Ocean color satellite atmospheric correction

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Acknowledgements: Zia Ahmad, Sean Bailey, Bryan Franz, Amir Ibrahim, and Fred Patt

2021 Ocean Optics Summer Course



Today's Lecture

The previous lecture illustrated the atmospheric correction problem



Today's lecture shows one solution

- work through the steps of ocean color atmospheric correction as used by NASA/OBPG
- identify the places where ancillary (external) data are required
- identify the places where in situ data and bio-optical models are used

For the details see

NASA/TM-2016-217551



Atmospheric Correction for Satellite Ocean Color Radiometry

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https://oceancolor.gsfc.nasa.gov/docs/technical/ NASA-TM-2016-217551.pdf

OCFANI OPTICS Web Book

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Light and Radiometry

- Overview of Optical Oceanography
- Absorption
- Scattering
- Optical Constituents of the Ocean
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Chapter 10 of The Ocean Optics Book

http://www.oceanopticsbook.info





where all radiances are defined at the top-of-atmosphere (TOA)



Processing Constraints

MODIS and VIIRS have about 14 orbits/day

The distance between terminators is ~20,000 km. The MODIS +/-50 degrees of scan angle gives ~1220 pixels, or about 24.4 Mpixels/orbit. For VIIRS the useful scan angle range corresponds to ~2480 pixels, and the along-track resolution is 0.75 km, so about 66.1 Mpixels/orbit. If we limit processing to SZA < 75 degrees, then the numbers are 5/6 of the above, which is 20.3 Mpix for MODIS and 55.1 for VIIRS.

Currently have 2 MODIS and 2 VIIRS in orbit (coming soon, PACE and 3 more VIIRS), so 151 Mpixels/orbit, times 14 orbits/day = 2.1 Gigapixels/day to be processed. **Every pixel gets its own atmospheric correction.**

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If atmospheric correction takes 1 second/pixel, this is 67 YEARS to process 1 DAY of collected data!

Atmospheric correction must be done in near real time. (One day to process one day's data.) This requires processing ~34,000 pixels/second (or 0.00003 sec/pixel for one computer; if have 100 computers, then 0.003 sec/pixel)

This is a severe constraint on what atmospheric correction algorithms are used. You WANT the AC algorithms to be as accurate as possible, but they MUST BE computationally FAST



where all radiances are defined at the top-of-atmosphere (TOA)

(2)
$$L_t = L_R + [L_a + L_{Ra}] + TL_g + tL_f + tL_w$$

where L_g , L_f , and L_w are now defined at the sea surface, and L_{sky} is accounted for in Rayleigh correction. T and t are the direct and diffuse transmittance.

Direct: one particular path connects the source & observer



Diffuse: radiance from all locations & directions can be scattered into the direction of interest



where all radiances are defined at the top-of-atmosphere (TOA)

(2)
$$L_t = L_R + [L_a + L_{Ra}] + TL_g + tL_f + tL_w$$

where **L**_g, **L**_f, and **L**_w are now defined at the sea surface and **L**_{sky} is accounted for in Rayleigh correction. **T** and **t** are the direct and diffuse transmittance.

factor out

gaseous diffuse

transmissions: (3)
$$L_{\rm t} = \left(L_{\rm r} + [L_{\rm a} + L_{\rm ra}] + t_{\rm dv}L_{\rm wc} + t_{\rm dv}L_{\rm w}\right)t_{\rm gv}t_{\rm gs}f_{\rm p} + TL_g$$

 t_{gv} is the diffuse transmission by atmos. gases in the viewing direction t_{gs} is the diffuse transmission by atmos. gases in the Sun's direction t_{dv} is the diffuse transmission along viewing path of the sensor

This "factoring" is done for reasons of computational efficiency. Theoreticians often use (1) or (2); OBPG works with (3).

Justification

Consider the Rayleigh term: $L_{
m R} = L_{
m r} t_{
m gv} t_{
m gs} f_{
m p}$

The TOA Rayleigh contribution L_R depends on Sun and viewing geometry, absorbing and non-absorbing atmospheric gasses, sea-level pressure, and polarization. This is a serious RT calculation, including polarization.

The L_r term is a "standard" Rayleigh contribution computed using a standard atmosphere and only nonabsorbing gases N₂ and O₂, for various Sun & viewing geometries. This can be computed once and placed in a look-up table.

The gaseous transmittances are computed by use of gas absorption coefficients, computed path lengths, and gas concentrations for the various absorbing gases. Compute once and put in a LUT.

The f_p term is a polarization correction, which depends on atmosphere and surface polarization states (modeled Rayleigh and glint Stokes vectors) and the sensor-specific polarization sensitivity with viewing direction. Again, compute once for various inputs and make a LUT.

 L_r , the diffuse transmittances, and f_p are pre-computed and stored in look-up tables as functions of Sun and viewing geometry, gas concentrations, instrument polarization sensitivity, etc. Evaluation of L_R then requires no real-time radiative transfer calculations.

Ditto for the other terms in the L_t equation

Keep in Mind What We Want: R_{rs}

ocean color satellites measure top-of-atmosphere radiances $\lambda_{t} = \left(L_{r} + \left[L_{a} + L_{ra}\right] + t_{dv}L_{f} + t_{dv}L_{w}\right) t_{gv} t_{gs} f_{p} + TL_{g}$ $R_{rs} = \frac{(L_w)^{\leftarrow ----}}{F_0 \cos(q_s) t_{ds} f_s f_b f_l}$ desired we desire (normalized)

remote sensing reflectances

11

Now sequentially step through the meaning & evaluation of each term in these equations



top-of-atmosphere radiance

$$L_t = (L_r + [L_a + L_{ra}] + t_{dv}L_f + t_{dv}L_w)t_{gv}t_{gs}f_p + TL_g$$

Menghua Wang, IOCCG Report 10



 L_w is often <10% of L_t !

0.5% error in atmospheric correction or calibration \rightarrow 5% error in L_w

processing cadence: L_t





transmittances



gaseous transmittances



calculating gaseous transmittance requires ancillary data



example using OMI (Ozone Monitoring Instrument on Aura satellite) ozone measurements

 $au_{{\rm O}_3}(\lambda) = [{\rm O}_3] k_{{\rm O}_3}(\lambda)$ for nadir direction

$$t_{\rm O_3} = \exp\left[-\tau_{\rm O_3}\left(\frac{1}{\cos\theta_{\rm s}} + \frac{1}{\cos\theta_{\rm v}}\right)\right]$$



all ancillary data are not created equal

comparison of three ancillary sources of O_3 :

TOAST OMI EPTOMS and climatology

small differences in ancillary data can lead to big differences in geophysical products



TOAST: Total Ozone from Analysis of Stratospheric and Tropospheric components OMI: Ozone Mapping Instrument EPTOMS: Earth-Probe Total Ozone Monitoring Spectrometer ...and there are others

No surprise: Lots of NO₂ near NY City and Philadelphia



.80e+16 1.60e+16 1.40e+16 1.20e+16 1.00e+16 8.00e+15 -6.00e+15 4.00e+15 2.00e+15

RGB image of a MODIS-Aqua scene from 11 April 2005

OMI tropospheric NO2 amount on 11 April 2005

From Ahmad et al. (2007)

processing cadence: $L_t / t_{gv} / t_{gs}$





 t_{gs} (in the Sun's direction at 443)

0.996

tg_sol_443

0.99

0.993

tg_sen_443 0.99 0.993 0.996 1

t_{gv} (in the sensor viewing direction at 443)



instrument polarization sensitivity

instrument polarization correction factor (pre-launch measurement)

$$L_{t} = \left(L_{r} + [L_{a} + L_{ra}] + t_{dv}L_{f} + t_{dv}L_{w}\right)t_{gv}t_{gs}f_{p} + TL_{g}$$

$$R_{rs} = \frac{L_{w}}{F_{0}\cos(q_{s})t_{ds}f_{s}f_{b}f_{l}}$$
MODIS requires an additional correction

processing cadence: $L_t / t_{gv} / t_{gs} / f_p$





some satellite instruments include depolarizers in their fore optics, which mitigates instrument polarization sensitivity Atmospheric correction of ocean color sensors: analysis of the effects of residual instrument polarization sensitivity

Howard R. Gordon, Tao Du, and Tianming Zhang

APPLIED OPTICS / Vol. 36, No. 27 / 20 September 1997

Moderate-Resolution Imaging Spectroradiometer ocean color polarization correction

Gerhard Meister, Ewa J. Kwiatkowska, Bryan A. Franz, Frederick S. Patt, Gene C. Feldman, and Charles R. McClain

APPLIED OPTICS / Vol. 44, No. 26 / 10 September 2005

foam & whitecaps

$$L_{t} = \left(L_{r} + [L_{a} + L_{ra}] + t_{dv}L_{f} + t_{dv}L_{w}\right)t_{gv}t_{gs}f_{p} + TL_{g}$$

$$R_{rs} = \frac{L_{w}}{F_{0}\cos(q_{s})t_{ds}f_{s}f_{b}f_{l}}$$

foam & whitecaps



Hyperspectral Measurements, Parameterizations, and Atmospheric Correction of Whitecaps and Foam From Visible to Shortwave Infrared for Ocean Color Remote Sensing

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Department of Marine Sciences, University of Connecticut, Groton, CT, United States

 $\rho_{\rm f}$ = π L_f = A F_{wc}

A = 22% (11-33%) from Koepke 1984 + a correction for decreasing reflectance in red & NIR

 $\rho_{\rm f}(412) = \pi L_{\rm f}(412) = 1.925 \times 10^{-5} (U_{10} - 6.33)^3$

estimation of contribution of whitecaps & foam requires ancillary wind data (NCEP)



processing cadence: $L_t / t_{gv} / t_{gs} / f_p - tL_f$





tLf 443

0.015

0

(W m^-2 um^-1 sr^-1)

0.03

0.045

0.06





molecular (Rayleigh) scattering

$$L_{t} = \left(L_{r}\right) + \left[L_{a} + L_{ra}\right] + t_{dv}L_{f} + t_{dv}L_{w}\right)t_{gv}t_{gs}f_{p} + TL_{g}$$

$$R_{rs} = \frac{L_{w}}{F_{0}\cos(q_{s})t_{ds}f_{s}f_{b}f_{/}}$$

molecular (Rayleigh) scattering

- elastic scattering of electromagnetic radiation by particles much smaller than the wavelength of light (atoms or molecules)
- Rayleigh scattering of sunlight in atmosphere causes diffuse sky radiation

 why the sky is blue and the Sun is yellow



• scattering phase function is symmetrical – equal forward & backward

Once more...

$$L_{\rm R} = L_{\rm r} t_{\rm gv} t_{\rm gs} f_{\rm p}$$

- Rayleigh TOA contribution (L_R) is factored into the product of a Rayleigh term (L_r) , diffuse transmittances, & a polarization correction factor
- develop & maintain one look-up table (LUT) for L_r, computed using a standard atmosphere & the non-absorbing gases N₂ and O₂ for various Sun & viewing geometries
- transmittances and f_p calculated separately
- computational efficiency: without doing this, the Rayleigh LUT would become a function of other gases affecting transmission (ozone, NO₂, & water vapor), making it too large for efficient operational use

molecular (Rayleigh) scattering

Rayleigh optical properties are calculable (to ~0.2%) – made challenging by a rough, reflective ocean (versus a flat, black ocean)

Rayleigh radiances (with polarization) are retrieved from look up tables given:

- solar & satellite viewing geometries
- wind speed (a proxy for surface roughness (influences L_{sky}))
- atmospheric pressure (\propto # gas molecules, adjusts Rayleigh optical thickness: $\tau_{\rm R}(P,\lambda) = \frac{P}{P_{\rm o}} \tau_{\rm Ro}(P_{\rm o},\lambda)$



processing cadence: $L_t / t_{gv} / t_{gs} / f_p - tL_f - L_r$











Sun glint

$$L_{t} = \left(L_{r} + [L_{a} + L_{ra}] + t_{dv}L_{f} + t_{dv}L_{w}\right)t_{gv}t_{gs}f_{p} + T_{L_{g}}$$

$$R_{rs} = \frac{L_{w}}{F_{0}\cos(q_{s})t_{ds}f_{s}f_{b}f_{l}}$$

we cannot see "ocean color" through Sun glint



Ground to Space: A Glittering Path of San Francisco Sunglint https://earthobservatory.nasa.gov/blogs/earthmatters/2016/11/09/ground-to-space-a-glittering-path-of-san-francisco-sunglint/



MODIS-Aqua (without tilt) SeaWiFS (with tilt)

Sun glint



two step iteration since we don't know τ_a :

(1) $[L_t, \tau_a', W] \rightarrow L_t^{(1)} = L_t - L_g \rightarrow \tau_a^{(1)}$ (2) $[L_t^{(1)}, \tau_a^{(1)}, W] \rightarrow L_t^{(2)} = L_t^{(1)} - L_g \rightarrow \tau_a^{(2)}$

with initial guess of $\tau_a' \sim 0.1$ (additional logic included to prevent overcorrection)

processing cadence: $L_t / t_{gv} / t_{gs} / f_p - tL_f - L_r - TL_g$









when *L_{GN}* > 0.005 sr⁻¹ mask the pixel as HIGH GLINT

when $L_{GN} \le 0.005 \text{ sr}^{-1}$ remove TL_q from L_t



aerosols: the hard part

$$L_{t} = \left(L_{r} + \begin{bmatrix}L_{a} + L_{ra}\end{bmatrix} + t_{dv}L_{f} + t_{dv}L_{w}\right)t_{gv}t_{gs}f_{p} + TL_{g}$$

$$R_{rs} = \frac{L_{w}}{F_{0}\cos(q_{s})t_{ds}f_{s}f_{b}f_{l}}$$
aerosol tables

- aerosol properties can be characterized by their particle size distribution (PSD) & their complex index of refraction (m)
- given a PSD & m (& assuming sphericity), aerosol optical properties can be computed using Mie theory:
 - \circ scattering phase function (\tilde{b})
 - \circ single scattering albedo ($\omega = b / c$)
 - \circ extinction coefficient (c = a + b)
- aerosol optical thickness relates to extinction coefficient $^{\circ} \ \tau_a = \int_0^{TOA} c(z) \ dz$
- aerosol tables are generated for various PSDs (& m's) & are
 - o defined by \tilde{D} , ω , τ_a (& other variables)
 - o navigated using solar & satellite viewing geometries

aerosol tables

- we assume each PSD to be represented by 2 lognormal distributions
 - o fine particles (continental & sometimes absorbing)
 - coarse particles (oceanic / sea salt & non-absorbing)



- each PSD modulated by varying relative humidity
 - \circ humidity changes particle size
 - \circ $\,$ requires ancillary data from NCEP $\,$
- 80 aerosol tables total, built from AERONET measurements
 - o 10 PSDs
 - o 8 relative humidities

see Ahmad et al., Applied Optics, 2010

aerosol tables

- the Angstrom exponent (α) provides an estimator of particle size
 - \circ high α = small particles
 - \circ low α = large particles

$$\circ \quad \text{defined via} \ \frac{\tau_a(\lambda)}{\tau_a(\lambda_o)} = \left(\frac{\lambda_o}{\lambda}\right)^{\alpha}$$

 aerosol models often defined by epsilon (ε)

•
$$e(748, 869) = \frac{L_a(748)}{L_a(869)}$$

do we know these values yet???



black pixel assumption



in the open ocean, we can assume (???) that L_w in the nearinfrared (NIR) is = 0 (i.e., the ocean is *black* in the NIR)

thus, in the NIR (e.g., 748 and 869 nm): $L_a(NIR) + L_{ra}(NIR) = L_t(NIR) -$ the terms we computed

aerosol selection

 $L_w(NIR) = 0$, so $L_a(NIR) + L_{ra}(NIR) = L_t(NIR) - (everything previously computed)$ how do we estimate $L_a(visible) + L_{ra}(visible)$?

- let's refer to [L_a + L_{ra}] simply as L_a & ignore single- vs. multi-scattering issues
- select the 10 aerosol tables that match the observed NCEP relative humidity
- compute epsilon values for the 10 tables [ϵ (748,869) = L_a(748) / L_a(869)]
- perform an iterative determination of the mean ϵ (748,869) value & select a final bounding 2 aerosol models
- using 2 bounding models, calculate $\varepsilon(\lambda, 869)$ from $\varepsilon(748, 869)$
- calculate $L_a(\lambda) = \varepsilon$ (λ ,869) $L_a(869)$

see Gordon & Wang, Applied Optics, 1994

final retrieval of $L_a(\lambda)$ is more accurate than that of τ_a and α ; not unlike retrievals of $a(\lambda)$ being more accurate than $a_{dg}(\lambda)$ & $a_{ph}(\lambda)$ in inversion models



is the black pixel assumption valid?

Is R_{rs}(NIR) really black?





is the black pixel assumption valid?

Is R_{rs}(NIR) really black?





is the black pixel assumption valid?

what happens when we don't account for $R_{rs}(NIR) > 0$?



retrievals using the "black pixel" assumption (e.g., SeaWiFS 1997-2000)

how to proceed with the black pixel assumption? what to do when $R_{rs}(NIR) > 0$?

many approaches exist, here are a few examples:



use a coupled ocean-atmosphere optimization

e.g., Chomko & Gordon 2001, Stamnes et al. 2003, Kuchinke et al. 2009

The Bailey et al. (2010) iterative procedure

- 1. Assume that $R_{\rm rs}(765)$ and $R_{\rm rs}(865)$ are both 0, i.e. make the black-pixel assumption for both NIR reference bands.
- 2. Complete the atmospheric correction process as described in §9.2. This gives the initial estimate of $\rho_{\rm w}(\lambda)$, or equivalently $R_{\rm rs}(\lambda)$.
- 3. Use $R_{\rm rs}(443)$ and $R_{\rm rs}(555)$ from the initial estimate of $R_{\rm rs}(\lambda)$ to get η by the empirical relationship (Lee et al. (2010, Eq. 8); Bailey et al. (2010, Eq. 3))

$$\eta = 2 \left[1 - 1.2 \exp\left(0.9 \frac{R_{\rm rs}(443)}{R_{\rm rs}(555)}\right) \right]. \tag{9.10} \quad \begin{array}{l} \text{bio-optica} \\ \text{model} \end{array}$$

- 4. Use the initial $R_{\rm rs}(\lambda)$ to get an initial estimate of the chlorophyll concentration *Chl*. The particular algorithm used to obtain *Chl* from $R_{\rm rs}(\lambda)$ depends on the sensor.
- 5. Use this Chl to obtain a(670) via the empirical relationship (Bailey et al., 2010, Eq. 4)

$$a(670) = \exp[0.9389\ln(Chl) - 3.7589] + a_w(670).$$
(9.11) bio-optical model

where the $a_w(670) = 0.439 \text{ m}^{-1}$ is the absorption by pure water.

6. Use a(670) and $R_{\rm rs}(670)$ in Eq. (9.9) to solve for $b_{\rm b}(670) = b_{\rm bw}(670) + b_{\rm bp}(670)$, where $b_{\rm bw}(670) = 4.26 \times 10^{-4} \text{ m}^{-1}$ is the backscatter coefficient for pure sea water.

 $R_{\rm rs}(\lambda) = \frac{f(\lambda)}{Q(\lambda)} \frac{b_{\rm b}(\lambda)}{a(\lambda) + b_{\rm b}(\lambda)} \quad \text{Eq. (9.9)} \quad \begin{array}{l} \text{f/Q is modeled for Case 1 water as a function of} \\ \text{Chl and is assumed known} \quad \begin{array}{l} \text{model} \end{array}$

The Bailey et al. (2010) iterative procedure

7. Use η from Eq. (9.10) and (Bailey et al., 2010, Eqs. 2b,3) to compute $b_{\rm b}(765)$:

$$b_{\rm b}(765) = b_{\rm bw}(765) + b_{\rm pb}(670) \left(\frac{670}{765}\right)^{\eta}$$
 (9.12) bio-optical model

where $b_{bw}(765) = 2.38 \times 10^{-4} \text{ m}^{-1}$. $b_{b}(865)$ is computed in the same manner using $b_{bw}(865) = 1.41 \times 10^{-4} \text{ m}^{-1}$.

- 8. Use this $b_{\rm b}(765)$ and $a(765) = a_{\rm w}(765) = 2.85 \text{ m}^{-1}$ to get $R_{rs}(765)$ from Eq. (9.9). Similarly, compute $R_{rs}(865)$ using $b_{\rm b}(865)$ and $a_{\rm w}(865) = 4.61 \text{ m}^{-1}$.
- 9. Use the new, non-zero value of $R_{\rm rs}(765)$ (i.e. $\rho_{\rm w}(765)$) to remove the non-zero $\rho_{\rm w}(765)$ contribution to $\rho_t(765)$. Do the same calculation for 865 nm.
- 10. Return to Step 2 and repeat the atmospheric correction using the black-pixel algorithm. This will give a new (hopefully better) estimate of $R_{\rm rs}(\lambda)$, thus an new estimate of the other parameters, and finally new estimates of $R_{\rm rs}(765)$ and $R_{\rm rs}(865)$ at Step 8. After using the new values of $\rho_{\rm w}(765)$ and $\rho_{\rm w}(865)$ to correct for the non-zero water contribution to $\rho_{\rm t}(765)$ and $\rho_{\rm t}(865)$, return to Step 2 for a new iteration. Continue iterating until the change in $R_{\rm rs}(765)$ from one iteration to the next is less than 2%, which typically takes 2-4 iterations, or when 10 iterations have been made.

The Bailey et al. (2010) iterative procedure

If this process doesn't converge after 10 iterations, try once more assuming that all NIR reflectance is due to water. If still no convergence, flag the pixel as "atmospheric correction warning." (The pixel might still be useable; your call.)

The above iteration is not done if Chl < 0.3 mg m⁻³, and is always done if Chl > 0.3. In between use linear combination of the two results.

The two NIR wavelength bands depend on the sensor:

Band Label	Wavelengths [nm]	Nominal Wavelength [nm]
SeaWiFS		
7	745-785	$\lambda_1 = 765$
8	845-855	$\lambda_2 = 865$
MODIS		
15	743-753	$\lambda_1 = 748$
16	862-877	$\lambda_2 = 869$
VIIRS		
M6	739-754	$\lambda_1 = 745$
M7	846-885	$\lambda_2 = 862$

Table 9.1: NIR bands used for aerosol correction.

black pixel assumption – a bio-optical model



black = land; grey = Chl < 0.3 mg m⁻³; white Chl > 0.3 mg m⁻³

not applied when Chl < 0.3 mg m⁻³ weighted application when 0.3 < Chl < 0.7 mg m⁻³ fully applied when Chl > 0.7 mg m⁻³

Bailey et al., Optics Express, 2010

black pixel assumption – a bio-optical model correction of non-negligible R_{rs}(NIR)



estimate R_{rs}(NIR) using a bio-optical model

operational SeaWiFS & MODIS processing ~ 2000-present

example alternative aerosol selection schemes

ORIGINAL RESEARCH ARTICLE

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Multiband Atmospheric Correction Algorithm for Ocean Color Retrievals

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ORIGINAL RESEARCH ARTICLE Provisionally accepted The full-text will be published soon. So Notify me

Front. Earth Sci. | doi: 10.3389/feart.2019.00100

Modeling atmosphere-ocean radiative transfer: A PACE mission perspective.

Jacek Chowdhary1*,Pengwang Zhai2,Emmanuel Boss3,Heidi M. Dierssen4,Robert J.Frouin5,Amir I. Ibrahim6,Zhongping Lee7,Lorraine A. Remer6,Michael Twardowski8,Feng Xu9,Xiaodong Zhang10,Matteo Ottaviani11,William R. Espinosa6 andDidierRamon12

- "MBAC"
- uses multiple NIR/SWIR bands instead of 2
- GW94, but no assumption of single-scattering
- Will be used for PACE with its extra wavelengths
- "POLYMER"
- spectral matching approach
- https://www.hygeos.com/polymer
- Popular in Europe and better for Sun-glint regions
- radiative transfer for coupled oceanatmosphere systems
- The ultimate goal but computationally expensive

atmospheric correction (currently) assumes no absorbing aerosols



Q: Why not use a wavelength near 350 or 400 nm, which could distinguish between absorbing and nonabsorbing aerosols?

atmospheric correction (currently) assumes no absorbing aerosols



Q: Why not use a wavelength near 350 or 400 nm, which could distinguish between absorbing and nonabsorbing aerosols? A: the water isn't black

Why not use a wavelength < 300 nm, where the ocean is again black due to high CDOM and water absorption?

atmospheric correction (currently) assumes no absorbing aerosols



Q: Why not use a wavelength near 350 or 400 nm, which could distinguish between absorbing and nonabsorbing aerosols? A: The water isn't black

Q: Why not use a wavelength < 300 nm, where the ocean is again black due to high CDOM and water absorption?

A: There ain't no sunlight



processing cadence: $L_t / t_{gv} / t_{gs} / f_p - tL_f - L_r - TL_g - L_a$



spectral bandpass correction

$$L_{t} = \left(L_{r} + [L_{a} + L_{ra}] + t_{dv}L_{f} + t_{dv}L_{w}\right)t_{gv}t_{gs}f_{p} + TL_{g}$$

$$R_{rs} = \frac{L_{w}}{F_{0}\cos(q_{s})t_{ds}f_{s}f_{b}f_{l}}$$

instrument spectral bandpasses



however, sensor band response overlaps enough to cause problems

sensor bands appear well separated on a linear axis

correction for light that comes from outside the nominal wavelength band



processing cadence: spectral bandpass correction



take care when executing satellite-to-in situ match-ups

Example for heritage multi-spectral satellite instruments:

when using multispectral in situ radiometers: enable the bandpass adjustment

when using hyperspectral in situ radiometers:

enable the adjustment when applying 10-nm filter to in situ $R_{\rm rs}$ disable the adjustment when applying full-spectral-response to in situ $R_{\rm rs}$

$$L_{t} = \left(L_{r} + [L_{a} + L_{ra}] + t_{dv}L_{f} + t_{dv}L_{w}\right)t_{gv}t_{gs}f_{p} + TL_{g}$$

$$R_{rs} = \frac{L_{w}}{F_{0}\cos(q_{s})t_{ds}f_{s}(f_{b})f_{f}}$$







we normalize R_{rs} to account for Sun's changing position in the sky:

- pathlengths through atmosphere
- transmission of light through air-sea & sea-air interfaces
- angular features of in-water volume scattering functions

$$[L_{\mathbf{w}}]_{\mathbf{N}}^{\mathbf{ex}} \equiv [L_{\mathbf{w}}(\theta_{\mathbf{v}}, \phi)]_{\mathbf{N}} \frac{\mathfrak{R}_{\mathbf{o}}(W)}{\mathfrak{R}(\theta_{\mathbf{v}}', W)} \frac{f_{\mathbf{o}}(\mathrm{ATM}, W, \mathrm{IOP})}{Q_{\mathbf{o}}(\mathrm{ATM}, W, \mathrm{IOP})} \left[\frac{f(\theta_{\mathrm{s}}, \mathrm{ATM}, W, \mathrm{IOP})}{Q(\theta_{\mathrm{s}}, \theta_{\mathbf{v}}', \phi, \mathrm{ATM}, W, \mathrm{IOP})}\right]^{-1}$$

Morel et al., Applied Optics, 2002

 $\Re, \Re_0, f, f_0, Q, Q_0$ from look-up-tables based on Chl & geometries of Sun & sensor to normalize all measurements (no subscript) to condition of overhead Sun (subscript $_0$) and no atmospheric losses (not "no atmosphere")



Morel et al., Applied Optics, 2002

 $\Re, \Re_0, f, f_0, Q, Q_0$ from look-up-tables based on Chl & geometries of Sun & sensor

to normalize all measurements (no subscript) to condition of overhead Sun (subscript ₀) and no atmospheric losses (not "no atmosphere")

processing cadence: BRDF correction





so there you have it – perfect R_{rs}

$$L_{t} = \left(L_{r} + [L_{a} + L_{ra}] + t_{dv}L_{f} + t_{dv}L_{w}\right)t_{gv}t_{gs}f_{p} + TL_{g}$$

$$R_{rs} = \frac{L_{w}}{F_{0}\cos(q_{s})t_{ds}f_{s}f_{b}f_{f}}$$

ancillary data requirements

ancillary data

ancillary source

uses

atmospheric pressure water vapor relative humidity wind speed ozone NO₂ sea surface temperature sea ice NCEP NCEP NCEP OMI/TOMS Sciamachy/OMI/GOME Reynolds NSIDC Rayleigh transmittance aerosol models white caps, Sun glint, Rayleigh transmittance transmittance bio-optical algorithms masking

look-up tables, coefficients

aerosol models Rayleigh Rayleigh optical thickness ozone absorption NO₂ absorption pure seawater absorption, scattering, index of refraction (temp/sal dependent) f/Q (bidirectional reflectance distributions) others ... All of this actually does work pretty well most of the time!!



"I know of no competent biologist who considers this an important problem."

The opinion of an "eminent biologist" who reviewed the proposed Coastal Zone Color Scanner (1978-1986) back in the 1970s.

-- related to me by Howard Gordon

Thank you! Questions?



single vs. multiple scattering



single scattering solution historically used to simplify math (CZCS – present)

in the atmospheric correction algorithm, the relationship between single (SS) and multiple scattered (MS) reflectances is defined as:

 $ln(\rho_{\rm ms}) = a_0 + a_1 ln(\rho_{\rm ss}) + a_2[ln(\rho_{\rm ss})]^2$

see Gordon & Wang, Applied Optics, 1994




single vs. multiple scattering

why bring this up?

- satellites "observe" multi-scattering
- operational atmospheric corrections schemes convert to single-scattering

- what errors are introduced? (plus, error propagation made difficult)- is this still necessary?