative RMSE of 16.5% and a slope of 1.013 (figure 5(*b*)).



Figure 5. Scatter plots of HICO<sup>TM</sup>-derived products and *in situ* concentrations measured in 9 stations (cf. figure 1 red dots): (*a*) the SPM plot and (*b*) the chl-*a* plot. The statistics of fitting are given as correlation coefficient (*r*), coefficient of determination ( $R^2$ ), relative root means square error (RMSE) and mean absolute error (MAE). In both plots, the number of samples (*n*) is 9 and the 1:1 line is plotted as dotted lines.

#### 4. Conclusions

A HICO<sup>TM</sup> image of the northern Adriatic Sea was acquired on 12 May 2012 and was used for assessing water-quality parameters. Physically-based methods were used to convert the at-the-sensor radiance into  $R_{rs}$  values (HICO@CRI, Bassani *et al.* 2010) and to retrieve synchronous water-quality parameters from image data (BOMBER, Giardino *et al.* 2012). These methods were parameterised using *in situ* measurements to characterize both the optical properties of atmosphere and the specific inherent optical properties of water.

The performance of the proposed method was measured by comparing the  $R_{rs}$  values obtained by forward modelling, *in situ* measurements and atmospherically corrected HICO<sup>TM</sup> data (including a correction for adjacent effects). The match of the  $R_{rs}$  spectra was sufficient for the purpose of this study, but further investigations are needed at the shortest wavelengths to obtain a better consistency of the HICO<sup>TM</sup> data.

This study confirms that the HICO<sup>TM</sup> has a good enough SNR and spatial resolution for assessing the complexity of coastal features, enabling local phenomena to be distinguished. While we await the next generation of space-borne hyperspectral imagers (e.g. SENTINELs, HYSPIRY, PRISMA, EnMap), the HICO<sup>TM</sup> facility has given us an opportunity to simulate and verify the new sensors' capabilities.

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